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Observational Study
Real-world data on the infliximab biosimilar CT-P13 (Remsima®) in inflammatory bowel disease

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Abstract
BACKGROUND
CT-P13 (Inflectra™, Remsima™) monitoring in patients with inflammatory bowel disease (IBD) is a promising step to reduce treatment costs. Since monitoring of Remicade™ serum trough levels and anti-Remicade™ immunogenicity hold an important significance in treatment modalities, no data about monitoring of drug serum trough ...

Cited by: 7  
Author: Kornelius Schulze, Nadine Koppka, F...

Switching from Remicade® to Remsima® is well Tolerated ...

Abstract. Background and aims: A biosimilar version of infliximab [CT-P13/Remsima®] recently entered the European market. The clinical data on its use in inflammatory bowel disease [IBD] are sparse, especially on switching from the originator Remicade®. In this study, we aimed to prospectively investigate the feasibility, safety and immunogenicity of switching from Remicade to Remsima in a real-life ...

Cited by: 97  
Author: Lydia C. T. Buer, Lydia C. T. Buer, Bjö...

Clinical experience with infliximab biosimilar Remsima (CT ...

Based on current data, CT-P13 seems to be efficacious and generally well tolerated in IBD especially in patients who are naïve to biological therapy. Knowledge with regard to interchangeability between CT-P13 and the originator infliximab is however, still rather sparse and more data are desired.

Cited by: 21  
Author: Jørgen Jahnsen, Jørgen Jahnsen

Post-marketing analysis for biosimilar CT-P13 in ...
Infliximab biosimilar CT-P13 is interchangeable with its ...

Background/aims: An interim analysis of post-marketing surveillance of CT-P13, an infliximab biosimilar, was performed to evaluate its safety and efficacy in Japanese patients with inflammatory bowel disease. Methods: Patients were prospectively enrolled between November 2014 and March 2017, after the launch of CT-P13 in Japan, and case report forms of patients followed for at least 4 ...
**Infliximab biosimilar CT-P13 is interchangeable with its ...**


Background/aims: An interim analysis of post-marketing surveillance of CT-P13, an infliximab biosimilar, was performed to evaluate its safety and efficacy in Japanese patients with inflammatory bowel disease.

Methods: Patients were prospectively enrolled between November 2014 and March 2017, after the launch of CT-P13 in Japan, and case report forms of patients followed for at least 4 months ...

**CT-P13 (Inflectra™, Remsima™) monitoring in patients with ...**


The approval of infliximab biosimilars Remsima™ and Inflectra™ (CT-P13) for patients with inflammatory bowel disease (IBD) is a promising step to reduce treatment costs. Since monitoring of Remicade™ serum trough levels and anti-Remicade™ immunogenicity hold an important significance in treatment modalities, no data about monitoring of drug serum trough levels or anti-drug antibody ...

**Clinical experience with infliximab biosimilar Remsima (CT ...**


Based on current data, CT-P13 seems to be efficacious and generally well tolerated in IBD especially in patients who are naïve to biological therapy. Knowledge with regard to interchangeability between CT-P13 and the originator infliximab is however, still rather sparse and more data are desired.

**Switching from infliximab to biosimilar in inflammatory ...**

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Apr 15, 2019 · In 2017, the extension study of PLANETRA and PLANETAS showed that switching from RP to its biosimilar CT-P13 is possible without negative effects on safety or efficacy in patients with AS. 65,66 Furthermore, switching from the IFX RP to CT-P13 after 1 year of IFX RP treatment showed continued comparable efficacy, immunogenicity and safety, to maintenance of CT-P13 treatment during the second year of the treatment. 66 In PLANETRA ...