



# Polio eradication: How technologies are helping us travel the last mile

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# What is Polio?

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Only one serotype of Wild Poliovirus (WPV1) is still in circulation globally



Polio is a crippling and potentially deadly infectious disease caused by the poliovirus. Spreads person-to-person and causes paralysis in 1 out of 200 of infections

Two vaccines provide protection against the virus



Improved sanitation and polio vaccination have resulted in eradication of Type 2 (September 2015) and Type 3 (October 2019)

**6** CORE PARTNERS

**200** COUNTRIES INVOLVED

**20** MILLION VOLUNTEERS

over **2.5** BILLION CHILDREN  
VACCINATED

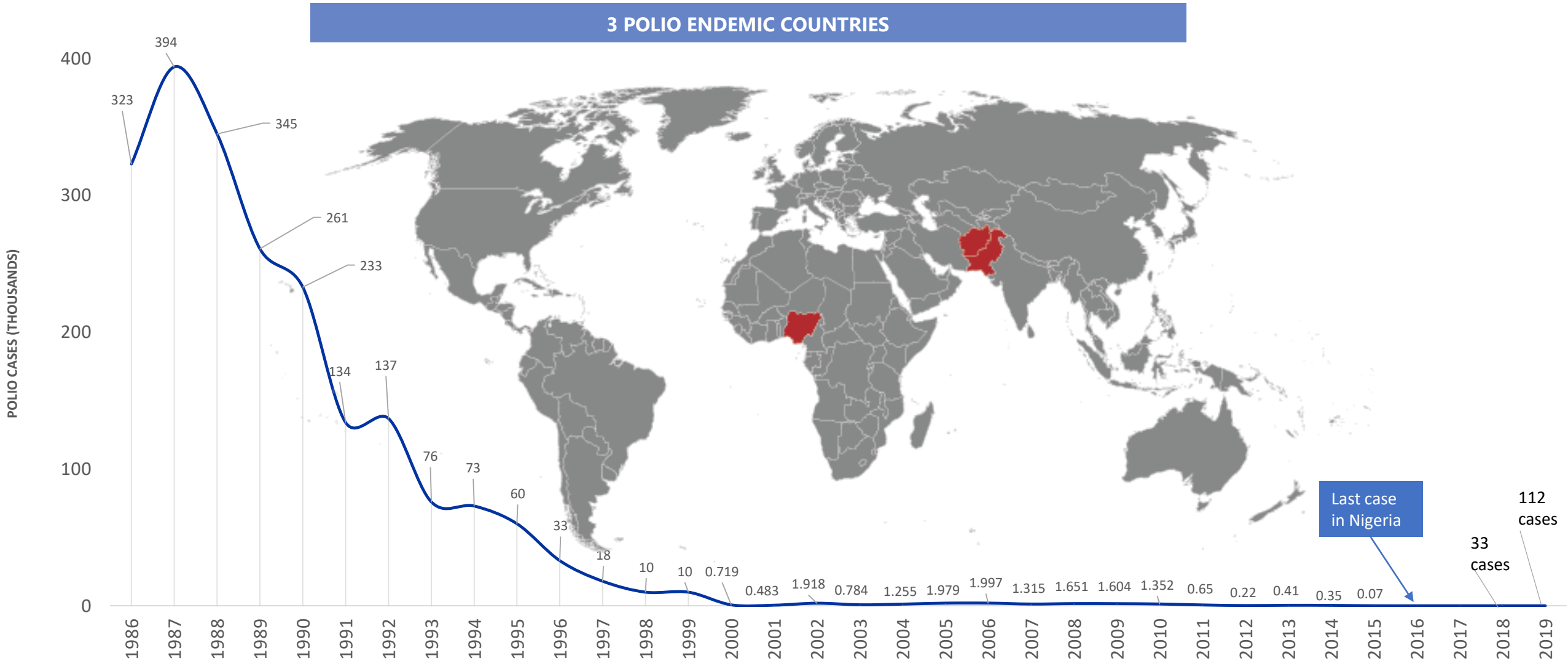
US\$ **17** BILLION  
INTERNATIONAL  
INVESTMENT

**GOAL** A POLIO – FREE  
WORLD

In 1988, GPEI started upon the bold mission to eradicate polio |



# Wild Poliovirus is currently circulating in two countries: Pakistan & Afghanistan



Reaching the last child and traveling the last mile requires new innovations

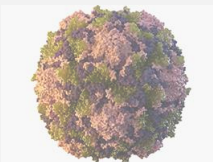
## The Eradication Program Faces Challenges



