



# DTP1-3 and MCV1 vaccination coverage in the Democratic Republic of the Congo: mapping zero-dose and under-vaccinated children

C. Edson Utazi, PhD, FHEA

(Joint work with H. R. Chamberlain, K. Kunnumpurathu-Sasi, A. Cunningham and A. N. Lazar)

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VaxPop WorldPop



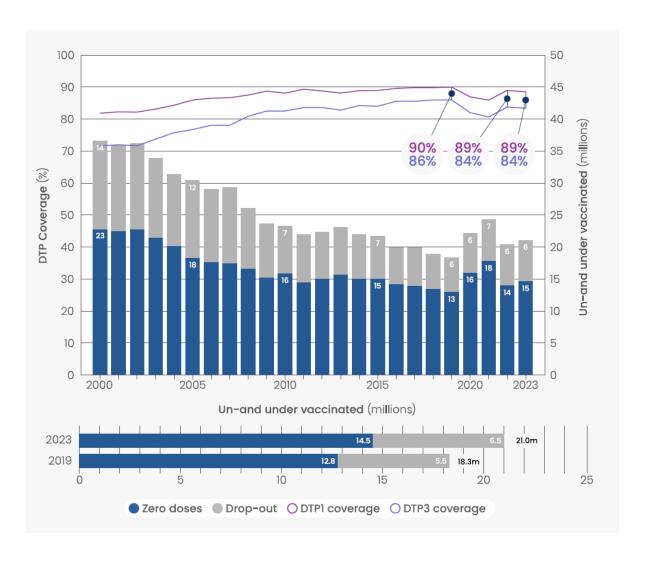




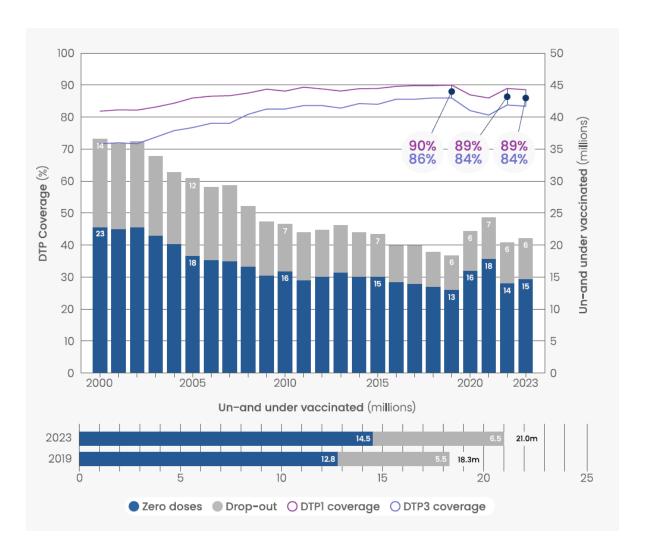


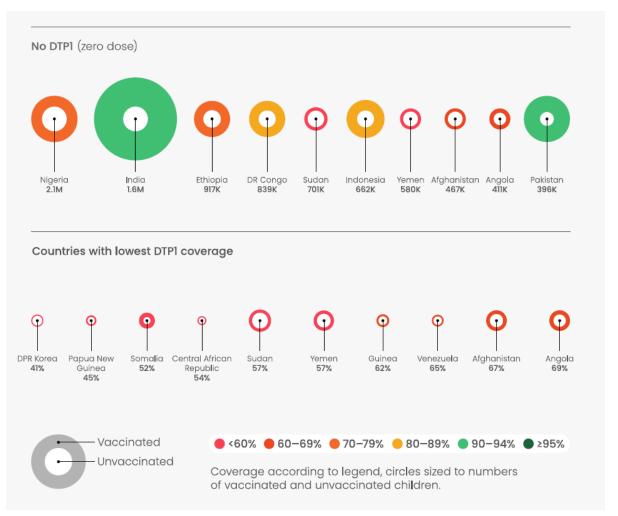


Conflicts of interest
All authors declare no competing interests.

