MEDICAID PHARMACY STEP THERAPY PROGRAM

Based on the recommendations of the Drug Utilization Review Board (DURB), prescribers should adhere to the following step therapy parameters. These step therapy parameters have been instituted to ensure clinically appropriate and cost effective use of these drugs and drug classes.

For more information on DUR Board recommendations please refer to DUR meeting summaries at http://nyhealth.gov/health_care/medicaid/program/dur/index.htm

Acthar H.P. Gel (repository corticotropin injection)

| Trial of first-line therapy for all FDA-approved indications other | er than infantile spasms. See below: |
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| Injectable <u>Actha</u> r: FDA Indication | First Line Therapy |
| Corticosteroid or plasmapheresis | MS exacerbations |
| Corticosteroid | Polymyositis/dermatomyositis |
| ACE inhibitor, diuretic, corticosteroid (and for refractory patients: an immunosuppressive) | Idiopathic nephritic syndrome |
| Immunosuppressive, corticosteroid, or ACE inhibitor | Nephrotic syndrome due to systemic lupus erythematosus (SLE) |
| Corticosteroid, topical retinoid, biologic disease- modifying antiheumatic drug, non-biologic DMARD, or a non-steroidal anti-inflammatory (NSAIDS) | Rheumatic disorders (specifically: psoriatic arthritis, juvenile rheumatoid arthritis, ankylosing spondylitis) |
| Corticosteriod, anti-malarial, or cytotoxic/immunosuppressive agent | Systemic lupus erythematosus (SLE) |
| Corticosteroid or analgesic | Dermatologic diseases (specifically Stevens-Johnson syndrome and erythema multiforme) |
| Topical or oral corticosteroid, antihistamine, or an NSAID | Allergic states (specifically serum sickness) |
| Analgesic, anti-infective agent, and agents to reduce inflammation, such as NSAIDS and steroids. | Ophthalmic diseases (keratitis, iritis, iridocyclitits, diffuse posterior uveitis/choroiditis, optic neuritis, choriorentinitis, anterior segment inflammation) |
| Oral corticosteroid or an immunosuppressive | Respiratory diseases (systemic sarcoidosis) |

Confirm diagnosis for Medicaid covered uses. <u>Acthar</u> is first line therapy for infantile spasms in children less than 2 years of age –step therapy not required.

Date Effective 3/21/13

Angiotensin Converting Enzyme Inhibitors (ACEI)/Angiotension Receptor Blockers (ARB)/Direct Renin Inhibitors (DRI) and Related Combination Products

Trial of a product containing an ACEI prior to initiating a preferred ARB. Trial of a product containing either an ACE inhibitor or an ARB prior to initiating a preferred DRI.

View; **Preferred Drug List**

Amitiza

Step therapy with trials of both a bulking-agent and an osmotic laxative prior (defined as within 89 days) to lubiprostone.

Carisoprodol Containing Products

Trial with one (1) preferred analgesic and two (2) preferred skeletal muscle relaxants prior to use of <u>carisoprodol</u> containing products;

carisoprodol carisoprodol/ASA carisoprodol/ASA/codeine Soma® Soma Compound®

Soma Compound with Codeine®

See also; Frequency/Quantity/Duration, Preferred Drug List

Cymbalta

Trial with a tricylic antidepressant OR gabapentin for treatment of Diabetic Peripheral Neuropathy (DPN).

View; Preferred Drug List

Doxycycline delayed-release

Prescribers should attempt treatment with a more cost effective <u>immediate-release doxycycline</u> first before progressing to <u>delayed-release doxycycline</u>.

See also; Frequency/Quantity/Duration, Preferred Drug List

Forteo

Require a trial with a preferred oral bisphosphonate prior to teriparatide.

See also; Frequency/Quantity/Duration

Glucagon-like Peptide-1 Agonists (GLP-1)

Requires a trial with metformin plus another oral antidiabetic agent prior to a GLP-1 agonist.

Prior authorization is required with lack of covered diagnosis in medical history.

See also; Preferred Drug List

Invega®

Trial with <u>risperidone</u> prior to initiation of <u>paliperidone</u> therapy.

See also; Frequency/Quantity/Duration, Preferred Drug List

Lyrica®

Trial with a tricylic antidepressant OR gabapentin for treatment of Diabetic Peripheral Neuropathy (DPN).