**Insecurity:** limits the reach of vaccination campaigns and threatens the safety of polio volunteers



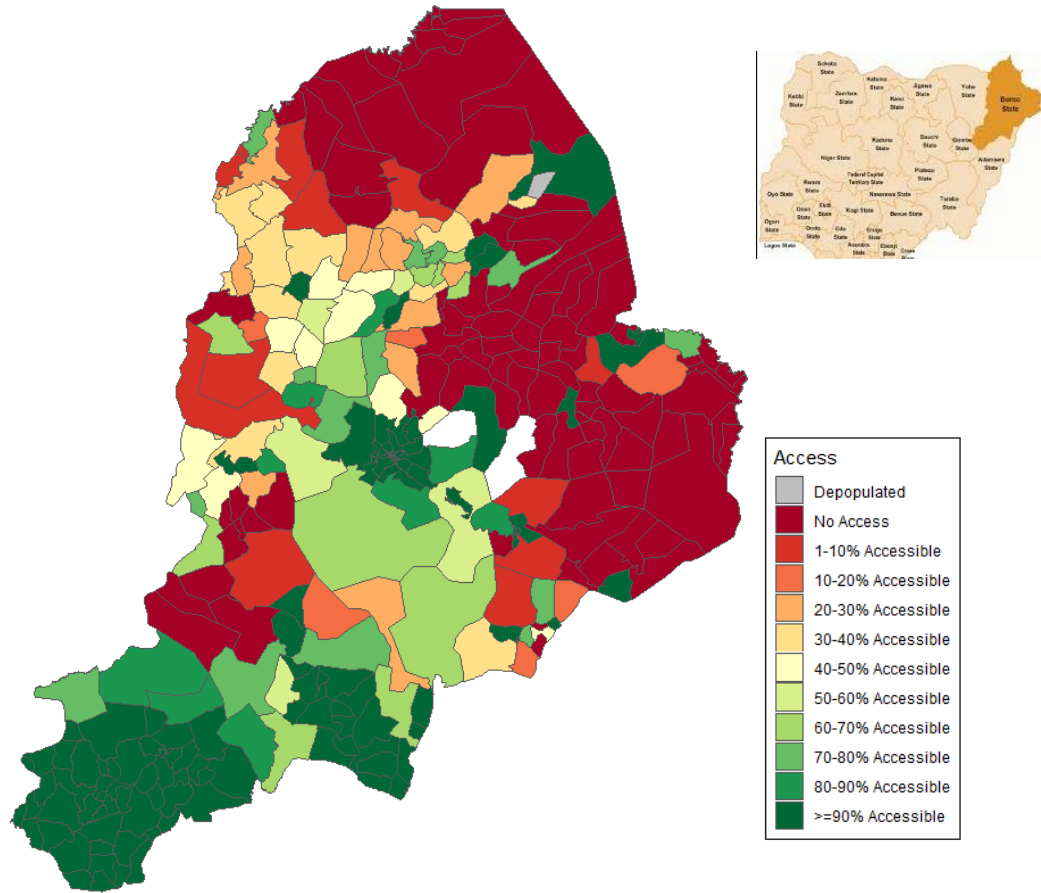
**Poor health infrastructure:** Limits vaccine coverage through routine immunization allowing continued transmission



**Changing epidemiology:** new outbreaks of cVDPV2 stretch the resources of the program



## Ward Level Access in Borno, 2016



**Challenge:** Insecurity in North East Nigeria allowed transmission of Polio to continue undetected for years

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Inhabited through 2016 rainy season

Abandoned between 2016 and 2017 rainy seasons

	2016									2017														
Month	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep	Oct	Nov	Dec	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep	Oct	Nov	Dec
Avg Precip (mm)	0	0	0.3	13	30.5	73.8	147.1	193.2	83	11.1	0	0.1	0	0	0.3	13	30.5	73.8	147.1	193.2	83	11.1	0	0.1

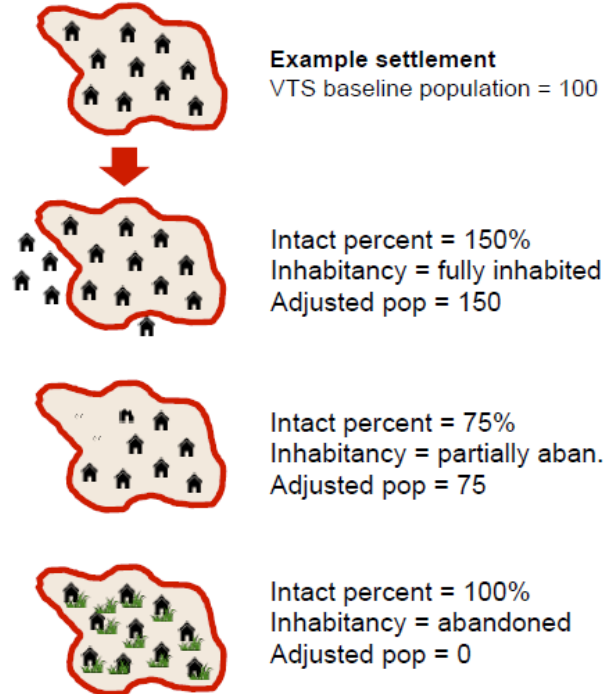


**Solution:** satellite imagery with machine learning reviews thousands of settlements annually

Program now using machine learning to estimate percentage of structures intact



CDC data integrated into VTS  
Assessments modify VTS settlement level  
population estimates:



Satellite images used to adjust the size of the population



**Borno State:  
Inhabited Settlements\* and Vaccination Contacts\*\*  
August 2016 - March 2018**

\*Settlements with signs of potential habitation visible in satellite imagery  
\*\*Areas with GIS evidence of visits by vaccination teams in separate rounds