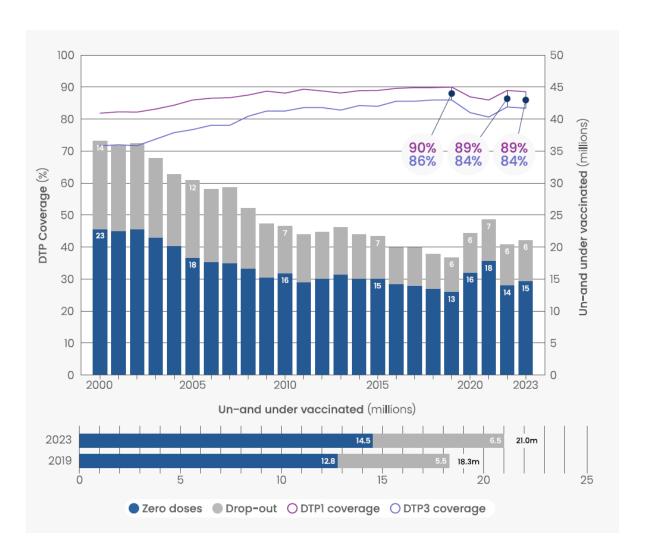


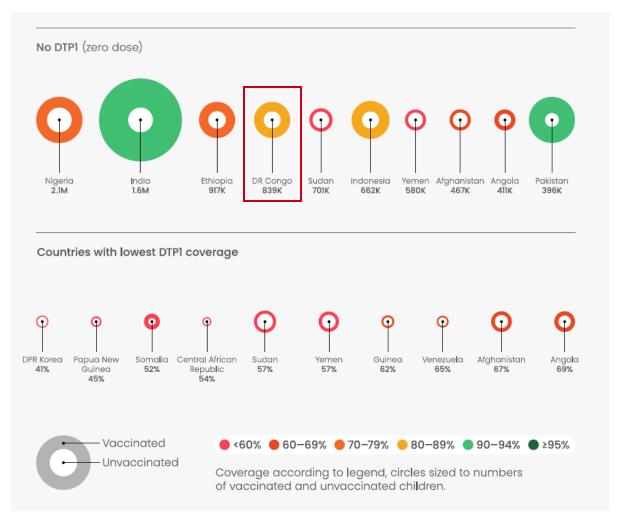




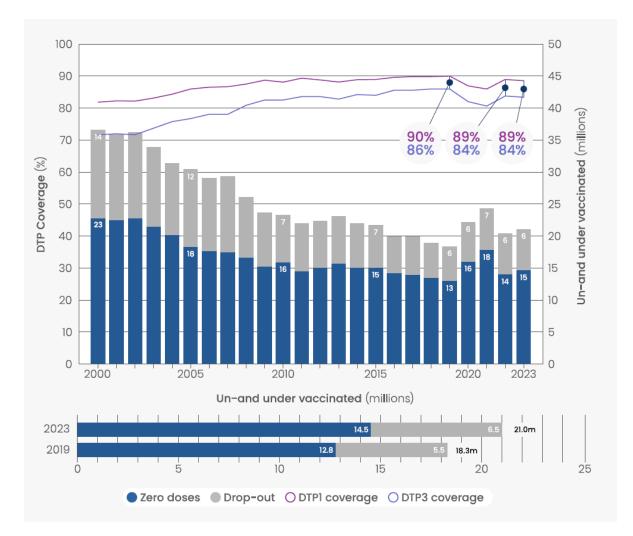


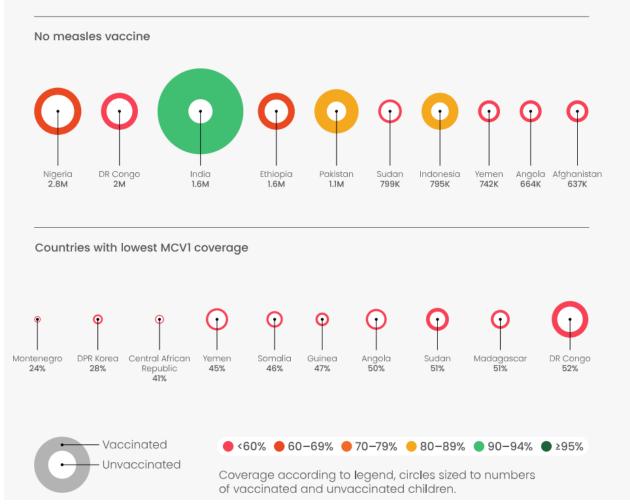




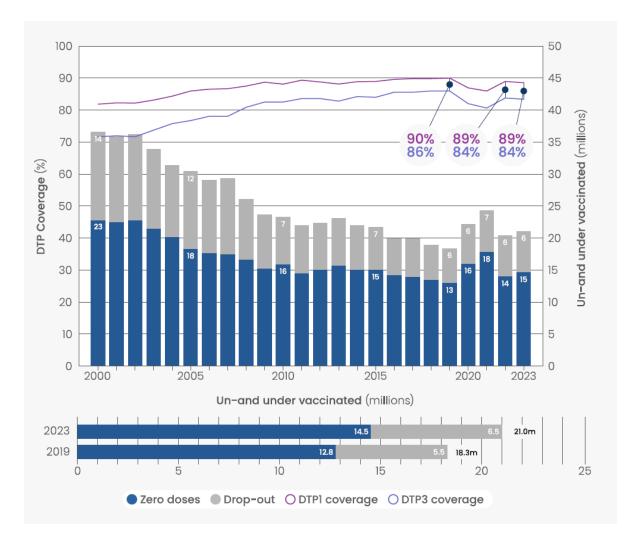


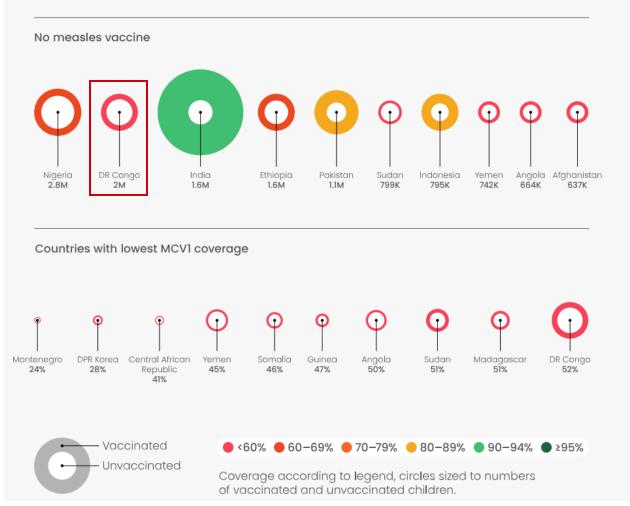




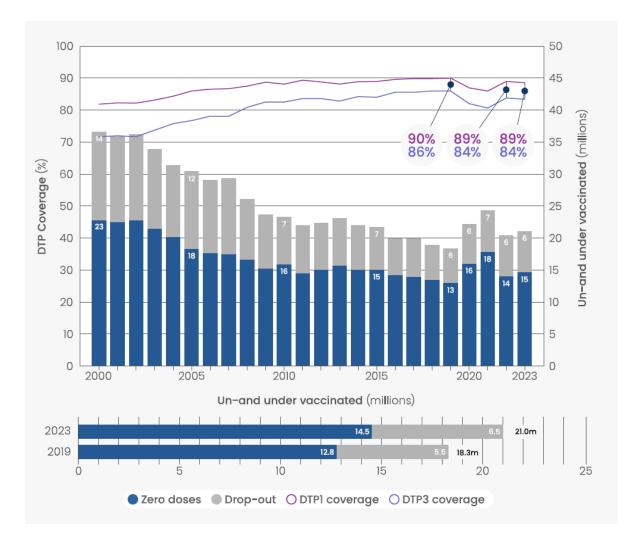


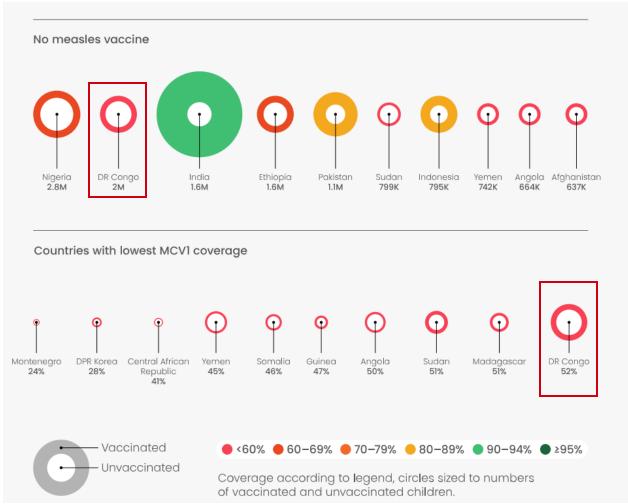






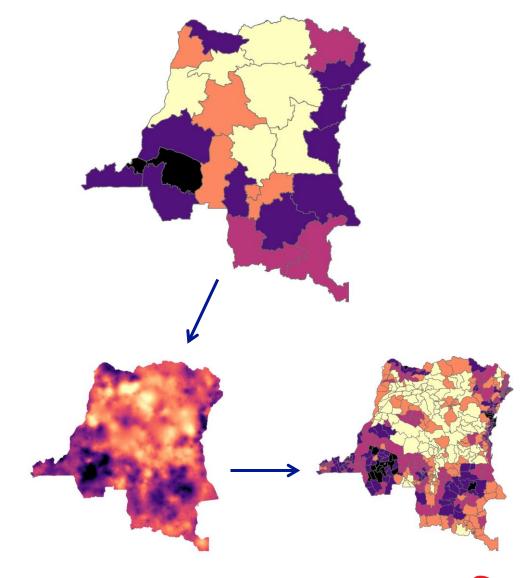








- Over 300,000 measles cases and nearly 6,000 deaths in 2023
