

July 14, 2025

From: Kansas Department of Health and Environment – Bureau of Disease Control and Prevention

To: Health Care Providers and Local Health Departments

RE: Pfizer Voluntary Recall – Bicillin L-A

Summary

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	270131		
		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		

King Pharmaceuticals LLC., a subsidiary of Pfizer, is voluntarily recalling the above referenced lots of **Bicillin® L-A (Penicillin G Benzathine Injectable Suspension)**. Pfizer is initiating this recall due to particulates identified during visual inspection. Pfizer has completed a Health Hazard Assessment which indicated that the potential risk to patients from use of the impacted product is medium.

Pfizer advises that you check your stock immediately against the table above. If you have any of the affected lots in your inventory, please discontinue use, stop distribution, quarantine the product immediately, and return all the affected products in accordance with the attached recall notification.

Please review the attached information from Pfizer regarding the voluntary recall and next steps to return the recalled lots.

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