**Estimated Population (VTS +satellite)**

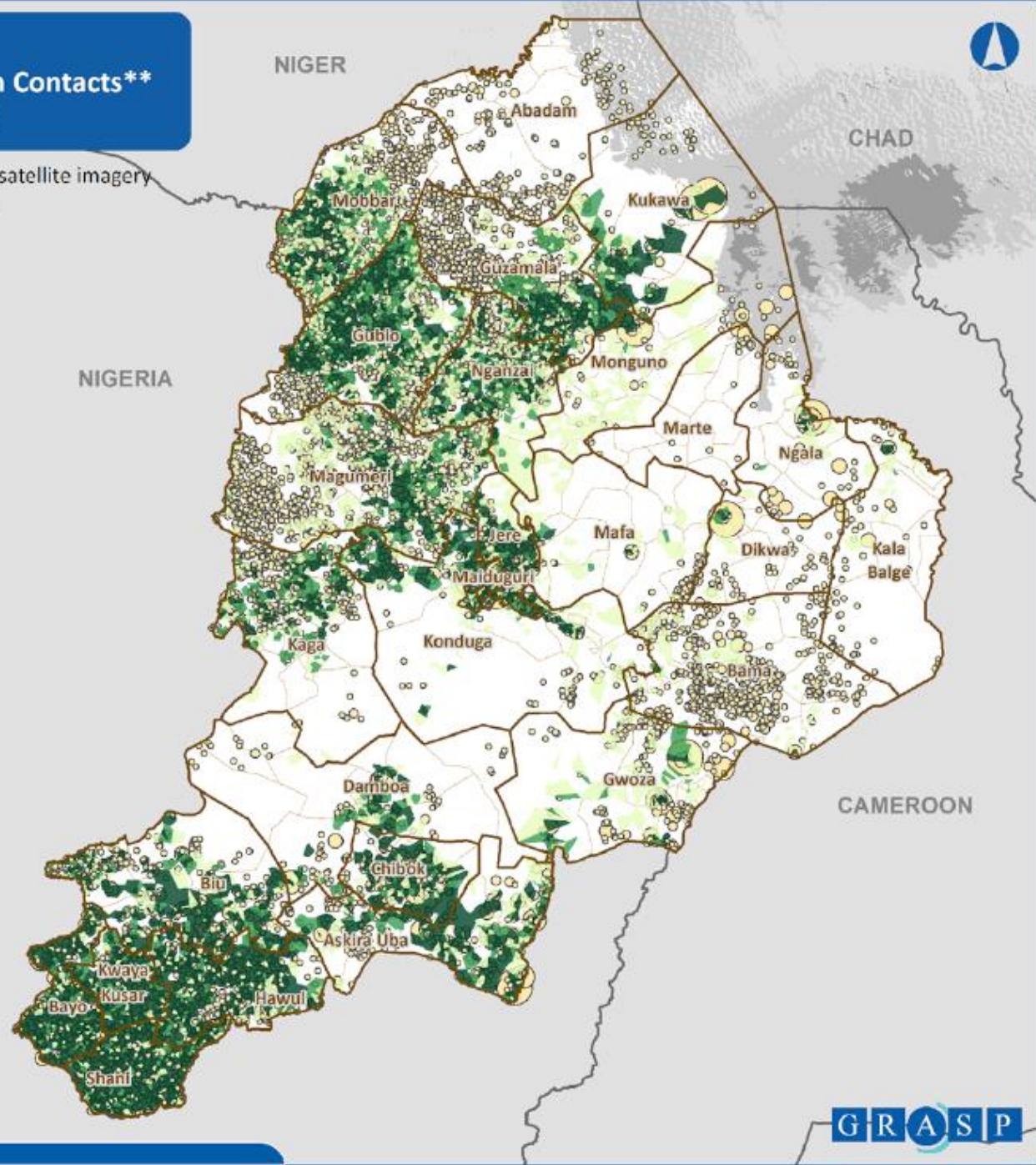
- 1 - 500
- 501 - 2000
- 2001 - 10000
- 10001 - 30000
- 30001 - 200000

**Vaccination Contacts**

- 0
- 1 - 2
- 3 - 5
- 6+

**Estimated u5 children by # of contacts**

0 contacts:	132,546
1-2 contacts:	44,888
3-5 contacts:	42,849
6+contacts:	652,224



Using GPS, we now track which settlements are visited by vaccination campaigns and estimate the size of the population missed





Challenge: Poor infrastructure creates barriers for even basic tasks



## Vaccines



## Surveillance specimens



Solution: Drones can help overcome challenges with landscape and maintaining cold chain to help deliver vaccines and return surveillance specimens



