EXHIBIT FOUR

2014 OPTUM CONTRACT WITH CITY OF CINCINNATI

PRESCRIPTION DRUG BENEFIT ADMINISTRATION AGREEMENT

This Prescription Drug Benefit Administration Agreement, effective January 1, 2014 ("*Effective Date*"), is between City of Cincinnati ("*Client*" or "*City*") and OptumRx, Inc., a California corporation ("*Administrator*").

The parties agree as follows:

1. PRESCRIPTION DRUG BENEFIT SERVICES

- 1.1 Engagement. Client engages Administrator as its exclusive provider of the core prescription drug benefit plan services, mail order pharmacy services, specialty pharmacy services and retiree drug subsidy services set forth on Exhibit B ("Services") to support Client's Benefit Plans, and Administrator accepts this engagement, subject to the terms of this agreement.
- 1.2 **Performance Standards**. Administrator will perform the Services in a diligent and professional manner by personnel or contractors who are trained, qualified and competent to perform or deliver the Services, supervise and monitor the performance and satisfy all requirements that apply to the Services as set forth in this agreement, on condition that Client performs or complies with its obligations under this agreement. Administrator will provide the performance guarantees set forth on **Exhibit E**.
- 1.3 Additional Services. If Client asks, and Administrator agrees to perform any service in addition to the Services ("Service Change"), then the parties will amend this agreement to include the Service Change and the increase in rates, fees and reimbursements to be charged to Client. Administrator will not be obligated to perform the Service Change until the amendment for the Service Change is executed. A Service Change will not arise from an obligation required by Administrator or its subcontractor (including a Network Pharmacy) for the Services to comply with Laws and Regulations in effect as of the Effective Date. If a change in Laws and Regulations materially burdens Administrator, requires Administrator to increase payments for Covered Prescription Drugs or materially changes the Services, then the Services, rates, fees, reimbursements or Rebates will be modified appropriately so the parties are returned to their comparable economic position as of the Effective Date.
- 1.4 Compliance with Laws. Each party will comply with all Laws and Regulations applicable to its respective business and the performance of its obligations under this agreement, including maintaining any necessary licenses and permits. If a party's performance as required by this agreement is prohibited by or conflicts with any applicable Laws and Regulations, then the party whose performance is owed or required will be required to perform, but only to the extent permitted by applicable Laws and Regulations. Any provisions now or hereafter required to be included in this agreement by applicable Laws and Regulations or any Governmental Authority will be binding and be enforceable against the parties and deemed incorporated in this agreement, irrespective of whether these provisions are expressly provided for in this agreement.
 - 1.4.1 Equal Employment Opportunity Program. This agreement is subject to the Client's Equal Employment Opportunity Program contained in Chapter 325 of the Cincinnati Municipal Code. Said chapter is hereby incorporated by reference into this agreement.
 - 1.4.2 Small Business Enterprise Program. This agreement is subject to the provisions of the Small Business Enterprise Program contained in Chapter 323 of the Cincinnati Municipal Code. Section 323-99 of the Cincinnati Municipal Code is hereby incorporated into this agreement.

Details concerning this program can be obtained from the Office of Contract Compliance, Two Centennial Plaza, 805 Central Avenue, Suite 222, Cincinnati, Ohio 45202, (513) 352-3144.

Administrator shall utilize best efforts to recruit and maximize the participation of all qualified segments of the business community in subcontracting work, including the utilization of small, minority, and women business enterprises. This includes the use of practices such as assuring the inclusion of qualified small business enterprises in bid solicitation and dividing large contracts into small contracts when economically feasible.

1.5 Use of Subcontractors. Administrator will not subcontract any Service to a subcontractor without Client's prior notification, which shall not be unreasonably withheld. unless the subcontract satisfies the requirements of this section. The subcontractor will perform the subcontracted Services in accordance with the terms of this agreement and all applicable Laws and Regulations. Administrator will provide Client with a list of all subcontractors, which list will be updated annually.

2. TERM AND TERMINATION

- 2.1 Term. The initial term of this agreement begins on the Effective Date and expires on December 31, 2016. After the initial term, this agreement automatically renews for two successive 12-month renewal periods on each applicable anniversary date (the renewal periods together with the initial term, the "Term"), unless either party provides the other party with notice of non-renewal no later than 90 days before the end of the initial term or a renewal period.
- 2.2 **Termination**. This agreement may be terminated as set forth in this section or as specified elsewhere in this agreement.
 - 2.2.1 For Cause. If either party materially defaults in performing any of its material obligations under this agreement (a "Default"), the party not in Default, at its election, may terminate this agreement either in its entirety or only for the affected Services; unless the party in Default cures the Default within 30 days following notice of the Default given by the party not in Default. The notice will specify in reasonable detail the nature of the Default, the actions required to cure the Default, if the Default is curable, and whether the party not in Default is seeking to terminate either this entire agreement or only the affected Services. If the party in Default does not cure the Default to the reasonable satisfaction of the other party by the end of the 30-day cure period, then this agreement or the applicable Services, as the case may be, will terminate upon expiration of the 30-day cure period without any further notice or other action by the party not in Default. If the Default is cured before the 30-day cure period expires, then this agreement will remain in effect. Notwithstanding the foregoing, the 30-day opportunity to cure a Default will not apply to a payment default described in section 2.2.2.
 - 2.2.2 <u>Payment Default</u>. If Client fails to pay any amount due on a validly submitted invoice (for which no objection is filed in good faith in accordance with section 3.3) within five business days after the applicable Payment Due Date (as defined in section 3.2), then Administrator may immediately upon written notice to Client terminate this agreement in its entirety or only for the applicable, affected Services or withhold or suspend Services until payment is received.
 - 2.2.3 <u>Automatic Bankruptcy Termination</u>. This agreement will terminate automatically in the event of a Bankruptcy Event affecting either Administrator or Client. "*Bankruptcy Event*" means that Client or Administrator: (a) cannot pay its debts generally as they become due; (b) makes a voluntary assignment for the benefit of creditors; (c) is declared insolvent in any proceeding; (d) commences a voluntary case or other proceeding seeking liquidation, reorganization or other relief for itself, any of its property, assets or debts

under any bankruptcy, insolvency or other similar laws now or hereafter in effect or petitions or applies to any tribunal for the appointment of a receiver, liquidator, custodian or trustee for the party under any bankruptcy, reorganization, arrangement, insolvency, readjustment of debt, liquidation, or dissolution law of any jurisdiction now or hereafter in effect; (e) is named as a debtor or party in the petition, application, case or proceeding and indicates its approval thereof, consents thereto, acquiesces therein or acts in furtherance thereof, or if the petition, application, case or proceeding is not dismissed or stayed for 60 days after it begins, or is the subject of any order appointing any such receiver, liquidator, custodian-or-trustee-or-approving-the-petition-in-any-such-case-or-proceeding; or (f) the sum of the party's debts (including contingent obligations) exceeds the fair market value of the party's assets, exclusive of any property transferred, concealed, or removed with the intent to hinder, delay or defraud the party's creditors.

- 2.2.4 <u>Adverse Legal Determination</u>. Subject to section 7.3, either party may terminate this agreement immediately upon notice to the other party (a) following a Judgment (as defined in section 6.1) or change in any applicable Laws and Regulations that would make performance of this agreement, in all material respects, unlawful or illegal for the terminating party, or (b) if a Governmental Authority requires either party to terminate this agreement.
- 2.2.5 For Convenience. After the first anniversary of the Effective Date, either party may terminate this agreement without cause upon 90 days prior notice of termination to the other party. If Client terminates without cause within the first three years of the Term, (a) Administrator will have no obligation under any guarantees under this agreement for the contract year (i.e., each 12-month period following the Effective Date) in which Client terminates, if the portion of the contract year before the effective date of Client's termination is less than 12 full months and (b) Client will reimburse Administrator no later than 30 days before Client's termination for Administrator's implementation costs (maximum of \$40,000) and the implementation allowance paid or credited to Client by Administrator under Exhibit C. The amount owed by Client for Administrator's implementation costs and the implementation allowance paid or credited to Client by Administrator under Exhibit C will both be prorated monthly over the first three years of the Term.
- 2.3 Effect of Termination. Upon termination of this agreement for any reason, Client will pay Administrator all undisputed amounts for Services performed through the date this agreement terminates in its entirety. Termination of this agreement will not alter Client's obligation to pay any disputed amounts owing to Administrator under this agreement that are subsequently resolved. Termination of this agreement for any reason will not relieve either party from liability for any obligations required of the party before this agreement terminates.
- 2.4 Transition Assistance Following Termination. Upon termination of this agreement for any reason, Administrator will, as directed by Client, provide Client or its designee with the following files: (a) existing Mail Order or Specialty Pharmacy open refill transfer files for Members, as based upon Client's most current eligibility files; (b) Client's claims history file; (c) Client's prior authorization files; and (d) Client accumulator files. Each file will be sent using Administrator's standard format and delivered using a media agreed to by the parties. Administrator will send up to three transmissions (i.e., one full test file, one production file, and one lag file) of each file. After the first set of files transmitted Client will pay or reimburse Administrator \$1,500 for each file transferred.
- 2.5 Phase-Out Period for Network Pharmacies. Despite section 2.4, and only to the extent required by any applicable Laws and Regulations and if requested by Client, Administrator will continue to provide Covered Prescription Services through Network Pharmacies and Administrator's Mail Order Pharmacies and Specialty Pharmacy for up to 180 days, or longer if required by applicable Laws and Regulations, following, as applicable, (a) the date this

agreement was to have terminated in its entirety as described in sections 2.1 or 2.2 or (b) the date Administrator no longer owns or operates a Mail Order Pharmacy or Specialty Pharmacy.

3 COMPENSATION AND BILLING

- 3.1 Compensation. Client will pay Administrator the rates, fees and reimbursements set forth on **Exhibit C** for the Services. If the total number of Members on the Effective Date reduces by 10% or more during the Term, Administrator may modify the rates, fees or guarantees in Exhibit C, effective the date of the change, upon notice to Client.
- 3.2 Payment Terms. Administrator will invoice Client at semi-monthly billing cycles that run from the 1st through the 15th and from the 16th through the end of the month. Administrator will submit invoices to Client that accurately reflect the Services performed during the invoice period and include Prescription Claims information to support the involced amounts at no charge. At Client's expense, Administrator may provide electronic claims files to Client's third party service provider, subject to the third party's execution of Administrator's form confidentiality agreement. Client will pay Administrator all undisputed involced amounts, via electronic fund transfer or be and other reliable means, no later than five business days after Client receives the invoice and supporting claims detail file (Payment Due Date").
- 3.3 Timely Notice of Overpayment. Client may object to any amounts on Administrator's invoices that Client believes do not comply with the Pharmacy Plan Specifications. Client must notify Administrator of Client's objection no later than 60 days after the invoice date stating the disputed charges. If Client fails to object within this 60-day period, Client will be deemed to have acknowledged that the invoiced amounts comply with the Pharmacy Plan Specifications. This section will not preclude Client's right to audit Administrator's Services described in section 4.2.
- 3.4 Late Payments and Late Fees. Any amounts Client owes under this agreement that are not paid by the Payment Due Date will bear interest from the Payment Due Date pursuant to artis de la Chapter 319 of the Gincinnati Municipal Code.

电子多数形式 杨州城市,"我国的政治,但是这是多大,是我们的"统治"的,"这种国际的",在这种政治和政治

- 3.5 Right of Recoupment. Administrator may withhold, deduct, net or recoup from future amounts owed or reimbursable to Client under this agreement any undisputed amounts Client owes to Administrator that are outstanding beyond their applicable Payment Due Date. 지수는 경기 맛있는 경우는 그렇게 가장 그리를 하는 것이 없다.
- THE WAR THE STATE OF THE PARTY OF THE STATE 3.6 Payment from Members Except as permitted by applicable Laws and Regulations, Administrator will not bill, charge, collect a deposit from, have recourse against or otherwise seek payment from a Member for Covered Prescription Services or amounts due to Administrator from Client, other than Cost-Sharing Amounts, returned checks or collection costs, Administrator will require, under the terms of its Network Pharmacy Agreements, each Network Pharmacy to comply with the requirements of this section. ingvin Barron Polencia
- 3.7 Claims Processor Fees, Administrator may retain as part of its compensation under this agreement any claims processor fees received from Network Pharmacies in connection with the Prescription Drugs dispensed to Members under the Benefit Plans, including: (a) a per claim communications charge for on-line electronic claims processing by point-of-service communication; (b) a charge for each claim submitted to Administrator via paper, tape or a medium other than point-of-service communication; (c) surcharges for canceled or reversed claims; and (d) a charge if a Network Pharmacy requests an evidence of benefits report in a tape medium.

4. MAINTENANCE OF RECORDS; AUDITS

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4.1 Records. Administrator will maintain accurate, complete and timely books and records of all transactions occurring from Administrator performing the Services to Members, including Covered Prescription Services, health care cost, Repate data (including Rebate calculation and,

- if applicable, Rebate allocation between Client and Administrator), encounter data and other data, based upon information available to Administrator at the time of data collection or calculation. Client will keep and maintain accurate, complete and timely books and records relating to operation of the Benefit Plans. Administrator and Client each will retain these books and records during the Term and for seven years following the date of their creation or for a longer time period, if required by applicable Laws and Regulations or an on-going audit or investigation of Administrator by Client, a Governmental Authority or another person or entity, and will make these books and records available to a Governmental Authority to the extent required by applicable Laws and Regulations.
- Client Audits. Upon reasonable advance notice and at reasonable times, Client may audit once annually Administrator's performance of the Services, including concurrent eligibility, Formulary compliance and, when applicable, Rebates, for the period not to exceed 24 months immediately preceding the audit. No later than 45 days after receipt of Client's written audit request, Administrator will compile and prepare all claim detail information Client requires to perform its requested audit and furnish this information to Client in an agreed upon format. Client may audit Administrator through an audit firm of its choice, so long as: (a) the auditor does not have a conflict of interest with Administrator; (b) the audit firm executes Administrator's form Auditor Protocol and Confidentiality Agreement; (c) Client pays all costs associated with the audit, excluding Administrator's cost in compiling, copying, and making available the claim detail information necessary for the audit; and (d) Client does not compensate the audit firm, in whole or in part, on a basis that is contingent upon the results of the review of Administrator's records or the contents of the audit report. No audits may be initiated or conducted during December or January because of annual renewal period demands. Administrator will provide Client's auditor with access to all relevant data, records, contracts, files, personnel, books and other information reasonably necessary for Client's auditors to audit Administrator, subject to Administrator's third party confidentiality obligations. The audit information Administrator provides will be limited to Client-specific information necessary for Client to verify Administrator's performance under this agreement. Other documentation (e.g. policies and procedures) requested during the course of an audit, other than that needed to determine the accuracy of Client claims payments, will be provided at Administrator's reasonable discretion. Any Client requests for an auditor to audit will constitute Client's direction and authorization to Administrator to disclose this Client-specific information, including Member information and PHI, to the auditor.

5. DATA PROTECTION AND OWNERSHIP

- 5.1 Data Ownership. Client owns and will continue to own Client Data and Administrator owns and will continue to own Administrator Data, despite data use or possession by the other party or its subcontractor in accordance with an authorized subcontract. Each party will use commercially reasonable efforts to maintain the proprietary character of the other party's respective Client Data or Administrator Data.
- Data Use. Each party grants the other party a non-exclusive, nontransferable, non-sublicensable, royalty free license to use its Marks and Content in furtherance of this agreement, except that neither party is granted any license to, and will not be permitted to use, the other party's Marks or Content except as pre-approved in writing by the other party. The parties will agree on use of the other's Marks, Content or words or phrases identifying the other party in any promotional or other materials, any advertisements identifying the other party, and in connection with Client identifying the Benefit Plans, or in any public announcement or press release, including agreeing on the timing and content of any public release. Despite any contrary provisions in this agreement, during the Term and for a reasonable period after termination of this agreement, Client grants Administrator the right to use and disclose to third parties Member drug and related medical data to perform Administrator's responsibilities under this agreement and to use in Administrator's research, cost analyses, and cost comparison studies. All research, cost analyses, cost comparisons and other similar studies or reports Administrator conducts or prepares will be Administrator's sole and exclusive property.

Administrator may aggregate this information with that of other clients and de-identify it to protect Client and Member confidentiality.

5.3 Confidentiality

- 5.3.1 <u>Confidentiality Obligations</u>. Each party ("*Recipient*") will, and will use commercially reasonable efforts to cause each of its Representatives to, keep confidential the Confidential Information of the other party ("*Discloser*") and not disclose any Confidential Information without Discloser's prior written consent or as permitted by this Agreement. Confidential Information may be disclosed to either party's employees, contractors or another third party ("*Representative*") as reasonably necessary to carry out the purposes of this agreement, on condition that the Representative agreed to keep confidential the Confidential information with obligations at least as comprehensive as the obligations in this agreement. Recipient will be responsible for any breach of this agreement by any Representative to which it discloses Confidential Information.
- 5.3.2 <u>Definition of Confidential Information</u>. "Confidential Information" means: (a) the terms of this agreement; (b) all Discloser material, non-public information, materials or data, in any form, that Recipient knows or has reason to know is confidential or proprietary to Discloser, including Client Data or Administrator Data; (c) any other information that Discloser marks or designates clearly as confidential or proprietary; and (d) Discloser trade secrets, know how, inventions, current and future business plans, marketing plans and strategies, financial and operational plans, business methods and practices, customer or prospect data, records, information and profiles, supplier or vendor information and data, historical or prospective financial information, budgets, cost and expense data, employment records and contracts and personnel information as well as software, technology, inventions (whether or not patentable) that Discloser owns, licenses or uses. Confidential Information will not include information that: (a) is generally available to the public; (b) becomes available to Recipient on a non-confidential basis from a source, other than Discloser or its Affiliates or agents, not bound by a confidentiality agreement; or (c) that is required to be disclosed by law or pursuant to court order.
- 5.3.3 Exceptions to Confidentiality Obligations. The obligations in this section 5.3 will not restrict or limit disclosures by Recipient: (a) to offer or administer the Benefit Plans; (b) to comply with Rebate reporting or other data collection, maintenance, security or submission requirements; (c) to perform functions or responsibilities required by applicable Laws and Regulations; (d) as required or compelled by applicable Laws and Regulations or a Governmental Authority with competent jurisdiction over Recipient, on condition that Recipient will: (i) give prompt notice to Discloser after learning of the need to disclose (if allowed by applicable Laws and Regulations); (ii) disclose only that portion of Discloser's Confidential Information that Recipient's outside legal counsel advises is legally necessary to comply with the Laws and Regulations or Governmental Authority order; and (iii) assist Discloser if it objects to the disclosure; or (e) as provided in the next paragraph.

The parties acknowledge that Client is governed by the Ohio Public Records Laws and agree to comply with all applicable provisions of the Ohio Public Records Laws, despite contrary provisions in this agreement. Records (as defined by Ohio Revised Code §§ 149.011 and 149.43) related to this agreement may be subject to disclosure under the Ohio Public Records Laws. Client shall have no duty to defend the rights of Administrator or any of its agents or affiliates in any records requested to be disclosed. Upon receipt of a public records request, Client will notify Administrator in accordance with Section 7.1 of this agreement of its intent to release records to the requestor. Beginning with the date it receives notification, Administrator shall have the lesser of: (a) ten (10) business days or (b) a "reasonable amount of time" as that term has been interpreted by Supreme Court of Ohio and as notified by Client to Administrator in order to respond to Client by either accommodating the requestor or pursuing legal remedies to stop the Client's release of

requested information. Administrator and its agents and affiliates shall have the right to pursue legal and/or equitable remedies to stop or limit disclosure at their sole expense. Administrator will defend and hold harmless Client, including any reasonable and actual attorneys' fees awarded against Client under O.R.C. section 149.43 to the extent the claim is related to Administrator's designation of certain records as exempt from public disclosure as a trade secret ("Claim"). Client will promptly notify Administrator of the Claim, and will allow Administrator to control the defense and settlement thereof.

- 5.4 Return of Confidential Information. Upon Discloser's request, for any reason whatsoever and at any time, Recipient will return immediately all Discloser's Confidential Information within Recipient's possession or control. If any Confidential Information is contained in analyses, compilations, studies or other documents prepared by Recipient or its contractors, Recipient will promptly destroy, and instruct its Authorized Representatives to destroy, these items and certify to Discloser that this destruction has occurred. Recipient and its Authorized Representatives may retain one copy of Discloser's Confidential Information for archival purposes or as otherwise required by applicable Laws and Regulations.
- 5.5 **Protected Health Information**. The parties will execute and abide by the Business Associate Agreement in the form of **Exhibit D**, which outlines the parties' obligations for use and disclosure of PHI.
- 5.6 **Equitable Relief.** The parties acknowledge that it would be difficult to measure damages resulting from any breach of their respective obligations in this section 5, injury from this breach would be impossible to calculate and money damages would be an inadequate remedy. Consequently, in addition to any other rights or remedies available under this agreement, the parties may seek injunctive and other equitable relief, without bond or other security, for a party's actual or threatened breach of this section 5. The obligations, rights and remedies of the parties under this section 5 are cumulative and in addition to, and not in lieu of, all obligations, rights or remedies created by applicable patent, copyright or other laws, including statutory and common laws governing unfair competition and misappropriation or theft of trade secrets, proprietary rights or confidential information.

6. INDEMNIFICATION; INSURANCE; LIMITATION OF LIABILITY

Indemnification. Each party ("Indemnitor") will be solely financially responsible for, and will defend and indemnify the other party, its Affiliates and their respective directors, officers, employees, representatives, agents, successors, successors-in-interest and assigns ("Indemnitee") from and against all claims, legal or equitable causes of action, suits, litigation, proceedings (including regulatory or administrative proceedings), grievances, complaints, demands, charges, investigations, audits, arbitrations, mediation or other process for settling disputes or disagreements, including any of the foregoing processes or procedures in which injunctive or equitable relief is sought ("Claims") made by a third party against Indemnitee arising or resulting from, or to the extent attributable to, Indemnitor's material breach of this agreement or its negligence or intentional misconduct (including fraud), except to the extent the liability results from Indemnitee's negligence, willful misconduct or breach of this agreement. Indemnitor will pay promptly and satisfy fully in connection with an indemnified Claim all (a) losses, damages of any kind or nature, assessments, fines, penalties, deficiencies, interest, payments, expenses, costs, debts, obligations, liabilities, liens or Judgments that are sustained, incurred or accrued; (b) judgments, writs, orders, injunctions or other orders for equitable relief, awards or decrees of or by any Governmental Authority ("Judgments"); and (c) costs, expenses and fees, including settlement costs, attorneys' fees, accounting fees and expert costs and fees incurred in connection with Claims. Each party will provide prompt notice to the other party upon learning of any occurrence or event that may result in an obligation of the other party under this section. A party's failure to provide prompt notice of a Claim will not relieve the other party of its obligations under this section, except to the extent that the omission results in a failure of actual notice to the other party and the other party suffers damages because of the failure to notify.

- 6.1.1 Notwithstanding any statement in section 6.1 (Indemnification) above, the City's obligation to indemnify Administrator shall be limited to the remaining amount of funds, if any, which have been previously certified to this contract by the City's Director of Finance pursuant to O.R.C § 5705.41 and C.M.C. § 301-1. The City is under no obligation to certify additional funds to this contract to meet any request for indemnification.
- 6.2 Insurance Requirements. The parties acknowledge that Client does not maintain commercial insurance and is self-insured for workers compensation insurance purposes. Administrator will maintain: (a) during and for a reasonable period of time after the Term, reasonable and customary insurance (whether through third party carriers or self-insured arrangements or retentions), as to type, policy limits and other coverage terms, to cover the risks of loss faced by companies similar to the party in size, industry and business operations; and (b) all insurance coverage, bonds, security and financial assurances as applicable Laws and Regulations may require from time-to-time. All authorized Administrator subcontracts will require the subcontractors to maintain adequate and customary insurance. Administrator, at its sole cost and expense, will procure and maintain (a) workers compensation insurance coverage, (b) comprehensive general liability insurance, including personal injury, and (c) automobile liability, including non-owned and hired auto coverage, of not less than \$1,000,000 per occurrence. Administrator shall have Client named as an additional insured on Administrator's general liability and automobile liability insurance policies.
- 6.3 **Limitation of Liability**. The parties' liability to each other under this agreement will not exceed the actual damages caused by breach of this agreement. The parties will have no liability under this agreement for any loss of profit or revenue or for any consequential, indirect, incidental, special or punitive damages, even if they are aware of the possibility of the loss or damages.

7. GENERAL TERMS

7.1 Notices. All notices, requests, consents, demands or other communications under this agreement will be in writing and deemed to have been duly given either (a) when delivered, if delivered by hand, sent by United States registered or certified mail (return receipt requested), delivered personally by commercial courier or (b) on the second following business day, if sent by United States Express Mail or a nationally recognized commercial overnight courier; and in each case to the parties at the following addresses (or at other addresses as specified by a notice) with applicable postage or delivery charges prepaid.

If to Administrator:

OptumRx, Inc. 2300 Main Street Irvine, California 92614 Attn: Vice President, Client Management

Copy to:

OptumRx, Inc. 2300 Main Street Irvine, California 92614 Attn: General Counsel

If to Client:

City of Cincinnati Risk Management 805 Central Avenue Cincinnati, OH 45202 Attn: Risk Manager

- 7.2 Amendment. Except as otherwise provided in this section or elsewhere in this agreement, this agreement may be modified, changed or amended only by a dated written instrument executed by the parties. If, despite section 1.4, any Governmental Authority or Laws and Regulations require that this agreement be amended, including to incorporate specific required terms, Administrator may amend this agreement to comply with this requirement by providing 30-days prior notice to Client. This amendment will become effective at the end of the 30-day notice period or a shorter period if necessary to comply with the requirement, unless Client can demonstrate conclusively in writing that the amendment is not necessary to comply with the Governmental Authority or Laws and Regulations.
- 7.3 Waiver; Severability. The failure of any party to insist in any one or more instances upon performance of any term of this agreement will not be construed as a waiver of future performance of the term, and the party's obligations for the term will continue in full force and effect. The provisions of this agreement are severable. The invalidity or unenforceability of any term or provision in any jurisdiction will be construed and enforced as if it has been narrowly drawn so as not to be invalid, illegal or unenforceable to the extent possible and will in no way affect the validity or enforceability of any other terms or provisions in that jurisdiction or of this entire agreement in that jurisdiction.
- 7.4 Assignment. A party may not assign, delegate or transfer this agreement without the prior written consent of the other party, except that Administrator may assign this agreement to any Affiliate upon 30-day notice to Client, so long as Administrator remains obligated under this agreement. This agreement will bind the parties and their respective successors and assigns and will inure to the benefit of the parties and their respective permitted successors and assigns.
- 7.5 Governing Law. This agreement and each party's rights and obligations under it will be governed by and construed in accordance with the laws of Ohio, without giving effect to conflicts of law principles.
- 7.6 **Certification as to Non-Debarment.** Administrator certifies that neither it nor its principals is presently debarred, suspended, proposed for debarment, declared ineligible, or voluntarily excluded from participation in the transaction covered by this Agreement. Administrator acknowledges and agrees that if it or its principals is/are presently debarred then it will promptly return to Client any funds received pursuant to this Agreement. In such event, any materials received by Client pursuant to this agreement will be retained as liquidated damages.
- 7.7 Force Majeure. If any party is prevented from performing or cannot perform any of its obligations under this agreement because of any cause beyond the reasonable control of and not the fault of the party invoking this section, including any act of God, fire, casualty, flood, earthquake, war, strike, lockout, epidemic, destruction of production facilities, riot, insurrection or material unavailability, and if the non-performing party has been unable to avoid or overcome its effects through the exercise of commercially reasonable efforts, this party will give prompt notice to the other party, its performance will be excused, and the time for its performance will be extended for the period of delay or inability to perform due to such occurrences, except that if performance is extended under this section for more than 60 days, then at any time before reinstatement of the performance, the other party may terminate this agreement upon notice to the non-performing party. Administrator will maintain commercially reasonable business continuity and disaster recovery plans.
- 7.8 Relationship of the Parties; Third Party Beneficiaries. The sole relationship between the parties is that of independent contractors. This agreement will not create a joint venture,

- partnership, agency, employment or other relationship between the parties. Nothing in this agreement will be construed to create any rights or obligations except among the parties; no person or entity will be regarded as a third party beneficiary of this agreement.
- 7.9 **Taxes.** Member, and not Administrator, will be responsible for any transfer, consumption, sales, use or other tax levied on the transfer of items dispensed or on any of the Services if assessed or required by local, city, state or other government authority.
- 7.10 **Survival**. Any term of this agreement that contemplates performance after termination of this agreement will survive expiration or termination and continue until fully satisfied, including section 5, which will survive so long as the information is Confidential Information or the data is proprietary to either party or its successors, successors-in-interest or assigns, and section 6, which will survive indefinitely.
- 7.11 Dispute Resolution. If a dispute occurs between the parties, the complaining party may request a meeting by executive officers of each party who will attempt to resolve the dispute in good faith before beginning a legal action, except for matters subject to injunctive relief. If the parties' executive officers do not resolve the dispute within 45 days after the notice, each party will retain all rights to bring an action regarding such matter in accordance with law.
- 7.12 Integrated Agreement; Interpretation; Execution. This agreement constitutes the final entire agreement between the parties regarding its subject matter and supersedes all prior or contemporaneous written or oral agreements, representations, negotiations or understandings between the parties regarding its subject matter. The language in this agreement will be construed in accordance with its fair meaning, as if prepared by all parties and not strictly for or against any party. The legal doctrine of construction of ambiguities against the drafting party will not be employed in any interpretation of this agreement. Whenever approval of any party is required under this agreement, the approval will not be unreasonably withheld or delayed. For all terms in this agreement, unless otherwise specified: (a) a term has the meaning assigned to it in the Schedule of Definitions attached as Exhibit A or elsewhere in this agreement; (b) "or" is not exclusive; (c) "including" means including without limitation; (d) "party" and "parties" refer only to a named party to this agreement; and (e) any reference to an agreement, instrument or statute means that agreement, instrument or statute as from time-to-time amended, modified or supplemented and any applicable corresponding provisions of successor statutes or regulations. The headings in this agreement are provided for convenience only and do not affect its meaning. An electronic signature of this agreement, or a signature on a copy of this agreement that a party receives by facsimile, email or other means, is binding as an original, and the parties will treat an electronic or photo copy of this signed agreement as an original. The parties may sign this agreement in two or more counterparts, and as so signed this agreement will constitute one and the same agreement binding on the parties.

[signature page follows]

The parties' duly authorized representatives are signing this Prescription Drug Benefit Administration Agreement as of the Effective Date.

CITY OF CINCINNATI

OPTUMRX, INC.

By:

By:

Name:

Name:

Name:

Title:

APPROVED FOR COMPLIANCE

Karen Ader, Risk Manager

Administration Agreement as of the Effective Date.

OPTUMRX, INC.

By:

APPROVED FOR COMPLIANCE

Approved For Compliance Officer

Contract Compliance Officer

APPROVED AS TO FORM

Assistant City Solicitor

CERTIFICATION OF FUNDS

Date: DEC 3 1 2013

Funding: 7/1×135×3521×

Amount: \$2,064,000.

Reginale Zeno Finance Director

Exhibits:

Exhibit A, Schedule of Definitions

Exhibit B, Services

Exhibit C, Compensation

Exhibit D, Business Associate Addendum

Exhibit E, Performance Guarantees

EXHIBIT A

SCHEDULE OF DEFINITIONS

Capitalized terms used in this agreement are defined below or elsewhere in this agreement.

"Administrator Data" means: (a) all data and information Administrator submits or transmits to Client regarding Administrator or its formulary advisory committee, Administrator's formularies, Network Pharmacies or Pharmacy Network, (b) all data, records and information generated in Administrator's business or operations; (c) all information pertaining to any programs, services or products Administrator or any of its clients market or offer; (d) all Administrator Content, Marks and Intellectual Property, together with all derivative works of the Administrator Content, Marks and Intellectual Property; (e) Administrator's software and any tangible or readable embodiments of such software, (f) Member specific information received or generated by Administrator's Mail Order or Specialty Pharmacies in connection with dispensing Prescription Drugs; and (g) for any matters referenced in the foregoing clauses (a) through (f), data, records or information occurring in any form, including written, graphic, electronic, visual or fixed in any tangible medium of expression and whether developed, generated, stored, possessed or used by Administrator, Client or a third party. Administrator Data does not include any claims data or data or information that relates exclusively to Client or its business, operations or activities or to another Client customer or contractor or the customer's or contractor's business, operations or activities.

"Affiliate" means for any person or entity, any other person or entity that directly or indirectly controls, is controlled by or is under common control with this person or entity.

"AWP" means the average wholesale price of a Prescription Drug or other pharmaceutical products or supplies based on the Pricing Source. For Prescription Drugs or other pharmaceutical products or supplies not dispensed by Administrator's Mail Order Pharmacy or Specialty Pharmacy, AWP is based on the NDC of the drug dispensed. For Prescription Drugs or other pharmaceutical products or supplies dispensed by Administrator's Mail Order Pharmacy or Specialty Pharmacy, AWP is based on a package size of 100 units for pills, capsules and tablets and 16 ounce quantities for liquids (or the next closest package size if these quantities or sizes are not available) or the manufacturer's individual prepackaged item (e.g., tube, drop dispenser, etc.). Administrator will rely on the Pricing Source as updated by Administrator no less frequently than every seven days to determine AWP for purposes of establishing the pricing provided to Client under this agreement. Administrator will not establish AWP, and Administrator will have no liability to Client arising from use of the Pricing Source. If Administrator decides to use a pricing benchmark other than AWP or is required to do so because the Pricing Source discontinues publication of AWP and the change would materially affect Client's economic benefit under this agreement, then Administrator will provide Client with proposed modified pricing terms at least 60 days before the effective date of the change. If the parties fail to agree upon the modified pricing terms before the effective date of the modified pricing terms, then Administrator's proposed modified pricing terms will apply until the parties otherwise agree. If the parties are unable to agree to modified pricing terms, then either party may terminate this agreement upon 60 days prior notice to the other party.

"Benefit Plan" means the certificate of coverage, summary plan description, or other document or agreement, whether delivered in paper, electronic, or other format, under which Client is obligated to provide Covered Prescription Services. Benefit Plan coverage includes any deductible or co-insurance provided for under the coverage.

"Brand Drug" means a Prescription Drug, pharmaceutical product or supply designated as "M," "N" or "O" in Medi-Span's Generic Product Indicator.

"Client Data" means (a) all data and information Client submits or transmits to Administrator, including information about Benefit Plans, Pharmacy Plan Specifications, Members, and Client's other programs, services, products and plans; (b) any data and information submitted or transmitted to Administrator by a Governmental Authority or a third party about Client or Benefit Plans; (c) data, records

and information Administrator generates that relates directly to Administrator performing Services for Client under this agreement, exclusive of information or documentation Administrator generates for use in Administrator's business generally or for use with multiple clients; (d) all Client Content, Marks and Intellectual Property, together with all derivative works of the Client Content, Marks and Intellectual Property; (e) data, records and information Administrator generates about Client's business or operations; and (f) for any matters referenced in the foregoing clauses (a) through (f), data, records or information occurring in any form, including written, graphic, electronic, visual or fixed in any tangible medium of expression and whether developed, generated, stored, possessed or used by Client, Administrator or a third party. Client Data will not include data or information that is generated in or relates exclusively to: (a) Administrator or its business, operations or activities; (b) another Administrator client or contractor or the client's or contractor's business, operations or activities; (c) Administrator's or its personnel or contractor's use other than in performing this agreement; or (d) data or information disclosed, sold, assigned, leased or otherwise provided to third parties in a form that the Client Data has been aggregated with other client's data and cannot be distinguished as Client Data.

"Compound Prescription Drug" means a Prescription Drug that is prepared by a pharmacist who mixes or adjusts one or more Prescription Drugs to customize a medication to meet a Member's individual medical needs. Client's payment to Administrator for providing a Compound Prescription Drug to a Member will include the Network Pharmacy contracted rate for each Prescription Drug included in the medication and one contracted dispensing fee minus any Cost-Sharing amount.

"Content" means any text, graphics, photographs, video, audio or other data or information, including any advertisements used by the applicable party, or in the case of Client by it or its vendors, in its business, operations or in connection with the offering of its products, services, programs or plans.

"Cost-Sharing Amount" means the coinsurance, copay or other cost sharing amount that a pharmacy may collect from a Member for Covered Prescription Services in accordance with the Member's Benefit Plan.

"Covered Prescription Services" means Prescription Drugs or other pharmaceutical products, services or supplies dispensed by a pharmacy to a Member for which coverage is provided in accordance with the Member's Benefit Plan.

"Drug Manufacturer" means a person or entity that manufactures, sells, markets or distributes Prescription Drugs.

"FDA" means the United States Food and Drug Administration or any successor Governmental Authority.

"Formulary" means the list of Prescription Drugs covered by the applicable Benefit Plan as developed by Administrator and approved and adopted by Client for use with the Benefit Plans. The Formulary will be made available to physicians, pharmacies and other healthcare persons or entities to guide the prescribing, dispensing, sale and coverage of Covered Prescription Services.

"Generic Drug" means a Prescription Drug, pharmaceutical product or supply designated as "Y" in Medi-Span's Generic Product Indicator.

"Governmental Authority" means the Federal government, any state, county, municipal or local government or any governmental department, political subdivision, agency, bureau, commission, authority, body or instrumentality or court that regulates the party's activities or operations.

