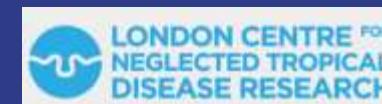




Genomic impact of mass drug administration on human infective *Schistosoma mansoni*

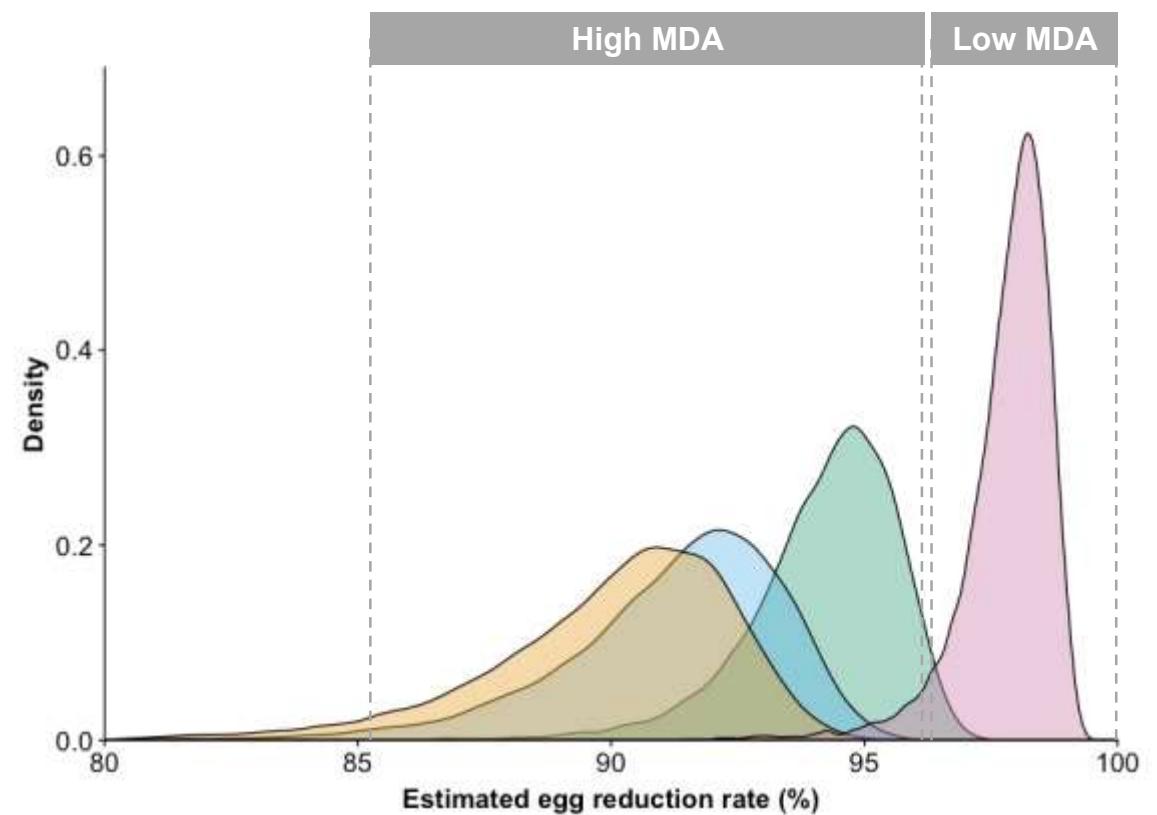
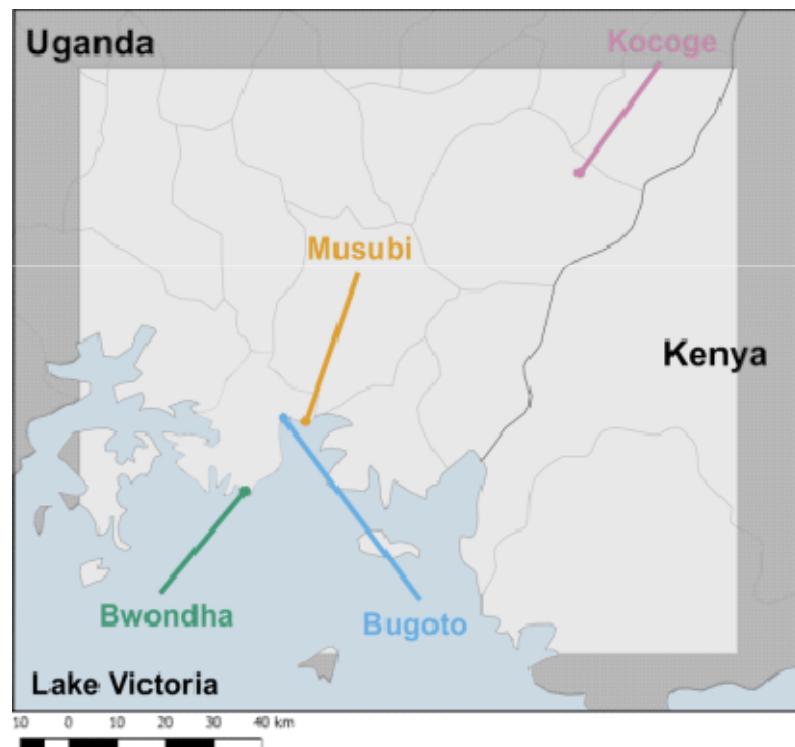
Duncan Berger, Thomas Crelen, Jennifer Noonan, Poppy Lamberton, Magda Lotkowska, Nancy Holroyd, Gabriel Rinaldi, Narcis Kabatereine, Edridah Tukahebwa, Fiona Allan, James Cotton, Joanne Webster, Matthew Berriman