# Currently the technology is being piloted in Papua New Guinea and Cameroon



## **Phase I (PNG)**

Building foundations for learning, capacity building

Introduce stakeholders to technology

Test how quickly drone team could be deployed and flight permission secured

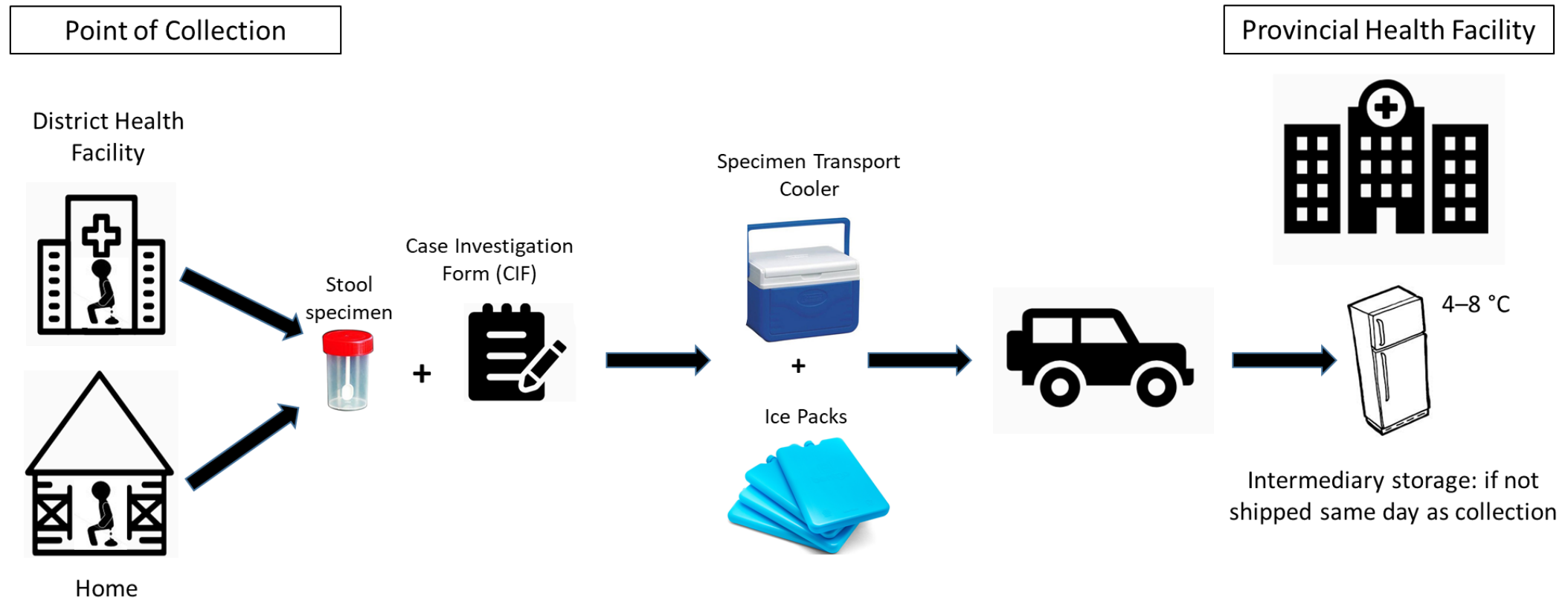
## **Phase II (Cameroon)**

Implement flight routes

Collect a lot of data to evaluate value and inform future project planning

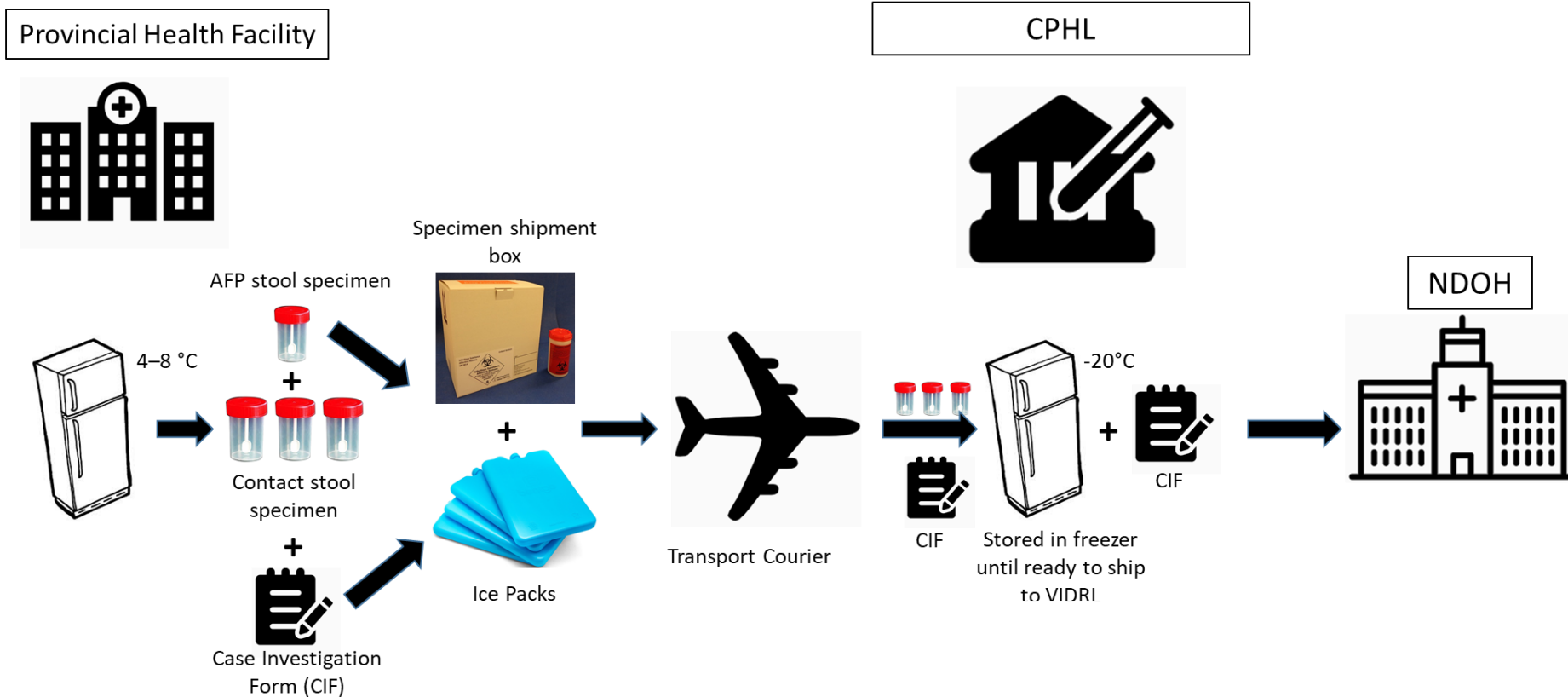
Plan for sustainability and scale up

# Challenge: Poor understanding of weak links in the complex cold chain for specimen shipment and vaccines



From point of collection to province-level health facility