"Intellectual Property" means any patent, invention, discovery, know-how, technology, software, copyright, authorship, trade secret, trademark, trade dress, service mark, confidentiality, proprietary, privacy, intellectual property or similar rights (including rights in applications, registrations, filings and renewals) that are now or hereafter protected or legally enforceable under state or Federal common laws or statutory laws or under laws of foreign jurisdictions.

- "Laws and Regulations" means all common law and any and all state, Federal or local statutes, ordinances, codes, rules, regulations, restrictions, orders, procedures, standards, directives, guidelines, instructions, bulletins, policies or requirements enacted, adopted, promulgated, applied, followed or imposed by any Governmental Authority, including the Financial Modernization Act of 1999, also known as the Gramm-Leach-Billey Act, and the Health Insurance Portability and Accountability Act of 1996 and its implementing regulations, as well as any of the preceding Laws and Regulations that from time-to-time may be amended, modified, revised or replaced, interpreted or enforced by any Governmental Authority.
- **"MAC"** means the maximum allowable cost of a Generic Drug as specified on a list established by Administrator. Administrator may have multiple MAC lists, each of which is subject to Administrator's periodic review and modification. The MAC used at Mail Order Pharmacy will perform in the aggregate better than or equal to the MAC used at retail pharmacy.
- "Mail Order Pharmacy" means a facility that is duly licensed to operate as a pharmacy at its location and to dispense Prescription Drugs via postal or commercial courier delivery to individuals, including Members. Mail Order Pharmacy includes pharmacies that Administrator owns or operates.
- "Marks" means the names, logos and other proprietary symbols and phrases belonging to a person or entity.
 - "Member" means an eligible individual legitimately enrolled in a Benefit Plan.
 - "NCPDP" means that National Counsel for Prescription Drug Programs.
- "NDC" means the National Drug Code that is the identifying Prescription Drug number maintained by the FDA.
- "Network Pharmacy" means a retail pharmacy, Mail Order Pharmacy, Specialty Pharmacy or other facility that is duly licensed to operate as a pharmacy at its location and to dispense Prescription Drugs to individuals, including Members, and has entered into a Network Pharmacy Agreement. Administrator in its capacity as a Mail Order Pharmacy or Specialty Pharmacy is a Network Pharmacy of Client.
- "Network Pharmacy Agreement" means a Prescription Drug Services Agreement between a Network Pharmacy and Administrator or Client to provide Covered Prescription Services.
- "Paid Claim" means a Prescription Drug claim that is approved for payment during Administrator's semi-monthly billing cycle or is a reversal during this semi-monthly billing cycle of a Prescription Drug Claim that was approved for payment during a prior semi-monthly billing cycle. A rejected or denied claim or a claim approved for payment and reversed during the same semi-monthly billing cycle is not a Paid Claim.
- "Pharmacy Plan Specifications" means the written Benefit Plan descriptions, Member information and instructions Administrator needs to carry out its obligations under this agreement, including Member eligibility and identification requirements, benefit definitions, Formulary, Pharmacy Network, utilization management programs, applicable Cost-Sharing Amounts, number of days' supply for acute and maintenance medications, dispensing and other limitations, manuals and other Benefit Plan or Member information. All Pharmacy Plan Specifications will be either provided by Client or prepared by Administrator and approved by Client.
 - "PHI" means Protected Health Information, as defined in 45 C.F.R. § 160.103.
- "Prescription Claim" means a single request for payment for, or a bill or invoice relating to, a Covered Prescription Service that a Network Pharmacy, other health care provider or Member submits, whether the request, bill or invoice is paid or denied.

- "Prescription Drug" means a Generic Drug or Brand Drug that is approved by the FDA and required under applicable Laws and Regulations to be dispensed only as authorized by a written or oral order to dispense a Prescription Drug by an appropriately licensed and qualified health care professional in accordance with applicable Laws and Regulations.
- "Prescription Drug Compensation" means the applicable reimbursement, remuneration, compensation or other payment paid by Client to Administrator for the provision of Covered Prescription Services to a Member as described in **Exhibit C**.
- "Pricing Source" means the Medi-Span Prescription Pricing Guide (with supplements) or another nationally recognized pricing source determined by Administrator.
- "Rebate" means any discount, price concession or other direct or indirect remuneration Administrator receives from a Drug Manufacturer under a Rebate Agreement that is contingent upon and related directly to Member use of a Prescription Drug during the Term. Rebate does not include any discount, price concession or other direct or indirect remuneration Administrator receives from a Drug Manufacturer for direct purchase of a Prescription Drug or any amounts a Drug Manufacturer pays Administrator for providing any products or services, including fees for managing and administering Administrator's rebate program.
- "Rebate Agreement" means an agreement, other than a purchase agreement, by Administrator and a Drug Manufacturer for price concessions, direct or indirect remunerations or reimbursements based on projected or actual use of the Drug Manufacturer's Prescription Drugs by members of prescription drug benefit plans that are Administrator clients, including Client. Drug Manufacturer purchase agreements that include volume discounts or rebates resulting from Prescription Drug purchases, rather than member use, will not be Rebate Agreements.
- "Specialty Drugs" means the Prescription Drugs available at Administrator's Specialty Pharmacy, including: (a) biotechnology drugs; (b) orphan drugs used to treat rare diseases; (c) typically high-cost drugs; (d) drugs administered by oral or injectable routes, including infusions in any outpatient setting; (e) drugs requiring on-going frequent patient management or monitoring; and (f) drugs that require specialized coordination, handling and distribution services for appropriate medication administration.
- "Specialty Pharmacy" means a facility that is duly licensed to operate as a pharmacy at its location and to dispense Specialty Drugs to individuals, including Members. Specialty Pharmacy includes pharmacies that Administrator owns or operates.
- "Usual and Customary Charge" means the price, including all applicable customer discounts, such as special customer, senior citizen and frequent shopper discounts, that a cash paying customer pays a pharmacy for Prescription Drugs.
- "WAC" means the wholesale acquisition cost of medication drugs or ancillary supplies, as applicable, as dispensed and set forth in the Pricing Source.
- "Zero Balance Due Claims" means claims for which Client will not be billed an amount by Administrator. Member is responsible for the full cost of these claims as required in the applicable pharmacy benefit coverage criteria.

EXHIBIT B

SERVICES

1. CORE PRESCRIPTION DRUG BENEFIT SERVICES

1.1 Administrative Support

- 1.1.1 General. Administrator will provide administrative, management, consultative, claims processing and other general pharmacy benefit management support services to Client in conjunction with administration and operation of the Benefit Plans as set forth in this exhibit. Administrator will administer and support the Benefit Plans in accordance with the most current Pharmacy Plan Specifications that Client has provided to Administrator as required by this agreement.
- 1.1.2 <u>Reporting</u>. Administrator will provide Client with Administrator's standard reporting package and reports.
- 1.1.3 Benefit Plan Responsibility. Although Administrator will perform Services under this agreement to support the Benefit Plans, Client retains complete and exclusive discretionary authority over the Benefit Plans and is responsible ultimately for administering, managing and operating the Benefit Plans, including establishing and amending the Formulary, controlling or directing appeals conducted by an independent outside party or independent review organization ("IRO") and determining, interpreting and amending all Benefit Plan structures and terms. Except as the parties specifically agree in writing and despite any contrary provisions in this agreement, (a) neither Administrator nor its Affiliates is acting on behalf of any employee welfare benefit plan (as defined in 29 U.S.C. § 1002(1)) or participants or beneficiaries in any such plan, or on behalf of a fiduciary (as defined in 29 U.S.C. § 1002(21)(A)) of any such plan; (b) Client will not name or deem Administrator as a fiduciary for any purpose; (c) Administrator's role in all respects will be limited to that of a provider of "ministerial functions" (as described in 29 C.F.R. § 2509.75-8, D-2) and will be performed within the framework of policies and interpretations established by Client, such that the Services Administrator performs under this agreement will not include the power to exercise discretionary authority over any Benefit Plan's management or operations or plan assets (if any); (d) Client has selected and is solely responsible for each Benefit Plan's benefits and design; and (e) Client retains all discretionary authority for each Benefit Plan, Benefit Plan assets (If any) and administration of each Benefit Plan.
- 1.1.4 Benefit Plan Eligibility Data. Client will provide Administrator with electronic eligibility data in NCPDP format, or another format agreed to by the parties, for all Members who are entitled to Covered Prescription Services under the Benefit Plans. Administrator will load correctly formatted Member eligibility data no later than three business days after receipt from Client. Administrator will be entitled to rely on the accuracy and completeness of the Member eligibility data. Client will be solely responsible for any errors in Member eligibility data that Client furnishes to Administrator.
- 1.1.5 <u>Member Notification</u>. Client will apprise Members of the type, scope, restrictions, limitations and duration of Covered Prescription Services to which Members are entitled under an applicable Benefit Plan.
- 1.1.6 <u>Pharmacy Plan Specifications</u>. Client will provide Administrator with the technical assistance and information Administrator reasonably needs to perform the Services, including information regarding Members, Benefit Plans and Pharmacy Plan Specifications. Client will provide Administrator with the Pharmacy Plan Specifications no later than 45 days before the Effective Date. Client may amend or terminate the Pharmacy

Plan Specifications upon 45 days prior notice to Administrator, unless a Governmental Authority requires that the amendment or termination occur in a shorter time period. Any Client-initiated change to the Pharmacy Plan Specifications, including the Benefit Plan, Formulary, Pharmacy Network or a utilization management program, that impacts Administrator's compensation, cost to provide services or ability to satisfy a guarantee under this Agreement will be a Service Change and Administrator may adjust the rates, fees or guarantees in **Exhibit C**, effective the date of the change, in accordance with section 1.3 of the agreement. Client's failure to provide Pharmacy Plan Specifications within the time periods stated in this section may delay Administrator's implementation of the Services and any applicable performance guarantees. Client is responsible for the accuracy, completeness and timeliness of all Pharmacy Plan Specifications provided to Administrator and acknowledges Administrator's reasonable reliance on the Pharmacy Plan Specifications.

1.2 Pharmacy Network Administration

- 1.2.1 Pharmacy Network. Administrator will establish and maintain a network of pharmacies to provide the Services to Client ("Pharmacy Network"). Upon request, Administrator will make available to Client a current list of Network Pharmacies in the Pharmacy Network. Administrator may add or remove Network Pharmacies from the Pharmacy Network. Administrator shall make every effort to maintain its Pharmacy Network so that it is substantially the same in terms of number and locations of Network Pharmacies as were in the Pharmacy Network as of the effective date of this Agreement, excluding changes based on pharmacy closures, pharmacy self-terminations, and pharmacies removed for contract violations,. Administrator will notify Client if any pharmacy chain with 4,000 or more stores leaves or is removed from the Pharmacy Network. Client shall have the right to terminate this agreement if, in Client's determination, Administrator's Pharmacy Network is significantly altered during the term of this agreement, by following the provision in Section 2.2.1 of this Agreement.
- 1.2.2 Network Pharmacy Credentialing. Administrator will establish and maintain a reasonable process for credentialing Network Pharmacles that includes verifying the good standing of the license of the pharmacy, adopting policies and procedures required by applicable Laws and Regulations. Administrator will use best efforts to contractually require each Network Pharmacy and their pharmacists dispensing Covered Prescription Services to Members to be duly licensed in accordance with all applicable Laws and Regulations in the state or other jurisdictions in which the Network Pharmacy furnishes Covered Prescription Services.
- 1.2.3 Network Pharmacy Agreements. Administrator has entered into Network Pharmacy Agreements to secure Network Pharmacies for the Pharmacy Network. Administrator will use best efforts to contractually require each Network Pharmacy (a) to comply with all applicable Laws and Regulations and (b) to collect from Members for the provision of Covered Prescription Services a charge that is the lesser of the Cost-Sharing Amount, Usual and Customary Charge or Prescription Drug Compensation, as applicable. The amount Administrator pays to Network Pharmacies for providing Covered Prescription Services to Members shall be the same amount Client pays Administrator for these services under this agreement
- 1.2.4 <u>Customer Service</u>. Administrator will maintain one or more call centers to provide customer service assistance for Members in connection with Administrator's Mail Order Pharmacy and Network Pharmacies.
- 1.2.5 <u>Desk and On-Site Audits.</u> Administrator will, as required by applicable Laws and Regulations and at its own expense, conduct real-time and retrospective desk audits and selected on-site audits of the Network Pharmacies to determine whether the Network

Pharmacies are submitting appropriate billings for payment by Client or Members. Administrator will report the results of the audits to Client. Administrator will pay Client, or apply as a credit to invoices payable by Client to Administrator, the amounts Administrator recovers from these audits. Client will be financially responsible for all expenses incurred in connection with audits of Network Pharmacies requested by Client that are not required by applicable Laws and Regulations.

1.3 Claims Process

- 1.3.1 Claims Adjudication. Administrator, directly or through a third party claims processor with which Administrator may contract, will adjudicate, process or pay Prescription Claims for Covered Prescription Services by application of the Benefit Plan rules determining eligibility for Covered Prescription Services. Administrator will pay, on Client's behalf, only Clean Claims (a) submitted by the Network Pharmacies in a timely manner through Administrator's point-of-service system in accordance with NCPDP guidelines and the Pharmacy Plan Specifications and (b) properly submitted by Members as requests for reimbursement for Covered Prescription Services. For each Clean Claim submitted by a Network Pharmacy, Administrator will reimburse the Network Pharmacy the amount specified in the Network Pharmacy Agreement for the dispensed Prescription Drug less any Cost-Sharing Amounts. "Clean Claim" means a Prescription Claim prepared in accordance with the NCPDP-promulgated standard format that contains all information necessary for processing for a Prescription Claim and submitted for payment no later than 30 days after the date of service, or a longer period of time if required by applicable Laws and Regulations. Administrator will reimburse Network Pharmacies for each Clean Claim no later than 30 days after Administrator's receipt of the Clean Claim, or a lesser period of time if required in the Network Pharmacy Agreement. Administrator will deny Prescription Claims that are not Clean Claims at point-of-service no later than 60 days after request for payment, or a lesser time period if required in the Network Pharmacy Agreement. Administrator will use reasonable efforts to advise the Network Pharmacy of the basis that a Prescription Claim is ineligible for payment and specify any additional information required for Administrator to pay the Prescription Claim. Administrator will not be financially responsible for paying claims submitted by Network Pharmacies, except that Administrator will be financially responsible for claim liabilities to the extent they arise from Administrator error.
- 1.3.2 <u>Delays</u>, Administrator will not be responsible for any loss, omission or delay of any Prescription Claim by a Network Pharmacy (other than Administrator's Mail Order Pharmacy or Specialty Pharmacy) or other health care professional.
- 1.3.3 <u>Administrative Grievances and Appeals</u>. At Client's request, and subject to section 1.1.3, Administrator will process initial Benefit Plan coverage determinations and exception requests and support Client in connection with Benefit Plan appeals and grievances in accordance with Pharmacy Plan Specifications and this section 1.3 and to the extent required by applicable Laws and Regulations.

1.3.4 Prior Authorization Appeals

- 1.3.4.1. <u>Internal Appeals</u>. Administrator will not conduct Member appeals of prior authorization denials. For a fee, Administrator will facilitate Member appeals of prior authorization denials with an IRO to perform internal appeals. Client will accept and abide by the IRO's process and appeal decisions. After receipt of the IRO's appeal decision, Administrator will remit the appeal decision to the Member on Client's behalf.
- 1.3.4.2. <u>External Appeals</u>. For all external appeals, Client will contract directly with an independent outside party or, where permitted by Laws and Regulations, if Client

requests and at an additional cost to Client, Administrator will coordinate Client's appeals to an IRO for external review. Administrator contracts with three IROs to perform external appeals for non-insurance clients. The parties will agree upon additional fees for the Client or IRO for external appeals. Client will provide Administrator appropriate and necessary plan documentation needed for the IRO to review the adverse benefit determination (within the meaning of 29 C.F.R. § 2560.503-1(m)(4)).

1.4 Benefits Administration and Support

1.4.1 Utilization Management Program

- 1.4.1.1. Development and Support. Administrator will implement its standard utilization management policies, procedures, guidelines and programs for the Benefit Plans to promote cost-effective drug utilization management and to discourage Prescription Drug over and under-utilization. Administrator may, on behalf of Client, (a) communicate with Members to describe health-related products or services (or payment for the products or services) provided by or included in the Plan through the Services, including communications about Network Pharmacies, replacement or enhancement to the Plan, and health-related products or services available only to Members that add value to and are not part of the Plan; (b) conduct population-based activities relating to improving the health of Members and reducing their healthcare costs; and (c) contact Members with health education information and information about Prescription Drugs, treatment alternatives, and related functions. Upon Client's request and at an additional charge to Client, Administrator, in consultation with Client, will develop nonstandard utilization management policies, procedures, guidelines or programs for the Benefit Plans. Upon Client's request, Administrator will communicate Client's utilization program requirements to Members through Client-approved information and outreach materials. Although Administrator will recommend utilization management standards and programs that it believes may be appropriate for the Benefit Plans, Client retains complete and exclusive discretionary authority over its utilization management standards and programs and is responsible ultimately for these standards and programs.
- 1.4.1.2. <u>Administrator's Prior Authorization Services</u>. Administrator will respond to properly submitted prior authorization requests from providers, Members or pharmacies using utilization management standards and guidelines established in accordance with section 1.4.1.1 of this exhibit. Client retains complete and exclusive discretionary authority over approval of prior authorization requests, including Benefit Plan overrides; however, to the extent that Client overrides impact Administrator's compensation, cost to provide Services, or ability to satisfy a guarantee under this agreement, this will be a Service Change and Administrator may adjust the rates, fees or guarantees in Exhibit C in accordance with section 1.3 of the agreement.
- 1.4.2 <u>Client Prior Authorization and Overrides</u>. If Client chooses to perform prior authorizations or benefit overrides, then Administrator will provide Client access to the information in Administrator's computer systems that Client needs to perform these functions.
- 1.4.3 Quality Assurance Program. Administrator will establish a quality assurance program for the Benefit Plans that includes quality measures and reporting systems targeted at reducing medical errors and adverse drug interactions. Administrator will assist in implementing Client's quality assurance and patient safety programs. Administrator will perform activities to support Client's quality assurance requirements under applicable Laws and Regulations. In addition, Administrator will develop and implement systems or

- require Network Pharmacies to implement systems to: (a) offer Member counseling, when appropriate; (b) identify and reduce internal medication errors; and (c) maintain up-to-date Member quality assurance and patient safety program information. Upon Client's request, Administrator will communicate Client's quality assurance standards and programs to Governmental Authorities in the manner prescribed by applicable Laws and Regulations.
- 1.4.4 <u>Targeted Disease Intervention Program.</u> Upon Client's request and for an additional charge to Client, Administrator will help Client develop and operate a targeted disease intervention program for the Benefit Plans that is designed to promote appropriate use of medications and improve therapeutic outcomes for targeted Members. Administrator, on Client's behalf, will coordinate and Implement the targeted disease intervention program. Also, upon Client's request and at an additional cost to Client, Administrator will communicate with Members about the targeted disease intervention program through Client-approved information and outreach materials. Notwithstanding anything in this Section 1.4.4, Administrator will not charge Client for any services provided to Client or Members related to Members' participation in Client's diabetic and hypertension coaching program.
- 1.4.5 Other Clinical Services. Upon Client's request and for an additional charge to Client, Administrator will help Client develop and implement additional quality initiatives, intervention programs or other clinical services.

1.5 Formulary

- 1.5.1 <u>Formulary Adoption</u>. Client will adopt as the Formulary one or more of Administrator's formularies that are developed and maintained by Administrator's formulary advisory committee, as described in section 1.5.4 of this exhibit.
- 1.5.2 <u>Formulary Management</u>. Administrator will provide Client copies of the Formulary to distribute to plan providers and other appropriate parties semi-annually. Except as provided in this agreement, Client will not copy, distribute, sell or otherwise provide Administrator's formularies, including the Formulary, to another party without Administrator's prior written approval.
- 1.5.3 Formulary Changes. Administrator will include in the Formulary new FDA-approved medications as required by the Pharmacy Plan Specifications according to the following schedule: (a) if an open formulary, per the Pharmacy Plan Specifications, all new covered FDA-approved medications (formulary and non-formulary) will be included in the Formulary upon publication in the Medi-Span pricing index and loading into Administrator's systems or (b) if a closed formulary, per the Pharmacy Plan Specifications, all new covered FDA-approved medications (formulary only) will be included in the Formulary after review and addition to the Formulary by Administrator's formulary advisory committee. Following changes to the Formulary, Administrator, at Client's request, will provide or make available appropriate notifications of Formulary changes to Client, Members, prescribing physicians, Network Pharmacies and state pharmaceutical assistance programs as required by applicable Laws and Regulations and agreed to by the parties. If Client makes any change to its Formulary, not initiated by Administrator, or Benefit Plan, or adopts any formulary or utilization management program other than one of the options offered by Administrator under its formulary or utilization management programs, Administrator may adjust the rates, fees or guarantees in Exhibit C, effective the date of the change.
- 1.5.4 Formulary Advisory Committee. Administrator's formulary advisory committee will develop and maintain Administrator's formularies by: (a) selecting Prescription Drugs to include in Administrator's formularies based upon objective evaluation of the therapeutic merits, safety and cost of the Prescription Drug; (b) periodically revisiting Administrator's

formularies, evaluating new and therapeutically equivalent Prescription Drugs for inclusion in the formularies; (c) establishing programs and procedures to address cost-effective drug therapy; (d) reviewing requests to include non-formulary Prescription Drugs in Administrator's formularies; (e) implementing client educational programs; (f) advising Administrator on other matters about the use of Prescription Drugs; (g) overseeing client drug utilization review programs or quality assurance programs or auditing and reviewing the programs' results; and (h) reviewing adverse drug reactions and making recommendations to minimize their occurrence. Administrator's formulary advisory committee's functions, deliberations and results, including development and maintenance of Administrator's formulary, constitute opinions only of Administrator's formulary advisory committee and will not bind Administrator.

1.5.5 No Endorsement. Administrator's development and maintenance of its formularies will not be construed as an endorsement of any prescription drug product or drug manufacturer. Administrator will not be responsible for any actions or omissions of its formulary advisory committee or any adverse consequences that may relate, directly or indirectly, to Client's or a Member's reliance on Administrator's formulary advisory committee.

1.6 Rebate Management

- 1.6.1 Rebate Eligibility. Client will have a claim against Administrator for a Rebate if: (a) Exhibit C specifies that Client will be eligible for Rebates; (b) Client satisfies the minimum Rebate contract criteria and has included the Drug Manufacturer's Prescription Drug on its Formulary, and (c) Administrator has received Rebates resulting directly from Client's satisfaction of the foregoing clause (b). Administrator, in its sole and absolute discretion, may enter into Rebate Agreements with Drug Manufacturers that have Prescription Drugs on Administrator's or its clients' formulary. Many factors affect the amount of Rebates, including benefit design, arrangements with Drug Manufacturers, volume of Prescription Claims, formulary structure and Administrator's overall business strategy. Claims that will not be submitted to Drug Manufacturers for Rebates include Prescription Claims: (a) with invalid service provider identification or prescription numbers; (b) where, after meeting the deductible, the Member's Cost-Sharing Amount under the applicable Benefit Plan requires the Member to pay more than 50% of the Prescription Claim; (c) that are manufacturer negotiated fee products not listed on Client's Formulary for devices without a Prescription Drug component; (d) that are re-packaged NDCs; (e) from 340B pharmacies or other entities eligible for federal supply schedule prices (e.g., Department of Veterans Affairs, U.S. Public Health Service, Department of Defense); or (f) that are not for Prescription Drugs (except for insulin or diabetic supplies).
- 1.6.2 <u>Rebate Guarantees</u>. Except for any Rebate guarantees described in **Exhibit C**, Administrator has no obligation to obtain any particular amount of Rebates for Client. Rebate guarantees are subject to Client's eligibility for Rebates and the Rebate guarantee contingencies described in this section 1.6 and **Exhibit C**.
- 1.6.3 Collection. Administrator will use commercially reasonable efforts to collect Rebates. Administrator will not be responsible for any non-payments or partial payments by Drug Manufacturers of amounts owing under a Rebate Agreement. Adjustments to Rebates or Rebate guarantees may result from patent expirations or Client changes to Formulary or Benefit Plan design effective the date the expiration or change occurs. To the extent of any overpayment or erroneous payment to Client by Administrator, Client will refund immediately the payment or Administrator may recoup the payment from other sums due Client in accordance with section 3.5 of this agreement. Administrator may dispute any overpayment or erroneous payment to Client no later than 180 days after the payment date, except for Rebate repayments resulting from Drug Manufacturer audits.

- 1.6.4 <u>Disbursement</u>. Provided there is no payment default under section 2.2.2 of this agreement, Administrator will disburse, apply and allocate all applicable estimated or actual amounts earned on account of Rebates received by Administrator from Drug Manufacturers based upon the provisions set forth in this exhibit. Administrator will pay Client all Rebates within 90 days after the close of each calendar quarter based on cash received and applied during the quarter. Administrator shall reconcile within 180 days after each calendar year. Administrator will pay Client all Rebates collected within 24 months after termination of this agreement.
- 1.7 Client Incentives and Purchase Discounts. If Client, or its Affiliate or agents, contracts with another party, including a Drug Manufacturer, for a discount, utilization limit, rebate or other incentive associated with the utilization of a Prescription Drug, Client will be in material breach of this agreement, and Administrator, in addition to any other remedies available to it under this agreement, may determine in its sole discretion that Client will not be eligible for any applicable Rebates or ASP Guarantees (as defined in Exhibit C) under this agreement and adjust or eliminate any guarantees, including Rebate guarantees or ASP Guarantees, described in Exhibit C. Client will accept only amounts due under this agreement on account of eligible and legitimate Members. Upon request, Client will cooperate fully with Administrator or a Drug Manufacturer to verify Client's participation in any Rebate program and that all Rebate-related payments were made solely for Covered Prescription Services to eligible and legitimate Members. Administrator, in its capacity as a Mail Order Pharmacy or a Specialty Pharmacy, purchases Prescription Drugs from Drug Manufacturers and receives certain discounts and purchase rebates from Drug Manufacturers in connection with these purchases. Administrator retains these discounts and purchase rebates and does not pass them on to Client.
- 1.8 **E-Prescribing**. Upon Client's request and at an additional charge to Client, Administrator will provide prescribers with electronic access to Member Benefit Plan information, including: (a) Member eligibility status: (b) Member medication history; (c) Formulary status of the Prescription Drug being prescribed; (d) listing of Generic Drug or Brand Name Formulary alternative medications; (e) Member coverage information where applicable; (f) applicable Cost-Sharing Amount; and (g) drug classification information required by the Centers for Medicare & Medicaid Services or successor Governmental Authority.

2. MAIL ORDER PHARMACY SERVICES

- 2.1 Mail Order Services. Administrator, in its capacity as a Mail Order Pharmacy, will provide Client with Mail Order Pharmacy Covered Prescription Services to Members in accordance with the Pharmacy Plan Specifications for the Prescription Drug Compensation established in Exhibit C. Once a prescription for a Covered Prescription Service has been transmitted to Administrator, in its capacity as Mail Order Pharmacy, Administrator will promptly prepare, package and ship the applicable Covered Prescription Service to the Member or other authorized person or entity. Administrator will provide customer service support for Members who use Mail Order Pharmacy Services. Upon request, Administrator will make available to Client mail service brochures for distribution to Members.
- Standards and Professional Judgment. Administrator's Mail Order Pharmacies will be duly licensed under applicable Laws and Regulations of the state of the pharmacies' geographic locations and any other jurisdiction as necessary to furnish Covered Prescription Services to Members. Administrator will comply with all Laws and Regulations promulgated by the Board of Pharmacies that apply to the Administrator Mail Order Pharmacies. Administrator will notify promptly Client if the required licensure of Administrator's Mail Order Pharmacies is lost, suspended, limited or conditioned. Duly licensed personnel will provide all Covered Prescription Services at the Mail Order Pharmacies in accordance with applicable Laws and Regulations and generally accepted standards of practice in the local community of pharmacists. Each Mail Order Pharmacy must use independent professional judgment when dispensing Covered

- Prescription Services and may refuse to dispense any Prescription Drug based upon the pharmacist's professional judgment.
- 2.3 Control of Administrator. Administrator will solely and exclusively control and supervise the operation and maintenance of Administrator's Mail Order Pharmacies and their respective facilities and equipment and provision of Covered Prescription Services. All decisions respecting the provision of Covered Prescription Services by Administrator's Mail Order Pharmacies will be made solely by Administrator and its duly authorized personnel, and not by Client. The relationship between a Member and a Mail Order Pharmacy will be subject to the rules, limitations and privileges incident to the pharmacist-patient relationship. Administrator may exclude from coverage under this agreement a Prescription Drug that cannot be dispensed in accordance with Administrator's mail order pharmacy dispensing protocols or requires special record-keeping procedures.
- 2.4 Mall Order Rates. Prices stated for Prescription Drugs dispensed by the Mail Order Pharmacy are based on the average days supply specified in Exhibit C. Mail Order Pharmacy Prescription Drugs dispensed in smaller amounts will be compensated at the retail pharmacy compensation rates stated in Exhibit C. Specialty Drugs are not available at mail order rates, even if dispensed by a Mall Order Pharmacy. If Client requests or requires expedited or alternative shipping methods other than Administrator's standard method, Client will be solely responsible for those costs. If USPS rates increase, Administrator may pass these cost increases on to Client.

3. SPECIALTY PHARMACY SERVICES

3.1 Specialty Services. Administrator, in its capacity as a Specialty Pharmacy, will provide Client with Exclusive Specialty Drug Covered Prescription Services to Members as specified in Exhibit C. Client will receive the Specialty Drugs specified in Exhibit C as a Covered Prescription Service exclusively from Administrator's Specialty Pharmacy and not from any other retail, mail, specialty or other pharmacy, including a Network Pharmacy.

3.2 Addition of Newly Acquired or Approved Specialty Drugs

- 3.2.1 From the date a newly acquired or approved Specialty Drug ("New Specialty Drug") becomes available until Client rejects the New Specialty Drug as specified in section 3.2.2 of this exhibit, Client authorizes and directs Administrator to make the New Specialty Drug available to Members as part of the Specialty Drug Covered Prescription Services and during this period will compensate Administrator for the New Specialty Drug at the rate specified in Exhibit C. Administrator will not be required to make available to Client or Members a New Specialty Drug that has limited distribution or market access, such as a New Specialty Drug with one distributor or manufacturer.
- 3.2.2 On a periodic basis, Administrator will review the Specialty Drugs listed in Exhibit C and notify Client of the name and price of any New Specialty Drugs to be added to this list of Specialty Drugs. From the date of Client's receipt of this notice, Client will have 30 days to provide Administrator with notice of rejection of additions to the Specialty Drugs listed in Exhibit C.
- 3.2.3 No later than 45 days after Administrator's receipt of Client's notice of rejection of New Specialty Drugs, Administrator will remove the New Specialty Drugs to Exhibit C and dispense the New Specialty Drugs to Members at the pricing specified in Administrator's notice. If Client does not notify Administrator of its rejection of the New Specialty Drugs, Administrator will continue to include the New Specialty Drugs as a Specialty Drug made available to Members.

- 3.2.4 If Client requests that a Prescription Drug be handled as a Covered Prescription Service, but does not want Administrator to handle the Prescription Drug as a Specialty Drug, the parties will consider the request a Service Change and follow the procedures in section 1.3 of this agreement.
- 3.3 Standards and Professional Judgment. Administrator's Specialty Pharmacies will be duly licensed under applicable Laws and Regulations of the state of the pharmacies' geographic locations and any other jurisdiction as necessary to furnish Covered Prescription Services to Members. Administrator will comply with all Laws and Regulations promulgated by the boards of pharmacies that apply to the Administrator's Specialty Pharmacies. Administrator will notify promptly Client if the required licensure of Administrator's Specialty Pharmacies is lost, suspended, limited or conditioned. Duly licensed personnel will provide all Covered Prescription Services at the Specialty Pharmacies in accordance with applicable Laws and Regulations and generally accepted standards of practice in the local community of pharmacists. Each Specialty Pharmacy must use independent professional judgment when dispensing Covered Prescription Services and may refuse to dispense any Prescription Drug based upon the pharmacist's professional judgment.
- 3.4 Control of Administrator. Administrator will solely and exclusively control and supervise the operation and maintenance of Administrator's Specialty Pharmacies and their respective facilities and equipment and provision of Covered Prescription Services. All decisions respecting the provision of Covered Prescription Services by Administrator's Specialty Pharmacies will be made solely by Administrator and its duly authorized personnel, and not by Client. The relationship between a Member and a Specialty Pharmacy will be subject to the rules, limitations and privileges incident to the pharmacist-patient relationship.

4. MEDICARE PART D RETIREMENT DRUG SUBSIDY ("RDS") SERVICES

- 4.1. Administrator's Responsibilities. Administrator will provide Client with RDS Services as follows:
 - 4.1.1.Administrator will provide Client's monthly data files to Client for Client's submission to Centers for Medicare and Medicaid Services ("CMS") in support of Client's claims for federal subsidy payments pursuant to 42 U.S.C. § 1395w-132. Administrator will upload all files and data to CMS through the CMS RDS computer system.
 - 4.1.2.Administrator will, upon request, provide Client with standard reports that meet CMS requirements.
 - 4.1.3.Administrator will submit to Client costs data that reflects any discounts, chargebacks, rebates, and other price concessions given by the manufacturer or pharmacy in the aggregate that are attributable to gross retiree costs between the cost threshold and cost limit as required by CMS.
 - 4.1.4.Administrator is bound by all applicable federal Laws and Regulations, guidance and authorities pertaining to claims and debt collections.
 - 4.1.5.Administrator will maintain the records for Client's RDS Program for six years after the expiration of Client's plan year in which the costs were incurred. CMS or the Office of Inspector General, or their designees, will be provided access to such records upon request.
 - 4.1.6.Administrator agrees that Client's Medicare Drug Plan Members are afforded protection from liability for payment of fees that are Administrator's obligation in accordance with 42 CFR § 423.505(g).

- 4.2. Client Responsibilities. Client will be responsible for the following obligations in connection with Administrator providing Client with RDS Services:
 - 4.2.1.Client will pay administration fees as listed on RDS Compensation Exhibit for Services associated with Client's claims for federal subsidy payments pursuant to 42 USC § 1395w-132.
 - 4.2.2.Client will complete and submit any required updates, revisions, or changes to the application to CMS for approval of subsidy payments pursuant to 42 USC § 1395w-132.
 - 4.2.3. Client will upload all files and data to CMS through the CMS RDS computer system.
 - 4.2.4.Client will upload Member eligibility information and data to CMS. Client will provide Administrator with the final eligibility file generated by CMS.
 - 4.2.5. Client will create and distribute Creditable Coverage statements to Members.
 - 4.2.6. Client will submit information in compliance with the requirements that govern payments set forth in 42 C.F.R. § 423.888.
 - 4.2.7.Client will. at all times, be in compliance with all requirements for continued approval of the federal subsidy program for prescription drug benefits to its Medicare-eligible Members for whom subsidy payments are received from CMS and all Laws and Regulations related to its subsidized pharmacy benefit program.

4.3. Additional RDS Provisions

- 4.3.1. <u>Business Integrity</u>. Administrator will be bound by the provisions set forth at 45 CFR Part 76. In addition to the foregoing, Administrator represents and warrants that neither Administrator nor any personnel furnishing Prescription Drug Services to Medicare Drug Plan Members have been or will be (a) listed as debarred, excluded or otherwise ineligible for participation in federal health care programs or (b) convicted of a criminal felony. If at any time Administrator becomes aware of any violation of this representation and warranty, Administrator will notify Client in writing immediately. If Administrator becomes debarred or ineligible then Client may terminate this agreement immediately upon notice to Administrator without liability to Client or take such other corrective or remedial action as warranted under the circumstances.
- 4.3.2. Federal Policies; Flow Down Provisions. Because Administrator is furnishing Prescription Drug Services to Medicare Drug Plan Members that are the subject of a contract between Client and CMS, the following obligations are imposed upon Administrator with which Administrator will comply: Title VI of the Civil Rights Act of 1964, as amended (42 USC § 2000d et seq.); Sections 503 and 504 of the Rehabilitation Act of 1973, as amended (29 USC § 793 and 794); Title IX of the Education Amendments of 1972, as amended (20 USC § 1681 et seq.); Section 654 of the Omnibus Budget Reconciliation Act of 1981, as amended (41 USC § 9849); the Americans with Disabilities Act (42 USC § 12101 et seq.); and the Age Discrimination Act of 1975, as amended (42 USC § 6101 et seq.); the Vietnam Era Veterans Readjustment Assistant Act (38 USC § 4212); together with all applicable implementing regulations, rules, guidelines and standards as from time-to-time are promulgated thereunder by applicable Governmental Authorities.
- 4.3.3. Nondiscrimination. The Prescription Drug Services furnished to Medicare Drug Plan Members will be rendered without regard to health status, race, religion, color, creed, national origin, ancestry, religion, physical handicap, medical condition (including HIV status), mental status, age (except as provided by law), marital status, sex, sexual orientation or gender identity. In addition, Administrator will not unlawfully discriminate

- against any employee or applicant for employment because of race, religion, color, creed, national origin, ancestry, religion, physical handicap, medical condition (including HIV status), mental status, age, marital status, sex, sexual orientation or gender identity. The evaluation and treatment of Administrator employees and applicants for employment are, and will be, free from this unlawful discrimination. Administrator will comply with all Laws and Regulations relating to equal and fair employment.
- 4.3.4. Equal Opportunity Employer. As an equal opportunity employer, Administrator will abide by all applicable provisions of Executive Order 11246, as amended (Equal Opportunity/Affirmative Action), 38 USC § 4212, as amended, (Vietnam Era Veterans Readjustment Act), and Section 503 of the Rehabilitation Act of 1973, as amended (Handicapped Regulations), together with the implementing regulations (found at 41 CFR §§ 60-1, & 60-2, 41 CFR § 60-250, and 41 CFR § 60-741, respectively), rules guidelines and standards, as from time-to-time are promulgated thereunder by applicable Governmental Authorities and which are incorporated by reference into this agreement.
- 4.3.5. Other Laws and Regulations. Administrator will comply with (a) applicable federal Laws and Regulations designed to prevent fraud, waste and abuse, including but not limited to applicable provisions of Federal criminal law, the False Claims Act (31 USC §§ 3729 et seq.), and the anti-kickback provision of section 1128B of the Social Security Act; (b) applicable HIPAA Administrative Simplification Security and Privacy rules at 45 CFR parts 160, 162, and 164; and (c) all other applicable federal Laws and Regulations."

