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A phase I trial of PRN1008, a novel reversible covalent inhibitor of Bruton's tyrosine kinase, in healthy volunteers

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British Journal of Clinical Pharmacology, 2017, 83, 2367-2376.

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#	Paper	IF	Citations
70	A phase I trial of PRN1008, a novel reversible covalent inhibitor of Bruton's tyrosine kinase, in healthy volunteers. <i>British Journal of Clinical Pharmacology</i> , <b>2017</b> , 83, 2367-2376	3.8	55
69	Are BTK and PLCG2 mutations necessary and sufficient for ibrutinib resistance in chronic lymphocytic leukemia?. <i>Expert Review of Hematology</i> , <b>2018</b> , 11, 185-194	2.8	38
68	The development of Bruton's tyrosine kinase (BTK) inhibitors from 2012 to 2017: A mini-review. <i>European Journal of Medicinal Chemistry</i> , <b>2018</b> , 151, 315-326	6.8	84
67	Perspective From the 5th International Pemphigus and Pemphigoid Foundation Scientific Conference. <i>Frontiers in Medicine</i> , <b>2018</b> , 5, 306	4.9	15
66	Versatile Bioconjugation Chemistries of ortho-Boronyl Aryl Ketones and Aldehydes. <i>Accounts of Chemical Research</i> , <b>2018</b> , 51, 2198-2206	24.3	36
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57	Bruton's tyrosine kinase (BTK) inhibitors in treating cancer: a patent review (2010-2018). <i>Expert Opinion on Therapeutic Patents</i> , <b>2019</b> , 29, 217-241	6.8	29
56	Discovery of Branebrutinib (BMS-986195): A Strategy for Identifying a Highly Potent and Selective Covalent Inhibitor Providing Rapid in Vivo Inactivation of Bruton's Tyrosine Kinase (BTK). <i>Journal of Medicinal Chemistry</i> , <b>2019</b> , 62, 3228-3250	8.3	47
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