

HPTN Protocols Snapshot:

| Protocol # | Title | Sites | Study Status | IND | Research Area | Study Population | Open to Accrual (projected) | 1 st Participant Enrolled (projected) | Closed to Accrual (projected) | Closed to Follow Up (projected) | Target Accrual | Actual Accrual as of report date |
|------------------------------|--|---------|-------------------|-----|------------------------|---------------------------------------|-----------------------------------|--|-------------------------------------|---------------------------------------|-------------------|--|
| HPTN 119 | DoxyPEP Africa | INTL | In Development | N | Integrated Strategy | TBD | TBD | TBD | TBD | TBD | TBD | N/A |
| HPTN 117 | PK and adherence of Doxycycline | US | In Development | N | Integrated Strategy | Healthy Adults | March 2026 | March 2026 | Jan 2027 | TBD | 16 | N/A |
| DMID 24- 0020/HPTN 116 | Neurosyphilis Study | TBD | In Development | N | STI | TBD | Aug 2025 | Aug 2025 | Dec 2028 | Dec 2029 | TBD | N/A |
| HPTN 115/ATN 173 | Doxycycline Prophylaxis for Prevention of Sexually Transmitted Infections Among Adolescent and Young Women in the United States | us | In Development | N | PEP | Adolescents Women | Feb 2025 | Feb 2025 | June 2026 | June 2027 | 760 | N/A |
| HPTN 113 | Duo: A Phase IIIb Individual-Level Randomized Controlled Trial of an Integrated Strategy of HIV PrEP and STI PEP for Young Men who have Sex with Men in the Americas | US/INTL | Pending | N | Integrated Strategy | Young Men who have Sex with Men | Jan 2026 | Jan 2026 | June 2027 | June 2028 | 500 | N/A |

Updated: 3 November 2025

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| HVTN 141/HPTN 105 | A phase 1 clinical trial to evaluate the safety, tolerability, pharmacokinetics and in vitro neutralization of VRC01.23LS, ePGDM1400v9-LS and ePGT121v1-LS alone and in combination in healthy, HIV-uninfected adult participants | TBD | Pending | Y | PrEP | HIV Uninfected Adults | Nov 2025 | Nov 2025 | May 2026 | June 2027 | 136 | N/A |
| HPTN 104 | To evaluate adherence to a single dual prevention pill, DPP (co-formulated TDF/FTC + ethynyl estradiol/levonorgestrel oral contraceptive pill (OCP), compared with the two tablets with daily oral TDF/FTC + ethynyl estradiol/levonorgestrel oral contraceptive pill for pre-exposure prophylaxis and pregnancy prevention in HIV-uninfected women | INTL | Pending | Y | МРТ | HIV uninfected women | TBD | TBD | TBD | TBD | 300 | N/A |
| HPTN 096 Post Pilot | Testing the Efficacy of an Integrated Intervention Strategy in the American South | US | Pending | N | Integrated Strategy | MSM in the Southern US | Nov 2025 | Nov 2025 | Sept 2028 | Sept 2028 | N/A | N/A |

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| HVTN 206/HPTN 114 | A phase 2 clinical trial to evaluate the safety, tolerability, pharmacokinetics and neutralization of VRC07-523LS, PGT121.414.LS and PGDM1400LS broadly neutralizing monoclonal antibodies in adult participants without HIV and in overall good health. | US/INTL | Enrolling | Y | mAb | HIV uninfected adults | 14 Jan 2025 | 7 Mar 2025 | Nov 2025 | Nov 2026 | 200 | 106 |
| HPTN 106 | A Phase 2 Crossover Study Of On-Demand Prep Formulations Comparing Rectal And Oral Tenofovir- Based Prep Evaluating Extended Safety, Acceptability, And Pharmacokinetics/Pharma codynamics | US | Enrolling | Υ | PrEP | HIV Uninfected MSM | 25 Sept 2024 | 29 Oct 2024 | Mar 2026 | Aug 2026 | 150 | 131 |
| A5416/HVTN 806/HPTN 108 | A Phase I, Open-Label Study of the Safety, Antiviral & Immunomodulatory of Broadly Neutralizing Antibodies 3BNC117-LS-J and 10-1074-LS-J in Combination in ART- treated Adults in sub- Saharan Africa Living with HIV during a Monitored Analytical Treatment Interruption | INTL | Closed to Accrual | TBD | Antibody Mediated Prevention | Adult participants living with HIV | 26 Apr 2024 | 28 May 2024 | 5 Feb 2025 | Mar 2026 | 48 | 32 |

