



Progress in the control NTDs (human helminth infections) – moving from morbidity control to transmission elimination



Whipworm



Roundworm



Hookworm

Roy Anderson

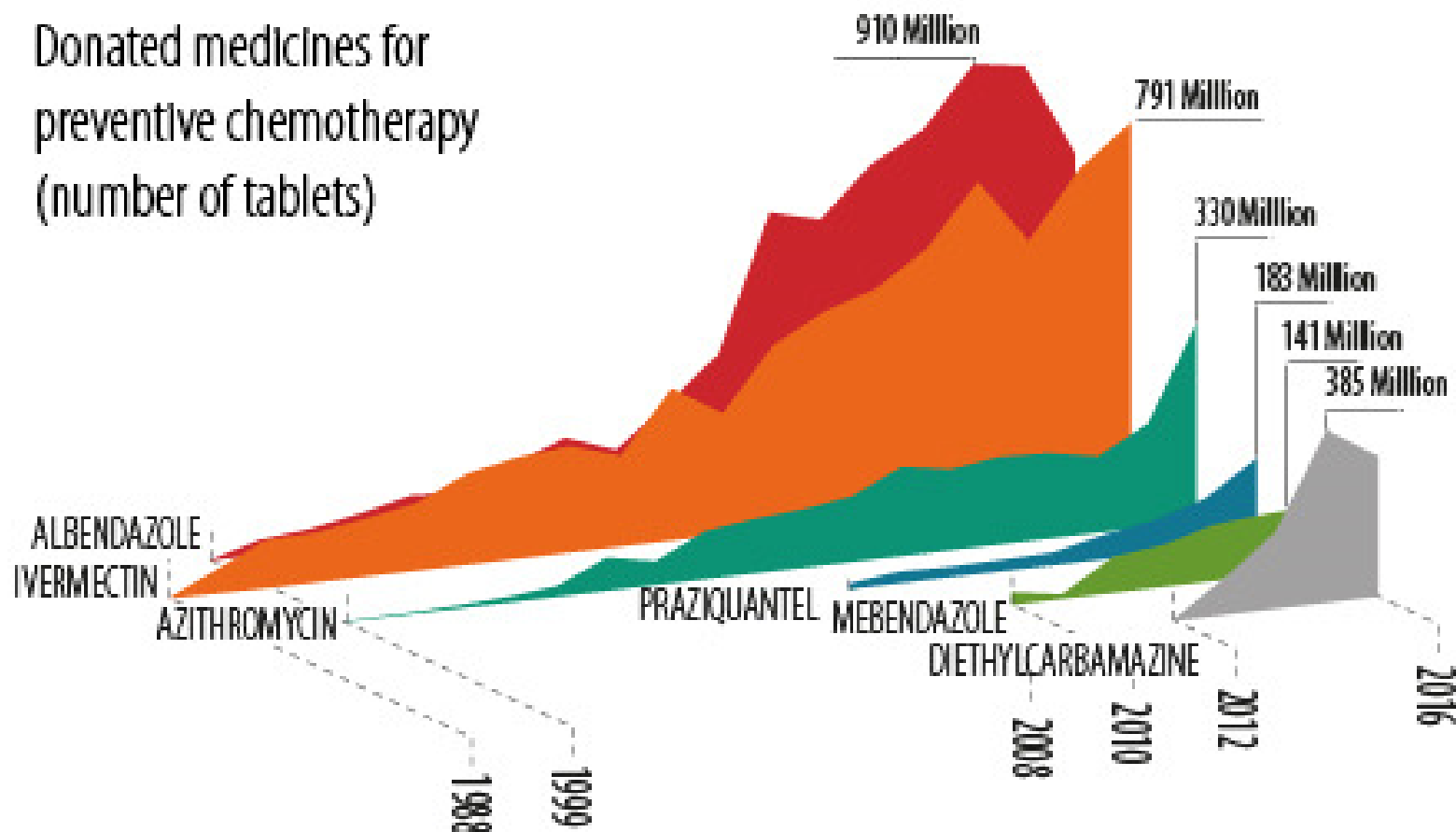
Department of Infectious Disease Epidemiology (DIDE) and the London Centre for Neglected Tropical Disease Research (LCNTDR).

