

Report of suspected adverse reaction due to **health products*** marketed in Canada

PROTECTED B**
 (when completed)

- The form should be printed and faxed toll free to: 1 866 678-6789 or mailed as per instructions below.
- La version française de ce document est disponible à: http://www.hc-sc.gc.ca/hpfb-dgpsa/tpd-dpt/adverse_f.pdf

A. Patient Information (See "Confidentiality" section below)					
1. Identifier		3. Sex		5. Weight	
2. Age at time of reaction		<input type="checkbox"/> Male		_____ feet	
		<input type="checkbox"/> Female		or _____ cm	
B. Adverse Reaction					
1. Outcome attributed to adverse reaction (check all that apply)					
<input type="checkbox"/> Death _____ (dd/mm/yyyy) <input type="checkbox"/> Disability <input type="checkbox"/> Life-threatening <input type="checkbox"/> Congenital malformation <input type="checkbox"/> Hospitalization <input type="checkbox"/> Required intervention to prevent damage/permanent impairment <input type="checkbox"/> Hospitalization - prolonged <input type="checkbox"/> Other : _____					
2. Date of reaction			3. Date of this report		
DD	MM	YYYY	DD	MM	YYYY
4. Describe reaction or problem					
5. Relevant tests / laboratory data (including dates (dd/mm/yyyy))					
6. Other relevant history, including pre-existing medical conditions (e.g. allergies, pregnancy, smoking and alcohol use, hepatic / renal dysfunction)					

C. Suspected Health Product(s) (See "How to report" section below)			
1. Name (give labeled strength & manufacturer, if known)			
# 1 _____			
# 2 _____			
2. Dose, frequency & route used		3. Therapy dates (if unknown, give duration)	
# 1		# 1 From (dd/ mm/yyyy - To (dd/ mm/yyyy)	
# 2		# 2	
4. Indication for use of suspected health product		5. Reaction abated after use stopped or dose reduced	
# 1		# 1 Yes No Doesn't apply	
# 2		# 2 Yes No Doesn't apply	
6. Lot # (if known)	7. Exp. date (if known)	8. Reaction reappeared after reintroduction	
# 1	# 1 (dd /mm/yyyy)	# 1 Yes No Doesn't apply	
# 2	# 2	# 2 Yes No Doesn't apply	
9. Concomitant health products (name, dose, frequency and route used), and therapy dates (dd/mm/yyyy) (exclude treatment of reaction)			
10. Treatment of adverse reaction (medications and / or other therapy), include dates (dd/mm/yyyy)			
D. Reporter Information (See "Confidentiality" section below)			
1. Name, address & phone number			
2. Health professional?		3. Occupation	4. Also reported to manufacturer?
Yes No			Yes No

Submission of a report does not constitute an admission that medical personnel or the product caused or contributed to the adverse reaction.
 * Use this form to report suspected adverse reactions to pharmaceuticals, biologics (including fractionated blood products, as well as therapeutic and diagnostic vaccines), natural health products or radiopharmaceuticals.
 ** As per the Treasury Board of Canada Secretariat Government Security Policy.

Return this form to the Adverse Reaction (AR) Centre listed below for your region

VOLUNTARY ADVERSE REACTION (AR) REPORTING GUIDELINES

Confidentiality of adverse reaction information

ARs to Canadian marketed health products, including prescription and non-prescription pharmaceuticals, biologics (including fractionated blood products, as well as therapeutic and diagnostic vaccines), natural health products and radiopharmaceuticals are collected by the Canadian Adverse Drug Reaction Monitoring Program (CADRMP). An AR is a harmful and unintended response to a health product. This includes any undesirable patient effect suspected to be associated with health product use. Unintended effect, health product abuse, overdose, interaction (including drug-drug and drug-food interactions) and unusual lack of therapeutic efficacy are all considered to be reportable ARs.

Any information related to the identity of the patient and/or the reporter of the AR is kept strictly confidential. For the “ identifier” box, provide some type of identifier that will allow you, the reporter, to readily locate the case if you are contacted for more information; do not use the patient’s name.

Privacy Notice Statement: Individuals have access to and protection of any provided personal information under the provisions of the *Access to Information Act* and the *Privacy Act*. Suspected health product-related AR information is submitted on a voluntary basis, and is maintained in a computerized database. AR information is used for the monitoring of marketed health products, and may contribute to the detection of potential product-related safety issues as well as to the benefit-risk assessments of these products. For more details with regard to personal information collected under this program, visit the Personal Information Bank; Health Canada; Health Products and Food Branch; Branch Incident Reporting System; PIB# PPU 088 at: http://infosource.gc.ca/inst/shc/fed07_e.asp.

What to report?

AR reports are, for the most part, only *suspected* associations. A temporal or possible association is sufficient for a report to be made. Reporting of an AR does not imply a definitive causal link.

