Final Report to the Minister

To: Canada Vigilance Program
Marketed Health Products Safety and Effectiveness Information Bureau
Marketed Health Products Directorate
Health Products and Food Branch
Health Canada
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From: **Name of Investigating Facility:** _____

[Subject to Blood Regulations section 115, on completion of the investigation, the investigating facility must file a final report with the Minister.]

This is a written final report to Health Canada for the serious and/or unexpected adverse recipient reaction concluded to be attributable to the quality and/or safety of a blood component(s) and related to a Health Canada-regulated activity carried out by our establishment.

Attached is the completed Saskatchewan Transfusion Adverse Event Report Form. *[Edit the patient identifier so that only the patient's age, gender and date of transfusion/reaction are sent.]*

File/Case #:			
Results of the Investigation: Was the root cause of the adverse recipient reaction attributable to an activity carried out in the facility?			
☐ Yes	☐ No		
Final Disposition of Implicated Blood Component			
☐ Transfused	☐ Quarantined	☐ Expired	☐ Discarded
Reason(s) for Disposition:			
Corrective actions taken and any other changes that are recommended to be made to relevant processes:			
Name of Other Establishments Notified:			
Report Prepared By:		**Date Reported:**	