

the **DBL** report

Issue #69, September 2012

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

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Brief Summary of Drug Review Activities

The Expert Committee on Drug Evaluation and Therapeutics met on July 19, 2012. The Committee reviewed Manufacturer submissions for 34 Drug Products for potential listing, or change in listing status, on the AHWDBL.

In addition to these Drug Products, 32 generic Drug Products underwent Expedited Review for listing on the AHWDBL effective September 1, 2012.

The following are highlights of recent changes to the AHWDBL. A complete list of changes and the full AHWDBL can be found at

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

Highlights of Products Added to the AHWDBL

- **VFEND** (voriconazole) (PFI) **40 mg/mL oral suspension** was submitted as a Line Extension to the existing **50 mg & 200 mg tablets**. The Expert Committee recommended **VFEND** oral suspension be listed on the AHWDBL via Special Authorization as it offers a therapeutic advantage for patients who require an alternative formulation to tablets. Please refer to the current AHWDBL for a full listing of coverage criteria.
- **ELIQUIS** (apixaban) (BMS) **2.5 mg tablet** was initially reviewed via the CDR process. **ELIQUIS** is indicated for the prevention of venous thromboembolic events (VTE) in adult patients who have undergone elective knee or hip replacement surgery. In keeping with the recommendations from the CDR, this Drug Product has been added to the AHWDBL with a listing via Restricted Benefit. Please refer to the current AHWDBL for a full listing of coverage criteria.

A complete list of changes, as well as the full AHWDBL may be accessed at <https://www.ab.bluecross.ca/dbl/publications.html>.

Please refer to the current AHWDBL for explanations of coverage, including a listing of coverage criteria (where applicable).

Highlights of IC Products Added

The following are established IC Groupings Drug Product, added to the AHWDBL, effective September 1, 2012:

- **JAMP-RIZATRIPTAN** (rizatriptan benzoate) (JPC)
5 mg & 10 mg tablets
- **MYLAN-ZOLMITRIPTAN ODT** (zolmitriptan) (MYP)
2.5 mg orally disintegrating tablet

Highlights of Products Not Added

- **MYLAN-BISOPROLOL** (bisoprolol fumarate) (MYP)
5 mg & 10 mg tablets were not recommended to be added to the AHWDBL as the Manufacturer failed to provide sufficient evidence of interchangeability with an Innovator Drug Product.

Highlights of Changes to the Multiple Sclerosis (MS) Drug Coverage

On September 1, 2012, the MS Drug Product **GILENYA** (fingolimod hydrochloride) **0.5 mg capsule** was added as a Special Authorization benefit to the AHWDBL. In addition, changes were made to Section 2 of the AHWDBL: Multiple Sclerosis (MS) Drug Coverage, and the coverage criteria for MS drugs. Please refer to the September AHWDBL update (https://www.ab.bluecross.ca/dbl/pdfs/ahw_september.pdf) for details on the changes.

Products Originally Reviewed via the Common Drug Review (CDR)

In keeping with the recommendations from the CDR, the following Drug Products have not been added to the AHWDBL:

- **BENLYSTA** (belimumab) (GKC) **120 mg/vial & 400 mg/vial injections**
- **EFFIENT** (prasugrel hydrochloride) (BMS) **10 mg tablet**
- **GELNIQUE** (oxybutynin chloride) (WAT) **100 mg/g topical gel**
- **OZURDEX** (dexamethasone) (ALL) **0.7 mg intravitreal implant**
- **SAPHRIS** (asenapine maleate) (BAI) **5 mg & 10 mg sublingual tablets (for the treatment of schizophrenia)**

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Please refer to the current AHWDBL for explanations of coverage, including a listing of coverage criteria (where applicable).