- MSF provided vaccination to over 1.6 million children and treatment to over 46,000 patients in 2023
- Survey estimates of vaccination coverage are often available at the national and provincial levels
- To effectively target vaccination campaigns and interventions, spatially detailed estimates are required
- Our work uses geospatial modelling approaches to address this problem





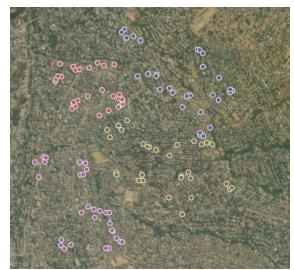
## Methodology - Input vaccination coverage data

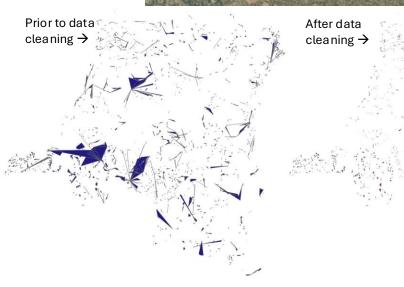
- Geolocated data were obtained from the 2023 Enquete de Couverture Vaccinale (ECV) survey conducted by the Kinshasa School of Public Health (KSPH) across all provinces in DRC
- 80,000+ households surveyed across 2,700+ clusters. Approximately 30 households per cluster.
- Data cleaned to address inconsistences related to individual children aged 12-23 months (nearly 50,000) and errors in household geolocation
- Data were aggregated to the cluster level for DTP1-3 and MCV1



