

Tables of Instructions

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Table 1. Instructions for Completion of the Healthcare Personnel Safety Monthly Reporting Plan Form (CDC 57.203)

This form collects data on which options and which months a facility intends to participate in NHSN Healthcare Personnel Safety (HPS) Component. This form should be completed for every month that the facility will participate in the HPS Component.

Data Field	Instructions for Data Collection
Facility ID #	Required. The NHSN-assigned facility ID will be auto-
	entered by the application.
Month/Year	Required. Enter the month and year for the surveillance
	plan being recorded.
No NHSN Healthcare Personnel Safety	Conditionally required. Check this box if you do <u>not</u> plan
Modules Followed this Month	to follow any of the NHSN Healthcare Personnel Safety
	Modules during the month and year selected.
Healthcare Pe	ersonnel Exposure Module
Blood/Body Fluid Exposure Only	Conditionally required. Check this box if you plan to
Blood/Body I fuld Exposure Only	follow blood/body fluid exposures only, without
	following exposure management during the month and
	year selected.
Blood/Body Fluid Exposure with Exposure	Conditionally required. Check this box if you plan to
Management Management	follow blood/body fluid exposure with exposure
Management	management during the month and year selected.
Influenza Exposure Management	Conditionally required. Check this box if you plan to
Influenza Exposure Management	follow influenza exposure management (i.e., antiviral
	chemoprophylaxis and/or treatment)
Healthcare Per	sonnel Vaccination Module
Influenza Vaccination Summary	Conditionally required. Check this box if you plan to
Influenza Vaccination Summary	follow the influenza vaccination summary option. Once
	the influenza vaccination summary is selected on the
	reporting plan, it is automatically updated with this
	information for the entire NHSN-defined influenza
	season (July 1 to June 30).



Table 2. Instructions for Completion of the Healthcare Worker Demographic Data Form (CDC 57.204)

This form must be completed for all HCP who have information recorded in HPS component of NHSN (e.g., exposure to blood or body fluid or influenza vaccination.) Alternatively, data for all or selected personnel can be imported from the facility's personnel database at facility enrollment.

Data Field	Instructions for Data Collection		
Facility ID#	Required. The NHSN-assigned facility ID will be auto-entered by the application.		
HCW ID #	Required. Enter the healthcare worker's (HCW) alphanumeric identification		
	number. This identifier is unique to the healthcare facility.		
Social Security #	Optional. Enter the HCW's Social Security Number.		
Secondary ID #	Optional. Enter the HCW's secondary ID number. This could be the employee's		
	medical record # or some other unique identifier.		
HCW Name:	Optional. Enter demographic information for the HCW.		
Last, First, Middle			
Street Address			
City			
State			
Zip Code			
Home Phone			
E-mail Address			
Gender	Required. Indicate the gender of the HCW by checking F (Female) or M (Male).		
Date of birth	Required. Enter the date of birth of the HCW using the format: mm/dd/yyyy.		
Born in the U.S.?	Optional. Select Yes, No, or Unknown.		
Ethnicity	Optional. Select one ethnicity of the HCW.		
Race	Optional. Select the race of the HCW. Check all that apply.		
Work Phone	Optional. Enter the work phone number of the HCW.		
Start Date	Required. Enter the date the HCW began employment or affiliation with the facility		
	(use format: mm/dd/yyyy).		
Work Status	Required. Select Active, Inactive, or No longer affiliated.		
Type of Employment	Required. Select from Full-time, Part-time, Contract, Volunteer, Other (please		
	specify).		
Work Location	Required. Select the code that best describes the HCW's current permanent work		
	location. This refers to physical work location rather than to department		
	assignment. For example, a radiology technician who spends most of his/her time		
	performing portable x-rays throughout the facility works at multiple locations. In		
	general, most interns/residents are not considered to work at a single location		
	because they rotate every month or every few months. For HCP who do not work		
	at least 75% of the time at a single location, the work location code for 'float'		
	should be entered. Location codes must be customized to the facility and set up prior to entering HCW records. The work location must be mapped to a CDC		
	Location (http://www.cdc.gov/nhsn/PDFs/master-locations-descriptions.pdf).		
Department	Optional. Enter the department in which the HCW works (facility defined).		
Department	Optional. Enter the department in which the fiew works (facility defined).		



Data Field	Instructions for Data Collection	
Supervisor	Optional. Enter the name of the HCW's supervisor (facility defined).	
Occupation	Required. Select the occupation code that most appropriately describes the HCW's	
	job. These must be customized to the facility and set up prior to entering HCW	
	records. The occupation must be mapped to a CDC Occupation Code.	
Title	Conditionally required. Required only for HCP designated as Influenza	
	Vaccinators if the facility intends on using NHSN to fulfill federal recordkeeping	
	requirements for administration of vaccine covered by the Vaccine Injury	
	Compensation Program. Enter the HCW's job title.	
Clinical specialty	Conditionally required. If Occupation is physician, fellow or intern/resident, select	
	the appropriate clinical specialty.	
Performs direct	Conditionally required. Required only when the HCW has influenza vaccination	
patient care	and/or influenza chemoprophylaxis/treatment records. Select Y (Yes) if the HCW	
	provides direct patient care (i.e., hands on, face-to-face contact with patients for the	
	purpose of diagnosis, treatment and monitoring); otherwise select N (No).	
Custom Fields	Optional. Up to two date fields, two numeric fields, and 10 alphanumeric fields that	
	may be customized for local use. NOTE: Each Custom Field must be set up in the	
Facility/Custom Options section of the application before the field can		
	for use.	
Comments	Optional. Enter any information about the HCW. This information cannot be	
	analyzed.	



