



State regulations require reporting of all HIV infections diagnosed or treated in New Mexico.

Reports may be securely faxed to 505-827-0013, or mailed to:

New Mexico Department of Health  
1190 St. Francis Dr., N 1359  
Santa Fe, NM 87502-6110  
Attn: Surveillance Coordinator

## 1. PROVIDER/FACILITY INFORMATION

▲ PERSON COMPLETING FORM

▲ PHONE

▲ DATE COMPLETED

▲ PHYSICIAN

▲ PHYSICIAN PHONE

▲ FACILITY NAME

▲ FACILITY PHONE

▲ FACILITY ADDRESS

▲ CITY/STATE/ZIP

### FACILITY TYPE

INPATIENT      OUTPATIENT

☐ HOSPITAL    ☐ PRIVATE PHYSICIAN

☐ OTHER        ☐ ADULT HIV CLINIC

☐ OTHER: \_\_\_\_\_

### SCREENING, DIAGNOSTIC, REFERRAL AGENCY

☐ STD CLINIC    ☐ OTHER

OTHER FACILITY ☐ ER    ☐ LAB    ☐ CORRECTIONS    ☐ UNKNOWN

☐ OTHER: \_\_\_\_\_

## CONFIDENTIAL PROVIDER HIV/AIDS ADULT CASE REPORT

### 2. PATIENT INFORMATION

▲ PATIENT LAST NAME      ▲ FIRST NAME      ▲ MIDDLE NAME

▲ AKA (CHOSEN NAME, PREFERRED NAME, NICKNAME, PREVIOUS LAST NAME, ETC.)

ADDRESS TYPE    ☐ RESIDENTIAL      ☐ HOMELESS  
☐ CORRECTIONAL FACILITY    ☐ POSTAL  
☐ FOSTER HOME      ☐ TEMPORARY  
☐ MILITARY      ☐ OTHER

▲ CURRENT STREET ADDRESS

▲ CITY      ▲ STATE      ▲ ZIP CODE

▲ PHONE NUMBER

▲ DATE OF BIRTH

▲ SOCIAL SECURITY NUMBER

▲ MEDICAL RECORD NUMBER

VITAL STATUS    ☐ ALIVE    ☐ DEAD

▲ DATE OF DEATH

▲ STATE OF DEATH

STATUS    ☐ HIV    ☐ AIDS

COUNTRY OF BIRTH    ☐ U.S.    ☐ OTHER/U.S. DEPENDENCY

#### ▲ SPECIFY

SEX ASSIGNED AT BIRTH    ☐ MALE    ☐ FEMALE    ☐ OTHER

#### CURRENT GENDER IDENTITY

☐ MALE    ☐ TRANSGENDER MAN (FEMALE-TO-MALE)

☐ FEMALE    ☐ TRANSGENDER WOMAN (MALE-TO-FEMALE)

☐ NON-BINARY/GENDER NONCONFORMING    ☐ UNKNOWN

☐ TWO-SPIRIT    ☐ ADDITIONAL GENDER IDENTITY: \_\_\_\_\_

#### ▲ SPECIFY

RACE (CHECK ALL THAT APPLY):    ☐ WHITE    ☐ BLACK    ☐ ASIAN

☐ AMERICAN INDIAN/ALASKAN NATIVE    ☐ PACIFIC ISLANDER

ETHNICITY    ☐ HISPANIC/LATINX    ☐ NOT HISPANIC/LATINX

☐ UNKNOWN

▲ EXPANDED RACE/ETHNICITY

### 3. RESIDENCE/FACILITY AT HIV/AIDS DIAGNOSIS

☐ Check if patient address/facility at HIV diagnosis are same as current (if checked, leave the rest of this section blank)

▲ ADDRESS AT TIME OF DIAGNOSIS IF DIFFERENT THAN CURRENT ADDRESS:

▲ FACILITY OF HIV DIAGNOSIS

▲ PHONE

▲ FACILITY ADDRESS

▲ CITY/STATE/ZIP

#### FACILITY TYPE

INPATIENT    ☐ HOSPITAL      ☐ OTHER

OUTPATIENT    ☐ PRIVATE PHYSICIAN    ☐ ADULT HIV CLINIC

☐ OTHER: \_\_\_\_\_

SCREENING, ETC    ☐ STD CLINIC    ☐ OTHER: \_\_\_\_\_

OTHER

☐ ER

☐ LAB

☐ CORRECTIONS

☐ UNKNOWN

▲ EARLIEST HIV DIAGNOSIS

EVER PROGRESSED TO STAGE-3?    ☐ YES    ☐ NO

### 4. PATIENT HISTORY & RISK FACTORS

#### CHECK ALL THAT APPLY:

SEX WITH MALE      ☐ YES    ☐ NO    ☐ UNKNOWN

SEX WITH FEMALE      ☐ YES    ☐ NO    ☐ UNKNOWN

INJECTION DRUG USE      ☐ YES    ☐ NO    ☐ UNKNOWN

PERINATAL INFECTION WITH HIV      ☐ YES    ☐ NO    ☐ UNKNOWN

#### HETEROSEXUAL RELATIONS WITH:

INJECTION DRUG USER      ☐ YES    ☐ NO    ☐ UNKNOWN

BISEXUAL MALE      ☐ YES    ☐ NO    ☐ UNKNOWN

PERSON /DOCUMENTED HIV/AIDS    ☐ YES    ☐ NO    ☐ UNKNOWN

▲ OTHER DOCUMENTED RISK (SPECIFY): \_\_\_\_\_

PLEASE COMPLETE  
BOTH SIDES OF THIS FORM.