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| HPTN 107 | A Phase II randomized, observer-blind, placebo- controlled study, to assess efficacy of meningococcal Group B vaccine rMenB+OMV NZ (Bexsero) in preventing gonococcal infection. | US | Closed to Accrual | Y | STI | Adults at risk of STI | 19 Nov 2020 | 29 Dec 2020 | Late 2024 | Feb 2026 | 2200 | 2606 |
| HPTN 103 | A Phase 2, Open-Label, Multicenter, Randomized Clinical Trial to Evaluate the Feasibility, Safety, and Acceptability of Long- Acting Subcutaneous Lenacapavir vs. Daily Oral Emtricitabine/Tenofovir Disoproxil Fumarate for Pre-Exposure Prophylaxis Among People who Inject Drugs | US | Closed to Accrual | Y | PrEP | HIV Uninfected PWID | 4 June 2024 | 20 Aug 2024 | 1 Aug 2025 | May 2028 | 180 | 181 |
| HPTN 102 | A Phase 2, Open-Label, Multicenter, Randomized Study to Evaluate the Pharmacokinetics, Safety, and Acceptability and Use of Twice Yearly Long- Acting Subcutaneous Lenacapavir for Pre- Exposure Prophylaxis Among Women in the United States | US | Closed to Accrual | Y | PrEP | HIV Uninfected Women | 23 Apr 2024 | 31 May 2024 | 29 July 2025 | Jan 2028 | 250 | 253 |

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| HPTN 084 Pregnancy | A Phase 3 Double Blind Safety and Efficacy Study of Long-Acting Injectable Cabotegravir Compared to Daily Oral TDF/FTC for Pre- Exposure Prophylaxis in HIV-Uninfected Women. | INTL | Closed to Accrual | Υ | PrEP | HIV-uninfected women | 1 Jan 2022 | 1 Jan 2022 | 5 Apr 2025 | Dec 2025 | N/A | 462 |
| HPTN 113-01 | Focus Group Discussion prior to HPTN 113 | US/INTL | Closed to Follow Up | N | Integrated Strategy | Men who have Sex with Men | 26 Sept 2024 | 26 Sept 2024 | 30 Apr 2025 | 31 May 2025 | | 63 |
| HTPN 112 | Improving HIV prevention among heterosexual men seeking STI services in sub-Saharan Africa: examining the feasibility, acceptability, and associated costs of a systems-navigator-delivered integrated prevention package. | INTL | Closed to Follow Up | N | Integrated Strategy | Heterosexual Men | 21 Mar 2024 | 2 Apr 2024 | 15 Nov 2024 | 1 Sept 2025 | 200 | 203 |
| HPTN 111 | Uptake of HIV Self-testing and Linkage to Prevention and Care among Heterosexual Men attending Barbershops in Uganda: A Cluster Randomized Trial. | INTL | Closed to Follow Up | N | Integrated Strategy | Heterosexual Men | 10 Mar 2024 | 13 Mar 2024 | 27 Jun 2024 | 30 June 2025 | 250 | 249 |

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| HVTN 140/HPTN 101 | A phase 1 dose-escalation clinical trial to evaluate the safety, tolerability, and pharmacokinetics of PGDM1400LS alone and in combination with VRC07-523LS and PGT121.414.LS in healthy, HIV-uninfected adult participants | US/INTL | Closed to Follow Up | Υ | Antibody Mediated Prevention | HIV Uninfected Adults | 20 Oct 2021 | 15 Nov 2021 | 5 Oct 2022 | 19 July 2023 | 95 | 95 |
| HVTN 804/HPTN 095 | Antiretroviral analytical treatment interruption (ATI) to assess immunologic and virologic responses in participants who received VRC01 or placebo and became HIV-infected during HVTN 704/HPTN 085 | US/INTL | Closed to Follow-Up | N | Antibody Mediated Prevention | HIV-infected MSM | 5 Feb 2020 | 22 Aug 2022 | 26 Jun 2023 | 22 July 2024 | 46 | 18 |
| HVTN 805/HPTN 093 | Antiretroviral analytical treatment interruption (ATI) to assess immunologic and virologic responses in participants who received VRC01 or placebo and became HIV infected during HVTN 703/HPTN 081 | INTL | Closed to Follow-Up | N | Antibody Mediated Prevention | HIV-infected women | 02 Apr 2021 | 28 May 2021 | 14 Sept 2022 | 2 Feb 2024 | 61 | 13 |