Imperial College London



Drug donations - WHO (2017)

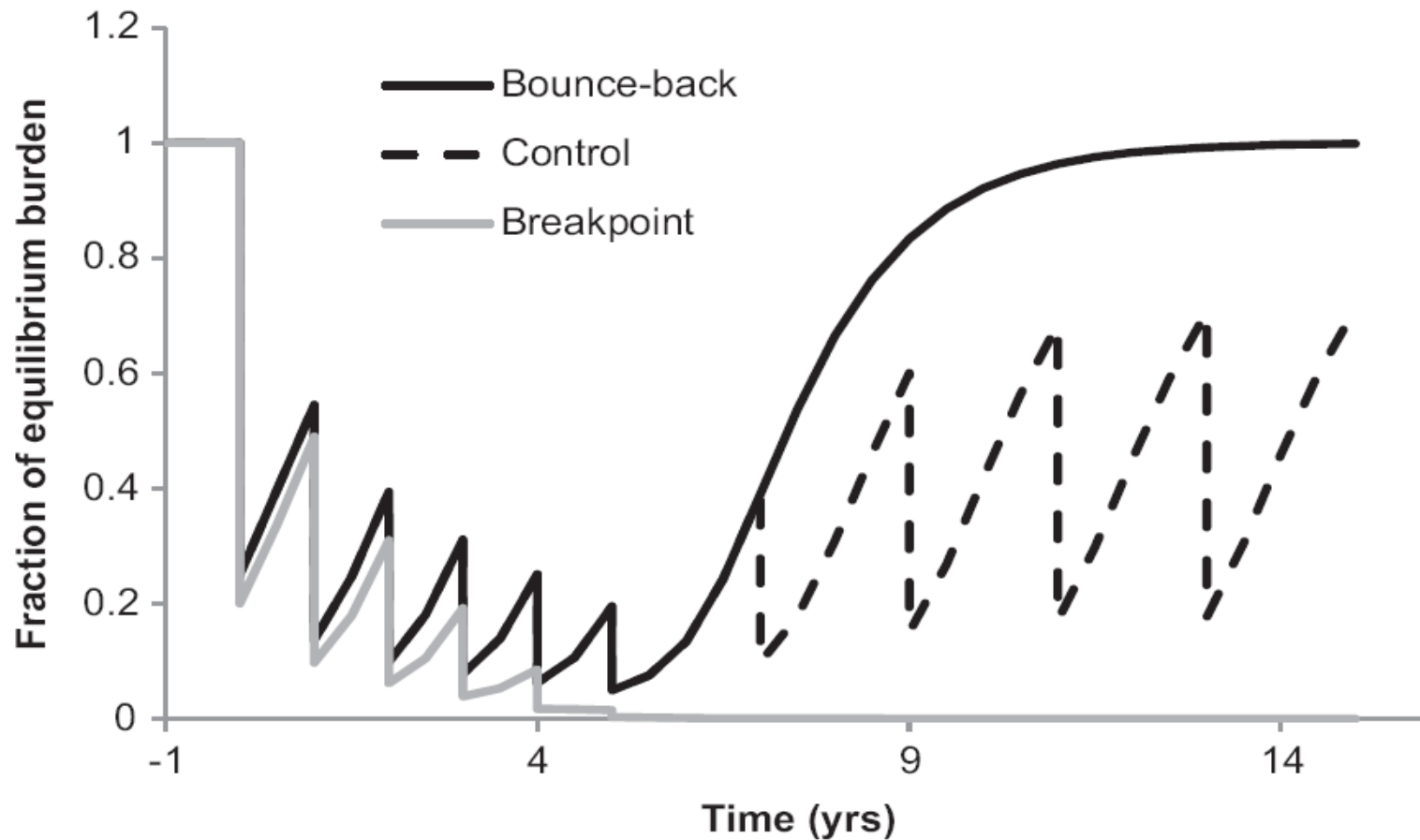
Donated medicines for
preventive chemotherapy
(number of tablets)



