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 Information: NON-HIV ANTIVIRALS:
 ZOVIRAX: 00168082530: ACYCLOVIR
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Clavulanate 2:1® tablet grade, for Direct
Compression, oral

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comparable quality, safety and efficacy

to existing biologics that introduces
much-needed competition. ... Our
products, which are sold in more than
160 countries, reach more than 500
million patients and our aspiration is to
reach one billion. In the UK we ...

Our Portfolio | Sandoz UK

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Drugs.com**

A biosimilar is a type of biologic product

that is licensed by the FDA because it is highly similar to a biological product that is already FDA-approved. This originally approved product is known as the biological reference product (or reference product), and the biosimilar has been shown to have no clinically meaningful differences in potency, safety, and purity compared with that of its ...

Sandoz Biosimilars | Home

Finished Dosage Forms (Partner Label)
Anti-Infectives. ORAL EUROPE LATIN
AMERICA APMA. Aciclovir•. Amoxicillin
Trihydrate•. Amoxicillin Trihydrate +
Potassium Clavulanate•. Azithromycin
Dihydrate•.

Finished Dosage Forms (Partner

Label), Active ... - Sandoz

The 2017 Sustainability Report is a snapshot of the current situation at Sandoz GmbH. As a manufacturer of medicinal products we work to promote people's wellbeing. We help to ensure that people around the world are and remain healthy. With this in mind, we are committed to high product quality, innovation and the efficient use of resources.

Sustainability Report 2017 - Sandoz

products such as Promacta/Revolade and Tafinlar + Mekinist. Sandoz is expanding access to biosimilars, and we have a leading portfolio with five biosimilars now on the market. However, Sandoz net sales were down 2% cc due to fierce price competition in the US.

Alcon made significant progress on its turnaround, returning to

Novartis Annual Report 2017

At Sandoz, we discover new ways to improve and extend people's lives. We pioneer novel approaches to help people around the world access high-quality medicine. UK/MKT/SDZ/17-0041(2) June 2019

Sandoz UK

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Mylan/Biocon's application for a Neulasta biosimilar was rejected in October 2017. Biocon has stated that it does not expect the rejection to impact the timing of the commercial launch of its product.

Sandoz reported this year that it intends to resubmit its pegfilgrastim biosimilar application to FDA in 2019. FDA rejected Sandoz's ...

US Biosimilar Approvals Soar in 2017 | Biologics Blog

List of nationally authorised medicinal products EMA/178876/2017 Page 4/16
Product Name (in authorisation country)
MRP/DCP Authorisation number National
Authorisation Number MAH of product in
the member state Member State where
product is authorised Mitoxantrona
Sandoz DK/H/0439/001 4996583
SANDOZ FARMACÊUTICA LDA. PT

List of nationally authorised medicinal products

The oral solid dose manufacturing facility

in Broomfield, Colorado, run by Novartis division Sandoz, is to close its doors within the next two years, with 450 jobs affected. Sandoz spokesman Tim Willeford told this publication increased

competition in the generic drug market has caused “above-average pricing pressure” and drove the decision to shutter the site and reduce its US product ...