

PATIENT INFORMATION						
Name of Patient (Last, First, MI)					Date of Report	
Patient Street Address			City	County	State	Zip Code
Patient Identifiers:	Medical Record Number	Prison ID Number	Patient ID Number	Other ID Type	ID No.	
Sex <input type="checkbox"/> Male <input type="checkbox"/> Female	Patient Birthdate ____ / ____ / ____	Ethnicity (Select One) <input type="checkbox"/> Hispanic <input type="checkbox"/> Not Hispanic <input type="checkbox"/> Unknown	Race (Select one or more) <input type="checkbox"/> Amer. Indian/Alaska Native <input type="checkbox"/> Asian <input type="checkbox"/> White	<input type="checkbox"/> Black/African American <input type="checkbox"/> Native Hawaiian or Other Pac. Isl. <input type="checkbox"/> Unknown	Patient Pregnant <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> Not applicable	
NAME OF FACILITY OR PROVIDER PRACTICE THAT ORDERED TESTS						
Name of Facility/Provider					Accession Number	
Facility/Provider Full Address					Name of Contact Person	
City			State	Zip Code	Main Telephone Number	
NAME OF LABORATORY						
Name of Laboratory					CLIA Code	
Street Address					Name of Contact Person	
City			State	Zip Code	Telephone Number	

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

HIV Immunoassays (Nondifferentiating)	
TEST 1 <input type="checkbox"/> HIV-1 IA <input type="checkbox"/> HIV-1/2 IA <input type="checkbox"/> HIV-1/2 Ag/Ab <input type="checkbox"/> HIV-1 WB <input type="checkbox"/> HIV-1 IFA <input type="checkbox"/> HIV-2 IA <input type="checkbox"/> HIV-2 WB	
Test brand name/Manufacturer _____	Accession No. _____
Facility name _____	Specimen Type _____
Result <input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Indeterminate	Collection Date ____ / ____ / ____ <input type="checkbox"/> Point-of-care rapid test
TEST 2 <input type="checkbox"/> HIV-1 IA <input type="checkbox"/> HIV-1/2 IA <input type="checkbox"/> HIV-1/2 Ag/Ab <input type="checkbox"/> HIV-1 WB <input type="checkbox"/> HIV-1 IFA <input type="checkbox"/> HIV-2 IA <input type="checkbox"/> HIV-2 WB	
Test brand name/Manufacturer _____	Accession No. _____
Facility name _____	Specimen Type _____
Result <input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Indeterminate	Collection Date ____ / ____ / ____ <input type="checkbox"/> Point-of-care rapid test
HIV Immunoassays (Differentiating)	
<input type="checkbox"/> HIV-1/2 type-differentiating immunoassay (differentiates between HIV-1 Ab and HIV-2 Ab)	Role of test in diagnostic algorithm <input type="checkbox"/> Screening/initial test <input type="checkbox"/> Confirmatory/supplemental test
Test brand name/Manufacturer _____	Accession No. _____
Facility name _____	Specimen Type _____
Result ¹ Overall interpretation: <input type="checkbox"/> HIV-1 positive <input type="checkbox"/> HIV-2 positive <input type="checkbox"/> HIV positive, untypable <input type="checkbox"/> HIV-2 positive with HIV-1 cross-reactivity <input type="checkbox"/> HIV-1 indeterminate <input type="checkbox"/> HIV-2 indeterminate <input type="checkbox"/> HIV indeterminate <input type="checkbox"/> HIV negative	
Analyte results: HIV-1 Ab: <input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Indeterminate <input type="checkbox"/> Point-of-care rapid test HIV-2 Ab: <input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Indeterminate ¹ Always complete the overall interpretation. Complete the analyte results when available.	
<input type="checkbox"/> HIV-1/2 Ag/Ab differentiating immunoassay (differentiates between HIV Ag and HIV Ab)	
Test brand name/Manufacturer _____	Accession No. _____
Facility name _____	Specimen Type _____
Result <input type="checkbox"/> Ag positive <input type="checkbox"/> Ab positive <input type="checkbox"/> Both (Ag and Ab positive) <input type="checkbox"/> Negative <input type="checkbox"/> Invalid	Collection Date ____ / ____ / ____ <input type="checkbox"/> Point-of-care rapid test
<input type="checkbox"/> HIV-1/2 Ag/Ab and type-differentiating immunoassay (differentiates among HIV-1 Ag, HIV-1 Ab, and HIV-2 Ab)	
Test brand name/Manufacturer _____	Accession No. _____
Facility name _____	Specimen Type _____
Result ² Overall interpretation: <input type="checkbox"/> Reactive <input type="checkbox"/> Nonreactive <input type="checkbox"/> Index value _____	
Analyte results: HIV-1Ag: <input type="checkbox"/> Reactive <input type="checkbox"/> Nonreactive <input type="checkbox"/> Not reportable due to high Ab level <input type="checkbox"/> Index value _____ HIV-1 Ab: <input type="checkbox"/> Reactive <input type="checkbox"/> Nonreactive <input type="checkbox"/> Reactive undifferentiated <input type="checkbox"/> Index value _____ HIV-2 Ab: <input type="checkbox"/> Reactive <input type="checkbox"/> Nonreactive <input type="checkbox"/> Reactive undifferentiated <input type="checkbox"/> Index value _____	
Collection Date ____ / ____ / ____	<input type="checkbox"/> Point-of-care rapid test ² Complete the overall interpretation and the analyte results.
HIV Detection Tests (Qualitative)	
TEST <input type="checkbox"/> HIV-1 RNA/DNA NAAT (Qualitative) <input type="checkbox"/> HIV-1 culture <input type="checkbox"/> HIV-2 RNA/DNA NAAT (Qualitative) <input type="checkbox"/> HIV-2 culture	
Test brand name/Manufacturer _____	Accession No. _____
Facility name _____	Specimen Type _____
Result <input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Indeterminate	Collection Date ____ / ____ / ____
HIV Detection Tests (Quantitative viral load) Note: Include earliest test at or after diagnosis.	
TEST 1 <input type="checkbox"/> HIV-1 RNA/DNA NAAT (Quantitative viral load) <input type="checkbox"/> HIV-2 RNA/DNA NAAT (Quantitative viral load)	
Test brand name/Manufacturer _____	Accession No. _____
Facility name _____	Specimen Type _____
Result <input type="checkbox"/> Detectable <input type="checkbox"/> Undetectable <input type="checkbox"/> Copies/mL _____	Log _____ Collection Date ____ / ____ / ____
TEST 2 <input type="checkbox"/> HIV-1 RNA/DNA NAAT (Quantitative viral load) <input type="checkbox"/> HIV-2 RNA/DNA NAAT (Quantitative viral load)	
Test brand name/Manufacturer _____	Accession No. _____
Facility name _____	Specimen Type _____
Result <input type="checkbox"/> Detectable <input type="checkbox"/> Undetectable <input type="checkbox"/> Copies/mL _____	Log _____ Collection Date ____ / ____ / ____