EXHIBIT C

COMPENSATION

Price Summary OPTUMRX**

Confidential and Proprietary

Retail Network

The following administrative fees and rates are exclusive to City of Cincinnati and are based upon provided information and an estimated 13,997 Active, or 6,138 Retiree, or a combined total of 20,135 Active and Retiree eligible members or greater. Rates and fees are effective upon the implementation of services on 01/01/2014. This summary represents our Pass-Through Pricing model. All rates and fees are guaranteed and subject to the applicable terms in this cost proposal unless stated as otherwise.

Discounts and Dispensing Fees		Active	Retiree	Active + Retiree							
● Access to over 65,000 pharmacies nationwide	Brand:	2014: AWP-14.7%	2014: AWP-14.7%	2014: AWP-14.7%							
Rates exclude compound and DMR claims	Transport	2015: AWP-15.3%	2015: AWP-15.3%	2015; AWP-15.3%							
		2016: AWP-15.7%	2016: AWP-15.7%	2016: AWP-15.7%							
		\$1.39 DF	\$1.37 DF	\$1.38 DF							
	Generic:	MAC \$1.41 Dispensing Fee	MAC \$1.39 Dispensing Fee	MAC \$1.40 Dispensing Fee							
	Aggregate av	Aggregate average minimum discount off AWP for									
	MAC & non-N	AC generics:									
	2014 2015 2016	AWP –75.3% AWP –77.0% AWP –77.2%	AWP78.4% AWP80.2% AWP80.4%	AWP -79.7% AWP -81.4% AWP -81.7%							
OptumRx Mail Service											
Discounts and Dispensing Fees		Active	Retiree	Active + Retiree							
Postage included											
Rates may vary for claims not covered under	Brand:	AWP -22.0%	AWP -22.5%	AWP -23.0%							
pharmacy benefit Mail discounts and dispensing fees exclude		\$0.00 Dispensing Fee	\$0.00 Dispensing Fee	\$0.00 Dispensing Fee							
specialty and certain non-specialty injectable products											
	Generic:	MAC	MAC	MAC							
		\$0.00 Dispensing Fee	\$0.00 Dispensing Fee	\$0.00 Dispensing Fee							
	Aggregate av	erage minimum disco	unt off AWP for								
	MAC & non-N	AC generics:	•								
	2014	AWP -80.4%	AWP80.4%	AWP -82.7%							
	2015	AWP -81.1%	AWP -82,2%	AWP -83.4%							
	2016	AWP -81.2%	AWP -82.4%	AWP -83.7%							
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Price Summary OPTUMRX**

ptumRx Specialty Pharmacy				
Discounts and Dispensing Fees		Active	Retiree	Active + Retiree
Postage included				
 Rates for these products dispensed from 	Brand:	AWP -15.5%	AWP -16.0%	AWP16.0%
from OptumRx's mail pharmacies are listed in the		\$0.00 Dispensing	\$0.00 Dispensing	\$0.00 Dispensing
specialty exhibit		Fee	Fee	Fee
17) albada 100 maar aan 4				
Rebate Management				
Rebate guarantees are contingent upon the following terms:		Active	Retiree	Active + Retiree
City of Cincinnati's adoption of OptumRx's		Active	Notifico	
formulary, formulary management, and	Retail:	100% Pass-Through	100% Pass-Through	100% Pass-Through
utilization management	(with Specialty)			Zaluman.
OptumRx's collection and distribution of funds		Minimum Guarantee:	Minimum Guarantee:	Minimum Guarantee:
received	2014	\$28.53 / Brand Claim	\$27.21 / Brand Claim	\$27,74 / Brand Claim
Rebate ineligible paid claims such as those from	2015	\$32.21 / Brand Claim	\$30.40 / Brand Claim	\$31.14 / Brand Claim
340B pharmacies or entities eligible for federal supply	2016	\$32.63 / Brand Claim	\$29.03 / Brand Claim	\$30,52 / Brand Claim
schedule prices (e.g., Dept. of Veterans Affairs, US				
blic Health Service, Dept. of Defense) are excluded				
nom rebate guarantees	Mail:	100% Pass-Through	100% Pass-Through	100% Pass-Through
A minimum of \$10 difference in copayment or 10%	(Non-Specialty)			Minimum Oversatas:
difference in coinsurance between preferred and				Minimum Guarantee: \$86.54 / Brand Claim
non-preferred branded drugs		•	***************************************	\$95.57 / Brand Claim
 Rebates may not include Prescription Claims that require the Members to pay more than 50% of the total 			1	\$88.91 / Brand Claim
annual cost for all Prescription Claims under the	2010	φον.ου γ Diqiiu Qiaiiii	WOS.40 / DIGING CIGINI	poolo () Diana Olam
applicable Benefit Plan				
Any deviations to the Administrator's Formulary and	Mail	100% Pass-Through	100% Pass-Through	100% Pass-Through
Utilization Management that adversely impacts rebates	Specialty:	g	_	
will result in a proportional adjustment to the		Minimum Guarantee:	Minimum Guarantee:	Minimum Guarantee:
corresponding rebate guarantees	2014	\$505.13 / Brand Claim	\$315.96 / Brand Claim	\$409.75 / Brand Claim
 Unrestricted access to 90 days supply scripts 		g ·	\$456.04 / Brand Claim	
in retail	2016	\$667.31 / Brand Claim	\$431.37 / Brand Claim	\$545.80 / Brand Claim
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Price Summary OPTUMRX

Connucinal and Prophetary			
Standard Services Dedicated Implementation and Client Management Team Client Manager Project Manager Client Service Representative Pharmacist Business Analyst Help Desks – Toll-free access for members, physicians, and pharmacies DUR and System Edits – Standard Concurrent DUR and flexible plan designs Communication Materials - Welcome Package and standard ID cards Internet Direct Access Real time access to claims and eligibility system Accounts set up for up to two users Real-Time Audit – Filters 100% of claims before payment–outiliers sent to audit team Eligibility Maintenance – Via FTP or encrypted e-mail Website Access – www.optumrx.com Safety Notifications for Providers and/or Members (e.g., drug recalls) Standard Reporting Package – Integrated retail and mail claim data Online Reporting Tool - Software, training and maintenance costs for up to two users	\$1.50 per Paid Claim	\$1.50 per Paid Claim	\$0.95 per Paid Claim
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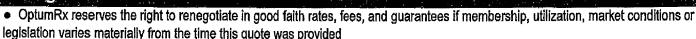
Ulinical Programs	
Bundled Clinical Programs Programs include: DIAP, Geriatric Monitor, Narcotic, and PolyPharmacy	\$0.02 PMPM for Bundle of 4 Programs
Generic Strategy Program	\$0.05 PMPM
Condition Specific Bundle Programs Includes: Statin Initiative, Asthma Program and Migraine Prophylaxis	\$0.05 PMPM per Bundle of 3 Programs or \$0.02 PMPM for each Program selected individually
Adherence Program Increases medication adherence rates in a number of chronic and high-impact disease states	\$0.02 PMPM
Other Standard Programs	\$0.02 - \$0.05 PMPM per Program selected
Health, Wellness, and Disease Education provided through www.optumrx.com	Included
Customized Clinical Programs	Quoted Separately Upon Request. Client claims data required for custom analysis and presentation.
Clinical Prior Authorization Overrides requiring clinical intervention or evaluation	\$30 per Authorization
Physician Reviewed Prior Authorization	\$225 per Authorization
Clinical Appeals Services	\$550 per Level
Administrative Appeals Services	\$180 per Level
Additional Services	
Custom Programming/Report Generation Minimum \$500.00	\$150 per Hour
E-Prescribing	Included
Non-Standard or Manual Eligibility Maintenance	\$1.50 per Member
Direct Member Reimbursement (DMR) Entered by OptumRx, includes creation and mailing of letters for denied claims, in accordance with state or federal requirements	\$4.50 per claim + Postage
Credits and Allowances	
Implementation Allowance ■ Based upon documented actual out of pocket implementation costs incurred by Plan	Up to \$4.00 Credit per Transitioned Member
Pre-implementation Audit Allowance Based upon documented actual out of pocket costs incurred by Plan	Up to \$30,000 for Audit Up to \$3,000 for Travel

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Price Summary OPTUMRX

Confidential and Proprietary

Pricing Terms



- All rates and fees are contingent upon the selection of OptumRx as the exclusive mail provider, and a similar benefit design as applicable to the historical data provided for the purpose of this cost proposal that includes an exclusive Specialty arrangement. All rates and fees are subject to change otherwise
- The rates, fees or guarantees provided to City of Cincinnati for this cost proposal were based on a set of assumptions used by OptumRx ("Administrator") from the data and other pertinent information provided by City of Cincinnati to Administrator. In the event that the data/information is deficient or inaccurate, resulting in a material change to the relative economics of Administrator or City of Cincinnati, some of the rates, fees or guarantees may no longer apply. If such an event were to occur, Administrator will make every attempt to negotiate with City of Cincinnati in good faith to arrive at a solution that is mutually acceptable to both parties involved
- Any reduction in rebates earned by OptumRx as a result of changes to formulary, days supply, and/or benefit design not initiated or approved by OptumRx may result in adjustments to rebate payments or guarantees effective concurrently with such occurrences
- Rebate guarantees and generic AWP discounts may be adjusted proportional to the impact of unexpected releases of generic products to market, or the withdrawal/recall of existing branded products
- Mail discounts and dispensing fees exclude specialty and certain non-specialty injectable products. Rates for these products dispensed from OptumRx's mail pharmacies are listed in the specialty exhibit
- Mail Service rates are based on an average days supply of 84 or greater for all claims with the exception of all specialty and certain provided

Specialty Pharmacy Price Summary



Confidential and Proprietary

The rates and terms quoted in this price summary and attached Specialty Pharmacy Pricing Schedule are subject to the accuracy and completeness of the information provided by Client, including the minimum number of Members, and the parties entering into a definitive agreement for the provision of pharmacy benefit management services by the implementation Date.

Date:	6/20/2013
Client:	City of Cincinnati
Members:	20,135
Implementation Date:	1/1/2014

Drug Pricing OptumRx dispenses all drug label names on the pricing schedule provided. A comprehensive list of NDCs can be provided upon request.	See attached Specialty Pharmacy Pricing Schedule
Specialty Drugs may include: • Ancillary supplies • Needles • Syringes • Sharp containers	Included at no extra charge
Value Added Services Monthly Member contact by patient care coordinators Access to pharmacist around the clock Distribution of medications to place of choice within the U.S. and its territories Refill reminder program Patient assistance programs	Included at no extra charge
Clinical Management Program Programs include Multiple Sclerosis, Inflammatory Conditions, Transplant, Oral Oncology, HIV/Aids, Hepatitis C, Hemophilia Member customized care plans Continuity of care with minimal clinician changes for Members Personal one-on-one Member phone assessment/reassessment consultation	Included at no extra charge

Specialty Pharmacy Price Summary



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Utilization Management Prior authorization or case review -Review requests for Specialty Drugs to meet Client's utilization management program (including step therapy) and request additional information if needed -Accept Specialty Drug authorization by phone or fax -Verify Member eligibility -Mail denial letters to Members and providers as required by applicable Laws and Regulations	\$55 per case
Physician review of submitted documentation, if needed, as required by applicable Laws and Regulations	\$390 per physician reviewed case
Dose optimization/waste avoidance	Included at no extra charge
Compliance Management Refill reminder process with three calls to Members starting five days before refill Notify providers for non-compliance management	Included at no extra charge
Fulfiliment Process • Shipping to location of choice within the U.S. and its territories • Monitoring and tracking shipping process • Postage	Included at no extra charge
Standard Reports • Prior authorization / return on investment savings analysis • Member adherence by percent compliance • Annual report • Quarterly utilization reports	Included at no extra charge
Online Reporting Tool • Software and training for up to three users	Included at no extra charge
Client-requested custom system or reporting configurations	\$150 per hour



Specialty Pharmacy Pricing Schedule Confidential and Proprietary

The rates and terms quoted in this price summary and attached Specially Pharmacy Pricing Schedule are subject to the accuracy and completeness of the information provided by Client, including the minimum number of Members, and the parties entering into a definitive agreement for the provision of pharmacy benefit management services by the Implementation Date. OptumRx has available in its Specialty Pharmacies the listed Specially Drugs regardless of package size or NDC. A comprehensive list of NDCs can be provided upon request. Products listed below that are not available to be dispensed directly by OptumRx will be dispensed through a third party vendor at the specified rates.

620/2013 City of Cincinnati 20,135 01/01/2014 Exclusive Spedalty Pharmacy Program Implementation Date: Specially Service: Members: Date: Client

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Service Control of the Control of th	AS OF THE OWN	-16.0%	-13.5%	-13.5%	-13.5%	.15.0%	-11.0%	-47 090	-16.5%	-15.0%	-15.0%	-15 0%	15.0%	-15.0%	-15.0%	-15.0%	.15.0%	-16.5%	-18.5%	-15.0%	20.0	50%	45.0%	45.0%	-45.0%	-15.0%	-33.0%	-13.5%	-16.5%	-18.5%	-16.5%	-15,5%	-13.5%	-13,5%	-18.0%	-18.0%	-18.0%	-18.0%	*18.0%	-47.592
	5646020002120	6800005000202D	64154090102020	64154090102010	64154090102030	23100030008920	44603060002120	16000005402170	07000070002520	83101010102015	83101010102053	83101010102058	83101010102050	83101010102020	83101010102065	83101010102040	83101010102D45	72170085003020	72170085000320	21755030002120	93400030001920	50280035192130	50250035102001	50250035102010	50250035102015	88300017102320	85100033006440	75800040002220	39480050200130	39480050200140	39480050200120	21755050102120	21765060002120	30907715002120	21758050002010	99392070000130	99392070000135	99392070000140	99392070000120	24500049304090
Removed	CANAKINUMAB FOR INJ 180 MG	PEGLOTICASE INJ 8 MGML (FOR IV INFUSION)	ZICONOTIDE ACETATE INTRATHECAL INJ 100 MCG/ML	ZICONOTIDE ACETATE INTRATHECAL INJ 500 MCG/ZOML (25 MCG/ML)	ZICONOTIDE ACETATE INTRATHECAL INJ 500 MCG/SML	TESTOSTERONE IMPLANT PELLETS 75 MG	OMALIZUMAB FOR INJ 150 MG	AZTREONAM LYSINE FOR INHAL SOLN 75 MG (BASE EQUIVALENT)	TOBRAMYCIN NEBU SOLN 300 MG/SML	DALTEPARIN SODIUM INJ 10000 UNITAM	DALTEPARIN SODEUM INJ 12500 UNITROSML	DALTEPARIN SODYUM INJ 15000 UNITO.BML	DALTEPARIN SODIUM INJ 18000 UNITIO ZZML	DALTEPARIN SOBIUM IN J. 2500 UNITRO. ZML	DALTEPARIN SODIUM IN J 25000 UNITML	DALTEPARIN SODIUM IN 5000 UNITRIZML	DALTEPARIN SODIUM INJ 7500 UNIT/0.3ML	MGABATRIN POWD PACK 500 MG	INGABATRIN TAB 500 MG	GLICARPIDASE FOR IV INJ 1000 UNIT	NALTREXONE FOR IM EXTENDED RELEASE SUSP 380 MG	FOSAPREPITANT DIMEGLUMINE FOR IN INFUSION 150 MG (BASE EQ)	GRANISETRON HCL INJ 0.1 MG/ML	GRANISETRON INCLINA 1 MG/ML	GRANISETRON MCL INJ 4 MG/4ML (1 MG/ML)	FLUOCINOLONE ACETONIDE INTRAVITREAL IMPLANT 0.59 MG	FACTOR XIII CONCENTRATE (HUMAN) FOR INJ KIT 1000-1800 UNIT	HYLAN INTRA-ARTICULAR INJ 8 MGML	LOMITAPIDE MESYLATE CAP 10 MG (BASE EQUIN)	LOMITAPIDE MESYLATE CAP 20 MG (BASE EQUIV)	LOMITAPIDE MESYLATE CAP 5 MG (BASE EQUIV)	LEVOLEUCOVORIN CALCIUM FOR IV INJ 50 MG (BASE EQUIN)	PALIFERMIN FOR IV INJ 6.25 MG	ALGLUCOSIDASE ALFA FOR IV SOLN 50 MG	MESNA INJ 100 MG/ML	THALIDOMIDE CAP 100 MG	THALIDOMIDE CAP 150 MG	THALIDOMIDE CAP 200 MG	THAUDOMIDE CAP 50 MG	PACI ITAXEL PROTEIN BOLD DARBOLI ES EDE IVISI ES AGRAGO
		KRYSTEXXA	PRINCT	PRIMLT	PRIALT	TESTOPEL	XOLAIR	CAYSTON	rosi	FRAGMIN	FRAGMIN	FRAGMIN	FRAGMIN	FRAGMIN	FRAGMIN	FRAGMIN	FRAGMIN	SABRL	SABRIL	VORAXAZE	VIVITROL	EMEND	GRANISETRON	GRANISETRON	GRANISETRON	RETISERT	CORFACT	SYNVISC	JUXTAPID	JUXTAPID	UXTAPID	FUSILEV	KEPIVANCE	LUMEYME	MESNA	THALOMID	THALOMID	THALOMID	THALOMID	ABSAYANE
R. Personal	Analgesic & Anesthetic Agents				& Anesthetic Agents		atic & Monoclonal Antibodies						Anticoagulants		Anticoagulants .	Anticoagulants		Anticonvulsant	Anticonvulsant		Antidotes	Antlemetic					c Agent											Antineoplastic & Adjunctive Therapies	inctive Therapies	Antineoplastic Agent
CHANGE SAN	ILARIS INJ 180MG	ଣା	- 1	- 1	PRALT INJ SOOMCG	ᆈ	۳I	CAYSTON INH 75MG	٣!	- 1	- 1		FRAGMIN INJ 18000UNT	ı	FRAGMIN IN125000ML	FRAGMIN INJ 5000/0.2	z	SABRIL POW SOOMS	SABRIL TAB 500MG	VORAXAZE INJ 1000UNIT	VIVITROL INJ 380MG	EMEND SOL 150MG	GRAMSETRON INJOINGML	GRANISETRON INJ 1MG/ML	ŒΙ	- 1	니				\sim	FUSILEY INJ SOMG	KEPIVANCE INJ 6.25MG	LUMIZYME INJ 50MG	MESNA INJ 16M			THALOMID CAP 200MG		ARRAXANE INJ 100MG

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≟	00'03	\$0.00	\$0,00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	20.00	20.00	20.00	20.00	20'05	9	30.00	\$0.00	\$0.00	\$0.00	\$0.03	20.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0,00	\$0.00	\$6.00	\$0.00	\$0.00	\$0.00	80.00	3 8	3 5	3 5	8 8	\$0.00	20.00	80,03	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	20.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	20.03	20.00
	-16.9%	-16.0%	-18,0%	-13,5%	-18.0%	-18.0%	-13,5%	-13.5%	-13.5%	-10,378	14.0%	-14.5%	44.5%	-10.09K	-12.0%	-16,0%	-13,5%	-13.5%	-13.5%	-13.5%	-18,0%	-16.0%	960'9-	-8.0%	-11.5%	-13.5%	-13.5%	-13.5%	-13,5%	-11,0%	-11.0%	-18.0%	-18.0%	-18.0%	460'81-	40.070	000,01-	-18.0%	-18.0%	-18.0%	-18.0%	-18.0%	-13.5%	-15.0%	-15.6%	-15.0%	-13,5%	-18.0%	-18.0%	-18.0%	-15.0%	-15.0%	-18.0%	-18.0%	*18.0%	-18.0%	-18,0%	-18.0%	-18.0%	-18.0%
	21355020202120	21200040102115	21200040102105	21200040102110	21200040102010	21300030002020	21532530000330	21532530000310	2153253000520	2435238606323	OLISOUGES PRODUCED	21300053102120	241040102110	21353045001360	21353045001320	21335020002025	21358070001320	21600040002220	21600040002240	21102010002105	21200010102105	21200010102115	21534012000320	21534012000340	21100010002020	21353010002040	21550040102030	21550040102035	21550040102025	21534085000320	21634085000340	21100015002035	21100015002120	21199915902940	21100010002110	24 thoughtones	2120000000000	2110020002025	21100020002030	21100020002020	96465858002900	21300007002010	21300008002020	21534013106470	21534013106480	21534013105460	21200020002105	21101020002125	21101020002130	21101020002120	96468809802900	96468309802900	21300010002105	21300010002040	21300010002115	21300010002011	21300010002120	21300010002110	21700020002105	21700020002110
		DOXORUBICIN HCL FOR INJ 50 MG	DOXORUBICIN HOL FOR INJ 10 MG	DOXORUBICIN HOL FOR INJ 20 MG	DOXORUBICIN HCL INJ 2 MG/ML			EVERUIMUS TAB 2.5 MG	EVERTURATION INDICATE OF THE TABLE AND THE T	CONTROLLING AND IN SOUTH	POWELINGS DESCRION FOR IN SOLIN 100 MG (SASE COUN)	FEMERICASED LASCIDION FOR IV SOLIN SOUNG (BASE EQUIV)	MELTINATAN TILL FOR INC 30 MS (5ASE CAUSY)	OFATIMINAME CONC. FOR IV INFLISION 1000 MEGANI	DEATUNDINAB CONC FOR IV INFUSION 100 MG/SML	BEVACIZUMAB IV SOLN 100 MG/KML (FOR INFUSION)	TOSITUMOMAB FOR IV INJ CONC 14 MG/ML	KODINE I 131 TOSITUMOMAB INJ 0.61 MCIMIL	IODINE 1131 TOSITUMOMAB INJ S.6 MCIML	CARMUSTINE FOR INJ 100 MG	BLEOMYCIN SULFATE FOR INJ 15 UNIT	BLEOMYCIN SULFATE FOR INJ 30 UNIT	BOSUTINDS TAB 100 MG	BOSUTINIB TAB 500 MG	BUSULFAN INJ 6 MGML	ALEMITUZUMAB IV INJ 30 MG/ML (FOR INFUSION)	IRINOTECAN HCL INJ 100 MG/SML (20 MG/ML)	IRINOTECAN HCL INJ 300 MG/15ML (20 MG/RL)	IRRNOTECAN HOL INJ 40 MG/ZML (20 MG/M)	VANDETANIB TAB 100 MG	VANDETANIB TAB 300 MG	CARBOPLATIN IV SOLN 150 MG/15ML	CARBOPLATIN IV FOR INJ 150 NG	CARBOPLATIN IV SOLN 450 MG/45/ML	CARBOPLATIN IV FOR INJOURNE	CARBUTAN IN SOLD OF MANAGEMENT	CARBOTELON TO SOLN SOUTH TO THE	CASPI ATIN TALL 100 MCHODAL (1 MCAM)	CISPLATIN IN 200 MGZOOML (1 MG/ML)	CISPLATIN IN 50 MG/SDML (1 MG/ML)	CISPLATIN (BULK) POWDER	CLADRIBINE IN 1 MG/ML	CLOFARABINE IV SOLN 1 MSML	CABOZANTINIB S-MAL CAP 1 X 80 MG & 1 X 20 MG (100 DOSE) KIT	CABOZANTINIB S-MAL CAP 1 X 50 MG & 3 X 20 MG (140 DOSE) KIT	CABOZANTINIB S-MALATE CAP 3 X 20 MG (60 MG DOSE) KIT	DACTINOMYCIN FOR INJ 0.5 MG	CYCLOPHOSPHAMIDE FOR INJ 1 GM	CYCLOPHOSPHAMIDE FOR INJ 2 GM	CYCL OPHOSPHAMIDE FOR INJ 500 MG				CYTARABINE INJ 100 MGML	CYTARABINE FOR INJ 1 GM	CYTARABINE IN PP 20 MG/ML	CYTARABINE FOR INJ Z GM	CYTARABINE FOR INJ 500 MG	DACARBAZINE FOR IN J 100 MG	DACARBAZINE FOR INJ 200 MG
	ADCETRIS	ADRIAMYC	ADRIAMYCIN	ADRIAMYCIN	ADRIAMYCIN	AURUCA	ALINIOR	AFINITOR	AFINITOR	ATINTA	MUMIN A	ALIMIA	ADDANON	ARZERRA	ARZERRA	AVASTIN	BEXXAR	BEXXAR 131 I	BEXXAR 131 1	BICNU	BLEOMYCIN	BLEOMYCIN	BOSULIF	BOSULIF	BUSULFEX	CAMPATH	CAMPTOSAR	CAMPTOSAR	CAMPTOSAR	CAPRELSA	CAPRELSA	CARBOPLATIN	CARBOPLATIN	CARBOPLATIN	CARBOPLA IN	CARBOPLA! IN	CARBOTCHIRA	CENDEDINE	CISPLATIN	CISPLATIN	CISPLATIN	CLADRIBINE	CLOLAR	COMETRIO	COMETRIA	COMETRIQ	COSMEGEN	СУСТОРНОЅРН	EYCLOPHOSPH	CYCLOPHOSPH	CYCLOPHOSPHA	CYCLOPHOSPHA	CYTARABINE	CYTARABINE	CYTARABINE	CYTARABINE	١	- [. 1	DACARBAZINE
	Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineoptastic Agent	Ankneoplastic Agent	Angneoplastic Agent	Antheopiasic Agent	Antheopyteste Agent	Authorities Agont	Annreoplastic Agent	Antineoplastic Agent	Authorities Agent	Anthropiastic Apen	Antineoplastic Apent	Antineoplastic Agent	Antheoplastic Agent	Animeoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Amineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineophastic Agent	Anthreoplastic Agent	Antineoplastic Agent	Artineoplastic Agent	Artineoplastic Agent	Antineoplastic Agent	Artimeoplastic Agent	Ammedyasuc Agent	Amineopalisac Agent	Authoritatic Apart	Anthropisatic Acent	Antineoplastic Acent	Antineoplastic Agent	Antineoplastic Agent	Antineophasic Agent	Antineoplastic Agent	Antineoplastic Agent	Antheoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Asineoplastic Agent	Artheoplastic Agent	Antheopiastic Agent	Antineoptastic Agont	Antineoptastic Agent	Antheoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineopiastic Agent	Antineoplastic Agent
	ADCETRIS INJ SOMG	ADRIANYC INJ SOMG	ADRIAMYCIN INJ 10MG	ADRIVATION IN ZOMG	ADRIAMYCIN INJEMISMI.	ADRUCIL INJ SOMESMA	AFINITOR TAB 10MG	AFINITOR TAB 2.5MG	THINITOR IABONG	APPLIANT INDIVISIONS	- 1	ALIMTA IN SOUNG	- 1	ADYCODA CON 100/50M3	F	AVASTEN INJ	EXXAR CON 14 MG/ML	SEXXAR 131 I INJ 0.5EMI.	BEXXAR 131 I BUI 5,6MCIM	SECNU INJ TOOMG	BLEOMYCRN INJ 15UNT	SLEOMYCH IN 30UNIT	SOSULIF TAB 100MG	SOSULIF TAB 500MG	BUSULFEX INJEMENT	CAMPATH IN 30MGML	CAMPTOSAR INJ 100/5ML	CAMPTOSAR INJ 300/15ML	CAMPTOSAR INJ 40MG/ZML	CAPRELSA TAB 100MG	CAPRELSA TAB 300MG	CARBOPLATIN IN 150/15ML	CARBOPLATIN INJ 150MG	CARBOPLATIN INJ 450/45ML	ARBOPLATIN INJ 50MG	CARECPLATIN INJ SOMG/SML	CARECPLAIN INJEGUAL	SECURIONE IN TOOKS	CISCLAIN INTERIOR	SIGNO ATIN IN LEGISTRE	SEPTIMENT TO SEPTI	CLADRIBINE INJ 1MG/ML	CLOLAR INJ TMG/ML	COMETRIQ KIT 100MG		COMETRIQ KIT 60MG	COSMEGEN INJUSMG	CYCLOPHOSPH INJ 16M	CYCLOPHOSPH INJ 28M	CYCLOPHOSPH INJ 500MG	CYCLOPHOSPHA POW	CYCLOPHOSPHA POW USP	CYTARABINE INJ 100MG	CYTARABINE INJ 100MG/ML	CYTARABINE INJ 1GM	CYTARABINE INJ 20MG/ML	CYTARABINE INJ2GM	CYTARABINE INJ SOOMG	DACARBAZINE INI 100MG	DACARBAZINE INJ 200MG

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	\$0.00	\$0.00	20.03	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	20.00	20.00	20.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	00'0\$	00'0\$	\$0.00	20.00	\$0.00	\$0.00	20.02	20.02	20.00	00.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0,00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	20,00	20,00	200	20.05	20.00	\$0.00	\$0.00	\$0.00	\$0,00	\$0.00	20.00	20,00	\$0,00	\$0,00
	-11.5%	-18,0%	-18.0%	-18.0%	-13.5%	-14.5%	-14.5%	-25.0%	-25,0%	-25.0%	-25.0%	-25.0%	-25.0%	-25.0%	-25,0%	-14.5%	-18.0%	-18.0%	-18.0%	-18.0%	-14,5%	-14.5%	-14.5%	-14.5%	-13.5%	-19.0%	-19.0%	-18.0%	-13.5%	449.04	-1000	-18.0%	-18.0%	-13.5%	-15.0%	-13.5%	-18.0%	-15.0%	-14.0%	-7.5%	-7.5%	-18.0%	-13.5%	-25.0%	-18.0%	- 18.00 - 18.00	-19.096	-18 094	-18.0%	-15.0%	-15.0%	-15.0%	-16.8%	-15.8%	-13.5%	-1-0% -7-0%	-75.0%	-15.5%	-(5.5%
	21300015002120	21200020002105	21200030102105	21200030102210	21200030052210	21500005002120	21500005002140	21500005002050	21500005001317	21500005001320	21500005002030	21500005001310	21500005001325	21500005001315	21500005002040	21200040402210	21200040102010	21200040102010	21200040102010	21200040102010	21405010156432	21405010206435	21405010256445	21405010108415	21200042102030	21100028002030	Z11000Z800Z035	Z11000Z80Z5	21200042102045	212000421030	21200042102140	21353025002020	21353025002025	2137007000120	21250010402120	21500010602120	21500010002020	96508045482900	21403530002024	21405525102130	21405525102120	21300020002105	21300028102120	21300025102120	21300025102020	Z130003000000	Standandana	213000300035	21300030002035	96526463632900	96526483832900	96526463632900	25300054002020	21300054002025	2130002002105	21305034102140 54305034165140	21300034102160	21300034102140	21300034102110
	DECITABINE FOR INJ 50 MG	AYCIN FOR INJ 0.5 MG	DALINORUBICIN HOL FOR IN 1 20 MG	DAUNORUBICIN HCL IN S MG/ML (BASE EQUIV)	DAUNORUBICIN CITRATE LIPOSOME IN 12 MGML	DOCETAXEL FOR INJ 20 MG	DOCETAXEL FOR INJ 80 MG	DOCETAXEL SOLN FOR IV INFUSION 160 MG/16ML	DOCETAXEL FOR INJ CONC 150 MG/MIL (20 MG/ML)	DOCETAXEL FOR INJ CONC 20 MG/D.5ML (40 MG/ML)	DOCETAXEL SOLN FOR IV INFUSION 20 MG/2ML	DOCETAXEL FOR INJ CONC 2D MG/ML	DOCETAXEL FOR INJ CONC 80 MG/ZML (40 MG/ML)	DOCETAXEL FOR INJ CONC 80 MG/AML (20 MG/ML)	DOCETAXEL SOLN FOR IV INFUSION 80 MG8ML	DOXORUBICIN HCL LIPOSOMAL INJ (FOR IN INFJISION) 2 MG/ML	DOXORUBICIN HOL INJ 2 MGML	DOXORUBICIN HCL IN 2 MGML	DOXORUBICIN HCL IN 2 MG/ML	DOXORUBICIN HCL IN 2 MGML	LEUPROLIDE ACETATE (3 MONTH) FOR SUBCUTANECUS INJ KIT 22.5MG	LEUPROLIDE ACETATE (4 MONTH) FOR SUBCUTANEOUS INJ KIT 30 MG	LEUPROLIDE ACETATE (6 MONTH) FOR SUBCUTANEOUS INJ KIT 45 MG	ILEUPROLIDE ACETATE FOR SUBCUTANEOUS INJERT 7,5 MG	EPIRUBICIN HCL. IN J SO MIGREM, (2 MGML)	DAALIPLATIN IV SOLN TOO MGZOML	CANTID ATHE PLOT NEW MORAWIT	OVACHED IN 19 SOUND DIRECTION.	EPIRUBICIN HOLINIZON MOYCOM, 22 MONOS	EPIREBICIN HOLINI SO MGDSML (2 MGML)	EPIRUBICIN HOL FOR IN 50 MG	CETUXIMAB IV SOLIN 100 MG/50ML (2 MG/ML)	CETUXIMAB IV SOLN 200 MG/100ML (2 MG/ML)	VISMODEGIB CAP 150 MG	ASPARAGINASE ERMINIA CHRYSANTHEMI FOR INJ 10000 UNIT	ETOPOSIDE PHOSPHATE IV FOR INJ 100 MG	ETOPOSIDE INJ 20 MSML	ETOPOSIDE (BULK) POWDER	FULVESTRANT IN 220 MG/SML	DEGARELIX ACETATE FOR INJ 129 MG (BASE EQUIV)	DEGARELIX ACETATE FOR INJ 80 MG (BASE EQUIV)	FLOXURIDINE FOR INJOS GM	FLUDARABINE PHOSPHATE FOR INJ 50 MG	FLUDARABINE PHOSPHATE FOR INJ 50 MG	PLUJARABINE PHOSPIPA E INJ 25 MOM.	FLUCKCONCONCIL IN 1 GRAZUMIL (SO MUSMIL)	PLUCKURACIL INJ. 22 GROUME (30 Metals)	SELECTION OF THE CAMPONI (SO NEW)	FLUOROURACIL INJ S GAVIDOML (50 MGML)	FLUOROURACIL (BULK) POWDER	PLUOROURACIL (BULK) POWDER	FLUGROURACIL (BULK) POWIDER	PRALATREXATE IV INJ 20 MGML	PRALATREXATE IV INJ 40 BIGIZNA.	PLOXURIDINE FOR INJ 0.5 GM	GENCILABINE HOLFOR INJ 1 6M	GENCITABINE HOLFOR IN J. 6M	GENCTABINE HOL FOR IN 1 GM	GEMOTTABINE HOL FOR INJ 200 MG
**************************************	DACOGEN	DACTINOMYCIN	MUNORUBICIN	MUNORUBICIN	DAUNOXOME	DOCEFREZ	DOCEFREZ	DOCETAXEL	DOCETAXEL.	DOCETAXEL	DOCETAXEL	DOCETAXEL	DOCETAXEL	DOCETAXEL	DOCETAXEL	DOXIL	DOXORUBICIN	DOXORUBICIN	DOXORUBICIN	DOXORUBICIN	ELIGARD	ELIGARD	LIGARD	ELIGARD	ı		NEVANIE		N N					ERWEDGE	PWINAZE	ETOPOPHOS	ETOPOSIDE .	ETOPOSIDE	FASLODEX	TRMAGON	FIRMAGON	FLOXURIDINE	FLUDARA	FLUDARABINE	FLUDARABINE	PLUOROURAGI.	FLUOROURAGE ST 1000110406	E LIDBOLIBACII	FLUOROURACIL	LUDROURACIL	FLUDROURACIL	-LUOROURACIL	FOLOTIN	FOLOTYN	FUDR	GEMCITABINE	GENCITABINE	SEMZAR	GEMZAR
		Antineopinstic Agent	Antineopiastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent			Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineoptastic Agent	Antineoplastic Agent	Antineoplastic Agent		Antineoplastic Agent			Antineoplastic Agent			Antineoplastic Agent				Antireophistic Agent						Antineophatic Agent .						Antineoplastic Agent					Antineoplastic Agent	Antineoptastic Agent	Antheoplastic Agets	Anthropiastic Agent	Authoritatic Agent	Authoritaelli Autol	Antineopiastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent			Antineoplastic Agent			
	DACOGEN INJ SOMG	DACTINOMYCEN IN LUMB	DAUNORUBICIN INJ 20MG	DAUNORUBICIN INJ SMOMIL	DAUNOXOME IN ZMGMI	DOCEFREZ INJ 20MG		DOCETAXEL INJ 160/16ML	DOCETAXEL INJ 160/8ML	DOCETAXEL INJ 2010.5ML	DOCETAXEL INJ 20MG/ZML	DOCETANDEL INJ ZOMG/MIL	DOCETAXEL INJ SOMG/ZML	DOCETAXEL INJ SOMGAML	DOCETAXEL INJ COMG/BML	DOXIL INJ 2MG/ML	DOXORUBICIN INJ 10MG	DOXORUBICIN IN 200MG	DOXORUBICIN INJUNEMIL	mI	ELIGARD IN 22.5MG	ELIGARD INJ 30MG		- 1	- 1	FLOXALIN INJUNIO	CLOXATIN INTERNA		EPIRUBEIN IN ZOOMG	EPIRUBECIN IN SOZSMI	EPIRUBICIN IN SOME	ERBITUX INJ 100MG	ERBITUX IN 200MG		ERWINAZE INJ 10000UNT	ETOPOPHOS INJ 100MG		ETOPOSIDE POW	FASLODEX INJ 250MG	FIRMAGON INJ 120MG	FIRMAGON INJ 80MG	싎	FLUDARA INJ 50MG	PLUDARABINE INJ 60NG	FEUDARABINE IN SOMBIZMI	FLUCKOURACE INJ TEMZUM	FLOOROUPACE, IN 2282508	ELCONOCIDE IN SOUGH	FLIOROLIPACIL IN J. SCANTOOM	FLIDROUBACE POW	FLUOROURACIL POW USP	FLUOROURACIL POW USPAN	FOLOTYN INJ ZOMGRAL	إ₹	FUDR INJ 0.5GM	GENCITABINE IN 16M	GENCITABINE IN ZOME		GENZAR INJ 200MG

