Spotlight on New PrEP Tools and Data

From R&D to Access

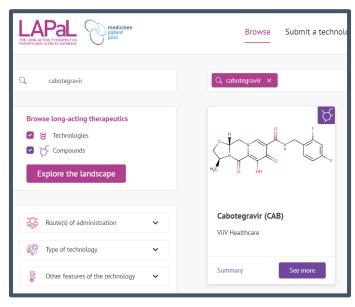
28 November 2023

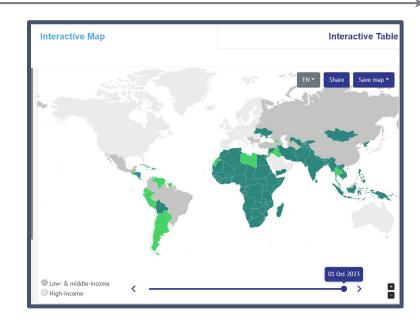




Welcome!











ACCESS TO MEDICINES TRACKER







Agenda

Welcome	Mitchell Warren, AVAC Cherise Scott, Unitaid
The Current PrEP Landscape	Catherine Verde Hashim, AVAC Sébastien Morin, MPP
PrEPWatch	Janki Tailor, AVAC Catherine Verde Hashim, AVAC
The Long-Acting Therapeutics Patents and Licenses Database	Lobna Gaayeb, MPP
Access to Medicines Tracker	Sébastien Morin, MPP
When do I use each tool?	Janki Tailor and Catherine Verde Hashim, AVAC Lobna Gaayeb, MPP
Interactive Quiz	Wawira Nyagah, AVAC Lobna Gaayeb, MPP
Q&A	Anne-Isabelle Cameron, Unitaid







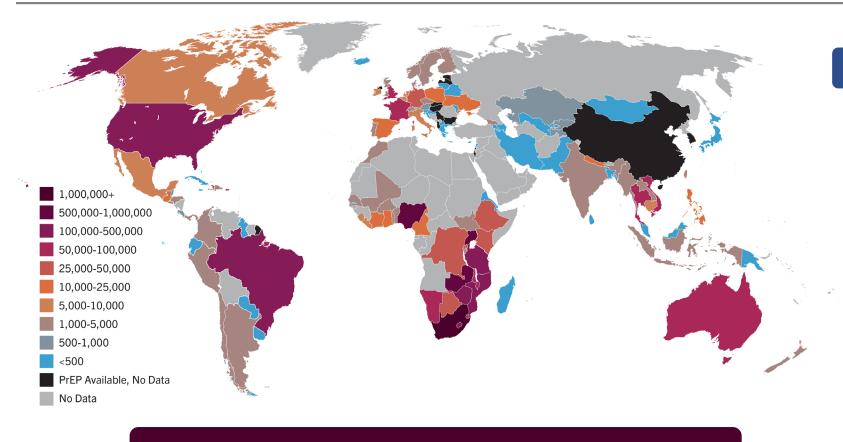
The Current PrEP Landscape



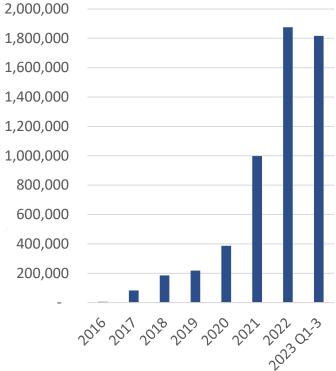




The PrEP Landscape - Initiations



PrEP Initiations Over Time



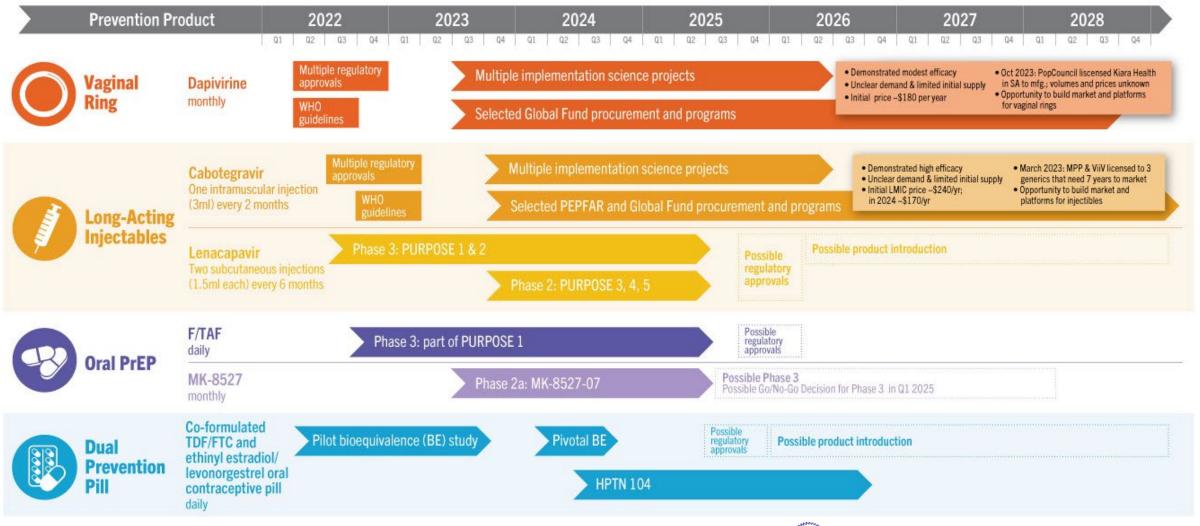
5.7m initiations to Sept 2023







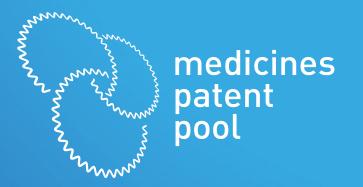
The PrEP Landscape - Pipeline











Access to generic CAB-LA for PrEP: Update on licensing and outlook for broad access

Webinar – Spotlight on new PrEP tools and data: From R&D to access
Convened by AVAC, MPP, and Unitaid on 28 November 2023

Presented by Sébastien Morin (Medicines Patent Pool)

MEDICINESPATENTPOOL.ORG

Spotlight on New PrEP Tools and Data: FROM R&D TO ACCESS



TUESDAY, NOVEMBER 28
8:30 TO 10:00 AM ET / 14:30 TO 16:00 PM CET

Join us to learn about three important online PrEP resources and to understand how they can support and enhance your work.

Register at tinyurl.com/PrEPTools





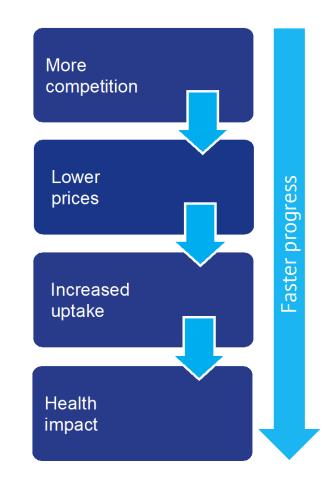


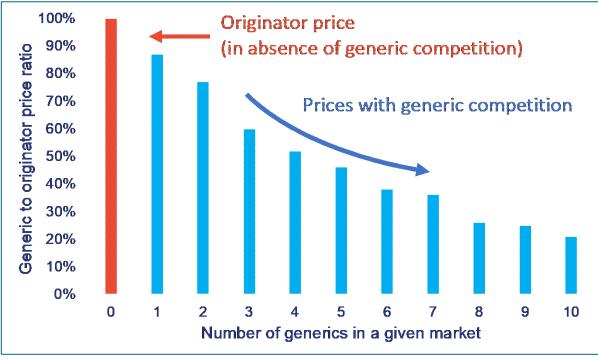


Accelerated access to generics and price reductions

Overview of the voluntary licensing model

- Driving prices down through generic competition
- Accelerating timelines for generic product availability





Generic competition induced pricing benchmarks: Dave et al. (2017) NEJM



Access to MPP-licensed medicines

We will present some of these tools today...

