



Fax completed form to: 1-855-278-5182

Tel: 1-888-398-0028 | Email: accesslink@bioscript.ca

National Vaccine Services for Immune-Modulating Therapy Patient Enrolment Form

Patient Information		
First Name: _____	Last Name: _____	DOB (MM/DD/YYYY): _____
Phone: _____	Email: _____	Health Card #: _____
Address: _____	City: _____	Prov: _____ Postal Code: _____
Insurance Name: _____	Group ID: _____	Certificate ID: _____
Is the patient pregnant? <input type="checkbox"/> Yes <input type="checkbox"/> No		Gestation Age (weeks): _____
Allergies: _____		Intended Therapy: _____
Best time for contact: <input type="checkbox"/> Morning <input type="checkbox"/> Afternoon <input type="checkbox"/> Evening		Can the AccessLink team leave you a message? <input type="checkbox"/> Yes <input type="checkbox"/> No
Authorized Representative (if applicable) Name: _____ Phone: _____ Email: _____		
Vaccine Prescription		
Hepatitis A and B <input type="checkbox"/> Twinrix (Rapid Dosing) - IM 1mL dose given on day 0, day 7 and day 21 followed by a booster at 12 months for a total of 4 doses.	Meningococcal <input type="checkbox"/> Menveo, Menquadfi - One 0.5 mL dose IM	
Hepatitis B <input type="checkbox"/> Engerix-B- 20 mcg/dose - To be administered as 20 mcg (1mL) dose IM at month 0, 1 and 6 months after initial dosing. <input type="checkbox"/> Engerix-B- 20 mcg/dose - To be administered as 20 mcg (1mL) dose IM at month 0, 1, 2 and 6 months after initial dosing. <input type="checkbox"/> Engerix-B- 40 mcg/dose - To be administered as 20 mcg (1mL) dose IM in each arm at month 0, 1 and 6 months after initial dosing.	Pneumococcal <input type="checkbox"/> Prevnar 20 - One 0.5 mL dose IM.	
Human Papillomavirus <input type="checkbox"/> Gardasil 9 - One 0.5mL dose IM at month 0, second 0.5mL dose IM 2 months later, third 0.5mL dose IM 6 months after the first dose.	Polio <input type="checkbox"/> Imovax - One 0.5ml SC dose followed by a second dose after 4-8 weeks. Third dose is between 6-12 months after the second dose. <input type="checkbox"/> Imovax - One 0.5ml SC dose as a booster in adults if more than 10 years has elapsed since primary vaccination.	
Measles, Mumps, Rubella (Live-attenuated Vaccine) <input type="checkbox"/> M-M-R II - One 0.5mL dose SC. <input type="checkbox"/> M-M-R II - One 0.5mL dose SC followed by a second 0.5mL SC dose (minimum of 4 weeks later).	Respiratory Syncytial Virus <input type="checkbox"/> Abrysvo - One 0.5mL dose IM.	
Varicella (Live-attenuated Vaccine) <input type="checkbox"/> Varivax - One 0.5mL dose SC. <input type="checkbox"/> Varivax - One 0.5mL dose SC followed by a second 0.5mL SC dose (minimum of 4 weeks later).	Shingles <input type="checkbox"/> Shingrix - One 50 mcg (0.5 mL) dose IM at month 0, followed by a second 50 mcg dose IM between 2 and 6 months later (while not preferred, the second dose may be administered as early as 4 weeks).	
Tetanus, Diphtheria, Pertussis <input type="checkbox"/> Adacel, Boostrix - One 0.5mL dose IM. <input type="checkbox"/> Adacel, Boostrix - One 0.5mL dose IM at week 0 and then at week 4 and 6 months after the first dose.		
Other Instructions: Is patient vaccine history confirmed to indicate appropriateness of the selected vaccines? <input type="checkbox"/> Yes <input type="checkbox"/> No Pharmacist to review vaccines prescribed and dispense as appropriate		
Medical Clearance		
<input type="checkbox"/> Patient requires TB-Quantiferon testing prior to starting therapy	<input type="checkbox"/> Patient requires viral serology testing prior to starting therapy	
Additional comments:		
Patient Consent and Signature		
<input type="checkbox"/> Patient or Authorized Representative has consented to being contacted by AccessLink for Drug Navigation Support and agrees to the terms in Section A and B of this Form.		
Patient (or Authorized Representative) Signature: _____		Date (MM/DD/YYYY): _____
Verbal Consent Received by: _____		
Prescriber Authorization		
By signing this form, Prescriber affirms the within information and certifies that: Prescriber has discussed the AccessLink Services with the Patient or Authorized Representative, and they consent to be contacted by AccessLink as described. Prescriber authorizes AccessLink to act as the designated agent for the purposes of forwarding the prescription to the patient's pharmacy of choice. This prescription represents the original of the prescription drug order. The original prescription has been invalidated and securely filed, and it will not be re-transmitted or used elsewhere.		
Prescriber Name (Printed): _____	Fax: _____	Phone: _____
Address: _____	Licence #: _____	
Prescriber Signature: _____	Date (MM/DD/YYYY): _____	