LCNTDR anniversary event - Beyond 2020: Research innovations for a new agenda



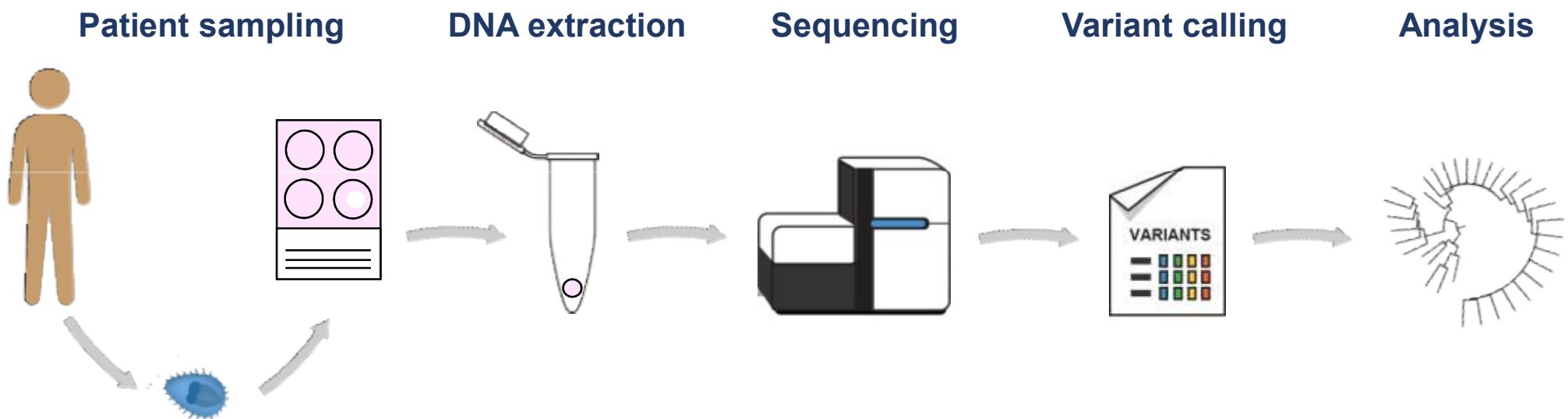


Reduced praziquantel efficacy is associated with multiple rounds of mass-drug administration