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		\$0.00	\$0.00	\$0.00	\$0,00	\$0.00	\$0.00	\$0.00	20,00	00.02	89.00	\$0.00	\$9.00	20.00	50.00	20.00	20.00	00.00	amas.	20,00	\$0.00	\$6.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0,00	\$0.00	\$0.00	20,00	\$0.00	\$0.00	\$0.08	\$0.0\$	\$0.03	\$0.00	80,03	\$0.00	20.00	\$0.00	\$0.00	\$0.00	50.60	\$0.00	\$0.00	\$0.00	20,00	\$0.00	\$0.00	20.00	240.00	20.00	30.00	30.00	200	200	200	200	900	\$0.00	\$0.00	\$0.00	
Specific to the Section of the Secti		-24.0%	-24.0%	-15.0%	-15,0%	-14.0%	-14.0%	-14.0%	-15.0%	-15.0%	-13,5%	-13.5%	-13.5%	-18.0%	18.00	4/0:01-	KA 07	KCO!	-13.0%	-18.0%	-17.0%	-(7.0%	-17.0%	-18.0%	-13.5%	-13,5%	-82.0%	-82.0%	-62.0%	-14.0%	-16.5%	-18.5%	-15.0%	-16.0%	-16,0%	-16.0%	-16,0%	-17.0%	-13.0%	-13.0%	-18,0%	-13,5%	-35.0%	-15.0%	-13,5%	-18.0%	-18.0%	-14,0%	-14,0%	-14.0%	-14.0%	-14.0%	-14.0%	-14,0%	240.95	14.0%	-18.0%	75056	45.000	-44 046	-13.5%	-13.5%	-18.0%	
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		IMATINIB MESYLATE TAB 100 MG (BASE EQUIVALENT)	IMATINIB MESYLATE TAB 400 MG (BASE EQUIVALENT)	ERIBULIN MESYLATE IN J 1 MG/ZML (0.5 MG/ML)	TRASTUZUMAB FOR IV SOLN 440 MG	TOPOTECAN HCL CAP 0.25 MG (BASE EQUIV)	TOPOTECAN HCL CAP 1 MG (BASE EQUIV)	TOPOTECAN HOL FOR INJ 4 MG	PONATINIB HOL TAB 15 MG (BASE EQUIV)	PONATINIB HCL TAB 45 MG (BASE EQLAV)	IDARUBICIN HCL IV INJ 16 MG/19ML (1 NG/ML)	IDARLIBICIN HOLIVINO MOZOWI (1 MG/ML)	JOAN DECK HOLD MORAL (1 MORAL)	IN A PRINCIPALITY OF THE PARTY	DATE PRODUCTION TO THE POST OF HOUSE	ILLARCHICITY TO LEVEL 20 MOSCOMIC (1 MOSIMIC)	-1.	31		IFOSFAMIDE FOR INJ 1 GM		IFOSFAMIDE FOR INJ 3 GM	JEOSFAMIDE IV INJ 3 GRUGGRIL (50 MIGML)	FOSFAMIDE & MESNA NJ KIT 1000-1000 MG	AXITINIB TAB 1 MG	AXITINIB TAB 5 MG	IRINOTECAN HOL INJ 100 MG/SML (20 MG/ML)	IRINOTECAN HCL IN 140 MG/ZML (20 MG/ML)	MINOTECAN HCL INJ 500 MSZSAM, (20 MGAN)	ROMIDEPSIN FOR IV IN 10 MG	IXABEPILONE FOR IV INFUSION 15 MG	IXABEPILONE FOR IV INFUSION 45 MG	RUXOLITINIS PHOSPHATE TAB 10 MG (BASE EQUIVALENT)	RUXOLITINIB PHOSPHATE TAB 15 MG (BASE EQUIVALENT)	RUXOLITINIB PHOSPHATE TAB 20 MG (BASE EQUIVALENT)	RUXOLITINIB PHOSPHATE TAB 25 MG (BASE EQUIVALENT)	RUXOLITINIB PHOSPHATE TAB 5 MG (BASE EQUIVALENT)	CABAZITAXEL IN 50 MG/1.5ML (FOR IV INFUSION)	ADO-TRASTLIZUMAB EMTANSINE FOR IV SOUN 100 MG	ADO-TRASTUZUMAB EMTANSINE FOR IV SOLN 160 MG	CARFILZOMIB FOR INJED MG	GRANISETRON YCL INJ 4 MG/4ML (1 MG/ML)	LEUPROLIDE ACETATE INJ KIT 5 MGML	LEUPROLIDE ACETATE (BULK) POWDER	CLADPABINE IN 1 MG/ML	DOXORUBICIN HCL LIPOSOMAL INJ (FOR IV INFUSION) Z MG/ML	DOXORUBICIN HOL LIPOSOMAL INJ (FOR IV INFUSION) 2 MGML.	LEUPROLIDE ACETATE FOR INJ PEDIATRIC KIT 11.25 MG	LEUPROLIDE ACETATE FOR INJ PEDIATRIC KIT 15 MG	LEUPROLIDE ACETATE (3 MONTH) FOR INJ PEDIATRIC KIT 30 MG	LEUPROLIDE ACETATE FOR INJ PEDIATRIC KIT 7,5 MG	LEUPROLIDE ACETATE (3 MONTH) FOR INJ KIT 11.25 MG	LEUPROUDE ACETATE (3 MONTH) FOR INJ KIT 22.5 MG	LEUPROLIDE ACETATE FOR INJKIT 3,75 MG	LEUPROLIDE ACETATE (4 MONTH) FOR INJ KIT 39 MG	LEUPROLIDE ACETATE (6 MONTH) FOR INJ KIT 45 MG	LEUPROLIDE ACETATE FOR IN KIT 7.5 MG	7	_	Т	BOLK CAEMICALS - POWDER"	MESNA INJ 100 MSIM.	MESNA TAB 400 MG	MITCMYCIN FOR INJ ZU MG
		S EEVEC	GLEEVEC	HALAVEN	HERCEPTIN	HYCAMTIN	HYCAMTIN	HYCAMTIN	CLUSIG	CLUSIG	IDAMYCIN PES	IDAMVCIN PES	IDAMYCIN DES	DADITEIN	Proposition and the second	DAKUBKIN	DARUBICIN	ΤĒΧ	颐	FOSFAMIDE	IFOSFAMIDE .	FOSFAMIDE	FOSFAMIDE	FOSFAMIDE	INLYTA	INLYTA	IRINOTECAN	RINOTECAN	RINOTECAN	STODAX	IXEMPRA KIT	KEMPRA KIT	JAKAFI	JAKAFT	JAKAFI	JAKAFI	JAKAFI	JEVTANA	KADCYLA	KADCYLA	KYPROLIS	KYTRIL	LEUPROLIDE	BOLORA	LEUSTATIN	UPODOX	UPODOX 50	LUPR DEP-PED		WPR DEP-PED	LUPR DEP-PED			LUPRON DEPOT	EUPRON DEPOT	LIPRON DEPOT	LUPRON DEPOT	MELPHALAN	MERCAPTOPUR	MERCAPTOPUR	MERCAPTOPUR	MESNEX	MESNEX	MITOMYCEN
		c Agent			1	Antineoplastic Agent	Antineoplastic Agent		Antineoplastic Agent										Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent	Antinecotastic Agent	Astingonistic Agent				Anthony September 1	Antineoplastic Agent				Antheoplastic Agent	Antineoplastic Agent	Antinaoplastic Agent							Antineoplastic Agent	Antineoplastic Agent								Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent					Antineoplastic Agent	Antineoplastic Agent	'DR Antineoplastic Agent	Animeoplastic Ageni	Antineoplastic Agent	Antineoplastic Agent	Antineoplastic Agent
		TAB 10	CE FEVEC TAB 400MG	HAI AVEN INU 1MGIZME	HERCEPTIN INJ 440MG	HYCAMTEN CAP 0.25MG	HYCMITIN CAP 1MG	HYCAMTIN INJAMG	ICLUSIG TAB 15MG	ICHISIG TABASMG	12	The same of the same of	ILAM TOTAL PAR LINE CASE OF THE PART LINE CO.	ILIAM TOTAL PES IN SMISSORIE	IDARUBICIN IN TOTOM	IDARUBICIN IN ZOZOMI.	IDARUBICIN INJ SMG/SML	IFEX INJ 1GM	FEX INJ3GM	IFOSFAMIDE IN 1GM	IFOSFAMIDE INJ 16MZDML	IFOSFAMIDE IN 39M	POSEAMIDE IN 3CAMBOML	IEOSEAMIDE KIT MESNA	INI VTA TARI (MG	1	INCLES TO CHE	INTROJECTIVE INTRODUCE	INTO TECHNICAL IN CONTROL	MINDLE WITHOUT	INCREDE ATT IN 14500	INEMPRANT INJASING	JAKAFI TAB 10MG	l	ı	1	1	JEVTANA INJ 60M SML	KADCYLA INJ 100MG	KADCYLA INJ 160MG		KYTRIL INJ 1MG/ML	12	I ELIDBOTIDE DOW ACETATE	CINCIPALIN IN THE SAME	INDOOR INI SMEARS	LIPODOX 50 INJ ZMG/ML	LUPR DEP.PED INJ 11.25MG	LUPR DEP PED INJ 15MG	LUPR DEP-PED INJ 30MG	LUPR DEP.PED INJ 7.5MG	LUPRON DEPOT INJ 11,25MG	LUPRON DEPOT IN J 22,5MG	LUPRON DEPOT INJ 3,75MG	LUPRON DEPOT INJ 30MG	LUPRON DEPOT INJ 45MG	LUPRON DEPOT INJ 7.5MG	MELPHALAN INJ 60MG	MERCAPTOPURI POW	MERCAPTOPURI POW MONOHYDR Antineophestic Agen	MERCAPTOPURI POW USP	MESNEX INJ 1GM	MESNEX TAB 400MG	MITOMYCIN INJ 20MG

Antineoplastic Agent MITOMYCIN		1 4 4 5 4 5 4 5 5 5 5 5 5 5 5 5 5 5 5 5		2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2			
Manuality Ant	Antineoplastic Agent	MITOMYCIN	MITCMYCIN FOR INJ 40 MS	21200050002120	-18,0%	\$0.00	
Administration Agent MITTATION MITTA	Antineoplastic Agent	MITOMYCIN	MITOMYCIN FOR INJSMG	21200050002105	-18.0%	\$0.00	\ \
Adjustable April Adjustable	Antineoplastic Agent	MITOMYCIN	MITOMYCIN (BULK) POWDER	96585860452900	-15.0%	\$0.00	Z
Management	Antineoplastic Agent	MITOMYCIN	MITOMYCIN (BULK) POWDER	96665860452900	-15.0%	\$0.00	z
Autocopies April Autocopies Autocopies	Antineoplastic Agent	MITOMYCING	MITOMYCIN (BULK) POWDER	96665850452900	-15,0%	\$0.0\$	Z
Accessive Age	Antineoplastic Agent	MITOXANTRON	MITOXANTRONE HCL INJ CONC 20 MG/10ML (2 MG/NL)	21200055001320	-18.0%	\$0.00	¥
Addressidate parts Marcia Marci Marcia Marcia Marcia Marcia Marcia Marcia Marcia	Artineoplastic Agent	MUSTARGEN	MECHLORETHAMINE HOL FOR INJ 10 MG	21101030102105	-13,5%	\$0.00	,
Addressing Action Addr	T	NAVEL BINE	WINDRELBINE TARTRATE IN 10 MGMI.	21500050802020	-26.0%	\$0.00	>
Accession of All		NAVELBINE	VINDRELEINE JARIKAIE IN 30 MGSML (10 MG/ML)	21500050802025	-26.0%	\$0.60	٠,
Administration of Americans Controller Service Processories Control Administration of Americans Control Processories Control Administration of Americans Control Control Control Administration of Americans Control <td>T</td> <td>NEWSON</td> <td>SONATENIS IOSILAIE IAS ZU MG (BASE EQUIVALENI)</td> <td>12533050404320</td> <td>-13.0%</td> <td>20.00</td> <td>λ.</td>	T	NEWSON	SONATENIS IOSILAIE IAS ZU MG (BASE EQUIVALENI)	12533050404320	-13.0%	20.00	λ.
Adiabachille, April CONTROLLE, ATTAN DESIGNATION CONTROLLE, ATTANDAR DESIGNATION CONTROLLE, ATTANDAR DESIGNATION CONTROLLE, ATTANDAR DESIGNATION CONTR		ONCASOOD	DECLACIONAL CONTROL INCOME.	02120004002120	-18.5%	20:00	>
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Accordance of the control of	T	NII WATER	CARLIFICATION IN SOCIAL TO INSTACTORIS	THIOTOGOGGGGG	-53.03e	90.03	إ ح
Molecopietic John Microbial Agenti Profit Delay (Street Medical Agenti Profit D		CAALPOOL	CAALIFICATION FOR IV INJ 30 MG	UZI ZGRGGGGGG	-36.0%	00'05	>
Authoristic Act POLYLINGER POLYLINGER		PACLIFACE.	PACELIANEL IV CONC 100 MG/16./ML (5 MG/ML)	225000120001335	-18.0%	\$0.00	X
Administration of the Control of t		PACELIAVEL	PACE TACK BY COME TO MICE SHEET OF CHANGE	21300012001340	-18.0%	\$0.00	٨
AMERICADER IN POLITIONER IN POLITIO		PACUTAXEL	PACLITAXEL IV CONC 300 MISSIBIL (8 MISMIL)	21500012001350	-18.0%	\$0.00	Υ.
Aminteoplatic Apetal PERLITY PERLITY <td>PACLITAXEL IN 30MG/SML Anlineoplastic Agent</td> <td>PACLITAXEL</td> <td>PACLITAXEL IV CONC 30 MG/SML (6 MG/ML)</td> <td>21500012001325</td> <td>-18.0%</td> <td>\$0.00</td> <td>λ</td>	PACLITAXEL IN 30MG/SML Anlineoplastic Agent	PACLITAXEL	PACLITAXEL IV CONC 30 MG/SML (6 MG/ML)	21500012001325	-18.0%	\$0.00	λ
Andmissionale Agent PERTITY PERTITY <td>Antineoplastic Agent</td> <td>PENTOSTATIN</td> <td>PENTOSTATIN FOR INJ 10 MG</td> <td>21700045002120</td> <td>-18,0%</td> <td>\$0.00</td> <td>Υ</td>	Antineoplastic Agent	PENTOSTATIN	PENTOSTATIN FOR INJ 10 MG	21700045002120	-18,0%	\$0.00	Υ
Интервете уден РОМИЛОТИТЕ ДО-1 100 2170001001(14 -12.55 Интервете уден РОМИЛОТИТЕ ДО-1 100 2170001001(14 -12.55 Интервете уден РОМИЛОТИТЕ ДО-1 100 2170001001(12 -12.55 Интервете уден РОМИЛОТИТЕ ДО-1 100 21700000001(22 -12.55 Интервете уден РОМИЛОБИТЕ ДО-1 100 217000000001(22 -12.55 Интервете уден РОМИЛОБИТЕ ДО-1 100 21700000001(22 -12.55 Интервете уден РОМИЛОБИТЕ ДО-1 100 21700000001(22 -12.55 Интервете уден РОМИЛОБИТЕ ДО-1 100 217000000001(22 -12.55 Интерве	Antineoplastic Agent	PERJETA	PERTUZUMAB SOLN FOR IV INFUSION 420 MG/14ML (30 MG/ML)	21353054002020	-12.5%	\$0.00	*
Attlimengaties Agent FOUNDALYST FOUNDALYST FOUNDALYST FOUNDALYST FOUNDALYST CHARLEDGRIDE CAP 1 NG ZESTSTERSBOODTIO -12.55 Affiniergaties Agent FOUNDALYST FOUNDALYST FOUNDALYST FOUNDALYST TOWALLOGUEIC CAP 2 NG ZESTSTERSBOODTIO -12.55 Affiniergaties Agent FOUNDALYST FOUNDALYST FOUNDALYST FOUNDALYST ACTIVE -12.55 Affiniergaties Agent FOUNDALYST FOUNDALYST FOUNDALYST FOUNDALYST -12.55 Affiniergaties Agent FOUNDALYST FOUNDALYST FOUNDALYST -12.55 -12.55 Affiniergaties Agent FOUNDALYST FOUNDALYST FOUNDALS -12.55 -12.55 Affiniergaties Agent FOUNDALYST FOUNDALS FOUNDALS -12.55 -12.55 Affiniergaties Agent FOUNDALS FOUNDALS FOUNDALS -12.55 -12.55 Affiniergaties Agent FOUNDALS FOUNDALS FOUNDALS -12.55 -12.55 Affiniergaties Agent FOUNDALS FOUNDALS FOUNDALS -12.55 <td>Antineoplastic Agent</td> <td>PHOTOFRIN</td> <td>PORFIMER SODIUM FOR INJ 75 MG</td> <td>21707070102140</td> <td>-13,5%</td> <td>\$0.00</td> <td>z</td>	Antineoplastic Agent	PHOTOFRIN	PORFIMER SODIUM FOR INJ 75 MG	21707070102140	-13,5%	\$0.00	z
Mathemaphities Agent	Antineoplastic Agent	İ.	POMALIDOMIDE CAP 1 MG	21450080000110	-13.0%	\$0.00	χ.
Administration Agent DOMALIVET POMALIVET POMALIDATION C. A. L. M.	Antineoplastic Agent		POMALIDOMIDE CAP 3 MG	21450080000120	-13.0%	\$0.00	×
Aditionsplate Agent POWALVER POWALVER POWALVER CATORISON OF THE AGENT ADDRESS ENGINE CAR A MAN AND AGENT AS A MAN AND AGENT AGENT ADDRESS AND AGENT AGENT ADDRESS AND AGENT A	Antineoplastic Agent		POMALIDOMIDE CAP 3 MG	21450080000120	-13.0%	\$0.00	
Adminishabilite Agent PROLE BENN ADMINISTREE CARD IN MICHIGATINE CARD IN SIGNATOROUND UNIT STREAMMENT CONTRIBUTION CONTRIBUTIO	Antineonizatie Agent		POMAI IDOMINE CAPANG	2145RRBDD0125	-13 090	69.00	
Authorografie Aget REAL MAD ENAL/COURTE COF 18 MG SERSELEGOOTTO C-12.097	Antinophistic Appoi	ı	A DESI FIXEN FOR M SOL M SAMMAN LINE	24703824200	140.00	20.00	- >
Antimospiale Agent RESOLUTION LEMALICORREE CARE 13 NO. RESOLUTION C-12.0F Antimospiale Agent RESOLUTION LEMALICORREE CARE 13 NO. RESOLUTIONEE CARE 13 NO. RESOLUTION CARE CARE 23 NO. RESOLUTION CARE 23 NO.	And and and and		CONTROLLED ON AS NO.	000000000000000000000000000000000000000	700.00	0000	-,
	Animacopus Agom	l		Deleganneces:	#0.61	AUC.DA	4
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Aditinoplatis Agent REAL ANION LEAVALIDORATIC CAP & SAME SESSAGISSORITION -13.54A Aditinoplatis Agent RITLAVAN RITLAV	Antineopiastic Agent	1	LENALISCANDE CAPS ZO MG	9535405000110	-13.0%	20.00	+
Andmooplatic Agent FEVAMID CLEANADORICE CAPE AND CASE OF CASE	Antineoplastic Agent		LENALIDOMIDE CAP 25 MG	9539405000150	-13.0%	\$0.00	>-
Authoopiatic Agent	Antineoplastic Agent	REVLWID	LENALIDOMIDE CAP 5 MG	99394050000120	-13.0%	80,08	٠
Antimeoplastic Agent PREVICE: DASANTHIST FOR IN LOUGE 10 MSAMI. ZISSAGDD00020 -14.5% Antimeoplastic Agent SPEYCEE: DASANTHIST FOR 140 MG ZISSAGDD00020 -14.5% Antimeoplastic Agent SPEYCEE: DASANTHIST FOR 140 MG ZISSAGDD00020 -14.5% Antimeoplastic Agent SPEYCEE: DASANTHIST FOR 140 MG ZISSAGD000020 -14.5% Antimeoplastic Agent SPEYCEE: DASANTHIST ARE 70 MG ZISSAGD0001250 -14.5% Antimeoplastic Agent SPEYCEE: DASANTHIST ARE 70 MG ZISSAGD0001250 -14.5% Antimeoplastic Agent SPEYCEE: DASANTHIST ARE 70 MG ZISSAGD0001250 -15.5% Antimeoplastic Agent SINTAMING MALTIE CAP 72 MG ZISSAGD0001250 -15.5% Antimeoplastic Agent SINTAMING MALTIE CAP 72 MG ZISSAGD0001250 -15.5% Antimeoplastic Agent SINTAMING MALTIE CAP 72 MG ZISSAGD0001250 -15.5% Antimeoplastic Agent SINTAMING MALTIE CAP 72 MG ZISSAGD0001250 -15.5% Antimeoplastic Agent SINTAMING MALTIE CAP 72 MG ZISSAGD0001250 -15.5%	Antineoptastic Agent	RITUXAN	RITUXIMAB FOR IV INJ CONC 10 MG/ML	21353060001310	-18.5%	20,00	٨
Antineoptatic Agent SPENCEZ. DASATINEE TAG DASA 2158/02/00/00/20 -14.5% Antineoptatic Agent SPENCEZ. DASATINEE TAG DASA 2158/02/00/20 -14.5% Antineoptatic Agent SPENCEZ. DASATINEE TAG DASA 2158/02/20/20 -14.5% Antineoptatic Agent SPENCEZ. DASATINEE TAG DASA 2158/02/20/20 -14.5% Antineoptatic Agent SPENCEZ. DASATINEE TAG DASA 2158/02/20/20/20 -14.5% Antineoptatic Agent SPENCEZ. DASATINEE TAG DASA 2158/02/20/20/20 -14.5% Antineoptatic Agent SPENCEZ. DASATINEE TAG DASA -14.5% -14.5% Antineoptatic Agent SUPPREDINIA HISTORIA MACEE/TE CAPPORTALINO 2158/02/20/20/20 -14.5% Antineoptatic Agent SUPPREDINIA HISTORIA MACEE/TE CAPPORTALINO 2158/02/20/20/20 -16.5% Antineoptatic Agent SUPPREDINIA HISTORIA MACEE/TE CAPPORTALINO 2158/02/20/20/20 -16.5% Antineoptatic Agent SUPPREDINA PARCENA BALLANDER DASA -16.5% Antineoptatic Agent TAGCENA BELOTINE T	Antineoplastic Agent	RATUXAN	RITUXIMAB FOR IV INJ CONC 10 MGMI.	21353060001310	-16.5%	00:0\$	٨
Andineoplatio Agent SPRYCEE DASATINIB TAB 64 MAG 2155402000020 -14.5% Andineoplatio Agent SPRYCEE DASATINIB TAG 20 MG 2155402000020 -14.5% Andineoplatio Agent SPRYCEE DASATINIB TAG 20 MG 2155402000020 -14.5% Andineoplatio Agent SPRYCEE DASATINIB TAG 20 MG 2155402000020 -14.5% Andineoplatio Agent STRYCEE DASATINIB TAG 20 MG 2155402000020 -14.5% Andineoplatic Agent STRYMER Andineoplatic Agent 2155402000020 -14.5% Andineoplatic Agent STRYMER Andineoplatic Agent 2155402000020 -15.5% Andineoplatic Agent STRYMER Andineoplatic Agent STRYMER Andineoplatic Agent 2155402000020 -15.5% Andineoplatic Agent STRYMER ANDIANT GAP 20 MG ANDIANT GAP 20 MG 21554070000020 -15.5% Andineoplatic Agent TAGCENA RECOPACE AGENT ALLAND 215540700000020 -15.5% Andineoplatic Agent TAGCENA ALLAND ONCERNALE GAP NA GARD CAP AGENT ALLAND	Antheoplastic Agent	SPRYCEL	DASATINIB TAB 100 MG	21534020000350	-14.5%	\$0.00	\
Voltacoplatic Agent SPRYCEL DASTINUE INE 20 No.	Antinecolastic Acest	SPACE	DASATINIB TAR 140 MG	2153402000340	14 SP.	50.05	
Anilhacoplastic Agent SPRYCEL DASANTHIST TARS 30 MG CASSALGADOGGS -14.5% Anilhacoplastic Agent SPRYCEL DASANTHIST TARS 10 MG 21534020000054 -14.5% Anilhacoplastic Agent STRYCEL DASANTHIST TARS 10 MG 21534020000055 -14.5% Anilhacoplastic Agent STRYCEL DASANTHIST TARS 10 MG 21534020000056 -14.5% Anilhacoplastic Agent SURPRELINIA HEIGH TARS 10 MG 21534020000056 -15.5% Anilhacoplastic Agent SUNTRIBIO SUNTRIBIO ANILHACOP 25 MG RASE ECAUPALENT) 21534020000056 -15.5% Anilhacoplastic Agent SUNTRIBO SUNTRIBO ANILHACOP 25 MG RASE ECAUPALENT) 21534020000050 -15.5% Anilhacoplastic Agent SUNTRIBO SUNTRIBO ANILHACOP 25 MG RASE ECAUPALENT) 21534020000050 -15.5% Anilhacoplastic Agent TARCENA RELCTINIB TAB 100 MG 21534020000050 -15.5% Anilhacoplastic Agent TARCENA RELCTINIB TAB 25 MG ANILHACOP 20 MG AGENT 215340200000050 -15.5% Anilhacoplastic Agent TARCENA RELCTINIB TAB 25 MG	Anticopianic Appril	THE STATE OF THE S	DASSETTING TAR ON INC.	DESTRUCTION OF THE PROPERTY OF	-14 GK	40.00	->
Antineoplasite Agent STRYCEL LOWATINEST TAB 370 14.55% 14.55% Antineoplasite Agent SERVICEL DASATINEST TAB 370 MA 21534020000155 -14.55% Antineoplasite Agent STIPARGA REGORDE-FUB TAB 40 MA 2154020000155 -14.55% Antineoplasite Agent STIPARGA REGORDE-FUB TAB 40 MA 2154020000155 -15.55% Antineoplasite Agent SULPRIA SINTENT SIANITINIB MALATE CAP 25 MG 604SE EGUIVALENT) 21553070300102 -16.55% Antineoplasite Agent SULPRIA SIANITINIB MALATE CAP 25 MG 604SE EGUIVALENT) 2155307030100 -16.55% Antineoplasite Agent SULPRIA SIANITINIB MALATE CAP 25 MG 604SE EGUIVALENT) 2155307030100 -16.55% Antineoplasite Agent TARCENA BRLOTINIB TAB 100 MG 2170040102120 -16.55% Antineoplasite Agent TARCENA BRLOTINIB TAB 100 MG 217005001020 -16.55% Antineoplasite Agent TARCENA BRLOTINIB TAB 100 MG 2170000001120 -16.55% Antineoplasite Agent TARCENA BRLOTINIB TAB 100 MG 21700000001120 -16.55%	Anticopiasiic Marii	STATISTIC		10000000000000000000000000000000000000	POPE .	2000	- ;
Antineoplastic Agant STRYCEL. DASATINB TAB 20 MAG CISSAGGGOOGTS -14.55% Antineoplastic Agant STRYCEL. DASATINB TAB 20 MAG ZISSAGGGOOGTS -14.55% Antineoplastic Agant STRYCEL. RECORAFENB TAB 40 MAG ZISSAGGGOOGTS -15.55% Antineoplastic Agant SURPTENIA SURPTINIB MALNE CAP 125 MG RASE EQUIVALENT) ZISSAGGGOOGTS -16.55% Antineoplastic Agant SURPTINIB MALNE CAP 125 MG RASE EQUIVALENT) ZISSAGGGOOGTS -16.55% Antineoplastic Agant SURPTINIB MALNE CAP 25 MG RASE EQUIVALENT) ZISSAGGGOOGTS -16.55% Antineoplastic Agant STRACES RECORNES RASE EQUIVALENT) ZISSAGGGOOGTS -16.55% Antineoplastic Agant TARCENA RELOTINIB TAB 100 MG RASE EQUIVALENT) ZISSAGGGOOGTS -16.55% Antineoplastic Agant TARCENA RELOTINIB TAB 100 MG ZISSAGGGOOGTS -16.55% Antineoplastic Agant TARCENA RELOTINIB TAB 100 MG ZISSAGGGOOGTS -16.55% Antineoplastic Agant TAXOTIERE DOCETAZEL FOR NI CONC 20 MGML ZISSAGGGOOGTS -16.55% Antineoplastic Agant	Antinooplastic Agent	SPATCEL	LASA IING JAB 20 MG	UPCUDAUCUPCCT2	-16.5%	30.00	¥
Anitheoplasite Agent STRYNCEL BASATHUBE TARE 80 MG 21543020000254 -14.554 Anitheoplasite Agent STRVARCA REGORGERINA FIRE 80 MG 20000045106459 -15.454 Anitheoplasite Agent SULPENT SILVIRIAN SILVIRIAN FILE AGENT -15.456 Anitheoplasite Agent SULPENT SILVIRIAND MALATE CAP 28 MG RASE EQUIVALENT) 2153020000120 -16.554 Anitheoplasite Agent SULPENT SILVIRIAND MALATE CAP 28 MG RASE EQUIVALENT) 2153020000120 -16.554 Anitheoplasite Agent SULPENT SILVIRIAND MALATE CAP 28 MG RASE EQUIVALENT) 215300000140 -16.554 Anitheoplasite Agent SULPENT SILVIRIAND MALATE CAP 28 MG RASE EQUIVALENT) 215300000140 -16.554 Anitheoplasite Agent TARCENA BILLOTINIB TAB 35 MG 2153000000140 -16.554 Anitheoplasite Agent TARCENA BILLOTINIB CAP 200 MG 2153000000105 -16.554 Anitheoplasite Agent TAXOTERE DOCETAXEL FOR NU CONC 20 MG/ML) 2150000000105 -15.054 Anitheoplasite Agent TAXOTERE DOCETAXEL FOR NU CONC 20 MG/ML 21104070000147	Antineoplastic Agent	SPRYCEL	DASATINIS TAR 70 MG	2153402000350	-14,5%	\$0.00	>-
Autheropisate Agent STIVARGA RECORA-FEBRA TAS AGENASE EXAUNALENTY 200000044064506 -15.554 Autheropisate Agent SULPENT ALLE CAP 125 MG (BASE EXUNALENTY) 21333073001020 -16.554 Autheropisate Agent SULPENT SUNITINAI MALATE CAP 25 MG (BASE EXUNALENTY) 21533073001030 -16.554 Autheropisate Agent SUNITINAI MALATE CAP 25 MG (BASE EXUNALENTY) 21533073001030 -16.554 Autheropisate Agent SUNITINAI MALATE CAP 25 MG (BASE EXUNALENTY) 21533073001030 -16.554 Autheropisate Agent SUNITINAI MALATE CAP 25 MG (BASE EXUNALENTY) 21534073001030 -16.554 Autheropisate Agent TARCENA ERUCINIBI TAB 35 MG 215340230000040 -16.554 Autheropisate Agent TARCENA ERUCINIBI TAB 35 MG 215340230000000000000000000000000000000000	Antineoplastic Agent	SPRYCEL	DASATINIB TAB 80 MG	21534020000354	-14.5%	\$0.00.	Υ
Antilaceopasite Agent SUPPREINTA HISTRELIN ACETATE (CPP) INPLANT BIT ACATATE (CPP) INPLANT BIT ACATATE (CPP) INPLANT (CAP 225 MG GASE EQUITALENT) 21533070300140 -16.5% Antilaceopasite Agent SUTENT SUNTENT SUTENT SUTENT<	Anthracplastic Agent	STIVARGA	REGORAFENIB TAB 40 MG	2153305000320	-13.5%	\$0.00	¥
Arithacopiastic Agent SUITENT SUITENT </td <td>Antipeconizatic Agent</td> <td>SUPPREMINIA</td> <td>N ACETATE (CPP)</td> <td>30080045106450</td> <td>-15.4%</td> <td>\$0.00</td> <td>></td>	Antipeconizatic Agent	SUPPREMINIA	N ACETATE (CPP)	30080045106450	-15.4%	\$0.00	>
Authreoplastic Agent SUITENT SUNTINIE MALATE CAP SS MG (RASE EQUITALIENT) ZISSA070300130 -18.5% Authreoplastic Agent SUITENT SUNTENT SUNTINIE MALATE CAP SS MG (RASE EQUITALIENT) ZISSA070300140 -16.5% Authreoplastic Agent STATOBOLIZAZIO -16.5% -16.5% Arthreoplastic Agent TARCEVA ERLOTINIB TAB 150 MG ZISSA025000360 -16.5% Arthreoplastic Agent TARCEVA ERLOTINIB TAB 150 MG ZISSA0000016 -16.5% Arthreoplastic Agent TARSICARA BILOTINIB TAB 150 MG ZISSA0000016 -16.5% Arthreoplastic Agent TARSICARA MILOTINIB CAP 160 MG ZISSA00000116 -15.0% Arthreoplastic Agent TARSICARA MILOTINIB CAP 160 MG ZISSO00000116 -15.0% Arthreoplastic Agent TARSICARA MILOTINIB CAP 160 MG ZISSO000000122 -15.0% Arthreoplastic Agent TARXOTIERE DOCETARE FOR IN J CONC 20 MG/ML ZISSO000000132 -15.0% Arthreoplastic Agent TEMODAR TEMODAR TEMODAR TEMODAR -11.00 MG/ML Arthreoplastic Agent	Antipopologic Apont	TNATIO	SHINTINIE MAI ATE CAP 12 5 MG PASE FOX IIVAI FINTI	21533070300120	-16.5%	\$0.05	*
Antineoplastic Agent STITERIT SIMITARIO SERVICIANTE FOR INJ. 3.5 MG ZITSONOSOGI 4G -(6.5% Antineoplastic Agent STREAT SIMITARIO SERVICIANTE FOR INJ. 3.5 MG ZITSONOSOGI 4G -(6.5% Antineoplastic Agent TARCENA ERLOTNIBI TAB 100 MG ZITSONOSOGI 4G -(6.5% Antineoplastic Agent TARCENA ERLOTNIBI TAB 100 MG ZITSONOSOGI 2G -(6.5% Antineoplastic Agent TARCENA INILOTINIB CAP 100 MG ZITSONOSOGI 2G -(6.5% Antineoplastic Agent TARCENA INILOTINIB CAP 100 MG ZITSONOSOGI 2G -(6.5% Antineoplastic Agent TARCENA INILOTINIB CAP 100 MG ZITSONOSOGI 2G -(6.5% Antineoplastic Agent TARCITERE DOCETAZEL FOR INJ CONC 20 MG/ML) ZITSONOSOGI 2G -(6.5% Antineoplastic Agent TAXOTERE DOCETAZEL FOR INJ CONC 20 MG/ML) ZITSONOSOGI 2G -(6.5% Antineoplastic Agent TAXOTERE DOCETAZEL FOR INJ CONC 20 MG/ML) ZITSONOSOGI 2G -(6.5% Antineoplastic Agent TEMODAR TEMOZOLOMIDE CAP 100 MG ZITSONOSOGI 2G -(7.0% Antineo	Andrew Control of the	SHOW	CINITINIO MAI ATE CAD 25 MG GASE EXITY ENT	O-Karanzahan dan	-4B 5%	\$	À
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Antihoplastic Agent TAXCENA ERLOTINIB TAB 100 Mis CATABORISED CATABORI	Primeoplastic Agent	STINDS		021201040102120	101 mg	0000	
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Aprilheoplastic Agent TASIGNA INLOTINUE CAP 160 MG 2155A450000115 -13.5% Aprilheoplastic Agent TASIGNAA INLOTINUE CAP 200 MG 2155A050000120 -13.5% Aprilheoplastic Agent TAXOFTERE DOCETAXEL FOR INJ CONC 20 MG/ML) 2150000500130 -15.0% Aprilheoplastic Agent TAXOFTERE DOCETAXEL FOR INJ CONC 20 MG/ML) 2150000500130 -15.0% Aprilheoplastic Agent TAXOFTERE DOCETAXEL FOR INJ CONC 20 MG/ML) 2150000500132 -15.0% Aprilheoplastic Agent TAXOFTERE DOCETAXEL FOR INJ CONC 20 MG/ML) 2150000500132 -15.0% Aprilheoplastic Agent TEMODAR TEMOZOLOMIDE CAP 140 MG 21104070000440 -17.0% Aprilheoplastic Agent TEMODAR TEMOZOLOMIDE CAP 140 MG 21104070000440 -17.0% Aprilheoplastic Agent TEMODAR TEMOZOLOMIDE CAP 100 MG 21104070000147 -17.0% Aprilheoplastic Agent TEMODAR TEMOZOLOMIDE CAP 250 MG 21104070000140 -17.0% Aprilheoplastic Agent TEMODAR TEMOZOLOMIDE CAP 250 MG 21104070000150 -17.0% Aprilheo	Antineoplastic Agent	TARCEVA	ERLOTINIB TAB 25 MG	21534025000320	-16.5%	50.05	¥
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Antineoplastic Agent TAXOTERE DOCETAXEL FOR NJ CONC 20 MG/ML) 21500065001320 -15.03% Antineoplastic Agent TAXOTERE DOCETAXEL FOR NJ CONC 20 MG/ML 21500005001310 -15.03% Antineoplastic Agent TAXOTERE DOCETAXEL FOR NJ CONC 20 MG/ML 21500005001326 -15.03% Antineoplastic Agent TEMODAR TEMOZOLOMIDE CAP 100 MG 21104070000740 -17.03% Antineoplastic Agent TEMODAR TEMOZOLOMIDE CAP 100 MG 21104070000740 -17.03% Antineoplastic Agent TEMODAR TEMOZOLOMIDE CAP 100 MG 21104070000140 -17.03% Antineoplastic Agent TEMODAR TEMOZOLOMIDE CAP 100 MG 21104070000140 -17.03% Antineoplastic Agent TEMODAR TEMOZOLOMIDE CAP 20 MG 21104070000150 -17.03% Antineoplastic Agent TEMOZOLOMIDE CAP 20 MG 21104070000150 -17.03% Antineoplastic Agent TEMOZOLOMIDE CAP 20 MG 21104070000150 -17.03%	Antineoplastic Agent	TASIGNA	INITION IN CAP 200 MG	21534060000120	-13.5%	\$9.00	χ.
Antineoplastic Agent TAXCOTERE DOCETAZEL FOR NA CONC 20 MG/ML 2150005501310 -15.05s Antineoplastic Agent TAXCOTERE DOCETAZEL FOR NA CONC 80 MG/ML Z1500005501325 -15.05s Antineoplastic Agent TAXCOTERE DOCETAZEL FOR NA CONC 80 MG/ML Z1104070600430 -17.05s Antineoplastic Agent TEMODAR TEMOZOLOMIDE CAP 140 MG Z1104070600440 -17.05s Antineoplastic Agent TEMODAR TEMOZOLOMIDE CAP 140 MG Z1104070600440 -17.05s Antineoplastic Agent TEMODAR TEMOZOLOMIDE CAP 120 MG Z1104070600147 -17.05s Antineoplastic Agent TEMODAR TEMOZOLOMIDE CAP 250 MG Z1104070600147 -17.05s Antineoplastic Agent TEMODAR TEMOZOLOMIDE CAP 250 MG Z1104070600150 -17.05s Antineoplastic Agent TEMODAR TEMOZOLOMIDE CAP 250 MG Z1104070600150 -17.05s	Antineoplastic Agent	TAXOTERE	DOCETAXEL FOR INJ CONC 20 MG/D.SML (40 MG/ML)	21500005001320	-15.0%	\$0.00	>
Artitheoplastic Agent TAXOTERE DOCETAXEL FOR NJ CONC 80 MGZML (20 MGM2) Z150005901325 -15.0% Artitheoplastic Agent TAXOTERE DOCETAXEL FOR INJ CONC 80 MGZML (20 MGM2) Z150005901315 -15.0% Artitheoplastic Agent TEMODAR TEMOZOLOMIDE CAP 140 MG Z1104070000440 -17.0% Artitheoplastic Agent TEMODAR TEMOZOLOMIDE CAP 160 MG Z1104070000140 -17.0% Artitheoplastic Agent TEMODAR TEMOZOLOMIDE CAP 160 MG Z1104070000120 -17.0% Artitheoplastic Agent TEMOZOLOMIDE CAP 250 MG Z1104070000120 -17.0% Artitheoplastic Agent TEMOZOLOMIDE CAP 250 MG Z1104070000150 -17.0% Artitheoplastic Agent TEMOZOLOMIDE CAP 35 MG Z1104070000150 -17.0%	Ī	TAXOTERE	DOCETAXEL FOR INJ CONC 20 MGML	21500005001310	-15.0%	\$0.00	٠
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Antineoplastic Agent TEMODAR TEMOZOLOMINE CAP 100 MG Z1104070000140 -17.0%	T	TAXOTERE	CONTESTANT TO IN LOOK SPINISHING TO MANAGE	2150005001315	-15.0%	\$0.00	>
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TEMODAR TEMOZOLOMIDE CAP 250 MG 21104G70400150 -17.0% TEMODAR TEMOZOLOMIDE CAP 5 MG 21104G70400110 -17.0%	Antineoplastic Agent	TEMODAR	TEMOZOLOMIDE CAP 20 MG	2110407000120	-17.0%	\$0.00	Y
TEMOZOLOMIDE CAP 5 MG 21104070000110 -17.0%	Antinecolectin Agent	TEMODAR	TEMOZOLOMIDE CAP 250 MG	21104070806150	-17.0%	\$0.00	À
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The second second	THERACYS	TICEBOO	TODOSAB	TOPOSAR	TOPOSAR	TOPOTECAN	TOPOTECAN	TORISE	TREANDA		TRES STAR DED		•	TRETINOIN	1	TYKERB	VANTAS	VECTIBIX	VECTIBIX	VELCADE	VDAZA.	VINBLASTINE	VANBLASTINE	WINCASAR PFS	VINCRISTINE	WINORELBINE	VINORELBINE	Ę			XALKORI			XXANDI	XWITHA SOLOF	YERVOY	YERVOY	ZALTRAP	ZALTRAP	ZANOSAR	ZELBORAF	ZEVALIN	ZEVALIN	ZOLADEX	ZOLADEX	ZOLINZA	ZYTIGA	١	STELARA				- 1	BARACLUDE	- 1	BARACLUDE	COMBIVIR	COMPLERA	CRIXIVAN	CRIXIVAN	THEORIGINA
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A Part of the Control	na I	TICE OF IN 1985	TODOSAB INI 400/5MI	TOPOSAR IN 16M/50M	TOPOSAR INJ 500/25ML	TOPOTECAN INJAMS	TOPOTECAN INJAME/AM	TORISEL SOLZSWGAL	TREANDA IN TOOMG	TREANDA IN JEWS	TREE STAR DEP IN 13 75MG	TOPE STAP A IN 14 25UR	TREI STAR MIX IN 1 22.5MG	TRETINOIN CAP 10MG	TRISENOX SOL 10MG/10M	TYNCERB TAB 250MG	VANTAS KIT SOMG	VECTIBIX INJ 100MS	VECTIBIX INJ 400MG	VELCADE INJ 3.5MG	VIDAZA INJ 100MG	VINBLASTINE INJ 10MG	VINBLASTINE INJ SMG/ML	WINCASAR PFS INJ IMGAM	VINCRISTINE INJ 1MG/ML	VINORELBINE INJ 10NG/ML	VINORELBINE INJ SOMG/SML	VOTRENT TAB 200MG	VUMON INJ SOMG/SME.	XALKOR! CAP 200MG	XALKOR! CAP 250MG	XELODA TAB 150MG	XELODA TAB 500MG	XTANDI CAP 40MG	XYNTHA SOLOF IN3 1000UNIT	YERVOY INJ 200MG	YERVOY INJ SOMG	ZALTRAP INJ 100/4ML	ZALTRAP INJ 20078ML	ZANOSAR INJ 1GM	9	ZEVALIN KIT IN-111	ZEVALIN KIT Y-90			ZOLINZA CAP 100MG	ZYTIGA TAB 250MG	STELARA IN345MGR0.5	STELARA INJ SOM GMIL		APTIVUS CAP 250MG	1	ATRIPLA TAB	BARACLUDE SOL.DSMG/ML	BARACLUDE TAB 0.5MG	BARACLUDE TAB IMG	COMBIVIR TAB 150-300	COMPLERA TAB	CREKIVAN CAP 200MG	CREXIVAN CAP 400MG	CONTRACTOR OF STREET

