Norman Stockbridge

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/10735630/publications.pdf

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72 papers 5,136 citations

32 h-index 91884 69 g-index

77 all docs

77
docs citations

times ranked

77

5790 citing authors

#	Article	IF	CITATIONS
1	Defining High Bleeding Risk in Patients Undergoing Percutaneous Coronary Intervention. Circulation, 2019, 140, 240-261.	1.6	428
2	The Comprehensive in Vitro Proarrhythmia Assay (CiPA) initiative $\hat{a} \in \text{``Update}$ on progress. Journal of Pharmacological and Toxicological Methods, 2016, 81, 15-20.	0.7	335
3	Defining high bleeding risk in patients undergoing percutaneous coronary intervention: a consensus document from the Academic Research Consortium for High Bleeding Risk. European Heart Journal, 2019, 40, 2632-2653.	2.2	335
4	Evolution of strategies to improve preclinical cardiac safety testing. Nature Reviews Drug Discovery, 2016, 15, 457-471.	46.4	323
5	Developing Therapies for Heart Failure WithÂPreservedÂEjection Fraction. JACC: Heart Failure, 2014, 2, 97-112.	4.1	267
6	International Multisite Study of Human-Induced Pluripotent Stem Cell-Derived Cardiomyocytes for Drug Proarrhythmic Potential Assessment. Cell Reports, 2018, 24, 3582-3592.	6.4	254
7	Comprehensive Translational Assessment of Human-Induced Pluripotent Stem Cell Derived Cardiomyocytes for Evaluating Drug-Induced Arrhythmias. Toxicological Sciences, 2017, 155, 234-247.	3.1	213
8	Concentrationâ€QT Relationships Play a Key Role in the Evaluation of Proarrhythmic Risk During Regulatory Review. Journal of Clinical Pharmacology, 2008, 48, 13-18.	2.0	206
9	Cardiovascular Drug Development. Journal of the American College of Cardiology, 2015, 65, 1567-1582.	2.8	168
10	Dealing with Global Safety Issues. Drug Safety, 2013, 36, 167-182.	3.2	134
11	Comprehensive T wave Morphology Assessment in a Randomized Clinical Study of Dofetilide, Quinidine, Ranolazine, and Verapamil. Journal of the American Heart Association, 2015, 4, .	3.7	115
12	Current challenges in the evaluation of cardiac safety during drug development: Translational medicine meets the Critical Path Initiative. American Heart Journal, 2009, 158, 317-326.	2.7	113
13	Mechanistic Modelâ€Informed Proarrhythmic Risk Assessment of Drugs: Review of the "CiPAâ€Initiative and Design of a Prospective Clinical Validation Study. Clinical Pharmacology and Therapeutics, 2018, 103, 54-66.	4.7	106
14	Heart Failure With Preserved Ejection Fraction Expert Panel Report. JACC: Heart Failure, 2018, 6, 619-632.	4.1	103
15	The IQâ€CSRC Prospective Clinical Phase 1 Study: "Can Early QT Assessment Using Exposure Response Analysis Replace the Thorough QT Study?― Annals of Noninvasive Electrocardiology, 2014, 19, 70-81.	1.1	92
16	Assessing proarrhythmic potential of drugs when optimal studies are infeasible. American Heart Journal, 2009, 157, 827-836.e1.	2.7	81
17	Novel oral anticoagulants and reversal agents: Considerations for clinical development. American Heart Journal, 2015, 169, 751-757.	2.7	69
18	The Evolving Roles of Human iPSC-Derived Cardiomyocytes in Drug Safety and Discovery. Cell Stem Cell, 2017, 21, 14-17.	11.1	69

