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An Official Accompaniment to the Alberta Drug Benefit List (ADBL) produced by Alberta Blue Cross The Expert Committee on Drug Evaluation and Therapeutics (ECDET)

EXPERT COMMITTEE MEMBERS:

Scott Klarenbach, MD, MSc (Health Econ), FRCPC (Chair) Saibal Nandy, MBBS, MRCPsych, FRCPC (Vice-Chair) Caitlin A. Clarke, BScPharm, PharmD Glen J. Pearson, BScPhm, PharmD, FCSHP Cheryl A. Sadowski, BSc (Pharm), PharmD, FCSHP Jeremy Slobodan, BSP

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ADMINISTRATIVE AND SCIENTIFIC SUPPORT:

Amanda Chung, BSc (Pharm) Sherry Dieleman, BSc (Pharm), MSc Connie Lussier, BSP, MA Carlyn Volume-Smith, BSc (Pharm), MSc, PhD Darcia Wasarab-Rolland, BSc (Pharm)

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Brief Summary of Drug Review Activities

The Expert Committee on Drug Evaluation and Therapeutics met on March 17, 2016. The Committee reviewed Manufacturer submissions for thirty-eight (38) Drug Products for potential listing, or change in listing, on the *ADBL*. The Committee also considered information for a number of supplementary assessments of the coverage status of fourteen (14) Drug Products.

In addition to Drug Products reviewed by the Expert Committee, thirty (30) Drug Products underwent Expedited Review for listing on the *ADBL* effective May 1, 2016.

The following are <u>highlights</u> of recent changes to the *ADBL* and other topics of general interest. A complete list of changes, as well as the full *ADBL* may be accessed at

https://www.ab.bluecross.ca/dbl/publications.html

Highlights of Products Originally Reviewed via the Common Drug Review (CDR)

The following Drug Product was review by CDR and the Expert Committee and added to the *ADBL* via Special Authorization (SA) for the indications of Ankylosing Spondylitis, Plaque Psoriasis, Psoriatic Arthritis and Rheumatoid Arthritis effective April 1, 2016:

• INFLECTRA* (infliximab) (CHH) 100 mg/vial injection

The following Drug Product was reviewed by CDR and the Expert Committee and added to the *ADBL* effective May 1, 2016:

• SPIRIVA RESPIMAT (tiotropium bromide monohydrate) (BOE) 2.5 mcg actuation inhalation solution

The following Drug Product was reviewed by CDR and the Expert Committee and added to the *ADBL* via SA effective May 1, 2016:

 INSPIOLTO RESPIMAT* (tiotropium bromide monohydrate/olodaterol hydrochloride) (BOE) 2.5 mcg/2.5 mcg actuation inhalation solution

A complete list of changes, as well as the full ADBL may be accessed at https://www.ab.bluecross.ca/dbl/publications.html. *Please refer to the current ADBL for explanations of coverage, including a listing of coverage criteria (where applicable).*

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Highlights of Non-Interchangeable Old Drug Products Added

The following Non-Interchangeable Old Drug Product has been added to the *Palliative Coverage Drug Benefit Supplement (PCDBS)* effective May 1, 2016:

EMOLAX (polyethylene glycol 3350) (JPC)
 1 g/g oral powder

Highlights of Line Extension Drug Products Reviewed for the ADBL

The following Drug Product was added to the *ADBL* as a Regular Benefit effective May 1, 2016:

 MINT-CITALOPRAM (citalopram hydrobromide) (MPI) 10 mg tablet

The following Drug Product was reviewed but **NOT** added as it did not offer a cost or therapeutic advantage:

 PLAN B (levonorgestrel) (PAL) 1.5 mg tablet

Highlights of Other Products Reviewed but Not Added

 ANTI-STATIC COMPACT SPACE CHAMBER PLUS (aerosol holding chamber/mask) (MPI) WITH MOUTHPIECE, and SMALL/MEDIUM/ LARGE MASKS were reviewed but not added as they fail to offer a cost or therapeutic advantage.

Highlights of Interchangeable Drug Products Reviewed

The following Drug Product was reviewed by the Expert Committee as a Resubmission and was added to the *ADBL* effective May 1, 2016:

• AZATHIOPRINE (azathioprine) (SNS) 50 mg tablet

The following Drug Product was reviewed by the Expert Committee and added via Optional Special Authorization effective May 1, 2016:

• APO-MOXIFLOXACIN* (moxifloxacin HCL) (APX) 400 mg tablet

Highlights of Other Products Added

The following Product was reviewed by the Expert Committee and added to the *ADBL* via SA effective May 1, 2016:

 NEOCATE WITH DHA & ARA* (NUN) oral powder infant formula

Please note that the coverage criteria for this product on the *Human Services Drug Benefit Supplement (HSDBS)* will remain unchanged.

Highlights of New Interchangeable (IC) Groupings

Addition of the following Drug Products to the *ADBL* has resulted in the creating of a New IC Grouping, effective May 1, 2016:

- TEVA-TOBRAMYCIN (tobramycin sulfate) (TEV) 60 mg/mL inhalation solution
- TOBRAMYCIN (tobramycin sulfate) (SDZ) 60 mg/mL inhalation solution

Special Authorization Criteria Changes

The Special Authorization criteria for coverage for the following Drug Products have been revised effective May 1, 2016:

- JAMP-VANCOMYCIN* (vancomycin HCL) (JPC) 125 mg & 250 mg capsules
- RITUXAN (rituximab) (HLR) 10 mg/mL injection
- VANCOCIN^{*} (vancomycin HCL) (MLI) 125 mg & 250 mg capsules
- VANCOMYCIN HYDROCHLORIDE* (vancomycin HCL) (FKC)
 125 mg & 250 mg capsules