A. Overview of AccessLink Service

AccessLink Drug Navigation Services (“**AccessLink**”) offered and administered by BioScript is aimed at simplifying access to patient care and medication access through assistance in administrative and drug reimbursement navigation support (the “**Services**”). As part of my participation in the Services, I understand that I will be offered confidential patient support services, at no cost, including but not limited to administrative, drug navigation and reimbursement support and may be contacted by phone, email or otherwise. Services offered may include:

- Benefits investigations and reimbursement support, including assistance in identifying potential coverage requirements and eligibility for financial support;
- Coordination of required paperwork and forms submission to facilitate coverage for prescribed medications;
- Coordination and enrolment with available patient support programs;
- Coordination of medical services including pharmacy, clinical and other paramedical services; and
- Such other services that AccessLink may offer you

The AccessLink Services do not provide medical advice or medical diagnosis. You agree to seek the advice of your treating physician or other qualified healthcare provider(s) if you have a health concern and not to disregard professional medical advice based on the information obtained from AccessLink. AccessLink reserves the right at any time, without notice, to modify, discontinue or terminate the Services.

You acknowledge that you have read the below Consent terms and you consent to the transfer of your personal information, health information, and the prescription (if applicable) to AccessLink.

B. Consent to Collection, Use and Disclosure of Personal Information

What information: You hereby authorize AccessLink to collect, use and disclose your personal information and health information to provide you the Services, including your:

- Health & Drug Insurance
- Prescription Information
- Drug Interactions
- Adverse Event Information
- Medical Conditions & Medical History
- Medication Shipment & Treatment Dates
- Personal Information (Name, Address, Contact Information)

Who may we interact with: AccessLink may collect, use and disclose your personal information and health information as needed to provide you the Services, including with your healthcare providers (physician, nurse practitioner, pharmacist, etc.), pharmacy of choice, public or private insurance or benefits provider and any Patient Support Program (“PSP”) that you are or will be enrolled in. You authorize AccessLink to collect and disclose your personal information and health information to and from the above listed individual(s) and organization(s).

For what purpose: The purpose of the collection, use and disclosure of your personal information and health information is to provide you the Services which may include drug reimbursement assistance, adverse drug event reporting, and to assist your PSP, healthcare providers, and pharmacy of choice to provide their services to you. AccessLink will use your information to provide the Services to you and may also use your information in an aggregate or de-identified form to improve its products and services.

Your personal information and health information collected as part of the Program will be protected by reasonable physical, administrative, and technical safeguards to protect it against loss, theft, and unauthorized access, communication, copying, use or alteration.

How long does this apply? This consent is effective from the day first written above and shall remain effective so long as you receive the Services and for a reasonable period of time thereafter. You may refuse to provide this consent to us or withdraw it at any time. If refuse consent, AccessLink will not be able to provide you with the Services. If you withdraw your consent, AccessLink will no longer be able to provide you with the Services, but such withdrawal will not be retroactive.

Your obligations:

- a) You must inform us if you cease to be enrolled in your PSP(s) for any reason or if there are any changes to your treating healthcare providers or pharmacy of choice.
- b) Provide us accurate information and updates so we can provide you the Services.

Acknowledgements:

- a) I understand that my personal information and health information may leave my province of residence and may be stored or processed outside of Canada. If this is the case, then my information would be subject to the laws of that country where it is stored and may be disclosed to that government under different circumstances than it would in Canada.
- b) I understand why I have been asked to provide consent to the disclosure of my personal information and health information, and I am aware of the risks and benefits of consenting or refusing to consent.
- c) I may ask any questions about privacy and compliance or exercise my privacy rights by contacting the Privacy Officer by email (privacyofficer@bioscript.ca) or telephone (1-888-734-3814).
- d) I understand I may withdraw this consent in writing at any time addressed to AccessLink using the contact information below.

AccessLink Drug Navigation Services

To enroll patients: Submit the completed form to AccessLink via fax at 1-855-278-5182. For assistance: email accesslink@bioscript.ca or phone 1-888-398-0028. Please note messages are checked daily and returned within two business days.