

South Dakota Department of Social Services

Medicaid P&T Committee Meeting

December 12, 2014





DEPARTMENT OF SOCIAL SERVICES

MEDICAL SERVICES
700 Governors Drive
Pierre, South Dakota 57501-2291
(605) 773-3495
FAX (605) 773-5246

**SOUTH DAKOTA
MEDICAID P&T COMMITTEE MEETING
AGENDA**

**Friday, December 12, 2014
1:00 – 3:00 PM**

**DDN Locations:
Sioux Falls
University Center
Room FADM253
4801 North Career Avenue**

**Pierre
Capitol Building
DDN Room A
500 E Capitol**

**Rapid City
SDSMT
Room CB109
501 E Joseph St.**

Call to Order

Approval of Minutes of Previous Meeting

Prior Authorization Update

Review of Top 15 Therapeutic Categories/Top 25 Drugs

Review of Drug Spend

Patent Expirations

Old Business

**Hepatitis C
Stimulant use in adults
Evzio
Otezla**

New Business

**High cost drugs
GLP-1 receptor agonists
Topical therapies for onychomycosis**

Oral Presentations and Comments by Manufacturers' Representatives

Next Meeting Date/Adjournment

**Minutes of the September 26, 2014
Pharmacy & Therapeutics (P&T) Committee Meeting
South Dakota Department of Social Services, Division of Medical Services**

Members Present

Richard Holm, MD; Bill Ladwig, RPh; Dana Darger, RPh; Mikel Holland, MD; James Engelbrecht, MD; Lenny Petrik; Michelle Baack, MD; Tim Soundy

DSS Staff Present

Mike Jockheck, RPh; Ann Schwartz, Dep. Director of Medical Services; Kirby Stone, Director of Medical Services

Administrative Business

The P&T meeting was called to order by D. Darger at 1:00 p.m. The minutes of the June 6, 2014 meeting were presented. M. Baack made a motion to approve. R. Holm seconded the motion. The motion was approved unanimously.

Prior Authorization Update and Statistics

The committee reviewed the prior authorization (PA) activity for August 2014. There were a total of 3,026 PAs processed in the month of August, with 99.70% of those requests responded to in less than eight hours. There were 2,344 (77%) requests received electronically and 682 (23%) requests received by fax.

Analysis of the Top 15 Therapeutic Classes

The committee reviewed the top 15 therapeutic classes by total cost of claims from 04/1/2014 – 06/30/2014. The top five classes were antipsychotics, respiratory and CNS stimulants, amphetamines, central nervous system agents, misc., and corticosteroids (respiratory tract). The top 15 therapeutic classes make up 40.41% of total claims. The committee also reviewed the top 25 drugs based on total claims cost and number of claims. The top 25 drugs by claims cost make up 13.48% of total claims.

Treatment for Hepatitis C Review

The committee reviewed utilization information, PA forms, and treatment agreements for hepatitis C therapy. A motion was made by M. Baack to accept the Treatment Agreement on page 14 of the P&T committee pack with the addendum, "I agree to have a blood test to monitor my viral load during and after treatment." R. Holm seconded the motion. The motion was approved unanimously. The committee requested a box be added to the Hepatitis C PA forms asking if the patient is treatment naïve. The committee also suggested that all combination therapy requests and all retreatment therapy requests be reviewed by the medical director. M. Puryear, representing Gilead, spoke regarding Sovaldi. Jennifer Stoffel, representing Janssen, spoke regarding Olysio. A motion was made by R. Holm that all new drugs used to treat hepatitis C will automatically require prior authorization. M. Baack seconded the motion. The motion was approved unanimously. A motion was made by B. Ladwig to include baseline HCV RNA and HCV RNA 4 weeks after treatment (SVR4) to the PA forms. R. Holm seconded the motion. The motion was approved unanimously.

Stimulant Use in Adults

The committee reviewed stimulant use in adults at the June meeting. It was requested that a review of the top prescribers be brought to the September meeting. J. Engelbrecht made a motion that a form be developed for stimulant use in adults. B. Ladwig seconded the motion. The motion was approved unanimously. A form will be brought to the December meeting for committee review.

Oral Allergen Extracts (Ragwitek, Grastek, Oralair)

The committee reviewed the oral allergen extracts prior authorization form. M. Baack made a motion to approve the form. R. Holm seconded the motion. The motion was approved unanimously.

Intuniv Review

The committee reviewed Intuniv clinical information. There was no public comment. This topic was tabled.

Transdermal Androgens Review

The committee reviewed transdermal androgens clinical information. There was no public comment. This topic was tabled.

Phosphate Binders Review

The committee reviewed phosphate binders clinical information. There was no public comment. This topic was tabled.

Zontivity Review

The committee reviewed Zontivity clinical information. There was no public comment. This topic was tabled.

Evzio Review

The committee reviewed Evzio clinical information. There was no public comment. B. Ladwig made a motion to place Evzio on prior authorization. T. Soundy seconded the motion. The motion was approved unanimously. A prior authorization form will be brought to the December meeting for committee review.

Otezla Review

The committee reviewed Otezla clinical information. There was no public comment. J. Engelbrecht made a motion to place Otezla on prior authorization. R. Holm seconded the motion. The motion was approved unanimously. A prior authorization form will be brought to the December meeting for committee review.

The next meeting is scheduled for December 12, 2014. R. Holm made a motion to adjourn the P&T Committee meeting. L. Petrik seconded the motion. The motion passed unanimously and the meeting was adjourned.



**South Dakota Medicaid
Monthly Prior Authorization Report
October 1, 2014 – October 31, 2014**

Time Ratio

| Total PAs | Response Under 8 Hours | Response Over 8 Hours | % Under 8 Hours | % Over 8 Hours |
|-----------|------------------------|-----------------------|-----------------|----------------|
| 3,409 | 3,387 | 22 | 99.35% | 0.65% |

By Form Type

| Form Type | Description | Approve | Deny |
|---------------|----------------------------|------------|-------------|
| ADP | Antidepressant | 163 | 348 |
| AFX | Amrix and Fexmid | 0 | 4 |
| ALT | Altabax | 1 | 8 |
| AMB | Ambien CR | 6 | 8 |
| ANF | Anti-Infectives | 0 | 75 |
| ANT | Antihistamines | 8 | 52 |
| APS | Antipsychotic | 444 | 408 |
| ARB | ARBS | 2 | 9 |
| COA | Oral Anticoagulants | 3 | 5 |
| DAW | Dispense As Written | 7 | 4 |
| EME | Antiemetics | 0 | 17 |
| FUN | Antifungals | 0 | 1 |
| GRH | Growth Hormone | 11 | 2 |
| GSM | Genitourinary SMR | 9 | 28 |
| HEP | Hepatitis Meds | 2 | 13 |
| HLM | Head Lice Medication | 10 | 67 |
| LID | Lidoderm | 1 | 101 |
| MAX | Max Units Override | 83 | 799 |
| NAR | Name Brand Narcotics | 2 | 4 |
| NUC | Opioids | 8 | 37 |
| ONF | Onfi | 8 | 12 |
| OPH | Ophthalmic Antihistamines | 1 | 26 |
| PPI | Proton Pump Inhibitors | 47 | 86 |
| SMR | Skeletal Muscle Relaxants | 2 | 19 |
| STE | Nasal Steroids | 9 | 90 |
| STI | Stimulants | 5 | 61 |
| SUB | Suboxone/Subutex | 5 | 17 |
| TIM | Targeted Immune Modulators | 12 | 17 |
| TOP | Topical Acne Agents | 21 | 155 |
| TRP | Triptans | 7 | 43 |
| ULT | Ultram ER | 2 | 1 |
| XIF | Xifaxan | 1 | 9 |
| XOI | Xanthine Oxidase Inhibitor | 1 | 2 |
| Totals | | 881 | 2528 |

**South Dakota Medicaid
Monthly Prior Authorization Report
October 1, 2014 – October 31, 2014**

By Request Type

| 10/01/14 - 10/31/14 | # of Requests | Electronic Requests | | Faxed Requests | |
|-----------------------------------|---------------|---------------------|------------|----------------|------------|
| | | # | % | # | % |
| Prior Authorizations: | | | | | |
| Antidepressant | 511 | 387 | 76% | 124 | 24% |
| Amrix and Fexmid | 4 | 4 | 100% | 0 | 0% |
| Altanax | 9 | 8 | 89% | 1 | 11% |
| Ambien CR | 14 | 9 | 64% | 5 | 36% |
| Anti-Infectives | 75 | 75 | 100% | 0 | 0% |
| Antihistamines | 60 | 54 | 90% | 6 | 10% |
| Antipsychotic | 852 | 518 | 61% | 334 | 39% |
| ARBS | 11 | 10 | 91% | 1 | 9% |
| Oral Anticoagulants | 8 | 6 | 75% | 2 | 25% |
| Dispense As Written | 11 | 0 | 0% | 11 | 100% |
| Antiemetics | 17 | 17 | 100% | 0 | 0% |
| Antifungals | 1 | 1 | 100% | 0 | 0% |
| Growth Hormone | 13 | 0 | 0% | 13 | 100% |
| Genitourinary SMR | 37 | 30 | 81% | 7 | 19% |
| Hepatitis Meds | 15 | 0 | 0% | 15 | 100% |
| Head Lice Medication | 77 | 55 | 71% | 22 | 29% |
| Lidoderm | 102 | 79 | 77% | 23 | 23% |
| Max Units Override | 882 | 769 | 87% | 113 | 13% |
| Name Brand Narcotics | 6 | 0 | 0% | 6 | 100% |
| Opioids | 45 | 42 | 93% | 3 | 7% |
| Onfi | 20 | 10 | 50% | 10 | 50% |
| Ophthalmic Antihistamines | 27 | 26 | 96% | 1 | 4% |
| Proton Pump Inhibitors | 133 | 101 | 76% | 32 | 24% |
| Skeletal Muscle Relaxants | 21 | 20 | 95% | 1 | 5% |
| Nasal Steroids | 99 | 78 | 79% | 21 | 21% |
| Stimulants | 66 | 60 | 91% | 6 | 9% |
| Suboxone/Subutex | 22 | 16 | 73% | 6 | 27% |
| Targeted Immune Modulators | 29 | 16 | 55% | 13 | 45% |
| Topical Acne Agents | 176 | 138 | 78% | 38 | 22% |
| Triptans | 50 | 40 | 80% | 10 | 20% |
| Ultram ER | 3 | 3 | 100% | 0 | 0% |
| Xifaxan | 10 | 8 | 80% | 2 | 20% |
| Xanthine Oxidase Inhibitor | 3 | 1 | 33% | 2 | 67% |
| Prior Authorization Totals | 3409 | 2581 | 76% | 828 | 24% |



**South Dakota Medicaid
Monthly Prior Authorization Report
October 1, 2014 – October 31, 2014**

Electronic PAs (Unique)

