

GILENYA EXCEPTION DRUG STATUS (EDS) APPLICATION

Check if:

- ☐ New Application (Complete Sections 1 thru 4 in full)
☐ Annual Renewal (Complete Sections 1, 3 and 5 in full)

Section 1 (Please print) :

PATIENT INFORMATION:	NEUROLOGIST INFORMATION:
Name: _____	Name: _____
Date of Birth: _____	Address: _____
HSN: _____	_____
Address: _____	Postal Code: _____
_____	Phone: _____ Fax: _____
Postal Code: _____	Date: _____
Phone: _____	Date of Most Recent Consultation: _____
Family Physician: _____	Neurologist's Signature: (required)
Patient Signature: (required) _____	_____ Date: _____

Section 2:

EDS approval will be given to patients with Relapsing Remitting Multiple Sclerosis (RRMS) who are assessed and meet **ALL** of the following criteria:

- i. Has failed to respond to a full and adequate course of at least one disease modifying therapy listed on the SK Formulary as initial therapy (trial of at least 6 months **AND** experienced at least one relapse attack while on treatment) ☐ YES ☐ NO

OR

- Has documented intolerance (serious side effect or contraindication) ☐ YES ☐ NO
to at least **TWO** disease modifying therapies listed on the SK Formulary as initial therapy

Name of Drug	Duration of Treatment

- ii. Has experienced one or more clinically disabling relapses in the previous year ☐ YES ☐ NO
- iii. Has shown evidence of a significant increase in T2 lesion load (i.e. 3 or more new lesions) compared to a previous MRI OR at least one gadolinium-enhancing lesion on MRI (include summary of MRI findings) ☐ YES ☐ NO
- iv. Has a current Extended Disability Status Scale (EDSS) score of 5.5 or less ☐ YES ☐ NO
Date of most recent EDSS score: (D / M / Y) _____ EDSS Score _____

Section 3:

Contraindications to treatment:

- | | | |
|-----------------------------------------------------------------------------------------------------------------------------------------------------------|------------------------------|-----------------------------|
| i. Patient has had a heart attack or stroke in the last 6 months | <input type="checkbox"/> YES | <input type="checkbox"/> NO |
| ii. History of sick sinus syndrome, atrioventricular block, significant QT prolongation, bradycardia, ischemic heart disease, or congestive heart failure | <input type="checkbox"/> YES | <input type="checkbox"/> NO |
| iii. Patient is currently taking class Ia or class III anti-arrhythmic medications | <input type="checkbox"/> YES | <input type="checkbox"/> NO |
| iv. Immunocompromised due to immunosuppressant or anti-neoplastic therapy or due to immunodeficiency syndrome (HIV, leukemia, lymphoma etc.) | <input type="checkbox"/> YES | <input type="checkbox"/> NO |
| v. Severe hepatic impairment | <input type="checkbox"/> YES | <input type="checkbox"/> NO |
| vi. Concurrent malignancies | <input type="checkbox"/> YES | <input type="checkbox"/> NO |
| vii. Pregnancy, anticipated pregnancy or breast-feeding | <input type="checkbox"/> YES | <input type="checkbox"/> NO |
| viii. Active infectious disease (such as tuberculosis, hepatitis) | <input type="checkbox"/> YES | <input type="checkbox"/> NO |
| ix. Patient is less than 18 years of age | <input type="checkbox"/> YES | <input type="checkbox"/> NO |

(Skin reactions at the site of injection do NOT qualify as a contraindication to injectable disease modifying therapy. Needle phobia or preference for oral therapy is not an appropriate indication for Gilenya)

Section 4:

- | | | |
|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------------------------|-----------------------------|
| Has a risk factor assessment been done?
(due to safety considerations with Gilenya, applications cannot be considered until the required investigations are completed. Please refer to the manufacturer's list of investigations as outlined on the Gilenya Treatment Checklist) | <input type="checkbox"/> YES | <input type="checkbox"/> NO |
| Have the benefits/risks of this medication been discussed with your patient? | <input type="checkbox"/> YES | <input type="checkbox"/> NO |
| Has your patient agreed to proceed with treatment with this medication? | <input type="checkbox"/> YES | <input type="checkbox"/> NO |
| Have arrangements been made for the patient to be monitored following drug administration according to the Health Canada guidelines outlined in the product monograph? | <input type="checkbox"/> YES | <input type="checkbox"/> NO |

Section 5:

Renewal of Coverage: Renewal will be given to patients who meet **ALL** of the following criteria:

- | | | |
|------------------------------------------------------------------------------------------------------------|------------------------------|-----------------------------|
| i. Patient has been stable or has experienced no more than one disabling attack/relapse in the past year | <input type="checkbox"/> YES | <input type="checkbox"/> NO |
| ii. Has an EDSS score of 5.5 or less
Date of most recent EDSS score: (D / M / Y) _____ EDSS Score _____ | <input type="checkbox"/> YES | <input type="checkbox"/> NO |

Please forward clinical history including:

- documentation of attacks, date of onset, date of diagnosis
- neurological findings (exam must have occurred within 90 days of the request), EDSS score
- MRI reports/summary of findings or other significant information
- complete medication profile

TO: Saskatchewan MS Drugs Program
Suite 7718 – 7th Floor
Saskatoon City Hospital
SASKATOON SK S7K 0M7

OR Fax: (306) 655-8404