Patent and licence information tools

- To support procurement agencies, national programmes, and treatment advocates
- Fully public quarterlyupdated country-bycountry information



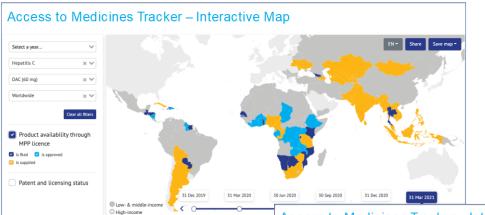
www.medspal.org

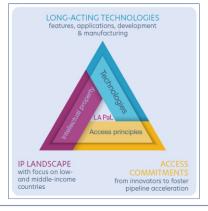


LAPaL
THE LONG-ACTING THERAPEUTICS
PATENTS AND LICENCES DATABASE

www.vaxpal.org

https://lapal.medicinespatentpool.org





Access to Medicines Tracker – Interactive Table



More at: https://medicinespatentpool.org/what-we-do/medspal
https://medicinespatentpool.org/progress-achievements/access-to-medicines-tracker



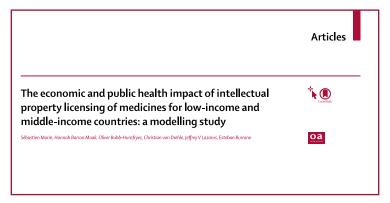
To learn more about the work of MPP...

MPP strategy for 2023-2025

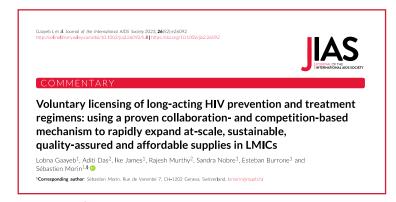


Greater access to medicines and health technologies for those who need them, 2023 https://medicinespatentpool.org/what-we-do/strategy

Peer-reviewed publications



The Lancet Public Health, 2022 https://doi.org/10.1016/S2468-2667(21)00202-4



Journal of the International AIDS Society, 2023 https://doi.org/10.1002/jia2.26092 https://www.youtube.com/watch?v=h57gXs4aQtg



The Lancet Global Health, 2023 https://doi.org/10.1016/S2214-109X(22)00460-0



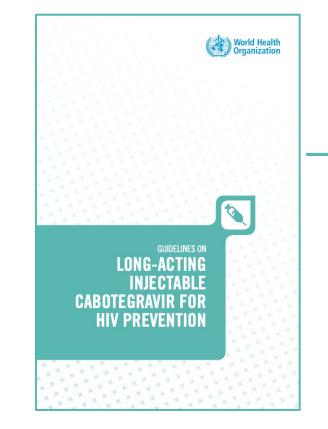


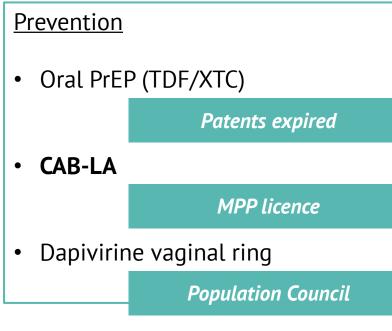
CAB-LA as one more HIV prevention option

"CAB-LA should be delivered as an additional choice alongside other PrEP options, including oral PrEP and the DVR, as part of a comprehensive HIV prevention approach."

CAB-LA for PrEP is

- ✓ Recommended by WHO
- Approved in several countries
- Effective, well tolerated, safe
- ✓ Taken as an injection every 2 months





Patented until 2031 but <u>licensed to MPP</u> - thanks in large parts to community advocacy efforts!



MPP-ViiV licence for CAB-LA for PrEP

The licence covers the patents on CAB-LA for PrEP, enabling the development and supply of generic versions

The market for CAB-LA for PrEP is uncertain and investments from generic manufacturers might be significant; hence the decision to have **a maximum of three licensees** (this could be revised at a later stage)

Technical and financial support to manufacturers is being explored, such as **technology transfer from ViiV to licensees**

Development and approval of generic versions will likely take 3-4 years from now; meanwhile, the only source of CAB-LA will be the originator product from ViiV Healthcare

- + Quality: WHO PQ or SRA approval required
- + Transparency: the licence is published in full on the MPP website

News & Publications » News & Press Releases » Press Releases

ViiV Healthcare and the Medicines Patent
Pool sign new voluntary licensing agreement
to expand access to innovative long-acting
HIV prevention medicine

28 July 2022

News & Publications » News & Press Releases » Press Releases

Medicines Patent Pool signs sublicences with Aurobindo, Cipla and Viatris to produce generic versions of ViiV Healthcare's innovative long-acting HIV prevention medicine

30 March 2023

- Licences should enable potentially millions of people living in areas most impacted by HIV to access this innovative prevention medicine through lowcost generic manufacturers
- Announcement includes potential for large scale manufacturing on the continent of Africa









Partnering with generics

Sub-licensee selection process

- MPP invites qualified product developers to **apply for licences** to manufacture and sell licensed medicines in LMICs; this is done through an Expression of Interest (EOI) process
- The assessment is rigorous and objective (with **blinded review** for short-listing), and responses are graded using a standardised tool and top-scoring applicants are selected
- The number of licences granted is informed by demand forecasts and other information

By granting multiple sublicences, MPP aims to support adequate supply and competitive pricing

For CAB-LA, there was also an onsite audit of short-listed candidate's manufacturing plant

Sub-licensee selection criteria

- Capacity, capabilities and track-record for manufacturing quality-assured medicines (including experience developing, manufacturing and/or marketing other products)
- **R&D**, financials, and regulatory compliance (including around required quality standards)
- **Specific plans** for the licensed product (regarding development, manufacturing, regulatory plan, distribution, and projected investments – viability of those plans, especially in case of projects requiring specific capital expenditure or investments)
- **Readiness with needed formulation technologies** (e.g., FDCs, paediatrics)
- Past experience and performance as MPP licensee



MPP licence for CAB-LA for PrEP– Understanding the effective territory

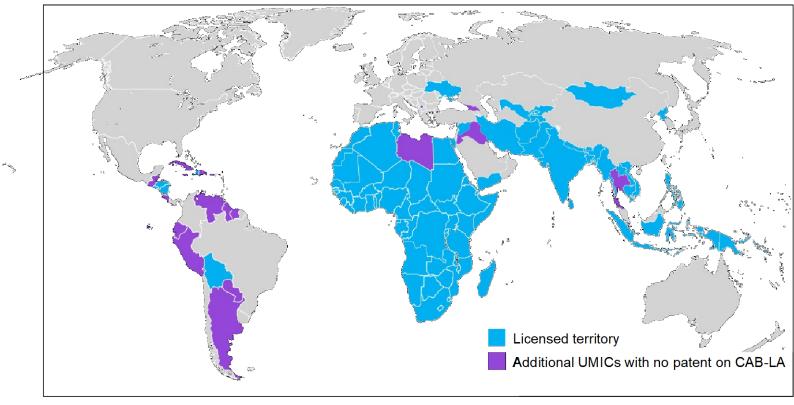
The effective licence territory covers: <u>all countries listed</u> + <u>countries without patents and where supply may be possible</u>

Countries listed (90)

- All low-income countries
- All lower middle-income countries
- All sub-Saharan African countries
- All least-developed countries

Countries without patents and where supply may be possible

 Based on data from MedsPaL (www.medspal.org)



Source: Beatriz Grinsztejn, Long-acting PrEP implementation: Fostering access and equity, AIDS 2022

More at: https://medicinespatentpool.org/progress-achievements/access-to-medicines-tracker



MPP licence for CAB-LA for PrEP Navigating the effective territory

CAB-LA has now been added to our interactive map! **Interactive Map** Save map ▼ 2023 XV HIV XV CAB LAI x V Worldwide XV Clear all filters Product availability through MPP licence Patent and licensing status Included in MPP and other relevant licences Outside of licences but no patents found (MPP) sublicensees may be able to supply) Outside of licences with compound patents Low- & middle-income Other complex patent situations High-income

MPP's interactive map focuses on low- and middle-income countries. Access in high-income countries is generally not included.



Closing message & ideas for future increased collaboration

Now is the time to (1) continue expanding access to oral PrEP and (2) plan for the introduction of CAB-LA for PrEP in anticipation of future availability of generic versions!



We can share information about access to MPP-licensed medicines in your region or country and support you in many ways

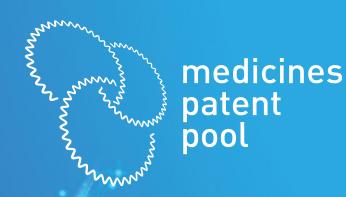
- Please tell us about your needs to increase generic competition for a given product
- Please notify us of stock-outs, quality issues, registration delays, or other challenges regarding MPP-licensed medicines

Do not hesitate to be in touch!