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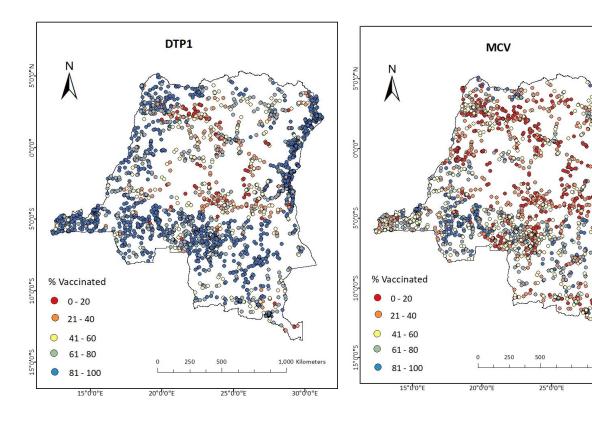






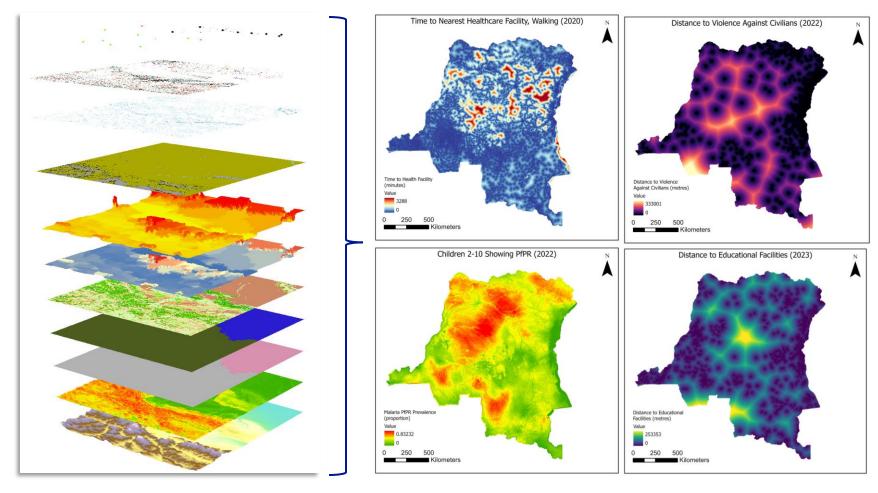
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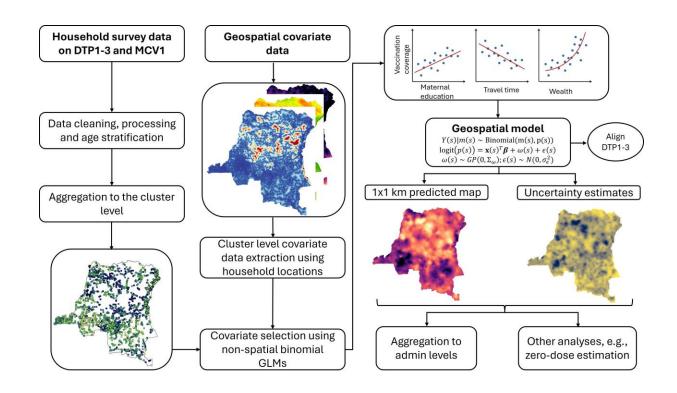
# Methodology - Geospatial covariate data



- 59 Geospatial covariate data were obtained from WorldPop and other sources
- The data were processed and harmonized to 1x1 km resolution.
- Covariate values were extracted at the cluster level using the corresponding processed household geocoordinates.



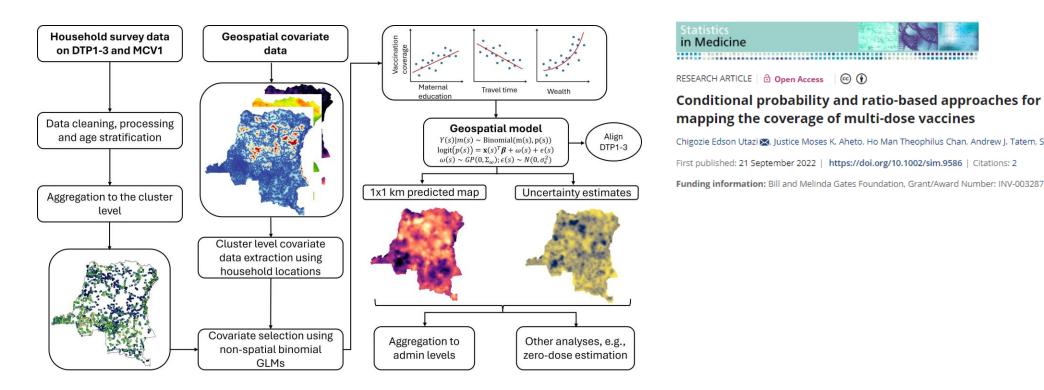
## Methodology - Geospatial model and prediction