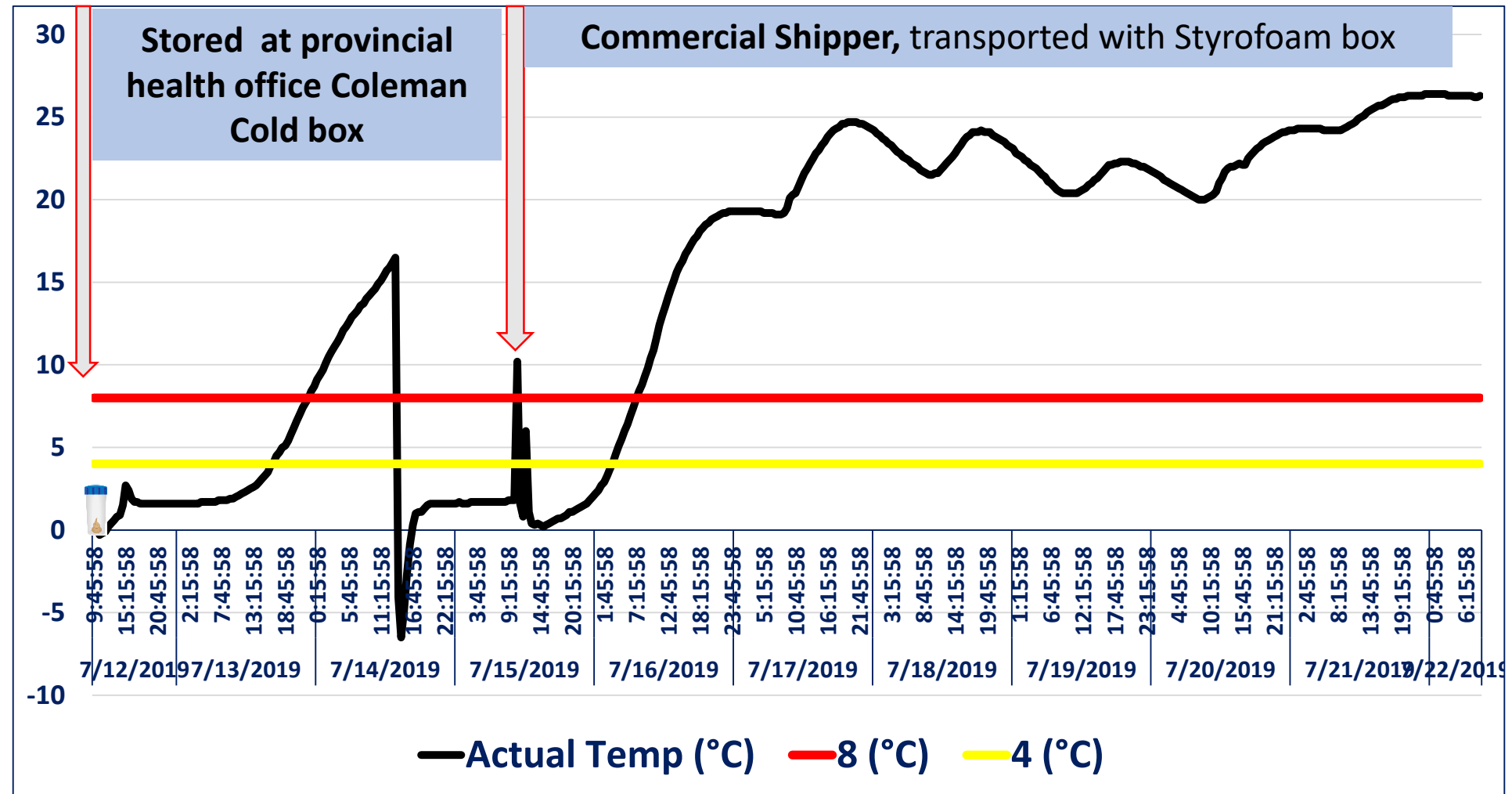
# Challenge: Poor understanding of weak links in the complex cold chain for specimen shipment and vaccines



From provincial facilities to the National reference laboratory (CPHL)



# Solution: Study using LogTags in Papua New Guinea identified commercial shippers as the weakest link in the cold chain







Finding used to inform conversations with commercial shippers to improve cold chain

**Solution:** Vaccine Vial Monitors (VVM) provide an easy way to monitor if vaccines have been outside cold chain too long

The sticker on the vials changes color when exposed to heat letting vaccinators know if they vaccine has lost its potency and should be discarded