**1.288 billion treatments were delivered
to 998 million people in 2015**

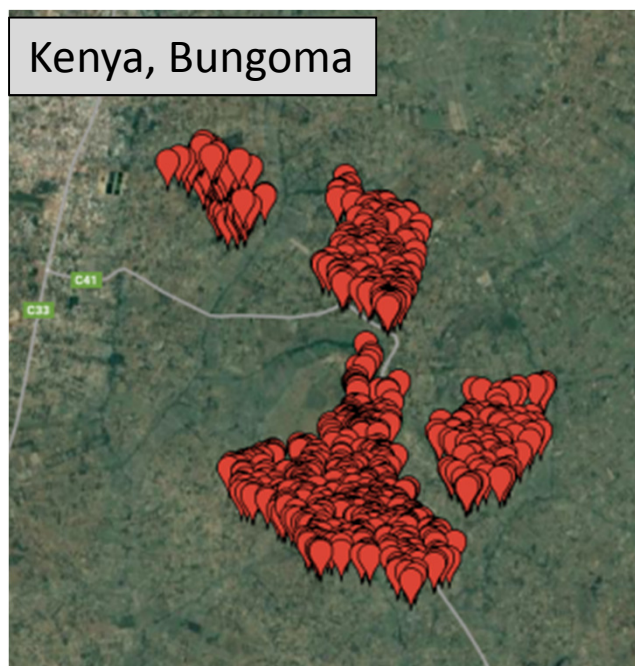


***The effect of mass drug treatment
on the intensity of infection – bounce back if
breakpoint is not crossed – no acquired immunity***

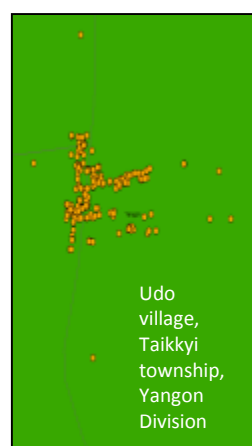




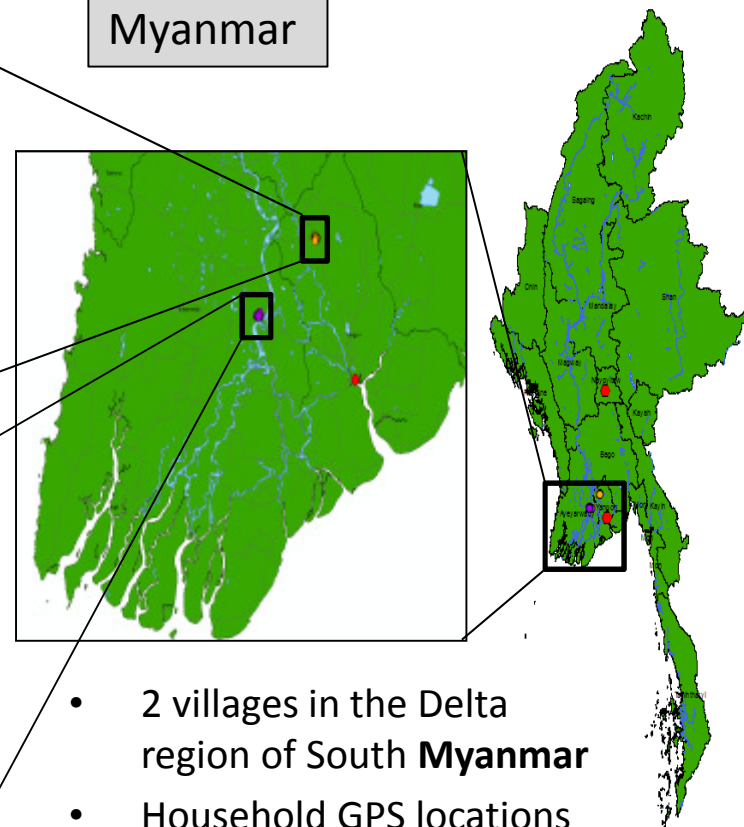
Study sites 1 & 2 in Kenya, Bungoma *[who infects whom], and South Myanmar* *(transmission in heavily treated communities)*