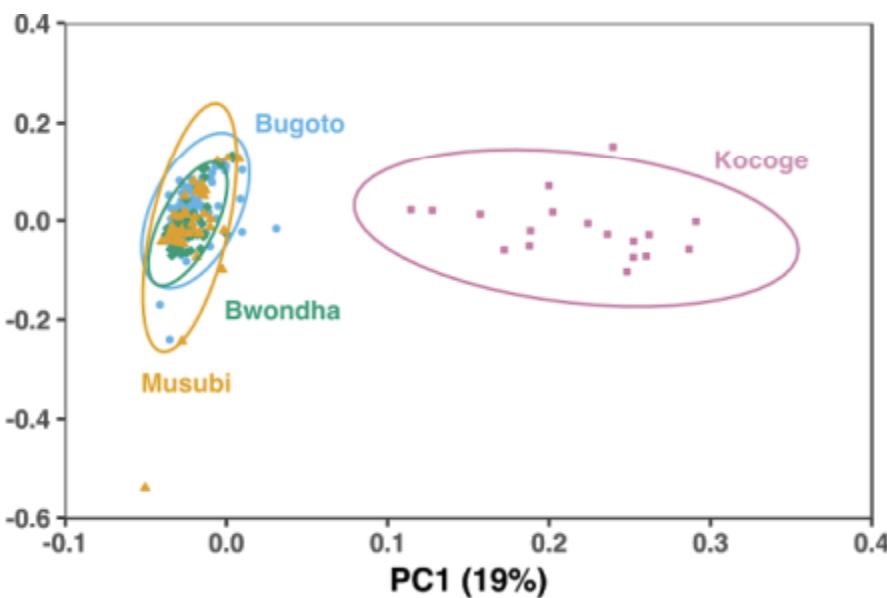
Can whole-genome sequencing provide insights?