Sébastien Morin, PhD
Policy and Advocacy Manager
Medicines Patent Pool
Geneva, Switzerland



smorin@mppf.ch





Thanks for your attention!

I will be happy to continue the discussion, including over emails (smorin@mppf.ch)





Swiss Agency for Development







MEDICINESPATENTPOOL.ORG



YOUTUBE.COM/USER/MEDICINESPATENTPOOL



in www.linkedin.com/company/medicines-patent-pool/



► YOUTUBE.COM/USER/MEDICINESPATENTPOOL

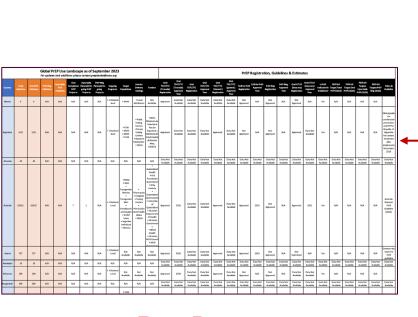




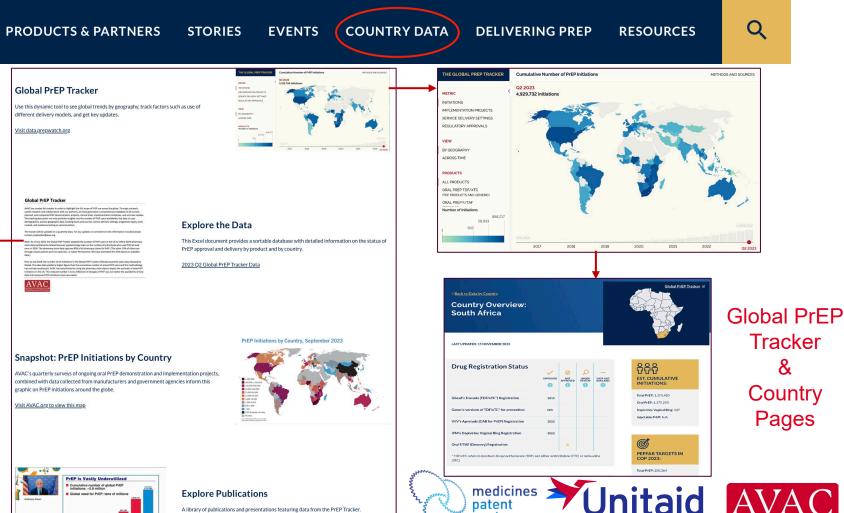


Country Data Resources

PrepWatch An initiative of AVAC.



Raw Data

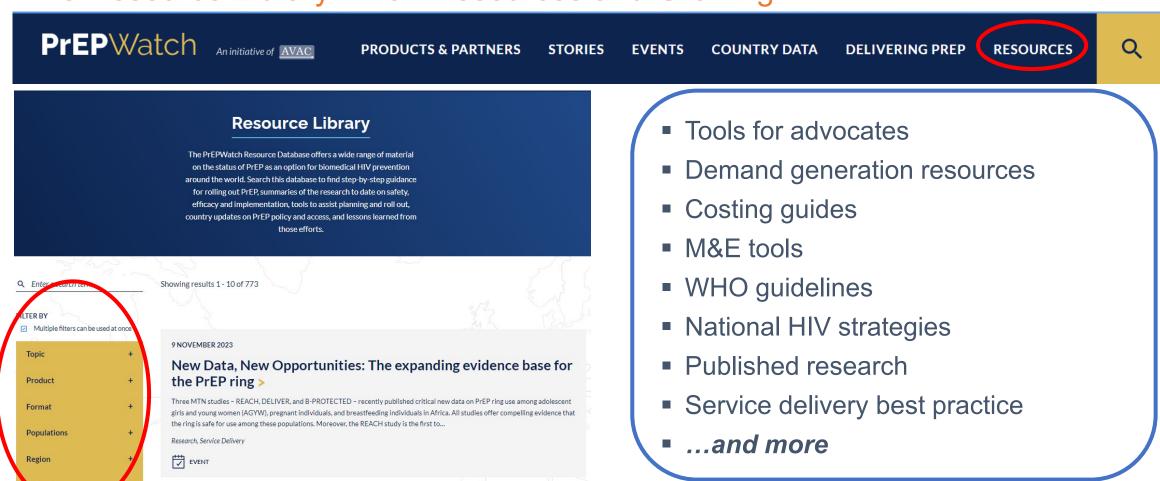


Click here for links to publications that are citing data from the PrEP Tracker.

pool

SAVE LIVES FASTER

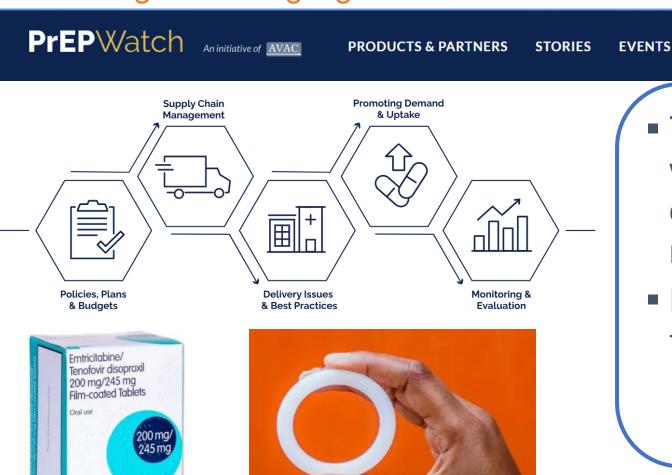
The Resource Library-770+ Resources and Growing







Delivering PrEP- Highlights from the Resource Library



 The most important and widely-used resources, organised by stage of the rollout process

DELIVERING PREP

- Product-specific rollout toolkits:
 - Plan 4 Oral PrEP
 - Plan 4 Ring

COUNTRY DATA





Q

RESOURCES

Dashboards

PrEPWatch

initiative of AVAC

PRODUCTS & PARTNERS

STORIES

EVENTS COUNTRY DATA

DELIVERING PREP



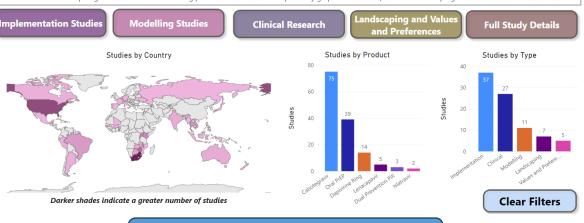


Integrated Study Dashboard for New PrEP Options

This dashboard, produced under the <u>BioPIC</u> project, reflects all currently known activities relating to implementation research, modelling, clinical research, and landscaping for new late stage biomedical HIV PrEP options, including cabotegravir for PrEP and the dapivirine vaginal ring. To update or add any activities missing from the dashboard, or if you have questions on using the dashboard, please contact <u>Catherine Verde Hashim</u>.

Click the buttons below to navigate to the dashboards, organised by study type, or click "Full Study Details" for a table of all studies. You can also navigate through the dashboards using the arrows at the bottom of the screen. The graphs below summarise all studies by location, product, and study type.

All graphs are interactive, and clicking on a graph element will filter all other graphs on the page accordingly. To clear a filter, either click on the graph element again, or click the "Clear Filters" button. For additional details on any of the studies, right click on the relevant graph element and select "Drill through > Full Study Details." For example, right click on South Africa on the map to get details of all studies taking place in South Africa. To expand any graph to full screen, hover over the top right corner and click "Focus mode."



AVAC's **Country Planning for Product Introduction Matrix** tracks the regulatory status of cabotegravir for PrEP and the dapivirine vaginal ring, along with related late stage clinic trials and implementation research, procurement plans, and oral PrEP provision. Use the buttons on the left to filter by region. Right click on any number in the Implementation Studies columns and select "Drill through > Study Details" for full study details. Note that "Nya" indicates year of oral PrEP approval not available.

