National Diabetes Services Scheme
An Australian Government Initiative

NDSS Helpline 1800 637 700
ndss.com.au

‘Other’ diabetes eligible to register with the NDSS

Category A. Genetic defects of beta-cell function	
Chromosome 20, HNF-4alpha (MODY 1)	Chromosome 11, INS (MODY 10)
Chromosome 7, glucokinase (MODY 2)	Chromosome 8, BLK (MODY 11)
Chromosome 12, HNF-1alpha (MODY 3)	Monogenic
Chromosome 13, IPF-1 (MODY 4)	Chromosome 11, ABCC8
Chromosome 17, HNF-1beta (MODY 5)	Chromosome 11, KCNJ11
Chromosome 2, NeuroD1 (MODY 6)	Mitochondrial DNA
Chromosome 2, KLF11 (MODY 7)	Permanent neonatal diabetes
Chromosome 9, CEL (MODY 8)	Transient neonatal diabetes
Chromosome 7, PAX4 (MODY 9)	

Category B. Genetic defects in insulin action	
Leprechaunism or Donohue syndrome	Rabson-Mendenhall syndrome
Lipoatrophic diabetes	Type A insulin resistance

Category C. Diseases of the exocrine pancreas	
Cancer of the pancreas	Pancreatectomy / Whipple’s procedure
Cystic fibrosis (mucoviscidosis)	Pancreas trauma or surgically removed
Cystic Fibrosis Related Diabetes (CFRD)	Pancreatitis-induced
Fibrocalculous pancreatopathy (FCPP)	Pancreatogenic
Haemochromatosis or Iron overload disease induced	
Insulinoma	
Neoplasia	

Category D. Endocrinopathies	
Acromegaly or Somatotroph adenoma	Glucagonoma
Aldosteronoma	Pheochromocytoma
Cushing's syndrome or Hypercortisolism	Somatostatinoma

Category E. Drug or chemical induced	
Alpha-interferon	Highly active antiretroviral therapy (HAART)
Calcineurin inhibitors	NODAT or Post renal transplant
Chemically-induced	Pentamidine
Chemotherapy-induced	Post liver transplant
Diazoxide	Steroid-induced
Fluoroquinolones	Streptozotocin-induced diabetes (SID)
Glucocorticoids	Vacor

Category F. Infections	
Congenital rubella	Cytomegalovirus
Coxsackie	

Category G. Uncommon forms of immune-mediated diabetes
Anti-insulin receptor antibodies
"Stiff-man" syndrome / Stiff person syndrome (SPS)

Category H. Other genetic syndromes sometimes associated with diabetes*	
Acid Maltase Deficiency Myopathy (AMD) or Pompe disease	Laurence-Moon-Bardet-Biedl syndrome
Alstrom syndrome	Myotonic dystrophy / Myotonic Muscular Disorder (MMD)
Down syndrome	Porphyria
Friedreich's ataxia	Prader-Willi syndrome
Glycogen storage disease	Turner syndrome or Monosomy X
Huntington chorea	Wolfram syndrome or DIDMOAD
Klinefelter syndrome	

Miscellaneous conditions*	
Congenital hyperinsulinism (CHI) / Persistent hyperinsulinaemic hypoglycaemia of infancy (PHHI)**	Sarcoidosis
Russel-Silver syndrome	Other rare genetic conditions
*These conditions require further information to register with the NDSS. Please contact the NDSS Helpline on **1800 637 700** for further information.	
**Congenital hyperinsulinism (CHI)/Persistent hyperinsulinaemic hypoglycaemia of infancy (PHHI) is an eligible condition for NDSS registration.	