National HIV Surveillance System (NHSS)

Attachment 3a.

Adult HIV Confidential Case Report Form

Patient Identification (reco *First Name		*Middle Na		1	*La	*Last Name			Last Name Soundex		
Alternate Name Type (ex: Alias, Ma	s, Married)		*First Name		* M i	*Middle Name		*	*Last Name		
Address Type Residential Bac Bac Foster home He Postal Shelter	omeless	s □ Military		*Curren	t Address, S	treet				Address Date	
*Phone City			County		Sta	te/Countr	у		*ZII	P Code	
*Medical Record Number			<u> </u>	*Other ID T	ype			*Num	ber		
U.S. Department of Health and Human Services	(Patie		HIV Col)	Centers for Disease Control and Prevention (CDC)	
Health Department Use Or		cord all da					Form	approved	OMB no	. NNNN-NNNN Exp. MM/DD/YYY	
Date Received at Health Departme	ent		eHARS Do	ocument UI	D	State Number			ımber		
Reporting Health Dept—City/Cour	nty		-	City/County Nur			Number	mber			
Document Source				Surveillance Method □ Active □ Passive □ Follow up □ Reabstraction □ Unknown							
Did this report initiate a new case investigation? ☐ Yes ☐ No ☐ Unknown			Report Me □ 1-Field v		2-Mailed □ 3-Faxed □ 4-Phone □ 5-Electro			ectronic t	ransfer □ 6-CD/disk		
Facility Providing Informat	ion (r	ecord all d	lates as m	m/dd/yyyy)						
Facility Name								*Phone			
*Street Address											
City	County	/			State/Cour	itry		*ZIP Cod	е		
Facility <u>Inpatient</u> : Type □ Hospital □ Other, specify		Adult HIV clin			Screening, D □ CTS □ S □ Other, spec	TD clinic			□ Labora	cility: ☐ Emergency room tory ☐ Corrections ☐ Unknown specify	
Date Form Completed	/_		*Person Co	mpleting Fo	orm			*Phone			
Patient Demographics (rec	ord al	l dates as	mm/dd/yyy	/y)							
Sex Assigned at Birth ☐ Male ☐ Female ☐ Unknown				try of Birth	IS dependen	ov (plaasa	cpocify)				
Date of Birth / /			03	□ Other/C	Alias Date		specify) _	/			
							State (of Death			
Current Gender Identity Male	□ Fe	male 🗆 Tra	ansgender m		e (MTF)	ransgend			FTM) □	Unknown	
□ Additional gender identity (specify) Ethnicity □ Hispanic/Latino □ Not Hispanic/Latino □ Unknown Expanded Ethnicity											
		dian/Alaska aiian/Other F	Native Pacific Islande		Black/Africar nite □ Unkı		Expan	ded Race)		
Residence at Diagnosis (ad	dd add	itional ad	dresses in	Comment	s) (record	all dates	s as mm/	dd/yyyy))		
Address Event Type) = [Danislaman at	t HIV diagnos	sis □ Resid	lence at stage	3 (AIDS) o	diagnosis	□ Check	if SAME	as current address	
(check all that apply to address below	w) ⊔ r	residence a	critic alagnos								
(check all that apply to address below Address Type ☐ Residential ☐ B				y □ Foster	home 🗆 Ho		Military				
				y □ Foster	home □ Ho		Military				

Public reporting burden of this collection of information is estimated to average 20 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, Project Clearance Officer, 1600 Clifton Road, MS D-74, Atlanta, GA 30333, ATTN: PRA (0920-0573). **Do not send the completed form to this address.**

