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Effectiveness and safety of switching from **originator** to **biosimilar** adalimumab in patients with **psoriasis**.
Nielsen VW, Lund TT, Gniadecki R, Olsen RH, Nissen CV, Schwensen JFB, Thomsen SF.
Dermatol Ther. 2020 Nov;33(6):e14258. doi: 10.1111/dth.14258. Epub 2020 Sep 11.
PMID: 32865289 No abstract available.

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Biosimilar Drugs for Psoriasis: Principles, Present, and Near Future.
Carrascosa JM, Jacobs I, Petersel D, Strohal R.
Dermatol Ther (Heidelb). 2018 Jun;8(2):173-194. doi: 10.1007/s13555-018-0230-9. Epub 2018 Mar 16.
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Currently, several anti-TNF **biosimilars** are available for use in patients with **psoriasis**, and other monoclonal antibodies are in development. This review provides dermatologists and those who treat and/or manage **psoriasis** with a working knowledge of the scien ...

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Effectiveness and safety of switching from **originator** to **biosimilar** adalimumab in patients with **psoriasis**.

Nielsen VW, Lund TT, Gniadecki R, Olsen RH, Nissen CV, Schwensen JFB, Thomsen SF. Dermatol Ther. 2020 Nov;33(6):e14258. doi: 10.1111/dth.14258. Epub 2020 Sep 11. PMID: 32865289 No abstract available.

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A single-centre, observational, retrospective, real-life study evaluating adalimumab **biosimilar** ABP 501 in the treatment of plaque-type **psoriasis** and psoriatic arthritis in **originator-naive** patients and in patients undergoing non-medical switch from **originator**.

Giunta A, Zangrilli A, Bavetta M, Manfreda V, Pensa C, Bianchi L. Curr Med Res Opin. 2021 Jul;37(7):1099-1102. doi: 10.1080/03007995.2021.1923467. Epub 2021 May 20.

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83

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RESULTS BY YEAR



1 Cite



Real-world Drug Survival of **Biosimilar SB5** vs GP2017 Following a Mandatory Non-medical Switch from Adalimumab **Originator** for **Psoriasis**: A Nationwide Cohort Study.

Sieborg J, Maul JT, Wu JJ, Loft NN, Skov L, Bryld LE, Rasmussen MK, Dam TN, Bertelsen T, Ajgeiy KK, Thomsen SF, Thyssen JP, Egeberg A, Thein D.

Acta Derm Venereol. 2025 Mar 3;105:adv42572. doi: 10.2340/actadv.v105.42572.

PMID: 40026110 [Free PMC article.](#)

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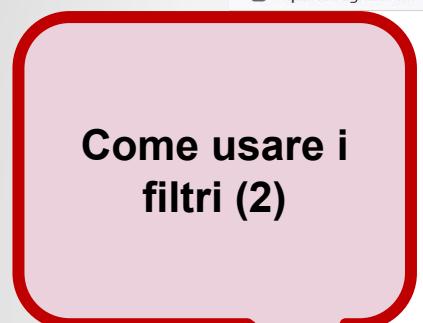
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10

Biosimilars are designed to be analogous to the biologic **originators**. However, comprehensive comparisons between **biosimilars** are limited and lacking for patients with **psoriasis**. This study's objective was to compare 1-year drug survival of the 2 adalim ...



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Books and Documents
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 Meta-Analysis
 Randomized Controlled Trial
 Review

2 Cite

CMAJ OPEN

Mandatory nonmedical switching from **originator** to **biosimilar** infliximab in patients with inflammatory arthritis and **psoriasis** in British Columbia: a cohort study.
Fisher A, Kim JD, Dormuth CR.
CMAJ Open. 2022 Feb 15;10(1):E109-E118. doi: 10.9778/cmajo.20200319. Print 2022 Jan-Mar.
PMID: 35168933 Free PMC article.

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BACKGROUND: In 2019, British Columbia's public drug plan, PharmaCare, was the first in Canada to implement a nonmedical switching policy from **originator** infliximab to its **biosimilar**, for patients with inflammatory arthritis or **psoriasis**. ...METHODS: We constr ...

3 Cite

BioDrugs

Cross-Switch from Etanercept **Originator** to **Biosimilar** SB4 and to GP2015 in Patients with Chronic Plaque **Psoriasis**.
Piaserico S, Conti A, Messina F, Meneguzzo A, Odorici G, Bellinato F, Gisondi P.
BioDrugs. 2021 Jul;35(4):469-471. doi: 10.1007/s40259-021-00485-8. Epub 2021 May 11.
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Book

Biosimilars Use in Medicine for Inflammatory Diseases

Inderbir S. Padda ¹, Rajat Bhatt ², Mayur Parmar ³

In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2025 Jan. 2023 Aug 28.

