A novel mutation in KVLQT1 is the molecular basis of inherited long QT syndrome in a near-drowning patient's family

Ackerman MJ, Schroeder JJ, Berry R, Schaid DJ, Porter CJ, Michels VV, Thibodeau SN.
Pediatric research
1998; 44(2):148-153

ARTICLE IDENTIFIERS
DOI: unavailable
PMID: 9702906
PMCID: not available

JOURNAL IDENTIFIERS
LCCN: sf 80001040
pISSN: 0031-3998
eISSN: 1530-0447
OCLC ID: 01761994
CONS ID: sn 78005036
US National Library of Medicine ID: 0100714