

# Zent2U announces completion of bioequivalence trials on Apremilast

Zent2U is delighted to announce the successful completion of the pivotal bioequivalence trial that was conducted under fasting conditions. Our trial was carried out on **Apremilast 30mg film-coated tablets** developed by Zentiva and study endpoints perfectly matched the necessary bioequivalence parameters. The completion of this bioequivalence study enables us to reach a significant milestone in terms of development.

We will progress into manufacturing registration batches and compilation of EU dossiers which will be available for review in October 2021. This target date is giving sufficient time for registration filing on the data exclusivity expiry date in January 2023. We believe that having a dossier in place in 2021 grants Zent2U and its partners the great position to register this dossier in all major European markets at the earliest possible time, and therefore launch the product indicated for psoriasis and psoriatic arthritis as early as possible after the expiration of the marketing exclusivity.

Partner up now and  
connect with our team  
today!



**Tomas Pilarcik**

Key Account Manager  
[tomas.pilarcik@zentiva.com](mailto:tomas.pilarcik@zentiva.com)



**Thomas Koene**

Head of Growth  
Partnerships  
[thomas.koene@zentiva.com](mailto:thomas.koene@zentiva.com)

**Disclaimer:** Apremilast which is subject to patent protection is currently not offered or made available in countries where patents are in force.