Click here to visit the PrEPTracker for full PrEP initiation data and click here to visit the study dashboard for full late-stage PrEP research data.

nical	AVAC
	Global Advocacy for HIV Prevention

Country	CAB for PrEP Regulatory Status	DVR Regulatory Status	HPTN 083/084 Studies	Ring/ASPIRE Studies	PURPOSE 1/2 Studies	Year Oral PrEP approved	Oral PrEP Initiations in 2022	PEPFAR CAB Procurement 2023- 2025 (# people)	Gle Pri Fu
Argentina						N/A			П
Australia	Approved					2016	12,196		
Austria	Approved					2016			
Belgium	Approved					2016			
Botswana	Approved	Approved				2016	10,214		
Brazil	Approved					2017	41,589		
Bulgaria	Approved					2016			
Burkina Faso						N/A	391		
Canada	Pending					2016			
China	Pending					2020			
Colombia	Pending					N/A			
Cote D'Ivoire	Pending					N/A	7,080		
Croatia	Approved					2016			
Cyprus	Approved					2016			
Czech Republic	Approved					2016			
Denmark	Approved					2016			
Estonia	Approved					2016			
Eswatini	Via South Africa	Via South Africa				N/A	28,603	Starting 2024/25	
Ethiopia		To file in 2023				N/A	15,141		
Finland	Approved					2016	150		
France	Approved					2016	4,547		
Germany	Approved					2016			

Integrated Study Tracker

Product Introduction Matrix







PRODUCTS & PARTNERS

Also on PrEPWatch...

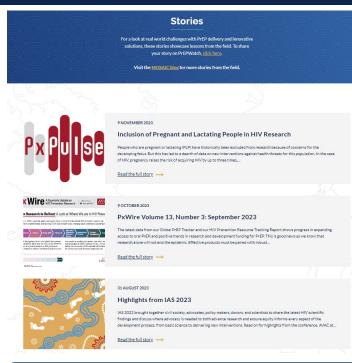
PrEPWatch



LAST UPDATED: 1 NOVEMBER 2023

The first topical HIV preve vaginal ring (DVR) is a silico (IPM). The ring is inserted in the v the course of one month to drug is absorbed elsewhere the bloodstream and other development of HIV resista increased resistance to non Evidence and Research for the ARV drug to which dapiviri Dapivirine Vaginal Ring Click here to access the PrE Learn more -> ring, including basic inform experience. Click here to access the PrE-

research in East and Southern Africa, including trial overviews and key findings.



STORIES

EVENTS COUNTRY DATA

Scroll down for a list of webinars, workshops, conferences, and other convenings where the field is coming together to learn, connect and ensure that PrEP reaches everyone who needs it!

Events

For more events, visit the Global PrEP Learning Network and The Choice Agenda.

Tuesday
28
November
© 13:30 - 15:00 GMT

VIRTUAL

DELIVERING PREP

Spotlight on New Tools and Data: From R&D to Access

Between the recent accelerated growth in global PFEP initiations, and the introduction of new PFEP products like cabotegravir and the daplivirine vaginal ring, the field of PFEP data has never been more exciting or more complex. Staying on top of the latest advances is key for advocate, researchers, funders, and others working in HIV prevention.

RESOURCES

Wednesday
29
November
© 13:00 - 14:00 GMT

VIRTUAL

New Data, New Opportunities: The expanding evidence base for the PrEP ring >

Three MTN studies – REACH, DELIVER, and B-PROTECTED – recently published critical new data on PrEP ring use among adolescent girls and young women (AGTW), pregnant individuals, and breastfeeding individuals in Africa, All studies offer compelling evidence that the ring is as fer or use among these populations. Moreover, the REACH study is the first to...

Lessons from the Field

Webinars and Other Events







Detailed Product Overviews



The PrEPWatch Team

Please do be in touch with any questions at prepwatch@avac.org or

Catherine Verde Hashim

catherine@avac.org



Janki Tailor janki@avac.org





THE LONG-ACTING THERAPEUTICS PATENTS AND LICENCES DATABASE

A collaborative tool to support access to innovative long-acting therapeutics

LAPaL is a free online resource that provides innovation and access support on technical features, development status and intellectual property status of selected long-acting therapeutics that could have impact in low- and middle-income countries.

www.lapal.ch





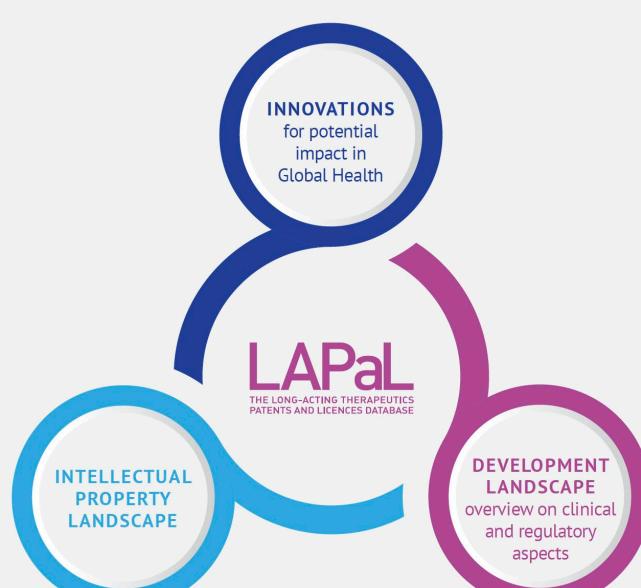






WHAT ARE LAPaL'S MAIN COMPONENTS?

Long-acting technologies and compounds



THE LONG-ACTING THERAPEUTICS ONE-STOP-SHOP



Long-acting therapeutics development landscape with technologies, compounds and their co-formulations



Visualisation dashboard:

- Insights on clinical trials
- Overview of regulatory status

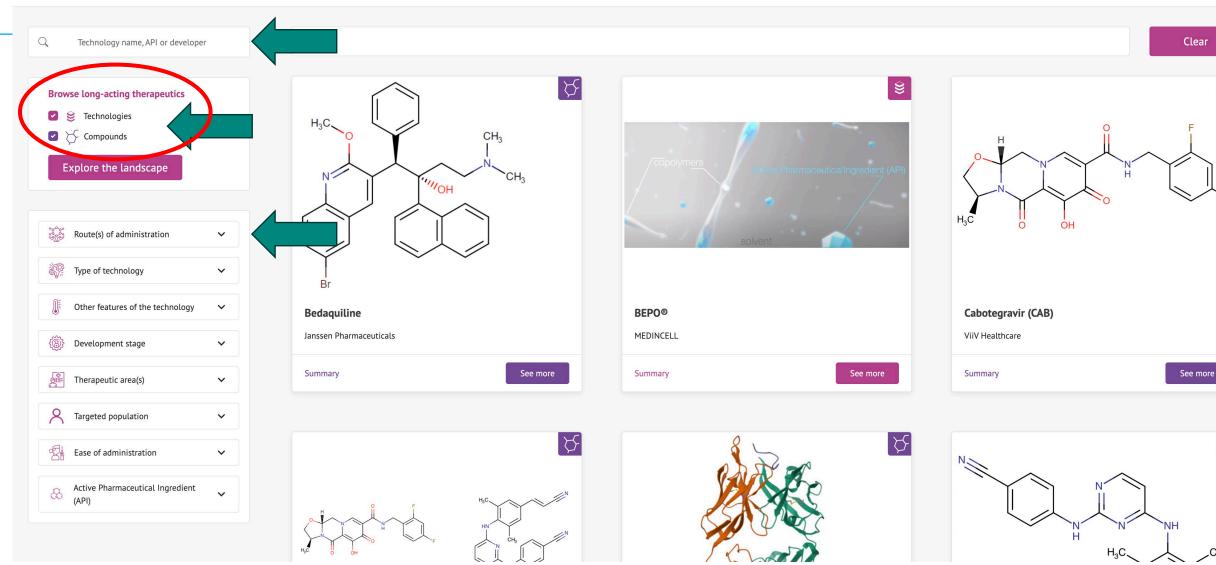


Intellectual property landscape



Access commitments for pipeline acceleration















Drug Combination Nanoparticles (DcNP)

Verified by the innovator, on Jul 2021



Active Pharmaceutical Ingredient
Tenofovir disproxil fumarate (TDF),
Lamivudine (3TC), Dolutegravir (DTG),
Lopinavir and ritonavir (LPV/r),

Tenofovir (TFV)