- We fitted Bayesian binomial geostatistical models using the INLA-SPDE approach implemented in the R-INLA package.
- We used the ratio-based approach (Utazi et al., 2022) to ensure that  $p(DTP1) \ge p(DTP2) \ge p(DTP3)$ .
- K-fold cross-validation was used to evaluate the accuracy of the modelled estimates at the cluster-level. The modelled estimates were also compared with direct survey estimates at the provincial level.
- Details of model-fitting, validation and prediction can be found in these publications: <u>Utazi et al., 2022</u>; <u>Utazi et al.,</u> 2021.



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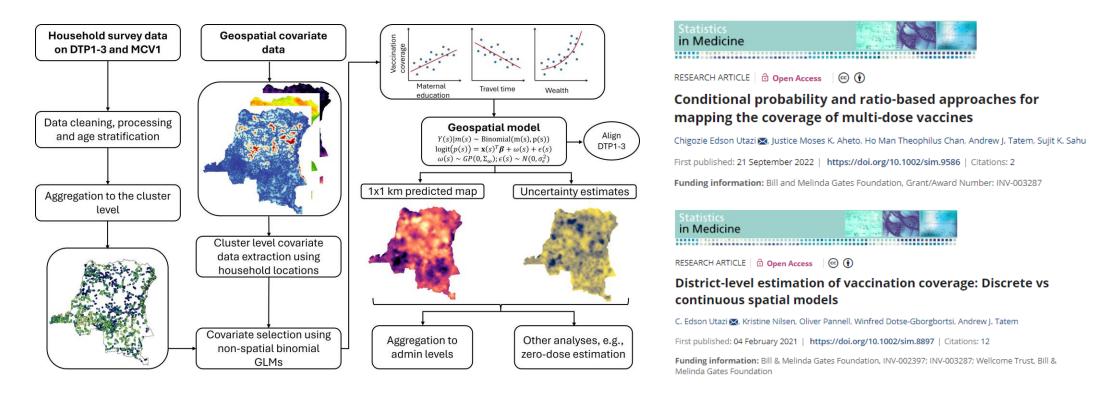