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Biosimilars Use in Medicine for Inflammatory Diseases

Inderbir S. Padda; Rajat Bhatt; Mayur Parmar.
Author Information and Affiliations
Last Update: August 28, 2023.

Continuing Education Activity Go to:

Biosimilars is FDA approved for the treatment of rheumatoid arthritis (RA), polyarticular juvenile idiopathic arthritis (JIA), psoriatic arthritis (PA), ankylosing spondylitis (AS), plaque psoriasis (PsO), inflammatory bowel disease (IBD): adult Crohn disease (CD), ulcerative colitis (UC), granulomatosis with polyangiitis (GPA) (Wegener granulomatosis), microscopic polyangiitis (MPA), type 1 diabetes mellitus (DM), non-squamous non-small cell lung cancer, metastatic colorectal cancer, metastatic renal cell carcinoma, glioblastoma, recurrent or metastatic cervical cancer, HER2 (human epidermal growth factor receptor 2) associated breast cancer, and HER2 associated gastric (metastatic) or gastroesophageal junction adenocarcinoma, chronic lymphocytic leukemia (CLL), non-Hodgkin lymphoma (NHL). Biosimilars are biotherapeutic agents similar to their respective original biologic product (reference product). This activity will highlight the mechanism of action, adverse event profile, and other key factors pertinent to interprofessional team members in managing patients with inflammatory diseases and the use of biosimilars.

Objectives:

- Identify the mechanism of action of all biosimilars.

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FDA's Approach to Regulating Biosimilars. [Clin Cancer Res. 2017]
Application of the FDA Biosimilar Extrapolation Framework to Make Off-Label Determinations [J Manag Care Spec Pharm. 2017]
Review The First Interchangeable Biosimilar Insulin: Insulin Glargine-yfgn. [J Diabetes Sci Technol. 2023]
The US approach to biosimilars: the long-awaited FDA approval pathway. [BioDrugs. 2012]
Review The totality of evidence approach in the development of AVT02 (adalimumab), a biosimilar to [Ther Adv Chronic Dis. 2024]



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- 1 There has been significant interest in this topic as a number of biosimilars will be available in 2023. OBJECTIVE: This review aims to evaluate the safety and efficacy of biosimilars used in dermatology and the experience with biosimilars used in dermatology.
- 2 A review of biosimilars in psoriasis: impact on access, and a first-hand look at biosimilar costs from the perspective of the department of veterans affairs.
Reese R, Nanavath SR, Martin J, Travers JB, Rohan CA.
J Dermatolog Treat. 2024 Dec;35(1):2402912. doi: 10.1080/09546854.2024.2402912. Epub 2024 Sep 15.
PMID: 39278830
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- 3 However, access to these drugs has been limited due to their high cost. Biosimilars utilize reverse engineering to create a highly similar product to an originator drug following patent expiration and provide an avenue to reduce costs of biologic treatment. ...Addit ...
TNFalpha inhibitor biosimilar associated with polychondritis. A case-based review.
Venetsanopoulou AI, Voulgari PV, Drosos AA.
Rheumatol Int. 2023 Sep;43(9):1751-1754. doi: 10.1007/s00296-023-05364-9. Epub 2023 Jun 15.
PMID: 37316633
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- 4 This report presents a 43-year-old man with psoriatic arthritis treated with ABP-501 (Amgevita), an adalimumab (ADA) biosimilar and who developed RP, 8 months after the initiation of the treatment. This is the first report of RP development during TNFalpha inhibitors b ...
Biosimilars for the Treatment of Psoriasis: A Systematic Review of Clinical Trials and Observational Studies

La review è un tipo di articolo in cui gli autori esaminano in modo critico, e possibilmente esaustivo, gli studi pubblicati su un determinato argomento.



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1. Review (Revisione narrativa):

- È una sintesi generale della letteratura su un argomento.
- Non segue necessariamente un metodo sistematico o rigoroso.
- Può essere soggettiva nella selezione e interpretazione degli studi.

2. Systematic Review (Revisione sistematica):

- Segue un protocollo rigoroso e predefinito.
- Ha criteri di inclusione ed esclusione chiari per gli studi.
- Mira a identificare, valutare e sintetizzare tutti gli studi rilevanti su una domanda specifica.
- È più oggettiva e riproducibile rispetto a una review narrativa.