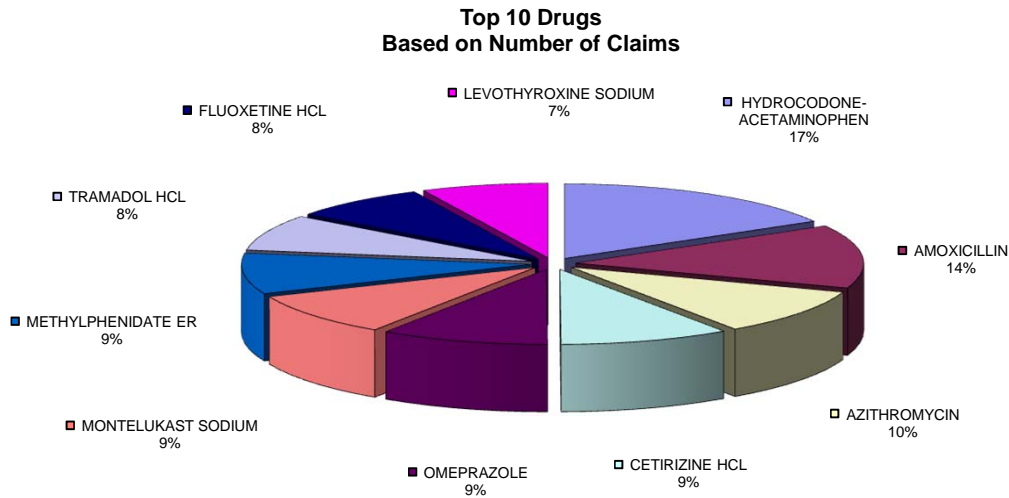
| 10/01/14 - 10/31/14 | # Unique Approved | # Unique Denied | # Unique Incomplete | Unique Total | Approval % | Total Transactions |
|------------------------------|----------------------|-----------------------|------------------------|-----------------|---------------|-----------------------|
| Prior Authorizations: | | | | | | |
| Antidepressant | 106 | 269 | 0 | 375 | 28.30% | 387 |
| Amrix and Fexmid | 0 | 4 | 0 | 4 | 0.00% | 4 |
| Altabax | 1 | 7 | 0 | 8 | 12.50% | 8 |
| Ambien CR | 4 | 5 | 0 | 9 | 44.40% | 9 |
| Anti-Infectives | 0 | 75 | 0 | 75 | 0.00% | 75 |
| Antihistamines | 4 | 50 | 0 | 54 | 7.40% | 54 |
| Antipsychotic | 178 | 331 | 0 | 509 | 35.00% | 518 |
| ARBS | 2 | 8 | 0 | 10 | 20.00% | 10 |
| Oral Anticoagulants | 1 | 5 | 0 | 6 | 16.70% | 6 |
| Antiemetics | 0 | 17 | 0 | 17 | 0.00% | 17 |
| Antifungals | 0 | 1 | 0 | 1 | 0.00% | 1 |
| Genitourinary SMR | 5 | 25 | 0 | 30 | 16.70% | 30 |
| Head Lice Medication | 0 | 52 | 0 | 52 | 0.00% | 55 |
| Lidoderm | 0 | 77 | 0 | 77 | 0.00% | 79 |
| Max Units Override | 26 | 699 | 0 | 725 | 3.60% | 769 |
| Opioids | 7 | 35 | 0 | 42 | 16.70% | 42 |
| Onfi | 0 | 10 | 0 | 10 | 0.00% | 10 |
| Ophthalmic Antihistamines | 1 | 25 | 0 | 26 | 3.80% | 26 |
| Proton Pump Inhibitors | 28 | 70 | 0 | 98 | 28.60% | 101 |
| Skeletal Muscle Relaxants | 1 | 15 | 0 | 16 | 6.30% | 20 |
| Nasal Steroids | 6 | 67 | 0 | 73 | 8.20% | 78 |
| Stimulants | 1 | 57 | 0 | 58 | 1.70% | 60 |
| Suboxone/Subutex | 0 | 15 | 0 | 15 | 0.00% | 16 |
| Targeted Immune Modulators | 1 | 15 | 0 | 16 | 6.30% | 16 |
| Topical Acne Agents | 10 | 123 | 0 | 133 | 7.50% | 138 |
| Triptans | 3 | 35 | 0 | 38 | 7.90% | 40 |
| Ultram ER | 2 | 1 | 0 | 3 | 66.70% | 3 |
| Xifaxan | 0 | 8 | 0 | 8 | 0.00% | 8 |
| Xanthine Oxidase Inhibitor | 0 | 1 | 0 | 1 | 0.00% | 1 |
| TOTALS | 387 | 2102 | 0 | 2489 | 15.50% | 2581 |

**South Dakota Medicaid
Cost Management Analysis**

TOP 25 DRUGS BASED ON NUMBER OF CLAIMS FROM 07/01/2014 - 09/30/2014

| Drug | AHFS Therapeutic Class | Rx | Paid | Paid/Rx | % Total Claims |
|-------------------------------|--|---------------|------------------------|-----------------|----------------|
| HYDROCODONE-ACETAMINOPHEN | OPIATE AGONISTS | 5,897 | \$ 103,860.09 | \$ 17.61 | 3.33% |
| AMOXICILLIN | PENICILLINS | 4,852 | \$ 40,802.29 | \$ 8.41 | 2.74% |
| AZITHROMYCIN | MACROLIDES | 3,536 | \$ 51,002.67 | \$ 14.42 | 2.00% |
| CETIRIZINE HCL | SECOND GENERATION ANTIHISTAMINES | 3,363 | \$ 25,027.37 | \$ 7.44 | 1.90% |
| OMEPRAZOLE | PROTON-PUMP INHIBITORS | 3,330 | \$ 38,827.42 | \$ 11.66 | 1.88% |
| MONTELUKAST SODIUM | LEUKOTRIENE MODIFIERS | 3,262 | \$ 68,457.12 | \$ 20.99 | 1.84% |
| METHYLPHENIDATE ER | RESPIRATORY AND CNS STIMULANTS | 3,121 | \$ 475,652.89 | \$ 152.40 | 1.77% |
| TRAMADOL HCL | OPIATE AGONISTS | 2,765 | \$ 21,531.92 | \$ 7.79 | 1.56% |
| FLUOXETINE HCL | ANTIDEPRESSANTS | 2,704 | \$ 21,446.77 | \$ 7.93 | 1.53% |
| LEVOTHYROXINE SODIUM | THYROID AGENTS | 2,510 | \$ 28,122.62 | \$ 11.20 | 1.42% |
| VYVANSE | AMPHETAMINES | 2,458 | \$ 473,057.96 | \$ 192.46 | 1.39% |
| SERTRALINE HCL | ANTIDEPRESSANTS | 2,416 | \$ 18,084.73 | \$ 7.49 | 1.37% |
| ALBUTEROL SULFATE | BETA-ADRENERGIC AGONISTS | 2,163 | \$ 38,265.63 | \$ 17.69 | 1.22% |
| VENTOLIN HFA | BETA-ADRENERGIC AGONISTS | 2,066 | \$ 99,097.29 | \$ 47.97 | 1.17% |
| TRAZODONE HCL | ANTIDEPRESSANTS | 1,938 | \$ 11,685.16 | \$ 6.03 | 1.10% |
| LORATADINE | SECOND GENERATION ANTIHISTAMINES | 1,930 | \$ 11,244.31 | \$ 5.83 | 1.09% |
| LISINAPRIL | ANGIOTENSIN-CONVERTING ENZYME INHIBITORS | 1,789 | \$ 9,998.04 | \$ 5.59 | 1.01% |
| INTUNIV | CENTRAL NERVOUS SYSTEM AGENTS, MISC. | 1,789 | \$ 488,694.99 | \$ 273.17 | 1.01% |
| DEXTROAMPHETAMINE-AMPHET ER | AMPHETAMINES | 1,785 | \$ 236,967.03 | \$ 132.75 | 1.01% |
| CEPHALEXIN | CEPHALOSPORINS | 1,710 | \$ 23,376.00 | \$ 13.67 | 0.97% |
| FLUTICASON PROPRIONATE | CORTICOSTEROIDS (EENT) | 1,668 | \$ 25,757.41 | \$ 15.44 | 0.94% |
| GABAPENTIN | ANTICONVULSANTS, MISCELLANEOUS | 1,582 | \$ 28,442.14 | \$ 17.98 | 0.89% |
| CLONIDINE HCL | CENTRAL ALPHA-AGONISTS | 1,558 | \$ 10,803.17 | \$ 6.93 | 0.88% |
| CLONAZEPAM | BENZODIAZEPINES (ANTICONVULSANTS) | 1,547 | \$ 11,967.70 | \$ 7.74 | 0.87% |
| SULFAMETHOXAZOLE-TRIMETHOPRIM | SULFONAMIDES (SYSTEMIC) | 1,485 | \$ 20,527.34 | \$ 13.82 | 0.84% |
| TOTAL TOP 25 | | 63,224 | \$ 2,382,700.06 | \$ 37.69 | 35.75% |

| | |
|---|---------|
| Total Rx Claims From 07/01/2014 - 09/30/2014 | 176,826 |
|---|---------|



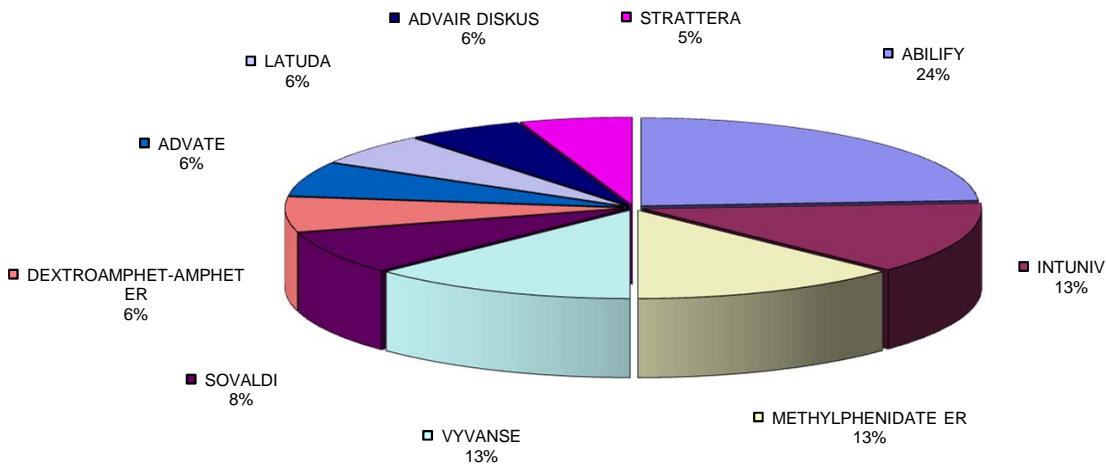
**South Dakota Medicaid
Cost Management Analysis**

