The information provided in the following table is from reports included in the Adverse Event Management System (TGA's internal adverse event database) but not included in the Database of Adverse Event Notifications (DAEN). The reports that do not appear in the DAEN may have insufficient information and/or no reasonable temporal association and/or the relationship between the medicine/vaccine and the adverse event appears to not be related as judged by a health professional and/or the report may have been received by the TGA within the last 90 days and/or the report may be a duplicate of a case included in the DAEN.

Case Number	Report Entry Date	Gender	Age	Medicine (Onset Time)	Reaction
433257	14/05/2018	Male	75	Fluzone High Dose (Influenza virus haemagglutinin) - Suspect (-); Minipress (Prazosin hydrochloride) - Concomitant (-); Trade Name Not Specified (atorvastatin) - Concomitant (-)	Death
433774	18/05/2018	Female	76	Fluzone High Dose (Influenza virus haemagglutinin) - Suspect (0 days)	Death
434588	30/05/2018	Female	76	Fluzone High Dose (Influenza virus haemagglutinin) - Suspect (0 days)	Pulmonary oedema
470313	4/07/2019	Male	34	FluQuadri Vaccine (Influenza virus haemagglutinin) - Suspect (-); Fluzone High Dose (Influenza virus haemagglutinin) - Suspect (-); Valaciclovir (valaciclovir hydrochloride) - Concomitant (-)	Pyrexia