Table 3. Instructions for Completion of the Exposure to Blood/Body Fluids Form (CDC 57.205)

Information for all blood/body fluid exposures should be recorded using this form. The variables to be entered depend upon whether the facility selects the exposure event only reporting or exposure reporting and management.

*Demographic data auto-entered by application if part of an existing HCW Demographic Data record (CDC 57.204).

Data Field Instructions for Data Collection Event Only Management			_	Exposure Event
Facility ID #	D 4 E' 11		Exposure	and Exposure
entered by the application. Exposure Event # The exposure event number will be autogenerated by the application. HCW ID Enter the HCW's alphanumeric identification number. This identifier is unique to the healthcare facility. *HCW Name: Last, First, Middle Gender Indicate the gender of the HCW by checking F (Female) or M (Male). *Date of Birth Enter the date of birth of the HCW using the format: mm/dd/yyyy. *Work Location Required. Select the code that best describes the HCW's current permanent work location. This refers to physical work location rather than to department assignment. Location codes are customized to the facility and set up prior to entering HCW records. See Table 2 for more details. *Occupation Required. Select the occupation code that most appropriately describes the HCW's job. Occupation codes are customized to the facility and set up prior to entering HCW records. See Table 2 for more details. Clinical Specialty If Occupation is physician, fellow or intern/resident, enter the appropriate clinical specialty. The list of clinical specialties can be found on Form CDC 57.204. Exposure Type The default setting is auto-entered by the application as Blood/Body Fluids. Section I – General Exposure Information 1. Did the exposure Choose Y (Yes) or N (No). Required Required Required Required Required Required				
Exposure Event # The exposure event number will be autogenerated by the application. HCW ID Enter the HCW's alphanumeric identification number. This identifier is unique to the healthcare facility. *HCW Name: Enter the HCW's name. Last, First, Middle *Gender Indicate the gender of the HCW by checking F (Female) or M (Male). *Date of Birth Enter the date of birth of the HCW using the format: mm/dd/yyyy. *Work Location Required. Select the code that best describes the HCW's current permanent work location. This refers to physical work location rather than to department assignment. Location codes are customized to the facility and set up prior to entering HCW records. See Table 2 for more details. *Occupation Required. Select the cocupation code that most appropriately describes the HCW's job. Occupation codes are customized to the facility and set up prior to entering HCW records. See Table 2 for more details. Clinical Specialty If Occupation is physician, fellow or intern/resident, enter the appropriate clinical specialty. The list of clinical specialties can be found on Form CDC 57.204. Exposure Type The default setting is auto-entered by the application as Blood/Body Fluids. Section I – General Exposure Information I. Did the exposure Choose Y (Yes) or N (No). Required Required	Facility ID #		Required	Required
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			Required	Required
			1	1



		12	Exposure Event
Data Field	Instructions for Data Collection	Exposure Event Only	and Exposure Management
1a. If No, specify the	If the exposure did not occur at the reporting	Conditionally	Conditionally
name of facility in which exposure	facility, enter the name of the facility where the event occurred.	required	required
occurred	event occurred.		
2. Date of exposure	Enter date of exposure in mm/dd/yyyy format.	Required	Required
3. Time of exposure	Enter the time the exposure occurred and whether it was AM or PM.	Required	Required
4. Number of hours on duty	Enter the number of hours the HCW had been on duty when the exposure occurred.	Optional	Optional
5. Is exposed person a temp/agency employee?	Choose Y (Yes) or N (No).	Optional	Optional
6. Location where exposure occurred	Choose the appropriate code for the physical location where the event took place. (This is customized to the facility).	Required	Required
7. Type of Exposure	Check the appropriate exposure type. Check all that apply.	Required	Required
7a. Percutaneous:	If Type of Exposure was Percutaneous, then check this item.	Conditionally required	Conditionally required
Did the exposure involve a clean, unused needle or sharp object?	If percutaneous is checked, then select Yes or No to indicate whether the exposure involved a clean, unused needle or sharp object. If the incident involved a clean, unused needle or sharp object you may not need to report this as an exposure (see your protocol for more information). If not, check No and complete Q8, Q9 and Section II. If following the protocol for exposure management also complete Sections V-XI.	Conditionally required	Conditionally required
7b. Mucous membrane	If Type of Exposure was Mucous Membrane, then check this item and complete Q8, Q9 and Section III. If following the protocol for exposure management also complete Sections V-XI.	Conditionally required	Conditionally required
7c. Skin:	If Type of Exposure was Skin, then check this item.	Conditionally required	Conditionally required
Was skin intact?	If Skin is checked, then indicate Y (Yes), N (No) or (U) Unknown for whether the skin remained intact during the exposure. If the answer is No, complete Q8, Q9 and Section III. If following the protocol for exposure management also complete Sections V-XI.	Conditionally required	Conditionally required