Route(s) of administration Subcutaneous

Type of technology

Based on other organic particles, Aqueous drug particle suspension

Other features of the technology
Biodegradable, Non-removable, Room
temperature storage, At least 1 year
shelf life

Development stage
Pre-clinical

Therapeutic area(s)

Ease of administration

Administered by a community health worker, Administered by a nurse,

Administered by a specialty health worker, Self-administered

Access principles





Targeted & Long-acting drug Combination (TLC) Program, University of Washington

Summary

See more











Drug Combination Nanoparticles (DcNP)

×

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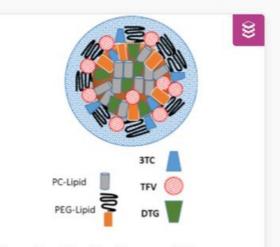
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Access principles



Drug Combination Nanoparticles (DcNP)

Targeted & Long-acting drug Combination (TLC) Program, University of Washington

Summary

See more













< Back Verified by the innovator, on Jul 2021



Technology name

Main developer(s)

Sponsor(s)









Unitaid

Technology information

Potential application(s)

Clinical trials

Patent info

Supporting material

Access principles

Partnerships & sponsors

Therapeutic area(s)

HIV

Use case(s)

Treatment

Potential associated API(s)

- Tenofovir disproxil fumarate (TDF), Lamivudine (3TC), Dolutegravir (DTG)
- Lopinavir and ritonavir (LPV/r), Tenofovir (TFV)

Use of technology

Ease of administration

- · Administered by a community health worker
- Administered by a nurse
- · Administered by a specialty health worker
- Self-administered

User acceptance Potentially good Frequency of administration

Monthly, Once every 8 weeks

Targeted user groups

Age Cohort Adults

Genders Male

- Female
- Cisgender female
- Cisgender male
- Transgender female
- Transgender male
- Intersex
- · Gender non-binary
- All

Yes

Pregnant individuals

Yes

Lactating individuals

Healthy individuals

Not provided

Comment

Tenofovir disproxil fumarate (TDF), Lamivudine (3TC), Dolutegravir (DTG)

Class(es)

Antiviral

Foreseen user group

People living with HIV (Chronic HIV positive patients)

Lopinavir and ritonavir (LPV/r), Tenofovir (TFV)

Class(es) antiviral Foreseen user group

People living with HIV















Technology name

Main developer(s)

Sponsor(s)



<u>_</u>

Drug Combination Nanoparticles (DcNP)



Unitaid
Innovation in Global Health

Technology information

Potential application(s)

Clinical trials

Patent info

Supporting material

Access principles

Partnerships & sponsors

Use case **♦**

Use case

Clinical trials (1 result)

Name

First in Human Clinical Trial of a Next Generation, Long-acting Injectable, Combination Antiretroviral
Therapy Platform TLC-ART 101 (ACTU 2001)

Identifier

NCT05850728

Phase Status

Purpose

Recruiting Safety, tolerability and PK of single subcutaneous injection of TLC-ART 101 in healthy adults

Phase Filt

Filter by Phase \$

Filter by Status

Status

Treatment

Use case





Clinical tria





Phase











Global information

Trials dates

Studied population

Study design

Use of drug candidate

Key results

Clinical trial general information



Name

First in Human Clinical Trial of a Next Generation, Long-acting Injectable, Combination Antiretroviral Therapy Platform TLC-ART 101 (ACTU 2001)



Link

https://clinicaltrials.gov/study/NCT05850728?intr=TLC-ART&rank=1

Sponsor

Technology info

NIH

Purpose

Clinical trials (

Safety, tolerability and PK of single subcutaneous injection of TLC-ART 101 in healthy adults

. .

Countries

United States

Name

Sites / Institutions

First in Human Cl Therapy Platform Seattle, Washington

Identifier

NCT05850728

Phase Status

Phase I Recruiting

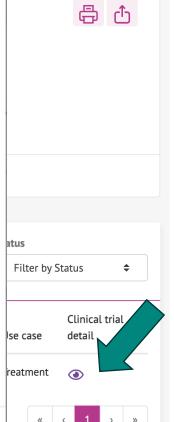
More details

This study is a prospective, open-label, single-site, first-in-human study of a long-acting, injectable combination antiretroviral therapy platform, with a pharmacologically-guided adaptive design for dose escalation, de-escalation, and study duration. The study is designed to learn whether the formulation can be used as a platform for other drugs for treatment of HIV. The formulation is a drug combination nanoparticle (DCNP). The study will be conducted by UW Positive Research. The sample size for this study is 12-16. The study population consists of healthy adults without HIV. The study duration is 57 days per participant at the start of the study.

Interventions

• TLC-ART 101 containing lopinavir 15.6mg, ritonavir 4.2 mg, and tenofovir 9.15 mg in a combination nanoparticle suspension of 1.5 mL

















Expiry date: 7 Jan 2041

Technology information	Potential app	olication(s) CI	inical trials Par	ent info	Supporting materi	ial A	ccess principles	Partnerships & sponsors			
Technology patent families											
Patent description		Representative paten	t Categories	Patent ho	older	Abstract	Patent status	Licence with N	1PP	Licence details	Patent source
Drug Combination Nanoparticles for drugs extended release	multiple	WO2016205384	Long-acting platfor	m The Univ Washing	-	Ш	(~		(Company
Expiry date: 15 Jun 2036											
Combination pharmaceutical composincluding a combination of hydrophi hydrophobic therapeutic agents		WO2020146788	Long-acting platfor	m Universit Washingt	•	Ш		~			Company
Expiry date: 10 Jan 2040											
Antiviral compositions that transform acting therapeutic agents into long-injectable forms that lasts for many administration	acting	WO2021142150	Long-acting platfor	m Universit Washingt	-	Ш	(~		(Company













Expiry date: 7 Jan 2041

Technology information Potential a	pplication(s) Clin	ical trials Patent	info Supporting materi	al Access p	rinciples Partn	erships & sponsors		
Technology patent families								
Patent description	Representative patent	Categories	Patent holder	Abstract	Patent status	Licence with MPP	Licence details	Patent source
Drug Combination Nanoparticles for multiple drugs extended release	WO2016205384	Long-acting platform	The University Of Washington		(~	(Company
Expiry date: 15 Jun 2036								
Combination pharmaceutical compositions including a combination of hydrophilic and hydrophobic therapeutic agents	WO2020146788	Long-acting platform	University of Washington	Ш		Ý		Company
Expiry date: 10 Jan 2040								
Antiviral compositions that transform short- acting therapeutic agents into long-acting injectable forms that lasts for many weeks per administration	WO2021142150	Long-acting platform	University of Washington	Ш	(~	(Company











Combination pharmaceutical co University of Washington, (10 Ja Source: Company	ompositions including a combination of hydrophilic and hydrophob an 2040)	ic therapeutic agents, WO2020146788, Long-acting platform, ×
Patent status	Low, low-middle and upper-middle countries	High income countries
Granted	South Africa	
Filed	China, Brazil, India, Nigeria	United States of America
Not in force	World Intellectual Property Organization (WIPO)	World Intellectual Property Organization (WIPO)
		Close

s for













Expiry date: 7 Jan 2041

Technology information Potential app	plication(s) Clini	cal trials Patent	info Supporting materia	l Access pri	nciples Partne	rships & sponsors		
Technology patent families								
Patent description	Representative patent	Categories	Patent holder	Abstract	Patent status	Licence with MPP	Licence details	Patent source
Drug Combination Nanoparticles for multiple drugs extended release	WO2016205384	Long-acting platform	The University Of Washington	Ш		~	(Company
Expiry date: 15 Jun 2036								
Combination pharmaceutical compositions including a combination of hydrophilic and hydrophobic therapeutic agents	WO2020146788	Long-acting platform	University of Washington	Ш	(~	0	Company
Expiry date: 10 Jan 2040								
Antiviral compositions that transform short- acting therapeutic agents into long-acting injectable forms that lasts for many weeks per administration	WO2021142150	Long-acting platform	University of Washington	Ш	(~	(Company











MPP/Uni Washington licence on Drug Combination Nanoparticles (DcNP) - HIV prevention/treatment