### The vaccine vial monitor...

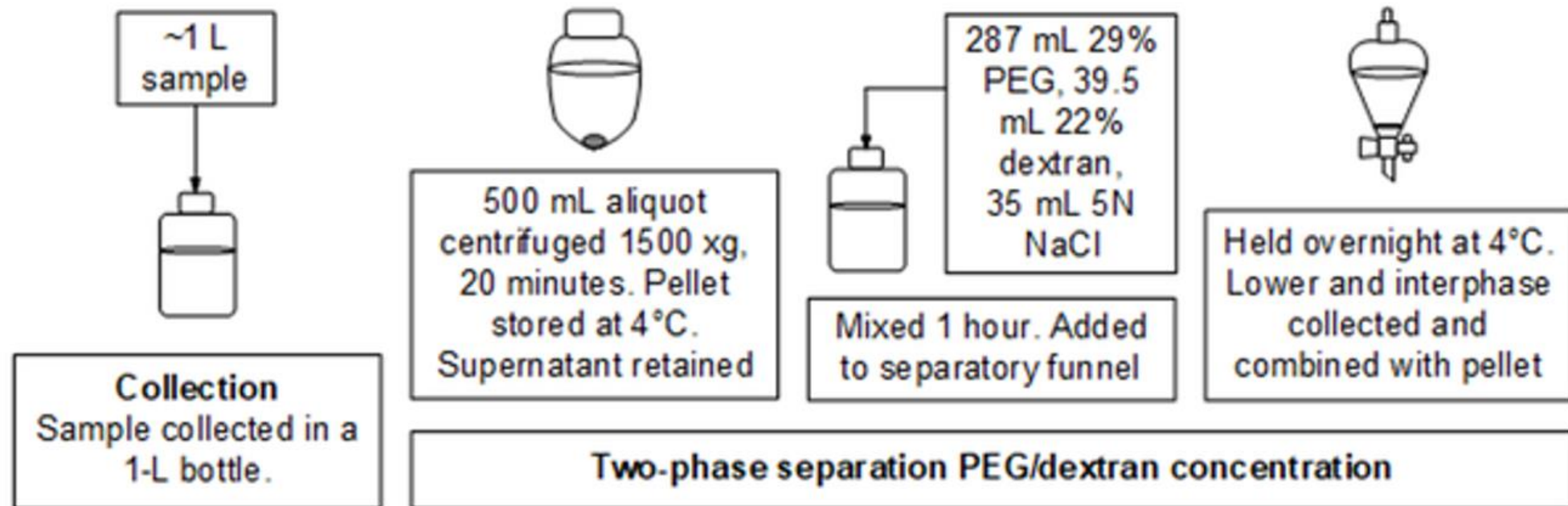
-  ✓ Inner square lighter than outer ring. If the expiry date has not been passed, USE the vaccine.
-  ✓ At a later time, inner square still lighter than outer ring. If the expiry date has not been passed, USE the vaccine.
-  ✗ **Discard point:** Inner square matches colour of outer ring. DO NOT use the vaccine.
-  ✗ **Beyond the discard point:** Inner square darker than outer ring. DO NOT use the vaccine.



**Challenge:** As we move to eradication, we need more environmental surveillance to detect virus without reliance solely on surveillance of paralytic cases



program uses the “Two Phase” method for separation of ES samples



# Solution: Development of cheaper, more sensitive CaFÉ method

11  $\mu\text{m}$  filter  
Collect sediment and extract  
virus



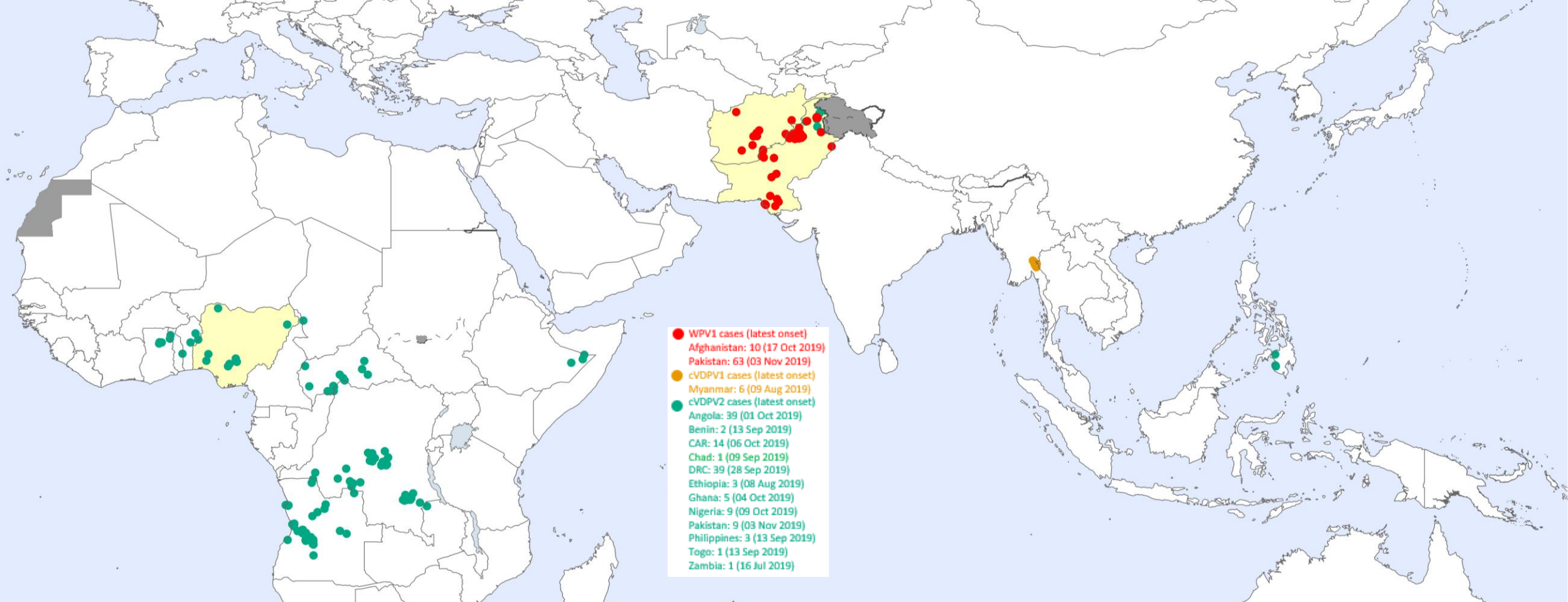
# CaFÉ method is more sensitive than traditional Two Phase testing (n=119)