Data Field	Instructions for Data Collection	Exposure Event Only	Exposure Event and Exposure Management
7d. Bite	If Type of Exposure was Bite, then check this item and complete Q9 and Section IV. If following the protocol for exposure management also complete Sections V-XI.	Conditionally required	Conditionally required
8. Type of fluid/tissue involved in exposure	Select the Type of fluid/tissue from the list. If Solutions or Body fluids are checked, indicate whether visibly bloody or not visibly bloody. For Body Fluids, indicate the primary body fluid type implicated in the exposure from the list.	Required Conditionally required	Required Conditionally required
	If Other is selected for either the Type of Fluid/Tissue involved in the exposure or the Body Fluid Type, please specify the type. (Make sure it is not a body fluid that is already listed in the box on the right side of the form).	Conditionally required	Conditionally required
9. Body site of exposure	Check body site of exposure from the list. Check all sites that were exposed.	Required	Required
	If the Body site of exposure was (Other), please specify the site.	Conditionally required	Conditionally required
Section II - Percutan	eous Injury		
1. Was the needle or sharp object visibly contaminated with blood prior to exposure?	Choose Y (Yes) or N (No).	Required	Required
2. Depth of the injury (check one)	Indicate the depth of the injury from the needle or sharp object using the list provided. Exposures that are not obviously superficial (e.g., scratch) or deep (e.g., "muscle contracted" or "touched bone"), should be classified as moderate.	Conditionally required	Conditionally required



Data Field	Instructions for Data Collection	Exposure Event Only	Exposure Event and Exposure Management
3. What needle or sharp object caused the injury?	Select one of the following categories: Device, Non-Device Sharp Object, or Unknown Sharp Object. If you select Device in the application you will be provided with a Device button that will take you to a screen to enter manufacturer, model, etc. Once a device has been entered you will be able to select it from the drop down list.	Conditionally required	Conditionally required
	If a Non-Device Sharp is selected, please describe the item or object.	Conditionally required	Conditionally required
	Within Devices, there are six categories: Hollow-bore needles, Suture needles, Other solid sharps, Glass, Plastic, Non-sharp safety devices, and Other devices.		
	If Other known device is selected, please specify.	Conditionally required	Conditionally required
4. Manufacturer and model	Enter the brand name and model of the device used. If the brand and model are unknown, generic device descriptors can be entered.	Conditionally required	Conditionally required
5. Did the needle or other sharp object involved in the injury have a safety feature?	Choose Y (Yes) or N (No). If Yes, answer 5a and 5b. If No, skip to Q6.	Conditionally required	Conditionally required
5a. If Yes, indicate the type of safety feature	If above is Y (Yes), choose one item from the list of safety devices.	Conditionally required	Conditionally required
5b. If the device had a safety feature, when did the injury occur?	Choose the timing of the injury event with relation to the use of the safety device. Check one item from the list provided.	Conditionally required	Conditionally required



		_	Exposure Event
Data Field	Instructions for Data Collection	Exposure Event Only	and Exposure Management
6. When did the	Choose the timing of the injury event from the	Conditionally	Conditionally
injury occur?	list provided.	required	required
(check one) Before use of the item	Injuries that occurred prior to intended use and usually involve clean needles or sharp objects. It may also include injuries that occurred with a clean device that passed through bloody gloves.		
During use of the item	Injuries that occurred during the use of the needle or sharp object. It also includes surgical or other invasive procedures with many steps.		
After use of item, before disposal	Injuries that occurred while in transit to disposal, cleaning instrument or recapping.		
During or after disposal	Injuries that occurred during or after the process of disposal or because of improper disposal of a needle or other sharp object.		
<u>Unknown</u>	Time of injury relative to the use of the device or object is unknown.		
7. For what purpose or activity was the sharp device being	Choose from the lists provided. If Other specify the purpose in the space provided.	Conditionally required	Conditionally required
used?	Select Unknown if injury was a result of contact with discarded or uncontrolled sharps, or in circumstances where the intent of device or object use is unknown or cannot be ascertained.		
8. What was the activity at the time of injury?	Choose the activity being performed at the time of injury involving the sharp object or needle. If the activity being performed at the time of the injury was different than the purpose indicated in Q7, select the activity at the time the actual injury event took place.	Conditionally required	Conditionally required
9. Who was holding the device at the time the injury occurred?	Select one answer.	Conditionally required	Conditionally required
10. What happened when the injury occurred?	Choose one item from the list. If Other, please record details in the space provided.	Conditionally required	Conditionally required
	Membrane and/or Skin Exposure		
1. Estimate the	Select the estimated amount of blood or body	Conditionally	Conditionally
amount of blood/body fluid exposure	fluid involved in the mucous membrane or skin exposure. Indicate Unknown if unable to estimate the amount.	required	required



