

DPP-4/SGLT2 INHIBITORS / **GLP-1 RECEPTOR AGONISTS** SPECIAL AUTHORIZATION REQUEST FORM

Please complete all required sections to allow your request to be processed.

Patients may or may not meet eligibility requirements as established by Alberta government-sponsored drug programs.

PATIENT INFO	PRMATION						COVER	AGE TYPE			
PATIENT LAST NAME		FIRST NAME	FIRST NAME I		INITIAL	Alberta Blue Cross					
BIRTH DATE (YYYY/MM/DD)		ALBERTA PERSONA	ALBERTA PERSONAL HEALTH N		IMBER	?	1 =	ta Human Se -	rvices		
·		ALBERTATI ERROTT					☐ Other				
STREET ADDRESS		CITY	PF	ROV POSTAL CODE		ID/CLIENT/COVERAGE NUMBER		AGE NUMBER			
PRESCRIBER	INFORMATION										
PRESCRIBER LAST NAME FIRST NAME INITIAL						PRESCRIBER PROFESSIONAL ASSOCIATION REGISTRATION					
					☐ CPSA ☐ ACO REGISTRATION NUMBER						
STREET ADDRESS				☐ CARNA ☐ ADA+C ☐ ACP ☐ Other							
CITY, PROVINCE				PHONE FAX							
POSTAL CODE				FAX NUMBER MUST BE PROVIDED WITH EAC					CH REQUEST SUBMITTED		
Indicate which drug is requested											
For the treatment of Type 2 diabetes					a for c	coverage*	Complete sections				
CANAgliflozin (e.g. Invokana) SAXAgliptin + metformin				First-line drug products: metformin				·			
LINAgliptin (e.g. Trajenta) (e.g. Komboglyze)				Second-line drug products: sulfonylureas							
☐ LINAgliptin + metformin ☐ SEMAglutide (e.g. Ozempic)				And where insulin is not			an option				
(e.g. Jent						Sections 1 and 2					
SAXAglipt											
(e.g. Janumet, Janumet XR) LIXIsenatide (e.g. Adlyxine)					e dru	g products					
ElAlbertatide (c.g. Adiyante)				Second-line drug products: sulfonylureas					Sections 1 and 2		
					sulin						
For the treatment of Type 2 diabetes OR Type 2 diabetes and established CV diseases as defined in the criteria for coverage				Criteria for coverage*					Complete sections		
☐ EMPAglifl		EMPAgliflozin + metform	nin	*See page 2 for complet			te criteria		Sections 1 and/or 2		
		(e.g. Synjardy)							(as applicable)		
Section 1 Please indicate if metformin was tried for at least 6 months											
	☐ Yes ☐ No, specify reason										
Section 2 Please indicate if a sulfonylurea was tried											
	☐ Yes ☐ No, specify reason										
Please indicate if insulin was tried											
☐ Yes ☐ No, indicate why insulin is not an option for this patient ☐ Cognitive impairment ☐ Manual dexterity concerns											
□ Needle phobia □ Visual impairment											
Other, specify											
Additional in	formation relating to rec	juest									
PRESCRIBER'S SIGNATURE DA		DATE (YYYY-MM-DD)			d this r	equest to					
		==/	Alberta Blue Cross, Clinical Drug Services								
			10009-108 Street NW, Edmonton, Alberta T5J 3C5 FAX: 780-498-8384 in Edmonton • 1-877-828-4106 toll free all other areas								
ONCE YOUR REQUEST HAS SUCCESSFULLY TRANSMITTED, PLEASE DO NOT MAIL OR RE-FAX YOUR REQUEST.											

The information on this form is being collected and pursuant to sections 20, 21 and 22 of the Health Information Act, and sections 33 and 34 of the Freedom of Information and Protection of Privacy Act, for the purposes of determining or verifying eligibility to participate in a program or receive a benefit, product or health service. If you have any questions regarding the collection or use of this information, please contact an Alberta Blue Cross privacy matters representative toll-free at 1-855-498-7302 or write to Privacy Matters, Alberta Blue Cross, 10009 108 Street, Edmonton AB T5J 3C5.





Criteria for coverage

DPP-4/SGLT2 INHIBITORS / GLP-1 RECEPTOR AGONISTS SPECIAL AUTHORIZATION CRITERIA

Patients may or may not meet eligibility requirements as established by Alberta government-sponsored drug programs.

CANAgliflozin (e.g. Invokana), LINAgliptin (e.g. Trajenta), LINAgliptin + metformin (e.g. Jentadueto), LIXIsenatide (e.g. Adlyxine), SAXAgliptin (e.g. Onglyza), SAXAgliptin + metformin (e.g. Komboglyze), SEMAglutide (e.g. Ozempic), SITAgliptin (e.g. Januvia) and SITAgliptin + metformin (e.g. Janumet, Janumet XR) special authorization criteria

FIRST-LINE DRUG PRODUCTS: METFORMIN SECOND-LINE DRUG PRODUCTS: SULFONYLUREAS AND WHERE INSULIN IS NOT AN OPTION

As add-on therapy for the treatment of Type 2 diabetes in patients with intolerance to and/or inadequate glycemic control on

- a sufficient trial (such as a minimum of 6 months) of metformin. AND
- a sulfonylurea, AND
- for whom insulin is not an option.

Or, for whom these products are contraindicated.

Special authorization may be granted for 24 months.

EMPAgliflozin (e.g. Jardiance) and EMPAgliflozin + metformin (e.g. Synjardy) special authorization criteria

FIRST-LINE DRUG PRODUCTS: METFORMIN

As add-on therapy for the treatment of Type 2 diabetes in patients with intolerance to and/or inadequate glycemic control on

- a sufficient trial (such as a minimum of 6 months) of metformin, AND
- a sulfonylurea, AND
- for whom insulin is not an option.

Or, for whom these products are contraindicated.

As an adjunct to diet, exercise, and standard care therapy to reduce the incidence of cardiovascular (CV) death in patients with Type 2 diabetes and established cardiovascular diseases who have an inadequate glycemic control, if the following criteria are met:

- a sufficient trial (such as a minimum of 6 months) of metformin. AND
- established cardiovascular disease* as defined in the EMPA-REG OUTCOME trial.
- * Established cardiovascular disease is defined on the basis of 1 of the following:
- 1) History of myocardial infarction.
- 2) Multi-vessel coronary artery disease in 2 or more major coronary arteries (irrespective of revascularization status).
- 3) Single-vessel coronary artery disease with significant stenosis and either a positive non-invasive stress or discharged from hospital with a documented diagnosis of unstable angina within the last 12 months.
- 4) Last episode of unstable angina greater than 2 months prior with confirmed evidence of coronary multi-vessel or single-vessel disease.
- 5) History of ischemic or hemorrhagic stroke.
- 6) Occlusive peripheral artery disease.

Special authorization may be granted for 24 months.

LIXIsenatide (e.g. Adlyxine) special authorization criteria

FIRST-LINE DRUG PRODUCTS: METFORMIN SECOND-LINE DRUG PRODUCTS: SULFONYLUREAS AND INSULIN

"As add-on therapy for the treatment of Type 2 diabetes in patients with intolerance to and/or inadequate glycemic control on

- a sufficient trial (such as a minimum of 6 months) of metformin, AND
- a sulfonylurea, AND
- insulin.

Or, for whom these products are contraindicated.

Special authorization may be granted for 24 months.

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