TOP 25 DRUGS BASED ON TOTAL CLAIMS COST FROM 07/01/2014 - 09/30/2014

| Drug | AHFS Therapeutic Class | Rx | Paid | Paid/Rx | % Total Claims |
|------------------------|--|--------|-----------------|--------------|----------------|
| ABILIFY | ANTIPSYCHOTIC AGENTS | 1,232 | \$ 890,224.87 | \$ 722.59 | 0.70% |
| INTUNIV | CENTRAL NERVOUS SYSTEM AGENTS, MISC. | 1,789 | \$ 488,694.99 | \$ 273.17 | 1.01% |
| METHYLPHENIDATE ER | RESPIRATORY AND CNS STIMULANTS | 3,121 | \$ 475,652.89 | \$ 152.40 | 1.77% |
| VYVANSE | AMPHETAMINES | 2,458 | \$ 473,057.96 | \$ 192.46 | 1.39% |
| SOVALDI | HCV ANTIVIRALS | 10 | \$ 292,331.00 | \$ 29,233.10 | 0.01% |
| DEXTROAMPHET-AMPHET ER | AMPHETAMINES | 1,785 | \$ 236,967.03 | \$ 132.75 | 1.01% |
| ADVATE | HEMOSTATICS | 5 | \$ 232,608.36 | \$ 46,521.67 | 0.00% |
| LATUDA | ANTIPSYCHOTIC AGENTS | 297 | \$ 215,932.28 | \$ 727.04 | 0.17% |
| ADVAIR DISKUS | CORTICOSTEROIDS (RESPIRATORY TRACT) | 698 | \$ 205,179.35 | \$ 293.95 | 0.39% |
| STRATTERA | CENTRAL NERVOUS SYSTEM AGENTS, MISC. | 769 | \$ 198,626.93 | \$ 258.29 | 0.43% |
| HUMIRA | DISEASE-MODIFYING ANTIRHEUMATIC AGENTS | 68 | \$ 194,527.02 | \$ 2,860.69 | 0.04% |
| LYRICA | ANTICONSULSANTS, MISCELLANEOUS | 614 | \$ 188,771.10 | \$ 307.44 | 0.35% |
| FOCALIN XR | RESPIRATORY AND CNS STIMULANTS | 749 | \$ 179,316.61 | \$ 239.41 | 0.42% |
| PULMOZYME | MUCOLYTIC AGENTS | 57 | \$ 174,121.47 | \$ 3,054.76 | 0.03% |
| INVEGA SUSTENNA | ANTIPSYCHOTIC AGENTS | 118 | \$ 171,293.75 | \$ 1,451.64 | 0.07% |
| LANTUS SOLOSTAR | INSULINS | 441 | \$ 160,638.32 | \$ 364.26 | 0.25% |
| FLOVENT HFA | CORTICOSTEROIDS (RESPIRATORY TRACT) | 778 | \$ 139,643.11 | \$ 179.49 | 0.44% |
| PREVACID | PROTON-PUMP INHIBITORS | 460 | \$ 137,909.96 | \$ 299.80 | 0.26% |
| COPAXONE | IMMUNOMODULATORY AGENTS | 27 | \$ 137,627.83 | \$ 5,097.33 | 0.02% |
| ENBREL | DISEASE-MODIFYING ANTIRHEUMATIC AGENTS | 45 | \$ 132,000.71 | \$ 2,933.35 | 0.03% |
| OXYCONTIN | OPIATE AGONISTS | 433 | \$ 130,937.08 | \$ 302.40 | 0.24% |
| NOVOLOG | INSULINS | 355 | \$ 117,697.81 | \$ 331.54 | 0.20% |
| RECOMBINATE | HEMOSTATICS | 3 | \$ 108,620.18 | \$ 36,206.73 | 0.00% |
| NOVOLOG FLEXPEN | INSULINS | 273 | \$ 108,603.97 | \$ 397.82 | 0.15% |
| KALYDECO | CYSTIC FIBROSIS (CFTR) POTENTIATORS | 4 | \$ 106,935.72 | \$ 26,733.93 | 0.00% |
| TOTAL TOP 25 | | 16,589 | \$ 5,897,920.30 | \$ 355.53 | 9.38% |

| | |
|---|---------|
| Total Rx Claims From 07/01/2014 - 09/30/2014 | 176,826 |
|---|---------|

**Top 10 Drugs
Based on Total Claims Cost**



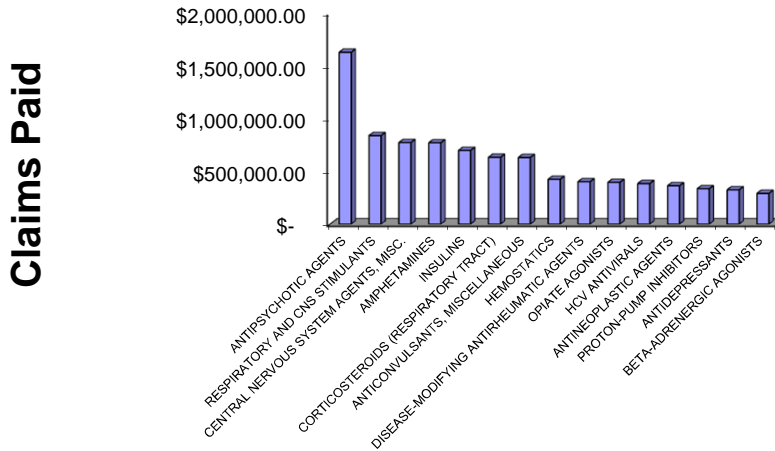
**SOUTH DAKOTA MEDICAID
Cost Management Analysis**

TOP 15 THERAPEUTIC CLASSES BY TOTAL COST OF CLAIMS FROM 07/01/2014 - 09/30/2014

| AHFS Therapeutic Class | Rx | Paid | Paid/Rx | % Total Claims |
|--|---------------|------------------------|------------------|----------------|
| ANTIPSYCHOTIC AGENTS | 5,758 | \$ 1,627,306.03 | \$ 282.62 | 3.26% |
| RESPIRATORY AND CNS STIMULANTS | 5,491 | \$ 839,850.10 | \$ 152.95 | 3.11% |
| CENTRAL NERVOUS SYSTEM AGENTS, MISC. | 2,607 | \$ 773,986.69 | \$ 296.89 | 1.47% |
| AMPHETAMINES | 5,059 | \$ 771,686.30 | \$ 152.54 | 2.86% |
| INSULINS | 2,036 | \$ 699,176.20 | \$ 343.41 | 1.15% |
| CORTICOSTEROIDS (RESPIRATORY TRACT) | 2,626 | \$ 633,378.60 | \$ 241.20 | 1.49% |
| ANTICONVULSANTS, MISCELLANEOUS | 7,719 | \$ 632,234.68 | \$ 81.91 | 4.37% |
| HEMOSTATICS | 21 | \$ 425,249.20 | \$ 20,249.96 | 0.01% |
| DISEASE-MODIFYING ANTIRHEUMATIC AGENTS | 162 | \$ 402,125.88 | \$ 2,482.26 | 0.09% |
| OPIATE AGONISTS | 12,759 | \$ 396,246.33 | \$ 31.06 | 7.22% |
| HCV ANTIVIRALS | 14 | \$ 384,708.52 | \$ 27,479.18 | 0.01% |
| ANTINEOPLASTIC AGENTS | 435 | \$ 363,754.63 | \$ 836.22 | 0.25% |
| PROTON-PUMP INHIBITORS | 5,241 | \$ 336,150.67 | \$ 64.14 | 2.96% |
| ANTIDEPRESSANTS | 14,834 | \$ 325,384.36 | \$ 21.94 | 8.39% |
| BETA-ADRENERGIC AGONISTS | 6,504 | \$ 291,533.07 | \$ 44.82 | 3.68% |
| TOTAL TOP 15 | 71,266 | \$ 8,902,771.26 | \$ 124.92 | 40.30% |

| | |
|---|---------|
| Total Rx Claims From 07/01/2014 - 09/30/2014 | 176,826 |
|---|---------|

**Top 15 Therapeutic Classes
Based on Total Cost of Claims**



SD Medicaid Drug Spend 2012 - 2014

Drug Spend 2012

| Total | 201201 | 201202 | 201203 | 201204 | 201205 | 201206 | 201207 | 201208 | 201209 | 201210 | 201211 | 201212 | Row SubTotal |
|-----------------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|-----------------|
| Rx_Dollars | \$4,849,391.09 | \$4,997,072.08 | \$5,257,537.69 | \$4,879,456.62 | \$4,919,253.17 | \$4,913,164.11 | \$3,845,419.78 | \$4,863,874.68 | \$4,014,447.34 | \$4,823,060.40 | \$4,576,545.82 | \$4,443,459.76 | \$56,382,682.54 |
| Rx_Count | 77,559 | 77,426 | 79,761 | 74,004 | 74,773 | 75,503 | 57,785 | 72,290 | 63,542 | 77,195 | 73,717 | 71,219 | 874,774 |
| Average_Rx_Cost | \$62.53 | \$64.54 | \$65.92 | \$65.94 | \$65.79 | \$65.07 | \$66.55 | \$67.28 | \$63.18 | \$62.48 | \$62.08 | \$62.39 | \$64.45 |
| Recip_Count | 29,127 | 30,437 | 29,893 | 28,110 | 27,215 | 23,029 | 22,803 | 26,411 | 25,434 | 28,741 | 28,017 | 27,141 | |
| Recip_Average_Rx_Cost | \$166.49 | \$164.18 | \$175.88 | \$173.58 | \$180.76 | \$213.35 | \$168.64 | \$184.16 | \$157.84 | \$167.81 | \$163.35 | \$163.72 | \$172.76 |

Drug Spend 2013

| Total | 201301 | 201302 | 201303 | 201304 | 201305 | 201306 | 201307 | 201308 | 201309 | 201310 | 201311 | 201312 | Row SubTotal |
|-----------------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|-----------------|
| Rx_Dollars | \$5,098,169.75 | \$4,387,497.68 | \$3,483,772.72 | \$4,587,727.94 | \$4,954,424.54 | \$4,080,090.86 | \$4,746,275.35 | \$4,785,301.34 | \$4,365,068.62 | \$5,013,651.58 | \$4,581,380.72 | \$4,062,526.21 | \$54,145,887.31 |
| Rx_Count | 75,726 | 68,046 | 54,790 | 68,602 | 68,081 | 57,927 | 65,531 | 66,324 | 62,203 | 72,334 | 65,858 | 56,581 | 782,003 |
| Average_Rx_Cost | \$67.32 | \$64.48 | \$63.58 | \$66.87 | \$72.77 | \$70.44 | \$72.43 | \$72.15 | \$70.17 | \$69.31 | \$69.56 | \$71.80 | \$69.24 |
| Recip_Count | 27,719 | 25,992 | 22,293 | 24,675 | 24,016 | 21,411 | 22,580 | 23,563 | 23,753 | 25,748 | 24,484 | 21,797 | |
| Recip_Average_Rx_Cost | \$183.92 | \$168.80 | \$156.27 | \$185.93 | \$206.30 | \$190.56 | \$210.20 | \$203.09 | \$183.77 | \$194.72 | \$187.12 | \$183.38 | \$187.99 |

Drug Spend 2014

| Total | 201401 | 201402 | 201403 | 201404 | 201405 | 201406 | 201407 | 201408 | 201409 | 201410 | 201411 | 201412 | Row SubTotal |
|-----------------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|--------|--------|-----------------|
| Rx_Dollars | \$5,148,590.25 | \$5,032,584.04 | \$5,434,106.10 | \$5,247,601.16 | \$5,247,835.49 | \$4,403,564.85 | \$4,929,653.23 | \$4,735,906.92 | \$4,496,590.80 | \$5,345,154.78 | \$0.00 | \$0.00 | \$50,021,587.62 |
| Rx_Count | 68,517 | 68,225 | 71,659 | 71,967 | 67,546 | 54,749 | 61,792 | 61,046 | 53,988 | 67,332 | 0 | 0 | 646,821 |
| Average_Rx_Cost | \$75.14 | \$73.76 | \$75.83 | \$72.92 | \$77.69 | \$80.43 | \$79.78 | \$77.58 | \$83.29 | \$79.39 | \$0.00 | \$0.00 | \$77.33 |
| Recip_Count | 25,130 | 25,540 | 26,526 | 25,766 | 24,319 | 20,659 | 22,345 | 22,689 | 21,874 | 24,405 | 0 | 0 | |
| Recip_Average_Rx_Cost | \$204.88 | \$197.05 | \$204.86 | \$203.66 | \$215.79 | \$213.15 | \$220.62 | \$208.73 | \$205.57 | \$219.02 | \$0.00 | \$0.00 | \$209.07 |

Anticipated Availability of First-Time Generics

—To help explain the benefits of generic drugs to your patients, the FDA has patient education materials available at
<http://www.fda.gov/Drugs/ResourcesForYou/Consumers/BuyingUsingMedicineSafely/UnderstandingGenericDrugs/ucm169209.htm>—

| Brand^a (Manufacturer) | Generic Name | Generic Manufacturer(s)^{b,1} | Anticipated Availability^c |
|---|---|---|---|
| <i>Actonel</i> (Warner Chilcott) | Risedronate tablets | Actavis, Apotex, Mylan | Generic now available |
| <i>Avelox</i> (Bayer) | Moxifloxacin | Alvogon, Aurobindo, Dr. Reddy's, Major, Teva, Torrent | Generic now available |
| <i>Baraclude</i> (Bristol-Myers Squibb) | Entecavir | Teva | Generic now available |
| <i>Boniva</i> (Roche) | Ibandronate injection | Caraco, Sagent | Generic now available |
| <i>Bromday</i> (Bausch & Lomb) | Bromfenac | Apotex, Hi-Tech Pharmacal | Generic now available |
| <i>Cipro</i> (Bayer) | Ciprofloxacin for oral suspension | Lupin | Generic now available |
| <i>Detrol LA</i> (Pfizer) | Tolterodine extended-release capsule | Mylan | Generic now available |
| <i>Differin</i> (Galderma) | Adapalene 0.3% gel | Sandoz (Tolmar) | Generic now available |
| <i>Diovan</i> (Novartis) | Valsartan | Ohm Laboratories, Sandoz | Generic now available |
| <i>Epivir-HBV</i> (GlaxoSmithKline) | Lamivudine | Apotex, Camber (Hetero) | Generic now available |
| <i>Evista</i> (Lilly) | Raloxifene | Teva | Generic now available |
| <i>Exalgo</i> (Mallinckrodt) | Hydromorphone extended-release tablet 8 mg, 12 mg, 16 mg | Actavis | Generic now available |

