

STATE OF WEST VIRGINIA DEPARTMENT OF HEALTH AND HUMAN RESOURCES

Joe Manchin III Governor

Bureau for Medical Services Office of Pharmacy Services 350 Capitol Street - Room 251 Charleston, West Virginia 25301-3706 Phone: (304) 558-1700 - Fax: (304) 558-1542

Patsy A. Hardy, FACHE, MSN, MBA **Cabinet Secretary**

Pharmaceutical and Therapeutics (P&T) Committee August 25, 2010 Diamond Building Rooms B10 & B11

Charleston, West Virginia

MINUTES

Members Present:

David Avery, M.D. James D. Bartsch, R.Ph. Rodney L. Fink, D.O. Scott Brown, R.Ph. Michael Grome, PA-C Robert Stanton, Pharm.D. Teresa Dunsworth, Pharm.D. Teresa Frazer, M.D., FAAP Steven R. Matulis, M.D.

Members Not Present:

Jeffrey V. Ashley, M.D. Harriet Nottingham, R.Ph. Barbara Koster, N.P.

DHHR/BMS Staff Present:

Peggy King, R.Ph., Pharmacy Director Gail Goodnight, R.Ph. Rebate Coordinator Vicki Cunningham, R.Ph., DUR Coordinator William Hopkins, Pharmacy Operations Manager Lynda Edwards, Secretary

Contract Staff/GHS Staff Present:

Laureen Biczak, D.O. Tim Clifford, M.D. Chad Bissell, Pharm.D. Kim Curtis, R. Ph. Shelagh Harvard Theresa Thompson

Other Contract Staff/State Staff Present:

Stephen Small, R.Ph., M.S., Rational Drug Therapy Program Eric Sears, R.Ph., Unisys

I. Call to Order

Dr. David Avery, M.D., Chairperson, called the meeting to order at 2:10 p.m.

II. Welcome and Introductions

All parties seated at the table introduced themselves and gave a brief statement about their professional credentials and affiliations.

III. Healthcare Reform: Impact on Medicaid Drug Rebates and PDL

Dr. Clifford reviewed Healthcare Reform and the implications for West Virginia Medicaid drug rebates.

IV. Public Comments

<u>Dr. Mark Casdorph, Novartis</u>, spoke in favor of keeping a preferred long-acting methylphenidate product on the PDL, specifically Focalin XR, that could be sprinkled onto food for patients who had difficulty swallowing tablets or capsules.

Dr. Philip Fisher, Endo, spoke in favor of Opana ER.

Linda Lauer, AstraZeneca, spoke in favor of Seroquel XR.

V. Executive Session

The Committee adjourned to Executive Session at 2:37 and resumed open session at 3:35.

VI. Recommended PDL Changes Based on Healthcare Reform: Line Extension Drugs

The following drugs were called for extraction: Focalin XR, Keppra XR, Opana ER, and Seroquel XR.

Dr. Avery called for a motion on those drugs that were not extracted. A motion was made to accept the proposed PDL status of all non-extracted drugs as recommended by GHS. The motion was seconded and passed.

P&T Committee Minutes August 25, 2010 Page 3 of 6

The following PDL changes were approved:

A. Aricept ODT to Non-preferred

PREFERRED AGENTS	NON-PREFERRED AGENTS
ARICEPT (donepezil) EXELON (rivastigmine)	ARICEPT ODT(donepezil) COGNEX (tacrine) galantamine galantamine ER RAZADYNE (galantamine)
	RAZADYNE ER (galantamine) rivastigmine

B. Detrol LA and Sanctura XR to Non-preferred, and Toviaz to Preferred

PREFERRED AGENTS	NON-PREFERRED AGENTS
ENABLEX (darifenacin)	DETROL (tolterodine)
oxybutynin	DETROL LA (tolterodine)
oxybutynin ER	DITROPAN (oxybutynin)
SANCTURA (trospium)	DITROPAN XL (oxybutynin)
TOVIAZ (fesoterodine)	GELNIQUE (oxybutynin)
VESICARE (solifenacin)	OXYTROL (oxybutynin)
	SANCTURA XR (trospium)