			Exposure Event
Doto Field	Instructions for Data Collection	Exposure	and Exposure
Data Field	Instructions for Data Collection	Event Only	Management
2. Activity/event when exposure	Select the activity or event at the time mucous membrane or skin exposure occurred.	Conditionally required	Conditionally required
occurred	memorane of skin exposure occurred.	required	required
occurred	If Other is selected record details of the activity	Conditionally	Conditionally
	or event in the space provided.	required	required
3. Barriers used by	Check all that apply.	Conditionally	Conditionally
the worker at the		required	required
time of exposure	If Other is selected, list other barriers in the	Conditionally	Conditionally
	space provided.	required	required
Section IV – Bite			<u> </u>
1. Wound	Select the description of the bite wound from	Conditionally	Conditionally
description	the list provided.	required	required
2. Activity/event	Choose the activity or event when the bite	Conditionally	Conditionally
when exposure	occurred.	required	required
occurred	If Other, specify the event in the space	Conditionally	Conditionally
	provided.	required	required
Sections V	– IX are required when following the protocols for		
Section V – Source In		Exposure man	
1. Was the source	Choose Y (Yes) if the source of the exposure	Optional	Required
patient known?	(patient) is known. Otherwise, select N (No).		1
2. Was HIV status	Indicate Y (Yes) if the source patient's	Optional	Required
known at time of	serostatus was known at the time of exposure.		
exposure?			
3. Check the test	Use codes: P= positive, N= negative,	Optional	Required
results for the source	I=Indeterminate, U=Unknown, R=Refused and		
patient:	NT=Not tested.		
Hepatitis B	Indicate the results of any tests performed prior		
ĤbsAg	to the exposure (as found in the medical record)		
HBeAg	or performed immediately after the exposure. If		
Total anti-HBc	the source is not known, check U. If the source		
anti-HBs Hepatitis C	refuses to be tested, check R. Not all tests listed		
anti-HCV EIA	on the form need to be offered after all		
anti-HCV suppl	exposures.		
PCR-HCV RNA			
HIV			
HIV EIA, ELISA			
Rapid HIV			
Confirmatory HIV Section VI – For HIV	Infected Source		
1. Stage of Disease	Indicate the stage of HIV disease of the source	Optional	Conditionally
1. Stage of Discuse	patient. Use CDC surveillance definitions. For	optionui	required
	end stage AIDS and acute HIV illness, use		4
	definitions as defined in the protocol.		



Data Field	Instructions for Data Collection	Exposure Event Only	Exposure Event and Exposure Management
2. Is the source	Indicate if the source patient is was taking anti-	Optional	Conditionally
patient taking anti-	retroviral drugs at the time of the exposure, Y (Yes), N (No), or U (Unknown).	Parama	required
retroviral drugs? 2a. If Yes, indicate	If the source patient was taking anti-retroviral	Optional	Conditionally
drug(s)	drugs at the time of the exposure, list them here. Drug codes are listed in Chapter 7 and will be in a drop down list in the application.	Optional	required
3. Most recent CD4	If available, indicate the most recent CD4 count	Optional	Conditionally
count	in mm ³ for the source patient.		required
Date	Enter the month and year of the test for the source patient.		
4. Viral Load	If available, indicate the most recent HIV viral	Optional	Conditionally
	load (# of copies per ml) or Undetectable for the		required
Data	source patient.		
Date Section VII: Initial C	Enter the month and year of the test. Care Given to Healthcare Worker		
1. HIV postexposure	are Given to Healthcare worker		
prophylaxis			.
Offered?	Choose Y (Yes), N (No), or U (Unknown) if antiretroviral drugs were offered to the HCW following this exposure.	Optional	Required
Taken?	Choose Y (Yes), N (No), or U (Unknown) if antiretroviral drugs were taken by the HCW. If Yes is selected, complete Post-Exposure Prophylaxis/Treatment form (CDC form 57.206).	Optional	Required
2. HBIG given?	Choose Y (Yes), N (No), or U Unknown) for whether Hepatitis B immunoglobulin was given.	Optional	Required
Date administered	Enter date HBIG prophylaxis pertaining to this exposure was administered. Use mm/dd/yyyy format.	Optional	Conditionally Required
3. Hepatitis B	Choose Y (Yes), N (No), or U. (Unknown) for	Optional	Required
vaccine given?	whether Hepatitis B vaccine was given after exposure.		
Date first dose		Optional	Conditionally
administered	Enter date of first dose of Hepatitis B vaccine (mm/dd/yyyy format). This and subsequent doses to complete the HBV series should be recorded in the HCW's file.		Required