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	-40.0%	-40.0%	-40.0%	-10.0%	-7.0%	-7.0%	-5.0%	-5.0%	-5.0%	-5.0%	-5.0%	-13.0%	-15.6%	-15.5%	-12.0%	-12,0%	-12.0%	-12.5%	-12.5%	-11.5%	-11.5%	-11.5%	-11.0%	-11.0%	-11.0%	280.6%	-20.0%	-20.0%	-125%	-12.5%	265.0%	-95.0%	-5.0%	-5.0%	-5.0%	-13.0%	-13.0%	-13.0%	-13.0%	-13,0%	360°S-	3.0%	-3.0%	-3.0%	-3.0%	-3.0%	-13.0%	-13.0%	-13.0%	-13.0%	-13.5%	-13.5%	-38.0%	-38.0%	-38.0%	-38,0%	-38.0%	-15.5%	-10.0%	-10.0%	-10.0%	-14.5%
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1. 2. 4. 4. 4. 4. 4.	DIDANDSINE DELAYED RELEASE CAPSULE 200 MG	DIDANOSINE DELAYED RELEASE CAPSULE 250 MG	DIDANOSINE DELAYED RELEASE CAPSULE 400 MG	RIL PIVIRINE HOL TAB 25 MG (BASE EQUIVALENT)	EMTRICITABINE CAPS 200 MG	EMTRICITABINE SOLN 10 MG/ML	LAMINUDINE ORAL SOLN TO MG/ML	LAMINADINE TAB 150 MG	LAMINUDINE TAB 300 MG	LAMINUDINE ORAL SOLN 5 MG/ML	LAMINUDINE TAB 100 MG	ABACAVIR SULFATE LAMINUDINE TAB 500-300 MG	ENFUMERIDE FOR INJ 90 MG	ENFUVIRTIDE FOR INJ KIT 90 MG	ETRAVIRINE TAB 100 MG	ETRAVIRONE TAB 200 MG		SAQUINAVIR MESYLATE CAP 200 MG		RALTEGRAVIR POTASSIUM CHEW TAB 100 MG (BASE EQUIV)	RALTEGRAVIR POTASSIUM CHEW TAB 25 MG (BASE EQUIV)	RALTEGRAVIR POTASSILM TAB 400 MG (BASE EQUIV)	LOPINAVIR-RITONAVIR SOLN 480-100 MG/SML (80-20 MG/ML)	LOPINAVIR-RITONAVIR TAB 100-25 MG	I OPINAVIR-RITONAVIR TAB 200-50 MG	A STANDARD CONTRACTOR TAR 140-300 MG	I AMA IDINE TAR 150 MG	I AMA DINE TAR AM MC	EDSAMPRENAME CALCILIES SIND SO MICHIEL PASSE FOLIM	ENAVIR CALCILIM TAR 700 MG (BA)			RITONAVAR CAP 100 MG	(MICHAEL DE LOS INCOMPANIONS)	RITONAVIR TAB 100 MC	DARI MANG STHANDI ATETAR 150 MG MASE BOUND	DARUNAVAR ETHANOLATE TAB 400 MG (BASE EQUIV)	DARUNAVIR ETHANOLATE TAB 600 MG (BASE EQUIV)	DARUNAVIR ETHANOLATETAB 76 KG (BASE EQUIV)	DARUNAVIR ETHANOLATE TAB 800 MG (BASE EQUIV)	DELAVIRDINE MESYLATE TAB 100 MS	DELAVIRDINE MESYLATE TAB 200 MS	ZIDOVUDINE CAP 100 MG	ZIDDWIDINE IV SOLN 10 MG/ML	ZIDOVUDINE SYRUP 10 MG/ML	ZIDOWIDINE TAB 300 MG	ATAZANAVIR SULFATE CAP 100 MG (BASE EQUIV)	ATAZANAVIR SULFATE CAP 150 MG IBASE EQUIVI	ATAZANAVIR SUI FATE CAP 200 MG IBASE EQUIVI	ATAZANAVIR SI II FATE CAP 300 MG RASE FOUN	MARAVIROC TAB 150 MG	WARAVIROC TAR 300 MG	STAVIDINE CAP 15 MG	STAVIDINE CAP 20 MG	STAVUDINE CAP 30 MG	STAVUDINE CAP 40 MG	STAVUDINE FOR ORAL SOLN 1 MGML	ELVITEGRAV.COBICIS-EMTRICITAB-TENOFOV TAB 150-150-200-300 MG	EFAVIRENZ CAP 200 MG	EFAVIRENZ CAP 50 MG	ECAMPENY TAR AM MC	ABACAVIES IN EATER AMINITIONE, ZIDOM IDINE TAR 300-150-300-MG
	DIDANOSINE	DIDANOSINE	DIDANOSINE	EDURANT	EMTRIVA	EMTRIVA	EPIVIR	EPMR	EPIVIR	EPWIR HBV	EPIVIR HBV	EPZICOM	FUZEON	FUZEON	INTELENCE	INTELENCE	INTELENCE	INVIRASE	INVIRASE	SENTRESS	ISENTRESS	ISENTRESS	KALETRA	KALETRA	KAIETRA	AMAIDON	AMPAIDINE	I AMEN ININE	I EXNA	I EXTINA	ACAINO AONIC	NEVIRONE	NOTEVIE	NOBINE				PREZISTA		١	1	RESCRIPTOR		RETROVIR	RETROVIR	RETROVIR	REYATAZ	REYATAZ	BEYATAZ	DEVATAZ	SEI ZENTRY	CEI ZENTEY	STAVIDINE	STAVEIDINE	STAVUDINE	STAMIDINE	STAVIDINE	STRIBILD	SELISTIVA	SUSTIVA	STIETTVA	OAII SAS
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	DIDANOSINE CAP 200KG	DIDANGSINE CAP 250MG	DIDANOSINE CAP 400MG	EDURANT TAB 25MG	EMTRIVA CAP 200MG	EMTRIVA SOL TOMGRAL		EPIVIR TAB 150MG	EPIVIR TAB 300MG	EPIVIR HEW SOLEMGINL	EPIVIR HBV TAB 100MG	EPZICOM TAB	FUZEON INJ 90MG	FUZEON KIT	INTELENCE TAB 100MG	INTELENCE TAB 200MG	INTELENCE TAB 25MG	INVIRASE CAP 200MG	INVIRASE TAR SOUMS	CHURCH CHURCH	SCHOOL OF THE STATE	CHOCK THE CONTROL	ISENIALES IND TUMES	WALETON TED AND SENS	CALCIDA TAB 200 E014C	KALEIKA IAB ZUUSUMUS	CAMINUDIZING TAN 16040	LAMINODINE IAB TOURS	LAMINOUINE LAB SUUMO	CHANGE SUBSTANT	LEXIVA IAB /ddwls	NEVIRAPINE SUS SUMSTANL	NEVIKAPINE LAB ZWANG	MUNICIPAL CONTROLL		•	PREZION INDIANA	DEED'STA TAR ANDUG			-15-	DESCRIPTION TAN 200110	RESCRIPTION INSTRUMENT	MCINCVIN CAL MORE	CETTOONED CODE AND TOTAL		PETROVIK IND SUMMS	RETAINS CAN INOMO	KETALAZ CAP 150MG	REYATAZ CAP ZOUMG	REYAIAZ CAP 300MG	SELZENIKY IAB 130MG	SELZENTRY TAB 300MG	STAVUDINE CAP 15MG	STAVDUNE CAP ANNO		STAVULINE CAP ANTIC	≆l.	STRBILD TAB	- 1	SUSTIVA CAP SOMG	SUSTIVA TAB 600MG

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=	20.00	50 63	Special	80.00	80.08	20.00	900	300	30.00	Bin	\$0,00	20.03	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	20.00	OTTOR	200	\$0.00	89.93	\$0.00	\$0.00	80.08	80.03	\$0.00	300	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	20.00	20.08	\$0.00	\$0.00	80.08	8,8	B 8	200	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	40.00	\$0.00	
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The second secon	EMITRICITABINE-TENOFOVIR DISOPROXIL FUMARATE TAB 200-300 MG	DIDANOSINE FOR SOLN 2 GM	DIDANOSINE FOR SOLN 4 GM	DIDANDSINE DELAYED RELEASE CAPSULE 125 MG	DIDANOSINE DELAYED RELEASE CAPSULE 200 MG	DIDANOSINE DELAYED RELEASE CAPSULE 250 MG	DIDANOSINE DELAYED RELEASE CAPSULE 400 MG	INELFINAVIR MESYLATE TAB 250 MG	NELFINAVIR MESYLATE TAB 625 MG	NEVIRAPINE SUSP 50 MG/SML		NEWPAPINE TAR SPOUND AND LICE	201 EDDONIC ACTI IN COMO ESS PANIS ISTAN ASSESSED	TENTED AND DISCOOKE THE STATE OF STATE	TENDERARD DISCREDE TOWARD CONTRACTOR TO MORER 40 MORGIN	TENDED OF DISCOPROSE FUNDANCE TAB 150 MG	HENDEDIAR DISORDAY FINADATE TAB SEATO	TENDERAL DISCONDENSE EN MANAGER AND SALVE	STAVIDING CAP 15 MG	STAVIDANE CAP 20 MG	STAVUDINE CAP 30 MG	STAVUDNE CAP 40 MG	STANDDINE FOR ORAL SOLN 1 MG/MIL	ABACAVIR SULFATE SOLN 20 MGML (BASE EQUIV)	ABACAVIR SULFATE TAB 300 MG (BASE ECUIV)	ZIDOVUDINE CAP 100 MG	ZIDOVUDINE SYRUP 10 MG/ML	AZIDOVDEJINE TAB 300 MG	ARATACED FOR MISON MACING INCOME.	TOFACITIVE CITEATE TAR SIMS (BASS ECONOMIC ENTI-	ADALIMUMAB INJ KIT 20 MG/0, 4ML	ADALIMUMAB INJIGT 40 MGD.8ML (50 MG)ML)	ADALIMUMAB INJ KIT 40 MG/D, BML (50 MG/ML)	ADALINUMAB INJ KIT 40 MG/0,8ML (50 MG/ML)	ADALIMUMAB INJ KIT 40 MG/D.BML (60 MG/ML)	INFLIXIMAB FOR IV INJ 100 MG	TELADREMISTANCE		INTERFERON ALFACON-1 INJ 9 MCGROSML		PEGINTERFERON ALFA-28 FOR INJ KIT 120 MCG/0,5ML	PEGINTERFERON ALFA-2B FOR INJ KIT 150 MCG/0.5ML	PEGNI ENFERON ALFR-28 FOR INJINI 159 MCGRUSML DEGNITORERON ALEADED IN FRIT EN SCOR SUF	PEGINTERFERON ALFA-78 FOR IN KIT SO MOSA SMI	PEGINTERFERON ALFA-28 FOR INJ KIT 30 MCG/0.6ML	PEGINTERFERON ALFA-28 FOR INJ KIT 80 MCGRO, SML,					KICKAVIKIN I AB 600 MS	KIDAVIRIN I AB 400 MG	RIBAMRN TAB 200 MG	RIBAVIRIN TAB 400 MG	RIBAVIRIN TAB 600 MG	
Sui I - Section	TRUVADA	MDEX	MDEX	MDEXEC	WIDEXEC	MOEXEC	MDEXEC	MRACEPT	WRACEPT	VIRAMUNE	VIRAMUNE	VIRAMINE XR	ZOLEDRONIC	MEEAN	ì	WAREAD						Ĺ			ZIAGEN	ZIDOVUDINE	ZIDOWODINE	ZIDOVODINE	ORENCIA	XELLANZ	HUMIRA	HUMIRA	HUMBRA PEN	HUMIRAPEN	HUMIRA PEN	REMICADE	NOVEK	INFERGEN		z		PEG-INTRON	PEG-INTRON	PEC-INTRON	PEC-INTRON	PEG-INTRON	REBETOL	REBETOL	RIBAPAK	RIBAPAK	RIBAPAR	REASONERS	RIBASPHERE		ERE	
And local control Amand	Aniretrowisa Agent	Antiretroviral Agent	Antiretroviral Agent	Municipality Agent	Anneuconia Agent	Amianowial Agent	Antiretrovital Agent.	Artiretroviral Agent	Anthetrovital Agent	Antiretroviral Agent	Antifetroviral Agent	Anthetroviral Agent	Antiretroviral Agant	Anthetroviral Agent	Antiphowing Agent	Animittonini Ageni	Antretroviral Agent	Antiretrovinsi Agent	Antiretroviral Agent	Antiretroviral Agent	Antiretrovinal Agent	Antiretroviral Agent	Antiretroviral Agent	Antiretroviral Agent	Antiretrovkal Agent	Antreiroviral Agent	Authorities Agent	Authornalic	Antimeamatic	Antithermelic	Antimesumatic Agent	Antimeumatic Agent	Antimeumatic Agent	Antithoumatic Agent	Antimetralic Agent	Anthreumanic Agent	Antiviral Agent	Anthiral Agent	Antiviral Agent	Antiviral Agent	Antiviral Agent	Aminima Agent	Antiviral Agent	Antiviral Agent	Antiviral Agent	Antiviral Agent	Antiviral Agent	Antivinal Agent	Antiviral Agent	Andvari Agent	Anthrie Anan	Antiviral Agent	Antiviral Agent	Antiviral Agent	Antiviral Agent	Robbits Annual
And TAB	RUMANA IAB	SULZOM	"1	WINEY COT IZEMS	1	- 1	- [VIRACEPT TAB 625MG	VIRAMUNE SUS SOMG/SME.	VIRAMUNE TAB 200MG	MUNE XR TAB	MUNE XR TAB 400MG	AD POW 40MG/GM	AD TAB 150MG	VIREAD TAB 200MG	VIREAD TAB 250MG	1	ERIT CAP 15MG	SRIT CAP ZUMG	ERIT CAP 30MG			IAGEN SOL 20MG/ML	AGEN TAB 300MG	IDOVUDINE CAP 100MG	ZIDOM BANE TAR ADMIC	ORENCIA INJ 125MG/M	ORENCIA INJ 250MB			HUMIRA KIT 40MG/D.8	HUMIRA PEN KIT 40MG/0.8	HUMIRA PEN KIT CROHNS	CAPEN KIT PSORIASI	COPEGUS TAB 200MG	NCIVEK TAB 375MG	-	NFERGEN INJONCO	PEG-INTRON KIT 120 RP	PEG-INTRON KIT 120MCG	PECULTON KIT 150MCG	NTRON KIT SOMCG	NTRON KIT SOMCG RP	PEG-INTRON KIT BOMCG	NTRON KIT BOMCG RP	REBETOL CAP 20014G		RIBAPAK MIS 600/DAY		PAK AMINAY	İ	TAB 200MG	BASPHERE TAB 400MG	RIBASPHERE TAB BOOMG	DIDATAD DAY ADDOMNAY

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the Control	RIBAVIRIN CAP 200 MG	RIBAVIRIN (BULK) POWDER	RIBAVIRIN TAB 200 MS	OCCUPACION CAP ZOURIG	ACAPTOR TAB 150 MG	SAPROPTERIN DAYDROCHLORIDE SOLUBLE TAB 100 MG	TROMBOPAG OLAMINE TAB 12.5 MG (BASE EQUIV)	ELTROMBOPAG OLAMINE TAB 25 MG (BASE EQUIV)	TROMBOPAG OLAMINE TAB 50 MG (BASE EQUIV)	ELTROMBOPAG OLAMINE TAB 75 MG (BASE EQUIV)	ELBINUDINE TAB 600 MG	DEXTRANOMER-SODIUM HYALURONATE INJ GEL 60-15 MG/ML	TERFERON GAMMA-18 INJ 100 MCGNO.5ML (2000000 UNITIO.5ML)	PEGADEMASE BOVINE IN 250 UNITM.	WALHOWATE SOLVIAM IN SOLIN 3 MIGRAIL (BASE EQUIVALENT) HEDDANIC BOTH NYOTH A MCHARMI	DI EDRONG ACID IN CONC. FOR DAINE ISON A MAKEU	ALPHA1-PROTEINASE INHIBITOR (HUMAN) FOR IV SOLN 1000 MG	PHA1-PROTEINASE INHIBITOR (HUMAN) FOR IV SOLN 400 MG	ALPHA1-PROTEINASE INHIBITOR (HUMAN) FOR IN SOLN 500 MG	ALPHA1-PROTEINASE INHIBITOR (HUMAN) FOR IV SOLN 800 MG	TESTERASE INFIBITOR (HUMAN) FOR IV INJ. KIT 500 UNIT	A COTEDACE INITIONALLY INVESTIGATION OF THE CONTRACT CONT	LENTANDE INTERPRETENCE (HUMAN) FOR IV IN 3000 UPS	RHO D IMMUNE GLOBULIN (HUMAN) IM INJ 50 MCG	IMUNE GLOBULIN (RUMAN) IV SOLN 10 GMZ00ML	IMMIUNE GLOBULIN GRUMAN) IV SOLN 1 GMZZOM.	IMUNE GLOBULIN (HUMAN) IV SOLN Z.5 GHYSOMI.	MACINE GLOBULIN (RUMAN) IV SOLN 23 GWISOGAL MICINE GLOBULIN ALUMAN) IV SOLN 5 GWISOGAL	EGINESATIDE ACETATE SOLN IN 10 MGML	EGINESATIDE ACETATE SOLN IN 20 MGZML	IMMUNE GEOBULIN (HUMAN) IV SOLN 10 GM/100ML	RAILINE GLOBULIN (HUMAN) IV SOLN 20 GWIZDOML	MANUNE GLOBULIN (HUMAN) IV SOLN 5 GAZGALL ALOUAT DOOTENHEED HANDETOD JAHRAN FOR BY SOLN 1995 NO.	LPHAT-PROTEINASE INHIBITOR (HUMAN) FOR IV SOLN 500 MG	ALPHA1-PROTEINASE INHIBITOR (HUMAN) FOR IV SOLN 1000 MG	LPHA1-PROTEINASE INHIBITOR (HUMAN) FOR IV SOLN 1000 MG	GRUNE GLOBULIN (FURIAN) IV OR SUBCUTANEOUS SOLN 10 GM/1004R.	MACUNE GLOBULIN (HIMAN) IV ON SUBCUTANEOUS SOLN 1 GAN IMPER MACUNE GLOBULIN (HIMAN) IV OR SUBCUTANEOUS SOLN 2 6 GANZENI	IMMUNE GLOBULIN (HUMAN) IN OR SUBCUTANEOUS SOLN 20 GMIZGONIL	MININE GLOBULIN (HUMAN) IV OR SUBCUTANEOUS SOLN 5 GWISHM.	IMMUNE GLOBULIN (FUNKAN) IN SOLN 10 GMZGOML	IMMUNE GLUBULIN (HUMAN) IN SOLIN ZE SMIDUM.	IMMUNE GLOBULIN MUMANI SUBCITANEOUS IN 1 GMISHI	IMMUNE GLOBULIN (HUNAN) SUBCUTANEOUS IN 2 GM/10ML	MAILINE GLOBULIN (HUMAN) SUBCUTANEOUS INJ 4 GINZOMI.	EXBAZOXANE FOR INJ 250 MG	EXRAZOXANE FOR INJ 500 MG	DEXTRAZOXANE FOR INJ 500 MG	POPROSTENOL SODIUM FOR INJ 1.5 MS	TREPROSTINIL INFALATION SOLUTION 0.5 MG/ML	TREPROSTINIL INHALATION SOLUTION 0.5 MG/MIL	TREPROSTINIL INHALATION SOLUTION O'S MIGML	ADALAFIL TAB 20 MG (PAH)	ALEGORIAN TANDERS OF THE PROPERTY OF THE PROPE
# * * * * * * * * * * * * * * * * * * *	RIBAVIRIN		FUBAVIRIN R		L		CTA		PROMACTA E				ACTIMMUNE		ZOMETA 22	Ī				9	BERINERT		GLASSIA	MPL.			l	OCTAGAM IN					PRIVICEN IN	T	PROLASTIN-C A			Τ	Γ	GAMMAKED	T	GAMMAPLEX III	,	HIZENTRA			OXANE	TOTECT			REFIL	TYVASO START 1		
	Antiviral Agent	Antiviral Agent	Andring Agent	Amin'ni Agenti	Antivirals	Antivirals	Antivirals	Antivirals	Antivirals	Antivirals	Antivitals	Assorted Classes	Biologic Response Modizers	Biologicals	Bisphospironate Definalive	Bischosobonala Darivaliva	Blood Product Derivative	Blood Product Darlyative	Blood Product Derivative	Skod Product Derivative	Blood Praduct Derivative	Blood Product Detroites	Blood Product Derivative	Blood Product Derivativa	Blood Product Derivetive	Blood Product Derivative	Blood Product Derivative	Blood Product Derivative	Blood Product Derivative	Blood Product Derivative	Blood Product Derivative	Blood Product Derivative	Blood Product Derivative	Blood Product Derivative	Blood Product Derivative	Blood Product Dertvetive	Blood Product Derrivative	Blood Product Demyalive	Blood Product Derrivative	Blood Product Denivativa	Blood Product Derrivative	Blood Product Demivative	Blood Product Deminative	Stood Product Dentvalive	Blood Product Derivativa	Cardioprotectant	Cardloprotectant	Cardioprotectant	Cardiovascular Apart	Cardiovascular Agent	Cardiovascular Agent	Cardiovascular Agent	Cardiovascular Agents	Continue of the Asset of
	RIBAVIRIN CAP 200MG			VICINEEDS CAP ZOOMS	KALYDECO TAB 150MG	RUVAN TAB TOOMG	PROMACTA TAB 12.5MG	PROMACTA TAB 25MG	PROMACTA TAB SOMG	PROMACTA TAB 75MG	TYZEKA TAB 600MG	SOLESTA INJS0-15ML	ACTIMMUNE INJ 2MU/0.5	ADAGEN INJ 250ML	ZOWETA IN LANCHOR	ZOMETA INTAKCEMI	ARALAST NP INJ 1000MG	ARALAST NP INJ 400MG	ARALAST NP INJ SDOMG .	ARALAST NP INJ 800MG	BERNERT INJSORUNIT	CHURCAM ANJ TON	GLASSIA INJ	MICRHOGAM PL. INJ SOMCG	l		- 1	OCTAGAM INJEGM	OMONTYS INJ 10MG/ML		PRIVICEN INJ 10GRAMS	PRIVICEN IN 20GRAMS	PRIVICEN INJ 5 GRAMS	PROLESTIN BY DOOMS	PROLASTIN-C INJ 1009MG	ZEMAIRA INJ 1000MG	GAKMAKED INJ 10GM/100	ı	ı	ll	GARMAPLEX IN TOOM	GANMAPLEX IN 25GM	HIZENTRA IN I SCHISM	HIZENTRA INJ 2GM/10ML	HIZENTRA DU 46MZ0ML	DEXRAZOXANE INJ 250MG	DEXRAZOXANE IN 500MG	TOTECT INJ SOOMG	FLOLAN INJUSTICE	TYVASO SOLDENGME	100	TYVASO START SOL O BINGMIL	ADCIRCA TAB 20MG	

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	18 50°	25.01-	-17.0%	-17.0%	-17.0%	10.0%	-16.0%	-13.0%	-13.0%	-15.0%	%0"21-	*60'21-	%0'8-	-15.0%	-18,0%	-13.5%	-13.5%	-13.5%	-11,0%	-11.0%	-8.5%	-9.5%	-11,0%	-11.0%	-13.5%	-13.5%	-15.0%	-8,5%	-17,0%	-13.5%	-18.0%	-15.1%	-15,1%	-15,1%	-15.1%	-15.1%	-15.1%	-16.0%	-18.0%	-16.0%	-16.0%	-16.0%	25.0%	-25.0%	-25.0%	-25.0%	-1.0%	-18.0%	20.00 p	8.0%	-8.0%	-13.5%	-13.5%	-14.6%	-15.5%
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	AMBRISENTAN TAB 5 MG	TREPROSTINIL SODIUM INJ 10 MG/ML (BASE EQUIV)	TREPROSTINIL SODIUM INJ 1 MGMIL (BASE EQUIV)	TREPROSTINIL SODIUM IN 12.5 MG/ML (BASE EQUIV)	TREPROSTINIL SODIUM INJ 5 MGML (BASE EQUIV)	SILDENAFIL CITRATE IV SOLN 10 MGH2.5ML (BASE EQUIVALENT)		A CHARLES INTALATION SOLUTION 10 MCGML	ILOPROST INHALATION SOLUTION 20 MCGARL	SIPULEUCEE-1 SUBPENSION FOR IV INFUSION	I EINABENAZINE TAB 125 MG	FERNAMENACINE TAB 25 MG	SCULLIN CAYBALE DRAL SOLUTION 500 MG/ML	UEFERIPRONE TAB 500 MG	AMERICATIVE CRISIALINE FOR IN 500 MG	DEXRAZOXANE POR IN 150 MG	DEXRAZOXANE FOR INJ 600 MG	ETONOGESTREL SUBDERMAL IMPLANT 68 MG		ETONOGESTREL SUBDERMAL IMPL	DIVENTU	ONABOTUL	CAPSAIGN PAICH 8% & CLEANSING GEL KIT	OPPORTOR PALCH 5% & CLEANSING GEL XIT	DEFERASROX TAB FOR ORAL SUSP 250 MG	DEFERASIROX TAB FOR ORAL SUSP 500 MG	CORTICORELIN OVINE TRIFLUTATE FOR IV INJ 100 MCG	CODITION ALFA FOR IN 1.1 MG	PAMIDRONATE DISORRIMEDE IN 139 MG	CHORONIC GONADOTROPIN FOR IN JADGO LINIT	CHORIONIC GONADOTROPIN (BULK) POMDER	FOLLITROPIN BETA INJ 150 UNITROSME		FOLLITROPIN BETA IN 500 UNITA 22M.	FOLD ROPIN BETA IN 16 UNITY AND	FOLLISTIM PEN FOR RXSOL MAIL SERVICE	FOLLITROPIN ALFA FOR INJ 1050 UNIT	IN ALFA FO	FOLLITROPIN ALFA INJ 300 UNITO.6ML	FOLLITROPIN ALFA IN J 450 UNITO 75ML	FOLLING ALTA FOR INJ 19 UNIT	OCTREOTIDE ACETATE IN 1000 MCGALL (1 MGAR.)	OCTRECTIDE ACETATE INJ 100 MCGML (0.1 MG/ML)	OCTREOTIDE ACETATE IN 200 MCGAIL (0.2 MGAL)	OCTRECTIDE ACETATE IN 500 MCGML (0.5 MG/ML)	OCTREOTIDE ACETATE IN 50 MCGARL (0.05 MGARL)	CHORIOGONADOTROPIN ALFA INJ 250 MCGROSM.	MENOTROPINS FOR IN 75 INIT	PARICALCITOL CAP 1 MCG	17-1	PARICALCITOL CAP 4 MCG	PARICALCITOL IV SOLN 2 MCGMA.	PARICALCITOL IV SOLN S MCGML	LARONIDASE SOLN FOR IN INFUSION 29 MG/SML	UROFOLLITROPIN PURIFIED FOR INJ 75 UNIT
) 	LETAIRIS	REMODULIN	REMODULIN	REMODULIN	REMODULIN	REVATIO	CENTRICO	VENTAND	VENIAVES	PROVERGE.	VENTANE	WOEst.	CCOSTOCA	AMIEDETINE	THAT INC	ZINECARD	ZINECARD	IMPLANON	MIRENA	NEXPLANON	BOTOX COSMET	BOTOX COSMET	AU ENCA	EXIANS	EXJADE	EXADE	ACTHREL	ACTHAR HD	AREDIA	CHOR GONADOT	CHORIONIC	FOLLISTIM AG	FOLLISTIM AQ	FOLLS IIM AG	FOLLISTIM AD	FOLLISTIM PEN	GONAL-F	GONALF	GONAL-F RFF	COMAL CORE	GONAL-F RFF	остяботоб	OCTREOTIDE	OCTREOTIDE	OCTREOTIDE	CHECHDE	DEFENVI	REPRONEX		ZEMPLAR	ZEMPLAR	ZEMPLAR	ZEMPLAR	ALDURAZYME	BRAVELLE
	Cardiovascular Agents	Cardiovascular Agents	Candovascular Agents	Cardovascular Agents	Certinovasculet Agains	Controversion Agents	Cardiate Annual	Cardiocas des Santo	Calledon franciscollander	Carded Management Design	Control Mondaning-Depleting Agent	Central Nervoire Styles Department	Chefulat Anen	Chemonylankie Apani	Champhersov Agents	Chemotherapy Agenta	Chemolherapy Agents	Contraceptive	Contraceptive	Contraceptive	Dermatologicals	Dermalologicals	Demotologisale	Detoxilcation Agent	Detoxilkation Agent	Defoxilication Agent	Disposite Agent	Endocine & Metabolic Apen	Endocine & Metabolic Agent	Endocrine & Metabolic Agent	Endocrine & Metabolic Agent	Endocrine & Metabolic Agent	Endocine & Metabolic Agent	Endocting & Melabolic Association	Endocrine & Melabolic Agent	Endocrine & Metabolic Agent	Endocrine & Melabolic Agent	Endocrine & Metabolic Agent	Endocrine & Metabolic Agent	Endorine & Malabolic Agent	Endocrine & Metabolic Agent	Endocrine & Metabolic Agent	Endocrine & Metabolic Agent	Endocrine & Metabolic Agent	Endocane & Metabolic Agent	Concerns & Meanboll Agent	Endocrine & Metabolic Agent	Endocrine & Metabolic Agent	Endocrine & Metabolic Agent	Endocrine & Metabolic Agent	Endocrine & Metabolic Agent	Endocine & Metabolic Agent	Endocrine & Metabolic Agent	Endocrine & Metabolic Agants	Endocrine & Metabolic Agents
Here a first	LETAIRIS TAB SMG	REMODULIN INJ 10MG/ML	SENODICE IN THIS SECOND	SCHOOL IN IN SHOW	DENATIO IN	1	10	VENTAVIS SOL SOUCEAU		XEMATINE TAR 45 GLAS	CENAZINE TAB 25MG	XYREM SOL SOUNGAM	FERIPROX TAR SOUNG	AMERICATIVE IN SOCIAL	ETHYOL INJ SOOMS		INECARD INJ SDOMG	IMPLANDN IMP 68MG	MIRENA HID SYSTEM	VEXPLANON IMP 68MG	BOTOX COSMET IN 166UNIT	DOLUMENTA KIT 84 1 DOLUMENT	OUTENZA KIT 8% 2-PCH	EXAMPLE TAB 125MG	1 1	EXJADE TAS 500MG	ACTHREL INJURIOR	CTHAR HP INJ BOUNT	AREDIA INJ 30MG	CHOR GONADOT INJ 10000UNT	CHORIONIC POW GONADOTR	FOLLISTIM AQ INJ 150UNIT	FOLLISTIM AC IN SOCIAL	FOLLISTIM AO INJ 75UMT	CALLISTIM AG INJ 900UNIT		GONAL-F INJ 1050UNIT	GONAL-F INJ 450UNIT	GONAL-FREF INJ 300	ONAL-FREE IN 751 NIT	GONAL-FRFF INJ 900	OCTREOTIDE INJ 1000MCG	OCTREOTIDE INJ 100MCG	OCTREOTIDE IN 200MCG	CTREATINE IN SOUNCE	COLUMN IN SUMPRIME		150	ZEMPLAR CAP 1MCG			ZEMPLAR INJ ZWCGMIL	CHIPAGAME IN SMCGMIL	ALDURACIME INJ. SMISISM	KAVELLE INJ COUNT

