

AvMed

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

Directions: The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; **fax to 1-305-671-0200.** No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. **If the information provided is not complete, correct, or legible, the authorization process can be delayed.**

Drug Requested: Korlym[®] (mifepristone 300mg)

MEMBER & PRESCRIBER INFORMATION: Authorization may be delayed if incomplete.

Member Name: _____

Member AvMed #: _____ Date of Birth: _____

Prescriber Name: _____

Prescriber Signature: _____ Date: _____

Office Contact Name: _____

Phone Number: _____ Fax Number: _____

DEA OR NPI #: _____

DRUG INFORMATION: Authorization may be delayed if incomplete.

Drug Form/Strength: _____

Dosing Schedule: _____ Length of Therapy: _____

Diagnosis: _____ ICD Code, if applicable: _____

Weight: _____ Date: _____

Diagnosis: _____ ICD Code, if applicable: _____

Quantity Limits: 120 tablets per 30 days

Recommended Dosage: Initiate therapy with 300mg/day and titrate dose every 2-4 weeks based on tolerability and symptom control, and daily dose will NOT exceed 20mg/kg/day, OR 1,200 mg once daily

CLINICAL CRITERIA: Check below all that apply. **All criteria must be met for approval.** To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

Initial Authorization: 6 months

- Member is 18 years of age or older
- Prescribing physician is an endocrinologist

(Continued on next page)

- Member has a diagnosis of Endogenous Cushing's Syndrome, and satisfies **ONE** of the following:
 - Diagnosis of Type 2 Diabetes Mellitus
 - Glucose intolerance noted by **ONE** of the following (**must submit documentation**): oral glucose tolerance test or Hemoglobin A1c test (HbA1c)
- Past medical history confirms **ONE** of the following:
 - Member has undergone surgery and has not been curative
 - Member is not a candidate for surgery
- Documentation of clinical failure (unable to normalize cortisol levels for the treatment of Cushing's Syndrome) to ketoconazole tablets taken along with **ONE** of the following:
 - An additional steroidogenesis inhibitor such as Metopirone (metyrapone capsules) or mitotane tablets
 - A pituitary-directed therapy such as cabergoline or Signifor LAR (pasireotide)
- Documentation of clinical failure to control glucose levels with Metformin **AND TWO** (2) of the following treatments:
 - Glucagon-Like Peptide-1 (GLP-1) Receptor Agonist (e.g., Trulicity, Ozempic)
 - Insulin (e.g., Humalog, Lantus)
 - Dipeptidyl Peptidase 4 (DPP-4) Inhibitor (e.g., Januvia, Onglyza)
 - Member is not also taking/ will not take strong inhibitors of CYP2A medications (e.g., simvastatin, lovastatin, cyclosporine, dihydroergotamine, ergotamine, fentanyl, pimozide, quinidine, sirolimus, tacrolimus)
- For females of reproductive potential: pregnancy has been excluded before initiation of treatment and plans for prevention are implemented during treatment and for one month after stopping

Reauthorization Approval: 12 months. Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

- Positive disease response has been achieved, demonstrated by improved glycemic control (decreased hemoglobin A1c) (**current labs must be submitted to document HbA1c**)

Medication being provided by a Specialty Pharmacy - PropriumRx

Not all drugs may be covered under every Plan.

If a drug is non-formulary on a Plan, documentation of medical necessity will be required.

*****Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.*****

****Previous therapies will be verified through pharmacy paid claims or submitted chart notes.****