Data Field	Instructions for Data Collection	Exposure Event Only	Exposure Event and Exposure Management	
4. Is the HCW	Indicate the pregnancy status of HCW. Choose	Optional	Conditionally	
pregnant?	Y (Yes), N (No), or U (Unknown).	1	required	
4a. If yes, which	Check 1 (1 st trimester), 2 (2 nd trimester), or 3	Optional	Conditionally	
trimester?	(3 rd trimester) at the time of exposure. If stage of		required	
	pregnancy is unknown, check U.			
Section VIII - Baselin				
Was baseline testing	Choose Y (Yes) or N (No) or U (Unknown).	Optional	Required	
performed on the	Baseline lab tests should be performed within			
HCW?	hours of the exposure.			
HIV EIA	Enter the dates for each test performed and the	Optional	Conditionally	
HIV confirmatory	result (Use codes: P= Positive, N= Negative,		required	
HepC anti-HCV EIA	I=Indeterminate, U=Unknown, R=Refused).			
HepC anti-HCV-supp HepC PCR HCV RNA				
HepB HBsAg				
HepB IgM anti-Hbc				
HepB Total anti-Hbc				
HepB Anti-HBs				
ALT	Additional baseline laboratory tests may be	Optional	Optional	
Amylase	completed to document potential physiologic			
Blood glucose Hematocrit	changes associated with a blood/body fluid			
Hemoglobin	exposure. Enter the date (in mm/dd/yyyy			
Platelets	format) and result, using the specified units.			
Blood cells in urine				
WBC				
Creatinine				
Other				
Section IX – Follow-u			n	
1. Is it recommended	Choose Y (Yes) or N (No).	Optional	Required	
that the HCW return				
for follow-up of this				
exposure?	Character V (Van) and N (N)	0	C111	
1a. If Yes, will	Choose Y (Yes) or N (No).	Optional	Conditionally	
follow-up be			Required	
performed at this				
facility? Section X – Narrative				
In the worker's		Optional	Ontional	
	Enter the narrative of the HCW's description of	Орионаг	Optional	
words, how did the	how the injury occurred.			
injury occur?	i an			
Section XI – Prevention				



Data Field	Instructions for Data Collection	Exposure Event Only	Exposure Event and Exposure Management
In the worker's words, what could have prevented the injury?	Enter the narrative of the HCW's assessment of how the injury might have been prevented.	Optional	Optional
Custom Fields	Up to two date fields, two numeric fields, and 10 alphanumeric fields that may be customized for local use. NOTE: Each Custom Field must be set up in the Facility/Custom Options section of the application before the field can be selected for use.	Optional	Optional
Comments	Enter any additional information about the HCW. CDC will not analyze this information.	Optional	Optional



Table 4. Instructions for Completion of the Healthcare Worker Prophylaxis/Treatment – BBF Postexposure Prophylaxis (PEP) Form (CDC 57.206)

Use this form if HIV postexposure prophylaxis (PEP) was administered to a healthcare worker following a blood or body fluid exposure.

*Demographic data auto-entered by application if part of an existing HCW Demographic Data record (CDC 57.204).

Data Field	Instructions for Data Collection
Facility ID #	Required. The NHSN-assigned facility ID will be auto-entered by the application.
MedAdmin ID#	Required. Medical administration number. Data will be auto-entered by the
	application.
HCW ID#	Required. Enter the HCW's alphanumeric identification number. This identifier is
	unique to the healthcare facility.
*HCW Name:	Optional. Enter the HCW's name.
Last, First, Middle	
*Gender	Required. Indicate the gender of the HCW by checking F (Female) or M (Male).
*Date of Birth	Required. Enter the date of birth of the HCW using the format: mm/dd/yyyy.
Infectious Agent	Required. Enter HIV on form. Select HIV in the application.
Exposure Event #	Required. The Exposure event number will be auto-entered by the system. Use the
	Link/Unlink button to find any exposures for the entered HCW, select, and link
	the exposure for which PEP is being administered. PEP records cannot be saved
	unless they are linked to an exposure. PEP records entered from the Blood and
	Body Fluid Exposure Form will automatically be linked to that exposure.
Initial PEP	Indication: Prophylaxis
Time between	Required. Enter the number of hours between the exposure and when the 1st dose
exposure and 1 st dose	of PEP was administered.
Drug	Required. Enter any drugs prescribed for prophylaxis. See Chapter 7 in the
	protocol for a list of individual drug codes.
Drug	Conditionally required. Enter any additional drugs prescribed for initial
	prophylaxis.
Drug	Conditionally required. Enter any additional drugs prescribed for prophylaxis.
Drug	Conditionally required. Enter any additional drugs prescribed for prophylaxis.
Date Started	Required. Enter the date the initial PEP regimen commenced (mm/dd/yyyy
	format). The start date will apply to all drugs selected as the initial PEP regimen.
	The date started must be on or after the exposure date.
Date Stopped	Required. Enter the date the initial PEP regimen was stopped (mm/dd/yyyy
	format).
	Note: If any drug(s) of a drug regimen are discontinued, the entire regimen is
	considered 'stopped.' If select drugs in the regimen continue to be used as
	prophylaxis (and if other drugs are added) enter them as drugs under a PEP
	change with a new start date.



