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Similar efficacy, safety and immunogenicity of adalimumab biosimilar BI 695501 and Humira reference product in patients with moderately to severely active rheumatoid arthritis: results from the phase III randomised VOLTAIRE-RA equivalence study

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#	Paper	IF	Citations
77	Biosimilar Drugs for Psoriasis: Principles, Present, and Near Future. <i>Dermatology and Therapy</i> , 2018 , 8, 173-194	4	32
76	The biosimilars journey: current status and ongoing challenges. <i>Drugs in Context</i> , 2018 , 7, 212543	5.2	8
75	Low Percentage of Signal Regulatory Protein α Memory B Cells in Blood Predicts Development of Anti-drug Antibodies (ADA) in Adalimumab-Treated Rheumatoid Arthritis Patients. <i>Frontiers in Immunology</i> , 2018 , 9, 2865	8.4	6
74	Similar Pharmacokinetics of the Adalimumab (Humira) Biosimilar BI 695501 Whether Administered via Subcutaneous Autoinjector or Prefilled Syringe (VOLTAIRE-AI and VOLTAIRE-TAI): Phase 1, Randomized, Open-Label, Parallel-Group Trials. <i>Rheumatology and Therapy</i> , 2018 , 5, 403-421	4.4	11
73	Research Techniques Made Simple: Sample Size Estimation and Power Calculation. <i>Journal of Investigative Dermatology</i> , 2018 , 138, 1678-1682	4.3	20
72	Review of Biosimilar Trials and Data on Adalimumab in Rheumatoid Arthritis. <i>Current Rheumatology Reports</i> , 2018 , 20, 57	4.9	28
71	Non-medical Switching from Originator Tumor Necrosis Factor Inhibitors to Their Biosimilars: Systematic Review of Randomized Controlled Trials and Real-World Studies. <i>Advances in Therapy</i> , 2018 , 35, 1295-1332	4.1	15
70	Long-term safety, efficacy, and immunogenicity of adalimumab biosimilar BI 695501 and adalimumab reference product in patients with moderately-to-severely active rheumatoid arthritis: results from a phase 3b extension study (VOLTAIRE-RAext). <i>Expert Opinion on Biological Therapy</i> , 2019 , 19, 1097-1105	5.4	9
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64	Switching to biosimilars: current perspectives in immune-mediated inflammatory diseases. <i>Expert Opinion on Biological Therapy</i> , 2019 , 19, 1001-1014	5.4	12
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62	Use of biosimilars in inflammatory bowel disease: a position update of the Italian Group for the Study of Inflammatory Bowel Disease (IG-IBD). <i>Digestive and Liver Disease</i> , 2019 , 51, 632-639	3.3	22
61	FKB327, an adalimumab biosimilar, versus the reference product: results of a randomized, Phase III, double-blind study, and its open-label extension. <i>Arthritis Research and Therapy</i> , 2019 , 21, 281	5.7	19

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59	Drug Discontinuation in Studies Including a Switch From an Originator to a Biosimilar Monoclonal Antibody: A Systematic Literature Review. <i>Clinical Therapeutics</i> , 2019 , 41, 155-173.e13	3.5	16
58	Biosimilars of adalimumab: the upcoming challenge in IBD. <i>Expert Opinion on Biological Therapy</i> , 2019 , 19, 1023-1030	5.4	13
57	The US Biosimilar Market: Stunted Growth and Possible Reforms. <i>Clinical Pharmacology and Therapeutics</i> , 2019 , 105, 92-100	6.1	33
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55	Clinical trial and 'real-world' data support switching from a bio-originator to its biosimilar. <i>Annals of the Rheumatic Diseases</i> , 2020 , 79, e44	2.4	3
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39	Similar efficacy, safety, and immunogenicity of the biosimilar BI 695501 and adalimumab reference product in patients with moderate-to-severe chronic plaque psoriasis: results from the randomized Phase III VOLTAIRE-PSO study. <i>Expert Opinion on Biological Therapy</i> , 2021 , 21, 87-96	5.4	2
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32	Efficacy and safety of biosimilar CT-P17 versus reference adalimumab in subjects with rheumatoid arthritis: 24-week results from a randomized study. <i>Arthritis Research and Therapy</i> , 2021 , 23, 51	5.7	11
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4	The economic burden of biologic disease-modifying antirheumatic drugs in rheumatoid arthritis patients in the United States.	0
3	Korean clinical practice guidelines on biologics for moderate to severe Crohn's disease.	0
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