Christopher D Breder

List of Publications by Year
in descending order

10	63	1936888 4	1588620 8
papers	citations	h-index	g-index

10	10	10	81
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Whatâ€™s the Regulatory Value of a Target Product Profile?. Trends in Biotechnology, 2017, 35, 576-579.	4.9	19
2	Exposureâ€“ and Doseâ€“response Analyses in Dose Selection and Labeling of FDA-approved Biologics. Clinical Therapeutics, 2018, 40, 95-102.e2.	1.1	14
3	Patterns of use and impact of standardised MedDRA query analyses on the safety evaluation and review of new drug and biologics license applications. PLoS ONE, 2017, 12, e0178104.	1.1	11
4	A Critical Evaluation of Safety Signal Analysis Using Algorithmic Standardised MedDRA Queries. Drug Safety, 2018, 41, 1375-1385.	1.4	8
5	Clinical Development of Biologics Approved by the US Food and Drug Administration, 2003-2016. Therapeutic Innovation and Regulatory Science, 2019, 53, 752-758.	0.8	5
6	Identification of factors associated with first-cycle drug approval rates and regulatory outcomes for new drug applications. Pharmacological Research, 2019, 139, 166-172.	3.1	3
7	A Method for Retrieval of Adverse Event Terms in Clinical Trial Databases Using Standardised MedDRA Queries. Pharmaceutical Medicine, 2016, 30, 103-108.	1.0	2
8	Investigation of Factors Associated With Immunogenicity Labeling Updates and Characteristics of Biologics License Applications. Clinical Pharmacology and Therapeutics, 2021, 110, 1381-1388.	2.3	1
9	Correlates of time to approval and other clinical development periods. Drug Discovery Today, 2019, 24, 1871-1876.	3.2	0
10	A Multifaceted Perspective of Pharmaceutical Innovation: A Consideration of the Regulatory Role. Therapeutic Innovation and Regulatory Science, 2021, 55, 262-269.	0.8	0