See also; Preferred Drug List

Metformin

Require a trial with <u>metformin</u> with or without insulin prior to initiation of other anti-diabetic agents (unless documented contraindication). Anti-diabetic classes affected:

Alpha-Glucosidase Inhibitors

Amylin Analogs

Dipeptidyl Peptidase-4 (DPP-4) Inhibitors

Glucagon-Like Peptide-1 (GLP-1) Agonists

Meglitinides

Sulfonylureas

Thiazolidinediones (TZDs)

See also; Preferred Drug List

Metozolv ODT®

Trial with conventional $\underline{\text{metoclopramide}}$ before $\underline{\text{metoclopramide ODT}}$. To be applied to patients without a diagnosis of diabetes.

See also; Frequency/Quantity/Duration

Date Effective 3/21/13

Moxatag®

Prescribers should attempt treatment with a more cost effective <u>immediate-release amoxicillin</u> first before progressing to extended-release amoxicillin.

See also; Frequency/Quantity/Duration

Non-Fluoroquinolone Ophthalmic Antibiotics

Prescribers should attempt treatment with a preferred product (of comparable coverage) before using a non preferred product.

View; **Preferred Drug List**

Nucynta®

Trial with <u>tramado</u>l and one (1) preferred opioid before <u>tapentadol immediate-release (IR).</u>

See also; Frequency/Quantity/Duration, Preferred Drug List

Nucynta ER®

Trial with tapentadol IR before tapentadol ER for patients who are naïve to a long-acting opioid.

See also; Frequency/Quantity/Duration, Preferred Drug List

Ophthalmic Antibiotic/Steroid Combinations

Prescribers should attempt treatment with a preferred product (of comparable coverage) before using a non preferred product.

View; Preferred Drug List

Ophthalmic Fluoroquinolones

For patients 21 yrs or younger, prescribers should attempt treatment with a non-fluoroquinolone ophthalmic antibiotic before progressing to the following products:

Besivance®

Ciloxan®

ciprofloxacin

IQUIX®

levofloxacin

Moxeza®

Ocuflox®

ofloxacin

Quixin®

Vigamox®

Zymar®

Zymaxid®

View; Preferred Drug List

Protease Inhibitors (Hepatitis C)

Telaprevir:

Step therapy will be applied to guarantee concomitant <u>peginterferon</u> and <u>ribavirin</u> therapy.

Boceprevir:

Step therapy will be applied to guarantee 4 consecutive weeks of <u>peginterferon</u> and <u>ribavirin</u> therapy immediately before initiation of boceprevir.

See also; Frequency/Quantity/Duration, Preferred Drug List

Restasis®

Prescribers are required to document a diagnosis to justify utilization as a first line agent or attempt treatment with an artificial tear/gel/ointment.

See also; Frequency/Quantity/Duration

Selective Serotonin Reuptake Inhibitors (SSRI)/Serotonin-Norepinephrine Reuptake Inhibitors (SNRI)

Require trial with an SSRI prior to an SNRI (exception for specific indications: Chronic musculoskeletal pain (CMP), Diabetic peripheral neuropathy (DPN) and Fibromyalgia (FM).

View; **Preferred Drug List**

Singulair®

For non-asthmatic patients, prescribers should attempt treatment with an intranasal corticosteroid or a 2nd generation oral antihistamine before progressing to <u>montelukast</u>.

View; Preferred Drug List

Topical Anti-Fungals

Prescriber should attempt treatment with a preferred product (of comparable coverage) before using a non-preferred product.

View; Preferred Drug List

Topical Corticosteroids

Prescriber should attempt treatment with a preferred product (of comparable potency) before using a non-preferred product.

View; Preferred Drug List

Tramadol ER

For <u>tramadol</u> naïve patients, prescribers should attempt treatment with the immediate release formulations before progressing to the following extended release formulations:

<u>Conzip</u>

Ryzolt®

tramadol ER

<u>Ultram ER</u>®

See also; Frequency/Quantity/Duration, Preferred Drug List

Triglyceride Lowering Agents

Trial with a fibric acid derivative OR niacin prior to progressing to the following products:

Lovaza®

Vascepa®:

See also; Frequency/Quantity/Duration, Preferred Drug List

Xifaxan (rifaximin)

Trial of a preferred fluoroquinolone before <u>rifaximin</u> for diagnosis of traveler's diarrhea

Date Effective 3/21/13