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:	LO US	5	20,03	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0,00	\$0.00	\$0.00	20.00	86.98	9 8	3.5	5 5	3 5	80.03	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	20.00	30.00	20.00	20.00	\$0.00	\$0.00	\$0.00	20.00	\$0.00	\$0,00	20.00	20.09	200	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$9.00	\$0.00	90.03	8 8	20.00
	-11.5%	-13.5%	-15.5%	-15.5%	-15.0%	-15.0%	-14,6%	*13.5%	-18.0%	-1.0%	-1.0%	-1,0%	-18.0%	-18.0%	-18.0%	-18.0%	-18.0%	-10.5%	-12.5%	80.11-	-17.0%	42.076	250 177	10.55	***************************************	-19.5%	-17.5%	-17,5%	-17.5%	-12.0%	-16.5%	-11.5%	-15.5%	-15.5%	-16.6%	- 20 OK	30.0%	-30.0%	-30.0%	-15,0%	-12.0%	-15,0%	-15.0%	-15.0%	-11.5%	-21.5%	-11.076	-13.6%	-13,5%	-40,5%	-40.5%	-40,5%	-40.5%	-40.5%	-40.5%	49'07-	-10.0%	-15.5%	%C'91-
	30090025106440	30906850002020	30903610402120	30903610102110	30090040102020	30062050002175	30907715002120	30907535002020	30052020002140	30904045000130	30804045000110	30904045000120	30042060102006	30042060102120	30042060102009	30042080102012	30042000102140	30044530002020	SUMEZUSUOUZUZU	10454060000320	20405225100320	30405225400320	30905225100340	30170050102040	30170050102025	30170050102030	30180060002120	30180060002130	30180060002140	30044530002630	82700080102120	99350035002120	43103030102045	4310303U102U2D	63403030403040	83403030405	83103030102020	83103030102035	83103030102040	51200080002030	50250070102020	52505020106420	52505020106440	52505020106450	52500050102020	SCHOOL WAS A	3010000010010K	85601010002120	85601010002110	85100010252140	85100010252150	85100010252170	85100010252120	85100010252180	85100010252185	85100010252130	02+00000000000000000000000000000000000	85100028202150	OZIZNZBZNANI GQ
	LIX ACETATE FOR INJ KIT	IDURSULFASE SOLN FOR IV INFUSION 6 MIGRAIL (2 MIGML)	AGALSIDASE BETA FOR IV SOLN 36 MG	AGALSIDASE BETA FOR IV SOUN 5 MG	GANIRELIX ACETATE IN 250 MCGROSM.	MENDIFICATIONS FOR SUBCUTANEOUS INJ 75 UNIT	ALGELOCUSIONSE ALFA FUR IV SULVA SU MIS	IGALSULFASE SOLN FOR IV INFUSION 1 MGML	ICHURICAIC GUNADO IROPIN POR INJ 10000 UNIT	MILIONOLIC CAP TV MG	MILIONOME CAS A MG	Introduction of the production of the state	PAMIDACAN E DECOMO IV SOLN S MOME	PAMERCINAL DISCUSING FOR ING 30 MG	PAMERICANA E DISCOMEM IV SOLN O MESMI.	Detail DONATE DISCOUNT IS SOLD STREAM.		DENCOUNTS OF MOTING	TO VAPTAN TAR 45 MG	TOLVAPTAN TAB 30 MG	CINACAL CET HOL TAB 30 MG RASE FOLITO	CINACALCET HC. TAB 80 MG (BASE FOLIV)	CINACALCET HOL TAB 30 MG (BASE EQUAY)	LANREOTIDE ACETATE EXTENDED RELEASE IN 120 MON.5ML	LANREOTIDE ACETATE EXTENDED RELEASE INJ 80 MGM; 2ML	LANREOTIDE ACETATE EXTENDED RELEASE IN 30 MG/0,3ML	PEGVISOMANT FOR IN 10 MG (AS PROTEIN)	FEGVISOMANT FOR INJ 15 MG (AS PROTEIN)	PEGVISOMANT FOR INJ 20 MG (AS PROTEIN)	DENOSUMAB IN 120 MG/L/7ML	I ALIGEDUCERANE ALTA FOR INJ 200 UNIT	COLLEGED OF CONTRACTOR AND TO MAKE THE PORT OF THE POR	CONOMINATION CONTINUES CARGO CAR	FONDADADINI Y SOUTH WELL STORY AND	FONDADARING SOUTHERN AND SERVICE SEE	FONDAPARINUX SODIUM IN 10 MG/0.8ML	FONDAPARINUX SODIUM IN 2.5 MG/0.5ML	FONDAPARINUX SODIUM INJ 5 MG/D,4ML	FONDAPARINUX SODIUM IN 17.5 MG/L/GML.	SACROSIDASE SOLN 8500 UNITRAL	PALCNOSETRON HOL IV SOLN 0.25 MG/SML (BASE EQUIVALENT)	CERTOLIZUMAB PEGOL FOR INJ KIT 2 X 200 MG	CERTOLIZUMAS PEGOL INJKIT 2 X 200 MG/ML	CERTOLIZUMAH PEGOL INJ KITB X200 MGML	METANTINE TO EVANUE DO JUDE IN 12 METANT AND STORY	METAN SALTERSONE PROMINE IN KIT 12 MAGARM	SOMATROPIN ROW REFRIGERATED FOR SUBCUTANEOUS IN BAKE	ALTEPLASE FOR INJ 160 MG	ALTEPLASE FOR INJ 50 MG	ANTIKEMOPHILIC FACTOR RAHF-PFM FOR INJ 1000 UNIT	ANTINEMOPHILIC FACTOR RAHF-PFM FOR INJ 1500 UNIT	ANTHEMOPHLIC FACTOR RAHF-PFM FOR INJ 2000 UNIT	ANTINEMOPHE IC FACTOR RAHE-PEM FOR INJ 250 UNIT	ANTHEMOPHILIC FACTOR RAHF-PFM FOR INJ 3000 UNIT	ANTIHEMOPHILIC FACTOR RAHE-PFM FOR INJ 4000 UNIT	ANTHEMOPHILE FACTOR RAHE-PEM FOR INJ 600 UNIT	COASCULATION FACTOR IX RECOMBINANT FOR IN 1900 UNIT	COAGULATION FACTOR IX RECOMBINANT FOR IN 2501 UNIT	
	CETROTIDE	ELAPRASE	FABRAZYME	FABRAZYME	GANIKELIX AC	MENOPOR	MI OCINIC	NAGCAZYME	NOVANE			1	TI WOOD ON THE	CAMIDEONATE	DANIDONATE	DAMEDONATE	PRO! 14	DEC! AST	SAMSCA	SAMSCA	SENSIPAR	SENSIPAR	SENSIPAR	SOMATULINE	SOMATULINE	SOMATULINE	SOMAVERT	SOMAVERT	8		STASI EV	ADMOA	ADMITTOR	ADIVIDA	ARIXTRA	FONDAPARINUX	FONDAPARINUX	FONDAPARINUX	FONDAPARINUX.	SUCRAID	ALOX	CIMZIA	CIMZIA	CIMZIA	NET INTO BE	REI ISTOR						ADVATE	ADVATE	ADVATE	ADVATE	ADVATE	DENEFIN	BENEFIX	DCINCTIO
	Endocrine & Metabolic Agents	Endocrine & Metabolic Agents	Endocrine & Metabolic Agents	Endocine & Metabolic Agents	Endocrate & Metabolic Agents	Codondos Photobolic Agents	Control of Materials Assets	Endocinia & Metabolic Agents	Endersing & Melabolic Agents	Cadomine & Malaballa Assair	Endoning & Metabolic Aponte			Endocrine & Metabolic Agents		Federate & Metabolic Apents	Endocripe & Metabolio Apenis			Endocrine & Metabolic Agents								Endocrine & Metabolic Agents	Endocrino & Metabolic Agems	Endocine & Metabolic Agents	England	Eactor Xv Ithilian	Earler X7 hibitor	Bactor Xv Ibiblior	Factor Xz (bb) lor	Factor Xz Inbitor				Gastrointestinal	Gastrointestinal Agents	Gastrointestinal Agents	Gastroinfestinal Agents	Castrollads that Agents	Control testing Acente	Gastrointesfingl Apends	Grawith Harmone	Hemstological Agent	Hamstological Agent	Hematological Agents	Hemutological Agents	Hematological Agents	Hematological Agents	Hematological Agents	Hematological Agents	Hematological Agents Hematological Anems	Hematological Agents Hematological Agents	Hemetological Agents	Following Chame
		ELAPRASE INJEMICIAMI	FABRAZYME IN 35MG	PABRACTME INJUNG	CONTRACTOR DESCRIPTION	MENOTON IN TOUR	MACI ADMIT THE ARCAN	NASCAZINE IN INGINE		ı	POR FAIR	DANING THE IN SOMEMI	DANIOSONATE IN SOUTH	DAVIDOONSTC IN SUCRUI	DAMINECNATE IN SOLION	PAMIDEONATE IN BOMG	PROLIA SOL SOMGAM	16	TAB 15MG	SAMSCA TAB 30MG	7 TAB 30MG	TAB 60MG	SENSIPAR TABBOMG	SOMATULINE INJ 120/5ML	SOMATULINE INJERIOZAL	SOMATULINE INJ 900:3ML	SOMAVERT INJ 10MG	SOMAVERT INJ 15MG	tal .	AGENO INTRACTAL	KINE EX INTO SHIP	APIXTRA SOF 100 A			ARIXTRA SOL 7.50.8	FONDAPARINUX SOL 10:0.8	FONDAPARINUX SOL 250.5	FONDAPARINUX SOL 5.00.4	FONDAPARINUX SOL 7.503.6	SUCRAID SOL 8500/ML	- 1	- 1	CIMZIA KIT ZOOMGAAL	CIMICA NII STANIEN	BELIETOP IN 180 AUP	RELISTOR KIT 12/0.5ML	SEROSTIM INCENS	ACTIVASE INJ 100MG	1			ADVATE IN 2000RNIT	ı	ıI		ADVATE IN SOUNIT		BENEFIX IN ZONOLNI	

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	50.03	2000	\$0.00	\$0.00	\$0,00	\$0.00	\$0.00	30.00	\$0.00	90.00	20.00		00.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$9.00	\$0.00	\$0.00	89	80.08	8.8	\$0.00	8 3	8 8	808	8 6	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	20,00	\$0.00	\$0.00	\$0.00	\$0,00	\$0.00	\$0.00	30.00		\$0.09	\$6.00	80.00	\$0.00	\$0.00	\$0.00	20.03	8 8	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00
	-16.5%	-16.5%	-18.0%	-18,0%	-18,0%	-18.0%	-15.0%	18,0%	-10.07	200 CC	-32 085	-13.5%	-13 0%	*18.0%	-18,0%	-18,0%	-18.0%	-18.0%	-18.0%	~18.0%	-18.0%	30,5%	-30.5%	30.5%	-33.5%	80.00-	8/5/50-	-26.0%	78092	-21,5%	-21.5%	-21.5%	21.5%	-21.5%	21.5%	74570	-140%	-17.5%	-17.5%	-17.5%	-17.5%	-17.5%	47.5%	-47.5%	-10,0%	-10.0%	-10.0%	-10.0%	-10.0%	-15.5%	-15.5%	-16,1%	-15.0%	-15.5%	-15.6%	-75,5%	-15,5%
	85100028202160	85100028202130	83101020102016	83101020102018	83101020102020	83101020102012	831010Z010Z050	RESTOTOCOTOCOTA	R3401020105015	85100020000100	85100020002100	\$5820040102020	85840030002020	83101020102016	83101020102018	83101020102020	53101020102012	83101020102050	63101020102013	63101020102014	63101020102015	85140428042180	65100028002160	85100028602770	85100010202155	astonographase	85100010202123	85100035002120	85100015102129	85100010266440	85100010286460	85100010266420	85100010266430	85199010266460	85100010268470	RS100010266220	82700065102120	82401015112043 ,	82401015112048	82401015112054	82401015112014	82401015112064	82401015112024	82401015112034	82401020002040	82401020002010	82401020002050	82401020002015	82401020002020	82402050002120	82402050002025	82404570002020	82403060002120	82401520002016	82401520002010	82401520002018	82401520002012
	COAGULATION FACTOR IX (RECOMBINANT) FOR INJ 3000 UNIT	COAGULATION FACTOR IX (RECOMBINANT) FOR INJ 500 UNIT	ENOXAPARIN SOCIUM IN 100 MGMI.	ENOXAPARIN SODIUM IN 1120 MG/0.8M.	ENCAPARIN SOCION IN 150 MEINE.	ENOMAPARIN SOCIETA IN 300 MORALI		ENOXAPARIN SODIUM INJ 60 MGR. 6M.	ENDXAPARIN SODIUM INJ 80 MGro, 691.	'ANTIINHIBITOR COAGULANT COMPLEX FOR IN."	"ANTIINHIBITOR COAGULANT COMPLEX FOR IN."	ICATIBANT ACETATE IN 30 MG3ML (BASE EQUIVALENT)	ECALLANTIDE IN 10 MGML	ENOXAPARIN SODIUM IN J 100 MGML	ENOXAPARIN SODIUM INJ 120 MG/0.8ML	ENOXAPARIN SODIUM INJ 150 MG/ML	ENOXAPARIN SODIUM IN J 30 MG/0.3ML	ENOXAPARIN SODIUM INJ 300 MG/3ML	ENDXAPARIN SODKIM INJ 40 MG/0.4ML	ENOXAPARIN SODIUM INJ 60 NGCLERIL	COASSI ATION EACTOR IN EACH IN 1 4000 IN THE	COACH ATION FACTOR IN FIGURIAL	COACULATION EACTOR IX COS IN COS INST	ANTHEMORIE OF EACTOR DECOMPRISATIONS OF LIVE	ANTHEMORPH IC FACTOR (RECOMBINANT) FOR BUILD COMP.	ANTHEMOPHILIC FACTOR (RECOMBINANT) FOR IN 1 401-800 LINIT	ANTIHEROPHILIC FACTOR (RECOMBINANT) FOR INJ 801-1249 UNIT	FIBRINGEN CONC (HUMAN) INJ APPROXIMATELY 1 GM (900-1300 MG)	ANTIHEMOPHILIC FACTOR/WIF (HUMAN) FOR INJ 500-500 UNIT	ANTHEMOPHILIC FACTOR RECOMBINANT PAF FOR INJ KIT 1000 UNIT	ANTIHEMOPHILIC FACTOR RECOMBINANT PAF FOR INJ KIT 2009 UNIT	ANTIHEMOPHILIC FACTOR RECOMBINANT PAF FOR INJERT 250 UNIT	ANTIHEMOPHILIC FACTOR RECOMBINANT PAFFOR IN KIT 500 UNIT	ANTHER COURT OF EACHOR RECOMBINANT PARTOR IN LIFE AND UNIT	ANTREMOPHIC FACTOR RECOMBINANT PAFFOR INJURIT SOLINIT	ANTINEMOPHILIC FACTOR RECOMBINANT PAF FOR INJKIT 250 UNIT	VELAGLUCERASE ALFA FOR INJ 400 UNIT	DARBEPOETIN ALFA-POLYSORBATE 80 SOLN INJ 100 MCG/0.5ML	DARBEPOETIN ALFA-POLYSORBATE 80 SOLN INJ 150 MCG/0.3ML.	DARBEPOETIN ALFA-POLYSORBATE 80 SOLN IN J 200 MCG/0.4ML	DARBEPOETIN ALFA-POLYSORBATE 80 SOLN IN 25 MICGIO 42M.	DARBEDOFTIN ALFA-POLYSCHEMIE BU SOLIN INJ 300 MCCROSM.	DARBEPOETIN ALFA-POLYSORBATE RO SOLN INJ 500 MCCANL	DARBEPOETIN ALFA-POLYSORBATE 80 SOLN INJ 60 MCG/D.3ML	EPOETIN ALFA INJ 10000 UNITAIL,	EPOETIN ALFA INJ 2000 UNITIME.	EPOETIN ALFA INJ 20000 UNITAIL	EPOETIN ALFA INJ 3000 LINITAN.	EPOETIN ALFA INJ 4000 UNITANI.	SARGRAMOSTIM LYOPHILIZED FOR INJ 250 MCG	SARGERAMUSTAM INLIBRO MICGIMI. TO EDIYARDO STIPOLITANEDIE INTERNACIONAL CONTINUE		OPRELVEKIN FOR INJ 5 MG	FILGRASTIM INJ 300 MCG/0.5ML (800 MCG/ML)	FILGRASTIM INJ 300 MCG/ML	FILGRASTIM INJ 480 MCG/0,8ML (800 MCGML)	FILGRASTIM INJ 480 MCG11.8ML (300 MCGML)
de en esta	BENEFIX	BENEFIX	ENOXAPARIN	ENOXAPARIN	CHONAPARIN	ENOXAPARIN	ENOXAPARIN	ENOXAPARIN	ENOXAPARIN	FEISANF	FEISA VH	FIRAZYR	KALBITOR	LOVENOX	LOVENOX	LOVENOX	OVENOX	LOVENOX	OVENOX	COVENCY	MONONINE	MONONINE	MONONINE	RECOMBINATE	RECOMBINATE	RECOMBINATE	RECOMBINATE	RIASTAP	MLATE	XYNTHA	AHLINX	XMTHA	ATNIES.	XVATTA SOLOF	XYNTHA SOLOF	XYNTHA SOLOF	VPRIV	ARANESP	ARANESP	ARANESP	ARANESP	ARANESP	ARANESP	ARANESP	EPOGEN	EPOGEN	EPOGEN	EPOGEN	POGEN	LEUKINE	LEUKINE	NEUL ASTA	NEUMEGA	NEUPOGEN	NEUPOGEN	NEUPOGEN	NEUPOGEN
	Hematological Agents			Hematological Agents	Hermitological County	Hematological Agents			Hematological Agents			Hematological Agents										Hemaloboical Acents			Hematological Agents	Hematological Agents	Hemafological Agents	Hemalological Agents	Hematological Agents				Hematological Agents	Hematological Appois	Hematological Agents							Hematopoletic Agent					Hemulopoletic Agent				Hematopoieuc Agent						Hematopoletic Agent
	H	BENEFIX INJ SOUDIT	ENOXAPARIN IN 100MG/ML	ENOXAPARIN INJ 1200.8	ENOXABABIN INTARAM	ENOXAPARIN INJ 300/3/M	ENOXAPARIN INJ 400.4ML	ENOXAPARIN INJ 60/0.6ML	ENOXAPARIN INJ 800.8ML	FEIBANF INJ	FEIBA VH INJ IMMUNO	FIRAZYR INJ 30MG/3ML				- 1	COVENOX IN 300.3ML	ı	LOVENOX IN AUGAM		Jш	MONONINE INJ 250UNIT		15	RECOMBINATE IN 220-460	RECOMBINATE INJ 401-800	RECOMBINATE INJ 801-1240	RIASTAP SOL 1GM	WRATE IN		-	1	XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX	XXXTHA SOLDE IN COORDINE	XYMTHA SOLOF INJ SOUNT	XYNTHA SOLOF KIT 250UNIT	VPRIV INJ 400UNIT		ARANESP INJ 150MCG	- 1	APANESP IN ZOMUS	1	APANESP INJ SOOMCG	1	ı	EPOGEN INJ 2000/ML	- 1	-1	- 1	LEUKINE IN 250kG3		NEULASTA INFEMGROEM		NEUPOGEN IN 300/0.5	NEUPOGEN IN 300IACG	NEUPOGEN INJ 480/0.6	NEUPOGEN INJ 480MCG

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	¢)	20.00	\$0,00	\$0.00	\$0.00	\$0.00	\$0.00	20.00	20.00	\$0.00	\$0.00	\$0.00	\$0.00	.00'0\$	00'08	2000	20.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	00.03	50.00	10.00	\$0.00	\$0.00	\$0.00	\$0.00	50.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	20.05	30.00	\$0.00	\$0,00	\$0.00	50.00	\$0.00	\$0,00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	80.02	20 09	40.00	\$0.00
	-13.1%	-13.0%	-13.0%	-13.0%	-13,0%	-13.0%	-13.0%	*0.51-	10.00 A	-13,5%	-43.0%	-13,8%	-13,8%	-13,8%	-25.0%	-16.0%	-46.5%	-46.5%	46.5%	-46.5%	-46.5%	-46.5%	46.5%	10.234 46.5%	46.5%	-48.5%	-46.5%	₩0,14-	-41.0%	30.5	*40°F-	-1.0%	-46,5%	48,5%	%5"9 ►	46.5%	26.5%	-46.5%	-48.5%	-46.5%	46.5%	-26,0%	-26.0%	-26.0%	-31.5%	-31.5%	-31.5%	-31,5%	-26,0%	-28.0%	-26.0%	-18.0%	1000 F	740 OK	20007 20007
	82405060002120	82405060002130	82401020002040	82401020002010	\$2401020002050	82401020002015	82401020002020	82200050005000	82700050002120	82700070000120	85100015102170	85100028002180	85100028002185	85100028002170	6510003000Z150	84100040002010	8510001020644D	85100010206450	85100010206420	85100010206460	85100010208430	351U0010206440	85100010206420	85100010002109	85100010002125	85100010002170	\$5100010002180	85100015102122	85100015102132	85100015102144 [85100010002140	185100010002140	85100010002130	85100010206440	85100010206440	65100010206450	85100010206450	25100010205420	85100010206460	\$5100010206460	65100010206430	85100010206430	85100010006480	85100010006410	85100016006430	85100025202117	85100025202126	65100026202145	65100026202160	85100030002110	85100030002115	ARTOOTOGASO	84100040002010	19100020102135	19100020102155	1120toSpootest
Section 1	ROMIPLOSTIM FOR INJ 250 MCG	ROMIPLOSTIM FOR INJ 500 MCG	EPOETIN ALFA INJ 10000 UNITML	EPOETIN ALFA INJ 2000 UNITAIL	EPOETIN ALFA INJ 20000 UNITANI.	EPOETIN ALFA IN 3000 UNITAR	EPOETRI ALFA IN LARGO UNITALI	MIGHTOGRASE FOR IN 200 LINIT	MIGLUCERASE FOR IN 400 UNIT		ANTIHEMOPHILIC FACTORAWF (HUMAN) FOR INJ 500 UNIT	COAGULATION FACTOR IX FOR INJ 1000 UNIT	COAGULATION FACTOR IX FOR INJ 1500 UNIT	COAGULATION FACTOR IX FOR INJ SOD UNIT	FACTOR IX COMPLEX FOR IX 1 200-1200 DMT	TRANEXAMIC ACID INJ 100 MGMIL.	ANTIHEMOPHILIC FACTOR (RECOMBINANT) FOR INJ KIT 1000 UNIT	ANTIHEMOPHILIC FACTOR (RECOMBINANT) FOR INJ KIT 2000 UNIT	ANTIHEMOPHILIC FACTOR (RECOMBINANT) FOR INJIKIT 250 UNIT	ANTIHEMOPHILIC FACTOR (RECOMBINANT) FOR INJKIT 3000 UNIT	ANTHEMOPHILIC FACTOR (RECOMBINANT) FOR INJ KIT 500 UNIT	ANTIDEMOPTIC PACTOR (RECUMBINATO) FOR INJURIOUS UNIT	ANTHEMOPHIC FACTOR (RECOMBINANT) FOR INJUST 250 UNIT	ANTIHEMOPHILIC FACTOR (HUMAN) FOR INJ 220-400 UNIT	ANTIHEMOPHILIC FACTOR (HUMAN) FOR INJ 401-800 UNIT	ANTIHEMOPHILIC FACTOR (HUMAN) FOR INJ 1501-2000 UNIT	ANTIHEMOPHILIC FACTOR (HUMAN) FOR INJ 804-1500 UNIT	ANTIHEMOPHILIC FACTORAWF (HUMAN) FOR INJ 250-600 UNIT	ANTHEMOPHILE FACTORAWF (HUMAN) FOR INJ 500-1200 UNIT	ANTHER CHILD FACTOR ARE MANNED IN LACK INC. TOUR LACT.	ANTHEMOCHIEC FACTOR OFTIMAN FOR IN 1550 UNIT	ANTIHEMOPHILIC FACTOR (HUMAN) FOR INJ 500 UNIT		ANTHEMOPHILIC FACTOR (RECOMBINANT) FOR INJ KIT 1000 UNIT	ANTIHEMOPHILIC FACTOR (RECOMBINANT) FOR INJ KIT 2000 UNIT	ANTHEROPHLIC FACTOR (RECOMBINANT) FOR IN KIT 2005 UNIT	ANTHEMORPH IC FACTOR (RECOMBINANT) FOR INJ KIT 250 UNIT	ANTHEMOPHILIC FACTOR (RECONBINANT) FOR INJ KIT 3000 UNIT	ANTHEMOPHILIC FACTOR (RECOMBINANT) FOR INJ. KIT 3000 UNIT	ANTIHEMOPHILIC FACTOR (RECOMBINANT) FOR INJ KIT 509 UNIT	ANTHEMOPHILIC FACTOR (RECOMBINANT) FOR INJ KIT 500 UNIT	ANTHERCEPHERS FACTOR (FRUMAN) FOR INJ NJ 1000 UNIT	ANTHEMOPHILIC FACTOR HUMAN FOR INJ KIT 250 UNIT	ANTIHEMOPHILIC FACTOR (HUMAN) FOR INJ KIT 500 UNIT	COAGULATION FACTOR VIIA (RECOMB) FOR INJ 1 MG (1000 MCG)		COAGULATION FACTOR VIIA (RECOMB) FOR INJ 5 MG (5000 MCG)	7	FACTOR IX COMPLEX FOR IN 1000 UNIT	FACTOR IX COMPLEX FOR INJ 1500 UNIT	ANTHEMORHI IS EASTED JECONOMISTOR IN THE STATE OF THE STA	TRANEXAMIC ACID IN 100 MONL	MAMINE GLOBULIN HUMANI IV FOR SOLN 12 GM	BILLIANDE ALCOULING HORIZON IN COLON COLLA LE COL	MAMOUNT CODE IN CHARANT IN CONTROL OF THE PRINT OF THE PR
	NPLATE	NPLATE	PROCRIT	PROCRIT	PROCRIT	PROCE	PROCEIT	CEREZYME	CEREZYME	ZAVESCA	ALPHANATE	ALPHANINE SD	ALPHANINE SD	ALPHANINE SD	REPUTIN VH	CYKLOKAPRON	HELIXATE FS	HELIXATE PS	HELIXATE PS	HELIXATE FS	HELIXATERS	HELIVATE CO	HELIXATEFS	HEMOFIL IN	HEMOFIL M	HEMOFILM	HEMOFIL M	HUMATE-P	HUMATEP	KOATE, DV	KOATE-DVI	KOATE-DVI	KOGENATE FS	KOGENATE FS	KOGENATE FS	KOGENATERS	KOGENATE FS	KOGENATE FS	KOGENATE FS	KOGENATE FS	KOGENATE FS	MONOCA ATER	MONOCLATEP	MONOCLATEP	NOVOSEVEN RT	NOVOSEVEN RT	NOVOSEVENRT	NOVOSEVEN RT	PROFILMINE	PROFILINE	PROFILMINE	TRANEXAMIC	CARIMINENE	A POINT INE NE	CALCIMONE IN
	Hemakopoletic Agent	Hematopoletic Agent	Hematopoletic Agent	Hematopoletic Agent	Hematopoletic Agent	Hematopoleuc Agent		95		Hematopolisic Agents	Hemostatics			Hemostatics	Herrostatics						Hemostalics			Hemostatics	Hemostatics	Hemostatics	Hemoslatics	Hemostatics	Hemoslatics	Homostoffes	Hemostatos	Hemostefics				Hemostatics Hemostatics	Hemostatics	Hemostatics	Hemostatics	Hemostatics	Hemostatics	Hemostants				Hemostatics	Hemostatics	Hemostatics	Hamostaßes	Hemostatics	Hemostalics	Hemostatics	Indianamirina Anard	Imministed Agent	manual Agent
	INPLATE IN J. 250MCG	l I		PROCRIT INJ 2000/ML		PROCEET IN AGORA	PROCEET IN ACCOUNT	CEREZYME IN ZODUNIT	CEREZYME INJ 400UNIT	ZAVESCA CAP 100MG	ALPHANATE INJ VWF/HUM	ALPHANINE SD INJ togotiniT	ALPHANINE SD INJ 1500UNIT	ALPHANINE SD INJ SOUCHT	BEBLEIN VH. INJ 200-1200	CYKLOKAPRON INJ 100MG/ML	HELIXATE FS INJ 1000UNIT	HELIXATE FS INJ 2000UNIT	HELIXATE FS INJ 250UNIT	HELIXATE FS IN 3000UNIT	HELIXALE ES INJ SOUCINIT	LELIVATE ES SOL JUDGUNII	HELIXATE ES SOL SOMINIT	HEMOFIL M INJ 220-400	HEMOFIL M INJ 401-800	HEMOFILM SOL. 501-2000	HEMOFIL M SOL 801-1500	HUMATE-P INJ SOOUNIT	HUMATE-P SOL 1200UNIT	KOATE-DVI IN LODGEST	KOATE-DVI IN ZSOUNIT	KOATE-DVI INJ SOOUNIT	KOGENATE FS INJ 1000/BS	KOCENATE PS INJ 1000UNIT	KOGENATE FS IN 2000/BS	KOSENATE ES INJ 2000UIT	KOGENATE ES INJ 250UNIT	KOGENATE FS INJ 3000/BS	KOCENATE PS IN 3000UNIT	KOGENATE FS INJ 500/BS	KOGENATE FS INJ SOOUNIT	MONOCLATE-P IN TOUGHIII	MONOCI ATEP IN I SEGUNIT	MONOCLATE-P INJ SOOUNIT	NOVOSEVEN RT INJ 1MG	NOVOSEVEN RT INJ 2MG	NOVOSEVEN RT INJ SIMG	NOVOSEVEN RT INJ 8MG	PROFILININE INJ 1000UNIT	PROFILNINE INJ 1500UNIT	PROFICHINE INJ SOUCHIT	TREFACTO INJ SUBLINI	ANDREADING IN TORING	CADIMINENT IN 125M	CARIMUNE Nº 444 SOUR

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	ALOVIRUS (MIMUNE GLOBULIN (AUMAN) IV INJ		OBULIN (HUMAN) IV SOIN 0.5 GM 10ML	SOLN 20 GM/400ML	OBULIN (HUMAN) INJ	OBULIN (MUMAN) IV OR SUBCUTANEOUS SON N 10 GM/10mul	IMMUNE GLOBULIN (HUMAN) IV OR SUBCUTANEOUS SOLN 1 GMYCM)	Ī				CO COLL COMPONING								IMMUNE GLOBULÍN (HUMAN) IV OR SUBCUTANEOUS SOLN 5 GM50ML			ECHI DI IMMUNE GLOBULIN (RUMAN) NU 1500 UNITZMI. (300 MCG/ZML) 191						RHO D IMMUNE GLOBULIN (HUMAN) INJ 5000 UNITRACML			RUME GLOBULIN (HUMAN) INJ 150 UNITAIL	TOCKLIZUMAB IV INJ 200 MG/10ML	ABIVING MOMENT	PT FOR IN 200 MG	EPT SUBCUTANEOUS INJ 25 Marc. SM.	EPT FOR SUBCUTANEOUS IN KIT 25 MG	EPT SUBCUTANEOUS IN J SO MIGME.	PT SUBCUTANEOUS INJ 50 MGML	SUBCUTANEOUS IN 100 MGK.67M.	GOLIMUMAB SUBCUTANEOUS INJ 60 MG/0,5ML	KA ATE KORETII CAD SENIO	NOLATE MOFETIL FOR ORAL SUSP 200 MGAM	VOLATE MOFETIL TAB 500 MG	JOLATE MOFETIL HOL FOR IV SOLN 500 MG (BASE EQUIV)		3AP 100 MG	RINE CAP 25 MG		CYCLOSPORINE MODIFIED CAP 50 MG				CACA OSPORING MODIFIED CAP TOURS		TACROLIMUS CAP D.5 MG BOAR	
	CYLOGAM	FLEBOGAWMA	FLEBOGAMMA	FLEBOGANIMA	GAMASTAN S/D	GAMMAGARD	GAMMAGARD	GAMMAGARD	GANNAGARD	GAMMAGARD	GAMMAGARD	GAMMAGARD SD	GAMMAGARD SD	GAMMAGARD SD	GAMUNEX	GAMUNEX-C	GAMUNEX-C	GAMINEX-C	GAMUNEX.C	GAMENEX-C	MOGAM RABIE	KHOLOWING AC	SOI IPIE	SYNAGIS	SYNAGIS	POS OFFINA	PAINTHO SDF	WANRHO SDF	WINRHO SDF	BENLYSTA	BENLYSTA	THE SECTION	ACTEMBA	ACTEMBA	ARCALYST	ENBREL	ENBREL	ENBREL	ENBREL SRCLK	KINERET	SIMPONE	CELCEPT	CELICEPT	CELLCEPT	CELCEPTIV	CYCLOSPORINE	CYCLOSPORINE	CYCLOSPORINE	CYCLOSPORINE	CYCLOSPORINE	CTCLUSPURINE	CVCIOCOOBINE	GENGRAE	GENGRAF	GENGRAF	HECORIA	
Imministra Awad	Mariantizating Agent	Institutiong Agent	Immunizing Agent	Immunizing Agent	(mmunizing Agent	Immunizing Agent	Immunizing Agent	lomenizing Agent	Immunizing Agent	Immunizing Agent	Immunizing Agent	Immunizing Agent	Immunizing Agent	Immunizing Agent	Immunizing Agenl	framunizing Agent	Immunizing Agent	Immunizing Agent	Immunicing Agent	Intribution Agent	International Agent	Imministra Annal	Immunizaci Acent	kmunizing Agent	Immunizing Agent	Immunizing Agent	Immunizing Agent	Immunizing Agent	Immunizing Agent	Immunizing Agents	Immunizing Agents	International regular	Internation Appears	Immunologic Agents	Immunologic Agents	Immunologic Agents	immunologic Agents	Immunologic Agents	Immunologic Agents	Immunologic Agents	Immunosuppessive Agent	Immunosuppressive Agent	Immunosuppressive Agent	Immunosuppressive Agent	Immunosuppressive Agent	Immunosuppressive Agent	immunosappiessive Agent	munitosuppressiva Agent	inimutesuppressive Agent	Interest appleasing Agent	from incention in the Agent	Impunosuppressing Agent	Immuosuomessiva Agent	Immunosuppressive Agent	Immunosuppressive Agent	Intrimusesuppressive Agent	
CCIOCAM IN	12	FLEBUSAMIRA INJ 10%	FLEBOGAMMA INJ5%	FLEBOGAMMA INJUNE 5%	GAMASTAN SID INJ	GAMMAGARD INJ 10GMJ100	GAMMAGARD INJ 1GM/10ML	GAMMAGARD INJ 2.5GM/25	GAMMAGARD INJ 20GMIZ00	GAMMAGARD INJ 30GM/300	GAMMAGARD INJ SGM/SOML	GAMMAGARD SD INJ 10GM HU	GAMMAGARD SD INJ 2.5GM HU	GAMMAGARD SD INJ 5GM HU	GAMUNEX BNJ 10%	GAMUNEX-C INJ 10GM/100	GAMUNEX-C IN 1GM/10ML	GAMUNEX.C INJ 2.5GM25	GAMMUNEX-C IN ZUGMZOU	BANGONEA-C INJOONADONE		l	SOURIS IN TOWGIME	SYNAGIS INJ 100MS/ML	SYNAGES INJEOMG	WINRHO SDF INJ 15000UNT	WARRED SDF INJ 1500UNIT	WINNERHO SDF INJ 2500UNIT	WINITERO SOF INJ 5000UNIT		HADEBRAR SID IN 1 1 SOOMS	ACTEMBA IN CONTROL	ACTEMBA IN 400200M		ARCALYST IN 1220MG	П		ENBREL INJ SOMSING	ENERGE SECUL IN SOMEONIL	SHOOM INTERES	ı	CELLCEPT CAP 250MG	CELLCEPT SUS 200MGAML	CELLCEPT TAB 500MG	CELLCEPT IV INJ 556MG	CYCLOSPURINE CAP TURKS	CYCLOSPONIE CAP TUNIS MU	CYCL CSPORINE CAP 25MG	CYCLOSPORINE CAP Zame mod	CVCI OSPORINE INI SOMOMI	CYCLOSPORINE SOIL SOMEONE	CYCLOSPORINE SOL MODIFIED	GENGRAF CAP 100MG	1		HECORIA CAP 0.5MG	ı