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

Drug Resistance Tests (Genotypic)			
TEST <input type="checkbox"/> HIV-1 Genotype (Unspecified)		Test brand name/Manufacturer _____	
Lab name _____		Accession No. _____	
Provider name _____		Specimen Type _____	
		Collection Date ____ / ____ / ____	
Immunologic Tests (CD4 count and percentage)			
CD4 at or closest to diagnosis: CD4 count _____ cells/μL		CD4 percentage _____ %	Collection Date ____ / ____ / ____
Test brand name/Manufacturer _____		Accession No. _____	
Facility name _____		Specimen Type _____	
First CD4 result <200 cells/μL or <14%: CD4 count _____ cells/μL		CD4 percentage _____ %	Collection Date ____ / ____ / ____
Test brand name/Manufacturer _____		Accession No. _____	
Facility name _____		Specimen Type _____	
Other CD4 result: CD4 count _____ cells/μL		CD4 percentage _____ %	Collection Date ____ / ____ / ____
Test brand name/Manufacturer _____		Accession No. _____	
Facility name _____		Specimen Type _____	
Documentation of Tests			
Did documented laboratory test results meet approved HIV diagnostic algorithm criteria? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown			
If YES, provide specimen collection date of earliest positive test for this algorithm ____ / ____ / ____			
<i>Complete the above only if none of the following were positive for HIV-1: Western blot, IFA, culture, viral load, qualitative NAAT (RNA or DNA), HIV-1/2 type-differentiating immunoassay (supplemental test), stand-alone p24 antigen, or nucleotide sequence.</i>			
If HIV laboratory tests were not documented, is HIV diagnosis documented by a physician? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown			
If YES, provide date of diagnosis ____ / ____ / ____			
Date of last documented negative HIV test (before HIV diagnosis date) ____ / ____ / ____			
Specify type of test: _____			