C. Lescol XL to Non-preferred

PREFERRED AGENTS	NON-PREFERRED AGENTS
CRESTOR (rosuvastatin)	ALTOPREV (lovastatin)
LESCOL (fluvastatin)	LESCOL XL (fluvastatin)
LIPITOR (atorvastatin)	LIVALO (pitavastatin)
lovastatin	MEVACOR (lovastatin)
pravastatin	PRAVACHOL (pravastatin)
simvastatin	ZOCOR (simvastatin)

D. Maxalt MLT to Non-preferred, and naratriptan to Preferred

PREFERRED AGENTS	NON-PREFERRED AGENTS
IMITREX NASAL SPRAY(sumatriptan) IMITREX INJECTION (sumatriptan) naratriptan sumatriptan	AMERGE (naratriptan) AXERT (almotriptan) FROVA (frovatriptan) IMITREX tablets (sumatriptan) MAXALT (rizatriptan) MAXALT MLT (rizatriptan)
	RELPAX (eletriptan) sumatriptan nasal spray/injection* ZOMIG (zolmitriptan)

E. Daytrana to Preferred

PREFERRED AGENTS	NON-PREFERRED AGENTS
CONCERTA (methylphenidate) DAYTRANA (methylphenidate) FOCALIN (dexmethylphenidate) FOCALIN XR (dexmethylphenidate) METADATE CD (methylphenidate) methylphenidate methylphenidate ER STRATTERA (atomoxetine)	dexmethylphenidate INTUNIV (guanfacine) METADATE ER (methylphenidate) NUVIGIL (armodafinil) pemoline PROVIGIL (modafinil) RITALIN (methylphenidate) RITALIN LA (methylphenidate)
OTTATIETA (atomoxetine)	RITALIN-SR (methylphenidate)

The following drugs were discussed:

A. Focalin XR

GHS initially recommended that Focalin XR be made a non-preferred drug in the Stimulants and Related Agents, Non-Amphetamine category. Based on updated information, GHS amended their recommendation to keep Focalin XR preferred for the time being. All strengths will be reviewed during the October 27, 2010 meeting. Dr. Fink spoke against the recommendation of GHS by expressing concern about the cost to the State due to line extensions. Dr. Frazer spoke in favor of keeping Focalin XR preferred to minimize the disruption to children during the school year. A motion was made to accept the amended recommendation of GHS. The motion was seconded, votes were taken and the motion carried.

PREFERRED AGENTS	NON-PREFERRED AGENTS
CONCERTA (methylphenidate)	dexmethylphenidate
DAYTRANA (methylphenidate)	INTUNIV (guanfacine)
FOCALIN (dexmethylphenidate)	METADATE ER (methylphenidate)
FOCALIN XR (dexmethylphenidate)	NUVIGIL (armodafinil)
METADATE CD (methylphenidate)	pemoline
methylphenidate	PROVIGIL (modafinil)
methylphenidate ER	RITALIN (methylphenidate)
STRATTERA (atomoxetine)	RITALIN LA (methylphenidate)
,	RITALIN-SR (methylphenidate)

B. Keppra XR

GHS recommended that Keppra XR be made a non-preferred drug in the Anticonvulsants, Adjuvants category. GHS further recommended grandfathering those patients with a seizure disorder. Mr. Bartsch asked whether there was a specific timeframe attached to the grandfathering recommendation. Dr. Biczak replied that there was no timeframe included. A motion was made to accept the recommendation of GHS with grandfathering recommendation to the Bureau. The motion was seconded, votes were taken and the motion carried.

