

UPDATE: Non-Safety Related Voluntary Recall of Specific Lots of Influenza A (H1N1) 2009 Monovalent Vaccine Live, Intranasal

On December 22, 2009 MedImmune announced that it is voluntarily recalling unused doses of 13 **specific lots** of Influenza A (H1N1) 2009 Monovalent Vaccine Live, Intranasal due to a slight decrease in potency. (See [Press Release](#)) **There is no safety concern with the lots that are being recalled and therefore no need to revaccinate anyone who had been immunized with vaccines from these specific lot numbers:**

- . 500754P
- . 500751P
- . 500756P
- . 500757P
- . 500758P
- . 500759P
- . 500760P
- . 500761P
- . 500762P
- . 500763P
- . 500764P
- . 500765P
- . 500776P

Providers who received H1N1 influenza vaccine directly from McKesson:

Providers should receive a recall information packet from MedImmune by Monday, January 4, 2010, COB local time. If you have not received a recall information packet by this date, please call **866-209-9273** to request a recall information packet.

Providers who received H1N1 influenza vaccine directly from your local county health department (i.e., not directly from McKesson):

H1N1 registered providers who received H1N1 influenza vaccine, live intranasal, from your local county health department and still have unused H1N1 influenza vaccine doses from the affected lots, please call **866-209-9273** to request a recall information packet.

Stericycle® is handling the return of affected vaccines for MedImmune, they can be reached at the toll-free number noted. All returns should be sent to: Stericycle®, Attention: Event 2073, 2670 Executive Drive, Suite A, Indianapolis, IN 46241.