Description
The adverse reaction to the MMRV of persistent, inconsolable crying lasting > 3 hours within 48 hours of dose is reported to the Immunization Registry using a Z22/VXU message.

Comments
No Comments

Pre Condition
An adverse reaction to the MMRV of persistent, inconsolable crying lasting > 3 hours within 48 hours of dose is recorded in the EMR.

Post Condition
The adverse reaction has been transmitted to the IIS.

Test Objectives
Transmit Standard Patient Immunization History Report: The EHR or other clinical software system directly or indirectly through an intermediary creates and transmits a report of a patient's immunization history to public health immunization registries. ***Identify Adverse Event:*** The EHR or other clinical software system enables capture of structured data regarding adverse events.

Evaluation Criteria
The VXU/Z22 message passes validation using the NIST Immunization VXU Validation Tool (Z22) (context-free). The content of the message correctly reflects the test data (context-based) in accordance with the Test Data Specification and the Message Content.

Notes for Testers
No Note