Sub-national estimation of surveillance sensitivity to inform declaration of elimination A retrospective validation against the elimination of wild poliovirus in Nigeria

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Near elimination, a surveillance system yields an increasing proportion of *negative* observations





■ Positive ■ Negative

Near elimination, a surveillance system yields an increasing proportion of *negative* observations





Positive Positive - undetected Negative

How confident are we that infection is absent given the system is returning negatives?





This is the **negative predictive value** of the surveillance system



To infer this, we need to understand the **sensitivity** of the surveillance system for detecting infection

For polio, the surveillance system comprises of both *case-based* and *environmental* surveillance

Indicators of "performance" are routinely monitored,
highlighting variability over time and space
➢ How can this be incorporated into interpretation?



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→ Extend an approach developed in animal health to model surveillance via scenario trees (*Martin et al. 2007*).





Components of polio surveillance







Acute Flaccid Paralysis

- Detect *symptomatic* infection in an individual
- Implemented nationally all notified AFP cases tested

Environmental

- Detect symptomatic and asymptomatic infection within catchment
- Limited population coverage















Time-/districtdependent and datainformed estimation of these probabilities



Probability that a single infection yields a <u>positive</u> <u>outcome</u> from the surveillance system







How low a prevalence does the surveillance system <u>need</u> to detect?

- Detecting lower prevalence demands higher sensitivity
- What prevalence do we expect near elimination?
- What prevalence is sufficient to interrupt transmission?

If the *country* is infected, we want sufficient sensitivity that infection is detected in **at least one district**.

If a *district* is infected, we want sufficient sensitivity to detect a **prevalence of 1 per 100,000**.

Example: Elimination of WPV1 in Nigeria



WPV1 was not detected in *any* AFP stool or environmental sample between August 2014 and June 2016.



- **27,600** AFP cases notified
- 1,027 env. samples analysed
 => Zero positives



WPV1 was not detected in *any* AFP stool or environmental sample between August 2014 and June 2016.

In July 2016, four WPV1+ paralytic cases were detected in the northeastern state of Borno.



Surveillance performance: AFP

Across the majority of LGAs, WHO thresholds for **AFP reporting** and **stool adequacy** were consistently met/exceeded.



Surveillance performance: Env

Despite overall expansion, **% LGA population within catchment** was <u>low</u> (~3%)

Detection of nonpolio enteroviruses (NPEV) was adequate on average (~50%), but low in some LGAs



LGAs with no active ES are shown in white

Surveillance sensitivity over time





Summarised across 1,000 iterations

🛱 AFP 🛱 ENV

Freedom from infection: 2014–2016





2016-2020

Since 2016, **ES expanded** while **AFP declined** (lower adequacy of stool collection)

Confidence in freedom from infection **consistent with official declaration**

- > 95% after 34 months
- ~99% by mid-2020 declaration





Undetected circulation in Borno was thought to be due to **conflict and resulting inaccessibility** in the region.

This disruption was not evident in routine surveillance indicators, so doesn't influence our estimates

Population **catchment beyond LGA boundaries** is not incorporated

Extent will likely vary depending on weather conditions and characteristics of the site



Conclusions



AFP and environmental surveillance are **complementary** approaches, and we aimed to quantify their **joint contribution** to evidence of elimination on a national scale.

With this framework, we integrate routine **indicators of performance** into the interpretation of **negative observations**.

- We draw conclusions that are **consistent** with both persistence during 2014-16 and elimination by 2020.
- Supports prospective use for inferring WPV elimination in remaining endemic countries (Afghanistan and Pakistan).

Thank you!



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Further considerations: Choice of prior



We need a starting distribution (prior) for the first time point

- How likely is it that the country is free from infection, given that a positive was observed *last month*?
- Judgment depends on the prior sequence of events



Influence of design prevalence





Influence of catchment radius





Catchment radius • 2km 5km

Static vs time-varying sensitivity





Sensitivity 🔶 Static 🔶 Time-varying