STATE/LOCAL USE ONL	Υ									
*Provider Name (Last, First, N						*Phone ()			
Hospital/Facility	,					`	,			
•										
Facility of Diagnosis (ad			(AIDO) = -	Objects if OANE	6	tallia ar ta farana	- 41			
Diagnosis Type (check all that a	apply to facility below	r) HIV Stage 3	(AIDS)	Check if <u>SAME</u> as			ation			
Facility Name					*Pho	ne ()				
*Street Address	0		la							
City		1 1				*ZIP Code	Facility: □ Emergency room			
Facility Type <u>Inpatient</u> : □ Ho □ Other, specify	spital <u>Outpatient</u> : □ □ Adult HIV o	Private physician's office	Screening, I	<i>Diagnostic, Referra</i> STD clinic		Other Facility ☐ Laboratory			Jnknown	
		cify		ecify		□ Other, spe				
*Provider Name	,	*Provider Phone ()			Spec	ialty				
				_						
Patient History (respond After 1977 and before the earli	<u> </u>	<u> </u>			Pediatric	RISK (ple	ease ent	er in Cor	nments	
	est known diagnos	is of filv infection, this p	Datient nau:							
Sex with male						□ Ye				
Sex with female						□ Ye				
Injected nonprescription drugs							s 🗆 No			
Received clotting factor for hemo Specify clotting factor:	philia/coagulation dis	sorder	Date red	ceived /	/	□ Y€	es 🗆 No	□ Unkr	iown	
HETEROSEXUAL relations with	h anv of the followi	ng:	Date let	ceived/						
HETEROSEXUAL contact with in						□ Ye	es □ No	□ Unkr	nown	
HETEROSEXUAL contact with hisexual male							es 🗆 No	□ Unkr	nown	
HETEROSEXUAL contact with person with hemophilia/coagulation disorder with documented HIV infection										
IETEROSEXUAL contact with person with hemophilia/coagulation disorder with documented HIV infection										
	IETEROSEXUAL contact with transplant recipient with documented HIV infection								nown	
	IETEROSEXUAL contact with person with documented HIV infection, risk not specified								nown	
Received transfusion of blood/blo			·	on in Comments)		□Ye	es 🗆 No	□ Unkr	nown	
First date received// Last date received// Received transplant of tissue/organs or artificial insemination							es 🗆 No	□ Unkr	nown	
Worked in a healthcare or clinical						□ Ye	es □ No	□ Unkr	nown	
If occupational exposure is being		idered								
as primary mode of exposure, specify occupation and setting:										
Other documented risk (please include detail in Comments)						□ Ye	s 🗆 No	□ Unkr	iown	
Clinical: Acute HIV Infec	ction and Oppo	rtunistic Illnesses	(record all	l dates as mm/	dd/yyyy)					
Suspect acute HIV infection? If				IIV test data in Labor	atory Data sec	ction, and	□ Yes	□ No □ l	Jnknowr	
enter patient or provider report of prev Clinical signs/symptoms consiste	ent with acute retrovi	ral syndrome (e.g., fever, i	malaise/fatio	uue. mvalaia. phar			□ Yes		Jnknown	
lymphadenopathy)? Date of sig Other evidence suggestive of ac	n/symptom onset _	1/				, 	l	□ No □ U		
Date of evidence///		II TES, piease describe.					l res	1 NO 11 C	TIKHOWH	
Opportunistic Illnesses	lp. p	le:	-		n					
Diagnosis Candidiasis, bronchi, trachea, or lungs	Dx Date	Diagnosis Herpes simplex: chronic ulcers			Diagnosis M. tuberculosis	. pulmonary ¹		Dx Date		
3		duration), bronchitis, pneumoni esophagitis				, , , , , , , , , , , , , , , , , , , ,				
Candidiasis, esophageal		Histoplasmosis, disseminated of	or		M. tuberculosis,		or			
Carcinoma, invasive cervical		extrapulmonary Isosporiasis, chronic intestinal	(>1 mo.		extrapulmonary ¹ Mycobacterium, of other/unidentified					
, 		duration)	, -		species, disseminated or extrapulmonary					
Coccidioidomycosis, disseminated or extrapulmonary		Kaposi's sarcoma			Pneumocystis p	orieumonia				
Cryptococcosis, extrapulmonary		Lymphoma, Burkitt's (or equiva			Pneumonia, red		o. period			
Cryptosporidiosis, chronic intestinal (>1 mo. duration)		Lymphoma, immunoblastic (or	equivalent)		Progressive muleukoencephalo					
Cytomegalovirus disease (other than in iver, spleen, or nodes)		Lymphoma, primary in brain			Salmonella sep	ticemia, recurr	ent			
Cytomegalovirus retinitis (with loss of		Mycobacterium avium complex			Toxoplasmosis	of brain, onset	at >1 mo.			
rision) HIV encephalopathy		kansasii, disseminated or extra	pulmonary		of age Wasting syndro	me due to HIV				
If a diagnosis data is entered for other t	phoroulogic diagnosis = 5	ove provide BVCT Cose Novelle			5 7					

