Belgian Genetic Tests database

ANALYTE:
LMNA

NAME:	lamin A/C
SYMBOL:	LMNA
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SYNONYMS:	HGPS MADA mandibuloacral dysplasia type A
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Cardiomyopathy, hereditary (gene panel)
- Cardiomyopathy: hypertrophic cardiomyopathy, dilated cardiomyopathy, restrictive cardiomyopathy, left ventricular non-compaction cardiomyopathy, arrhythmogenic right ventricular cardiomyopathy (gene panel)
- Cardiopathies, hereditary (gene panel)
- Charcot-Marie-Tooth (other than type 1A) (gene panel, IPN panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Congenital structural heart defects (gene panel)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Dilated Cardiomyopathy (Gene panel)
- Dilated cardiomyopathy
- Heart / Cardio disorders / Cardiopathy (gene panel)
- Hypertrophic cardiomyopathy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Lipodystrophy and/or hyperinsulinism (gene panel)
- Myopathy (gene panel)
- Myopathy (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Neuromuscular disorders (232 genes (= myopathy, metabolic myopathy, ion channel muscle diseases, muscular dystrophy, myotonic dystrophy, rhabdomyolysis, myasthenia)
- Neuromuscular disorders (548 genes)
- Neuromuscular disorders (gene panel)
- Neuromuscular disorders : congenital & distal myopathy, congenital muscle dystrophy / Limb-girdle muscular dystrophy / Rhabdomyolysis / Myopathy (with prominent contractures) / distal artrogryposis (gene panel)
- Neuropathy (gene panel)

- Neuropathy (gene panel)
- Peripheral neuropathy (gene panel)
- Primary Electrical disorders / Brugada syndrome / Long QT syndrome (LQT) / Short QT syndrome (SQT) / Arrhythmogenic right ventricular cardiomyopathy (ARVC) / Catecholaminergic polymorphic ventricular tachycardia (CPVT) (gene panel)
- Primary cardiac arrhythmias (Atrial fibrillation / Brugada syndome / Catech. polymorphic ventricular tachycardia / Early repolaristion syndrome / Ideopathic ventricular fibrillation / Long QT syndrome / Sick sinus syndrome / Short QT syndrome) (gene pane)
- Primary ciliary dyskinesia (PCD) Heterotaxyies (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skin disorders (gene panel)
- Stroke (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- Atypical Werner syndrome
- Autosomal dominant Emery-Dreifuss muscular dystrophy
- Autosomal dominant limb-girdle muscular dystrophy type 1B
- Autosomal recessive Emery-Dreifuss muscular dystrophy
- Autosomal semi-dominant severe lipodystrophic laminopathy
- Charcot-Marie-Tooth disease type 2B1
- Congenital muscular dystrophy due to LMNA mutation
- Dilated cardiomyopathy-hypergonadotropic hypogonadism syndrome
- Familial dilated cardiomyopathy with conduction defect due to LMNA mutation
- Familial isolated arrhythmogenic ventricular dysplasia, biventricular form
- Familial isolated arrhythmogenic ventricular dysplasia, left dominant form
- Familial isolated arrhythmogenic ventricular dysplasia, right dominant form
- Familial isolated dilated cardiomyopathy
- Familial partial lipodystrophy, Dunnigan type
- Familial partial lipodystrophy, Köbberling type
- Heart-hand syndrome, Slovenian type

- Hutchinson-Gilford progeria syndrome
- LMNA-related cardiocutaneous progeria syndrome
- Left ventricular noncompaction
- Mandibuloacral dysplasia with type A lipodystrophy
- Restrictive dermopathy

Related Gene Panels

- Inherited Peripheral Neuropathies gene panel (139 genes) - KUL
- Cardiomyopathy (genepanel) - UZA
- Cardiomyopathy, hereditary (208 genes) - VUB
- Cardiopathies, hereditary (102 genes) - KUL
- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Dilated Cardiomyopathy (79 genes) - IPG
- Dilated Cardiomyopathy (DCMP) - UGent
- Dilated cardiomyopathy - UGent
- Heterotaxie PCD - UGent
- Hypertrophic cardiomyopathy (75 genes) - IPG
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Lipodystrophy and/or hyperinsulinism (30 genes) - IPG
- Myopathy (332 genes) - IPG
- Myopathy (genepanel) - UZA
- Nephropathy panel - UGent
- Neuromuscular disorders (548 genes) - ULB
- Neuromuscular disorders (166 genes) - VUB
- Neuromuscular disorders (232 genes) - KUL
- Neuromuscular disorders - UGent
- Neuropathy (148 genes) - IPG

- Neuropathy (genepanel) - UZA
- Neuropathy panel - UGent
- Primary Electrical disorders/Brugada syndrome (genepanel) - UZA
- Primary cardiac arrhythmias (113 genes) - VUB
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Skin disorders - UGent
- Stroke - UGent
- cardiopathy panel - UGent

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