Global Polio Eradication: Progress and Prospects

ACIP Meeting February 26-27, 2020





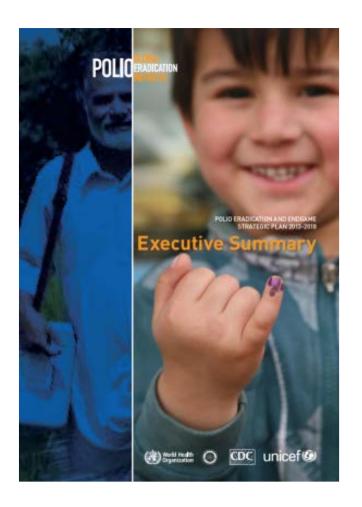






Polio Eradication and Endgame Strategy

- 1. Poliovirus detection & interruption
- 2. OPV2 withdrawal, IPV introduction, immunization system strengthening
- 3. Containment & Global Certification
- 4. Transition Planning

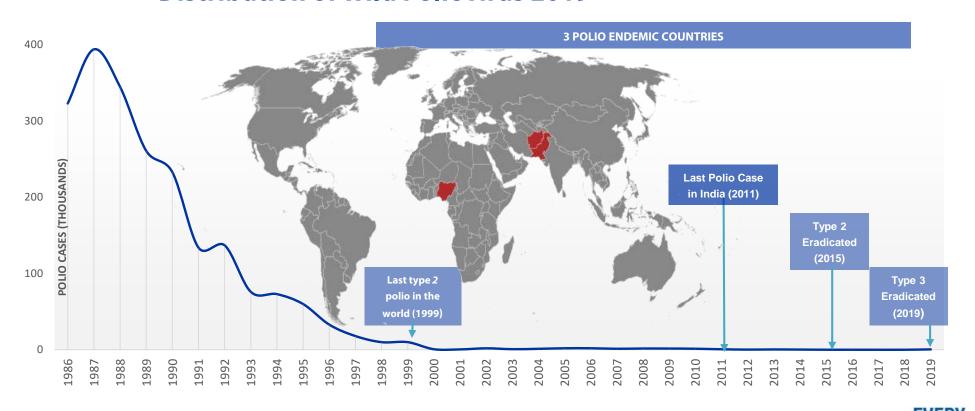






GLOBAL UPDATE

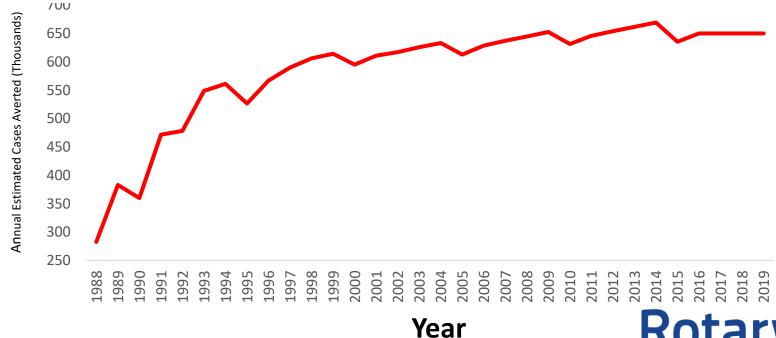
Distribution of Wild Poliovirus 2019





Annual Number of Polio Cases Averted Globally, 1988-2019







Source: WHO/CDC







WPV3 Eradication Certified





The "good" news

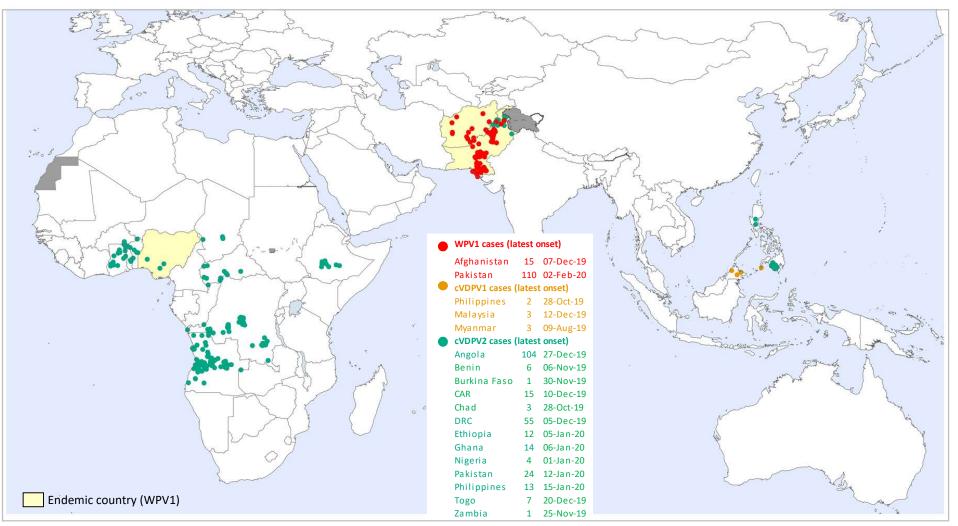
- 7+ years have passed without detection of wild poliovirus type 3
 - GCC certified WPV3 eradication on 17 October 2019
- The number of inaccessible children in formerly Boko Haram controlled areas in Borno State, Nigeria, has been drastically reduced
- 3+ years have passed with detection of any wild poliovirus in Africa, and African Regional Certification Commission will convene in June 2020 to determine regional certification
- IPV supplies are now sufficient for routine immunization, and catch-up of missed cohorts is in progress
- EURO, PAHO, SEARO remain polio-free (incl. cVDPV2)
- Gavi (the Vaccine Alliance) has joined GPEI





Global WPV1 & cVDPV Cases¹, Previous 6 Months²





¹Excludes viruses detected from environmental surveillance