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| HPTN 091 | Integrating HIV Prevention, Gender-Affirmative Medical Care, and Peer Health Navigation to Prevent HIV Acquisition and HIV Transmission for Transgender Women in the Americas: A Vanguard Feasibility and Acceptability Study | US/INTL | Closed to Follow-Up | N | Integrated Strategy | Transgender Women | 24 Feb 2021 | 26 Mar 2021 | 16 Dec 2022 | 16 Aug 2024 | 310 | 307 |
| HVTN 130/HPTN 089 | A Phase I clinical trial to evaluate the safety, pharmacokinetics, and functional activity of a combination of VRC07-523LS, PGT121, and PGDM1400 in healthy, HIV-1 uninfected adult participants. | us | Closed to Follow-Up | Υ | Antibody mediated prevention | HIV-uninfected adults | 17 July 2019 | 31 Jul 2019 | 17 Dec 2019 | 25 Mar 2021 | 27 | 27 |
| HPTN 084-01 | Safety, Tolerability and Acceptability of Long- Acting Cabotegravir (CAB LA) for the Prevention of HIV among Adolescent Females – A Sub-study of HPTN 084 | INTL | Closed to Follow-Up | Υ | PrEP | HIV-uninfected adolescents | 4 Nov 2020 | 3 Dec 2020 | 6 Aug 2021 | 10 Jan 2023 | 55 | 55 |
| HPTN 083-01 | Safety, Tolerability and Acceptability of Long- Acting Cabotegravir (CAB LA) for the Prevention of HIV among Adolescent Males – A sub-study of HPTN 083 | US | Closed to Follow Up | Υ | PrEP | HIV-uninfected adolescents | 19 Feb 2020 | 6 July 2020 | 10 Jan 2022 | 7 July 2023 | 55 | 9 |

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| HPTN 083 OLE | Phase 3 study of Cabotegravir LA vs daily oral TDF/FTC for PrEP in HIV uninfected men who have sex with men | US/INTL | Closed to Follow Up | Υ | PrEP | HIV-uninfected MSM | 5 Dec 2016 | 19 Dec 2016 | 16 Mar 2020 | 31 Mar 2025 | N/A | 2278 |
| HPTN 083 | Phase 3 study of Cabotegravir LA vs daily oral TDF/FTC for PrEP in HIV uninfected men who have sex with men | US/INTL | Closed to Follow Up | Y | PrEP | HIV-uninfected MSM | 5 Dec 2016 | 19 Dec 2016 | 16 Mar 2020 | 31 Mar 2025 | 5000 | 4570 |
| HVTN 136/HPTN 092 | A phase 1 dose-escalation clinical trial to evaluate the safety, tolerability, pharmacokinetics, and antiviral activity of the monoclonal antibody PGT121.414.LS administered alone and in combination with VRC07-523LS via intravenous infusion or via subcutaneous injections in healthy, HIV-uninfected adult participants | US | Participants of Study/Primary Analysis Complete | Y | Antibody Mediated Prevention | HIV-uninfected adults | 24 Aug 2020 | 10 Nov 2020 | 05 Oct 2021 | 18 Jan 2023 | 32 | 33 |
| HPTN 084 Blinded | A Phase 3 Double Blind Safety and Efficacy Study of Long-Acting Injectable Cabotegravir Compared to Daily Oral TDF/FTC for Pre- Exposure Prophylaxis in HIV-Uninfected Women. | INTL | Participants of Study/Primary Analysis Complete | Y | PrEP | HIV-uninfected women | 7 Nov 2017 | 27 Nov 2017 | 8 Nov 2020 | 17 Oct 2022 | 3200 | 3224 |