- 5 villages in rural Western **Kenya**
- Household GPS locations mapped



Myanmar

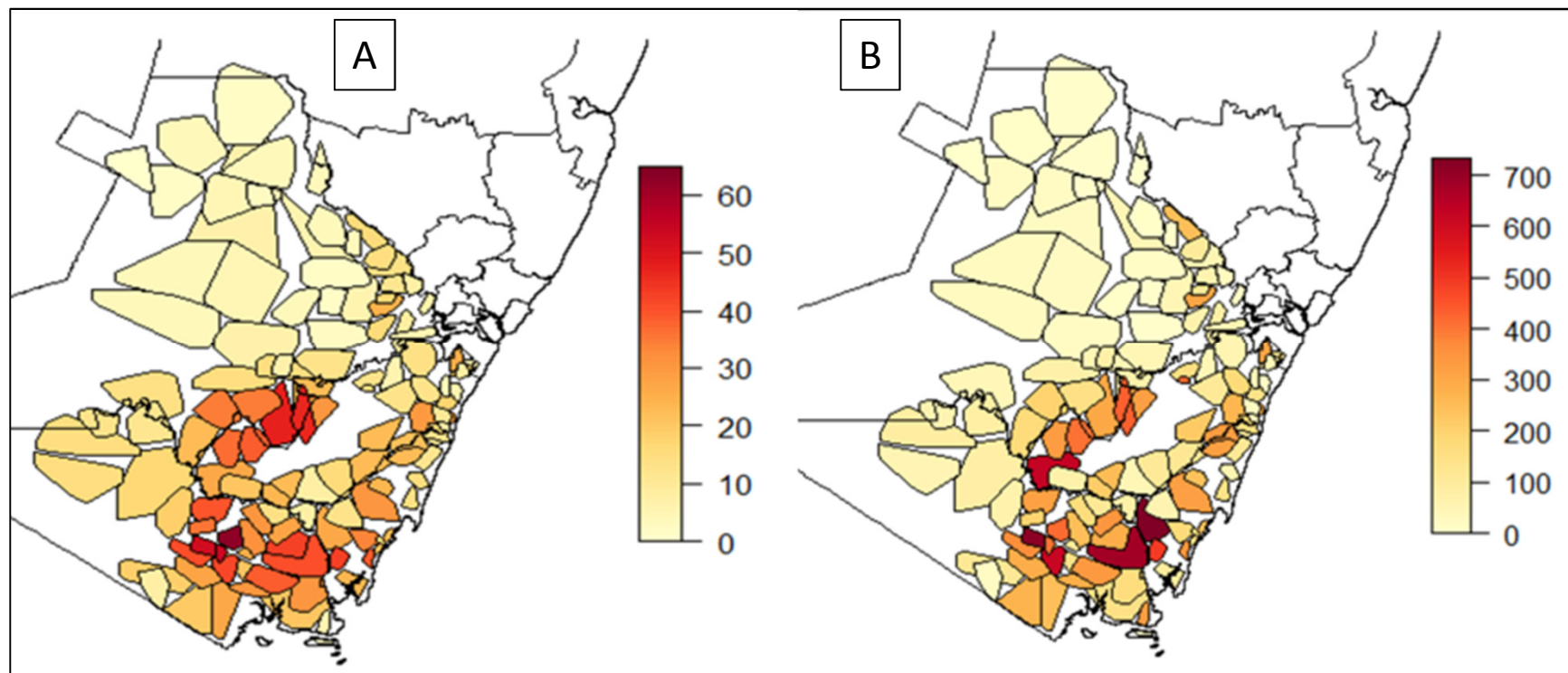


- 2 villages in the Delta region of South **Myanmar**
- Household GPS locations mapped



