

PRIOR AUTHORIZATION PROGRAM FORM

OZEMPIC AND GENERICS (semaglutide)

Instructions:

Section 1 to be completed by Plan Member / Patient
 Section 2 and 3 to be completed by Physician
 Section 4 to be completed by Physician and Pharmacist

Please submit the completed form to:

NexgenRx Formulary Management
 145 The West Mall P.O. Box 110 U,
 Toronto, Ontario, M8Z 5M4
by Fax: 1-877-639-4369
by email: providerservices@nexgenrx.com

The costs incurred to complete this form are the responsibility of the patient. The costs are not eligible for reimbursement under the plan

ALL FIELDS ARE MANDATORY: INCOMPLETE FORMS MAY RESULT IN YOUR APPLICATION BEING DECLINED

SECTION 1A: POLICY INFORMATION

Carrier ID: Policy Group Number (8 digits): Certificate Number (12 digits): Patient Code (3 digits):
 98

SECTION 1B: PATIENT INFORMATION

First Name: Last Name: Date of Birth (DD MMM YYYY): Gender:
☐ Male ☐ Female
☐ Prefer not to disclose

Street Address: City: Province: Postal Code:

Preferred Phone: Email: Relationship to Member:
☐ Employee ☐ Spouse ☐ Dependent

SECTION 1C: PLAN MEMBER INFORMATION

First Name: Last Name: Employer:

Preferred Phone: Email:

SECTION 1D: COORDINATION OF BENEFITS - PROVINCIAL COVERAGE

Has the patient applied for reimbursement under a provincial plan? ☐ Yes ☐ No
 What is the coverage decision? (Attach decision outcome letter) ☐ Approved ☐ Denied

SECTION 1E: COORDINATION OF BENEFITS - PRIVATE COVERAGE

Is the patient covered by any other drug plan? ☐ Yes ☐ No
 What is the coverage decision? (Attach decision outcome letter) ☐ Approved ☐ Denied

Plan Member First Name: Plan Member Last Name: Date of Birth (DD MMM YYYY): Relationship to Member:
☐ Employee ☐ Spouse ☐ Dependent

Coverage provided by: Carrier ID: Policy Group Number: Certificate Number:

SECTION 1F: COORDINATION OF BENEFITS - PATIENT SUPPORT PROGRAM

Is the patient enrolled in a Patient Support Program? ☐ Yes ☐ No
 Patient consents to NexgenRx contacting Patient Support Program? ☐ Yes ☐ No

Program Name: Program Patient Identifier: Program Fax:

Contact First Name: Contact Last Name: Phone: Email:

SECTION 1G: COORDINATION OF BENEFITS – DISABILITY

Is the patient currently on disability benefits for the condition being treated? ☐ Yes ☐ No

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SECTION 1H: COORDINATION OF BENEFITS – DRUG ACCESS NAVIGATOR

Is the patient currently being supported by a Drug Access Navigator?

☐ Yes

☐ No

Patient consents to NexgenRx contacting Drug Access Navigator

☐ Yes

☐ No

Organization Name:

Patient Identifier:

Fax:

Contact First Name:

Contact Last Name:

Phone:

Email:

Results of this request to be communicated to

☐ Patient

☐ Plan Member

☐ Patient Support Program *

☐ Drug Access Navigator *

☐ Pharmacy*

For each preferred recipient, email and patient consent **MUST** be provided in their respectful sections. Patient, Plan Member, Patient Support Program, and Drug Access Navigator details are found in Section 1. Pharmacy details are found in Section 4.

☐ Legal Guardian

Consent Provided:

E-MAIL (mandatory):

☐ Yes ☐ No

SECTION 1I: CONSENT

I authorize NexgenRx Inc. (a) to use the personal information disclosed on this form, and any other personal information known to NexgenRx Inc. regarding the above-named patient, for the purpose of participating in this clinical patient support program and (b) to contact, and to obtain any such personal information from and to disclose any such personal information to, any physician, pharmacist, other health care professional or agent having knowledge of such patient's health relevant to this request and any related claim. You give the authorization to have this form provided to a Licensed Authorized Supplier your Employer has agreed to upon request.

I consent to be contacted by any outside companies other than NexgenRx via phone, text or email and to the transfer of personal information by phone, fax or email between the Licensed Authorized Supplier, my insurer, and my health care provider(s) for the purpose of determining my eligibility for the program and the delivery of program services. Email and text may be used during the course of my participation in the program to inform me about my status in the program, program services, and to provide notifications and reminders. I acknowledge that neither email nor text is a secure method of communication. Information in emails and texts has the potential to be accessed and read by a third party. Electronic communication is at my option and I may withdraw this option to communicate electronically at any time.

* If the patient has another drug plan, this application may cover some or all of the portion not paid by the primary plan.

By signing below, you certify that all the information referred to in this form are genuine and that the information provide is true and complete and if any such claim concerns your spouse or any dependent that you have their consent to disclose their personal information to us for purpose referred to above. You also authorize us to obtain and exchange information with respect to this claim with any person having such relevant information including any health care provider, insurer, claims adjudicator or administrator or any privately or publicly funded benefit plan or program.

