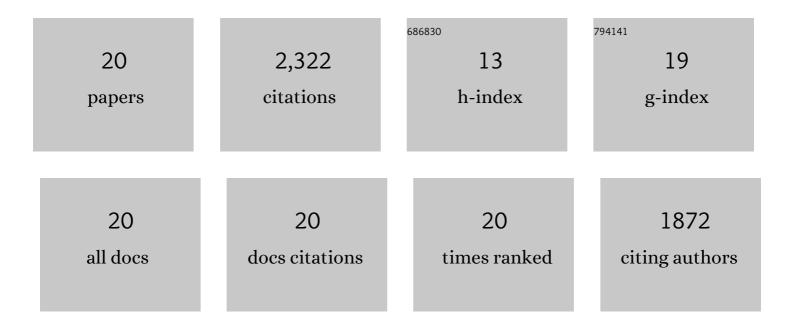
Li Zhang

List of Publications by Year in descending order

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LI ZHANC

| # | Article | IF | CITATIONS |
|----|--|-----|-----------|
| 1 | Hospitalized cancer patients with acquired long QT syndrome-a matched case-control study. Cardio-Oncology, 2020, 6, 3. | 0.8 | 5 |
| 2 | Unique ECG presentations and clinical management of a symptomatic LQT2 female carrying a novel de novo KCNH2 mutation. Journal of Electrocardiology, 2018, 51, 111-116. | 0.4 | 8 |
| 3 | The role of mexiletine in the management of long QT syndrome. Journal of Electrocardiology, 2018, 51, 1061-1065. | 0.4 | 26 |
| 4 | Acquired long QT syndrome in hospitalized patients. Heart Rhythm, 2017, 14, 974-978. | 0.3 | 42 |
| 5 | A New Formula for Estimating the True QT Interval in Left Bundle Branch Block. Journal of Cardiovascular Electrophysiology, 2017, 28, 684-689. | 0.8 | 18 |
| 6 | Isolated left ventricular arrhythmogenic cardiomyopathy: A case report. Journal of Electrocardiology, 2017, 50, 144-147. | 0.4 | 3 |
| 7 | Common Genotypes of Long QT Syndrome in China and the Role of ECG Prediction. Cardiology, 2016, 133, 73-78. | 0.6 | 16 |
| 8 | Right Ventricular Outflow Tract Tachycardia with Structural Abnormalities of the Right Ventricle and Left Ventricular Diverticulum. Case Reports in Cardiology, 2015, 2015, 1-3. | 0.1 | 2 |
| 9 | The Electrocardiographic Manifestations of Arrhythmogenic Right Ventricular Dysplasia. Current Cardiology Reviews, 2014, 10, 237-245. | 0.6 | 53 |
| 10 | Long QT syndrome, cardiovascular anomaly and findings in ECG-guided genetic testing. International Journal of Cardiology Heart & Vessels, 2014, 4, 122-128. | 0.5 | 6 |
| 11 | Is the Phenotype–Genotype Relationship Necessary to Understand Cardiomyopathies?. Circulation: Cardiovascular Genetics, 2014, 7, 405-406. | 5.1 | 0 |
| 12 | Inhibition of Late Sodium Current by Mexiletine. Circulation: Arrhythmia and Electrophysiology, 2013, 6, 614-622. | 2.1 | 75 |
| 13 | Nonsense-mediated mRNA decay caused by a frameshift mutation in a large kindred of type 2 long QT syndrome. Heart Rhythm, 2011, 8, 1200-1206. | 0.3 | 27 |
| 14 | Mutations of Plakophilin-2 in Chinese With Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy. American Journal of Cardiology, 2009, 103, 1439-1444. | 0.7 | 22 |
| 15 | Nonsense Mutations in hERG Cause a Decrease in Mutant mRNA Transcripts by Nonsense-Mediated mRNA Decay in Human Long-QT Syndrome. Circulation, 2007, 116, 17-24. | 1.6 | 111 |
| 16 | Long QT Syndrome in Adults. Journal of the American College of Cardiology, 2007, 49, 329-337. | 1.2 | 369 |
| 17 | Inaccurate electrocardiographic interpretation of long QT: The majority of physicians cannot recognize a long QT when they see one. Heart Rhythm, 2005, 2, 569-574. | 0.3 | 345 |
| 18 | An intronic mutation causes long QT syndrome. Journal of the American College of Cardiology, 2004, 44, 1283-1291. | 1.2 | 57 |

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| # | Article | IF | CITATIONS |
|----|--|------|-----------|
| 19 | Spectrum of ST-T–Wave Patterns and Repolarization Parameters in Congenital Long-QT Syndrome. Circulation, 2000, 102, 2849-2855. | 1.6 | 409 |
| 20 | Influence of the Genotype on the Clinical Course of the Long-QT Syndrome. New England Journal of Medicine, 1998, 339, 960-965. | 13.9 | 728 |