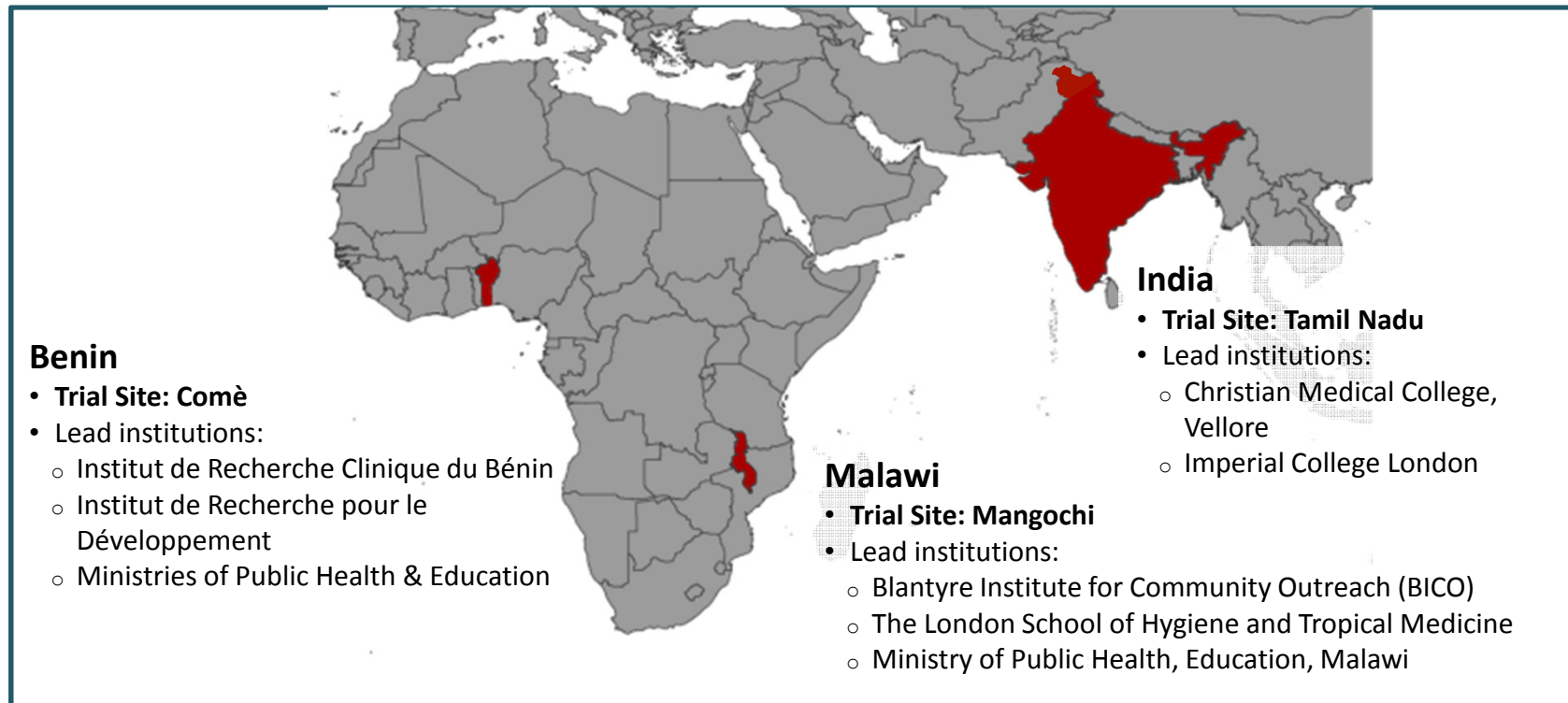
Study site 4 - the Tumikia project ***in South East Kenya (Pullen et al 2017)***