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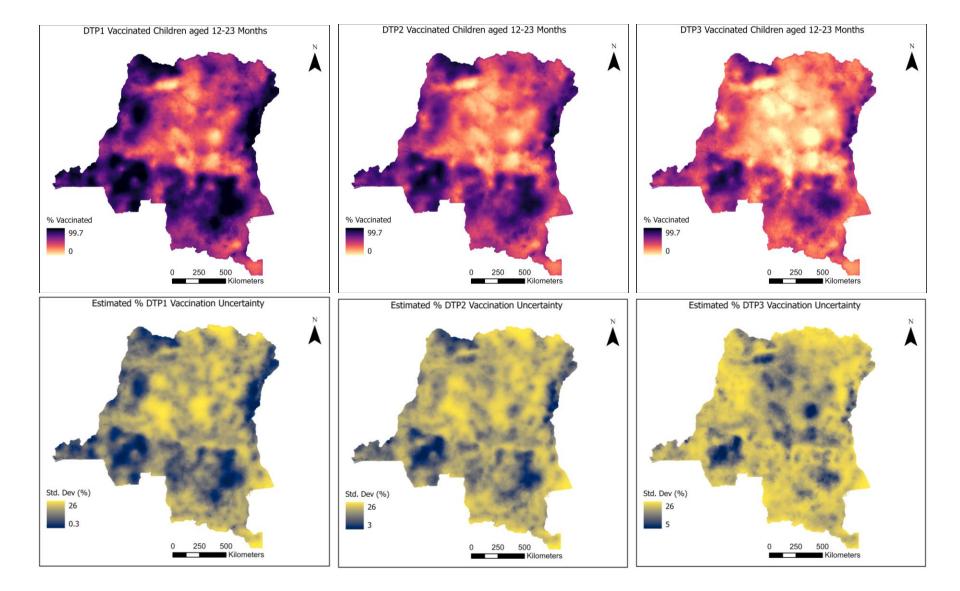
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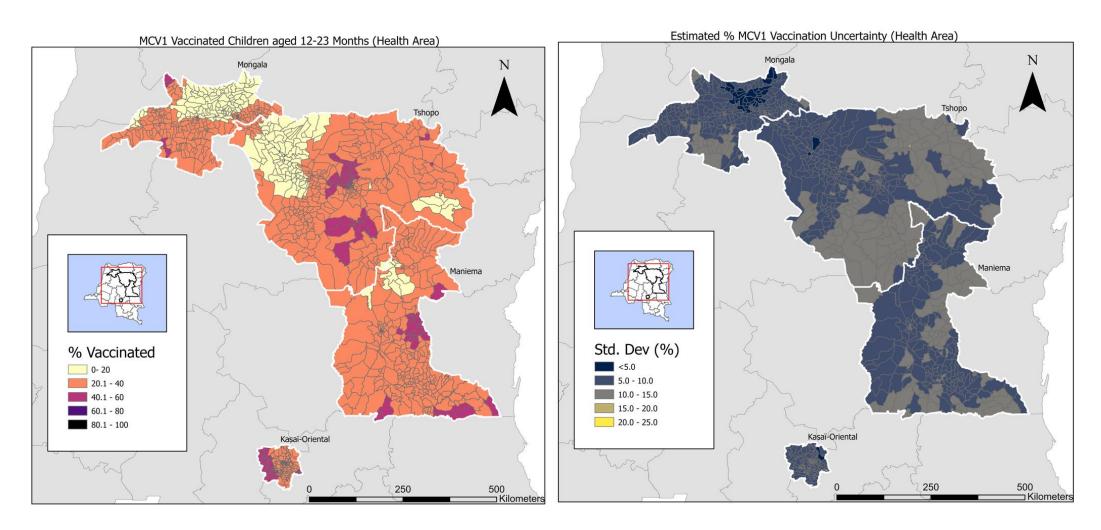


# Predicted DTP1-3 coverage map – 1x1 km resolution





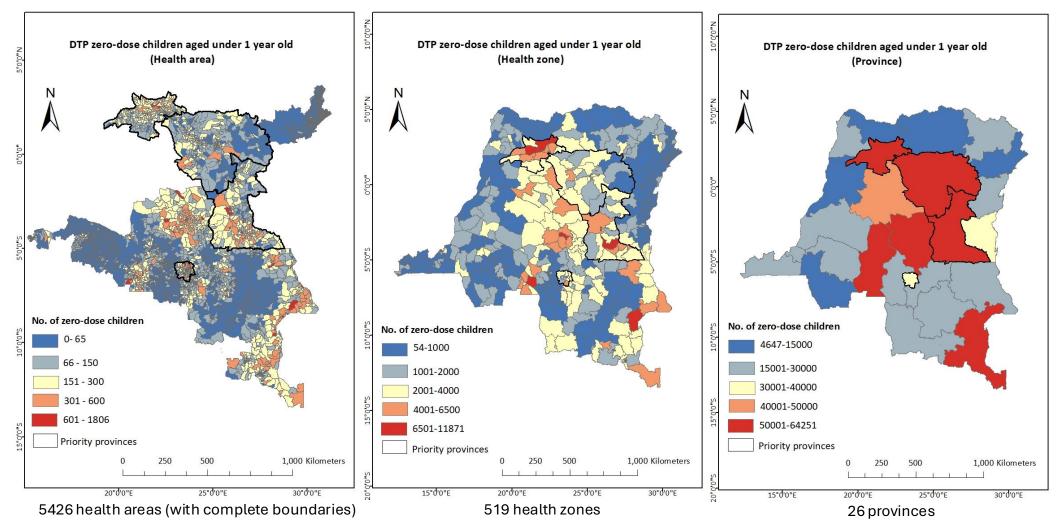
# Predicted MCV1 coverage map – Health areas (4 provinces)