Laboratory Data (record additional tests and tests not specified	d below in Comments) (record all dates as mm/dd/yyyy)
HIV Immunoassays (Nondifferentiating)	
TEST 1 - HIV-1 IA - HIV-1/2 IA - HIV-1/2 Ag/Ab - HIV-1 WB - HIV-1 I	
Test brand name/Manufacturer	
Facility name	Provider name
Result Positive Negative Indeterminate	
TEST 2 HIV-1 A HIV-1/2 A HIV-1/2 Ag/Ab HIV-1 WB HIV-1	
Test brand name/Manufacturer	
Result □ Positive □ Negative □ Indeterminate	Provider name
HIV Immunoassays (Differentiating)	Officerion Bate
☐ HIV-1/2 type-differentiating immunoassay	Role of test in diagnostic algorithm
(differentiates between HIV-1 Ab and HIV-2 Ab)	□ Screening/initial test □ Confirmatory/supplemental test
Test brand name/Manufacturer	Lab name
Facility name	
Result ¹ Overall interpretation: □ HIV-1 positive □ HIV-2 positive □ HIV p	
□ HIV-1 indeterminate □ HIV-2 indetermina	te
	Always complete the overall interpretation. Complete the analyte results when available.
□ HIV-1/2 Ag/Ab differentiating immunoassay (differentiates between HIV Ag	
Test brand name/Manufacturer	
Facility name	Provider name
Result □ Ag positive □ Ab positive □ Both (Ag and Ab positive) □ Negative	/e □ Invalid
Collection Date// Point-of-care rapid test	
☐ HIV-1/2 Ag/Ab and type-differentiating immunoassay (differentiates amon	g HIV-1 Ag, HIV-1 Ab, and HIV-2 Ab)
Test brand name/Manufacturer	
Facility name	
Result ² Overall interpretation: □ Reactive □ Nonreactive □ Index value	
Analyte results: HIV-1 Ag: ☐ Reactive ☐ Nonreactive ☐ Not report	
HIV-1 Ab: □ Reactive □ Nonreactive □ Reactive	
HIV-2 Ab: ☐ Reactive ☐ Nonreactive ☐ Reactive	
Collection Date/ Point-of-care rapid test 2	Complete the overall interpretation and the analyte results.
HIV Detection Tests (Qualitative) TEST □ HIV-1 RNA/DNA NAAT (Qualitative) □ HIV-1 culture □ HIV-2 RNA/	DNA NAAT (Qualitativa) HIV 2 aultura
Test brand name/Manufacturer	
Facility name	
Result □ Positive □ Negative □ Indeterminate	Collection Date / /
HIV Detection Tests (Quantitative viral load) Note: Include earliest test a	
TEST 1 ☐ HIV-1 RNA/DNA NAAT (Quantitative viral load) ☐ HIV-2 RNA/DNA	NAAT (Quantitative viral load)
Test brand name/Manufacturer	Lab name
Facility name	Provider name
	Log Collection Date / /
TEST 2 ☐ HIV-1 RNA/DNA NAAT (Quantitative viral load) ☐ HIV-2 RNA/DNA	
Test brand name/Manufacturer	
Facility name	Provider name
Result Detectable Undetectable Copies/mL	Log Collection Date / / /
Drug Resistance Tests (Genotypic) TEST □ HIV-1 Genotype (Unspecified)	Test brand name/Manufacturer
Lab name	
Provider name	Facility name
Immunologic Tests (CD4 count and percentage)	Concolion Bate
	. CD4 percentage % Collection Date / /
	Lab name
Facility name	Provider name
First CD4 result <200 cells/µL or <14%: CD4 count cells/µL	CD4 percentage% Collection Date//
Test brand name/Manufacturer	
Facility name	
	Provider name
Test brand name/Manufacturer	Provider name % Collection Date //
	Provider name % Collection Date //
Test brand name/Manufacturer	Provider name CD4 percentage % Collection Date //
Test brand name/Manufacturer Facility name Documentation of Tests Did documented laboratory test results meet approved HIV diagnostic algorithms also become collection date of earliest positive test for this also	Provider name CD4 percentage% Collection Date// Lab name Provider name orithm criteria? □ Yes □ No □ Unknown gorithm / /
Test brand name/Manufacturer Facility name Documentation of Tests Did documented laboratory test results meet approved HIV diagnostic algority YES, provide specimen collection date of earliest positive test for this all Complete the above only if none of the following were positive for HIV-1: Weste	Provider name CD4 percentage% Collection Date/
Test brand name/Manufacturer Facility name Documentation of Tests Did documented laboratory test results meet approved HIV diagnostic algorithms also become collection date of earliest positive test for this also	Provider name CD4 percentage
Test brand name/Manufacturer	Provider name CD4 percentage