Data Field	Instructions for Data Collection
Reason for Stopping	Required. Indicate the primary reason for stopping the initial PEP regimen by selecting the appropriate choice.
PEP Change 1	Indication: Prophylaxis
Drug	Required. Enter drugs prescribed for a second prophylaxis regimen. Note that the second PEP regimen may contain drugs that were included in the first regimen.
Drug	Conditionally required. Enter any additional drugs prescribed for prophylaxis.
Drug	Conditionally required. Enter any additional drugs prescribed for prophylaxis.
Drug	Conditionally required. Enter any additional drugs prescribed for prophylaxis.
Date Started	Conditionally required. Enter the date the second PEP regimen was started using mm/dd/yyyy format.
Date Stopped	Conditionally required. Enter the date the second PEP regimen was stopped using mm/dd/yyyy format.
	Note: If any drug(s) of a drug regimen are discontinued, the regimen is considered 'stopped.' Whatever drugs in the regimen are continued (and if other drugs are added) will constitute a new regimen and should be recorded as part of a new PEP regimen(s) with dates that resume from the last stop date.
Reason for Stopping	Conditionally required. Indicate the primary reason for stopping this PEP regimen by selecting the appropriate choice.
PEP Change 2	Indication: Prophylaxis
I DI Ciidiige I	indication: 1 tophytaxis
Drug	Conditionally required. Enter drugs prescribed for a third prophylaxis regimen. Note that the third PEP regimen may contain drugs that were included in previous
Drug	Conditionally required. Enter drugs prescribed for a third prophylaxis regimen. Note that the third PEP regimen may contain drugs that were included in previous regimens.
	Conditionally required. Enter drugs prescribed for a third prophylaxis regimen. Note that the third PEP regimen may contain drugs that were included in previous
Drug Drug	Conditionally required. Enter drugs prescribed for a third prophylaxis regimen. Note that the third PEP regimen may contain drugs that were included in previous regimens. Conditionally required. Enter any additional drugs prescribed for prophylaxis.
Drug Drug Drug	Conditionally required. Enter drugs prescribed for a third prophylaxis regimen. Note that the third PEP regimen may contain drugs that were included in previous regimens. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter any additional drugs prescribed for prophylaxis.
Drug Drug Drug Drug	Conditionally required. Enter drugs prescribed for a third prophylaxis regimen. Note that the third PEP regimen may contain drugs that were included in previous regimens. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter the date the new PEP regimen was started using
Drug Drug Drug Drug Date Started	Conditionally required. Enter drugs prescribed for a third prophylaxis regimen. Note that the third PEP regimen may contain drugs that were included in previous regimens. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter the date the new PEP regimen was started using mm/dd/yyyy format. Conditionally required. Enter the date the new PEP regimen was stopped using mm/dd/yyyy format. Note: If any drug(s) of a drug regimen are discontinued, the regimen is considered 'stopped.' Whatever drugs in the regimen are continued (and if other drugs are added) will constitute a new regimen and should be entered as such.
Drug Drug Drug Drug Date Started	Conditionally required. Enter drugs prescribed for a third prophylaxis regimen. Note that the third PEP regimen may contain drugs that were included in previous regimens. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter the date the new PEP regimen was started using mm/dd/yyyy format. Conditionally required. Enter the date the new PEP regimen was stopped using mm/dd/yyyy format. Note: If any drug(s) of a drug regimen are discontinued, the regimen is considered 'stopped.' Whatever drugs in the regimen are continued (and if other
Drug Drug Drug Drug Date Started Date Stopped	Conditionally required. Enter drugs prescribed for a third prophylaxis regimen. Note that the third PEP regimen may contain drugs that were included in previous regimens. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter the date the new PEP regimen was started using mm/dd/yyyy format. Conditionally required. Enter the date the new PEP regimen was stopped using mm/dd/yyyy format. Note: If any drug(s) of a drug regimen are discontinued, the regimen is considered 'stopped.' Whatever drugs in the regimen are continued (and if other drugs are added) will constitute a new regimen and should be entered as such. Conditionally required. Indicate the primary reason for stopping this PEP regimen
Drug Drug Drug Drug Date Started Date Stopped Reason for Stopping Adverse Reactions Signs or symptoms of	Conditionally required. Enter drugs prescribed for a third prophylaxis regimen. Note that the third PEP regimen may contain drugs that were included in previous regimens. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter the date the new PEP regimen was started using mm/dd/yyyy format. Conditionally required. Enter the date the new PEP regimen was stopped using mm/dd/yyyy format. Note: If any drug(s) of a drug regimen are discontinued, the regimen is considered 'stopped.' Whatever drugs in the regimen are continued (and if other drugs are added) will constitute a new regimen and should be entered as such. Conditionally required. Indicate the primary reason for stopping this PEP regimen by selecting the appropriate choice.
Drug Drug Drug Drug Date Started Date Stopped Reason for Stopping Adverse Reactions	Conditionally required. Enter drugs prescribed for a third prophylaxis regimen. Note that the third PEP regimen may contain drugs that were included in previous regimens. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter any additional drugs prescribed for prophylaxis. Conditionally required. Enter the date the new PEP regimen was started using mm/dd/yyyy format. Conditionally required. Enter the date the new PEP regimen was stopped using mm/dd/yyyy format. Note: If any drug(s) of a drug regimen are discontinued, the regimen is considered 'stopped.' Whatever drugs in the regimen are continued (and if other drugs are added) will constitute a new regimen and should be entered as such. Conditionally required. Indicate the primary reason for stopping this PEP regimen by selecting the appropriate choice.



