Potassium-aggravated myotonia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Potassium-aggravated myotonia. ORPHA:612

Potassium-aggravated myotonia (PAM) is a muscular channelopathy presenting with a pure myotonia dramatically aggravated by potassium ingestion, with variable cold sensitivity and no episodic weakness. This group includes three forms: myotonia fluctuans, myotonia permanens, and acetazolamide-responsive myotonia (see these terms).