

**PLEASE FAX TO YOUR JANSSEN BIOADVANCE® COORDINATOR UPON COMPLETION**

Janssen BioAdvance® Coordinator:		Tel.:	Fax:
<b>PATIENT INFORMATION</b>		<b>PHYSICIAN INFORMATION</b>	
Patient Name:		Physician Name:	
Address:		Other Office Contact:	
Language preference: <input type="checkbox"/> English <input type="checkbox"/> French <input type="checkbox"/> Other		Office Address:	
Tel. (Home):	Tel. (Other):	Email:	
Can leave a message at this phone number: <input type="checkbox"/> YES <input type="checkbox"/> NO		Tel. (Office):	
Email:		Fax (Office):	
Date of Birth:	Health Card #:		

<b>PRESCRIBING PHYSICIAN SECTION†</b>		Please <input checked="" type="checkbox"/> and complete the required information.	
Indication: <input type="checkbox"/> Plaque Psoriasis <input type="checkbox"/> Psoriatic Arthritis		Patient Weight:	
<input type="checkbox"/> <b>ADULT</b> (18 years old and above) <b>PRESCRIPTION</b>	<input type="checkbox"/> <b>STELARA® (ustekinumab) 45 mg subcutaneous injection</b>	<input type="checkbox"/> <b>STELARA® (ustekinumab) 90 mg subcutaneous injection</b>	<input type="checkbox"/> <b>STELARA® (ustekinumab) 45 mg subcutaneous injection x 2</b>
<input type="checkbox"/> <b>PEDIATRIC</b> (6-17 years old) <b>PRESCRIPTION†</b> (plaque psoriasis)	<input type="checkbox"/> <b>STELARA® (ustekinumab) 45 mg subcutaneous injection</b>	<input type="checkbox"/> <b>STELARA® (ustekinumab) 90 mg subcutaneous injection</b>	<input type="checkbox"/> <b>STELARA® (ustekinumab) 0.75 mg/kg (45 mg/0.5 ml vial)</b>
<b>DOSE</b>	<b>INITIAL weeks:</b> <input type="checkbox"/> 0 <input type="checkbox"/> 4 <input type="checkbox"/> 16 <input type="checkbox"/> 28	<b>MAINTENANCE: Q weeks:</b> <input type="checkbox"/> Weeks <input type="checkbox"/> Repeats <input type="checkbox"/> 52 weeks	Other Directives / Notes:
<b>AND / OR</b>			

<b>REIMBURSEMENT</b>	<b>PREVIOUS THERAPY</b>																					
<b>PSORIASIS ASSESSMENT DETAILS</b> (please complete if necessary)																						
BSA %: <input type="text"/> PASI: <input type="text"/> DLQI: <input type="text"/>																						
<input type="checkbox"/> Face <input type="checkbox"/> Hands <input type="checkbox"/> Feet <input type="checkbox"/> Genitals																						
<b>TUBERCULOSIS EVALUATION</b>																						
<input type="checkbox"/> Not Required																						
<input type="checkbox"/> Negative Result <input type="text"/> Date:																						
<input type="checkbox"/> Pending																						
<input type="checkbox"/> Positive Result <input type="text"/> Date:																						
<input type="checkbox"/> Ontario, Psoriasis only: Patient qualifies for coverage under LU code 419																						
	<table border="1" style="width:100%; border-collapse: collapse;"> <thead> <tr> <th></th> <th>Reason for discontinuation:</th> <th>Start/stop dates:</th> </tr> </thead> <tbody> <tr> <td><input type="checkbox"/> Methotrexate</td> <td><input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Failure</td> <td></td> </tr> <tr> <td><input type="checkbox"/> Acitretin</td> <td><input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Failure</td> <td></td> </tr> <tr> <td><input type="checkbox"/> Cyclosporine</td> <td><input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Failure</td> <td></td> </tr> <tr> <td><input type="checkbox"/> Topicals</td> <td><input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Failure</td> <td></td> </tr> <tr> <td><input type="checkbox"/> Phototherapy</td> <td><input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Failure <input type="checkbox"/> Not Available</td> <td></td> </tr> <tr> <td><input type="checkbox"/> Other</td> <td><input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Failure</td> <td></td> </tr> </tbody> </table>		Reason for discontinuation:	Start/stop dates:	<input type="checkbox"/> Methotrexate	<input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Failure		<input type="checkbox"/> Acitretin	<input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Failure		<input type="checkbox"/> Cyclosporine	<input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Failure		<input type="checkbox"/> Topicals	<input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Failure		<input type="checkbox"/> Phototherapy	<input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Failure <input type="checkbox"/> Not Available		<input type="checkbox"/> Other	<input type="checkbox"/> Intolerance <input type="checkbox"/> Contraindication <input type="checkbox"/> Failure	
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<b>PHYSICIAN</b>	† Effective date. Order(s) expire one year from the date of signature. Prescriber certification: I certify that this prescription is an original prescription and this pharmacy is the only receiver. The original will not be reused.	
Physician Signature:	College License #:	Date†:

<b>PATIENT</b>	I have read and understood the Patient Consent text printed on the back of this form and agree to the collection, use and disclosure of my personal information in accordance with these terms.
Patient Signature:	Date:

## PATIENT CONSENT

I agree to permit my healthcare provider(s), including physician(s) or nurse(s), to disclose my personal information to the BioAdvance® Coordinator assigned to managing my patient file, or his/her replacement (where applicable) (the “BAC”) in order to facilitate my enrolment in the BioAdvance® Program and facilitate the obtaining of my first STELARA® prescription, and I agree that the BAC can contact me for such purposes. Personal information may include my name, address, date of birth, phone numbers and any other personal information, including my personal health information, such as my diagnosis and the information included on my prescription or on this Patient Enrolment, Rx & Consent Form (the “Consent Form”).

My personal information will not be used or disclosed by the BAC for any purpose other than those described above unless information that identifies me directly is first removed or as is permitted or required by law. In addition, once I have been contacted by the BAC, if I elect to benefit from the services provided by the BAC, I will be required to sign another consent form with respect to the collection, use and disclosure of my personal information by the BAC.

I understand that:

- I do not have to sign this Consent Form, but if I do not, my healthcare provider(s) will not be able to disclose my personal information to the BAC and I will not be able to benefit from the services provided by the BAC (unless I contact the BAC directly myself);
- The medical treatment provided by my healthcare provider(s) will not be impacted by whether I sign this consent form or not;
- I may revoke (take back) this authorization at any time by mailing or faxing signed letter(s) of revocation to my healthcare provider(s), but if I do so, I will be unable to benefit from the services provided by the BAC;
- Revoking this authorization will prohibit disclosure of my health information by my healthcare provider(s) after the date my letter of revocation is received and processed, but will not affect the use or disclosure of information already received by the BAC;
- I am entitled to a copy of this Consent Form;
- If I want to access my patient file maintained by the BAC and/or make changes or corrections to it, I may do so by written request to the currently active BAC.

‡ In pediatric patients, it is recommended that STELARA® be administered by a healthcare provider.\*

\* STELARA® (ustekinumab) Product Monograph, Janssen Inc. Available at: <http://www.janssen.com/canada/>

Please see product monograph for full prescribing information for STELARA®.  
For more information, please contact Janssen Inc. Medical Information at 1-800-567-3331.