Form Approved OMB No. 0920-0573

Expiration Date: XX/XX/XXXX

National HIV Surveillance System (NHSS)

Attachment 3a.

Adult HIV Confidential Case Report Form

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/ATSDR Information Collection Review Office, 1600 Clifton Road NE, MS D-74, Atlanta, Georgia 30329; ATTN: PRA (0920-0573).

Patient Identification (red	ord all dates a	as mm/dd/yyyy	/)				
*First Name	*Middle Na	me		*Last Name		L	ast Name Soundex
Alternate Name Type (ex: Alias, Married)		*First Name		*Middle Name		*Last N	ame
Address Type □ Residential □ B □ Foster Home □ Homeless □ Pos			Current Addre	ss, Street			Address Date
*Phone City		County		State/Country		*	ZIP Code
*Medical Record Number		*Othe	er ID Type			* Number	
U.S. Department of Health & Human Services	Adult + (Patients ≥13 Years	IV Confide s of Age at Time of				to CDC	Centers for Disease Contro and Prevention
Health Department Use C		dates as mm	/dd/yyyy)		For	m approved ON	IB no. 0920-0573 Exp. XX/XX/XXX
Date Received at Health Departm	ent	eHARS Docui	ment UID			State Numbe	er
Reporting Health Dept - City/Cou	nty			City/County No	umber		
Document Source		Surveillance M	ethod 🗆 Acti	ve □ Passive □ Fo	ollow up	□ Reabstractio	n □ Unknown
Did this report initiate a new case ☐ Yes ☐ No ☐ Unknown	e investigation?	Report Medium	ı □ 1-Field \	/isit □ 2-Mailed □ 5-Electronic Tra			ne
Facility Providing Informa	ntion (record al	I dates as mn	n/dd/yyyy)				
Facility Name					*F	Phone (
*Street Address							
City	County		State/	Country	*	ZIP Code	
Facility Inpatient: Type □ Hospital □ Other, specify	□ Adult F	ent: □ Private Physici HIV Clinic specify	A	creening, Diagnosti gency: □ CTS □ Other, specify	STD Clin	nic 🗆 Laborato	lity: □ Emergency Room ry □ Corrections □ Unknown pecify
Date Form Completed/	_/	*Person Complet	ting Form		*F	Phone ()	
Patient Demographics (re	cord all dates	as mm/dd/yyy	y)				
Sex assigned at Birth Male	□ Female □ Unknow	vn Country of E	Birth □ US □	Other/US Depend	dency (p	lease specify)	
Date of Birth//		·	Alias Date o	f Birth/	/		
Vital Status □ 1-Alive □ 2-Dead	ı	Date of Death			State of	Death	
Current Genoer Identity	le □ Female □ Tra ditional gender ident	•	Female (MTF)	□ Transgender F	emale-to-	-Male (FTM)	Unknown
Ethnicity	o □ Not Hispanic/La	atino 🗆 Unknown			Expande	d Ethnicity _	
	erican Indian/Alaska tive Hawaiian/Other			rican American Jnknown	Expande	d Race	
Residence at Diagnosis (a	dd additional a	addresses in (Comments	(record all d	lates a	s mm/dd/y	(VY)
Address Type (Check all that apply to address be	low) □ Residence	at HIV diagnosis	□ Residence	at AIDS diagnosis	□ Che	ck if <u>SAME as</u>	Current Address
*Street Address							Address Date
City	County		State/Co	untry			*ZIP Code

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STATE/LOCAL USE ONLY					
*Provider Name (Last, First, M.I.)			*Phone ()		_
Hospital/Facility					
— — — — — — Facility of Diagnosis (add add	ditional	facilities in Commer			
Diagnosis Type (Check all that apply	to facility	below) □ HIV □ AIDS	☐ Check if <u>SAME as Facilit</u>	y Providing	nformation
Facility Name				*Phone ()
*Street Address					, <u> </u>
City	County		State/Country		*ZIP Code
Facility Inpatient: □ Hospital Type □ Other, specify	☐ Adult F	<u>nt:</u> □ Private Physician's Office IIV Clinic specify	Screening, Diagnostic, Referra □ CTS □ STD Clinic □ Other, specify	al Agency:	Other Facility: ☐ Emergency Room ☐ Laboratory ☐ Corrections ☐ Unknown ☐ Other, specify
*Provider Name		*Provider Phone () _		Specialty	y
Patient History (respond to all	questio	ns) (record all dates as	mm/dd/yyyy) 🗆 Pediatric	risk (pl	ease enter in Comments)
After 1977 and before the earliest kno	wn diagn	osis of HIV infection, this p	patient had:		
Sex with male					□ Yes □ No □ Unknown
Sex with female					□ Yes □ No □ Unknown
Injected non-prescription drugs					□ Yes □ No □ Unknown
Received clotting factor for hemophilia/ coagulation disorder		ecify clotting factor: ite received (mm/dd/yyyy):			□ Yes □ No □ Unknown
HETEROSEXUAL relations with any of	of the follo	wing:			
HETEROSEXUAL contact with intrave	nous/injed	tion drug user			□ Yes □ No □ Unknown
HETEROSEXUAL contact with bisexu	al male				□ Yes □ No □ Unknown
HETEROSEXUAL contact with person	with hem	ophilia/coagulation disorder v	with documented HIV infection		□ Yes □ No □ Unknown
HETEROSEXUAL contact with transfu	sion recip	ent with documented HIV inf	ection		□ Yes □ No □ Unknown
HETEROSEXUAL contact with transpl	ant recipie	nt with documented HIV infe	ction		□ Yes □ No □ Unknown
HETEROSEXUAL contact with person	with docu	mented HIV infection, risk no	ot specified		□ Yes □ No □ Unknown
Received transfusion of blood/blood cor	nponents	other than clotting factor) (do	ocument reason in Comments)		□ Yes □ No □ Unknown
First date received///	Las	st date received/	_/		
Received transplant of tissue/organs or	artificial in	semination			□ Yes □ No □ Unknown
Worked in a healthcare or clinical laboral foccupational exposure is being investing i		•	of exposure, specify occupation	and settir	g: Unknown

