Norman Stockbridge

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/10735630/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	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
2	Challenges of Cardio-Kidney Composite Outcomes in Large-Scale Clinical Trials. Circulation, 2021, 143, 949-958.	1.6	15
3	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
4	Conduct of Clinical Trials in the Era of COVID-19. Journal of the American College of Cardiology, 2020, 76, 2368-2378.	2.8	35
5	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
6	Trial Design Principles for Patients at HighÂBleeding Risk Undergoing PCI. Journal of the American College of Cardiology, 2020, 76, 1468-1483.	2.8	35
7	Effects of Electrical Stimulation on hiPSC-CM Responses to Classic Ion Channel Blockers. Toxicological Sciences, 2020, 174, 254-265.	3.1	12
8	Endpoints in HeartÂFailure DrugÂDevelopment. JACC: Heart Failure, 2020, 8, 429-440.	4.1	28
9	Ask the Expert: A Regulatory Perspective on Clinical Trials for Pulmonary Arterial Hypertension. Advances in Pulmonary Hypertension, 2020, 19, 62-65.	0.1	1
10	Design of a "Lean―Case Report Form for HeartÂFailure Therapeutic Development. JACC: Heart Failure, 2019, 7, 913-921.	4.1	6
11	Detection of T Wave Peak for Serial Comparisons of JTp Interval. Frontiers in Physiology, 2019, 10, 934.	2.8	12
12	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
13	Workshop Report. Circulation Research, 2019, 125, 855-867.	4.5	53
14	Heart Rate Correction of the J-to-Tpeak Interval. Scientific Reports, 2019, 9, 15060.	3.3	10
15	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
16	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
17	Defining High Bleeding Risk in Patients Undergoing Percutaneous Coronary Intervention. Circulation, 2019, 140, 240-261.	1.6	428
18	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

#	Article	IF	CITATIONS
19	Assessment of Multiâ€lon 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	Implications of Individual QT/RR Profiles—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
21	Implications of Individual QT/RR Profiles—Part 1: Inaccuracies and Problems of Population-Specific QT/Heart Rate Corrections. Drug Safety, 2019, 42, 401-414.	3.2	14
22	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
23	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
24	Improving Heart Failure Therapeutics Development in the United States. Journal of the American College of Cardiology, 2018, 71, 443-453.	2.8	40
25	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
26	Mechanistic Modelâ€Informed Proarrhythmic Risk Assessment of Drugs: Review of the "CiPA―Initiative and Design of a Prospective Clinical Validation Study. Clinical Pharmacology and Therapeutics, 2018, 103, 54-66.	4.7	106
27	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
28	Heart Failure With Preserved Ejection Fraction Expert Panel Report. JACC: Heart Failure, 2018, 6, 619-632.	4.1	103
29	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
30	Evaluation of Batch Variations in Induced Pluripotent Stem Cell-Derived Human Cardiomyocytes from 2 Major Suppliers. Toxicological Sciences, 2017, 156, kfw235.	3.1	45
31	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
32	The FDA in the 21st Century. JACC: Heart Failure, 2017, 5, 67-70.	4.1	2
33	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
34	Reassessing Phase II Heart Failure Clinical Trials. Circulation: Heart Failure, 2017, 10, .	3.9	14
35	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
36	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

#	Article	IF	CITATIONS
37	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
38	The Evolving Roles of Human iPSC-Derived Cardiomyocytes in Drug Safety and Discovery. Cell Stem Cell, 2017, 21, 14-17.	11.1	69
39	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
40	Resourcing Drug Development Commensurate With its PublicÂHealthÂImportance. JACC Basic To Translational Science, 2016, 1, 309-312.	4.1	4
41	Evolving regulatory paradigm for proarrhythmic risk assessment for new drugs. Journal of Electrocardiology, 2016, 49, 837-842.	0.9	24
42	Exploring New Endpoints for Patients With Heart Failure With Preserved Ejection Fraction. Circulation: Heart Failure, 2016, 9, .	3.9	46
43	The Cardiac Safety Research Consortium enters its second decade: An invitation to participate. American Heart Journal, 2016, 177, 96-101.	2.7	9
44	The Comprehensive in Vitro Proarrhythmia Assay (CiPA) initiative — Update on progress. Journal of Pharmacological and Toxicological Methods, 2016, 81, 15-20.	0.7	335
45	Evolution of strategies to improve preclinical cardiac safety testing. Nature Reviews Drug Discovery, 2016, 15, 457-471.	46.4	323
46	Universal Correction for QT/RR Hysteresis. Drug Safety, 2016, 39, 577-588.	3.2	33
47	A proposal for scientific framework enabling specific population drug dosing recommendations. Journal of Clinical Pharmacology, 2015, 55, 1073-1078.	2.0	39
48	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
49	Novel oral anticoagulants and reversal agents: Considerations for clinical development. American Heart Journal, 2015, 169, 751-757.	2.7	69
50	Early Drug Discovery Prediction of Proarrhythmia Potential and Its Covariates. AAPS Journal, 2015, 17, 1025-1032.	4.4	22
51	Sex differences in drug-induced changes in ventricular repolarization. Journal of Electrocardiology, 2015, 48, 1081-1087.	0.9	8
52	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
53	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
54	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

NORMAN STOCKBRIDGE

#	Article	IF	CITATIONS
55	Cardiovascular Safety Outcome Trials: A meeting report from the Cardiac Safety Research Consortium. American Heart Journal, 2015, 169, 486-495.	2.7	21
56	Cardiovascular Drug Development. Journal of the American College of Cardiology, 2015, 65, 1567-1582.	2.8	168
57	Implications of the IQ-CSRC Prospective Study: Time to Revise ICHÂE14. Drug Safety, 2015, 38, 773-780.	3.2	52
58	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
59	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
60	Lessons Learned From Hundreds of Thorough QT Studies. Therapeutic Innovation and Regulatory Science, 2015, 49, 392-397.	1.6	13
61	Personalized Cardiovascular Medicine Today. Circulation, 2015, 132, 1425-1432.	1.6	33
62	The IQ SRC 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
63	Developing Therapies for Heart Failure WithÂPreservedÂEjection Fraction. JACC: Heart Failure, 2014, 2, 97-112.	4.1	267
64	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
65	Dealing with Clobal Safety Issues. Drug Safety, 2013, 36, 167-182.	3.2	134
66	Thorough QT Studies and Indirect Causes of QTc Changes. PACE - Pacing and Clinical Electrophysiology, 2012, 35, 1411-1412.	1.2	3
67	Implications of geographical variation on clinical outcomes of cardiovascular trials. American Heart Journal, 2012, 164, 303-312.	2.7	44
68	Practice and challenges of thorough QT studies. Journal of Electrocardiology, 2012, 45, 582-587.	0.9	22
69	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
70	Assessing proarrhythmic potential of drugs when optimal studies are infeasible. American Heart Journal, 2009, 157, 827-836.e1.	2.7	81
71	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
72	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