Type: MPP Licence

Licensor: University of Washington

Jurisdictions: Afghanistan; Algeria; American Samoa; Angola; Argentina; Armenia; Azerbaijan; Bangladesh; Belarus; Belize; Benin; Bhutan; Bolivia (Plurinational State

of); Bosnia and Herzegovina; Botswana; Brazil; Bulgaria; Burkina Faso; Burundi; Cabo Verde; Cambodia; Cameroon; Central African Republic; Chad; China;

Colombia; Comoros; Congo; Congo, democratic Republic of the; Costa Rica; Côte d'Ivoire; Cuba; Djibouti; Dominica; Dominican Republic; Ecuador; Egypt;

El Salvador; Equatorial Guinea; Eritrea; Eswatini; Ethiopia; Fiji; Gabon; Gambia (the); Georgia; Ghana; Grenada; Guatemala; Guinea; Guinea-Bissau;

Guyana; Haiti; Honduras; India; Indonesia; Iran (Islamic Republic of); Iraq; Jamaica; Jordan; Kazakhstan; Kenya; Kiribati; Korea (Democratic People's

Republic of); Kosovo; Kyrgyzstan; Lao People's Democratic Republic (the); Lebanon; Lesotho; Liberia; Libya; Madagascar; Malawi; Malaysia; Maldives; Mali; Marshall Islands; Mauritania; Mexico; Micronesia (Federated States of); Moldova, Republic of; Mongolia; Montenegro; Morocco; Mozambique;

Myanmar; Namibia; Nepal; Nicaragua; Niger; Nigeria; North Macedonia; Pakistan; Papua New Guinea; Paraguay; Peru; Philippines; Russian Federation;

Rwanda; Saint Lucia; Saint Vincent and the Grenadines; Samoa; Sao Tome and Principe; Senegal; Serbia; Sierra Leone; Solomon Islands; Somalia; South

Africa; South Sudan; Sri Lanka; State of Palestine; Sudan; Suriname; Syrian Arab Republic; Tajikistan; Tanzania, United Republic of; Thailand; Timor-Leste;

Togo; Tonga; Tunisia; Türkiye; Turkmenistan; Tuvalu; Uganda; Ukraine; Uzbekistan; Vanuatu; Venezuela (Bolivarian Republic of); Viet Nam; Yemen; Zambia;

Zimbabwe

Licensee(s):

MPP licence: https://medicinespatentpool.org/licence-post/long-acting-injectable-drug-combination-for-hiv-treatment-prevention

Close











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LAPaL Landscape Clinical Trials Regulatory Status



rug Name	Trial Name	Trial ID	Trial Phase	Enrollment	Age Eligibility	Cohort	Current Status	Use Case
Cabotegravir	ECLAIR	NCT02076178	Phase II	127	18-65	Adults, Older Adults	Completed	PrEP
Cabotegravir	HPTN 077	NCT02178800	Phase II	199	18-65	Adults, Older Adults	Completed	PrEP
Cabotegravir	HPTN 083	NCT02720094	Phase II/III	4570	18+	Adults, Older Adults	Active Not Recruiting	PrEP
Cabotegravir	HPTN 084	NCT03164564	Phase III	3224	18-45	Adults	Active Not Recruiting	PrEP
Cabotegravir	EBONI	NCT05514509	Marketed	250	18+	Adults, Older Adults	Recruiting	PrEP







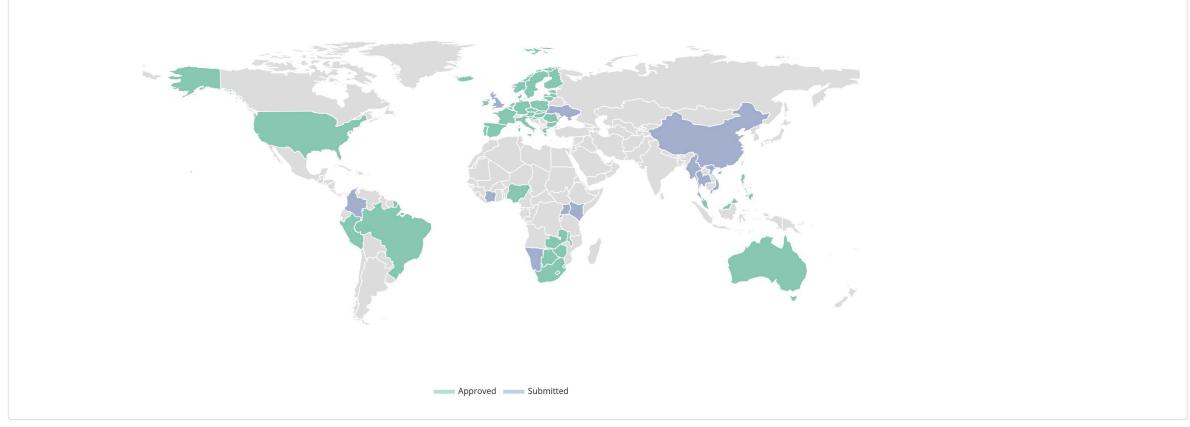




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Regulatory Stat	istics
Approved	44
Submitted	12

Date of Last Update:		
1st Nov 2023		







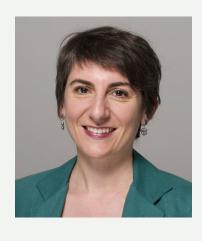








THE LONG-ACTING THERAPEUTICS PATENTS AND LICENCES DATABASE



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Thanks for your attention!

Please reach out to us with questions, proposals for improvements or any comments

www.lapal.ch

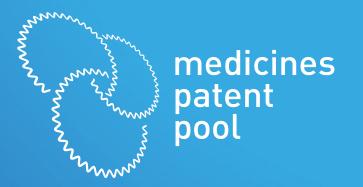












Access to Medicines Tracker: Interactive Map (and Table)

Webinar - Spotlight on new PrEP tools and data: From R&D to access

Convened by AVAC, MPP, and Unitaid on 28 November 2023

Presented by Sébastien Morin (Medicines Patent Pool)

Spotlight on New PrEP Tools and Data: FROM R&D TO ACCESS



TUESDAY, NOVEMBER 28
8:30 TO 10:00 AM ET / 14:30 TO 16:00 PM CET

Join us to learn about three important online PrEP resources and to understand how they can support and enhance your work.

Register at tinyurl.com/PrEPTools







MEDICINESPATENTPOOL.ORG



Access to Medicines Tracker: Interactive Map (and Table)

Step 1.

Navigate to the MPP website and select "Progress"/"Access to Medicines Tracker"





Step 2.

On the "Access to Medicines Tracker" webpage, scroll down to the "Interactive Map"



Step 3.

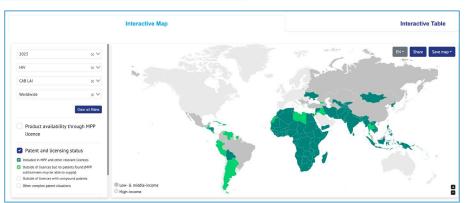
Explore country-level quarter-byquarter data on access, filing, approval, and supply of MPP-licensed medicines

More at: https://medicinespatentpool.org

https://medicinespatentpool.org/progress-achievements/access-to-medicines-tracker

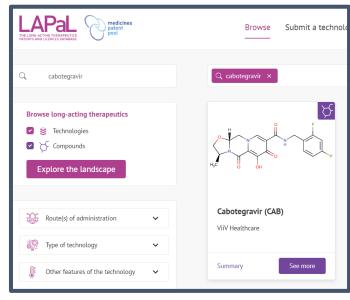
https://map.medicinespatentpool.org -> This is the link for direct access to the map in full window mode.

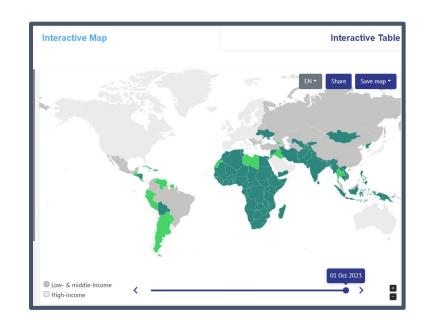
https://map.medicinespatentpool.org/table -> This is the link for direct access to the table in full window mode.



When do I use each tool?











ACCESS TO MEDICINES TRACKER







Cheat Sheet

Where do I go to find information on...?