²Onset of paralysis: 19 Aug.2019 – 18 Feb. 2020

Data in WHO HQ as of 18 Feb. 2020

The "bad" news

- Wild type 1 cases increased from 33 cases in 2018 to 173 cases in 2019
- The Taliban ban on <u>house-to-house</u> vaccination in Afghanistan is severely affecting the ability of the program to carry out campaigns
- In Pakistan, a new government is starting to provide national leadership but >6 months passed in 2nd half of 2019 without large-scale vaccination campaigns and wild polio cases surged
- AFRO, EMRO and WPRO battle outbreaks of type 2 circulating vaccine-derived poliovirus (cVDPV2)

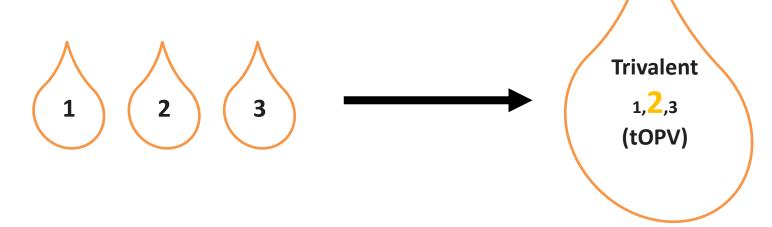
Polioviruses Can Rarely Regain Ability to Cause Paralysis



- Polioviruses in trivalent OPV are attenuated wild polioviruses (WPVs)
- Attenuation results in:
 - Markedly less ability to cause paralysis than WPV
 - Less capacity to pass from person to person than WPV
 - Similar induction of antibodies as WPV
- OPV polioviruses in areas with low polio vaccine coverage can rarely mutate during prolonged circulation and become vaccine-derived polioviruses (VDPVs) able to spread and cause paralysis (circulating VDPVs, cVDPV)



Preventing Circulating Type 2 Vaccine Derived Polioviruses ^

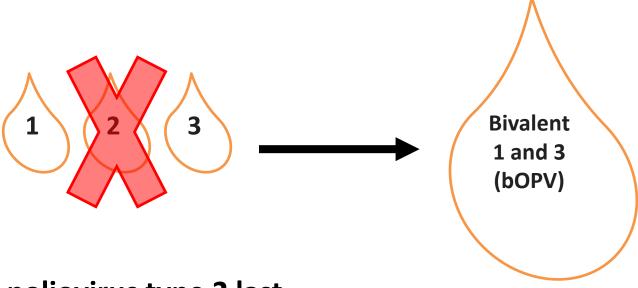


- 700 paralytic cases due to type 2 cVPDV polioviruses confirmed during 2001-2015
- Prompted strategic decision to withdraw OPV2 use in all routine and supplementary immunization activities