More . . .

| Brand^a (Manufacturer) | Generic Name | Generic Manufacturer(s)^{b,1} | Anticipated Availability^c |
|---|--|---|---|
| <i>Exforge</i> (Novartis) | Amlodipine/Valsartan | Par | Generic now available |
| <i>Hectorol</i> (Genzyme) | Doxercalciferol injection | Sandoz, West-Ward (Hikma) | Generic now available |
| <i>Hectorol</i> (Genzyme) | Doxercalciferol capsule | Roxane | Generic now available |
| <i>Klor-Con</i> (Upsher-Smith) | Potassium chloride extended-release tablet 8 mEq, 10 mEq | Mylan | Generic now available |
| <i>Locoid</i> (Onset Dermatologics) | Hydrocortisone butyrate 0.1% cream (lipophilic) | Glenmark | Generic now available |
| <i>Loestrin 24 Fe</i> (Warner Chilcott) | Ethinyl estradiol/Norethindrone acetate | Amneal | Generic now available |
| <i>Lovaza</i> (GlaxoSmithKline) | Omega-3-Acid Ethyl Esters | Par, Teva | Generic now available |
| <i>Lunesta</i> (Sunovion) | Eszopiclone | Avkare, Caraco (Sun), Dr. Reddy's, Glenmark, Lupin, Mylan, Roxane, Teva | Generic now available |
| <i>Malarone</i> (GlaxoSmithKline) | Atovaquone/Proguanil | Glenmark, Mylan | Generic now available |
| <i>Micardis HCT</i> (Boehringer Ingelheim) | Telmisartan/Hydrochlorothiazide | Lupin, Mylan, Qualitest (Alembic), Torrent | Generic now available |
| <i>Monodox</i> (Aqua) | Doxycycline capsule | Lupin, Par, Ranbaxy | Generic now available |
| <i>Mycobutin</i> (Pfizer) | Rifabutin | Lupin | Generic now available |
| <i>Nexium IV</i> (AstraZeneca) | Esomeprazole sodium injection | Caraco (Sun) | Generic now available |
| <i>Ortho-Evra</i> (Janssen) | Ethinyl estradiol/Norelgestromin patch | Mylan | Generic now available |
| <i>Oxsoralen-Ultra</i> (Valeant) | Methoxsalen soft gelatin capsules | Strides Arcolab | Generic now available |

| Brand^a (Manufacturer) | Generic Name | Generic Manufacturer(s)^{b,1} | Anticipated Availability^c |
|--|---|--|---|
| <i>Pennsaid</i> (Mallinckrodt) | Diclofenac 1.5% topical solution | Apotex | Generic now available |
| <i>Precedex</i> (Hospira) | Dexmedetomidine | Mylan, Par | Generic now available |
| <i>Rapamune</i> (Pfizer) | Sirolimus tablet | Zydus | Generic now available |
| <i>Renvela</i> (Genzyme) | Sevelamer carbonate tablet | Global (Impax)(authorized generic) | Generic now available |
| <i>Retin-A Micro</i> (Valeant) | Tretinoin gel, microspheres 0.1 and 0.04% | Spear | Generic now available |
| <i>Rhinocort Aqua</i> (AstraZeneca) | Budesonide nasal spray | Apotex | Generic now available |
| <i>Solaraze</i> (Fougera) | Diclofenac sodium gel 3% | Global (Impax; Tolmar) | Generic now available |
| <i>Taclonex</i> (Leo Pharma) | Betamethasone dipropionate/Calcipotriene hydrate ointment | Sandoz (Tolmar) | Generic now available |
| <i>Trizivir</i> (ViiV Healthcare) | Abacavir, Lamivudine, Zidovudine | Lupin | Generic now available |
| <i>Twynsta</i> (Boehringer Ingelheim) | Amlodipine/Telmisartan | Lupin, Torrent | Generic now available |
| <i>Urocit-K</i> (Mission Pharma) | Potassium citrate extended-release tablet | Zydus | Generic now available |
| <i>Vanos</i> (Medicis) | Fluocinonide 0.1% cream | Glenmark, Perrigo, Taro | Generic now available |
| <i>Vidaza</i> (Celgene) | Azacitidine injection | Dr. Reddy's | Generic now available |
| <i>Viramune XR</i> (Boehringer Ingelheim) | Nevirapine extended-release tablets | Apotex, Sandoz | Generic now available |
| <i>Vivelle-Dot</i> (Novartis) | Estradiol extended-release transdermal film | Mylan | Generic now available |

| Brand ^a (Manufacturer) | Generic Name | Generic Manufacturer(s) ^{b,1} | Anticipated Availability ^c |
|---|--|---|---------------------------------------|
| <i>Xeloda</i> (Hoffmann La Roche) | Capecitabine | Mylan, Teva | Generic now available |
| <i>Celebrex</i> (Pfizer) | Celecoxib | Actavis, Mylan, Teva | December 2014 ⁵⁵ |
| <i>Intuniv</i> (Shire) | Guanfacine extended-release tablet | Actavis, ^{75,h} Sandoz, Teva, TWI | December 2014 ⁷⁵ |
| <i>Namenda</i> (Forest) | Memantine tablet | Alembic, Amneal, Apotex, Aurobindo, Jubilant Life, Lupin, Macleods, Mylan, Upsher-Smith, Torrent, Unichem, Wockhardt | January 2015 ²¹ |
| <i>Sustiva</i> (Bristol-Myers Squibb) | Efavirenz | <u>Tablet</u> : Aurobindo, Cipla, Emcure, Hetero, Macleods, Matrix, Micro Labs, Par, Strides <u>Capsule</u> : Aurobindo, Cipla, Micro Labs | March 2015 |
| <i>Welchol</i> (Daiichi Sankyo) | Colesevelam tablet and oral suspension | Impax, ^{f,22} Actavis, ^{i,23,71} Glenmark ^{f,59} | March 2015 ²² |
| <i>Abilify</i> (Otsuka) | Aripiprazole | <u>Tablet</u> : Apotex, Alembic, Barr, Sun, Torrent <u>Orally disintegrating tablet</u> : Alembic, Barr, Zydus | April 2015 ⁴⁰ |
| <i>Generess Fe</i> (Actavis) | Norethindrone, Ethinyl estradiol, Ferrous fumarate | Family Care | April 2015 ^{107,h} |
| <i>Oxytrol</i> (Actavis) | Oxybutynin transdermal patch | Teva | April 2015 |
| <i>Factive</i> (Cornerstone) | Gemifloxacin tablet | Orchid | June 2015 |
| <i>Aggrenox</i> (Boehringer Ingelheim) | Aspirin/Dipyridamole | Barr | July 2015 ^{24,h} |
| <i>Gleevec</i> (Novartis) | Imatinib | Sun | July 2015 |

| Brand^a (Manufacturer) | Generic Name | Generic Manufacturer(s)^{b,1} | Anticipated Availability^c |
|---|---|--|---|
| <i>Targretin</i> (Eisai) | Bexarotene | Mylan ^{f,47} | July 2015 ⁴⁷ |
| <i>Valcyte</i> (Roche) | Valganciclovir tablet | Dr. Reddy's, Ranbaxy | September 2015 |
| <i>Asacol HD</i> (Warner Chilcott) | Mesalamine delayed-release tablet | Zydus ^{f,90} | November 2015 ⁹⁰ |
| <i>Avodart</i> (GlaxoSmithKline) | Dutasteride | Barr, Banner, Endo, Roxane, Sandoz | November 2015 |
| <i>Axert</i> (Ortho-McNeil-Janssen) | Almotriptan | Teva | November 2015 |
| <i>Frova</i> (Endo) | Frovatriptan | Mylan | November 2015 ^{37,h} |
| <i>Jalyn</i> (GlaxoSmithKline) | Dutasteride/Tamsulosin | Anchen, Impax, ^{i,13} Mylan, ^{i,13} Actavis ^{i,72} | November 2015 ^d |
| <i>Patanol</i> (Alcon) | Olopatadine 0.1% ophthalmic solution | Apotex, Sandoz, Wockhardt | December 2015 |
| <i>OxyContin</i> (Purdue) | Oxycodone extended-release tablet (new formulation) | Impax ^{f,12} | January 2016 ¹² |
| <i>Glumetza</i> (Santarus) | Metformin extended-release tablet | Lupin, ^{f,17} Sun | February 2016 ¹⁷ |
| <i>Crestor</i> (AstraZeneca) | Rosuvastatin calcium | Actavis, Apotex, Aurobindo, Glenmark, Mylan, Par, Sandoz, Sun, Teva | May 2016 ²⁶ |
| <i>Nuvigil</i> (Teva) | Armodafinil | Actavis, Mylan, Lupin | June 2016 ³¹ |
| <i>Zegerid/Zegerid OTC</i> (Santarus) | Omeprazole/Sodium bicarbonate capsule and oral suspension | Dr. Reddy's, ^{i,10} Par, Perrigo, Zydus ^{i,32} | July 2016 ^{20,d} |
| <i>Enablex</i> (Novartis) | Darifenacin | Anchen | August 2016 |
| <i>Oracea</i> (Galderma) | Doxycycline | Lupin, Mylan | August 2016 |

| Brand ^a (Manufacturer) | Generic Name | Generic Manufacturer(s) ^{b,1} | Anticipated Availability ^c |
|--|--|--|---------------------------------------|
| <i>Benicar/Benicar HCT</i> (Daiichi Sankyo) | Olmesartan/Olmesartan HCTZ | <u>Olmesartan</u> : Mylan, Sandoz <u>Olmesartan HCTZ</u> : Mylan, Teva | October 2016 |
| <i>Seroquel XR</i> (AstraZeneca) | Quetiapine extended-release tablet | Accord, Handa, Lupin, ^{f,43} Mylan, Osmotica, ^{f,53} Torrent ^{f,53} | November 2016 ⁷⁴ |
| <i>Kaletra</i> (Abbott) | Lopinavir/Ritonavir | <u>Tablet</u> : Aurobindo, Cipla, Hetero, Matrix, Mylan <u>Oral solution</u> : Cipla, Hetero | December 2016 ^{2,d} |
| <i>ProAir HFA</i> (Teva) | Albuterol | Perrigo, Catalent | December 2016 ³⁴ |
| <i>Zetia</i> (Merck) | Ezetimibe | Glenmark, Mylan | December 2016 ²⁸ |
| <i>Zyvox</i> (Pfizer) | Linezolid | <u>Injection</u> : Teva, Sandoz <u>Tablet</u> : Gate, Glenmark, Mylan, Teva <u>Oral suspension</u> : Roxane | Late 2016 ³⁶ |
| <i>Azilect</i> (Teva) | Rasagiline mesylate tablet | Apotex, Mylan, Orchid, ^{f,30} Sandoz ^{f,30} | February 2017 |
| <i>Vytorin</i> (Merck) | Ezetimibe/Simvastatin | Mylan | April 2017 ¹⁸ |
| <i>Strattera</i> (Lilly) | Atomoxetine | Actavis, ^{i,29} Apotex, Aurobindo, Dr. Reddy's, Glenmark, Mylan, Sandoz, Sun, Teva ^{i,29} | May 2017 ²⁹ |
| <i>Metozolv</i> (Salix) | Metoclopramide orally disintegrating tablet | Novel | July 2017 |
| <i>Reyataz</i> (Bristol-Myers Squibb) | Atazanavir | Aurobindo, Emcure, Matrix, Teva | July 2017 ⁵² |
| <i>Relpax</i> (Pfizer) | Eletriptan | Apotex, Teva | August 2017 ^{62,d} |
| <i>Treximet</i> (Pozen) | Sumatriptan/Naproxen | Alphapharm, ^{f,101} Dr. Reddy's, Mylan, Par, Teva ^{f,101} | August 2017 ¹⁰¹ |
| <i>Nasonex</i> (Merck) | Mometasone | Apotex, ^{i,95} Teva ^{i,95} | October 2017 ^{113,d} |

