Canada Vigilance

Summary of Reported Adverse Reactions

Report Runtime: {{report_runtime}} **Initial Received Date:** {{initial_received_date}} **Latest Received Date:** {{latest_received_date}} **Total Number of Reports:** {{no_of_reports}}

Report Information ***AER = Adverse Reaction Report

Adverse Reaction Report Number	Latest AER Version Number	Initial Received Date	Latest Received Date	Source of Report	Market Authorization Holder AER Number	Type of Report	Reporter Type
{{adverse_reaction_report_number}}	{{latest_aer_version_number}}	{{initial_received_date}}	{{latest_received_date}}	{{source_of_report}}	{{market_authorization_holder_aer_number}}	{{type_of_report}}	{{reporter_type}}

Serious Report
{{serious}}

Death	{{death}}	**Disability**	{{disability}}	**Congenital Anomaly**	{{anomaly}}
Life Threatening	{{life_threatening}}	**Hospitalization**	{{hospitalization}}	**Other Conditions**	{{other_conditions}}

Patient Information

Age	Gender	Height	Weight	Report Outcome
{{age}}	{{gender}}	{{height}}	{{weight}}	{{report_outcome}}

Link / Duplicate Report Information

Record Type	Link AER Number
{{record_type}}	{{link_aer_number}}

Product Information

Product Description	Health Product Role	Dosage Form	Route of Administration	Dose	Frequency	Therapy Duration	Indication(s)
{{product_description}}	{{health_product_role}}	{{dosage_form}}	{{route_of_administration}}	{{dose}}	{{frequency}}	{{therapy_duration}}	{{indication}}

Adverse Reaction Term Information

Adverse Reaction Term(s)	MedDRA Version	Reaction Duration
{{adverse_reaction_terms}}	{{meddra_version}}	{{reaction_duration}}