Globally Coordinated Switch from tOPV to bOPV in 2016



Wild poliovirus type 2 last isolated in 1999, certified eradicated in 2015

155 countries switch in April 2016



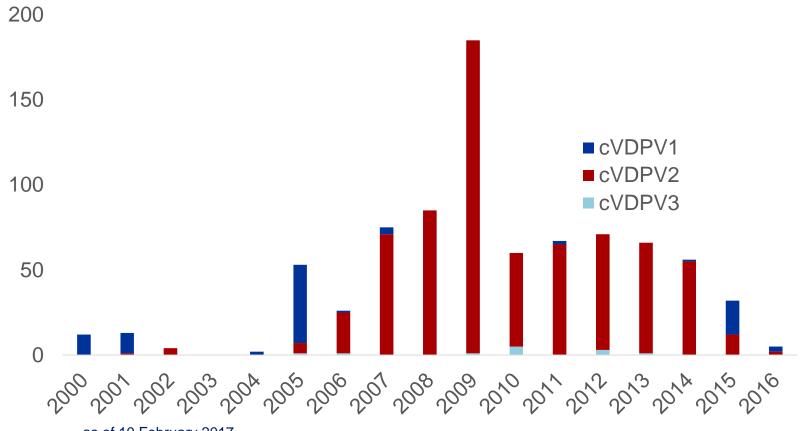


Why Introduce IPV?

- IPV complements tOPV by increasing immunity to all three types of polioviruses, prepares for withdrawal of all OPV
- After the switch:
 - IPV will provide protection against paralysis from type 2 polioviruses (in those reached and who seroconvert)
 - In previous OPV2 recipients, IPV will boost intestinal immunity to infections with type 2 polioviruses
 - Strategic use of IPV in response to type 2 poliovirus outbreaks alongside monovalent OPV 2 (mOPV2) will increase population protection from paralysis



In 2016, fewer cVDPVs than in over a decade



as of 10 February 2017

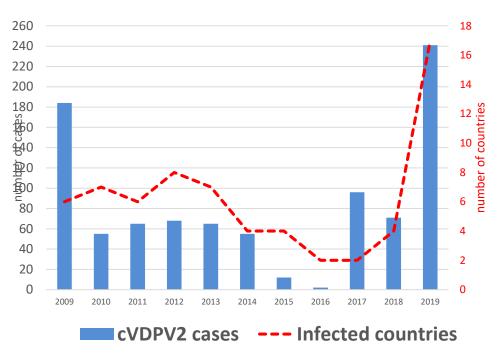
(current numbers: http://www.polioeradication.org/Dataandmonitoring/Poliothisweek.aspx)



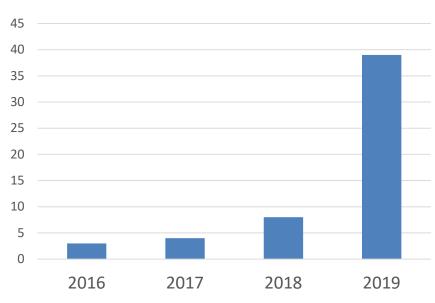


In 2019, the Number of cVDPV2 Cases and Infected Countries Sharply Increased

Number of cVDPV2 cases and infected countries, 2009-2019



Number of cVDPV2 outbreaks, 2016-2019



■ Number of outbreak (ongoing and new)





cVDPV2 outbreaks

- Several outbreaks have been terminated after successful implementation of at least two mOPV2 rounds
- However:
 - To stop outbreaks, many responses required >4 rounds
 - Many new emergences are occurring across the African region due to low quality responses with mOPV2
 - Increasingly, outbreaks are occurring in areas where mOPV2 has not been used
- Caused by :
 - Decreasing population mucosal immunity since OPV2 withdrawn in 2016
 - Population movement



More "bad" news An evolving new challenge

- The program is battling many outbreaks of cVDPV2 in Sub-Saharan Africa → and at risk of re-establishing poliovirus type 2 endemicity in Africa
- Detection of cVDPV2 outbreaks in Asia (China, Pakistan and the Philippines) may herald a global emerging problem
- Limited supply in global mOPV2 stockpile requires balancing use with availability of new shipments



HEALTH

'The switch' was supposed to be a major step toward eradicating polio. Now it's a quandary

By HELEN BRANSWELL @HelenBranswell / SEPTEMBER 13, 2019



A child is vaccinated against polio in Kajiado, Kenya.

hree years ago, the leaders of the international campaign to eradicate polio pulled off a landmark feat, <u>phasing out a problematic component of the vaccine</u> used in developing countries, and introducing a newer version that they hoped would put the world on a better footing to finally eliminate a global scourge.

Now, some organizers are weighing whether "the switch," as the process was known, needs to be reversed.

If it's not, some fear, the world could face a heightened risk of spread of the disease, currently confined to its last redoubt, Pakistan and Afghanistan.

