

¹For consistency this report follows the format used by the U.S. National Newborn Screening & Genetics Resource Center

Dot "●" indicates that screening for the condition is universally required by Law

A = universally offered but not yet required, **B** = offered to select populations or by request, **C** = testing required or offered universally but not yet implemented, **D** = likely to be detected (and reported) as a by-product of MRM screening (MS/MS) targeted by Law

Questions regarding the Canada Status Report should be directed to John Adams at john.adams7@sympatico.ca

Province/Territory (east to west/ west to east)	Core ¹ Conditions:									Additional Conditions Included (universally offered unless otherwise indicated)
	Hearing	Endocrine		Hemoglobin (HB)			Other			
	HEAR	CH	CAH	S/S	S/A	S/C	BIO	GALT	CF	
Province										
Alberta		A					A			
British Columbia	A	A						A		
Manitoba	A	A	A				A	A		DMD (B) (males)
New Brunswick	A	A								
Newfoundland & Labrador		A								
Nova Scotia	A	A								
Ontario	A	A	C	C	C	C	C	C		
Prince Edward Island	A	A								
Quebec	A	A								Amino acid transport disorders ²
Saskatchewan	B	●	C				C	B		
Yukon	A	A						A		
Territory										
Northwest Territories - eastern	A	A								
Northwest Territories - western	B	A					A			
Nunavut - eastern	A	A								
Nunavut - western	A	A	A				A	A		

¹Terminology consistent with ACMG report - Newborn Screening: Towards a Uniform Screening Panel and System. Genetics in Medicine. 2006; 8(Supp 1): S32-S33.

²Including Fanconi syndrome, Cystinurias, Hartnup syndrome, Prolidase deficiency, etc.

Deficiency/Disorder Abbreviations and Names

BIO	Biotinidase	CF	Cystic fibrosis	GALT	Transferase deficient galactosemia (Classical)	HB S/C	Sickle- C disease	HEAR	Hearing screening
CAH	Congenital adrenal hyperplasia	CH	Congenital hyperthyroidism	HB S/S	Sickle cell disease	HB S/A	S-beta thalassemia		

Additional Disorders

DMD	Duchenne Muscular Dystrophy
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Province/Territory (east to west; west to east)	Core ¹ Conditions: Metabolic																			
	Fatty Acid Disorders					Organic Acid Disorders								Amino Acid Disorders						
	CUD	LCHAD	MCAD	TFP	VLCAD	GA-I	HMG	IVA	3-MCC	Cbl-A,B	BKT	MUT	PROP	MCD	ASA	CIT	HCY	MSUD	PKU	TYR-I
Province																				
Alberta																	A	A	A	
British Columbia			A																A	
Manitoba						B													A	
New Brunswick	C	C	C	C	C	C		C										C	A	
Newfoundland & Labrador			A														A		A	A
Nova Scotia	A	A	A	A	A	A		A										A	A	
Ontario	C	C	A	C	C	C	C	C	C	C	C	C	C	C	C	C	C	C	A	C
Prince Edward Island	A	A	A	A	A	A		A									A	A		
Quebec						A	A	A	A	A		A			A	A			A	A
Saskatchewan	B	A	A	A	A	A	A	A	A	A	A	A	A	A	B	A	B	A	●	A
Yukon			A																A	
Territory																				
Northwest Territories - eastern																				A
Northwest Territories - western																				A
Nunavut - eastern						A	A	A	A	A		A			A	A			A	A
Nunavut - western																				A

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Deficiency/Disorder Abbreviations and Names (optional nomenclature)

3-MCC	3-Methylcrotonyl-CoA carboxylase	CUD	Carnitine uptake defect (Carnitine transport defect)	LCHAD	Long-chain hydroxyacyl-CoA dehydrogenase	PKU	Phenylketonuria/hyperphenylalaninemia
ASA	Argininosuccinate acidemia	GA-1	Glutaric acidemia type 1	MCAD	Medium-chain acyl-CoA dehydrogenase	PROP	Propionic acidemia (Propionyl-CoA carboxylase)
BKT	Beta ketothiolase (mitochondrial acetoacetyl-CoA thiolase ; short-chain ketoacyl thiolase; T2)	HCY	Homocystinuria (cystathionine beta synthase)	MCD	Multiple carboxylase (Holocarboxylase synthetase)	TFP	Trifunctional protein
CBL A,B	Methylmalonic acidemia (Vitamin B12 Disorders)	HMG	3-Hydroxy 3 - methylglutaric aciduria (3-Hydrox 3- methylglutaryl-CoA lyase)	MSUD	Maple syrup urine disease (branched-chain ketoacid dehydrogenase)	TYR-1	Tyrosinemia Type 1
CIT I	Citrullinemia type I (Argininosuccinate synthetase)	IVA	Isovaleric acidemia (Isovaleryl-CoA dehydrogenase)	MUT	Methylmalonic Acidemia (methylmalonyl-CoA mutase)	VLCAD	Very long-chain acyl-CoA dehydrogenase

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Province/ Territory (east to west; west to east)	Secondary Target ¹ Conditions																								
	Fatty Acid Disorders							Organic Acid Disorders						Amino Acid Disorders							Other Metabolic		Hgb		
	CACT	CPT-Ia	CPT-II	DE-RED.	GA-II	MCKAT	M/SCHAD	SCAD	2M3HBA	2MBG	3MGA	Cbl-C,D	IBG	MAL	ARG	BIOPT-BS	BIOPT-RG	CIT-II	H-PHE	MET	TYR-II	TYR-III	GALE	GALK	Variant hemoglobins
Province																									
Alberta																									
British Columbia																									
Manitoba		B																							
New Brunswick	C	C	C		C																				
Newfoundland & Labrador																					A				
Nova Scotia	A	A	A		A																				
Ontario																									
Prince Edward Island	A	A	A		A																				
Quebec											A			A			A								
Saskatchewan		A	A	B	A	A	A	A	A	A	A	A	A	A	B	B	A	A	A	A	A	B	B	B	
Yukon																									
Territory																									
Northwest T. - eastern																									
Northwest T. - western																									
Nunavut - eastern											A			A			A								
Nunavut - western																									

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Deficiency/Disorder Abbreviations and Names (optional names)

2M3HBA	2-Methyl-3-hydroxy butyric aciduria	CACT	Carnitine acylcarnitine translocase	GA-II	Glutaric acidemia type II	MAL	Malonic acidemia (Malonyl-CoA decarboxylase)
2MBG	2-Methylbutyryl-CoA dehydrogenase	CBL- C,D	Methylmalonic acidemia (Cbl C,D)	GALE	Galactose epimerase	MCKAT	Medium-chain ketoacyl-CoA thiolase
3MGA	3-Methylglutaconic aciduria	CIT-II	Citrullinemia type II	GALK	Galactokinase	MET	Hypermethioninemia
ARG	Arginemia Arginase Deficiency	CPT-Ia	Carnitine palmitoyltransferase I	H-PHE	Benign hyperphenylalaninemia	SCAD	Short-chain acyl-CoA dehydrogenase
BIOPT-BS	Defects of biopterin cofactor biosynthesis	CPT-II	Carnitine palmitoyltransferase II	IBG	Isobutyryl-CoA dehydrogenase	TYR-II	Tyrosinemia type II
BIOPT-REG	Defects of biopterin cofactor regeneration	De-Red	Dienoyl-CoA reductase	M/SCHAD	Medium/Short chain L-3-hydroxy acyl-CoA dehydrogenase	TYR-III	Tyrosinemia type III