The cluster level prevalence (A) and mean intensity of hookworm infection, eggs per gram (B). **20,842 individuals aged 1 to 99 years**



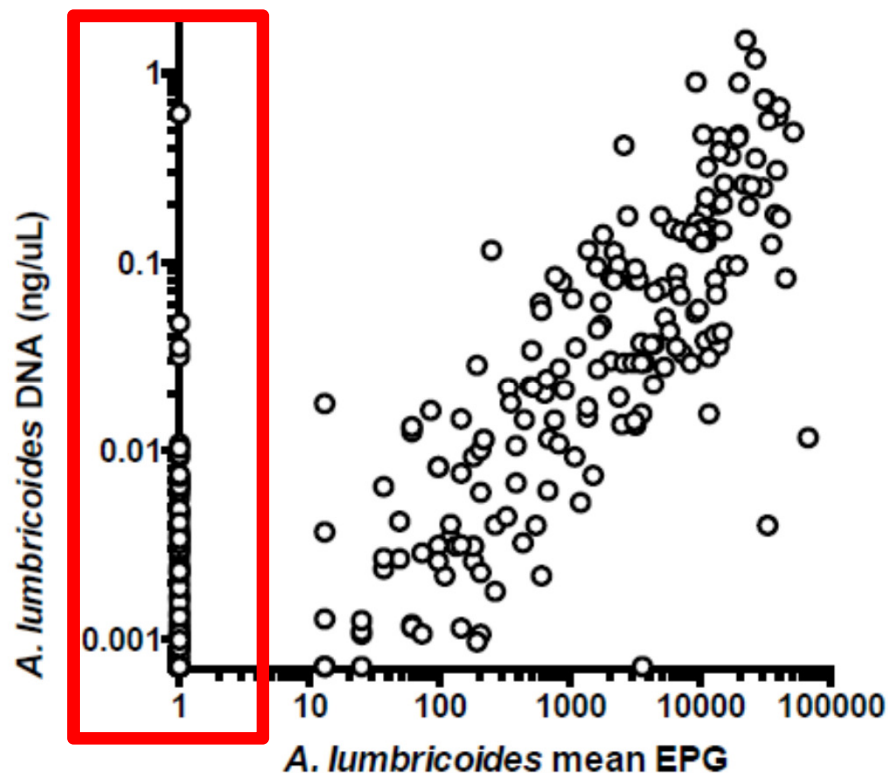


Study sites 5, 6 & 7 - randomized cluster
controlled trials of breaking transmission in BGMF project
entitled DeWorm3 with different treatment strategies