| Brand ^a (Manufacturer) | Generic Name | Generic Manufacturer(s) ^{b,1} | Anticipated Availability ^c |
|--------------------------------------|---|---|---------------------------------------|
| <i>Adcirca</i> (Lilly) | Tadalafil | Synthon | November 2017 |
| <i>Cubicin</i> (Cubist) | Daptomycin | Hospira, Teva ^{f,69} | December 2017 ⁶⁹ |
| <i>Viagra</i> (Pfizer) | Sildenafil | Actavis, Amneal, Apotex, Dr. Reddy's, Hetero, Macleods, Mylan, Teva | December 2017 ⁹¹ |
| <i>Viread</i> (Gilead) | Tenofovir disoproxil fumarate | Aurobindo, Cipla, InvaGen, Macleods, Matrix, Strides Arcolab, Teva | December 2017 ⁸ |
| <i>Sensipar</i> (Amgen) | Cinacalcet tablet | Teva | March 2018 ⁵¹ |
| <i>Zemplar</i> (AbbVie) | Paricalcitol injection | Dr.Reddy's, ^{i,96} Hospira, ^{i,44} Sandoz, Sun, ^{i,33} Teva | April 2018 ^{96,d} |
| <i>Acanya</i> (Valeant) | Benzoyl peroxide/Clindamycin phosphate | Actavis ^{f,87} | July 2018 ⁸⁷ |
| <i>Fentora</i> (Cephalon) | Fentanyl citrate buccal/sublingual tablet | Teva, ^{f,56} Impax ^{f,57} | October 2018 ⁵⁶ |
| <i>Levitra</i> (GlaxoSmithKline) | Vardenafil | Teva | October 2018 ⁴² |
| <i>Epzicom</i> (GlaxoSmithKline) | Abacavir sulfate/Lamivudine | <u>Tablet:</u> Aurobindo, Cipla, Matrix, Mylan, Teva <u>Tablet for Suspension:</u> Cipla | November 2018 |
| <i>Fortesta</i> (Endo) | Testosterone gel | Actavis ^{f,65} | November 2018 ^d |
| <i>Vesicare</i> (Astellas) | Solifenacin succinate | Teva | November 2018 |
| <i>Lyrica</i> (Pfizer) | Pregabalin | <u>Capsule:</u> Actavis, Apotex, Lupin, Mylan, Sandoz, Teva, Wockhardt <u>Oral solution:</u> Lupin | December 2018 ⁵⁰ |

| Brand ^a (Manufacturer) | Generic Name | Generic Manufacturer(s) ^{b,1} | Anticipated Availability ^c |
|--------------------------------------|---|--|---------------------------------------|
| <i>Exelon</i> (Novartis) | Rivastigmine transdermal patch | Actavis, ^{i,66} Noven ^{i,64} | January 2019 ^d |
| <i>Ranexa</i> (Gilead) | Ranolazine | Lupin | February 2019 ⁸¹ |
| <i>AzaSite</i> (InSite Vision) | Azithromycin 1% ophthalmic solution | Mylan, ^{i,16} Sandoz | March 2019 ^d |
| <i>Emend</i> (Merck) | Fosaprepitant injection | Accord, ^{i,35} Sandoz ^{i,35} | March 2019 ^d |
| <i>Angiomax</i> (Medicines Co) | Bivalirudin | Hospira, Teva ^{f,104} | June 2019 ¹⁰⁴ |
| <i>Prezista</i> (Janssen) | Darunavir | Hetero, ^{i,84} Lupin, ^{i,79} Mylan, Teva | December 2019 ^d |
| <i>Silenor</i> (Somaxon) | Doxepin | Actavis, Mylan, Par, Zydus | January 2020 ³⁹ |
| <i>Vigamox</i> (Alcon) | Moxifloxacin ophthalmic | Actavis, ^{i,6} Akorn, Apotex, Lupin, Teva | March 2020 ^d |
| <i>Safyral</i> (Bayer) | Drospirenone/Ethinyl estradiol/Levomefolate calcium | Actavis ^{i,46} | April 2020 ^d |
| <i>Chantix</i> (Pfizer) | Varenicline | Apotex, Mylan, Teva | May 2020 ^{49,d} |
| <i>Lialda</i> (Shire) | Mesalamine delayed-release tablet | Zydus ^{i,76} | June 2020 ^d |
| <i>Oleptro</i> (Angelini) | Trazodone extended-release tablet | Actavis ^{i,11} | June 2020 ^d |
| <i>AndroGel 1.62%</i> (AbbVie) | Testosterone | Perrigo, ^{i,58} Actavis ^{i,67} | August 2020 ^d |
| <i>Lescol XL</i> (Novartis) | Fluvastatin extended-release | Mylan, Par, Teva | October 2020 |
| <i>Tarceva</i> (OSI) | Erlotinib tablet | Mylan | November 2020 ^{73,h} |
| <i>Sutent</i> (Pfizer) | Sunitinib | Mylan | February 2021 ^{4,d} |

| Brand ^a (Manufacturer) | Generic Name | Generic Manufacturer(s) ^{b,1} | Anticipated Availability ^c |
|--------------------------------------|--|--|---------------------------------------|
| <i>Absorica</i> (Ranbaxy) | Isotretinoin | Actavis ^{i,88} | September 2021 ^d |
| <i>Atripla</i> (Gilead) | Efavirenz/Emtricitabine/Tenofovir disoproxil fumarate | Aurobindo, Cipla, Hetero, Matrix, Teva | September 2021 ^{68,d} |
| <i>Bystolic</i> (Forest) | Nebivolol | Actavis, ^{f,67} Alkem, ^{f,67} Amerigen, ^{f,67} Glenmark, ^{f,63} Hetero, ^{f,67} Indchemie, ^{f,67} Torrent ^{f,67} | September 2021 ⁶³ |
| <i>Emtriva</i> (Gilead) | Emtricitabine | Aurobindo, Cipla, Matrix | September 2021 ^{7,h} |
| <i>Truvada</i> (Gilead) | Emtricitabine/Tenofovir disoproxil fumarate | Aurobindo, Cipla, Hetero, Matrix, Strides Arcolab, Teva | September 2021 ^{68,d} |
| <i>Vimpat</i> (UCB) | Lacosamide | Actavis, ^{i,5} Mylan, ^{i,5} Others ^{i,5} | March 2022 ^d |
| <i>HalfLytely</i> (Braintree) | Polyethylene glycol 3350, potassium chloride, sodium bicarbonate, sodium chloride, bisacodyl | Novel | October 2022 |
| <i>Banzel</i> (Eisai) | Rufinamide | Glenmark, ^{i,80} Hetero, ^{i,80} Lupin, ^{i,80} Mylan, ^{i,80} Roxane ^{i,80} | November 2022 ^d |
| <i>Xyrem</i> (Jazz) | Sodium oxybate | Amneal ^{i,48} | December 2022 ^d |
| <i>Vyvanse</i> (Shire) | Lisdexamfetamine | Actavis, ^{i,109} Amneal, Mylan, Roxane, Sandoz ^{i,109} | June 2023 ^d |
| <i>Exforge HCT</i> (Novartis) | Amlodipine/Valsartan/ Hydrochlorothiazide | Lupin, Teva, Par, Torrent ^{i,38} | November 2023 ^d |
| <i>Gralise</i> (Depomed) | Gabapentin | Actavis, Incepta, ^{i,97} Zydus ^{i,97} | January 2024 ^{97,d} |
| <i>Pataday</i> (Alcon) | Olopatadine | Actavis, Apotex, Teva, Wockhart ^{i,99} | May 2024 ^{100,d} |
| <i>Aloxi</i> (Eisai) | Palonosetron injection | Dr. Reddy's, Sandoz, ^{i,78} Teva | July 2024 ^d |

| Brand^a (Manufacturer) | Generic Name | Generic Manufacturer(s)^{b,1} | Anticipated Availability^c |
|---|---|--|---|
| <i>Januvia</i> (Merck) | Sitagliptin | Actavis, Apotex, Mylan, Sandoz, Sun | April 2026 |
| <i>Natazia</i> (Bayer) | Dienogest/estradiol valerate | Lupin ^{i,61} | May 2026 ^d |
| <i>Aplenzin</i> (Biovail) | Bupropion hydrobromide extended-release tablet | Actavis, Paddock ^{i,83} | June 2026 ^d |
| <i>Nuedexta</i> (Avanir) | Dextromethorphan/Quinidine | Par, Sandoz, ^{f,102} Wockhardt ^{f,85} | July 2026 ^{85,102} |
| <i>Dexilant</i> (Takeda) | Dexlansoprazole | Dr. Reddy's, ^{i,103} Handa, ^{i,92} Impax, ^{i,92} Par | February 2027 ^d |
| <i>Solodyn</i> (Medicis) | Minocycline extended-release tablet 80 mg, 105 mg | Mylan | March 2027 |
| <i>Ampyra</i> (Acorda) | Dalfampridine | Accord, ^{i,110} Actavis, ^{i,110} Aurobindo, ^{i,110} Roxane, ^{i,110} Alkem, ^{i,110} Mylan ^{i,110} | May 2027 ^d |
| <i>Lumigan</i> (Allergan) | Bimatoprost ophthalmic solution 0.01% | Sandoz | June 2027 |
| <i>Axiron</i> (Eli Lilly) | Testosterone transdermal solution | Actavis ^{i,9} | July 2027 ^d |
| <i>Emend</i> (Merck) | Aprepitant capsule | Sandoz | September 2027 |
| <i>Janumet</i> (Merck) | Sitagliptin/Metformin | Apotex, Sandoz | July 2028 |
| <i>Onglyza</i> (Bristol-Myers Squibb) | Saxagliptin | Actavis ^{i,98} | November 2028 ^d |
| <i>Lo Loestrin Fe</i> (Actavis) | Ethinyl estradiol/Norethindrone acetate/Ferrous fumarate | Amneal, ^{i,94} Lupin ^{i,94} | February 2029 |
| <i>Qsymia</i> (Vivus) | Phentermine/Topiramate extended-release capsule | Actavis ^{i,41} | June 2029 ^d |
| <i>Mirapex ER</i> (Boehringer Ingelheim) | Pramipexole extended-release tablet | Anchen | September 2029 |

