

## Cervarix<sup>®</sup> [human papillomavirus bivalent (types 16 and 18) vaccine, recombinant] – Product Discontinuation

- GlaxoSmithKline announced the discontinuation of Cervarix [human papillomavirus bivalent (types 16 and 18) vaccine, recombinant] in the U.S. due to very low market demand.
  - The discontinuation is not due to product quality, safety, or efficacy concerns.
  - Existing stock of Cervarix can be distributed until the inventory is depleted. GSK's last ship date for Cervarix was August 31, 2016.
- Cervarix is indicated for the prevention of the following diseases caused by oncogenic human papillomavirus (HPV) types 16 and 18: cervical cancer, cervical intraepithelial neoplasia (CIN) grade 2 or worse and adenocarcinoma in situ, and CIN grade 1.
- For further questions, contact the GSK Vaccine Service Center at 1-866-475-8222.
- Other vaccines currently available for the prevention of HPV-related diseases include <u>Gardasil<sup>®</sup> [HPV quadrivalent (types 6, 11, 16 and 18) vaccine, recombinant]</u> and <u>Gardasil<sup>®</sup> 9 [HPV 9-valent vaccine, recombinant]</u>. Refer to the respective drug labels for detailed indication information.

## **Action Plan**

 The discontinuation of Cervarix will be presented at an upcoming Pharmacy & Therapeutics Committee meeting.



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