3. Metanalisi:

- È un tipo di systematic review che include un'analisi statistica.
- Combina i risultati di multipli studi per ottenere una stima quantitativa dell'effetto complessivo.
- Fornisce una sintesi statistica dei risultati degli studi inclusi.

In sintesi, una systematic review è più rigorosa di una review narrativa, mentre una metanalisi aggiunge un'analisi statistica a una systematic review.



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10 results Page 1 of 1

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Biosimilars in Dermatology Review.
Baker J, Kalb R.
J Psoriasis Psoriatic Arthritis. 2024 Jan;9(1):28-35. doi: 10.1177/24755303231212154.
Epub 2023 Nov 3.
PMID: 39301301

Review.

There has been significant interest in this topic as a number of **biosimilar** versions of adalimumab will be available in 2023. OBJECTIVE: This review will discuss **biosimilar** basics and the experience with **biosimilars** used in dermatology in the US, Asia, and Eu ...

A review of biosimilars in psoriasis: impacts on efficacy, safety, access, and a first-hand look at biosimilar cost savings within the department of veterans affairs.
Reese R, Nanavath SR, Martin J, Travers JB, Rohan CA.
J Dermatolog Treat. 2024 Dec;35(1):2402912. doi: 10.1080/09546634.2024.2402912. Epub 2024 Sep 15.
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However, access to these drugs has been limited due to their high cost. **Biosimilars** utilize

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Review > Drug Des Devel Ther. 2021 Jul 8:15:2987-2998. doi: 10.2147/DDDT.S317382.
eCollection 2021.

An Update Review of Biosimilars of Adalimumab in Psoriasis – Bioequivalence and Interchangeability

Xin Zhou ¹, Zhuo Chen ², Xinling Bi ¹

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PMID: 34267501 PMCID: PMC8275172 DOI: 10.2147/DDOT.S317382

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Abstract

Biologic drugs have revolutionized the treatment of psoriasis and other rheumatological diseases. In recent years, many biosimilar agents that are highly similar in structure and function to their originator products have been developed, including the tumor necrosis factor-alpha antagonist adalimumab. The considerably lower cost of these products has greatly cut the economic burden of the patients and increased the accessibility of biologic therapies worldwide. The US Food and Drug Administration and/or the European Medicines Agency have approved eight biosimilars of adalimumab (ABP 501/BI 695501/SBS/GP2017/FKB327/MSB11022/PF-06410293/CT-P17) for the treatment of psoriasis, and others are under review. Given that these agents showed pharmacokinetic, efficacy, safety, and immunogenicity profiles comparable to those of the originator, adalimumab biosimilars were licensed for all indications approved for reference adalimumab based on extrapolation; however, some of the equivalence studies were only conducted in one or two disease populations. This review discusses the bioequivalence of adalimumab biosimilars as demonstrated by various clinical trials, the extrapolation of indications, guidance and policies of the EU and US on interchangeability (nonmedical switching/automatic substitution) between biosimilars and originators, and the real-life practices of switching from reference adalimumab to the respective biosimilars. Further data from real-world studies and post-marketing analyses are needed better to address the efficacy and safety of the transition strategy.

Keywords: adalimumab; biosimilar; extrapolation of indications; guidance of interchangeability; psoriasis.

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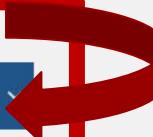
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2 Cite Health Benefits of Prebiotics, Probiotics, Synbiotics, and Postbiotics. Al-Habsi N, Al-Khalili M, Haque SA, Elias M, Olqi NA, Al Uraimi T. Nutrients. 2024 Nov 19;16(22):3955. doi: 10.3390/nu16223955. PMID: 39599742 Free PMC article. Review.

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[Arthritis, Psoriatic](#)

2. A type of inflammatory arthritis associated with **PSORIASIS**, often involving the axial joints and the peripheral terminal interphalangeal joints. It is characterized by the presence of HLA-B27-associated SPONDYLARTHROPATHY, and the absence of rheumatoid factor. Year introduced: 1990

[Tranebjærg Svegaard syndrome \[Supplementary Concept\]](#)

3. Date introduced: August 25, 2010

[PSORS1C1_protein,_human \[Supplementary Concept\]](#)

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Liu L, Li M, Hu M, Ge Z, Dong X.
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Ciprofloxacin and levofloxacin attenuate microglia inflammatory response via TLR4/NF- κ B pathway

Morena Zusso ¹, Valentina Lunardi ¹, Davide Franceschini ^{1,2}, Andrea Pagotto ¹, Rita Lo ³, Stefano Stifani ³, Anna Chiara Frigo ⁴, Pietro Giusti ⁵, Stefano Moro ¹

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