Estimated MCV1 coverage and associated uncertainties at the health area level



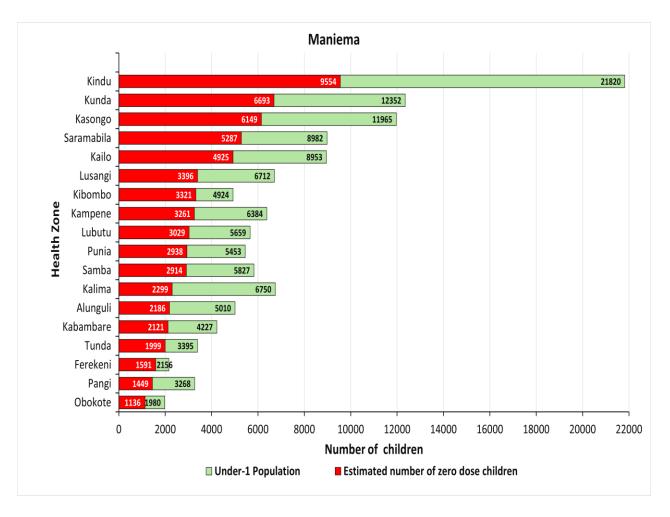
#### Estimates of numbers of DTP zero-dose children

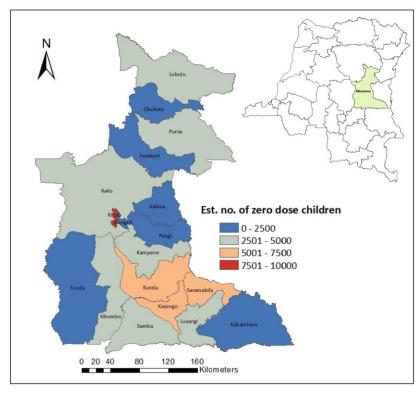






# **Example: Priority areas (health zones) for Maniema province**







## **Summary**

- Substantial heterogeneities exist in the spatial distributions of DTP1-3 and MCV1 coverage in DRC
- DTP1 coverage was lowest in health areas in Mongala, Tshopo, Maniema, Tshuapa and Sankuru provinces
- MCV1 coverage was lowest in areas Sud Ubangi, Equateur, Mongala, Bas Uele, Tshopo, Sankuru and Maniema provinces.
- We found greater dropout rates between DTP2&3 than between DTP1&2. Dropouts between DTP1&3
  were highest in areas of lower coverage in Mongala, Tshuapa, Tshopo, Sankuru and Maniema
  provinces
- Areas with the highest numbers of DTP zero-dose children were estimated in Mongala, Sankuru, Maniema, Kasaï, and Haut-Katanga provinces
- Areas with the highest numbers of MCV zero-dose children were estimated in Kinshasa, Maniema, Kasaï, Mongala, Haut-Katanga, Tanganyika, Sankuru, Kasaï-Oriental and Lomami provinces
- Additional analysis is needed to establish linkage with measles surveillance data and outbreak distribution
- Currently working with partners: GRID3, UNOPS and KSPH, to operationalise findings
- Challenges: (i) ECV not an internationally standardized survey like the Demographic and Health Surveys (DHS); data cleaning to correct GPS coordinates and remove duplicate records was time consuming. (ii) Selection of subnational age/sex pyramids to estimate under 1 population was challenging due to age heaping in available data sources. Four of these were compared and WorldPop projections were used.



# **Acknowledgements**



www.worldpop.org

Email: <u>c.e.utazi@soton.ac.uk</u>; X @chigedson