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草	MYCOPHENOLATE MOPETIL TAB 500 MG	MYCOPHENOLATE SODIUM TAB DR 180 MG (MYCOPHENOLIC ACID EQUIV)	MITCUPTENCIALE SCUIM 1AB DIT 350 MS (MYCOPHENOLIC ACID EQUIV)	CYCLOSPORINE MODIFIED CAP SAME	DYCLOSPORINE MODIFIED ORAL SOLIN 100 MG/ML	BELATACEPT FOR IV INFUSION 250 MG	TACROLIMUS CAP 0,5 MG	TACROLIMUS CAP 1 MG	TACROLIMUS CAP 5 MG	TACROLIMUS INJ 5 MG/ML	SIROLIMUS ORAL SOLN 1 MGML		SIROLIMUS TAB 1 MG	SIROLIMUS TAB 2 MG	CYCLOSPORINE CAP 100 MG	CYCLOSPORINE CAP 25 MG	CYCLOSPORINE IV SOLIN 50 MGML	CYCLOSPORINE ORAL SOLN 100 MGML	TACROCIMUS CAP 0.5 MG	TACROLIMUS CAP 1 MS	TACROLIMUS CAP 5 MG	TACROLIMUS (BULIQ POWDER	TACROLIKUS (BULK) POWDER	EVEROLIMUS TAB 0.25 MG	EVEROUMUS TAB 0.5 MG	EVEROLIMUS I AB U. O MG	INTERFERON ALEANS IN SAMAN HATAI	INTERFERENCE ETA-14 FOR IN INTERFERENCE CARDAMAIN	NTERFERON BETA-14 IM IN KIT 30 MCGG-5MI	INTERFERON BETA-1A IM INJ KIT 30 MCG/0,5/ML	INTERFERON ALFA-2B FOR INJ 10000000 UNIT	INTERFERON ALFA-ZB INJ GOMBOO UNITMIL	INTERFERON ALFA-28 INJ 10000000 UNITAIL	INTERFERON ALFA-28 FOR INJ 50000000 UNIT	PEGINTERFERON ALFA-2A INJ 180 MCG/0.5ML	PEGINTERFERON ALFA-24 IN J 180 MCGAIL	PEGINTERFERON ALFA-2A IN J 180 MCG/0,5ML	PEGINTERFERON ALFA-2A INJ KIT 180 MCG/0.5ML	PEGINTERFERON ALFA-2B FOR INJKIT 298 MCG	PEGINTERFERON ALFA-28 FOR INJKIT 444 MCS	PEGINTERFERON ALFA-2B FOR INJ KIT 888 MCG	CARGLUMIC ACID TAB 200 MG	TETANUS IMMUNE GLOBULIN (HUMAN) INJ 250 UNITML	CLOBAZAM TAB 10 MG	CLOBAZAM TAB 20 MG	CLOBAZAM TAB 5 MG	BOSENTAN TAB 125 MG	BOSENIAN IAB 62.5 MS	SOBIUM HYALURONATE INTRA-ARTICULAR INJ 20 MISZIMI.	APONICKTRINE MYLKOCHLOKICH ING NORMI	AN UNION THINK THE WILLIAM STATES THE TOWNS TO SECOND THE SECOND	SOUTHWAT TANDED IN TOWARD IN SOUTH TO SUMMERS AND SOUTHWAT THE SOUTHWAT TOO BUT SOUTHWIT	CANBOLDINOR CARA FOR IN 100 CALL	ADDROTT IN MINISTRAIN FOR IN 1990 INTE	ABOROTI BINI METONINA POR INI EGO INIT	DIMARCHI MINITORNE IN LANGO INITIALI	INTRODUCTULATION INTO THE SECOND CONTINUES.	RIMABOTULINUMTOXINB INJ 5000 UNITYML	
		MYFORTIC	_				щ		PROGRAF	PROGRAF	RAPAMUNE			1	١	-		1	١	TACROLIMUS	-		গু			2	AI FERON N			AVONEX PREFL		INTROMA	INTRON-A			PEGASYS							RTETSO				TRACLEER		EUFLEXXA	1	MINO NA	DOTON	AUTOR	BOIOX	DVebror	MYORIOC	MYORIOC	MYOBLOC	
			International particular American			Immunosuppressive Agent	kmmmosuppressive Agent		Immunosuppressive Agent	Innunosuppressive Agent	Immunosuppressive Agent				Immunosuppressive Agent		immundsuppressive Agent	Immunosuppressive Agent	Immunosuppressive Agent	Immacsuppressive Agent	Immunosuppressive Agent	Immunosuppressive Agent	7	7	manufaction Agent	annumount years Agent	Interferon					Interferon		Interferon					Interferon	interferon	Interferon	Metabolic Alkalosis Agent				Msc.			ıls	Notional Agent			wedlo)muschar Agents	Weuromuscular Agents	Management Agains	Marroquierider Arable	Metromuscular Agents	Neuromuscular Agents	
MYCOPHENOLAT CAP 250MG	MYCOPHENOLAT TAB 500MG	MYFORTIC TAB 180MG	NECODAL CAD 100MG		NEORAL SOL 100MG/MIL	NULOJIK INJ 250MG	Iъ.	ı	PROGRAF CAP SMG	1	PAPAMAINE SOLIMOMAL		RAPAMUNE TAB 1MG	RAPAMUNE TABIZMG	SANDIMINIONE CAP 100MG	SANDIMININE CAP ZEMIS	SANDIMUNE IN SOMEONE	SANDIMINUNE SOL 100MG/ML	TACROLAMUS CAP 0.5MG	TACROLIMUS CAP 1MG	TACROLIMUS CAP SING	TACROLIMUS POW	TACROLIMUS POW MONOHYD	ZORTRESS TAB 0.25MG	ZORINESS IAB OSMG	CORINESS IAB U. ORGAN	ALEERON N INJSMILM	AVONEX KIT 30MCB	AVONEX PEN XIT 30MCG	AVONEX PREPLIKIT 30MCG	INTRON-A INJ 16MU	INTRON-A INJ 18MU	INTRON-A INJ 25MU	INTRON-A INJSOMU	PEGASYS INJ		PEGASYS INJ PROCLICK	PEGASYS KIT	SYLATRON KIT 295MCG	SYLATRON KIT 444MCG	SYLATRON KIT 888MCG	CARBAGLU TAB 200MG	HYPERTET SID INJ 250ML	ONFI TAB 10MG	ONEI TAB 20MG	ONFI TAB SMG	TRACLEER TAB 125MG	IRACLEEK IAB 62,5MG	s١		APORTN INC TURBURAL	HALGAN IN ZUMGZONI		٦.	DISPURI IN SOUNII			MYOBLOC INJ SOOGNL	

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		40.02	45.001-	-13-078	42.0%	42 0%	-13.0%	.45.5%	-16.5%	-16.5%	-18.5%	+13.5%	-150%	-15.0%	-(5.0%	-15.0%	11.5%	-11.5%	-11.5%	-1.5%	-11.5%	-11.5%	-11.5%	-13.0%	-13.0%	-13.0%	-18.5%	-16.5%	-18.0%	-19.0%	-17.0%	-15.8%	-4.5%
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	Johnson State	SOMATROPIN FOR INJ 1.6 MG	SOMATROPIN FOR INJ 1.8 MG	SOMATROPIN FOR INJ 12 MG (13,8 MG OVERFILL)	SOMATROPIN FOR IN 1 MG	SOMATROPIN FOR INJ 2 MG	SOMATROPIN FOR SUBCUTANEOUS INJ SING	SOMATROPIN FOR INJ 12 MG (36 UNIT)	SOMATROPIN FOR IN 24 MG	SOMATROPIN FOR INJ 5 MG	SOMATROPIN FOR INJ 6 MG (18 UNIT)	MECASERMIN IN J 40 MG/AML (10 MG/ML)	SOMATROPIN INJ 10 MG/1.5ML	SOMATROPIN INJ 15 MG/1,5ML	SOMATROPIN IN 30 MG/2ML.	SOMATROPIN INJ S MGH. SML	SOMATROPIN FOR INJ 10 MG	SOMATROPIN FOR SUBCUTANEOUS INJ 5 MG	SOMATROPIN INJ 10 MG/2ML	SOMATROPIN INJ 20 MG/2MI.	SOMATROPIN INJ S MG/ZMI.	+STERILE TOWEL DRAPES 18X2 FOR RXSOL MAIL SERVICE	GROWTH HORMONE PEN KITS FOR RXSOL MAIL SERVICE	SOMATROPIN INJ 10 NG/1.5ML	SOMATROPIN FOR INJ S.B MG	SOMATROPIN IN 5 MG/1.5ME.	SOMATROPIN (NON-REFRIGERATED) FOR IN J 5 MG	SOMATROPIN (NON: REFRISERATED) FOR INJ 8,8 MG	SOMATROPIN (NON-REFRIGERATED) FOR SUBCUTANEOUS INJ 4 MG	SOMATROPIN (NOW-REFRIGERATED) FOR SUBCUTANEOUS INJ 5 MG	SOMATROPIN FOR SUBCUTANEOUS IN SMG	SOMATROPIN (NON-REFRIGERATED) FOR SUBCUTANEOUS INJUA MG	ALTEPLASE FOR IN 2 MG
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Price Summary



Confidential and Proprietary

The following administrative fees and rates are exclusive to City of Cincinnati. Rates and fees are effective upon the implementation be services; this offer expires in 90 days.

Retiree Drug Subsidy - Enhanced Services

Subsidy Services

- Interim Reporting
 - Download Covered Retiree List for eligible members from RDS*
 - Create cost summary reports based on drug utilization of members listed in Covered Retiree List as per RDS submission requirements and guidelines
 - Storage and archival of backup data per RDS guidelines for audit purposes
- Reconciliation
 - Create final cost summary data for the reconciliation plan year
 - · Provide back up claims and rebate data, if requested by client
 - Support B/D drug methodology, as per RDS guidelines
 - Provide calculations for ACA (Actual Cost Adjustment) using rebates
 - Storage and archival of backup data per RDS guidelines for audit purposes
 - Coordination of individual retiree cost:
 If our services are used, the client has to provide us with claim data and rebate information from the previous vendor. A separate one time charge of \$5,000 is associated with it.
- Audit
 - Provide backup data claims, rebates, and drug lists
 - Provide documentation to support methodology (Part B vs. Part D) used for calculating final cost data
 - Additional services required will be negotiable
- Transmission
 - Upload cost reports to RDS**

	<u> </u>
Account Setup for groups with fewer than 500 RDS Members	\$5,000
Creditable Coverage Determination	\$500 to \$1,500
Additional Subsidy Related Services	\$500 per hour - as negotiated
Actuarial Certification & Attestation	\$1,500 to \$3,500

Notes

- * Client must designate OptumRx as a designee
- ** Client must designate OptumRx as a designee or cost reporter

\$0.60 per Paid Claim in addition to Administrative Fee

06/20/2013 City of Cincinnati 1 / 1

EXHIBIT D

BUSINESS ASSOCIATE ADDENDUM

This Business Associate Addendum ("Addendum") is effective as of January 1, 2014, and is incorporated into and made part of the Prescription Drug Benefit Administration Agreement ("Agreement") by and between **OptumRx**, **Inc.** ("Business Associate") and **City of Cincinnati** ("Covered Entity") (each a "Party" and collectively the "Parties").

The Parties hereby agree as follows:

1. <u>DEFINITIONS</u>

- 1.1 Unless otherwise specified in this Addendum, all capitalized terms used in this Addendum not otherwise defined have the meanings established for purposes of the Health Insurance Portability and Accountability Act of 1996 and its implementing regulations as amended and supplemented by HITECH, as each is amended from time to time. (collectively, "HIPAA") Capitalized terms used in this Addendum that are not otherwise defined in this Addendum and that are defined in the Agreement shall have the respective meanings assigned to them in the Agreement.
- 1.2 "Affiliate", for purposes of this Addendum, means any entity that is a subsidiary of UnitedHealth Group.
- 1.3 "Breach" means the acquisition, access, use or disclosure of PHI in a manner not permitted by the Privacy Rule that compromises the security or privacy of the PHI as defined, and subject to the exclusions set forth, in 45 C.F.R. § 164.402.
- 1.4 "Breach Rule" means the federal breach regulations, as amended from time to time, issued pursuant to HIPAA and codified at 45 C.F.R. Part 164 (Subpart D).
- 1.5 "Compliance Date" means the later of September 23, 2013 or the effective date of the Agreement.
- 1.6 "Electronic Protected Health Information" ("ePHI") means PHI that is transmitted or maintained in Electronic Media.
- 1.7 "HITECH" means Subtitle D of the Health Information Technology for Economic and Clinical Health Act provisions of the American Recovery and Reinvestment Act of 2009, 42 U.S.C. §§ 17921-17954, and all associated existing and future implementing regulations, when and as each is effective.
- 1.8 "PHI" means Protected Health Information, as defined in 45 C.F.R. § 160.103, and is limited to the Protected Health Information received from, or received, maintained, created or transmitted on behalf of, Covered Entity by Business Associate in performance of the Services.
- 1.9 "Privacy Rule" means the federal privacy regulations, as amended from time to time, issued pursuant to HIPAA and codified at 45 C.F.R. Parts 160 and 164 (Subparts A & E).
- 1.9 "Security Rule" means the federal security regulations, as amended from time to time, issued pursuant to HIPAA and codified at 45 C.F.R. Parts 160 and 164 (Subparts A & C).
- 1.10 "Services" means, to the extent and only to the extent they involve the receipt, creation, maintenance, transmission, use or disclosure of PHI, the services provided by Business Associate to Covered Entity under the Agreement, including those set forth in this Addendum in Sections 4.3 through 4.7, as amended by written agreement of the Parties from time to time.

2. RESPONSIBILITIES OF BUSINESS ASSOCIATE

With regard to its use and/or disclosure of PHI, Business Associate agrees to:

- 2.1 not use and/or further disclose PHI except as necessary to provide the Services, as permitted or required by this Addendum, and in compliance with each applicable requirement of 45 C.F.R. § 164.504(e) or as otherwise Required by Law; provided that, to the extent Business Associate is to carry out Covered Entity's obligations under the Privacy Rule, Business Associate will comply with the requirements of the Privacy Rule that apply to Covered Entity in the performance of those obligations.
- 2.2 implement and use appropriate administrative, physical and technical safeguards and, as of the Compliance Date comply with applicable Security Rule requirements with respect to ePHI, to prevent use or disclosure of PHI other than as provided for by this Addendum.
- 2.3 without unreasonable delay, report to Covered Entity (i) any use or disclosure of PHI not provided for by this Addendum of which it becomes aware in accordance with 45 C.F.R. § 164.504(e)(2)(ii)(C); and/or (ii) any Security Incident of which Business Associate becomes aware in accordance with 45 C.F.R. § 164.314(a)(2)(i)(C).
- 2.4 with respect to any use or disclosure of Unsecured PHI not permitted by the Privacy Rule that is caused solely by Business Associate's failure to comply with one or more of its obligations under this Addendum, Covered Entity hereby delegates to Business Associate the responsibility for determining when any such incident is a Breach and for providing all legally required notifications to Individuals, HHS and/or the media, on behalf of Covered Entity. Business Associate shall provide these notifications in accordance with the notification requirements set forth in the Breach Rule, and shall pay for the reasonable and actual costs associated with those notifications. In the event of a Breach, without unreasonable delay, and in any event no later than sixty (60) calendar days after Discovery, Business Associate shall provide Covered Entity with written notification in accordance with 45 C.F.R. § 164.410 that includes a description of the Breach, a list of Individuals (unless Covered Entity is a plan sponsor ineligible to receive PHI) and, in the event the delegation set forth above has been triggered, a copy of the template notification letter to be sent to Individuals.
- 2.5 in accordance with 45 C.F.R. § 164.502(e)(1)(ii) and 45 C.F.R. § 164.308(b)(2), ensure that any subcontractors of Business Associate that create, receive, maintain or transmit PHI on behalf of Business Associate agree, in writing, to the same restrictions and conditions on the use and/or disclosure of PHI that apply to Business Associate with respect to that PHI, including complying with the applicable Security Rule requirements with respect to ePHI.
- 2.6 make available its internal practices, books, and records relating to the use and disclosure of PHI to the Secretary for purposes of determining Covered Entity's compliance with the Privacy Rule.
- 2.7 document, and within thirty (30) days after receiving a written request from Covered Entity, make available to Covered Entity, information necessary for Covered Entity to make an accounting of disclosures of PHI about an Individual, in accordance with 45 C.F.R. § 164.528.
- 2.8 provide access, within twenty days (20) days after receiving a written request from Covered Entity to PHI in a Designated Record Set about an Individual, to Covered Entity, sufficient to allow Covered Entity to comply with the requirements of 45 C.F.R. § 164.524.
- 2.9 to the extent that the PHI in Business Associate's possession constitutes a Designated Record Set, make available, within thirty (30) days after a written request by Covered Entity, PHI

for amendment and incorporate any amendments to the PHI as requested by Covered Entity, all in accordance with 45 C.F.R. § 164.526,

3. RESPONSIBILITIES OF COVERED ENTITY

In addition to any other obligations set forth in the Agreement, including in this Addendum, Covered Entity:

- 3.1 shall identify the records it furnishes to Business Associate it considers to be PHI for purposes of this Addendum.
- 3.2 shall provide to Business Associate only the minimum PHI necessary to accomplish the Services.
- 3.3 in the event that the Covered Entity honors a request to restrict the use or disclosure of PHI pursuant to 45 C.F.R. § 164.522(a) or makes revisions to its notice of privacy practices of Covered Entity in accordance with 45 C.F.R. § 164.520 that increase the limitations on uses or disclosures of PHI or agrees to a request by an Individual for confidential communications under 45 C.F.R. § 164.522(b), Covered Entity agrees not to provide Business Associate any PHI that is subject to any of those restrictions or limitations to the extent any may limit Business Associate's ability to use and/or disclose PHI as permitted or required under this Addendum unless Covered Entity notifies Business Associate of the restriction or limitation and Business Associate agrees to honor the restriction or limitation. In addition, if those limitations or revisions materially increase Business Associate's cost of providing services under the Agreement, including this Addendum, Covered Entity shall reimburse Business Associate for such increase in cost.
- 3.4 shall be responsible for using administrative, physical and technical safeguards at all times to maintain and ensure the confidentiality, privacy and security of PHI transmitted to Business Associate pursuant to the Agreement, including this Addendum, in accordance with the standards and requirements of HIPAA, before and during the transmission of such PHI to Business Associate.
- 3.5 shall obtain any consent or authorization that may be required by applicable federal or state laws and regulations prior to furnishing Business Associate the PHI for use and disclosure in accordance with this Addendum.
- represents that is has ensured, and has received certification from Plan Sponsor, that Plan Sponsor has taken the appropriate steps in accordance with 45 C.F.R. § 164.504(f) and 45 C.F.R. § 164.314(b) to enable Business Associate on behalf of Covered Entity to disclose PHI to Plan Sponsor, including but not limited to amending its plan documents to incorporate, and agreeing to, the requirements set forth in 45 C.F.R. § 164.504(f)(2) and 45 C.F.R. § 164.314(b). Covered Entity shall ensure that only employees authorized under 45 C.F.R. §164.504(f) shall have access to the PHI disclosed by Business Associate to Plan Sponsor.

4. PERMITTED USES AND DISCLOSURES OF PHI

Unless otherwise limited in this Addendum, in addition to any other uses and/or disclosures permitted or required by this Addendum, Business Associate may:

- 4.1 make any and all uses and disclosures of PHI necessary to provide the Services to Covered Entity.
- 4.2 use and disclose PHI for proper management and administration of Business Associate. In addition, to carry out the legal responsibilities of Business Associate, provided that the disclosures are Required by Law or any third party to which Business Associate discloses PHI for

those purposes provides written assurances in advance that: (i) the information will be held confidentially and used or further disclosed only for the purpose for which it was disclosed to the third party or as Required by Law; and (ii) the third party promptly will notify Business Associate of any instances of which it becomes aware in which the confidentiality of the information has been breached.

- 4.3 De-identify any and all PHI received or created by Business Associate under this Addendum, which De-identified information shall not be subject to this Addendum and may be used and disclosed on Business Associate's own behalf, all in accordance with the De-identification requirements of the Privacy Rule;
- 4.4 provide Data Aggregation services relating to the Health Care Operations of the Covered Entity in accordance with the Privacy Rule.
- identify Research projects conducted by Business Associate, its Affiliates or third parties for which PHI may be relevant; obtain on behalf of Covered Entity documentation of individual authorizations or an Institutional Review Board or privacy board waiver that meets the requirements of 45 C.F.R. § 164.512(i)(1) (each an "Authorization" or "Waiver") related to such projects; provide Covered Entity with copies of such Authorizations or Waivers, subject to confidentiality obligations ("Required Documentation"); and disclose PHI for such Research provided that Business Associate does not receive Covered Entity's disapproval in writing within ten (10) days of Covered Entity's receipt of Required Documentation.
- 4.6 make PHI available for reviews preparatory to Research and obtain and maintain written representations in accord with 45 C.F.R. § 164.512(i)(1)(ii) that the requested PHI is sought solely as necessary to prepare a Research protocol or for similar purposes preparatory to Research, that the PHI is necessary for the Research, and that no PHI will be removed in the course of the review.
- 4.7 use the PHI to create a Limited Data Set ("LDS") in compliance with 45 C.F.R. § 164.514(e).
- 4.8 use and disclose the LDS referenced in Section 4.7 solely for Research or Public Health purposes; provided that, Business Associate shall (i) not use or further disclose the information other than as permitted by this Section 4.8 or as otherwise Required by Law; (ii) use appropriate safeguards to prevent use or disclosure of the information other than as provided for by this Section 4.8; (iii) report to Covered Entity any use or disclosure of the information not provided for by this Section 4.8 of which Business Associate becomes aware; (iv) ensure that any agents to whom Business Associate provides the LDS agree to the same restrictions and conditions that apply to Business Associate with respect to such information; and (v) not identify the information or contact the Individuals.
- use and disclose PHI for Covered Entity's health care operations purposes in accordance with the Privacy Rule, including (a) conducting quality assessment and improvement activities with respect to the Benefit Plan services provided by Covered Entity through Business Associate; (b) conducting evaluations of Benefit Plan performance; (c) business planning and development; (d) conducting, on behalf of Covered Entity, population-based activities relating to improving the health of Members of Covered Entity's Benefit Plan and reducing their healthcare costs; (e) contacting Members of Covered Entity's Benefit Plan, on behalf of Covered Entity, with health education information and information about prescription drugs, treatment alternatives, and related functions; and (f) communicating with Members of Covered Entity's Benefit Plan, on behalf of Covered Entity, to describe health-related products or services (or payment for such products or services) provided by or included in Covered Entity's Benefit Plan through Business Associate's services, including communications about pharmacies participating in the Plan's network, replacement of or enhancement to the Plan, and health-related products or services available only to Members that add value to, but are not part of the Benefit Plan. Covered Entity

and Business Associate agree that these communications with Members constitute health care operations conducted on behalf of the Covered Entity.

5. TERMINATION AND COOPERATION

- 5.1 <u>Termination</u>. If either Party knows of a pattern of activity or practice of the other Party that constitutes a material breach or violation of this Addendum then the non-breaching Party shall provide written notice of the breach or violation to the other Party that specifies the nature of the breach or violation. The breaching Party must cure the breach or end the violation on or before thirty (30) days after receipt of the written notice. In the absence of a cure reasonably satisfactory to the non-breaching Party within the specified timeframe, or in the event the breach is reasonably incapable of cure, then the non-breaching Party may terminate the Agreement, and/or this Addendum.
- 5.2 <u>Effect of Termination or Expiration.</u> Within sixty (60) days after the expiration or termination for any reason of the Agreement and/or this Addendum, Business Associate shall return or destroy all PHI, if feasible to do so, including all PHI in possession of Business Associate's subcontractors. In the event that Business Associate determines that return or destruction of the PHI is not feasible, Business Associate shall notify Covered Entity in writing and may retain the PHI subject to this Section 5.2. Under any circumstances, Business Associate shall extend any and all protections, limitations and restrictions contained in this Addendum to Business Associate's use and/or disclosure of any PHI retained after the expiration or termination of the Agreement and/or this Addendum, and shall limit any further uses and/or disclosures solely to the purposes that make return or destruction of the PHI infeasible.
- 5.3 <u>Cooperation</u>. Each Party shall cooperate in good faith in all respects with the other Party in connection with any request by a federal or state governmental authority for additional information and documents or any governmental investigation, complaint, action or other inquiry.

6. MISCELLANEOUS

- 6.1 <u>Construction of Terms</u>. The terms of this Addendum to the extent they are unclear shall be construed to allow for compliance by Covered Entity and Business Associate with HIPAA.
- 6.2 <u>Survival</u>. Sections 5.2, 5.3, 6.1, 6.2, and 6.3 shall survive the expiration or termination for any reason of the Agreement and/or of this Addendum.
- 6.3 <u>No Third Party Beneficiaries</u>. Nothing in this Addendum shall confer upon any person other than the Parties and their respective successors or assigns, any rights, remedies, obligations, or liabilities whatsoever.
- 6.4 <u>Independent Contractor</u>. Business Associate and Covered Entity are and shall remain independent contractors throughout the term. Nothing in this Addendum or otherwise in the Agreement shall be construed to constitute Business Associate and Covered Entity as partners, joint venturers, agents or anything other than independent contractors.
- Motices. All notices given in connection with this Addendum shall be made in accordance with the applicable provisions of the Agreement. In addition, Covered Entity hereby directs Business Associate to send a copy of any notice or other communication given by Business Associate in connection with this Addendum to the following address (and/or, at Business Associate's discretion, provide verbal notice to the following telephone number) and/or to such other address(es) (or telephone number(s)) as Covered Entity may in the future designate in writing by proper notice. If such address(es) (or telephone number(s)) belongs to a third party, Covered Entity hereby acknowledges and agrees that Business Associate may rely on the direction in this section as being permissible under HIPAA and HITECH, and any other then-

effective laws or regulations relating to the use and/or disclosure of PHI, by virtue of a valid business associate relationship having been established between Covered Entity and such third party.

ADDRESS:

City of Cincinnati

805 Central Avenue

Cincinnati, OH 45202

PHONE NUMBER:

513-352-2551

[REMAINDER OF PAGE INTENTIONALLY LEFT BLANK. SIGNATURE PAGE FOLLOWS]

IN WITNESS WHEREOF, each of Covered Entity and Business Associate has executed in its name and on its behalf this Addendum effective as of the date first written above.

COVERED	ENTITY
City of Cinc	innati

Title:

BUSINESS ASSOCIATE OptumRx, Inc.

By: Del L. Holi

Name: David L. Holms

Ast an

Date: 12/3//13

By: Nagu 7/20E

Name: WAYNOMillSA

Title: SVP, Client STRUICOS

Date: 12.30.13

RECOMMENDED BY:

Karen Alder, Risk Manager

APPROVED AS TO FORM

Assistant City Solicitor

EXHIBIT E PERFORMANCE GUARANTEES

Service and Performance Guarantees

OptumRxTM is pleased to place an aggregate total of \$140,000 at risk annually for City of Cincinnati, of which \$56,000 will be applied to all implementation activities and \$84,000 will be applied to ongoing service standards in Contract Year 1. In subsequent contract years, 100% of all performance guarantee penalties will be applied to ongoing service standards.

Penalties will be assessed annually based upon aggregate annual average results. Penalty amounts per standards will be mutually agreed upon. Guarantees are to be monitored internally and reported quarterly.

				Live Case (see	N 700,3724 34,774,525 10,432,5
Network Pharmac	y Compliance				
Pharmacy Network Access	95% of members will have access to 1 pharmacy in 5 miles on average.	(Metric is Client- Specific)	\$4,800	\$8,000	\$8,000

RamilEnperObit	magaing and a	irfrire		0.5	
Retail Paper Claims Processing Time	97% of paper claims reimbursed or responded to within 10 business days of receipt.	Percent of paper claims reimbursed or responded to that do not require intervention (clean claims).	\$4,800	\$8,000	\$8,000
	All paper claims reimbursed or responded to within an average of 10 business days of receipt.	A minimum of 100 paper claims per year is required to qualify for this guarantee.			
MailiOrder:Claim	Processing Ti	(Metric is Client-Specific) me:			Eparting town

4. ·					
Mail Pharmacy Clean Turnaround	95% of clean prescription orders will be shipped within an average of 2.0 business days	Measured in whole business days from the date a prescription order is received by Administrator (either via mail, phone, fax, or Internet) to the date the prescription order is shipped. Calculated by taking the total number of whole days to ship divided the total number of prescription orders. (Metric is Client-Specific)	\$ 4 ,800	\$8,000	\$8,000

Mail Pharmacy Intervention Prescription Turnaround	100% of prescription orders requiring intervention (Problem) will be shipped within an average of 5.0 business days.	Measured in whole business days from the date a prescription order is received by Administrator (either via mail, phone, fax, or Internet) to the date the prescription order is shipped. Calculated by taking the total number of whole days to ship divided the total number of prescription orders. Contact with prescription orders. Contact with prescriber or customer not achieved as a result of unresponsiveness for an intervention prescription order will be excluded from calculation. (Metric is	\$4,800	\$8,000	
		Client-Specific)			

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Mail Pharmacy Dispensing Accuracy	99.99% of mail pharmacy and specialty pharmacy prescriptions dispensed accurately with no errors.	Accuracy Rate means (i) the number of all mail order prescriptions dispensed in a contract quarter less the number of those prescriptions dispensed in such contract quarter which are reported and verified as having been dispensed with the incorrect drug, strength, form, patient name, directions, packing non-conformance, or address causing medication to be delivered incorrectly divided by (ii) the number of all mail order prescriptions dispensed in such contract quarter. (Metric is total book of business driven)	\$4,800	\$8,000	\$8,000
Retail and Mail C	laims Frocessii	igeAccuracy and in			A CONTRACTOR OF THE STATE OF TH

1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1					
Claims Paid without Error	≥99.9%	Percent of all claims audited and found to be without error of any form, divided by all claims audited. Based on Administrator's internal quality review.	\$4, 800	\$8,000	\$8,000
		(Metric is total book of business driven)			

Gustômer Serv	ice				
Average Speed of Answer	30 seconds or less	The amount of time that elapses between the time a call is received into a member service queue to the time the phone is answered by a Customer Service Representative. Includes calls routed to IVR.	\$6,000	\$10,000	\$10,000
		(Metric is Client- Specific)			

Percent of Calls Abandoned	≤3%	Percentage of calls abandoned by the caller before call is answered by a Customer Service	\$6,000	\$10,000	\$10,000
		Calculated as the number of calls that are not answered, divided by the number of calls received.	·		
		Includes calls routed to IVR and calls abandoned within the first 15 seconds.		•	
	•	(Metric is Client- Specific)		١	
First Call Resolution	95% of calls resolved during initial call.	Calculated as the total calls to Administrator minus total number of unresolved calls, divided by the total number of calls received.	\$4,800	\$8,000	\$8,000
	,	Excludes calls routed to IVR.			
		(Metric is total book of business driven)			

Written Inquiry Response Time	97% of written inquiries received via e- mail will be responded to by e-mail within 5 business days. All written inquiries will be responded to within an average of 10 business days.	Member inquiries received via designated e-mail box. Response time for all written inquiries will be based on the number of calendar days subtracting the date received by Administrator from the date the	\$4,800	\$8,000	\$8,000
		response was sent. (Metric is Client- Specific)			
Member Satisfaction Survey	"Overall Member Satisfaction" survey results of "Satisfied" and "Very Satisfied" for 90% of respondents.	Member satisfaction results will be measured by the responses to Administrator's Voice of the Customer satisfaction survey.	\$ 4 ,800	\$8,000	\$8,000
·		(Metric is total book of business driven)			
Account Manag	oment				
Satisfaction with Account Management	"Overall Client Satisfaction" surve results of "Satisfied" and "Very Satisfied" wi be 7 or better on a 10 point scale Client must	measured by the response to Administrator's	\$4,800	\$8,000	\$8,000
	participate in the survey to qualify fo this guarantee.		, , , , , , , , , , , , , , , , , , , ,		

Plan	Administrator will	(Metric is	\$4,800	\$8,000	\$8,000
Management	provide attendance	Client-Specific)			
Meetings	by plan			•	
	representatives		į į		
	trained on Client's				
	plan benefits at				
	100% of meetings				
į	scheduled by Client,				
	for 100% of the				
	meeting's duration,				
	including all				
	Wellness and				
	Benefit Fairs, and				
	Client-sponsored				
	open enrollment				
	meetings.				

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Administration					
Electronic Eligibility Load -Standard	a between the state of the stat	Assumes complete and accurate information is received. This applies to maintenance loads only, not initial eligibility setup. Assumes use of an electronic interchange and Administrator's standard file format. (Metric is Client- Specific)	\$4,800	\$8,000	\$8,000

** - 1:					
Standard Financial and Clinical Reporting	45 days after the end of the quarter.	Measured as the time from the last day of the end of a reporting cycle to the day standard reports are sent. Ad hoc/custom reporting requests are excluded from this standard. (Metric is Client-Specific)	\$ 4 ,800	\$8,000	\$8,000
Ongoing Plan Design Set up —New Benefits or Updates	Within 7 business days.	Measured by Administrator's ability to implement and test new or revised plan design changes after receipt of signed documentation of new plan design. Any change considered rush or non-standard will be determined based upon a mutually agreed upon timeframe and excluded from this guarantee.	\$4,800	\$8,000	\$8,000
		(Metric is Client- Specific)			

Annual Benefit Plan Review	Maintain a documented quality control and pre- implementation document and provide it to Client for review and approval at least 15 days prior to implementation of any benefit or program change	to coincide with Client's plan implementation of benefit plan modifications. If such reviews identify any	\$4,800	\$8,000	\$8,000

Carroneo ar Warrania.	i Sings:	To the First of Contract			i Translation Proposition
New Client Impleme	ntation				
Implementation Client Satisfaction	"Post Implementati on" survey results of all implementatio n categories will be an average rating of 7 or better based on a 10 point scale.	survey results from all contacts who participate in implementation	\$8,000	N/A	N/A

Implementation Manager	Administrator will assign an implementatio n manager suitable to Client that will be mutually agreed upon between Client and Administrator	(Metric is Client-Specific)	\$8,000	N/A	N/A
Implementation Tasks	No later than 120 days prior to the effective date, tasks with deliverable dates, necessary to effectively install the program by the effective date, will be clearly defined by the Administrator and presented to Client.	(Metric is Client-Specific)	\$8,000	N/A	N/A

	•				
Benefit Set Up	Upon receipt of final sign- off from Client of plan parameters, Administrator will load, fully test, and release the plan benefit coding information for production within 6 weeks of Client's final sign off.	limited to member cost	\$8,000	N/A	N/A
		Client-Specific)			

-					
ID Card Production & Mailing	Accurate ID cards will be mailed at least 10 days prior to the effective date.	Assuming receipt of clean and accurate eligibility files from Client 30 business days prior to the effective date. (Metric is Client-Specific)	\$8,000	N/A	N/A
Pre-Implementation Audit	Administrator will fully support auditor requests for pre- implementatio n audit and will schedule on-site portion of audit at least 15 days prior to the effective date.	Provided the audit is conducted at least 6 weeks after Client's final sign-off.	\$8,000	N/A	N/A

1	2.4				
Contracting	A written redline response of the contract will be sent back to Client within 10 business days after receipt of a redline from Client. Administrator will have 10 business days to provide a written responsive redline to each redline received from Client from the time Administrator receives all the necessary information from Client or Aon Hewitt in order to complete the written responsive redline to the time Administrator transmits the responsive redline to Aon Hewitt.	failed to meet this requirement to the extent and proportion that such failure is due to circumstances caused by Client, Aon Hewitt or other third party and/or is otherwise not within	\$8,000	N/A	N/A

EXHIBIT FIVE

CAVANAUGH
MACDONALD

ANALYSIS OF

PROJECTED COST

IMPACTS OF SETTLEMENT

PROVISIONS



Projected Cost Impact of Items Included in the Settlement Agreement

The City of Cincinnati and various plaintiff groups representing certain active and retired members of the Cincinnati Retirement System (CRS) have agreed on a list of items that would affect the provision of retiree health care and pension benefits. These items are listed below along with the effect of each item on the actuarial accrued liabilities of the Retiree Health Care and Pension Trusts. The impact on the funding ratios are also shown and reflect projected results as of the 12/31/2013 valuation.