| Brand ^a (Manufacturer) | Generic Name | Generic Manufacturer(s) ^{b,1} | Anticipated Availability ^c |
|--|--|--|---------------------------------------|
| <i>Savella</i> (Cypress) | Milnacipran | Mylan ^{i,14} | September 2029 ^d |
| <i>Zubsolv</i> (Orexo) | Buprenorphine/Naloxone sublingual tablets | Actavis ^{i,111} | October 2029 ^d |
| <i>Suboxone</i> (Reckitt Benckiser) | Buprenorphine/Naloxone sublingual film | Actavis ^{i,86} | March 2030 ^d |
| <i>Copaxone</i> (Teva) | Glatiramer 40 mg | Dr. Reddy's ^{i,93} | September 2030 ^d |
| <i>Uloric</i> (Takeda) | Febuxostat | Mylan, ^{i,89} Roxane ^{i,105} | September 2031 ^d |
| <i>Vimovo</i> (AstraZeneca) | Naproxen/Esomeprazole magnesium delayed-release tablet | Actavis, ^{i,77} Anchen, ^{i,77} Dr. Reddy's, ^{i,77} Lupin, ^{i,77} Mylan ^{i,77} | May 2033 ^d |
| <i>Acova</i> (Pfizer) | Argatroban 100 mg/mL injection | Hikma, Mylan, Par | Uncertain |
| <i>Advicor</i> (Abbott) | Lovastatin/Niacin | Barr ^{f,106} | Uncertain |
| <i>Alocril</i> (Allergan) | Nedocromil ophthalmic | Akorn | Uncertain ^g |
| <i>AndroGel 1%</i> (AbbVie) | Testosterone | Perrigo | Uncertain ²⁵ |
| <i>Avandamet</i> (GlaxoSmithKline) | Rosiglitazone/Metformin | Teva | Uncertain ²⁷ |
| <i>Avandaryl</i> (GlaxoSmithKline) | Rosiglitazone/Glimepiride | Teva | Uncertain |
| <i>Avandia</i> (GlaxoSmithKline) | Rosiglitazone | Actavis, Dr. Reddy's, Mylan, Sandoz, Roxane, Teva, Westward (Hikma) | Uncertain |
| <i>CellCept</i> (Roche Palo) | Mycophenolate mofetil hydrochloride injection | Bedford Labs | Uncertain ^g |
| <i>Copaxone</i> (Teva) | Glatiramer injection | Dr. Reddy's, ^{70,e} Mylan, ^{70,e} Sandoz/Momenta, ^{70,e} Synthron ^{70,82,e} | Uncertain ^{70,82,e,g} |

| Brand ^a (Manufacturer) | Generic Name | Generic Manufacturer(s) ^{b,1} | Anticipated Availability ^c |
|--|---|--|---------------------------------------|
| <i>Dibenzyline</i> (Wellspring) | Phenoxybenzamine | Roxane | Uncertain ^g |
| <i>Exalgo</i> (Mallinckrodt) | Hydromorphone extended-release tablet 32 mg | Actavis ^{f,3} | Uncertain |
| <i>Exelon</i> (Novartis) | Rivastigmine oral solution | Ranbaxy | Uncertain ^g |
| <i>Lamictal ODT</i> (GlaxoSmithKline) | Lamotrigine orally disintegrating tablet | Impax | Uncertain ⁶⁰ |
| <i>Latisse</i> (Allergan) | Bimatoprost 0.03% | Apotex, Sandoz | Uncertain ¹⁰⁸ |
| <i>Megace ES</i> (Par) | Megestrol acetate oral suspension 125 mg/mL | TWi Pharmaceuticals | Uncertain ⁵⁴ |
| <i>Nexium</i> (AstraZeneca) | Esomeprazole magnesium delayed- release capsule | Ranbaxy | Uncertain ¹⁹ |
| <i>Orapred ODT</i> (Concordia) | Prednisolone sodium phosphate orally disintegrating tablet | Mylan | Uncertain ⁴⁵ |
| <i>Renagel</i> (Genzyme) | Sevelamer hydrochloride (oral suspension) | Impax ^{f,15} | Uncertain |
| <i>Renvela</i> (Genzyme) | Sevelamer carbonate (oral suspension) | Impax ^{f,15} | Uncertain |
| <i>Sarafem</i> (Warner Chilcott) | Fluoxetine tablet | Teva | Uncertain ^g |

- This list is not all-inclusive.
- Current through August 2014. These are manufacturers with either approval or tentative approval to market the generic version of the drug unless otherwise noted. For drugs already available, manufacturers with tentative approval are not listed.
- Generic availability is subject to change as a result of litigations and patent exclusivities.
- Ongoing litigation; availability may be sooner than patent expiration date.
- Availability uncertain due to ongoing litigation and regulatory issues. Generic manufacturer has not received approval or tentative approval.
- Generic manufacturer has not received approval or tentative approval from the FDA, but has settled patent litigation with the brand manufacturer.

- g. Patents have expired; however, generics are not yet available.
- h. Generic manufacturer has received approval or tentative approval from the FDA. Generic manufacturer has settled patent litigation with the brand manufacturer. Generic availability may be sooner than patent expiration date.
- i. Generic manufacturer has not received approval or tentative approval from the FDA, but has filed patent challenge.

Users of this PL Detail-Document are cautioned to use their own professional judgment and consult any other necessary or appropriate sources prior to making clinical judgments based on the content of this document. Our editors have researched the information with input from experts, government agencies, and national organizations. Information and internet links in this article were current as of the date of publication.



Project Leader in preparation of this PL Detail-Document: Melanie Cupp, Pharm.D., BCPS

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Evidence and Recommendations You Can Trust...



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**HARVONI
PRIOR AUTHORIZATION**
SD DEPARTMENT OF SOCIAL SERVICES
MEDICAL SERVICES DIVISION

Fax Completed Form to:
866-254-0761
For questions regarding this
Prior authorization, call
866-705-5391

SD Medicaid requires that patients receiving a new prescription for Harvoni must meet the following criteria:

- Patient must be ≥ 18 years old.
- Must have a diagnosis of chronic hepatitis C (genotype 1).
- Liver biopsy confirming a Metavir score of F3 or F4, unless medically contraindicated; or documentation of severe extrahepatic manifestations of hepatitis C infection.
- Must be prescribed by a hepatologist, gastroenterologist, or infectious disease specialist.
- Absence of renal impairment (eGFR must be >30mL/min/1.73m²) and absence of end stage renal disease (ESRD).
- Documentation showing that patient is drug and alcohol free for the past 6 months.
- The concomitant use of Harvoni and P-gp inducers (rifampin, St. John's wort), certain anticonvulsants, certain antiretrovirals, and rosuvastatin is not recommended.

Part I: RECIPIENT INFORMATION (To be completed by physician's representative or pharmacy):

| | | |
|-----------------|---------------------|--------------------------|
| RECIPIENT NAME: | MEDICAID ID NUMBER: | RECIPIENT DATE OF BIRTH: |
|-----------------|---------------------|--------------------------|

Part II: PHYSICIAN INFORMATION (To be completed by physician's representative or pharmacy):

| | | |
|-----------------|-----------------------|---------------------|
| PHYSICIAN NAME: | PHYSICIAN DEA NUMBER: | NAME OF SPECIALIST: |
| CITY: | PHONE: () | FAX: () |

Part III: TO BE COMPLETED BY PHYSICIAN:

| | | | |
|---|--|--|--|
| Requested Drug: <input type="checkbox"/> Harvoni Dosage: _____ | Diagnosis for this request: Genotype: | Documented liver fibrosis: <input type="checkbox"/> YES <input type="checkbox"/> NO | Patient is drug and alcohol free for past 6 months: <input type="checkbox"/> YES <input type="checkbox"/> NO eGFR: |
| Has the patient been previously treated for chronic hepatitis C? <input type="checkbox"/> YES <input type="checkbox"/> NO If yes, please indicate past treatment regimen(s), dates of treatment, and response to therapy: | | Baseline HCV RNA: HCV RNA 4 weeks after starting therapy: | |
| PHYSICIAN SIGNATURE: | | | DATE: |

Part IV: PHARMACY INFORMATION

| | |
|----------------|------------------------------|
| PHARMACY NAME: | SD MEDICAID PROVIDER NUMBER: |
| PHONE: (): | FAX: () |
| DRUG: | NDC# |

Part V: FOR OFFICIAL USE ONLY

| | |
|--|---|
| Date: / / | Initials: _____ |
| Approved - Effective dates of PA: From: / / | To: / / |
| Denied: (Reasons) | |



SOVALDI
PRIOR AUTHORIZATION
SD DEPARTMENT OF SOCIAL SERVICES
MEDICAL SERVICES DIVISION

Fax Completed Form to:
866-254-0761
For questions regarding this
Prior authorization, call
866-705-5391

SD Medicaid requires that patients receiving a new prescription for Sovaldi must meet the following criteria:

- Patient must be ≥ 18 years old.
Must have a diagnosis of chronic hepatitis C (genotypes 1, 2, 3, or 4).
Liver biopsy confirming a Metavir score of F3 or F4, unless medically contraindicated; or documentation of severe extrahepatic manifestations of hepatitis C infection.
Must be prescribed by a hepatologist, gastroenterologist, or infectious disease specialist.
Must be used in combination with ribavirin or in combination with pegylated interferon and ribavirin. (must not be used as monotherapy)
Female patients must have a negative pregnancy test within 30 days prior to initiation of therapy and monthly during treatment.
Absence of renal impairment (eGFR must be >30mL/min/1.73m²) and absence of end stage renal disease (ESRD).
Documentation showing that patient is drug and alcohol free for the past 6 months.

Part I: RECIPIENT INFORMATION (To be completed by physician’s representative or pharmacy):

Table with 3 columns: RECIPIENT NAME, MEDICAID ID NUMBER, RECIPIENT DATE OF BIRTH

Part II: PHYSICIAN INFORMATION (To be completed by physician’s representative or pharmacy):

Table with 3 columns: PHYSICIAN NAME, PHYSICIAN DEA NUMBER, NAME OF SPECIALIST; CITY, PHONE, FAX

Part III: TO BE COMPLETED BY PHYSICIAN:

Complex form with multiple rows and columns for drug details, diagnosis, liver fibrosis, pregnancy test, eGFR, treatment history, and signature/Date.

Part IV: PHARMACY INFORMATION

Table with 2 columns: PHARMACY NAME, PHONE, DRUG; SD MEDICAID PROVIDER NUMBER, FAX, NDC#

Part V: FOR OFFICIAL USE ONLY

Form with fields for Date, Initials, Approved - Effective dates of PA (From/To), Denied: (Reasons)



**OLYSIO
PRIOR AUTHORIZATION**
SD DEPARTMENT OF SOCIAL SERVICES
MEDICAL SERVICES DIVISION

**Fax Completed Form to:
866-254-0761**
**For questions regarding this
Prior authorization, call
866-705-5391**

SD Medicaid requires that patients receiving a new prescription for Olysio must meet the following criteria:

- Patient must be ≥ 18 years old.
- Must have a diagnosis of chronic hepatitis C, genotype 1.
- Liver biopsy confirming a Metavir score of F3 or F4, unless medically contraindicated; or documentation of severe extrahepatic manifestations of hepatitis C infection.
- Must be prescribed by a hepatologist, gastroenterologist, or infectious disease specialist.
- Must be used in combination with pegylated interferon and ribavirin. **(must not be used as monotherapy)**
- Alternative therapy should be considered for patients infected with HCV genotype 1a containing the Q80K polymorphism.
- Female patients must have a negative pregnancy test within 30 days prior to initiation of therapy and monthly during treatment.
- Documentation showing that patient is drug and alcohol free for the past 6 months.