Data Field	Instructions for Data Collection
Custom Fields	Optional. Up to two date fields, two numeric fields, and 10 alphanumeric fields that may be customized for local use. NOTE: Each Custom Field must be set up in the Facility/Custom Options section of the application before the field can be selected for use.
Comments	Optional. Enter any additional information about the HCW. CDC will not analyze this information.



Table 5: Instructions for Completion of Follow-Up Laboratory Testing Form (CDC 57.207)

This form should be completed for HCP who have additional laboratory testing done as a result of blood or body fluid exposures. These tests would occur after baseline laboratory testing had been completed.

Demographic data auto-entered by application if part of an existing HCW Demographic Data record (CDC 57.204).

Demograpine data data	b-entered by application if part of an existing HCW Demographic Data record (CDC 57.204).	
Data Field	Instructions for Data Collection	
Facility ID#	Required. The NHSN-assigned facility ID will be auto-entered by the application.	
Lab #	Required. The lab testing ID number will be auto-generated by the application.	
HCW ID#	Required. Enter the HCW's alphanumeric identification number. This identifier is unique to the healthcare facility.	
*HCW Name:	Optional. Enter the HCW's name.	
Last, First, Middle		
*Gender	Required. Indicate the gender of the HCW by checking F (Female) or M (Male).	
*Date of birth	Required. Enter the date of birth of the HCW using the format: mm/dd/yyyy.	
Exposure Event #	Required. The user is required to link the laboratory follow-up record to a blood and body fluid exposure record using the Link feature within the application. Once the exposure is selected and submitted, the form will display the message "Lab is Linked." Laboratory records must be linked to an exposure.	
Lab Results		
Lab Test	Required (At least one laboratory test and date are required). Multiple test results may be recorded on this form. Select lab test from dropdown menu:	
	HIV EIA HIV confirmatory Amylase HepC anti-HCV EIA HepC anti-HCV-supp Hematocrit HepC PCR HCV RNA Hemoglobin HepB HBsAg HepB IgM anti-Hbc HepB Total anti-Hbc HepB Anti-HBs Other	
Date	Required. Indicate date of test using mm/dd/yyyy format.	
Result	Conditionally required. Select one of the result codes: Use codes: P= positive, N= negative, I=Indeterminate, U=Unknown, R=Refused)	
Custom Fields	Optional. Up to two date fields, two numeric fields, and 10 alphanumeric fields that may be customized for local use. NOTE: Each Custom Field must be set up in the Facility/Custom Options section of the application before the field can be selected for use.	
Comments	Optional. Enter any additional information about the HCW. CDC will not analyze this information.	



Table 6. Instructions for Completion of the Healthcare Worker Prophylaxis/Treatment – Influenza Form (CDC 57.210)

This form should be completed when an HCW receives antiviral medications as influenza treatment or as chemoprophylaxis against influenza infection. It is used to collect information on which antiviral medications were administered, when, and what (if any) adverse reactions were experienced by the HCW.

*Demographic data auto-entered by application if part of an existing HCW Demographic Data record (CDC 57.204).

2 cm ograpme data date	-chered by application if part of all existing the wideling raphic Data record (eDe 37,204).
Data Field	Instructions for Data Collection
Facility ID #	Required. The NHSN-assigned facility ID will be auto-entered by the application.
Med Admin ID #	Required. The medication administration ID number will be auto-generated by the
	application.
HCW ID#	Required. Enter the HCW's alphanumeric identification number. This identifier is
	unique to the healthcare facility.
*HCW Name:	Optional. Enter the HCW's name.
Last, First, Middle	
*Gender	Required. Indicate the gender of the HCW by checking F (Female) or M (Male).
*Date of Birth	Required. Enter the date of birth of the HCW using the format: mm/dd/yyyy.
*Work Location	Required. Select the code that best describes the HCW's current permanent work
	location. This refers to physical work location rather than to department assignment.
	Location codes are customized to the facility and set up prior to entering HCW
	records. See Table 2 for more details.
*Occupation	Required. Select the occupation code that most appropriately describes the HCW's
	job. Occupation codes are customized to the facility and set up prior to entering
A	HCW records. See Table 2 for more details.
*Clinical Specialty	Conditionally required. If Occupation is physician, fellow or intern/resident, enter the
	appropriate clinical specialty. The list of clinical specialties can be found on Form
*Performs direct	CDC 57.204.
patient care	Required. Select Yes if the HCW provides direct patient care (i.e., hands on, face-to-face contact with patients for the purpose of diagnosis, treatment and monitoring);
patient care	otherwise select No.
Infectious agent	Required. Auto-filled on hard copy form. Select Influenza in application.
For season	Required. Select the vaccination season. Specify the year(s) during which this
1 of season	chemoprophylaxis or treatment date falls. For NHSN purposes, the vaccination
	"season" is 7/1 of the first year to 6/30 of the next calendar year.
#	Required. Indicate up to 10 antiviral medications given using sequential numbers
	starting with 1.
Indication	Required. Select Prophylaxis or Treatment as appropriate.
Influenza subtype	Required. Select the influenza subtype for which the HCW is receiving antiviral
	medications (for post-exposure chemoprophylaxis or for treatment). Select Unknown,
	if you do not know the specific subtype necessitating antiviral medication use.
Antiviral	Required. Enter the code of the antiviral medication that was administered to the
medication	HCW using the codes listed at the bottom of the form.
Start date	Required. Enter the start date of the antiviral using mm/dd/yyyy format.
Stop date	Conditionally required. Enter the stop date of the antiviral using mm/dd/yyyy format.



