

8.1.2 Approach to risk stratification and management

Risk stratification and management in Long QT Syndrome

Recommendations	Class ^a	Level ^b	Ref. ^c
<p>The following lifestyle changes are recommended in all patients with a diagnosis of LQTS:</p> <p>(a) Avoidance of QT-prolonging drugs (http://www.crediblemeds.org).</p> <p>(b) Correction of electrolyte abnormalities (hypokalaemia, hypomagnesaemia, hypocalcaemia) that may occur during diarrhoea, vomiting or metabolic conditions.</p> <p>(c) Avoidance of genotype-specific triggers for arrhythmias (strenuous swimming, especially in LQTS1, and exposure to loud noises in LQTS2 patients).</p>	I	B	434

Beta-blockers are recommended in patients with a clinical diagnosis of LQTS.	I	B	435
ICD implantation with the use of beta-blockers is recommended in LQTS patients with previous cardiac arrest.	I	B	436–438
Beta-blockers should be considered in carriers of a causative LQTS mutation and normal QT interval.	IIa	B	67
ICD implantation in addition to beta-blockers should be considered in LQTS patients who experienced syncope and/or VT while receiving an adequate dose of beta-blockers.	IIa	B	439
<p>Left cardiac sympathetic denervation should be considered in patients with symptomatic LQTS when</p> <p>(a) Beta-blockers are either not effective, not tolerated or contraindicated;</p> <p>(b) ICD therapy is contraindicated or refused;</p> <p>(c) Patients on beta-blockers with an ICD experience multiple shocks.</p>	IIa	C	440
Sodium channel blockers (mexiletine, flecainide or ranolazine) may be considered as add-on therapy to shorten the QT interval in LQTS3 patients with a QTc > 500 ms.	IIb	C	441–443
Implant of an ICD may be considered in addition to beta-blocker therapy in asymptomatic carriers of a pathogenic mutation in <i>KCNH2</i> or <i>SCN5A</i> when QTc is > 500 ms.	IIb	C	67