

## **Patient Vaccination**

OMB No. 0920-0666 Exp. Date: 09-30-2012

Safety Network		* required for saving			
Page 1 of 2	·	^ conditionally required			
*Facility ID:	*Event #:				
*Patient ID:	Social Security #:				
Secondary ID:	First.	Middle.			
Patient Name, Last:	First:	Middle:			
*Gender: F M Other		*Date of Birth:			
Ethnicity (specify):		Race (specify):			
*Event Type: FLUVAX					
*Influenza subtype: ☐ Seasonal ☐ Non-Seasonal		*Date Admitted to Facility:			
*Vaccine offered: ☐ Yes ☐ No		*Vaccine declined: ☐ Yes ☐ No			
Reason(s) vaccine declined (Check either sec		not both)			
		B. Personal reason(s) for declining (check all that apply):			
☐ Allergy to vaccine components		☐ Fear of needles/injections			
☐ History of Guillian-Barre syndrome within 6 weeks of		☐ Fear of side effects			
previous influenza vaccination		☐ Perceived ineffectiveness of vaccine			
☐ Current febrile illness (Temp > 101.5°F)		Religious or philosophical objections			
Other (specify):		☐ Concern for transmitting vaccine virus to contacts			
		☐ Other (specify):			
		a other (specify).			
*Vaccine administered: ☐ Yes ☐ No					
^Date Vaccine Administered:					
^Type of influenza vaccine administered:					
Seasonal: ☐ Afluria® ☐ Agriflu® ☐ Fluarix® ☐ FluLaval® ☐ Flumist®					
☐ Fluvirin® ☐ Fluzone® ☐	I Fluzone High-	Dose®			
Non-constant C Other (constitution					
Non-seasonal:   Other (specify)					
☐ Live attenuated influenza vaccine (LAIV) e.g., nasal ☐ Inactivated vaccine (TIV)					
^Manufacturer:	_ ^Lot nu	mber:			
^Route of administration:   Intramuscular   Intranasal   Subcutaneous					
Vaccine Information Statement (VIS) Provided to Patient:					
☐ Live Attenuated Influenza VIS ☐ Ir	nactivated Influ	enza VIS			
Edition Date:/					
Person Administering Vaccine:					
Vaccinator ID:	Title:				
Name: Last: First:		Middle:			
Work Address:					
City: State:		Zip code:			
	ot otherwise be disclosed o	would permit identification of any individual or institution is collected with a guarantee that it will released without the consent of the individual, or the institution in accordance with Sections 304,			
		ncluding the time for reviewing instructions, searching existing data sources, gathering and			

Public reporting burden of this collection of information is estimated to average 10 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC, Reports Clearance Officer, 1600 Clifton Rd., MS D-74, Atlanta, GA 30333, ATTN: PRA (0920-0666).



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Page 2 of 2

Custom Fields					
Label		Label			
Comments					