Part I: RECIPIENT INFORMATION (To be completed by physician's representative or pharmacy):

| | | |
|-----------------|---------------------|-------------------------|
| RECIPIENT NAME: | MEDICAID ID NUMBER: | RECIPIENT DATE OF BIRTH |
|-----------------|---------------------|-------------------------|

Part II: PHYSICIAN INFORMATION (To be completed by physician's representative or pharmacy):

| | | |
|-----------------|-----------------------|---------------------|
| PHYSICIAN NAME: | PHYSICIAN DEA NUMBER: | NAME OF SPECIALIST: |
| CITY: | PHONE: () | FAX: () |

Part III: TO BE COMPLETED BY PHYSICIAN:

| | | | |
|--|--|---|---|
| Requested Drug: <input type="checkbox"/> Olysio | Presence of Q80K polymorphism? <input type="checkbox"/> YES <input type="checkbox"/> NO | Diagnosis for this request: Genotype: | Patient is drug and alcohol free for past 6 months: <input type="checkbox"/> YES <input type="checkbox"/> NO |
| Dosage: _____ | Documented liver fibrosis: | Pegylated interferon dose: Ribavirin dose: | Negative pregnancy test in the past 30 days <input type="checkbox"/> YES <input type="checkbox"/> NO |
| Has the patient been previously treated for chronic hepatitis C? <input type="checkbox"/> YES <input type="checkbox"/> NO | | | Baseline HCV RNA: |
| If yes, please indicate past treatment regimen(s), dates of treatment, and response to therapy: | | | HCV RNA 4 weeks after starting therapy: |
| PHYSICIAN SIGNATURE: | | | DATE: |

Part IV: PHARMACY INFORMATION

| | |
|----------------|------------------------------|
| PHARMACY NAME: | SD MEDICAID PROVIDER NUMBER: |
| PHONE: (): | FAX: () |
| DRUG: | NDC#: |

Part V: FOR OFFICIAL USE ONLY

| | |
|--|---|
| Date: / / | Initials: _____ |
| Approved - Effective dates of PA: From: / / | To: / / |
| Denied: (Reasons) | |

| SD Medicaid Sovaldi/Olysio/Harvoni Utilization | | | |
|---|---------------|------------------------|----------------------------|
| 01/01/14 – 10/29/14 | | | |
| Label Name | Rx Num | Total Reimb Amt | Avg Cost per Script |
| SOVALDI 400 MG TABLET | 53 | \$1,548,718.96 | \$29,221.11 |
| OLYSIO 150 MG CAPSULE | 11 | \$255,698.30 | \$23,245.30 |
| HARVONI 90-400 MG TABLET | 0 | 0 | 0 |
| TOTAL | 64 | \$1,804,417.26 | 18 recipients |



**ADULT ADD/ADHD
PRIOR AUTHORIZATION**
SD DEPARTMENT OF SOCIAL SERVICES
MEDICAL SERVICES DIVISION

**Fax Completed Form to:
866-254-0761**
**For questions regarding this
Prior authorization, call
866-705-5391**

SD Medicaid requires that adult patients receiving a new prescription for ADHD therapies must meet the following criteria:

- Patient must be 18 years of age or older and have a documented diagnosis of adult ADD or ADHD
- Patient was diagnosed before the age of 16 and continues to have significant symptoms warranting treatment in adulthood
- Patient must have a specialist involved in therapy experienced in the diagnosis and treatment of Adult ADD/ADHD

Part I: RECIPIENT INFORMATION (To be completed by physician's representative or pharmacy):

| | | |
|-----------------|---------------------|-------------------------|
| RECIPIENT NAME: | MEDICAID ID NUMBER: | RECIPIENT DATE OF BIRTH |
| | | |

Part II: PHYSICIAN INFORMATION (To be completed by physician's representative or pharmacy):

| | | |
|-----------------|-----------------------|---------------------------------|
| PHYSICIAN NAME: | PHYSICIAN DEA NUMBER: | SPECIALIST INVOLVED IN THERAPY: |
| | | |
| CITY: | PHONE: () | FAX: () |
| | | |

Part III: TO BE COMPLETED BY PHYSICIAN:

| | | |
|---|-----------------------------|----------------------------------|
| Requested Drug: | Diagnosis for this Request: | Was the diagnosis before age 16: |
| | | |
| List symptoms significantly impacting, impairing, or compromising the patient's ability to function normally: | | |
| | | |
| PHYSICIAN SIGNATURE: | | DATE: |
| | | |

Part IV: PHARMACY INFORMATION

| | |
|----------------|------------------------------|
| PHARMACY NAME: | SD MEDICAID PROVIDER NUMBER: |
| | |
| PHONE: (): | FAX: () |
| | |
| DRUG: | NDC#: |
| | |

Part V: FOR OFFICIAL USE ONLY

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|---|------------------------|
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| Approved - | |
| Effective dates of PA: From: | / / |
| To: | / / |
| Denied: (Reasons) | |
| | |



EVZIO
PRIOR AUTHORIZATION
 SD DEPARTMENT OF SOCIAL SERVICES
 MEDICAL SERVICES DIVISION

| |
|--|
| Fax Completed Form to: 866-254-0761 For questions regarding this Prior authorization, call 866-705-5391 |
|--|

SD Medicaid requires that adult patients receiving a new prescription for Evzio must meet the following criteria:

- Patient must be taking opioids (over 100mg of morphine equivalents daily) or taking opioids with other interacting medications (benzos, muscle relaxants, etc.)

Part I: RECIPIENT INFORMATION (To be completed by physician's representative or pharmacy):

| | | |
|-----------------|---------------------|-------------------------|
| RECIPIENT NAME: | MEDICAID ID NUMBER: | RECIPIENT DATE OF BIRTH |
| | | |

Part II: PHYSICIAN INFORMATION (To be completed by physician's representative or pharmacy):

| | | |
|-----------------|--|--|
| PHYSICIAN NAME: | PHYSICIAN DEA NUMBER: | |
| | | |
| CITY: | PHONE: () () - () () | FAX: () () - () () |
| | | |

Part III: TO BE COMPLETED BY PHYSICIAN:

| | | |
|-------------------------------------|-----------------------------|--|
| Requested Drug: | Diagnosis for this Request: | Is patient opioid dependent and/or considered high risk for opioid overdose: <input type="checkbox"/> Yes <input type="checkbox"/> No |
| List other interacting medications: | | |
| PHYSICIAN SIGNATURE: | | DATE: |
| | | |

Part IV: PHARMACY INFORMATION

| | |
|---|---|
| PHARMACY NAME: | SD MEDICAID PROVIDER NUMBER: |
| | |
| PHONE: () () - () (): | FAX:: () () - () () |
| | |
| DRUG: | NDC#: |
| | |

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|---|---|
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| | |
| Approved - Effective dates of PA: From: / / | To: / / |
| Denied: (Reasons) | |



OTEZLA
PRIOR AUTHORIZATION
 SD DEPARTMENT OF SOCIAL SERVICES
 MEDICAL SERVICES DIVISION

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 For questions regarding this
 Prior authorization, call
866-705-5391

SD Medicaid requires that adult patients receiving a new prescription for Otezla must meet the following criteria:

- Patient must have psoriatic arthritis.
- Patient must be 18 years or older.
- Patient must first try Enbrel or Humira.

Part I: RECIPIENT INFORMATION (To be completed by physician's representative or pharmacy):

| | | |
|-----------------|---------------------|-------------------------|
| RECIPIENT NAME: | MEDICAID ID NUMBER: | RECIPIENT DATE OF BIRTH |
| | | |

Part II: PHYSICIAN INFORMATION (To be completed by physician's representative or pharmacy):

| | | |
|-----------------|-----------------------|-------------|
| PHYSICIAN NAME: | PHYSICIAN DEA NUMBER: | |
| | | |
| CITY: | PHONE: () | FAX: () |
| | | |

Part III: TO BE COMPLETED BY PHYSICIAN:

| | | |
|----------------------|-----------------------------|--|
| Requested Drug: | Diagnosis for this Request: | Previous trial: Drug: Date of trial: |
| | | |
| PHYSICIAN SIGNATURE: | | DATE: |
| | | |

Part IV: PHARMACY INFORMATION

| | |
|----------------|------------------------------|
| PHARMACY NAME: | SD MEDICAID PROVIDER NUMBER: |
| | |
| PHONE: (): | FAX:: () |
| | |
| DRUG: | NDC#: |
| | |

Part V: FOR OFFICIAL USE ONLY

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|---|---|
| Date: / / | Initials: _____ |
| Approved - Effective dates of PA: From: / / | To: / / |
| Denied: (Reasons) | |
| | |



**HIGH COST MEDICATIONS
PRIOR AUTHORIZATION**
SD DEPARTMENT OF SOCIAL SERVICES
MEDICAL SERVICES DIVISION

| |
|---|
| <p align="center">Fax Completed Form to: 866-254-0761 For questions regarding this Prior authorization, call 866-705-5391</p> |
|---|

SD Medicaid requires that patients receiving a new prescription that exceeds \$5,000 must meet the following criteria:

- The medication must be FDA approved and used for a FDA-approved indication listed on the manufacturer's package insert.
- The prescriber must document a clear clinical advantage for utilizing the new medication if existing treatments are available.
- If alternative treatments exist, the prescriber must document the patient's trial and failure of these existing treatments.
- The medication must be available from a SD Medicaid enrolled pharmacy.
- The patient must be able to self-administer the medication.
- The medication must not be experimental or available as part of a clinical trial.

Part I: RECIPIENT INFORMATION (To be completed by physician's representative or pharmacy):

| | | |
|-----------------|---------------------|-------------------------|
| RECIPIENT NAME: | MEDICAID ID NUMBER: | RECIPIENT DATE OF BIRTH |
| | | |

Part II: PHYSICIAN INFORMATION (To be completed by physician's representative or pharmacy):

| | | |
|-----------------|-----------------------|----------|
| PHYSICIAN NAME: | PHYSICIAN DEA NUMBER: | |
| | | |
| CITY: | PHONE: () | FAX: () |
| | | |

Part III: TO BE COMPLETED BY PHYSICIAN:

| | | |
|--|-----------------------------|---|
| Requested Drug: | Diagnosis for this Request: | Will patient self-administer this drug: <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | | |
| Document patient's trial and failure of existing alternative treatments: | | |
| | | |
| PHYSICIAN SIGNATURE: | DATE: | |
| | | |

Part IV: PHARMACY INFORMATION

| | |
|----------------|------------------------------|
| PHARMACY NAME: | SD MEDICAID PROVIDER NUMBER: |
| | |
| PHONE: (): | FAX: () |
| | |
| DRUG: | NDC#: |
| | |

Part V: FOR OFFICIAL USE ONLY

| | |
|---|---|
| Date: / / | Initials: _____ |
| Approved - Effective dates of PA: From: / / | To: / / |
| Denied: (Reasons) | |
| | |

PRODUCT DETAILS OF GLP-1 AGONISTS (GLUCAGON-LIKE PEPTIDE-1)

INDICATIONS AND USE: GLP-1 receptor agonists are indicated as adjuncts to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus. These agents should not be used as initial monotherapy. For most patients, GLP-1 agonists should be reserved for those who require two or more diabetes medications to maintain a desired A1C.

