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Glenmark bags marketing approval for Ryaltris in 13 countries across EU, UK

30 September 2021 | News

Ryaltris will be marketed in the rest of Europe by the Menarini Group as part of its exclusive licensing agreement with Glenmark



Glenmark Pharmaceuticals has received marketing approval for its fixed-dose combination nasal spray Ryaltris in 13 countries across the EU and UK. Glenmark is set to launch Ryaltris directly in the markets of Czech Republic, Slovakia, Poland, and the UK. Ryaltris will be marketed in the rest of Europe by the Menarini Group as part of its exclusive licensing agreement with Glenmark.

Ryaltris (olopatadine 665 ?g and mometasone furoate 25 ?g), is indicated for symptomatic treatment of seasonal and perennial allergic rhinitis in adults and children over 12 years of age.

Commenting on this development, Robert Crockart, Chief Commercial Officer, Glenmark Pharmaceuticals said, “The marketing approval will pave the way for the effective and timely treatment of allergic rhinitis for thousands of patients across Europe. We are already seeing its therapeutic benefits in other regions where Ryaltris has been launched, and we hope to extend this relief to more people across the world.”

Glenmark has also partnered with Hikma Pharmaceuticals and Bausch Health for the commercialisation of Ryaltris in the US and Canada respectively. In April this year, Glenmark concluded the DCP regulatory
procedure in Europe, enabling approval in 17 countries across the EU and the UK.