19 May 2022
EMA/275660/2022
EMEA/H/C/004889

Public statement

Zynquista

Withdrawal of the marketing authorisation in the European Union

On 22 March 2022, the European Commission withdrew the marketing authorisation for Zynquista (sotagliflozin) in the European Union (EU). The withdrawal was at the request of the marketing authorisation holder, Guidehouse Germany GmbH, which notified the European Commission of its decision not to market the product in the EU for commercial reasons.

Zynquista was granted marketing authorisation in the EU on 26 April 2019 for use as an adjunct to insulin therapy to improve glycaemic control in adults with type 1 diabetes mellitus. The marketing authorisation was initially valid for a 5-year period.

The European Public Assessment Report (EPAR) for Zynquista will be updated to indicate that the marketing authorisation is no longer valid.

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union