This report to the Centers for Disease Control and Prevention (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).

Other documented risk (please include detail in Comments)

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 \square Yes \square No \square Unknown

Laboratory Data (record additional tests and tests not specified below in Comments) (record all dates as mm/dd/yyyy)

HIV Immunoassays (Non-differentiating)
TEST 1: □ HIV-1 IA □ HIV-1/2 IA □ HIV-1/2 Ag/Ab □ HIV-1 WB □ HIV-1 IFA □ HIV-2 IA □ HIV-2 WB
Test Brand Name/Manufacturer:
RESULT: Positive/Reactive Negative/Nonreactive Indeterminate Collection Date:// Rapid Test (check if rapid)
TEST 2: 🗆 HIV-1 IA 🗆 HIV-1/2 IA 🗅 HIV-1/2 Ag/Ab 🗅 HIV-1 WB 🗀 HIV-1 IFA 🗀 HIV-2 IA 🗀 HIV-2 WB
Test Brand Name/Manufacturer:
RESULT: □ Positive/Reactive □ Negative/Nonreactive □ Indeterminate Collection Date:/ □ Rapid Test (check if rapid)
HIV Immunoassays (Differentiating)
□ HIV-1/2 Type-differentiating (Differentiates between HIV-1 Ab and HIV-2 Ab) Test Brand Name/Manufacturer:
RESULT: ☐ HIV-1 ☐ HIV-2 ☐ Both (undifferentiated) ☐ Neither (negative) ☐ Indeterminate Collection Date: ☐ / ☐ / ☐ ☐ ☐ Rapid Test (check if rapid)
□ HIV-1/2 Ag/Ab-differentiating (Differentiates between HIV Ag and HIV Ab) Test Brand Name/Manufacturer:
RESULT: □ Ag reactive □ Ab reactive □ Both (Ag and Ab reactive) □ Neither (negative) □ Invalid/Indeterminate Collection Date: □ / □ / □ □ □ Rapid Test (check if rapid)
□ HIV-1/2 Ag/Ab and Type-differentiating (Differentiates among HIV-1 Ag, HIV-1 Ab, HIV-2 Ab) Test Brand Name/Manufacturer:
RESULT*: HIV-1 Ag HIV-Ab
□ Reactive □ Nonreactive □ Not Reported □ HIV-1 Reactive □ HIV-2 Reactive □ Both Reactive, Undifferentiated □ Both Nonreactive *Select one result for HIV-1 Ag and one result for HIV Ab
HIV Detection Tests (Qualitative)
TEST: ☐ HIV-1 RNA/DNA NAAT (Qual) ☐ HIV-1 Culture ☐ HIV-2 RNA/DNA NAAT (Qual) ☐ HIV-2 Culture
RESULT: □ Positive/Reactive □ Negative/Nonreactive □ Indeterminate Collection Date: / /
HIV Detection Tests (Quantitative viral load) Note: Include earliest test at or after diagnosis
TEST 1: □ HIV-1 RNA/DNA NAAT (Quantitative viral load) □ HIV-2 RNA/DNA NAAT (Quantitative viral load)
RESULT: Detectable Undetectable Copies/mL: Log: Collection Date://
TEST 2: □ HIV-1 RNA/DNA NAAT (Quantitative viral load) □ HIV-2 RNA/DNA NAAT (Quantitative viral load)
RESULT: Detectable Undetectable Copies/mL: Log: Collection Date://
Immunologic Tests (CD4 count and percentage)
CD4 at or closest to diagnosis: CD4 count:cells/µL CD4 percentage:% Collection Date://
First CD4 result <200 cells/μL or <14%: CD4 count:cells/μL CD4 percentage:% Collection Date://
Other CD4 result: CD4 count:cells/µL CD4 percentage:% Collection Date://
Documentation of Tests
Did documented laboratory test results meet approved HIV diagnostic algorithm criteria? ☐ Yes ☐ No ☐ Unknown If YES, provide specimen collection date of earliest positive test for this algorithm: / /
Complete the above only if none of the following was positive: HIV-1 Western blot, IFA, culture, viral load, or qualitative NAAT [RNA or DNA]
If HIV laboratory tests were not documented, is HIV diagnosis documented by a physician? Yes No Unknown If YES, provide date of diagnosis: / //
Date of last documented negative HIV test (before HIV diagnosis date):// Specify type of test:/

