SENTARA HEALTH PLANS

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions:</u> 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; <u>fax to 1-800-750-9692</u>. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. <u>If information provided is not complete, correct, or legible, authorization may be delayed.</u>

<u>Drug Requested</u>: **Korlym**[®] (mifepristone 300mg)

MEMBER & PRESCRIBER INFORMAT	ION: Authorization may be delayed if incomplete.
Member Name:	
Member Sentara #:	Date of Birth:
Prescriber Name:	
Prescriber Signature:	Date:
Office Contact Name:	
Phone Number:	
DEA OR NPI #:	
DRUG INFORMATION: Authorization may	be delayed if incomplete.
Drug Form/Strength:	
Dosing Schedule:	
Diagnosis:	ICD Code, if applicable:
Weight:	Date:
Quantity Limits : 120 tablets per 30 days	
Recommended Dosage : Initiate therapy with 300 tolerability and symptom control, and daily dose will I	· ·
CLINICAL CRITERIA: Check below all that support each line checked, all documentation, including provided or request may be denied.	apply. All criteria must be met for approval. To ing lab results, diagnostics, and/or chart notes, must be
Initial Authorization : 6 months	
☐ Member is 18 years of age or older	
☐ Prescribing physician is an endocrinologist	
	ng's Syndrome, and satisfies ONE of the following:
☐ Diagnosis of Type 2 Diabetes Mellitus	
Glucose intolerance noted by ONE of the fitter tolerance test or Hemoglobin A1c test (Hb.	following (must submit documentation): oral glucose A1c)

(Continued on next page)

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 <u>TWO</u> (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
uthorization Approval: 12 months. Check below all that apply. All criteria must be met for

R 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.*