

04/13/2023

Makena Products No Longer Covered as of April 7, 2023

Background:

On April 6, 2023, the U.S. Food and Drug Administration [announced the final decision](#) to withdraw approval of Makena.

Key Details:

HHSC will remove the following Makena products and their clinical prior authorization as of April 7, 2023.

NDC	Drug Name
64011030103	MAKENA 275 MG/1.1 ML AUTOINJCT
00517176701	HYDROXYPROGEST 250 MG/ML VIAL
55150030901	HYDROXYPROGEST 250 MG/ML VIAL
64011030103	MAKENA 275 MG/1.1 ML AUTOINJCT
67457096701	HYDROXYPROGEST 250 MG/ML VIAL
69238179701	HYDROXYPROGEST 250 MG/ML VIAL
71225010401	HYDROXYPROGEST 1,250 MG/5 ML
71225010501	HYDROXYPROGEST 250 MG/ML VIAL

Contact: vdp-formulary@hhsc.state.tx.us

Type: Informational

To: CHIP; STAR; STAR+PLUS; STARHEALTH; STAR_KIDS

From: VDP