LGMD: ICD-10 codes

ICD-10 Code	Subtype, if applicable	Description
G71.031	LGMD1/D	Autosomal dominant LGMD
G71.032	LGMD2A/R1	Autosomal recessive LGMD due to calpain-3 dysfunction (calpainopathy)
G71.033	LGMD2B/R2	LGMD due to dysferlin dysfunction (dysferlinopathy)
G71.0340		LGMD due to sarcoglycan dysfunction, unspecified (sarcoglycanopathy)
G71.0341	LGMD2D/R3	LGMD due to alpha sarcoglycan dysfunction (alpha-sarcoglycanopathy)
G71.0342	LGMD2E/R4	LGMD due to beta sarcoglycan dysfunction (beta-sarcoglycanopathy)
G71.0349	LGMD2C/R5* LGMD2F/R6*	LGMD due to other sarcoglycan dysfunction
G71.035	LGMD2L/R12	LGMD due to anoctamin-5 dysfunction (anoctaminopathy)
G71.038		Other LGMD**
G71.039		LGMD, unspecified***

*LGMD2C/R5 is caused by mutations in the SGCG gene, which encodes gamma-sarcoglycan. LGMD2F/R6 is caused by mutations in the SGCD gene, which encodes delta-sarcoglycan.

**G71.038 is intended for all other forms of autosomal recessive LGMD, such as LGMD2I/R9 (caused by mutations in fukutin-related protein gene, FKRP).

***G71.039 is intended for patients that do not have a genetically confirmed LGMD diagnosis. If a patient has not yet received genetic testing to confirm their LGMD diagnosis, please visit limbgirdle.com/genetic-testing for more information.

Limb-Girdle Muscular Dystrophy: ICD-10 codes

SAREPTA THERAPEUTICS

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