Patient / Legal Guardian Name:

Signature of Patient / Legal Guardian:

Date (DD MMM YYYY):

SECTION 2: MEDICATION REQUESTED (to be completed by a Physician)

DRUG NAME, STRENGTH AND DOSE	
Drug Identification Number (DIN)	
Drug Type Requested	☐ Brand name or generic equivalent *Brand name eligible cost will be based on the generic alternative, and any price difference will be paid by the patient. ☐ Brand name – No generic substitution ** Please stop and do not proceed further. Complete the Brand Name Request Form, available for download on our website: https://priorauthforms.okrx.ca/Nexgen/EN . To be eligible for approval, the patient must have tried two generic equivalents with the same molecule, along with documented medical necessity for the brand-name medication. If approved, the plan will cover the cost of the brand-name medication. ☐ Generic equivalent
Indication	☐ Type 2 Diabetes Mellitus ☐ OFF-LABEL USE: (Please note that only the “Indications” published in the Product Monograph submitted to Health Canada will be considered) ☐ Type 1 Diabetes Mellitus ☐ Obesity and Chronic Weight Management: Weight: _____ Height: _____ BMI: _____ ☐ Other (please specify): _____
Anticipated Duration of Treatment	
Quantity prescribed or attach a copy of a prescription	Quantity Prescribed / Dose: _____ ☐ Weekly ☐ Monthly ☐ Other frequency (please specify): _____
Start Date:	
Treatment Administration Site (Specify where the medication will be Administered)	☐ HOME (administered by patient) ☐ Clinic (administered by health care professional) ☐ Hospital ☐ Other (please specify) _____
Has the patient previously used this medication?	☐ Yes, Renewal. Start of treatment date: _____ ☐ No. New drug Therapy
FOR RENEWAL: Clinical response:	☐ Improved ☐ Stable ☐ No response
FOR RENEWAL: Any adverse effects?	☐ Yes ☐ No
Concurrent medications with the requested drug, if any:	
If alternative therapies have previously been considered or initiated, please list the treatments that have been trialed, including the clinical response or outcomes observed, as well as the reasons for discontinuation (e.g., lack of efficacy, adverse effects, or patient preference).	

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SECTION 3: CLINICAL INFORMATION

Eligibility for drug coverage is dependent upon the patient meeting at least one of the qualifying criteria listed below. If the patient has another drug plan, prior authorization may cover some or all of the excess not paid by the primary plan.

Please indicate if the patient meets at least one of the following qualifying criteria for drug coverage:

- ☐ Indicated for the once-weekly treatment of adult patients with **TYPE 2 DIABETES** to improve glycemic control, in combination with:
 - ☐ Diet and exercise in patients for whom metformin is inappropriate due to contraindication or intolerance.
 - ☐ Metformin, when diet and exercise plus maximal tolerated dose of metformin do not achieve adequate glycemic control.
 - ☐ Metformin and a sulfonylurea, when diet and exercise plus dual therapy with metformin and a sulfonylurea do not achieve adequate glycemic control.
 - ☐ Metformin or a sulfonylurea and a sodium-glucose cotransporter 2 inhibitor (SGLT2i), when diet and exercise plus metformin or a sulfonylurea, in addition to an SGLT2i, do not achieve adequate glycemic control.
 - ☐ Basal insulin with metformin, when diet and exercise plus basal insulin with or without metformin do not achieve adequate glycemic control.
- ☐ Indicated as an adjunct to diet, exercise, and standard of care to reduce the risk of major adverse cardiovascular events (cardiovascular death, non-fatal myocardial infarction, or non-fatal stroke) in adults with type 2 diabetes and established cardiovascular disease and/or chronic kidney disease.
- ☐ Indicated to reduce the risk of sustained eGFR decline, end-stage kidney disease, and cardiovascular death in adults with type 2 diabetes and chronic kidney disease.

FOR TYPE 2 DIABETES MELLITUS

Target HbA1C	
Baseline HbA1C	
Most recent HbA1C	
Maximally tolerated Metformin dose used:	
Date Metformin Was Initially Initiated:	
If Metformin was discontinued or not tried, please justify	

Any Additional clinical information to support the request:

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SECTION 4A: MEDICAL PRACTITIONER'S INFORMATION

Physician First Name:	Physician Last Name:	License Number:	Specialty:
Clinic / Organization Name:			
Street Address:	City:	Province:	Postal Code:
Phone:	Fax:	Email:	
Signature of Physician:		Date (DD MMM YYYY):	

SECTION 4B: PHARMACIST'S INFORMATION

Pharmacist First Name:	Pharmacist Last Name:	License Number:
Pharmacy Name:	Provider Number:	
Street Address:	City:	Province: Postal Code:
Phone:	Fax:	Email:
Patient consents to NexgenRx contacting the Pharmacy? ☐ Yes ☐ No		
Signature of Pharmacist:	Date (DD MMM YYYY):	

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INTERNAL OFFICE USE

Date Received (DD/MM/YYYY):	Start Date (DD/MM/YYYY):	End Date (DD/MM/YYYY):	Date Approved (DD/MM/YYYY):