Technical info on PrEP- applicable LA platforms & compounds (incl. publications & documentation)

Scaled-up manufacturing information

Intellectual property information (patents and licences)

PrEP research studies (incl. implementation studies, modelling studies)

Information on PrEP product approvals

Country MoH & WHO PrEP guidelines

Innovators access commitments

Resources to support PrEP rollout process

PrEP initiations & PEPFAR targets

ACCESS TO MEDICINES TRACKER

Intellectual property access information (patents & licences) for MPP-licensed products (such as CAB-LA)

Information on filings, approvals, supplies, & suppliers of MPP-licensed products (such as CAB-LA)

Quarter-by-quarter country-bycountry data (with possibility to explore historical data points starting from 2019)

Finding and Using Data

Mentimeter

Click here to launch Mentimeter interactive quiz

Find the answers here:

PrEPWatch

The Long-Acting Therapeutics Patents and Licenses Database

Access to Medicines Tracker







Q&A







Thank you!

Further Resources

- PrEPWatch
- PrEPTracker
- Integrated Study Dashboard
- Country Planning for Product Introduction Matrix
- The Long-Acting Therapeutics Patents and Licenses Database
- Access to Medicines Tracker
- MedsPaL







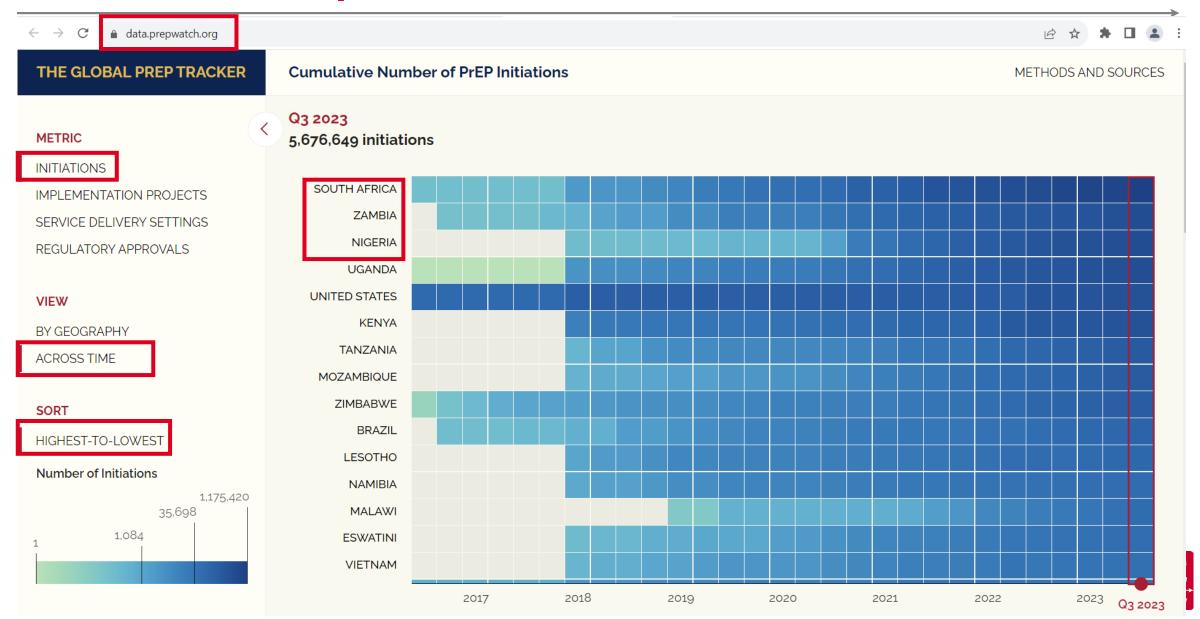
Quiz Questions and Answers



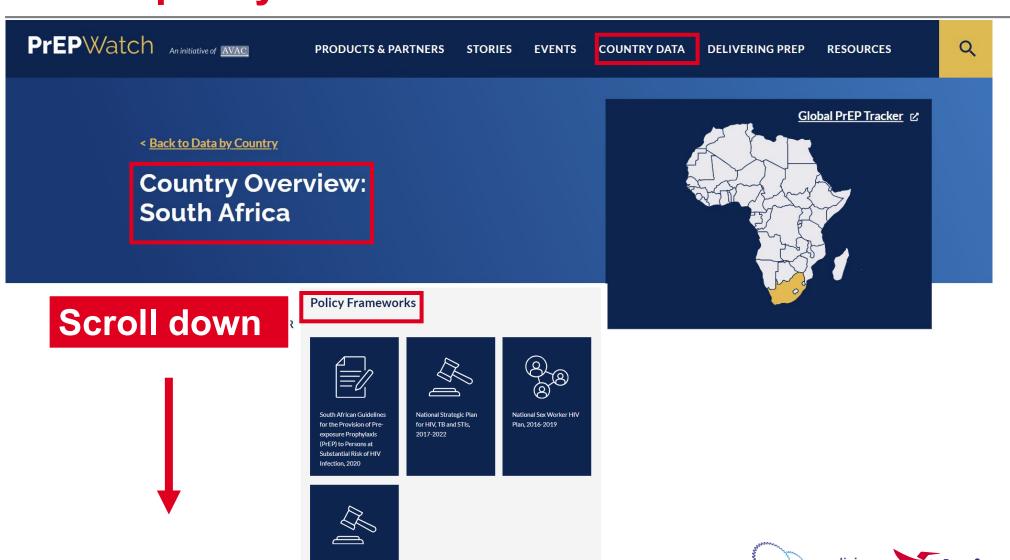




What are the top three countries for PrEP Initiations?



Which policy frameworks are available for South Africa?

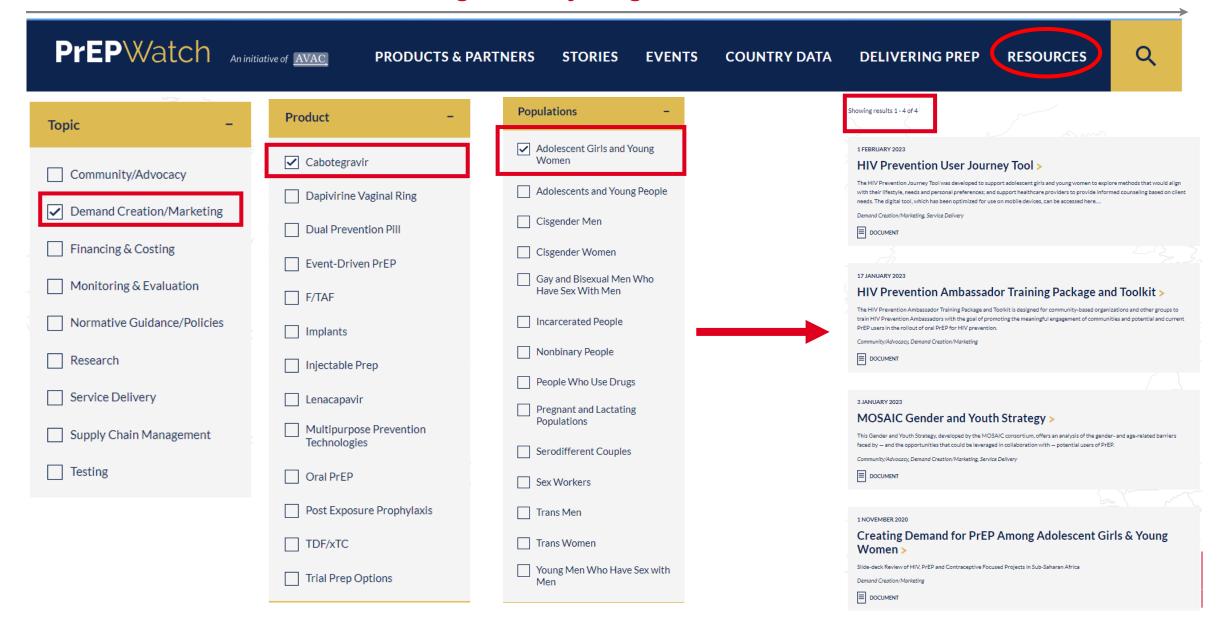








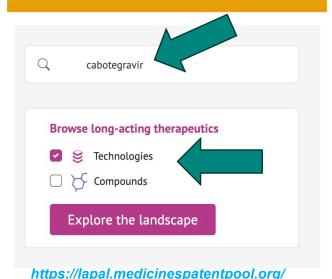
How many resources does the PrEPWatch library have on demand generation materials for CAB for PrEP aimed at adolescent girls and young women?

