Patient Identification (red	ord all dates a	s mm/dd/yyyy	<i>(</i>)				
*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 □ Post		out of a county	urrent Addres	ss, Street			Address Date
*Phone City		County		State/Country		*	ZIP Code
*Medical Record Number		*Othe	er ID Type	Social Secu	ırity * [*] •	lumber	
J.S. Department of Health & Human Services	(Patients ≥13 Years		Diagnosis) *I		ransmitted to C		Centers for Disease Contra and Prevention
Health Department Use 0		dates as mm	/dd/yyyy)		l l	oproved Oiv	IB 110. 0920-0373 Exp. 00/30/2019
Date Received at Health Departm	ent	eHARS Docur	ment UID		Stat	te Numbe	er
Reporting Health Dept - City/Cou	nty			City/County N	Number		
Document Source		Surveillance Me	ethod Activ	ve □ Passive □ F	Follow up □ R	eabstractio	n □ Unknown
Did this report initiate a new case ☐ Yes ☐ No ☐ Unknown	e investigation?	Report Medium		′isit □ 2-Mailed □ 5-Electronic Tı			ne
Facility Providing Informa	ntion (record al	l dates as mn	n/dd/yyyy)				
Facility Name					*Pho	ne (
*Street Address			-				
City	County		State/0	Country	* ZIP	Code	
Facility Inpatient: Type □ Hospital □ Other, specify	□ Adult I	e <u>nt:</u> □ Private Physici HV Clinic specify	A	creening, Diagnos gency: □ CTS Other, specify	☐ STD Clinic	□ Laborato	lity: □ Emergency Room ry □ Corrections □ Unknown pecify
Date Form Completed/	_/	*Person Complet	ing Form		*Pho	ne ()	
Patient Demographics (re	cord all dates	as mm/dd/yyy	у)				
Sex assigned at Birth ☐ Male ☐	☐ Female ☐ Unknov	vn Country of E	Birth □ US □	Other/US Deper	ndency (pleas	se specify)	
Date of Birth//			Alias Date of	f Birth/_	/	_	
Vital Status ☐ 1-Alive ☐ 2-Dead	ı	Date of Death	_//		State of Dea	ath	
	le □ Female □ Tra ditional gender ident		Female (MTF)	□ Transgender I	Female-to-Ma	le (FTM) [] Unknown
Ethnicity	o □ Not Hispanic/La	atino 🗆 Unknown			Expanded E	thnicity _	
	erican Indian/Alaska tive Hawaiian/Other			ican American Jnknown	Expanded R	lace	
Residence at Diagnosis (a	dd additional a	addresses in C	Comments)	(record all	dates as n	nm/dd/y	/yy)
Address Type (Check all that apply to address be	low) □ Residence	at HIV diagnosis	□ Residence a	at AIDS diagnosis	s □ Check if	SAME as	Current Address
*Street Address	,	. 3		1 10 10			Address Date
City	County		State/Cou	untry			*ZIP Code

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 ONLY					
*Provider Name (Last, First, M.I.)			*DI(
Hospital/Facility			*Phone ()		_
Facility of Diagnosis (add ad	ditional	facilities in Comme	 nts)		
Diagnosis Type (Check all that apply	y to facility	below) □ HIV □ AIDS	□ Check if <u>SAME as Facili</u>	ty Providing	g Information
Facility Name				*Phone ()
*Street Address					
City	County		State/Country		*ZIP Code
Facility Inpatient: ☐ Hospital Type ☐ Other, specify	☐ Adult F	<u>nt:</u> □ Private Physician's Office	Screening, Diagnostic, Referr □ CTS □ STD Clinic □ Other, specify		Other Facility: □ Emergency Room □ Laboratory □ Corrections □ Unknown □ Other, specify
*Provider Name		*Provider Phone ()		Specialty	
Patient History (respond to all	questio	ns) (record all dates as	mm/dd/yyyy) □ Pediatrio	c risk (pl	ease enter in Comments)
After 1977 and before the earliest known	own diagr	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		pecify clotting factor: ate received (mm/dd/yyyy):			□ Yes □ No □ Unknown
HETEROSEXUAL relations with any	of the follo	owing:			
HETEROSEXUAL contact with intrave	enous/injed	tion drug user			□ Yes □ No □ Unknown
HETEROSEXUAL contact with bisexu	al male				□ Yes □ No □ Unknown
HETEROSEXUAL contact with persor	n with hem	ophilia/coagulation disorder v	with documented HIV infection		□ Yes □ No □ Unknown
HETEROSEXUAL contact with transfu	usion recip	ient with documented HIV inf	ection		□ Yes □ No □ Unknown
HETEROSEXUAL contact with transp	lant recipie	ent with documented HIV infe	ection		□ Yes □ No □ Unknown
HETEROSEXUAL contact with persor	n with docu	mented HIV infection, risk no	ot specified		□ Yes □ No □ Unknown
Received transfusion of blood/blood cor	mponents	(other than clotting factor) (de	ocument reason in Comments))	□ Yes □ No □ Unknown
First date received / /	La	st date received/	_/		
Received transplant of tissue/organs or	artificial in	semination			□ Yes □ No □ Unknown
Worked in a healthcare or clinical laboral for occupational exposure is being investigation.			of exposure, specify occupatio	n and settir	□ Yes □ No □ Unknown
Other documented risk (please include	detail in C	omments)			☐ Yes ☐ No ☐ 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).

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:
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 Both (undifferentiated) Neither (negative) Indeterminate 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 Both (Ag and Ab reactive) Neither (negative) Invalid/Indeterminate 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 □ Reactive □ Nonreactive □ Not Reported Collection Date: / / = 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** This patient is receiving or has been referred for gynecological or Has this patient delivered live-born infants? Is this patient currently pregnant? 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 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: ___ Date of last use: ___/__/___/ □ PEP Date began: ___/__/___/ ARV medications: ___ □ PMTCT ARV medications: _____ Date began: ___/__/__/ Date of last use: __ /_ /_ _ /___ Date began: __ /__ /__ /____ Date of last use: ___/__/__/ □ HBV Tx ARV medications: _____ □ Other Date began: ___/__/___/ Date of last use: __ /_ /__ /_ ARV medications: 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 # Comments **Check OOS** State: *Local/Optional Fields NIR Status: PRISM # DOC# NIR OP NIR OP Date NIR CL NIR CL Date / Link with e-HARS stateno(s): NIR RE _ Other Risks: A B/C M NIR RE Date Initials (3) Source Code A Hepatitis: A Other **UNKnown** If pregnant, list EDD (due date)