COMPARISON:

| GLP-1 Agonist | ~ A1C Decrease | ~ Weight Loss | ~ Cost/month | Dosing Frequency |
|-------------------------|----------------|---------------|--------------|------------------|
| Byetta (exenatide) | 1% | 4 lbs | \$430 | Twice daily |
| Bydureon (exenatide ER) | 1.5% | 6 lbs | \$440 | Once weekly |
| Tanzeum (albiglutide) | 1% | 2 lbs | \$330 | Once weekly |
| Trulicity (dulaglutide) | 1.5% | 6 lbs | \$500 | Once weekly |
| Victoza (liraglutide) | 1.5% | 6 lbs | \$400-600 | Once daily |

HOW SUPPLIED:

| GLP-1 Agonist | Package Size |
|-------------------------|---|
| Byetta (exenatide) | 5 mcg per dose, 60 doses (1.2 mL prefilled pen). 10 mcg per dose, 60 doses (2.4 mL prefilled pen). |
| Bydureon (exenatide ER) | Four single-dose trays containing 2 mg vial. Four single-dose pens containing 2 mg pen. |
| Tanzeum (albiglutide) | 30 mg single-dose pen and 50 mg single-dose pen. (carton of 4) |
| Trulicity (dulaglutide) | Carton of 4 single-dose pen or prefilled syringe: (0.75 mg/0.5 mL and 1.5 mg/0.5 mL). |
| Victoza (liraglutide) | Disposable, pre-filled, multi-dose pens delivering doses of 0.6mg, 1.2mg, or 1.8mg (6 mg/mL, 3mL). |

DOSAGE AND ADMINISTRATION:

| GLP-1 Agonist | Dosage and Administration |
|-------------------------|---|
| Byetta (exenatide) | <ul style="list-style-type: none"> Inject subcutaneously within 60 minutes prior to morning and evening meals (or before the two main meals of the day, approximately 6 hours or more apart). Initiate at 5 mcg per dose twice daily; increase to 10 mcg twice daily after 1 month based on clinical response. |
| Bydureon (exenatide ER) | <ul style="list-style-type: none"> Administer 2 mg by subcutaneous injection once every seven days (weekly), at any time of day and with or without meals. Administer immediately after the dose is prepared. |
| Tanzeum (albiglutide) | <ul style="list-style-type: none"> Administer once weekly at any time of day, without regard to meals. Inject subcutaneously in the abdomen, thigh, or upper arm. Initiate at 30 mg subcutaneously once weekly. Dose can be increased to 50 mg once weekly in patients requiring additional glycemic control. If a dose is missed, administer within 3 days of missed dose. |
| Trulicity (dulaglutide) | <ul style="list-style-type: none"> Administer once weekly at any time of day. Inject subcutaneously in the abdomen, thigh, or upper arm. Initiate at 0.75 mg subcutaneously once weekly. Dose can be increased to 1.5 mg once weekly for additional glycemic control. If a dose is missed, administer within 3 days of missed dose. |

| GLP-1 Agonist | Dosage and Administration |
|-----------------------|--|
| Victoza (liraglutide) | <ul style="list-style-type: none"> Administer once daily at any time of day, independently of meals. Inject subcutaneously in the abdomen, thigh, or upper arm. The injection site and timing can be changed without dose adjustment. Initiate at 0.6 mg per day for one week. This dose is intended to reduce gastrointestinal symptoms during initial titration, and is not effective for glycemic control. After one week, increase the dose to 1.2 mg. If the 1.2 mg dose does not result in acceptable glycemic control, the dose can be increased to 1.8 mg. |

SPECIAL POPULATIONS:

| GLP-1 Agonist | Special Populations |
|-------------------------|--|
| Byetta (exenatide) | <ul style="list-style-type: none"> Byetta should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus. Caution should be exercised when Byetta is administered to a nursing woman. |
| Bydureon (exenatide ER) | <ul style="list-style-type: none"> Bydureon should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus. Caution should be exercised when Bydureon is administered to a nursing woman. |
| Tanzeum (albiglutide) | <ul style="list-style-type: none"> Tanzeum should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus. Discontinue nursing or discontinue Tanzeum. Monitor renal function in patients with renal impairment reporting severe adverse gastrointestinal reactions. |
| Trulicity (dulaglutide) | <ul style="list-style-type: none"> Trulicity should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus. Discontinue nursing or discontinue Trulicity. Monitor renal function in patients with renal impairment reporting severe adverse gastrointestinal reactions. |
| Victoza (liraglutide) | <ul style="list-style-type: none"> Victoza should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus. Discontinue nursing or discontinue Victoza. |

WARNINGS AND PRECAUTIONS:

| GLP-1 Agonist | Warnings and Precautions |
|--------------------|--|
| Byetta (exenatide) | <ul style="list-style-type: none"> Never share a Byetta pen between patients, even if the needle is changed. Postmarketing reports with Byetta include fatal and non-fatal hemorrhagic or necrotizing pancreatitis. Discontinue promptly. Consider other antidiabetic therapies in patients with a history of pancreatitis. Increased risk of hypoglycemia when used in combination with medications known to cause hypoglycemia (e.g., insulin or insulin secretagogue). Consider reducing the dose of insulin or insulin secretagogue. Byetta should not be used in patients with severe renal impairment or end-stage renal disease and should be used with caution in patients with renal transplantation. Caution should be applied when initiating Byetta in patients with moderate renal failure. |

| GLP-1 Agonist | Warnings and Precautions |
|-------------------------|--|
| | <ul style="list-style-type: none"> • Use of Byetta is not recommended in patients with severe gastrointestinal disease. • Patient should discontinue Byetta if symptoms of hypersensitivity reactions (e.g., anaphylaxis and angioedema) arise. |
| Bydureon (exenatide ER) | <ul style="list-style-type: none"> • Counsel patients regarding the risk of medullary thyroid carcinoma and the symptoms of thyroid tumors. • Discontinue promptly if pancreatitis is suspected. Do not restart if pancreatitis is confirmed. • Increased risk of hypoglycemia when Bydureon is used in combination with a sulfonylurea. Consider reducing the sulfonylurea dose. • Not recommended if patient has severe renal impairment or end-stage renal disease. Use with caution in patients with renal-transplantation or moderate renal impairment. • Not recommended in patients with severe gastrointestinal disease. • Patient should discontinue Bydureon if symptoms of hypersensitivity reactions (e.g., anaphylaxis and angioedema) arise. |
| Tanzeum (albiglutide) | <ul style="list-style-type: none"> • Discontinue promptly if pancreatitis is suspected. Do not restart if confirmed. • Hypoglycemia can occur when used in combination with insulin secretagogues (e.g., sulfonylureas) or insulin. Consider lowering sulfonylurea or insulin dosage when starting Tanzeum. • Discontinue Tanzeum if hypersensitivity reactions are suspected. • Monitor renal function in patients with renal impairment reporting severe adverse gastrointestinal reactions. |
| Trulicity (dulaglutide) | <ul style="list-style-type: none"> • Counsel patients regarding the risk of medullary thyroid carcinoma and the symptoms of thyroid tumors. • Discontinue promptly if pancreatitis is suspected. Do not restart if pancreatitis is confirmed. • Increased risk of hypoglycemia when Trulicity is used with an insulin secretagogue or insulin, consider lowering the dose of sulfonylurea or insulin to reduce the risk. • Discontinue Trulicity if hypersensitivity reactions are suspected. • Monitor renal function in patients with renal impairment reporting severe adverse gastrointestinal reactions. |
| Victoza (liraglutide) | <ul style="list-style-type: none"> • Counsel patients regarding the risk of medullary thyroid carcinoma and the symptoms of thyroid tumors. • Discontinue promptly if pancreatitis is suspected. Do not restart if pancreatitis is confirmed. • Increased risk of hypoglycemia when Victoza is used with an insulin secretagogue or insulin, consider lowering the dose of the insulin secretagogue or insulin to reduce the risk. • Use caution when initiating or escalating doses of Victoza in patients with renal impairment. • Discontinue Victoza if hypersensitivity reactions are suspected. |

ADVERSE REACTIONS:

| GLP-1 Agonist | Adverse Reactions |
|-------------------------|--|
| Byetta (exenatide) | <ul style="list-style-type: none"> • Most common ($\geq 5\%$) and occurring more frequently than placebo in clinical trials: nausea, hypoglycemia, vomiting, diarrhea, feeling jittery, dizziness, headache, dyspepsia, constipation, and asthenia. • Postmarketing reports with exenatide of increased international normalized ratio (INR) with concomitant use of warfarin, sometimes with bleeding. |
| Bydureon (exenatide ER) | <ul style="list-style-type: none"> • Most common ($\geq 5\%$) and occurring more frequently than comparator in clinical trials: nausea, diarrhea, headache, vomiting, constipation, injection-site pruritus, injection-site nodule, and dyspepsia. |
| Tanzeum (albiglutide) | <ul style="list-style-type: none"> • Adverse reactions, reported in $\geq 10\%$ of patients and more frequently than in patients on placebo, were upper respiratory tract infection, diarrhea, nausea, and injection site reaction. |
| Trulicity (dulaglutide) | <ul style="list-style-type: none"> • The most common adverse reactions, reported in $\geq 5\%$ of patients are nausea, diarrhea, vomiting, abdominal pain, and decreased appetite. |
| Victoza (liraglutide) | <ul style="list-style-type: none"> • The most common adverse reactions, reported in $\geq 5\%$ of patients and occurring more frequently than in patients treated with placebo, were headache, nausea, diarrhea, and anti-liraglutide antibody formation. • Immunogenicity-related events, including urticarial, were more common among Victoza-treated patients (0.8%) than among comparator-treated patients (0.4%) in clinical trials. |

UTILIZATION:

| Label Name | Rx Num | Total Reimb Amt | Avg Cost per Script |
|------------------------|---------------|------------------------|----------------------------|
| Byetta 5 mcg Dose Pen | 5 | \$1,994.49 | \$398.90 |
| Byetta 10 mcg Dose Pen | 37 | \$15,149.39 | \$409.44 |
| Bydureon 2 mg Pen | 2 | \$921.02 | \$460.51 |
| Bydureon 2 mg Vial | 87 | \$37,112.78 | \$426.58 |
| Tanzeum 30 mg Pen | 2 | \$660.60 | \$330.30 |
| Victoza 2-Pak | 129 | \$48,173.07 | \$373.43 |
| Victoza 3-Pak | 220 | \$123,903.33 | \$563.20 |
| Total (97 recipients) | 482 | \$227,914.68 | |

References:

1. PL Detail-Document, Comparison of GLP-1 Agonists. Pharmacist's Letter/Prescriber's Letter. December 2014.
2. Byetta [package insert]. Princeton, NJ: Bristol-Myers Squibb Company; August 2014.
3. Bydureon [package insert]. Wilmington, DE: AstraZeneca Pharmaceuticals LP; June 2014.
4. Tanzeum [package insert]. Research Triangle Park, NC: GlaxoSmithKline LLC; June 2014.
5. Trulicity [package insert]. Indianapolis, IN: Eli Lilly and Company; September 2014.
6. Victoza [package insert]. Plainsboro, NY: Novo Nordisk Inc.; July 2013.

PRODUCT DETAILS OF NEW TOPICAL THERAPIES FOR ONYCHOMYCOSIS

INDICATIONS AND USE:

| | |
|------------------------|---|
| Jublia (efinaconazole) | Topical treatment of onychomycosis of the toenails due to <i>Trichophyton rubrum</i> and <i>Trichophyton mentagrophytes</i> . |
| Kerydin (tavaborole) | Topical treatment of onychomycosis of the toenails due to <i>Trichophyton rubrum</i> or <i>Trichophyton mentagrophytes</i> . |

COMPARISON:

| Drug | Dose | Efficacy | ~ Cost | Comments |
|------------------------|---|--|------------|---|
| Jublia (efinaconazole) | Apply to affected toenails once daily for 48 weeks. | Toenail complete cure rate 17% at week 52. | \$450/4 mL | Consider for patients who can't use oral therapy. |
| Kerydin (tavaborole) | Apply to affected toenails once daily for 48 weeks. | Toenail complete cure rate < 10% at week 52. | \$450/4 mL | Consider for patients who can't use oral therapy. |

HOW SUPPLIED:

| | |
|------------------------|------------------------------------|
| Jublia (efinaconazole) | 4 mL and 8 mL 10% topical solution |
| Kerydin (tavaborole) | 4 mL and 10 mL 5% topical solution |

ADVERSE REACTIONS:

| Drug | Adverse Reactions |
|---------|---|
| Jublia | <ul style="list-style-type: none"> The most common adverse reactions (incidence >1%) were ingrown toenails, application site dermatitis, application site vesicles, and application site pain. |
| Kerydin | <ul style="list-style-type: none"> Common adverse reactions occurring in ≥1% in subjects included application site exfoliation, ingrown toenail, application site erythema, and application site dermatitis. |

References:

1. PL Detail-Document, Comparison of Topical Therapies for Onychomycosis. Pharmacist's Letter/Prescriber's Letter. July 2014.
2. Jublia [package insert]. Bridgewater, NY: Valeant Pharmaceuticals North America, LLC; June 2014.
3. Kerydin [package insert]. Palo Alto, CA: Anacor Pharmaceuticals, Inc.; July 2014.