IU Health

Procedure Code Listing by Specialty

Report Prepared: February 11, 2011 at 8:42 AM

Specialty	Code	Procedure	Code
Metabolism-Pediatrics	MET-P		
		Amino Acid Disorders	AMINOACI
		Beckwith Syndrome	BECKWITH
		Beckwith-Wiedemann Syndrome	BECKWIED
		Biotinidase Deficiency	BIOTINID
		Classis Galactosemia	GALACTOS
		Cysturnia	CYSTINUR
		Erythropoietic Protoporphyria	ERYTHPRO
		Fatty Acid Oxidation Defects	FATTYACI
		Gaucher Disease	GAUCHER
		Glycogen Storage Disorders	GLYCSTOR
		Hereditary Fructose Intolerance	HERFRUCT
		Homocystinuria	HOMOCYST
		Hunter Syndrome	HUNTERSY
		Hurler Syndrome	HURLER
		Hyperammonemia	HYPERAMM
		Hypercholesterolemia	HYPERCHO
		Hypophosphatemic Rickets	HYPORICK
		Inborn Metabolism Errors	INBORNME
		Ketotic Hypoglycemia	KETOTHYP
		Leigh Syndrome	LEIGH
		Lipid Metabolism	LIPIDMET
		Lowe Syndrome	LOWE
		Maple Syrup Urine Disorder	MAPLESYR
		Med. Chain Acetyl-CoA Dehyd.&Car. Defici	MEDCHAIN
		Metabolic Disorders	METABDIS
		Metabolism	METABOLI
		Methylmalonic Acidemia	METHYLMA
		Mitochondrial Disorders	MITOCHON
		Molecular Genetics	MOLGEN
		Molecular Genetics of Fatty Acid Metabo.	MOLGENFA
		Mucopolysaccharidoses	MUCOPOLY
		Oxidation Phosphorylation Defects	OXIDATIO
		Phenylketonuria	PHENYLKE
		Prader-Willi Syndrome	PRADERWI
		Propionic Acidemia	PROPACID
		Sanfilippo Syndrome	SANFILIP
		Tay Sachs Disease	TAYSACHS
		Tyrosinemia	TYROS
		Urea cycle disorders	UREACYCL

Clarian Health Partners - Marketing Group

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Specialty	Code	Procedure	Code
Metabolism-Pediatrics	MET-P		
		Zellweger Syndrome	ZELLWEGE