## 5. CLINICAL: ACUTE HIV INFECTION AND OPPORTUNISTIC ILLNESSES

SUSPECT ACUTE HIV? ☐ YES ☐ NO ☐ UNKNOWN

Clinical signs/symptoms consistent with acute retroviral

syndrome? (e.g., fever, malaise/fatigue, myalgia, pharyngitis, rash, lymphadenopathy) ☐ YES ☐ NO ☐ UNKNOWN

▲ IF YES, DATE OF SIGN/SYMPTOM ONSET

▲ OPPORTUNISTIC ILLNESS

▲ DIAGNOSIS DATE

## 6. PREGNANCY

IS PATIENT CURRENTLY PREGNANT? ☐ YES ☐ NO ☐ UNKNOWN

▲ EXPECTED DELIVERY DATE

IS PATIENT IN PRE-NATAL CARE? ☐ YES ☐ NO ☐ UNKNOWN

## 7. HIV TESTS

### DOCUMENTATION OF TESTS

**Required:** Attach copies of all relevant laboratory results for HIV diagnosis and indicate that labs are attached:

☐ **Labs are attached** (If checked, the results fields in this section can be left blank)

▲ DATE/LAST DOCUMENTED NEGATIVE HIV TEST (BEFORE HIV DIAGNOSIS DATE)

▲ SPECIFY TYPE OF TEST

If HIV lab tests were **not** documented, is HIV diagnosis confirmed by a clinician? ☐ YES ☐ NO ☐ UNKNOWN

▲ IF YES, DATE DOCUMENTATION BY CARE PROVIDER

Was the first positive test from a self-test performed by the patient? ☐ YES ☐ NO ☐ UNKNOWN

### HIV IMMUNOASSAYS (NON-DIFFERENTIATING)

#### ☐ HIV-1/2 AG/AB

▲ COLLECTION DATE

☐ RAPID TEST

☐ POS/REACTIVE ☐ NEG/NON-REACTIVE ☐ INDETERMINATE

## 7. HIV TESTS (CONTINUED)

### ☐ HIV-1/2 RNA/NDA NAAT (QUAL)

▲ COLLECTION DATE

☐ POS/REACTIVE ☐ NEG/NON-REACTIVE ☐ INDETERMINATE

▲ COLLECTION DATE, MOST RECENT HIV VIRAL LOAD

CHOOSE ONE: < = >

▲ COPIES/ML

▲ LOG

▲ COLLECTION DATE, MOST RECENT CD4 ☐ COUNT (CELLS/  $\mu$ L) ☐ %

▲ COLLECTION DATE, FIRST CD4 <200  $\mu$ L ☐ COUNT (CELLS/  $\mu$ L) ☐ %

### HIV IMMUNOASSAYS (TYPE-DIFFERENTIATING)

#### ☐ HIV-1/2 AG/AB AND TYPE DIFFERENTIATING

▲ COLLECTION DATE

☐ RAPID TEST

Overall Interpretation

☐ REACTIVE

☐ NON-REACTIVE

HIV-1 AG

☐ REACTIVE

☐ NON-REACTIVE

HIV-1 AB

☐ REACTIVE

☐ NON-REACTIVE

☐ REACTIVE, NON-DIFFERENTIATING

#### ☐ HIV-1/2 TYPE DIFFERENTIATING

▲ COLLECTION DATE

Role of test in diagnostic algorithm:

☐ SCREENING/INITIAL ☐ CONFIRMATORY/SUPPLEMENTAL

☐ RAPID TEST

Overall Interpretation:

☐ HIV-1 POSITIVE ☐ HIV-2 POSITIVE ☐ HIV NEGATIVE

☐ HIV POSITIVE, UNTYPABLE

☐ HIV-2 POSITIVE WITH HIV-1 CROSS-REACTIVITY

## 8. HIV TESTING & TREATMENT HISTORY

Ever taken **any** antiretroviral medications (ARVs)?

☐ YES ☐ NO ☐ UNKNOWN

IF YES, REASON FOR ARV USE (SELECT ALL THAT APPLY):

☐ FOR **HIV TREATMENT?** ☐ YES ☐ NO ☐ UNKNOWN

▲ ARV MED

▲ DATE BEGUN

▲ DATE OF LAST USE

☐ FOR **PrEP?**

☐ YES

☐ NO

☐ UNKNOWN

▲ ARV MED

▲ DATE BEGUN

▲ DATE OF LAST USE

☐ FOR **PEP?**

☐ YES

☐ NO

☐ UNKNOWN

▲ ARV MED

▲ DATE BEGUN

▲ DATE OF LAST USE

☐ FOR **PREGNANCY?**

☐ YES

☐ NO

☐ UNKNOWN

▲ ARV MED

▲ DATE BEGUN

▲ DATE OF LAST USE

☐ **OTHER**

☐ YES

☐ NO

☐ UNKNOWN

▲ ARV MED

▲ DATE BEGUN

▲ DATE OF LAST USE

The primary objective of HIV surveillance is to identify emerging trends so that prevention and control measures can be applied to effectively minimize disease burden. The data collected also help set priorities and develop targeted interventions for all affected by HIV. To meet these needs, NMDOH relies upon timely and complete reporting by all providers.

Any medical provider, laboratory, or organization that offers HIV testing by name (confidential testing) or provides care to persons with HIV infection must report.

Questions about this form or requests for data can be directed to NMDOH at (505) 699-2912.