Clinical (record all dates as mm/dd/yyyy)

Diagnosis	Dx Date	Diagnosis	Dx Date	Diagnosis	Dx Date
Candidiasis, bronchi, trachea, or lungs		Herpes simplex: chronic ulcers (>1 mo. duration), bronchitis, pneumonitis, or esophagitis		M. tuberculosis, pulmonary [†]	
Candidiasis, esophageal		Histoplasmosis, disseminated or extrapulmonary		M. tuberculosis, disseminated or extrapulmonary [†]	
Carcinoma, invasive cervical		Isosporiasis, chronic intestinal (>1 mo. duration)		Mycobacterium, of other/unidentified species, disseminated or extrapulmonary	
Coccidioidomycosis, disseminated or extrapulmonary		Kaposi's sarcoma		Pneumocystis pneumonia	
Cryptococcosis, extrapulmonary		Lymphoma, Burkitt's (or equivalent)		Pneumonia, recurrent, in 12 mo. period	
Cryptosporidiosis, chronic intestinal (>1 mo. duration)		Lymphoma, immunoblastic (or equivalent)		Progressive multifocal leukoencephalopathy	
Cytomegalovirus disease (other than in liver, spleen, or nodes)		Lymphoma, primary in brain		Salmonella septicemia, recurrent	
Cytomegalovirus retinitis (with loss of vision)		Mycobacterium avium complex or M. kansasii, disseminated or extrapulmonary		Toxoplasmosis of brain, onset at >1 mo. of age	
HIV encephalopathy				Wasting syndrome due to HIV	

Treatment/Services Referrals (record all dates as mm/dd/yyyy) Has this patient been informed of his/her HIV infection? This patient's partners will be notified about their HIV exposure and counseled by: ☐ Yes ☐ No ☐ Unknown □ 1-Health Dept □ 2-Physician/Provider □ 3-Patient □ 9-Unknown **For Female Patient** Is this patient currently pregnant? This patient is receiving or has been referred for gynecological or Has this patient delivered live-born infants? obstetrical services: ☐ Yes ☐ No ☐ Unknown □ Yes □ No □ Unknown ☐ Yes ☐ No ☐ Unknown For Children of Patient (record most recent birth in these boxes; record additional or multiple births in Comments) *Child's Name Child's Last Name Child's Date of Birth Soundex *Child's Coded ID Child's State Number Facility Name of Birth (if child was born at home, enter "home birth") *Phone *ZIP Code Facility Type Inpatient: Outpatient: Other Facility: ☐ Emergency Room ☐ Hospital ☐ Other, specify _ □ Corrections □ Unknown ☐ Other, specify ___ □ Other, specify *Street Address County State/Country HIV Antiretroviral Use History (record all dates as mm/dd/yyyy) Main source of antiretroviral (ARV) use information (select one): Date patient reported information □ Medical Record Review □ Patient Interview ☐ Provider Report □ Other □ NHM&E Ever taken any ARVs? ☐ Yes ☐ No ☐ Unknown If yes, reason for ARV use (select all that apply): ☐ HIV Tx ARV medications: _ Date began: ___/__/___/ Date of last use: ___/__/__/ □ PrEP Date began: ___/__/___/ Date of last use: ___/__/__/ ARV medications: ___ □ PEP Date began: ___/__/___/ Date of last use: ___/__/___/ ARV medications: ___ Date of last use: ___/__/__/ □ PMTCT ARV medications: ___ Date began: ___/__/__/ Date began: __ /__ /__ /____ Date of last use: ___/__/__/ □ HBV Tx ARV medications: _____ □ Other ARV medications: Date began: ___/__/___/ Date of last use: __ /__ /__ HIV Testing History (record all dates as mm/dd/yyyy) Main source of testing history information (select one): Date patient reported information □ Patient Interview □ Medical Record Review □ Provider Report □ NHM&E □ Other ___/__/___ Date of first positive HIV test / / Ever had previous positive HIV test? ☐ Yes ☐ No ☐ Unknown Date of last negative HIV test (If date is from Ever had a negative HIV test? ☐ Yes ☐ No ☐ Unknown a lab test with test type, enter in Lab Data section) Number of negative HIV tests within 24 months before first positive test #_ □ Unknown Comments *Local/Optional Fields

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