All suspected adverse reactions should be reported, especially those that are:

- **unexpected**, regardless of their severity, i.e., not consistent with product information or labeling; or
- **serious**, whether expected or not; or
- reactions to **recently marketed health products** (on the market for less than five years), regardless of their nature or severity.

What is a serious adverse reaction?

A serious AR is one that requires in-patient hospitalization or prolongation of existing hospitalization, causes congenital malformation, results in persistent or significant disability or incapacity, is life-threatening or results in death. ARs that require significant medical intervention to prevent one of these listed outcomes are also considered to be serious.

How to report?

To report a suspected AR for health products marketed in Canada, health professionals or consumers (preferably in conjunction with their health professional, so that information about medical history can be included in order to make the reports more complete and scientifically valid) should complete a copy of the Report of Suspected Adverse Reaction Due to Health Products Marketed in Canada (HC/SC 4016). This form may be obtained from the Internet at http://www.hc-sc.gc.ca/hpfb-dgpsa/tpd-dpt/adverse_e.pdf, from your Regional AR Centre or from the National AR Centre (see contact information below), and is also available in the appendices of the Compendium of Pharmaceuticals and Specialties (CPS).

All applicable sections of the AR reporting form should be filled in as completely as possible. Use a separate form for each patient. Up to two suspected health products for a particular AR may be reported on one form. Attach an additional form if there are more than two suspected health products for the AR being reported. Additional pages may be attached if more space is required. The success of the program depends on the quality and accuracy of the information provided by the reporter.

To report an AR for a vaccine used in the prevention of infectious disease, the same criteria as stated in these guidelines are used. Health professionals should complete a copy of the Report of a Vaccine-Associated Adverse Event form. This form is available on the Internet at <http://www.phac-aspc.gc.ca/dird-dimr/pdf/hc4229e.pdf>, or in the appendices of the CPS. These forms also exist as customized Provincial/Territorial adverse event forms which can be obtained either from local public health departments or from the Provincial/Territorial health authorities.

For more information on CADRMP, additional copies of AR reporting forms or to report an AR, health professionals and consumers are invited to contact the Adverse Reaction Centre listed below for their region. The following toll-free numbers may be used by health professionals and consumers. Calls will be automatically routed to the appropriate Regional or National AR Centre based on the area code from which the call originates. Toll-free telephone: 1-866-234-2345 Toll-free fax: 1-866-678-6789.

British Columbia: British Columbia Regional AR Centre, c/o BC Drug and Poison Information Centre, 1081 Burrard Street, Vancouver, BC, V6Z 1Y6, adr@dpic.ca

Alberta: Alberta Regional Adverse Reaction Centre, c/o Suite 730, 9700 Jasper Ave, Edmonton, AB, T5J 4C3, Alberta_AR@hc-sc.gc.ca

Saskatchewan: Saskatchewan Regional AR Centre, c/o Saskatchewan Drug Information Service, College of Pharmacy and Nutrition, University of Saskatchewan, 110 Science Place, Saskatoon, SK, S7N 5C9, Sask_AR@usask.ca

Manitoba: Manitoba Regional Adverse Reaction Centre, Room 114, 510 Lagimodière Blvd, Winnipeg, MB, R2J 3Y1, Manitoba_AR@hc-sc.gc.ca

Ontario: Ontario Regional AR Centre, c/o LonDIS Drug Information Centre, London Health Sciences Centre, 339 Windermere Road, London, ON, N6A 5A5, adr@lhsc.on.ca

Québec: Québec Regional AR Centre, c/o Drug Information Centre, Hôpital du Sacré-Coeur de Montréal, 5400, boul. Gouin ouest, Montréal, QC, H4J 1C5, pharmacovigilance.hsc@ssss.gouv.qc.ca

Atlantic: Atlantic Regional AR Centre, For New Brunswick, Nova Scotia, Prince Edward Island, Newfoundland and Labrador, c/o Queen Elizabeth II Health Sciences Centre, Drug Information Centre, 1796 Summer Street, Rm 2421, Halifax, NS, B3H 3A7, adr@cdha.nshealth.ca

All Territories: National AR Centre, Marketed Health Products Safety and Effectiveness Information Division, Marketed Health Products Directorate, Tunney's Pasture Address locator: 0701C, Ottawa, ON, K1A 0K9, cadrm@hc-sc.gc.ca

How to deal with follow-up information for an AR that has already been reported?

Any follow-up information for an AR that has already been reported can be submitted using a new AR reporting form. It can be communicated by telephone, fax or e-mail to the appropriate Regional AR Centre or the National AR Centre (see contact information above). In order that this information can be matched with the original report, indicate that it is follow-up information, and if known, the date of the original report and the case report tracking number provided in the acknowledgement letter. It is very important that follow-up reports are identified and linked to the original report.

What about reporting ARs to the Market Authorization Holder (manufacturer)?

Health professionals and consumers may also report ARs to the market authorization holder (MAH). Indicate on your AR report sent to Health Canada if a case was also reported to the product’s MAH.