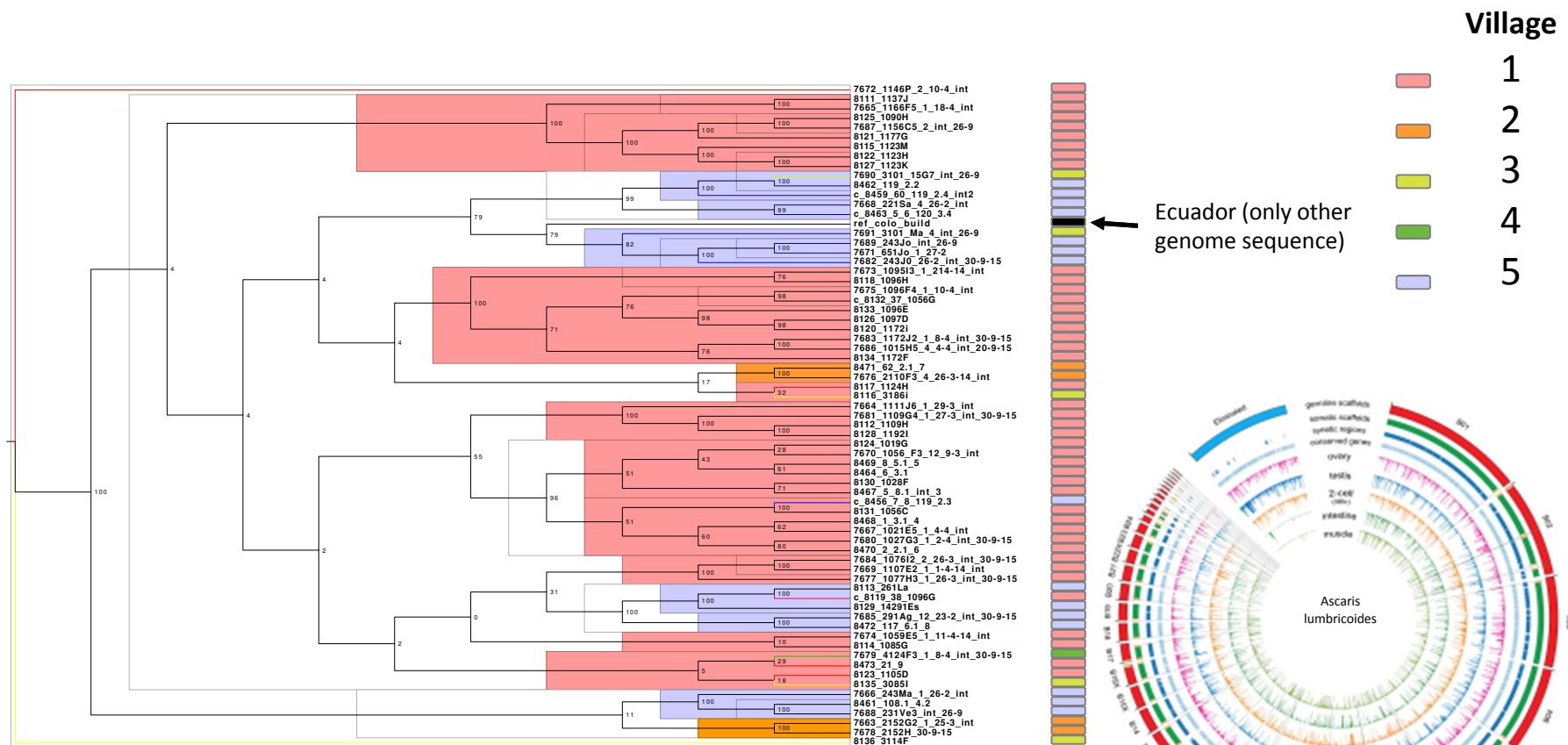
New diagnostic tools - Ascaris - Kato-Katz versus qPCR (Easton et al 2016)



46% more positive samples
when diagnosed with qPCR
compared with Kato-Katz (2
slides)



New genome methods help to determine ‘who infects whom?’



- >99% identity with *A. suum*
 - Differ only by SNPs
 - Same DNA break regions in DNA elimination
- Combined sequencing to generate two genomes
- Improvement on previous *A. lumbricoides* genome: down to 415 scaffolds compared to 31,720

273 Mb – *Ascaris* genome
3234 Mb – human genome



Conclusions

The research of staff at LCNTDR have been central in helping to define policy for the control of NTDs.

- They have defined key epidemiological concepts and been a template for the design of community based randomised treatment clinical trials.
- These trials were designed to test predictions of detailed analyses – so far so good - but a long way to go before they complete in 2021.
- Parameter uncertainty a key issue – need for better quality epidemiological studies.
- In achieving MDA coverage required to break transmission individual compliance to treatment is key – as is its accurate measurement.
- In the ‘End Game’, new tools, such as molecular epidemiological methods, will be key to helping to understand how infection persist at very low prevalence – may dictate a policy change towards targeted treatment.