



SMFM Notice: Bicillin® L-A Recall

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This guidance was developed by the Society for Maternal-Fetal Medicine (SMFM) Committee on Infectious Diseases and Emerging Threats with the assistance of Naima Joseph, MD, MPH and Brenna Hughes, MD, MSc.

This document has been updated as follows:

- Includes SMFM support of the use of Extencilline and Lentocilin® for treatment of patients with syphilis in pregnancy when Bicillin® L-A is unavailable.

Background

On July 10, 2025, King Pharmaceuticals LLC., a subsidiary of Pfizer, issued a notice informing customers about a new voluntary [recall](#) of selected lots of **Bicillin® L-A (Penicillin G Benzathine Injectable Suspension)** distributed from December 11, 2023 through June 24, 2025. Pfizer is initiating this recall due to particulates identified during visual inspection. The lots included are listed below.

Carton NDC	Syringe NDC	Lot Number	Expiration Date YYMMDD	Strength	Configuration/ Count
60793-701-10	60793-701-02	GL2954	270131	1,200,000 units/ 2 mL	10 (2 mL) syringes per carton, 24 cartons per shipping case
		HP6222	270131		
		HP6228	271031		
		HP6232	270930		
		HR9967	270531		
		HJ3235	260930		
		LT5190	270930		
60793-702-10	60793-702-04	GT2598	260930	2,400,000 units/ 4 mL	10 (4 mL) syringes per carton, 24 cartons per shipping case
		GT2599	260930		
		HR9969	270430		
		HK2909	270228		
		HR9984	270831		



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Extencilline (benzathine Benzylpenicillin) and Lentocilin® (Benzathine Benzylpenicillin Tetrahydrate) were approved by the Food & Drug Administration (FDA) for temporary importation due to prior shortages and remain available for those who are affected by this recall and have challenges procuring sufficient Bicillin® L-A to meet their needs.

SMFM supports the use of Extencilline and Lentocilin® for treatment of patients with syphilis in pregnancy when Bicillin® L-A is unavailable.