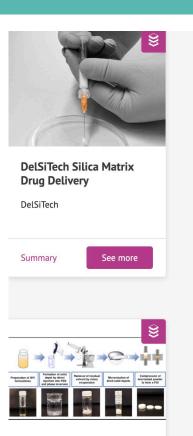




In LAPaL, how many long-acting technology platforms could potentially be associated with cabotegravir?

Five platform technologies listed on LAPaL could potentially be coupled with cabotegravir to extend its release



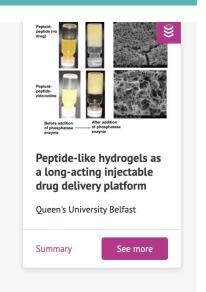


Tunable Biodegradable

Polymeric Solid Implant

Ultra-Long-Acting

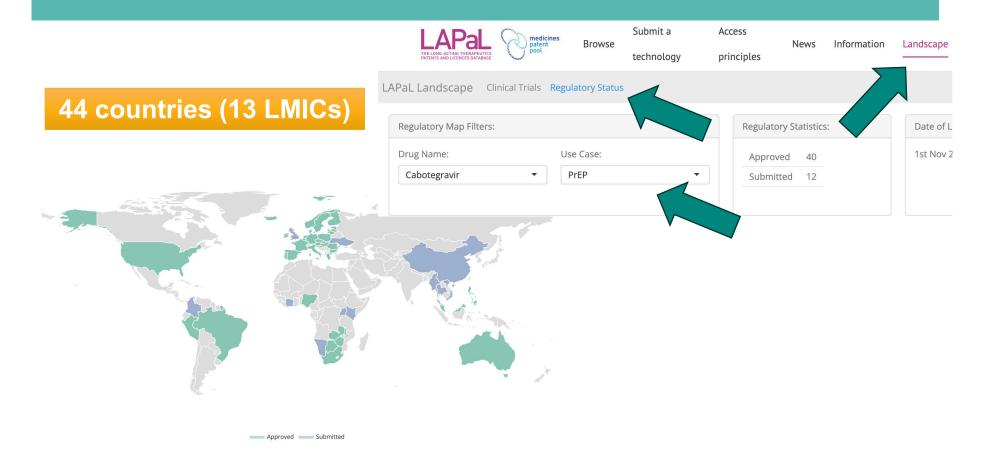








In how many countries has CAB-LA for PrEP received approval?





In the absence of a licence, when could generic quality assured versions of lenacapavir enter the market?

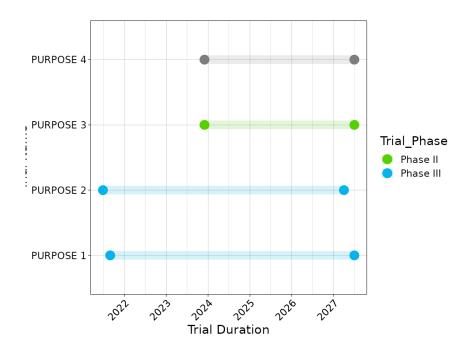
Not before 2034 or even 2037 in many LMICs

Patent description	Representative patent	Categories	Patent holder	Abstract	Patent status	Licence with MPP	Licence details	Patent source
Lenacapavir and analogues (Markush formula) and their use in HIV	WO2014134566	Compound	Gilead Sciences, Inc	Ш	(MPP Search, US FDA
Expiry date: 28 Feb 2034								
Lenacapavir compound and its use in HIV (oral and parenteral)	WO2018035359	Compound	Gilead Sciences, Inc	Ш	(MPP Search, US FDA
Expiry date: 17 Aug 2037								
Crystalline forms of Lenacapavir sodium salt	WO2019035904	Polymorphs	Gilead Sciences, Inc	Ш	(MPP Search, US FDA
Expiry date: 16 Aug 2038								



How many clinical trials are ongoing or planned for lenacapavir in the context of PrEP?

4 trials: PURPOSE 1 to 4



https://lapal.medicinespatentpool.org/landscape

About PURPOSE 1

- Phase 3 study of an investigational drug, lenacapavir, for PrEP and emtricitabine/tenofovir alafenamide (F/TAF) for PrEP in Adolescent Girls and Young Women
- This study will be conducted in South Africa and Uganda

PURPOSE 3

PK, Safety, and Acceptability of LEN for PrEP in Cisgender Women in the United States.

About PURPOSE 2

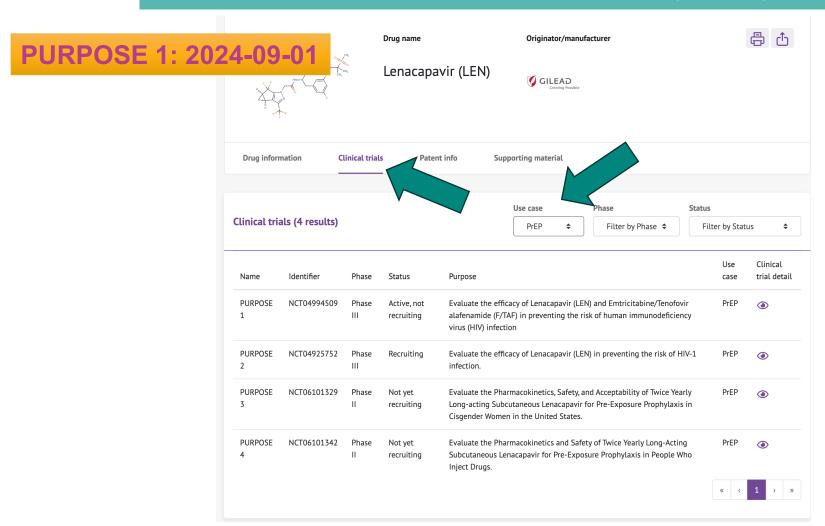
- Phase 3 study of an investigational drug, lenacapavir, for PrEP for Cisgender Men, Transgender Women, Transgender Men, and Gender Non-Binary individuals Who Have Sex With Partners Assigned Male at Birth
- This study will be conducted in the United States, South Africa, Peru, Brazil, Argentina, Mexico, and Thailand

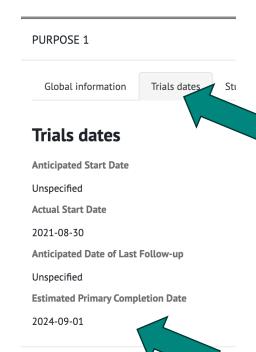
PURPOSE 4

PK, Safety, and Acceptability of LEN for PrEP in People Who Inject Drugs.



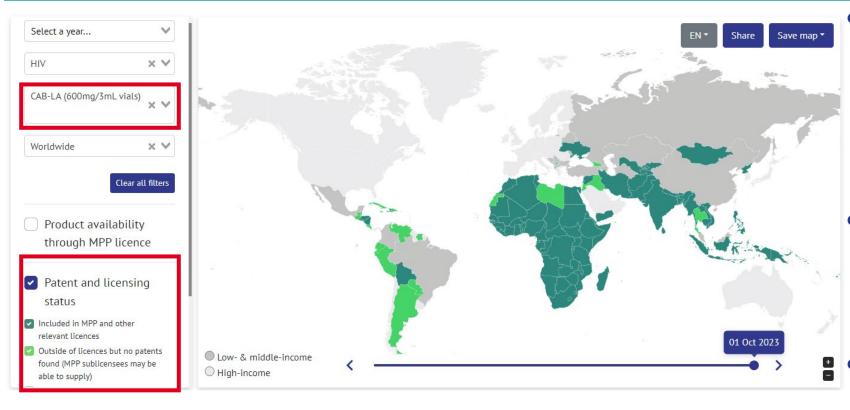
Which of these trials have the earliest primary completion date?







List three countries included in the MPP-licensed territory for generic cabotegravir long-acting injectable, and three countries not included in the licensed territory, but where access may be possible as there are no patents



- territory for generic cabotegravir long-acting injectable can be visualized on this interactive map and are also available from MedsPaL or on the CAB-LA licence page
- Countries not included in the licensed territory, but where access may be possible as there are no patents can be visualized on this interactive map
 - The full, effective overall territory can be visualized on this interactive map, it is also discussed in an online post by Brook Baker