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19	Assessment of Multi″on Channel Block in a Phase I Randomized Study Design: Results of the Ci∢scp>PA⟨/scp> Phase I ⟨scp>ECG⟨/scp> Biomarker Validation Study. Clinical Pharmacology and Therapeutics, 2019, 105, 943-953.	4.7	66
20	Workshop Report. Circulation Research, 2019, 125, 855-867.	4. 5	53
21	Implications of the IQ-CSRC Prospective Study: Time to Revise ICHÂE14. Drug Safety, 2015, 38, 773-780.	3.2	52
22	Prevalent and Incident Heart Failure inÂCardiovascular Outcome Trials of Patients With Type 2 Diabetes. Journal of the American College of Cardiology, 2018, 71, 1379-1390.	2.8	50
23	Exploring New Endpoints for Patients With Heart Failure With Preserved Ejection Fraction. Circulation: Heart Failure, 2016, 9, .	3.9	46
24	Evaluation of Batch Variations in Induced Pluripotent Stem Cell-Derived Human Cardiomyocytes from 2 Major Suppliers. Toxicological Sciences, 2017, 156, kfw235.	3.1	45
25	Implications of geographical variation on clinical outcomes of cardiovascular trials. American Heart Journal, 2012, 164, 303-312.	2.7	44
26	Improving Heart Failure Therapeutics Development in the United States. Journal of the American College of Cardiology, 2018, 71, 443-453.	2.8	40
27	A proposal for scientific framework enabling specific population drug dosing recommendations. Journal of Clinical Pharmacology, 2015, 55, 1073-1078.	2.0	39
28	Conduct of Clinical Trials in the Era of COVID-19. Journal of the American College of Cardiology, 2020, 76, 2368-2378.	2.8	35
29	Trial Design Principles for Patients at HighÂBleeding Risk Undergoing PCI. Journal of the American College of Cardiology, 2020, 76, 1468-1483.	2.8	35
30	Cardiovascular outcome trials in patients with chronic kidney disease: challenges associated with selection of patients and endpoints. European Heart Journal, 2019, 40, 880-886.	2.2	34
31	Personalized Cardiovascular Medicine Today. Circulation, 2015, 132, 1425-1432.	1.6	33
32	Universal Correction for QT/RR Hysteresis. Drug Safety, 2016, 39, 577-588.	3.2	33
33	Assessment of drug-induced increases in blood pressure during drug development: Report from the Cardiac Safety Research Consortium. American Heart Journal, 2013, 165, 477-488.	2.7	30
34	Drugâ€induced Proarrhythmia and Torsade de Pointes: A Primer for Students and Practitioners of Medicine and Pharmacy. Journal of Clinical Pharmacology, 2018, 58, 997-1012.	2.0	28
35	Endpoints in HeartÂFailure DrugÂDevelopment. JACC: Heart Failure, 2020, 8, 429-440.	4.1	28
36	The Cardiac Safety Research Consortium electrocardiogram warehouse: Thorough QT database specifications and principles of use for algorithm development and testing. American Heart Journal, 2010, 160, 1023-1028.	2.7	26

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37	Centralized adjudication of cardiovascular end points in cardiovascular and noncardiovascular pharmacologic trials: A report from the Cardiac Safety Research Consortium. American Heart Journal, 2015, 169, 197-204.	2.7	25
38	Evolving regulatory paradigm for proarrhythmic risk assessment for new drugs. Journal of Electrocardiology, 2016, 49, 837-842.	0.9	24
39	New Strategies for the Conduct of Clinical Trials in Pediatric Pulmonary Arterial Hypertension: Outcome of a Multistakeholder Meeting With Patients, Academia, Industry, and Regulators, Held at the European Medicines Agency on Monday, June 12, 2017. Journal of the American Heart Association, 2019, 8. e011306.	3.7	23
40	Standardized definitions for evaluation of heart failure therapies: scientific expert panel from the Heart Failure Collaboratory and Academic Research Consortium. European Journal of Heart Failure, 2020, 22, 2175-2186.	7.1	23
41	Practice and challenges of thorough QT studies. Journal of Electrocardiology, 2012, 45, 582-587.	0.9	22
42	Early Drug Discovery Prediction of Proarrhythmia Potential and Its Covariates. AAPS Journal, 2015, 17, 1025-1032.	4.4	22
43	Heart Failure End Points in Cardiovascular Outcome Trials of Sodium Glucose Cotransporter 2 Inhibitors in Patients With Type 2 Diabetes Mellitus. Circulation, 2019, 140, 2108-2118.	1.6	22
44	Cardiovascular Safety Outcome Trials: A meeting report from the Cardiac Safety Research Consortium. American Heart Journal, 2015, 169, 486-495.	2.7	21
45	Moxifloxacinâ€induced QT _c interval prolongations in healthy male Japanese and Caucasian volunteers: a direct comparison in a thorough QT study. British Journal of Clinical Pharmacology, 2015, 80, 446-459.	2.4	20
46	Improving cardiovascular clinical trials conduct in the United States: Recommendation from clinicians, researchers, sponsors, and regulators. American Heart Journal, 2015, 169, 305-314.	2.7	20
47	Can Bias Evaluation Provide Protection Against Falseâ€Negative Results in QT Studies Without a Positive Control Using Exposureâ€Response Analysis?. Journal of Clinical Pharmacology, 2017, 57, 85-95.	2.0	20
48	Errors of Fixed QT Heart Rate Corrections Used in the Assessment of Drug-Induced QTc Changes. Frontiers in Physiology, 2019, 10, 635.	2.8	18
49	Importance of QT/RR hysteresis correction in studies of drug-induced QTc interval changes. Journal of Pharmacokinetics and Pharmacodynamics, 2018, 45, 491-503.	1.8	15
50	Standardized Definitions for EvaluationÂofÂHeart Failure Therapies: Scientific Expert Panel From the HeartÂFailure Collaboratory and Academic Research Consortium. JACC: Heart Failure, 2020, 8, 961-972.	4.1	15
51	Challenges of Cardio-Kidney Composite Outcomes in Large-Scale Clinical Trials. Circulation, 2021, 143, 949-958.	1.6	15
52	Reassessing Phase II Heart Failure Clinical Trials. Circulation: Heart Failure, 2017, 10, .	3.9	14
53	Implications of Individual QT/RR Profilesâ€"Part 1: Inaccuracies and Problems of Population-Specific QT/Heart Rate Corrections. Drug Safety, 2019, 42, 401-414.	3.2	14
54	Lessons Learned From Hundreds of Thorough QT Studies. Therapeutic Innovation and Regulatory Science, 2015, 49, 392-397.	1.6	13