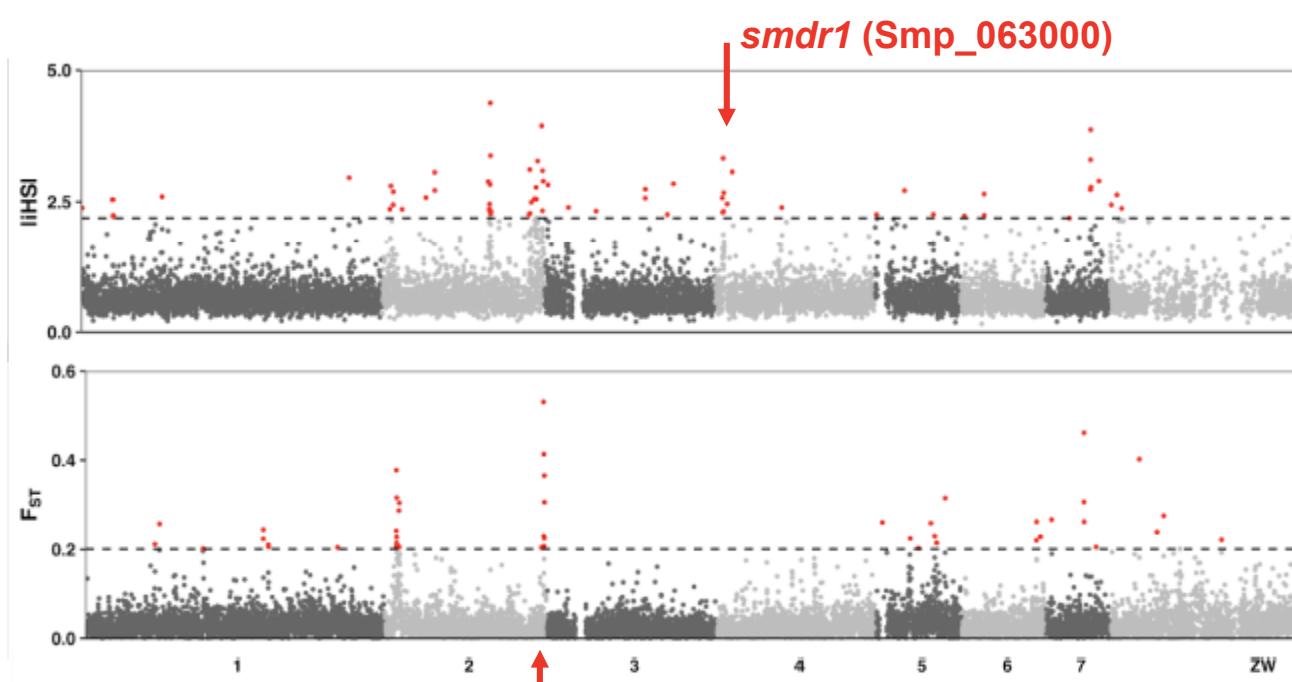


Population structure and signatures of selection

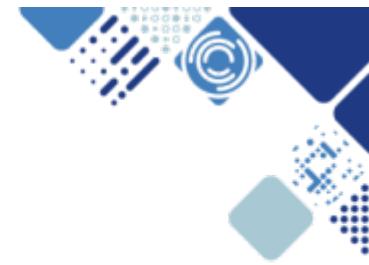
Population structure



Selection



Smp_170450, Smp_336070,
Smp_328710, Smp_094390



Beyond 2020: How can genomics help us understand schistosomiasis?

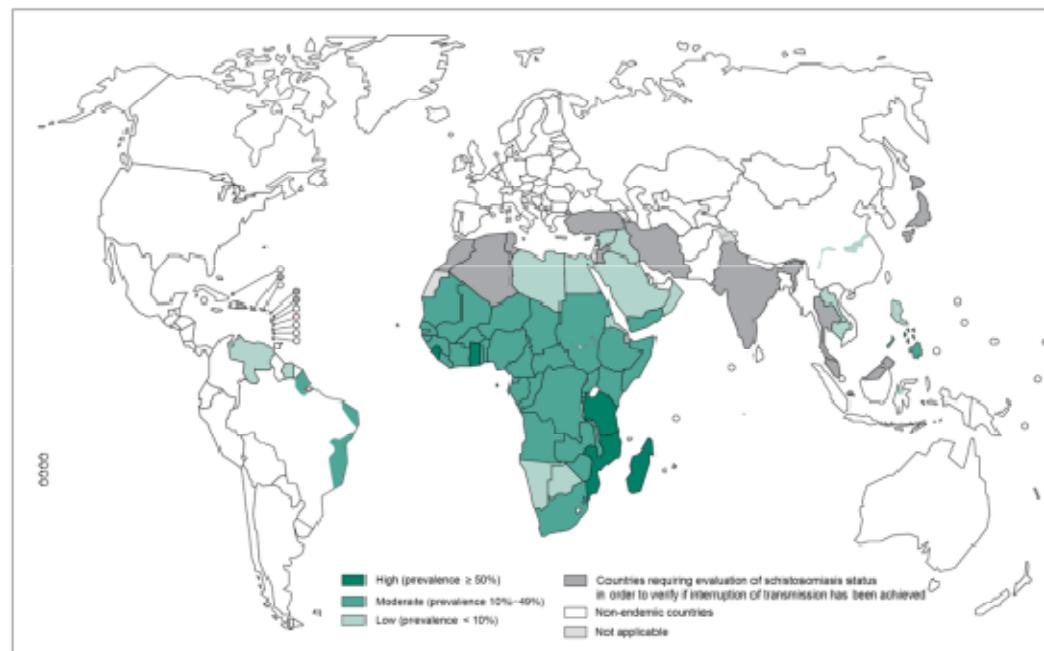
• Genomic surveillance

- Treatment efficacy and praziquantel resistance
- Hybridization (Kincaid-Smith et al., 2018; Oey et al., 2019)
- Diversity of host infrapopulations
- Global diversity

• Comparative genomics

- Evolution of parasitism (IHGC, 2019)
- Novel drug and vaccine targets (IHGC, 2019)

Distribution of schistosomiasis, worldwide, 2012



h, et al., 2018. *bioRxiv*.

2019. *PLoS Pathogens*, 15 (1): e1007513.

Helminth Genomes Consortium. 2019. *Nature Genetics* 51 (1): 163–74.

Schistosomiasis. World Health Organization, <https://www.who.int/schistosomiasis/en/>.





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Aiden Emery

Imperial College London / SCI

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Poppy Lamberton
Tom Crellan
Wendy Harrison
Alan Fenwick

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