Data Field	Instructions for Data Collection
Adverse reactions?	Required. Check Yes if the HCW had a severe adverse reaction attributable to the
	influenza antiviral medication; otherwise check No. If it is unknown whether or not
	the HCW experienced any adverse reactions, check Don't Know.
Adverse reactions	Conditionally required. If the HCW had a severe adverse reaction, check all reactions
to antiviral	that apply for each medication administered. Please correlate the antiviral medication
medication	# with the antiviral medication on page 1. If an adverse reaction is not listed, check
#1#10	Other and specify the adverse reaction in the space provided. All Other adverse
	reactions should be included if the reactions were severe enough to affect daily
	activities and/or resulted in the discontinuation of the antiviral medication.
Custom Fields	Optional. Up to two date fields, two numeric fields, and 10 alphanumeric fields that
	may be customized for local use. NOTE: Each Custom Field must be set up in the
	Facility/Custom Options section of the application before the field can be selected for
	use.
Comments	Optional. Enter any additional information about the HCW. CDC will not analyze
	this information.



Table 7. Instructions for Completion of Healthcare Personnel Safety Component – Annual Facility Survey (CDC 57.200)

This form must be completed once a year by any facility using the Healthcare Personnel Safety Component.

Data Field	Instructions for Data Collection/Entry
Tracking #	Required. The NHSN-assigned Tracking # will be auto-entered by the
	application.
Facility ID #	Required. The NHSN-assigned facility ID will be auto-entered by the
-	application.
Survey year	Required. Enter the year of the survey using the format: yyyy.
Total beds set up and staffed	Required. Enter the number of all active beds across specialties and intensive care units.
Patient admissions	Required. Enter the number of patients, excluding newborns, admitted for
	inpatient service.
Inpatient days	Required. Enter the number of adult and pediatric days of care, excluding
	newborn days of care, rendered during a specified reporting period.
Outpatient encounters	Required. Enter the number of visits by patients who are not admitted as
	inpatients to the hospital while receiving medical, dental, or other services.
Number of hours worked by	Optional. Number of hours worked is available from OSHA300 reporting
all employees	logs. The value can also be calculated by identifying the number of full
	time employees working in your facility within a year, multiply by the
	number of work hours for one full time employee in a year (typically
	ranges from 2000-2100 hours per year). Add in overtime hours and total
	hours worked by part-time, temporary, and contracted staff.
Number of HCWs	Required. HCWs are all persons who work in the hospital. Calculate the
	number of attending physicians by including only those who are active or
	associate staff (e.g. similar methodology to the American Hospital
	Association annual survey, if applicable). Do not include courtesy,
	consulting, honorary, provisional, or other attending physicians in this
	number. If you cannot determine the exact number for a particular category,
	please estimate it. If the facility does not have any HCP in a specific
	occupation, the user may enter 0. This is the denominator when used to
	calculate rates of particular exposure events per HCW.
Number of FTEs	Required. A subset of total number of HCP. FTEs are all HCP whose
	regularly scheduled workweek is 35 hours or more. To calculate the
	number of FTE's add the number of FTEs to ½ the number of part-time
	HCP (e.g., 2 part-time HCP = 1 FTE). If you cannot determine the exact
	number for a particular category, please estimate it. If the facility does not
	have any FTEs in a specific occupation, the user may enter 0. This is the
	denominator used to calculate rates of particular exposure events per FTE.



REFERENCES

The following CDC/PHS publications provide recommendations for management and follow-up of blood and body fluid exposures to HBV, HCV, and HIV:

- Updated U.S. Public Health Service Guidelines for the Management of Occupational Exposures to HBV, HCV, and HIV and Recommendations for Postexposure Prophylaxis. (MMWR, June 29, 2001 / 50(RR11); 1-42)
- Updated U.S. Public Health Service Guidelines for the Management of Occupational Exposures to HIV and Recommendations for Postexposure Prophylaxis (PEP regimens have been changed). (MMWR, September 30, 2005 / 54(RR09); 1-17)

The following CDC/PHS publication provides recommendations for the immunization of HCP:

- A Comprehensive Immunization Strategy to Eliminate Transmission of Hepatitis B Virus Infection in the United States. (MMWR, December 8, 2006 / 55(RR16); 1-25)
- Influenza Vaccination of Health-care Personnel. (MMWR, February 24, 2006 / 55(RR02); 1-16)
- Prevention and Control of Seasonal Influenza with Vaccines: Recommendations of the Advisory Committee on Immunization Practices (ACIP). (MMWR, July 29, 2009 / 58(Early Release); 1-52)