PREFERRED AGENTS	NON-PREFERRED AGENTS
ADJUVANTS	
carbamazepine	BANZEL(rufinamide)

CARBATROL (carbamazepine) DEPAKOTE SPRINKLE (divalproex) divalproex EC divalproex ER divalproex DR EPITOL (carbamazepine) FELBATOL (felbamate) gabapentin GABITRIL (tiagabine) levetiracetam lamotrigine lamotrigine chewable LYRICA (pregabalin) oxcarbazepine tablets topiramate TRILEPTAL SUSPENSION (oxcarbazepine) valproic acid zonisamide	carbamazepine XR DEPAKENE (valproic acid) DEPAKOTE (divalproex) DEPAKOTE ER (divalproex) EQUETRO (carbamazepine) FANATREX SUSPENSION (gabapentin) KEPPRA (levetiracetam) KEPPRA XR (levetiracetam) LAMICTAL (lamotrigine) LAMICTAL CHEWABLE (lamotrigine) LAMICTAL ODT (lamotrigine) LAMICTAL XR (lamotrigine) NEURONTIN (gabapentin) SABRIL (vigabatrin) STAVZOR (valproic acid) TEGRETOL (carbamazepine) TEGRETOL XR (carbamazepine) TOPAMAX (topiramate) TRILEPTAL TABLETS (oxcarbazepine) VIMPAT (lacosamide) ZONEGRAN (zonisamide)
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C. Opana ER

GHS recommended that Opana ER be made a non-preferred drug in the Analgesics, Narcotic - Long Acting (Non-parenteral) category. GHS further recommended grandfathering cancer patients for 60 days, allowing for a long transition to a preferred long-acting narcotic. Mr. Brown addressed the general need for more options in this PDL category. Dr. Avery agreed with Mr. Brown. A motion was made to accept the recommendation of GHS. The motion was seconded, votes were taken and the motion carried.

PREFERRED AGENTS	NON-PREFERRED AGENTS
fentanyl transdermal KADIAN (morphine) 10mg, 20mg, 30mg, 50mg, 60mg, 100mg methadone morphine ER	AVINZA (morphine) DOLOPHINE (methadone) DURAGESIC (fentanyl) EXALGO ER (hydromorphone) ^{NR} EMBEDA (morphine/naltrexone) KADIAN (morphine) 80mg, 200mg MS CONTIN (morphine) OPANA ER (oxymorphone) ORAMORPH SR (morphine) oxycodone ER
	OXYCONTIN (oxycodone) RYZOLT ER (tramadol)
	tramadol ER
	ULTRAM ER (tramadol)

D. Seroquel XR

GHS recommended that Seroquel XR be made a non-preferred drug in the Antipsychotics, Atypical (Oral) category. GHS further recommended grandfathering schizophrenia patients for 60 days. Mr. Bartsch inquired as to whether Ms. King felt that 60 days was enough time for patients with schizophrenia. Ms. King explained that the timeframe gives physicians 60 days from the October 1st implementation to request a PA for the patient to be transitioned. Mr. Grome asked if patients may remain on Seroquel XR. Ms. King replied

that it would be possible for patients to remain on the drug as long as their physician made a strong case for not moving the patient to another drug. Ms. King further explained that HID will send a targeted mailing to physicians containing a list of patients currently on the drug. A motion was made to accept the recommendation of GHS. The motion was seconded, votes were taken and the motion carried.

PREFERRED AGENTS	NON-PREFERRED AGENTS
ORAL	
clozapine GEODON (ziprasidone) INVEGA (paliperidone) risperidone risperidone ODT risperidone solution SEROQUEL (quetiapine)	ABILIFY (aripiprazole) CLOZARIL (clozapine) FANAPT (iloperidone) FAZACLO (clozapine) RISPERDAL (risperidone) RISPERDAL ODT (risperidone) RISPERDAL SOLUTION (risperidone) SAPHRIS (asenapine) SEROQUEL XR (quetiapine) ZYPREXA (olanzapine)

Dr. Avery stated that all of the changes voted on will take effect on October 1, 2010.

VII. Other Business

There was no other business.

VIII. Next Meeting Date

The next meeting of the P&T Committee will be held on October 27, 2010 at the Charleston Civic Center.

IX. Adjournment

A motion was made, was seconded, votes were taken and the motion carried to adjourn the meeting of the Pharmaceutical and Therapeutics Committee.