List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2752955/publications.pdf

Version: 2024-02-01

		1040056	1199594
14	281	9	12
papers	citations	h-index	g-index
15	15	15	185
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Sample size determination in clinical trials with multiple coâ€primary binary endpoints. Statistics in Medicine, 2010, 29, 2169-2179.	1.6	56
2	Sample Size Determination in Superiority Clinical Trials with Multiple Co-Primary Correlated Endpoints. Journal of Biopharmaceutical Statistics, 2011, 21, 650-668.	0.8	33
3	A convenient formula for sample size calculations in clinical trials with multiple coâ€primary continuous endpoints. Pharmaceutical Statistics, 2012, 11, 118-128.	1.3	32
4	Sample size determination in clinical trials with multiple coâ€primary endpoints including mixed continuous and binary variables. Biometrical Journal, 2012, 54, 716-729.	1.0	27
5	Group-Sequential Strategies in Clinical Trials with Multiple Co-Primary Outcomes. Statistics in Biopharmaceutical Research, 2015, 7, 36-54.	0.8	27
6	Sample size determination for clinical trials with coâ€primary outcomes: exponential event times. Pharmaceutical Statistics, 2013, 12, 28-34.	1.3	25
7	A logrank test-based method for sizing clinical trials with two co-primary time-to-event endpoints. Biostatistics, 2013, 14, 409-421.	1.5	24
8	Sample size determination in groupâ€sequential clinical trials with two coâ€primary endpoints. Statistics in Medicine, 2014, 33, 2897-2913.	1.6	24
9	Sizing clinical trials when comparing bivariate timeâ€toâ€event outcomes. Statistics in Medicine, 2017, 36, 1363-1382.	1.6	12
10	Sample Size Considerations in Clinical Trials When Comparing Two Interventions Using Multiple Co-Primary Binary Relative Risk Contrasts. Statistics in Biopharmaceutical Research, 2015, 7, 81-94.	0.8	10
11	Group-Sequential Designs When Considering Two Binary Outcomes as Co-Primary Endpoints. ICSA Book Series in Statistics, 2015, , 235-262.	0.2	7
12	Group-sequential logrank methods for trial designs using bivariate non-competing event-time outcomes. Lifetime Data Analysis, 2020, 26, 266-291.	0.9	4
13	Statistical Issues in Clinical Trials with Multiple Primary Endpoints. Japanese Journal of Biometrics, 2013, 34, 35-52.	0.0	0
14	Convenient Sample Size Formula. SpringerBriefs in Statistics, 2015, , 41-58.	0.4	0