CaFÉ	Two Phase NPEV*	
	Positive	Negative
Positive	63	23
Negative	2	13

CaFÉ	Two Phase Polio	
	Positive	Negative
Positive	3	14
Negative	1	101

The CaFÉ method detected 23 additional non-polio enteric virus and 14 additional polio viruses than the traditional Two-Phase method

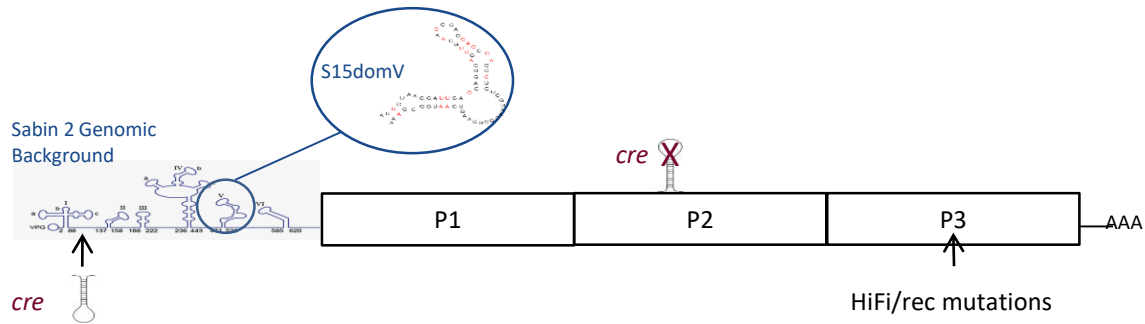




**Challenge:** Oral Polio Vaccine (OPV) can, in very rare instances, revert back to its neurovirulent form and cause outbreaks of cVDPV in under-immunized populations

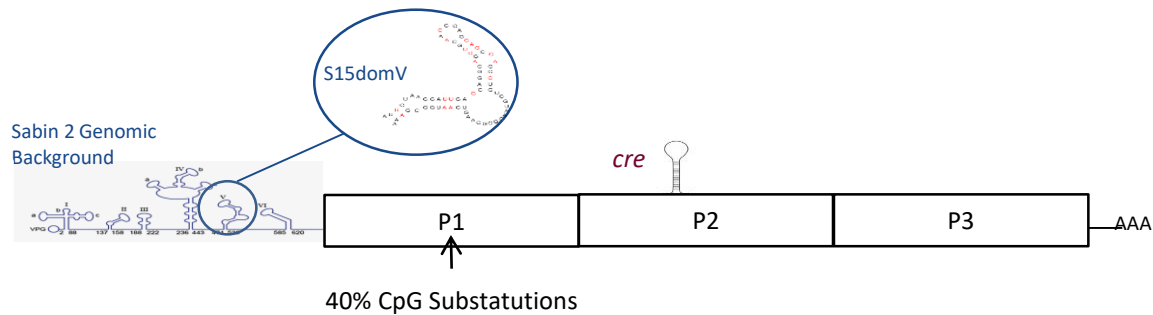
Globally we are responding to outbreaks in 16 countries due to cVDPV2