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55	Utility of Model-Based Approaches for Informing Dosing Recommendations in Specific Populations: Report From the Public AAPS Workshop. Journal of Clinical Pharmacology, 2017, 57, 105-109.	2.0	12
56	Detection of T Wave Peak for Serial Comparisons of JTp Interval. Frontiers in Physiology, 2019, 10, 934.	2.8	12
57	Effects of Electrical Stimulation on hiPSC-CM Responses to Classic Ion Channel Blockers. Toxicological Sciences, 2020, 174, 254-265.	3.1	12
58	Long-term electrocardiographic safety monitoring in clinical drug development: A report from the Cardiac Safety Research Consortium. American Heart Journal, 2017, 187, 156-169.	2.7	11
59	Heart Rate Correction of the J-to-Tpeak Interval. Scientific Reports, 2019, 9, 15060.	3.3	10
60	The Cardiac Safety Research Consortium enters its second decade: An invitation to participate. American Heart Journal, 2016, 177, 96-101.	2.7	9
61	Sex differences in drug-induced changes in ventricular repolarization. Journal of Electrocardiology, 2015, 48, 1081-1087.	0.9	8
62	Design of a "Lean―Case Report Form for HeartÂFailure Therapeutic Development. JACC: Heart Failure, 2019, 7, 913-921.	4.1	6
63	Implications of Individual QT/RR Profilesâ€"Part 2: Zero QTc/RR Correlations Do Not Prove QTc Correction Accuracy in Studies of QTc Changes. Drug Safety, 2019, 42, 415-426.	3.2	5
64	Resourcing Drug Development Commensurate With its PublicÂHealthÂImportance. JACC Basic To Translational Science, 2016, 1, 309-312.	4.1	4
65	Thorough QT Studies and Indirect Causes of QTc Changes. PACE - Pacing and Clinical Electrophysiology, 2012, 35, 1411-1412.	1.2	3
66	2017 ACC/AAP/AHA Health Policy Statement on Opportunities and Challenges in Pediatric Drug Development: Learning From Sildenafil. Circulation: Cardiovascular Quality and Outcomes, 2017, 10, .	2.2	3
67	The FDA in the 21st Century. JACC: Heart Failure, 2017, 5, 67-70.	4.1	2
68	2017 ACC/AAP/AHA Health Policy Statement on Opportunities and Challenges in Pediatric Drug Development: Learning From Sildenafil. Journal of the American College of Cardiology, 2017, 70, 495-503.	2.8	2
69	Methods for Employing Information About Uncertainty of Ascertainment of Events in Clinical Trials. Therapeutic Innovation and Regulatory Science, 2021, 55, 197-211.	1.6	2
70	Cardiac Safety Research Consortium (CSRC): Cardiovascular Safety and Adverse Event Case Report Forms. Therapeutic Innovation and Regulatory Science, 2015, 49, 511-513.	1.6	1
71	Topic of Timely Interest—Decision Criteria for Negative QT Assessment Using Exposure Response Analysis of Data From Early-Phase Clinical Studies: Letter to the Editor. Therapeutic Innovation and Regulatory Science, 2015, 49, 717-719.	1.6	1
72	Ask the Expert: A Regulatory Perspective on Clinical Trials for Pulmonary Arterial Hypertension. Advances in Pulmonary Hypertension, 2020, 19, 62-65.	0.1	1