Treatment/Services Referrals (record all dates	s as mm/dd/yyyy)	
Has this patient been informed of his/her HIV infection?		ed about their HIV exposure and counseled by
☐ Yes ☐ No ☐ Unknown Evidence of receipt of HIV medical care other than labora	☐ 1-Health dept ☐ 2-Physician/Prov	
to the contract of the contrac	Date of medical visit or prescription	,
For Female Patient		
This patient is receiving or has been referred for gynecol obstetrical services ☐ Yes ☐ No ☐ Unknown	logical or Is this patient currently pre	
For Children of Patient (record most recent birth in these	e boxes; record additional or multiple birth	ns in Comments)
*Child's Name		Child's Date of Birth
Child's Last Name Soundex	Child's State Number	
Facility Name of Birth		*Phone
(if child was born at home, enter "home birth") Facility Type Inpatient: Out	thatiant:	Other Facility D. Francesco
	· ·	Other Facility: ☐ Emergency room ☐ Corrections ☐ Unknown
☐ Other, specify	* 1 *	□ Other, specify
*Street Address		*ZIP Code
City	County	State/Country
Antiretroviral Use History (record all dates as i	mm/dd/vyvy)	
Main source of antiretroviral (ARV) use information (selection		Date patient reported information
☐ Patient interview ☐ Medical record review ☐ Pro	ovider report NHM&E Other	er//
Ever taken any ARVs? ☐ Yes ☐ No ☐ Unknown		
If yes, reason for ARV use (select all that apply)		
□ HIV Tx ARV medications		
□ PrEP ARV medications		
□ PEP ARV medications		
□ PMTCT ARV medications	Date began / /	
☐ HBV Tx ARV medications	Date began //	
□ Other (specify reason)		
ARV medications	Date began / /	
HIV Testing History (record all dates as mm/dd/	/vvv)	
Main source of testing history information (select one)		Date patient reported information
□ Patient interview □ Medical record review □ Provider	report □ NHM&E □ Other	
Ever had previous positive HIV test? Yes No L	Jnknown Date of first po	ositive HIV test//
Ever had a negative HIV test? Yes No Unknown	•	e HIV test (if date is from ne, enter in Lab Data section)///
Number of negative HIV tests within the 24 months befor	re the first positive test □ Unk	known
Comments		
Comments		
*Local/Optional Fields		

This report to CDC is authorized by law (Sections 304 and 306 of the Public Health Service Act, 42 USC 242b and 242k). Response in this case is voluntary for federal government purposes, but may be mandatory under state and local statutes. Your cooperation is necessary for the understanding and control of HIV. Information in CDC's National HIV Surveillance System that would permit identification of any individual on whom a record is maintained is collected with a guarantee that it will be held in confidence, will be used only for the purposes stated in the assurance on file at the local health department, and will not otherwise be disclosed or released without the consent of the individual in accordance with Section 308(d) of the Public Health Service Act (42 USC 242m).