Brugada syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Brugada syndrome. ORPHA:130

Brugada syndrome (BrS) manifests with ST segment elevation in right precordial leads (V1 to V3), incomplete or complete right bundle branch block, and susceptibility to ventricular tachyarrhythmia and sudden death. BrS is an electrical disorder without overt myocardial abnormalities.