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Genetic basis of cardiac ion channel diseases

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List of publications

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A common polymorphism in *KCNH2* (HERG) hastens cardiac repolarization.
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Sodium channel β 1-subunit mutations associated with Brugada syndrome and cardiac conduction disease.

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Biophysical properties of cardiac ion channels.

To be published as bookchapter in: *Ion Channel Biophysics and Diseases (Research Signpost, Kerala, India).*