Quandary (definition):

 a state of perplexity or uncertainty over what to do in a difficult situation



Way forward -- cVDPV2

- Prevent cVDPV2 spread into new geographies
 - Rapid deployment of mOPV2
 - Revised strategy guidance for control of cVDPV2 finalized in January 2020
 - Increase scope and quality of mOPV2 SIAs with surge in technical support
- Accelerate development & regulatory review & use of novel OPV2 → Emergency Use Listing (EUL)







- nOPV2 is a genetic modification of the existing OPV type 2
- The modifications made are designed to improve genetic stability of OPV
- This will in turn
 decrease the risk of
 seeding new cVDPVs
 and the risk of VAPP
 when deployed for
 cVDPV2 outbreak
 response

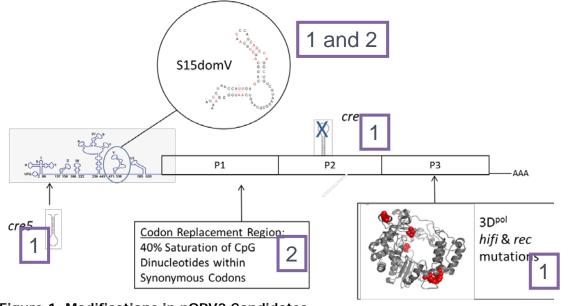


Figure 1: Modifications in nOPV2 Candidates

Sabin 2 genome is depicted showing the 5' untranslated region (UTR) in grey shading, polyproteins (P1-3), 3' UTR and polyA; locations of modifications within the genome are shown. Nucleotide differences between Sabin 2 and S15 domain V are shown in red.

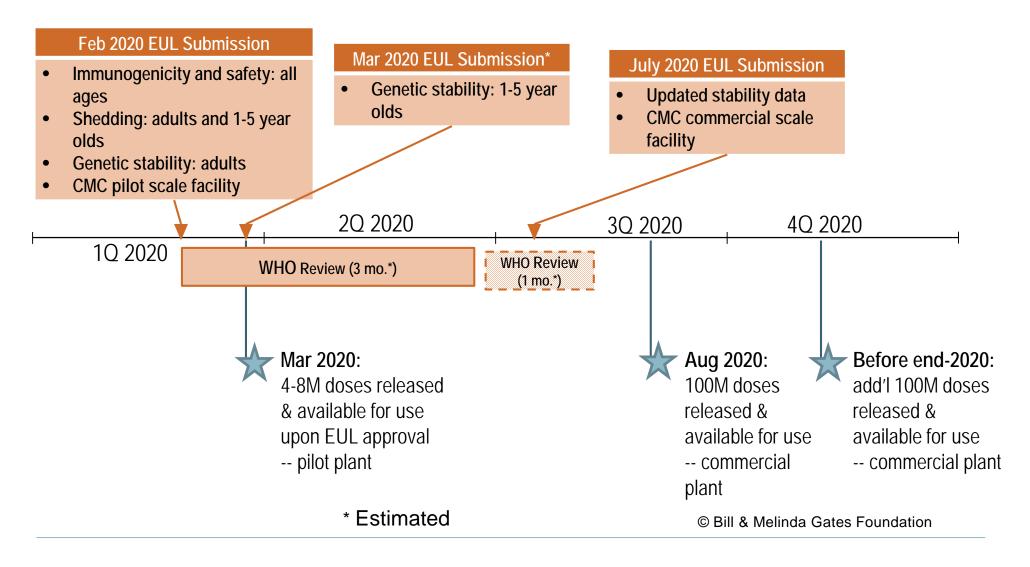


Accelerating Regulatory Approval through EUL

- Owner: WHO Essential Medicine Department (EMP, PQ)
- Goal: make "experimental" health products available for emergency response
- Products listed under EUL so far: 0
- Eligibility criteria nOPV2: poliovirus spread is a Public Health Emergency of International Concern (PHEIC)
- Fastest way to obtain regulatory review and approval



Ramp-up of nOPV2 Clinical Development and Production to Align with EUL Approval



Summary

- Polio eradication made some progress in 2019, but encountered serious challenges
- Wild poliovirus eradication requires access in Afghanistan and vaccination quality improvements/accountability in Pakistan
- cVDPV2 outbreaks threaten the success of "switch" and may lead to re-establishment of type 2 endemicity
- mOPV2 needs to be replaced as soon as feasible by genetically more stable novel OPV2
- A 2nd dose of IPV in Routine Immunization is under discussion when supplies allow
- Securing the funds to run the program is a very high priority



