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[Home](#) > Product recalled because stoppers for vials contain natural rubber latex

Generic Name:

Sodium chloride injection 0.9%

Trade Name:

No trade names

Company:

Fresenius Kabi USA

Notes:

Fresenius Kabi USA is voluntarily [recalling](#) 163 lots of sodium chloride injection 0.9%, 10 mL fill in a 10-mL vial; and sodium chloride injection 0.9%, 20 mL fill in a 20-mL vial. The product is being recalled because the stoppers contain natural rubber latex.

The product insert states that stoppers for both the 10-mL and the 20-mL vials do not contain natural rubber latex; the tray label for the two vial sizes and the vial label for the 20-mL vial also state that the stoppers do not contain latex.

For the population most at risk, those with a severe allergic reaction to latex, there is probability of an anaphylactic reaction that could result in hospitalization or death.

To date, Fresenius Kabi USA has not received any reports of adverse events related to this recall.

See the [tables](#) for a full list of the affected lots, including lot numbers and expiration dates.

Medication Monitor Categories:

[Alerts and Recalls](#)

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