Present Value of Future Benefits (PVFB): Value on a given date of the future payments expected to be paid to current retirees and current active members who are expected to become eligible for future retiree health care or pension benefits discounted to reflect the expected effects of the time value of money and probabilities of payment.

Actuarial Accrued Liability (AAL): The portion of the present value of future benefits that is expected to be paid in the future for current retirees, and a portion for current active members that is attributable to past service. <u>The AAL does not include liabilities for projected future accruals.</u>

Actuarial Value of Assets (AVA): The actuarial value of assets recognizes a portion of the difference between the market value of assets and the expected market value of assets, based on the assumed valuation rate of return. The amount recognized each year is 20% of the difference between market value and expected market value. Under this method, all investment gains and losses associated with a given year are recognized after five years. In addition, the actuarial value of assets cannot be less than 80% or more than 120% of the market value of assets.

Funding Ratio: The funding ratio is simply the AVA divided by either the AAL or the PVFB. The difference between the AAL and PVFB is the portion of liabilities allocated to the future service of active members funded via future normal cost contributions.

	Retiree Hea (\$ in milli		
	Inc	rease/(Decrease) in .	AAL
	Actives	Retirees	Total
	(\$19.5)	(\$58.3)	(\$77.8)
EGWP			
	(\$3.5)	(\$2.9)	(\$6.4)
MERP			
	(\$15.2)	\$0	(\$15.2)
New Eligibility			
Madical Dlan /Novy	(\$11.2)	\$0	(\$11.2)
Medical Plan/New Retirees	(\$11.2)	φU	(411.2)
Keurees			
Total*	(\$45.9)	(\$61.2)	(\$107.1)

^{*} Value of Plan changes are not additive of individual plan changes.

^{**} Assumes no change to any underlying assumptions from the 12/31/2013 valuation, including, but not limited to, discount rate, retiree health care benefit utilization, and pension benefit eligibility.

Retiree Health Care		
	<u>Funding R</u> 12/31	
	PVFB	AVA
Baseline as of 12/31/13	100.1%	109.1%
EGWP	114.5%	124.8%
MERP	101.3%	110.2%
New Eligibility	103.8%	111.8%
Medical Plan/New Retirees	102.3%	111.1%
Combined Changes w/No Asset Transfer*	122.8%	131.9%
Combined Changes w/Asset Transfer Out of \$215 Million 7/2016	90.0%	96.6%

^{*} Value of Plan changes are not additive of individual plan changes.

If all actuarial assumptions were realized, the projection results estimate a PVFB funding ratio of 86% in 2043 with an asset reduction of \$215 million in mid-2016 and if the plan was closed to new hires effective 1/1/2016. The future funding projections for Retiree Health Care will be highly sensitive to deviations from the actuarial assumptions such as medical inflation trends and investment returns. This sensitivity can be addressed through the development of a Retiree Health Care Funding & Benefits Policy.

^{**} Assumes no change to any underlying assumptions from the 12/31/2013 valuation, including, but not limited to, discount rate, retiree health care benefit utilization, and pension benefit eligibility.

	Pension in millions)		
	Increase	e/(Decrease) i	n AAL
	Actives	Retirees	Total
Change to 3% Simple COLA	\$25.2	(\$55.6)	(\$30.4)
3 Year COLA Suspension with one- time payment for Retiree Class in 2018 of 3% capped at \$1,000	(\$20.0)	(\$100.3)	(\$120.3)
Retirement Eligibility Changes	\$43.9	\$0	\$43.9
Increased Benefit Multiplier	\$5.3	\$0	\$5.3
Total*	\$48.1	(\$135.9)	(\$87.8)

^{*} Value of Plan changes <u>are not</u> additive of individual plan changes.

Pension	
	AAL Funding Ratio as of 12/31/13
Baseline as of 12/31/13	63.2%
Change to 3% Simple COLA	64.1%
3 Year COLA Suspension with one-time payment for Retiree Class in 2018 of 3% capped at \$1,000	66.8%
Retirement Eligibility Changes	62.0%
Increased Benefit Multiplier	63.1%
Combined Changes Before Asset Transfer*	65.8%
Combined Changes With Asset Transfer of \$215 Million 7/1/2016 and Lump Sum ERIP Payment of \$39.1 Million**	77.5%

^{*} Value of Plan changes are not additive of individual plan changes.

** Though the figures show the impact of the asset transfers (ERIP payment & \$215 million) as of the 12/31/2013 valuation, the true impact would be reflected on the 2015 and 2016 valuation results.

On a forward looking basis, and assuming all of the assumptions occur as expected, the Pension funding ratio is projected to reach 100% by 2043. This is due to the following three reasons:

- 1. As of the 12/31/2013 valuation, there are investment gains for actuarial smoothing still to be recognized.
- 2. The lower cost benefit structure of newer members will allow a greater portion of the fixed employer contribution rate to be applied towards paying down the unfunded actuarial liability in future years
- 3. The fixed employer contribution rate of 16.25% is higher than the projected annual required contribution in later years.

It is important to recognize that these figures and projections are based on many actuarial assumptions. They are likely to vary in future years to the extent that CRS actual experience varies from the expected actuarial assumptions.

The figures above were developed by the actuarial firm, Cavanaugh Macdonald. The firm is retained by the Cincinnati Board of Trustees.

EXHIBIT SIX

PROPOSED PRELIMINARY APPROVAL ORDER

UNITED STATES DISTRICT COURT SOUTHERN DISTRICT OF OHIO WESTERN DIVISION

NICK SUNYAK, et al.,

Case Nos.: 1:11-CV-445 and

1:12-cv-329

VS.

Judge Michael R. Barrett

CITY OF CINCINNATI, et al,

(City of Cincinnati Pension Litigation)

[PROPOSED]

ORDER GRANTING PRELIMINARY

APPROVAL OF CLASS ACTION

SETTLEMENT

WHEREAS, Current Employees Plaintiffs Nick Sunyak, Jeffery Harmon, Jill Allgeyer, Kim Kappel, Waleia Jackson, Finley Jones and Richard Ganulin, and Retiree Plaintiffs Thomas A. Gamel, Sr., Paul Smith, Mark K. Jones, Dennis Davis, Ely Ryder, and Ann DeGroot (collectively the "Named Plaintiffs"), individually and on behalf of two proposed Classes, and Defendants the City of Cincinnati, the Mayor of Cincinnati, the City Manager, the Vice-Mayor, the City Council Members, the Cincinnati Retirement System ("CRS"), and the appointed Board of Trustees of the CRS ("Board") have entered into a Collaborative Settlement Agreement (the "Agreement") intended to resolve certain litigation, including litigation pending in this Court; and

WHEREAS, the Settlement Agreement, together with supporting materials, sets forth the terms and conditions for the proposed settlement;

WHEREAS, the Court has before it the Parties' Motion for Preliminary Approval of Settlement and Memorandum in Support of Motion for Preliminary Approval of Settlement, together with the Agreement and related materials; and

WHEREAS, the Court is satisfied that the terms and conditions set forth in the Agreement and exhibits attached thereto were the result of good faith, arm's length settlement negotiations between competent and experienced counsel for both Named Plaintiffs and Defendants.

IT IS HEREBY ORDERED	this day of	, 2015, as follows:
II IO III IO I OID BILES		

- 1. The terms of the Agreement including all exhibits are hereby conditionally approved, subject to further consideration thereof at the Fairness Hearing provided for below.
- 2. The Agreement is adopted by the Court and made part of this Order as if set out in full herein.
- 3. The Agreement and the terms contained therein are hereby preliminarily approved as fair, reasonable, adequate, and in the best interests of the Current Employees Class and the Retirees Class.
- 4. The Court approves the proposed Notice Program set forth in Section 38 of the Agreement. The Notice Program is reasonably calculated to apprise Class Members of their right to object, constitutes due, adequate, and sufficient notice to all persons entitled to receive notice, is the best notice practicable under the circumstances, and meets all applicable requirements of the Federal Rules of Civil Procedure, the Class Action Fairness Act, the United States Constitution (including the Due Process Clause), the Rules of the Court, and any other applicable law. Subject to amendment if the need arises, the Notice Program shall be initiated within thirty (30) days of this Preliminary Approval Order and executed as set forth in the Settlement Agreement.
- 5. Pursuant to Federal Rule of Civil Procedure 23(a), (b)(1) and (b)(2), and for purposes of settlement only, the Court makes the following preliminary findings of fact and

conclusions of law:

- a. The Current Employees Class and the Retirees Class (as defined, respectively, in the Agreement) are sufficiently definite and identifiable;
- b. The Current Employees Class and the Retirees Class are so numerous that joinder of all Members is impracticable;
- c. There are questions of law and/or fact common within the Current Employees Class including but not limited to: (1) whether the Current Employees Class was fully vested in their CRS benefits on July 1, 2011; (2) whether Defendants improperly revoked and/ or impaired the Current Employees Class' vested CRS benefits when it enforced Ordinance No. 84-2011; (3) whether Defendants impaired contractual rights of the Current Employees Class when they enforced Ordinance No. 84-2011; (4) whether Defendants are estopped from enforcing Ordinance No. 84-2011 so as to prevent the revocation and/or impairment of the contractual rights of the Current Employees Class; and (5) whether Defendants' enforcement of Ordinance No. 84-2011 operated as an unconstitutional taking of the vested property interest of the Current Employees Class.
- d. There are questions of law and/or fact common within the Retirees Class including but not limited to: (1) whether Defendants' offer of retirement benefits to the Retirees Class created a fundamental property right, giving each of them a vested right in those retirement benefits which cannot be reduced, impaired, revoked, or eliminated; (2) whether Defendants' actions as explained in the Retirees Class Complaint constitute an unlawful taking of the Retirees Class Members' property rights in violation of the United States Constitution and/or the Ohio Constitution; (3) whether the Defendants have a contractual obligation to provide the Retirees Class with certain retirement benefits,

which cannot now or afterwards be reduced, impaired, revoked, or eliminated; (4) whether the unilateral reduction, impairment, revocation, and/or elimination of the Retirees Class Members' retirement benefits constitutes a breach of the Defendants' fiduciary duty; and (5) whether the Defendants are estopped from reducing, impairing, revoking, or eliminating the retirement benefits owed to the Retirees Class;

- e. The Current Employees Plaintiffs' claims are typical of the claims of the Members of the Current Employees Class as all subgroups were represented and no conflict exists between or among the subgroups, and the Retiree Plaintiffs' claims are typical of the claims of the Members of the Retirees Class;
- f. Current Employees Plaintiffs and the Current Employees Class Counsel have and will fairly and adequately represent and protect the interests of the Current Employees Class, and the Retiree Plaintiffs and the Retirees Class Counsel have and will fairly and adequately represent and protect the interests of the Retirees Class;
- g. Current Employees Plaintiffs' interests do not conflict with the interests of the Current Employees Class in the maintenance of this action and this Settlement, and the Retiree Plaintiffs' interests do not conflict with the interests of the Retirees Class in the maintenance of this action and this Settlement;
- h. The questions of law and/or fact common to the Current Employees Class and those common to the Retirees Class predominate over the questions affecting only individual members of those Classes;
- j. Certification of the Current Employees Class and the Retirees Class is appropriate because prosecuting separate actions by individual Members of these Classes would create a risk of inconsistent and varying adjudications with respect to individual

Members of the Classes that would establish incompatible standards of conduct for the Defendants;

- k. Certification of the Current Employees Class and the Retirees Class is appropriate because adjudications with respect to individual Members of the Classes, as a practical matter, would be dispositive of the interests of the other Members not parties thereto and would substantially impair or impede their ability to protect their interests; and
- I. Certification of the Current Employees Class and the Retirees Class is appropriate because the Defendants have acted or refused to act on grounds that apply generally to the Classes, so that final injunctive relief or corresponding declaratory relief as agreed to by the Parties is appropriate respecting the Classes as a whole.
- 6. Pursuant to Fed. R. Civ. P. 23 and for purposes of Settlement, the Court appoints the Named Plaintiffs as Class Representatives and conditionally certifies the following Classes:

Current Employees Class: All individuals (and/or their Dependents or Surviving Beneficiaries) who participated in the Cincinnati Retirement System with at least five years of creditable service and who were actively employed or otherwise qualified for benefits on July 1, 2011, and who are Members of "Group C," "Group D," "Group E," or "Group F" as these terms are defined by Cincinnati Municipal Code § 203-1-Ml (b), (c), (d), and (e).

Retirees Class: All individuals (and/or their Dependents or Surviving Beneficiaries) formerly employed by the City of Cincinnati, the University of Cincinnati, the University Hospital f/k/a General Hospital and Hamilton County, who retired on or before July 1, 2011 and have received retirement benefits from the City of Cincinnati and their Dependents and/or their Surviving Beneficiaries who are entitled to those benefits.

7. Pursuant to Fed. R. Civ. P. 23(g) the Court appoints Marc D. Mezibov, Robert D. Klausner, Jeffrey S. Goldenberg, and Christian A. Jenkins as Class Counsel for the Current

¹ The Current Employees Class also includes City of Cincinnati employees who had at least five years of creditable service prior to July 1, 2011 and who retired after July 1, 2011, as well as veterans who purchase service credit sufficient to satisfy the five years of service requirement prior to July 1, 2011.

Employees Class and Robert A. Pitcairn, Jr., James F. McCarthy, III, Peter O'Shea, and the law firm of Katz, Teller, Brant & Hild as Class Counsel for the Retirees Class. Class Counsel shall submit their applications for attorney fees and expenses no later than twenty-one (21) days prior to the date for Class Members to submit objections.

- 8. The City is authorized to retain Class Action Administration, Inc. as the Settlement Administrator to perform all functions and duties assigned to the Settlement Administrator in the Agreement, the cost of which shall be reimbursed by Defendant City of Cincinnati.
- 9. The Court directs the parties and Class Action Administration, Inc. to implement the Notice Program and to disseminate and/or publish the Notice referenced in Section 38 of the Agreement in accordance with this Order and the Agreement.
- adequacy of the Agreement, or to the request for attorneys' fees and expense reimbursement, must file with the Clerk of the Court and serve on designated Counsel, within 60 days of the Notice Date, a written statement of the objection as well as the specific reason(s), if any, for the objection, including any legal support the Class Member wishes to bring to the Court's attention and any evidence the Class Member wishes to introduce in support of the objection. Class Members may so object either on their own or through an attorney hired at their own expense.

Any attorney hired by a Class Member at that Class Member's expense for the purpose of objecting to the fairness, reasonableness, or adequacy of the Agreement, to any terms of the proposed Settlement, or to proposed attorneys' fees and expenses shall file with the Court and deliver to designated Counsel a Notice of Appearance no later than 60 days from the Notice Date.

Additionally, Class Members and/or their attorneys intending to make an appearance at the Fairness Hearing must by no later than 14 days prior to the Fairness Hearing:

- a. File a notice of intention to appear, with the Clerk of the Court, that contains the Class Member's and/or their attorney's name, address, and telephone number, as well as a description of all evidence the Class Member and/or Class Member's attorney will seek to introduce at the Fairness Hearing, including all documents to be introduced and witnesses to be called; and
- b. Serve a copy of such notice of intention to appear on counsel for the Parties as described in the Notice.

Any Class Member who files and serves a written objection in accordance with the procedure set forth above and in the Section 39 of the Settlement Agreement may appear at the Fairness Hearing to object to any aspect of the fairness, reasonableness, or adequacy of the Settlement. Class Members and/or their attorneys who do not timely comply with the procedures set forth above shall not be heard at the Fairness Hearing and waive any objection to the Settlement Agreement.

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representatives of their respective Class; (e) finally approve and appoint Current Employees

Class Counsel and Retirees Class Counsel to represent their respective Class; (f) determine and

approve the payment of reasonable attorneys' fees and expense reimbursements for Class

Counsel; (g) order the Settlement Administrator to process all payments due to Class Members

under the Agreement; and (h) order that the claims at issue in this litigation are fully and finally

resolved as of the date of Finality, as defined in the Agreement, and that Current Employees

Plaintiffs, the Current Employees Class, the Retirees Plaintiffs, and the Retirees Class are forever

barred and enjoined from filing, commencing, prosecuting, intervening in, participating in (as

Class Members or otherwise), or receiving any benefits or other relief from, any other lawsuit,

arbitration, or administrative, regulatory, or other proceeding or order in any jurisdiction based

on the claims at issue in this litigation, except as set forth in the Re-Opener provisions in the

Agreement and Consent Decree.

11 19 20 OKDEKED	IT	IS	SO	ORDERED
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Dated: _____

The Honorable Michael Barrett United States District Judge

5579117.2

KTBH: 4822-9768-6819, v. 2

8

EXHIBIT SEVEN

PROPOSED ORDER GRANTING FINAL APPROVAL

UNITED STATES DISTRICT COURT SOUTHERN DISTRICT OF OHIO WESTERN DIVISION

NICK SUNYAK, et al., : Case Nos.: 1:11-cv-445 and

1:12-cv-329

vs.

Judge Michael R. Barrett

CITY OF CINCINNATI, et al,

[PROPOSED]

(City of Cincinnati Pension Litigation)

ORDER GRANTING FINAL APPROVAL OF CLASS

ACTION SETTLEMENT

WHEREAS, this Court granted Preliminary Approval of the Class Action Settlement ("Settlement") of these Actions on _______, 2015 (Doc. #___).

WHEREAS, the Current Employees Plaintiffs (Nick Sunyak, Jeffery Harmon, Jill Allgeyer, Kim Kappel, Waleia Jackson, Richard Ganulin, and Finley Jones), the Retiree Plaintiffs (Thomas A. Gamel, Sr., Paul Smith, Mark K. Jones, Dennis Davis, Ely Ryder, and Ann DeGroot), the American Federation of State and Municipal Employees Ohio Council No. 8 ("AFSCME"), and the Defendants (The City of Cincinnati, Mayor John Cranley, City Manager Harry Black, Vice-Mayor David Mann, Cincinnati City Council Members, the Cincinnati Retirement System, and the Board of Trustees of the Cincinnati Retirement System, (collectively, the "Parties") have filed a motion seeking final approval of this Settlement ("Motion") (Doc. #__);

WHEREAS, the Parties appeared with their attorneys of record at a Fairness Hearing on _______, 2015 after all members of the Classes were given an opportunity to be heard in accordance with the Court's Preliminary Approval Order, and the Court has given due consideration to the Parties' Collaborative Settlement Agreement, including all attached exhibits and related materials, the Parties' Motion for Final Approval, including the attached

Memorandum and all other papers filed in support, all objections to the Settlement, the complete record in this litigation, the information and arguments presented at the _______, 2015 Fairness Hearing, and all other materials relevant to this matter including the Declaration of the Settlement Administrator on Implementation and Adequacy of Settlement Notice Program as well as the Declaration of ______ (insert name of actuarial expert) concerning the impact and benefits of this Settlement on the Cincinnati Retirement System ("CRS") and the members of the Classes;

WHEREAS, the Court recognizes that the Parties have litigated complex questions about the management of the CRS – and the respective rights of plan participants – for nearly five years, including issues related to benefits levels, eligibility requirements, healthcare packages, and funding mechanisms, and that while some of these lawsuits have been subject to conclusive appellate rulings, many pertinent legal and factual questions remain; and

WHEREAS, the Court is satisfied that the terms and conditions set forth in the Collaborative Settlement Agreement and related Consent Decree were the result of good faith, arm's length settlement negotiations between competent and experienced counsel for the Current Employees Plaintiffs, the Current Employees Class, the Retiree Plaintiffs, the Retirees Class, and Defendants;

NOW, THEREFORE, IT IS ORDERED THAT:

1. This Order approves, adopts, and incorporates by reference in their entirety the Collaborative Settlement Agreement and the Consent Decree which are attached hereto as Exhibits 1 and 2 respectively. The Collaborative Settlement Agreement and the Consent Decree are made part of this Order as if set out in full herein and shall be fully enforceable by this Court. Accordingly, the Parties are ordered to implement and comply with all the terms of the

Collaborative Settlement Agreement and the Consent Decree.

- 2. For purposes of this litigation, the Court has subject matter and personal jurisdiction over the Parties, including all Class Members, and has the power and authority to approve the Collaborative Settlement Agreement and Consent Decree, including all Exhibits thereto.
- 3. Pursuant to Federal Rules of Civil Procedure 23(a), 23(b)(1), and 23(b)(2), the Court confirms its previous certification, and for purposes of effectuating the Settlement, grants final approval to the following two Classes:

Current Employees Class: All individuals (and/or their Dependents or Surviving Beneficiaries) who participated in the Cincinnati Retirement System with at least five years of creditable service and who were actively employed or otherwise qualified for benefits on July 1, 2011, and who are Members of "Group C," "Group D," "Group E," or "Group F" as these terms are defined by Cincinnati Municipal Code § 203-1-Ml (b), (c), (d), and (e).

Retirees Class: All individuals (and/or their Dependents or Surviving Beneficiaries) formerly employed by the City of Cincinnati, the University of Cincinnati, the University Hospital f/k/a General Hospital and Hamilton County, who retired on or before July 1, 2011 and have received retirement benefits from the City of Cincinnati and their Dependents and/or their Surviving Beneficiaries who are entitled to those benefits.

- 4. Pursuant to Federal Rule of Civil Procedure 23(a), (b)(1), and (b)(2), and for purposes of settlement only, the Court makes the following findings of fact and conclusions of law:
 - a. The Current Employees Class and the Retirees Class are sufficiently definite and identifiable;
 - b. The Current Employees Class and the Retirees Class are so numerous that joinder of all members is impracticable;

¹ The Current Employees Class also includes City of Cincinnati employees who had at least five years of creditable service prior to July 1, 2011 and who retired after July 1, 2011, as well as veterans who purchase service credit sufficient to satisfy the five years of service requirement prior to July 1, 2011.

- c. There are questions of law and/or fact common within the Current Employees Class including but not limited to: (1) whether the members of the Current Employees Class were fully vested in their CRS benefits on July 1, 2011; (2) whether Defendants improperly revoked and/ or impaired Current Employees Class Members' vested CRS benefits when they enforced Ordinance No. 84-2011; (3) whether Defendants impaired contractual rights of the Current Employees Class when they enforced Ordinance No. 84-2011; (4) whether Defendants are estopped from enforcing Ordinance No. 84-2011 so as to revoke and/or impair the employment agreement with Current Employees Class Members; and (5) whether Defendants' enforcement of Ordinance No. 84-2011 operated as an unconstitutional taking of the vested property interest of Current Employees Class Members;
- d. There are questions of law and/or fact common within the Retirees Class including but not limited to: (1) whether Defendants' offer of retirement benefits to the Retirees Class Members created a fundamental property right, giving each of them a vested right in those retirement benefits which cannot be reduced, impaired, revoked, or eliminated; (2) whether Defendants' actions as explained in the Retirees Class Complaint constitute an unlawful taking of the Retirees Class Members' property rights in violation of the United States Constitution and/or the Ohio Constitution; (3) whether the Defendants have a contractual obligation to provide the Retirees Class Members with certain retirement benefits, which cannot now or afterwards be reduced, impaired, revoked, or eliminated; (4) whether the unilateral reduction, impairment, revocation, and/or elimination of the Retirees Class Members' retirement benefits constitutes a breach of the Defendants' fiduciary duty; and (5) whether the Defendants are estopped

from reducing, impairing, revoking, or eliminating the retirement benefits owed to the Retirees Class;

- e. The Current Employees Plaintiffs' claims are typical of the claims of the Members of the Current Employees Class as all subgroups were represented and no conflict exists between or among the subgroups, and the Retirees Plaintiffs' claims are typical of the claims of the Members of the Retirees Class;
- f. Current Employees Plaintiffs and the Current Employees Class Counsel have and will fairly and adequately represent and protect the interests of the Current Employees Class, and the Retiree Plaintiffs and the Retirees Class Counsel have and will fairly and adequately represent and protect the interests of the Retirees Class;
- g. Current Employees Plaintiffs' interests do not conflict with the interests of the Current Employees Class in the maintenance of this action and this Settlement, and the Retiree Plaintiffs' interests do not conflict with the interests of the Retirees Class in the maintenance of this action and this Settlement;
- h. Certification of the Current Employees Class and the Retirees Class is appropriate because prosecuting separate actions by individual members of these Classes would create a risk of inconsistent and varying adjudications with respect to individual members of the Classes that would establish incompatible standards of conduct for the Defendants;
- i. Certification of the Current Employees Class and the Retirees Class is appropriate because adjudications with respect to individual members of the Classes, as a practical matter, would be dispositive of the interests of the other Members not parties thereto and would substantially impair or impede their ability to protect their interests;

and

- j. Certification of the Current Employees Class and the Retirees Class is appropriate because the Defendants have acted or refused to act on grounds that apply generally to the Classes, so that final injunctive relief or corresponding declaratory relief as agreed to by the Parties is appropriate respecting the Classes as a whole.
- 5. The Collaborative Settlement Agreement, the Consent Decree, and the terms contained therein are hereby finally approved as fair, reasonable, adequate, in the best interests of the Current Employees Class and the Retirees Class, and in compliance with all applicable requirements of the Federal Rules of Civil Procedure, the Class Action Fairness Act, the United States Constitution (including the Due Process Clause), the Rules of the Court, and any other applicable law.
- 6. Pursuant to Rule 23(g), the following are hereby finally designated and approved as Current Employees Class Counsel: (1) Christian A. Jenkins, Esq., Minnillo & Jenkins, Co. LPA, 2712 Observatory Avenue, Cincinnati, Ohio 45208; (2) Marc D. Mezibov, Esq., Law Office of Marc Mezibov, 401 E. Court Street, Suite 600, Cincinnati, OH 45202; (3) Jeffrey S. Goldenberg, Esq., Goldenberg Schneider, LPA, One West Fourth Street, 18th Floor, Cincinnati, Ohio 45202; and (4) Robert D. Klausner, Esq., Klausner, Kaufman, Jensen & Levinson, 10059 Northwest 1st Court, Plantation, FL 33324. The Court's designation and approval of Current Employees Class Counsel is based upon: (1) the work they have done to identify and investigate the claims in this litigation; (2) their experience handling class actions and other complex litigation, including employee benefits litigation; (3) their knowledge of the applicable law and their familiarity with the complexities of this type of pension benefits litigation; and (4) the resources they committed and are willing to continue to commit to this litigation and the

implementation of the Consent Decree going forward.

- 7. Robert A. Pitcairn, Esq., James F. McCarthy, III, Esq., and Peter O'Shea, Esq. of the law firm of Katz Teller Brant & Hild, 255 East Fifth Street, Suite 2400, Cincinnati, Ohio, 45202 are hereby finally designated and approved as Retirees Class Counsel pursuant to Rule 23(g). The Court's designation and approval of Retirees Class Counsel is based upon: (1) the work performed to identify and investigate the claims in this litigation; (2) their experience handling class actions and other complex litigation, including employee benefits litigation; (3) their knowledge of the applicable law and their familiarity with the complexities of this type of pension benefits litigation; and (4) the resources they committed and are willing to continue to commit to this litigation and the implementation of the Consent Decree going forward.
- 8. Nick Sunyak, Jeffery Harmon, Jill Allgeyer, Kim Kappel, Waleia Jackson, Richard Ganulin, and Finley Jones are designated and granted final approval as the Current Employees Class Representatives. Jill Allgeyer is designated and granted final approval as the Sub-Class C representative. Kim Kappel, Waleia Jackson, and Richard Ganulin are designated and granted final approval as the Sub-Class D representatives. Finley Jones is designated and granted final approval as the Sub-Class E representative. Jeffrey Harmon and Nick Sunyak are designated and granted final approval as the Sub-Class F representatives.
- 9. Thomas A. Gamel, Sr., Paul Smith, Mark K. Jones, Dennis Davis, Ely Ryder, and Ann DeGroot are finally designated and approved as the Retirees Class Representatives.
- 10. The Parties have provided direct mail notice to the Classes in a manner consistent with the Order Granting Motion for Preliminary Approval of Class Action Settlement. The Notice Plan, as implemented, satisfied the requirements of due process and was the best notice practicable under the circumstances. The Notice Plan was reasonably calculated, under the

circumstances, to apprise Class Members of the terms of the proposed Settlement, their right to object or exclude themselves from the proposed Settlement, and their right to appear at the Fairness Hearing. Further, the notice was reasonable and constituted due, adequate, and sufficient notice to all persons entitled to receive notice. Also, Defendants, through Class Action Administration, Inc., notified the appropriate federal official (the Attorney General of the United States) and the appropriate State of Ohio officials (the Auditor of the State of Ohio and the Attorney General of the State of Ohio) pursuant to the Class Action Fairness Act of 2005 ("CAFA"), 28 U.S.C. § 1715. Accordingly, the Defendants' notification complies fully with its obligations under CAFA, and the notice met all applicable requirements under the Federal Rules of Civil Procedure, the United States Constitution (including the Due Process Clause), the Rules of the Court, and any other applicable rule or law.

- 11. Class Action Administration, Inc. ("Settlement Administrator") was retained to disseminate the Notice Plan in accordance with the terms of the Collaborative Settlement Agreement and the Court's Order Granting Motion for Preliminary Approval of Class Action Settlement. It is apparent from the Declaration of ______ that the Notice Plan was properly implemented and was effective.
- 12. The Court has determined that notice and full opportunity has been given to the Classes to object to the terms of the Settlement. The Court also has determined that notice and full opportunity has been given to the Classes to object to Current Employees Class Counsel's and Retirees Class Counsel's request for attorneys' fees and expenses. The Court has considered all of the objections to the Settlement that were submitted by members of the Classes as well as Class Counsel and Defendants' responses to those objections, and has determined as follows:
 - a. the Current Employees Plaintiffs, the Retiree Plaintiffs, the Current

Employees Class, and the Retirees Class face significant risks if this litigation were to proceed, including the real possibility of losing this litigation;

- b. the possibility of a greater ultimate recovery is highly speculative and any such recovery would only occur after considerable delay, if at all;
- c. the terms of the Collaborative Settlement Agreement and Consent Decree provide substantial and meaningful benefits to the Classes;
- d. the Collaborative Settlement Agreement and Consent Decree are the product of vigorous, highly-contested litigation that included meaningful investigation into the facts and the law underlying the claims at issue;
- e. the Settlement occurred after the litigation was substantially developed, including the exchange of voluminous actuarial data and other information during the mediation process and due diligence following the execution of the Memorandum of Understanding on December 31, 2014;
- f. the Settlement negotiations were extensive, arms-length, and under the direction of the Court through a collaborative and agreed-to process that occurred without any collusion;
- g. the reaction by the Classes has been overwhelmingly in favor of the Settlement; and
 - h. experienced Class Counsel support the Settlement.
- 13. Accordingly, having considered the foregoing, the costs and risks and delays of continued litigation versus the benefits provided by the Settlement, and based on this Court's knowledge of the Actions, the Court finds and concludes that the Settlement is in the best interests of the Classes and is fair, reasonable, and adequate to all members of the Classes.

- 14. This Settlement, including the terms of the Collaborative Settlement Agreement and Consent Decree, is accordingly granted final approval and is confirmed as fair, reasonable, adequate, and binding upon all members of the Classes.
- 15. The Parties are hereby directed to proceed with and complete the implementation of the Settlement. Therefore, the Court hereby orders and directs the Parties and their counsel to proceed with and complete the implementation and consummation of this Collaborative Settlement Agreement and Consent Decree according to its terms and provisions.
- 16. The Court enters judgment in accordance with the Collaborative Settlement Agreement and further declares the Collaborative Settlement Agreement binding on all the Parties.
- 17. Except as provided in the Collaborative Settlement Agreement and Consent Decree, all Parties are barred, estopped, and enjoined from asserting claims or interests arising under or out of, in connection with, or in any way relating to the claims set forth in the Litigation as defined in the Collaborative Settlement Agreement ("Barred Claims").
- 18. AFSCME warrants and acknowledges that it will dismiss with prejudice *State ex rel. Council 8 AFSCME*, et al. v. City of Cincinnati, et al., Case No. A 1 104791, within 10 days of Finality as defined in the Collaborative Settlement Agreement pursuant, and to the terms of the separate settlement agreement entered into between AFSCME and the Defendants.
- 19. All Parties are bound by this Order Granting Final Approval, the Collaborative Settlement Agreement, and the Consent Decree. The Court declares that the Collaborative Settlement Agreement and related Consent Decree are incorporated into this Order Granting Final Approval, each of which shall be binding on all Parties. Further, the Collaborative Settlement Agreement, the Consent Decree, and this Order shall be preclusive for the 30 years

following the Effective Date, as defined in the Collaborative Settlement Agreement, in all other pending and future lawsuits or other proceedings relating to the Barred Claims in these Actions.

- 20. Consistent with the above paragraph, the Court also orders that the Barred Claims are fully and finally resolved as of the date of Finality, as defined in the Collaborative Settlement Agreement, and that the City, CRS and related City Defendants are forever discharged and released from the Barred Claims and that the Current Employees Plaintiffs, the Retiree Plaintiffs, and the members of the Classes are permanently barred and enjoined from filing, commencing, prosecuting, intervening in, participating in (as Class Members or otherwise), or receiving any benefits or other relief from, any other lawsuit, arbitration, or administrative, regulatory, or other proceeding or order in any jurisdiction based on the Barred Claims, except as set forth in the Re-Opener provisions in the Collaborative Settlement Agreement and Consent Decree.
- 21. The Court, having considered the request of Current Employees Class Counsel for an award of attorneys' fees and reimbursement of expenses, hereby grants the request and awards Current Employees Class Counsel attorneys' fees in the sum of \$_____ which amount the Court concludes is fair and reasonable in light of the estimated \$_____ valuation of the benefits to the Current Employees Class resulting from this Settlement. The Court also approves and grants Current Employees Class Counsels' request for expense reimbursement in the amount of \$_____ for their reasonable expenses incurred in prosecuting this action and in implementing this Settlement. The fees shall be paid be paid by

^{22.} The Court, having considered the request of Retirees Class Counsel for an award of attorneys' fees and reimbursement of expenses, hereby grants the request and awards Retirees Class Counsel attorney's fees in the sum of \$_____ which amount the Court concludes is

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fair and reasonable. The Court also approves and grants Retirees Class Counsels' request for expense reimbursement in the amount of \$_____ for their reasonable expenses incurred in

prosecuting this action and in implementing this Settlement. The fees shall be paid by _____.

23. Without affecting the finality of this Final Order for purposes of appeal, if any,

the Court retains continuing and exclusive jurisdiction over the Parties for thirty years following

the Effective Date as to all matters relating to the administration, consummation, enforcement,

and interpretation of the Collaborative Settlement Agreement, the Consent Decree, and this

Order Granting Final Approval, and for any other necessary purpose related thereto, including

the entry of any additional orders as may be necessary and appropriate.

IT IS SO ORDERED.

DATED:	, 2015	
		The Henouskie Mic

The Honorable Michael R. Barrett United States District Judge

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EXHIBIT EIGHT

CURRENT CRS "POINT SYSTEM"

CINCINNATI RETIREMENT SYSTEM

2015 Retiree Healthcare Point System Matrix for Members Hired On or After January 9,1997

Points Full Years of Membership Service + Full Years of Age at Termination =

	cent of Month	reicent of Monthly neathicate riennum categories	eilluili categoi	<u>c</u>	
	90 + Points	80-89 Points	70-79 Points	60-69 Points	< 60 Points
	2%	25%	20%	%5/	75%
Tier of Coverage					
Retiree only - Medicare Eligible	\$23.73	\$118.66	\$237.33	\$355.99	\$355.99
Retiree only - Non Medicare	\$47.05	\$235.25	\$470.50	\$705.74	\$705.74
	(, ,	0000	(
Ketiree & Spouse - Medicare Eligible	\$46.23	\$231.14	\$462.29	\$693.43	Not Available
Retiree & Spouse - 1 Medicare	\$69.55	\$347.73	\$695.45	\$1,043.18	Not Available
Retiree & Spouse - Non Medicare	\$92.86	\$464.31	\$928.62	\$1,392.92	Not Available
Retiree & Child(ren) - Medicare Eligible	\$37.68	\$188.41	\$376.83	\$565.24	Not Available
Retiree & Child(ren) - Non Medicare	\$61.00	\$305.00	\$610.00	\$914.99	Not Available
Retiree & Spouse & Child(ren) - Medicare Eligible	\$65.04	\$325.18	\$650.37	\$975.55	Not Available
Retiree & Spouse & Child(ren) -1 Medicare Eligible	\$88.35	\$441.77	\$883.53	\$1,325.30	Not Available
Retiree & Spouse & Child(ren) -Non Medicare	\$111.67	\$558.35	\$1,116.70	\$1,675.05	Not Available