Patient Enrolment Form | Plaque Psoriasis



FAX FORM TO: 1-844-397-5635 OTEZLA ez Start Bridging Program app enrolment ID: _ _ _ _ _ _ _

PATIENT INFORMATION	NOTE: Asterisks (*)	denote mand	datory fields, daggers (†) de	note fields mandatory for br	idging program assessment.
First name:*	Last name:*			Date of birth (mm/dd/yy)	:/
Address:				Phone:*	\square OK to leave message
City:		Province: _	Postal code:	Email:	
Preferred method of contact:	Phone □ Email				
☐ Covered by private health in	surance [†]				
Insurance company:	□ Covered by spo	usal insurance			
Consent: By providing my email address, information and updates relating to my entime by providing notice to Innomar at: In	nrolment in the ez Start Pati	ient Support Pr	ogram (the "Program"). I underst	and that I may withdraw my conse	nt to such communications at any
☐ I consent to being contacted for confidential surveys about the contacting the ez Start Patient	e Program. I understan		•		
I HAVE READ, UNDERSTAND	AND AGREE TO THE	CONSENT	TERMS ON THE REVER	SE OF THIS FORM.	
Patient/patient representative signature: X Date (mm/dd/y)					y):/
If signed by patient representative, please include name and explain authority to act on behalf of the patient:					
OPTIONAL VERBAL CONSENT: To be c	ompleted by healthcare pr	ovider (if appli	cable). If healthcare provider is	unable to obtain written consen	t from patient, please document
when patient verbal consent was obtained. This will allow the Program to continue processing this enrolment while written consent is obtained by the Program.					
□ Verbal consent obtained by healthcare provider Name: Date (mm/dd/yy):/					
PRESCRIBER INFORMATIO	N*				
Name:			Phone:	Fax:	
Address:					
A SCESCIAENT FOR PRINCING		-			
ASSESSMENT FOR BRIDGING F Diagnosis:	ROGRAM COVERAG		Treatment history:	Additiona	l information:‡
☐ Plaque Psoriasis (PsO)†			rreachient history.	Additiona	ini oi macion.
Assessment parameters:					
BSA:†% OR PASI:†	Other:				
Affected areas:†					
PRESCRIPTION INFORMATION	 *				
Starter Pack (titration) Rx for C		Patient re	ceived Starter Pack	Maintenance Rx for OTE	ZLA
☐ Take as directed (x 14 days, 27 t☐ Other:		in office ☐ Yes ☐] No	☐ Take 30 mg PO BID x28 ☐ Other:	
Special instructions:					
PRESCRIBER AUTHORIZATION	 *				
I authorize Innomar to be my designated	— d agent to forward the pres				
pharmacy chosen by the above-named precipient and there are no others. The o	riginal prescription has bee	en invalidated a	and securely filed and it will not	be transmitted elsewhere at an	
Physician signature: X	D	ate (mm/dd/	/w/· /	License #•	

^{*} Mandatory field.

[†] Mandatory field for bridging program assessment.

[‡] For example, dosage, start of therapy, end of therapy, reason for discontinuation.

BID, twice a day; BSA, Body Surface Area; ID, identification; PASI, Psoriasis Area and Severity Index; PO, take by mouth; Rx, prescription; Tx, treatment.

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Enrolment and consent form for the ez Start Patient Support Program

By signing this form, I acknowledge that I have read and understand the information below and consent to the collection, use and disclosure of my personal information, including personal health information, by Innomar, Amgen and their authorized agents and service providers as explained. Furthermore, I acknowledge that the dispensing and delivery of my medication will be performed by a specialty pharmacy chosen by the ez Start Program unless I specify otherwise. I understand that I have the option to choose another pharmacy to dispense and deliver my medication. I further consent to being contacted from time to time by Innomar, Amgen or their authorized agents for the above-noted purposes.

The ez Start Patient Support Program ("Program") is sponsored by Amgen Canada Inc. ("Amgen") and administered by Innomar, a thirdparty service provider, on behalf of Amgen. Other service providers may be appointed by Amgen to administer the Program from time to time. The personal information that you and/or your healthcare providers (including your doctor and pharmacy), insurers or payers provide to the Program, including your name, contact information, and prescription information, will be used to manage and administer the Program, including provision of Program services to you, such as reimbursement assistance and administering, therapy training, medication delivery, and provision of information about the Program to you.

Amgen has a legal obligation to report adverse drug events to various local and international health authorities and to monitor product complaints. Personal information provided to the Program may be (i) monitored by Amgen or its service providers for safety-related data and product complaints in order to ensure compliance with these legal reporting requirements, and (ii) reported to local or international health authorities. Amgen may contact you or your physician for additional information to fulfill its reporting obligations. Your personal information may be combined with the information of others who participate in the Program in order to generate aggregated data that does not contain identifying information ("Aggregated Data"). Aggregated Data may be used by Amgen and its service providers to improve and/or refine the Program to design and implement other patient programs and for research purposes including health economic studies and analysis, the identification of trends such as product utilization, adherence or outcomes.

For these sole purposes, Innomar may on a confidential basis, collect your personal information and share it with your healthcare providers, insurers and/or other payers, Amgen and/or Amgen's agents and service providers (e.g., information technology providers). If, from time to time, another service provider is appointed by Amgen to administer the Program or provide Program services, your personal information will be transferred to this service provider to ensure the continuity of the Program services to you. Please note that Amgen and its service providers may store or process your personal information outside of Canada (including in the United States), where local laws may require the disclosure of personal information to governmental authorities under circumstances that are different than those that apply in Canada. In addition, your personal information may be used or disclosed to third parties when permitted or required by applicable laws, court orders or government regulations (collectively, "Applicable Laws").

Your personal information will be retained only for as long as is needed to fulfill the purposes for which it was collected and in order to comply with Applicable Laws. Industry standard safeguards will be used to protect the security of the personal information that is collected. You may contact the Program's Privacy Officer at any time to update or access your personal information, modify or withdraw your consent (in part or in full), express a privacy-related concern, or inquire about the privacy practices of the Program (including those related to foreign information processing). The Privacy Officer can be reached at privacycanada@amgen.com or Amgen Canada Inc., Attn: Chief Privacy Officer, 6775 Financial Drive, Suite 100, Mississauga, ON, L5N 0A4. Please note that if you modify or withdraw your consent, your ability to receive the Program services may be limited.

This document may contain private and confidential information and is intended only for the person(s) named on the reverse. If you are not a named addressee, you should not disseminate, distribute or copy this document. If you have received this document by mistake, please notify the sender immediately and then destroy this document. We thank you for your cooperation and assistance.









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