# Solution: Development of novel OPV (nOPV) which is genetically stable and less prone to reversion to neurovirulence



## nOPV2 – 1b

1. 5' UTR modification
2. Cre relocation
3. HiFi mutations
4. Rec mutations











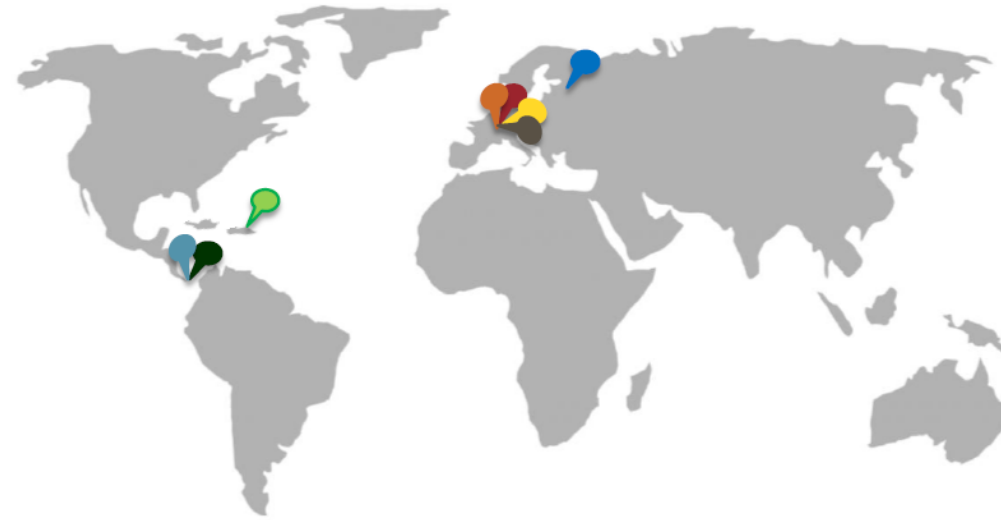
## nOPV2 – 2a

1. 5' UTR modification
2. CpG substitutions
3. Cre NOT moved

Two vaccine candidates use the original Sabin vaccine strain as a backbone

Attenuated virus is mutated to be more genetically stable

	Study	FSFV	LSLV
	<b>M1 *</b> <i>Belgium, adults</i>	Jan 25 2016	May 31 2016
	<b>M2 *</b> <i>Panama, kids/infants</i>	<b>Oct 23 2015</b>	Apr 27 2016
	<b>M3</b> <i>Lithuania, kids</i>	Jan 7 2016	Apr 30 2016
	<b>T1</b> <i>Belgium, adults</i>	Dec 7 2015	May 3 2016
	<b>T2</b> <i>Dominican Rep, kids/infants</i>	Nov 30 2015	Mar 2 2016
	<b>M4a *</b> <i>Belgium, adults, contained</i>	May 16 2017	Oct 27 2017
	<b>M4 *</b> <i>Belgium, adults</i>	Oct 15 2018	May 08 2019
	<b>M5 *</b> <i>Panama, kids/infants</i>	<b>Dec 7 2018</b>	Sep 28 2019



\* Clinical trials prioritized for clinical development, with focus on EUL submission

nOPV2 now in clinical trials around the world and Emergency Use  
Licensure for outbreaks targeted for mid 2020

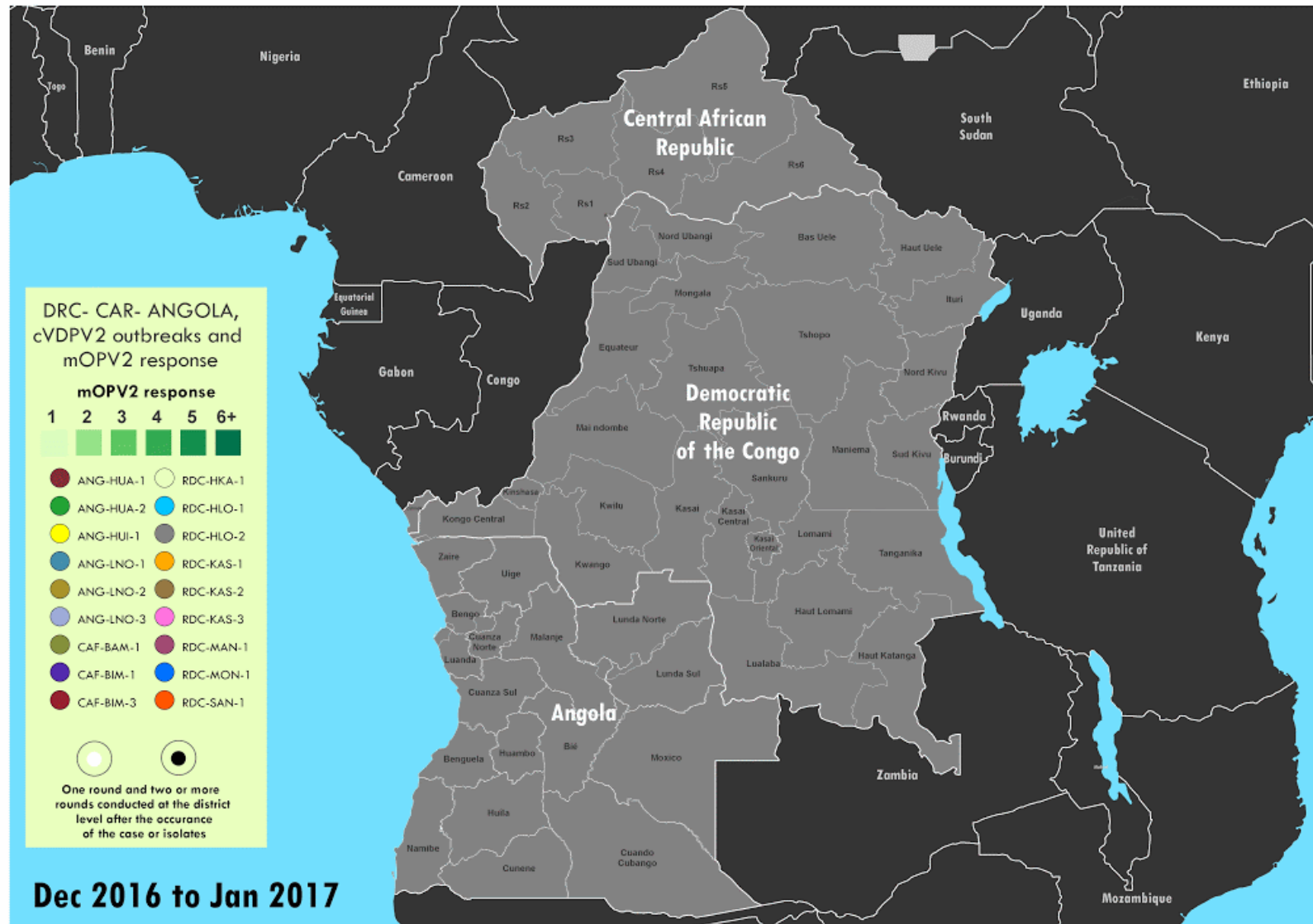
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**But, in the last mile challenges remain**



# Call to Action: Can we better anticipate cVDPV2 outbreaks?



Can machine learning help us predict the next emergence of cVDPV2?

Can big data solutions help us process data faster for improved decision making and more efficient responses?



Thank you

