

# Pooled Testing for TB – Evidence Gaps and Future Priorities

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# Start4All

# Current WHO Recommendations



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- 2026 WHO recommendations are conditional
- Proposes pools from 2-4 samples
- Only includes data from Xpert testing
- Adults and adolescents included
- Doesn't include PLHIV (limited evidence), children (insufficient evidence), or individuals at increased risk of drug resistance

'When resources are limited'

Department for HIV, Tuberculosis,  
Hepatitis and Sexually Transmitted  
Infections

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## Near point-of-care tests, tongue swabs, and sputum pooling for TB

▼ [Near point-of-care tests, tongue swabs, and sputum pooling for TB](#)

[NPOC-NAATs](#)

[Tongue swabs on LC-aNAATs](#)

[Sputum pooling](#)

# Future Work Needed



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- Pooling is disease and system agnostic – use with other platforms (Pluslife)
- Already underway as part of TB REACH-supported work in Cameroon (groups of three) and will be included in Start4All Phase 2
- Early results look promising, dilution with swab tests may be less of an issue
- Some work with Truenat pooled testing as well underway



# Future Work Needed



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- Pools of more than 4: in practice and in a few studies, this is being done
  - At low positivity rates, could pools of 5, 6 or have sufficient sensitivity?
- Pooling in different populations
  - Should not make a difference for specific populations – but rather for bacterial load in those populations.



# Future Work Needed



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- Pooled testing efficiency depends on positivity rates
- AI for CXR provides an abnormality score that increases conveys a likelihood of TB.
- Could using AI help decision making for deciding if people need individual or pooled testing?
- Initial modelling is promising and will be tested prospectively in Start4All Phase 2



RESEARCH

Open Access

Expanding molecular diagnostic coverage for tuberculosis by combining computer-aided chest radiography and sputum specimen pooling: a modeling study from four high-burden countries



Andrew James Codlin<sup>1,2†</sup>, Luan Nguyen Quang Vo<sup>1,2†</sup>, Tushar Garg<sup>3</sup>, Sayera Banu<sup>4</sup>, Shahriar Ahmed<sup>4</sup>,  
...  
1,2,3,4,5,6,7,8,9,10,11,12,13,14,15,16,17,18,19,20,21,22,23,24,25,26,27,28,29,30,31,32,33,34,35,36,37,38,39,40,41,42,43,44,45,46,47,48,49,50,51,52,53,54,55,56,57,58,59,60,61,62,63,64,65,66,67,68,69,70,71,72,73,74,75,76,77,78,79,80,81,82,83,84,85,86,87,88,89,90,91,92,93,94,95,96,97,98,99,100

# Future Work Needed



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- Everyone who has a positive test for TB should get an initial rapid DST (Xpert/Truenat)
  - In many settings RIF resistance is very low.
  - For people with positive Pluslife results, is pooled testing for RIF resistance feasible?
- Math is clear – how does this work in practice?
  - Documenting gains in access to molecular tests the real world, how transport systems are employed, what changes are needed to drive larger numbers of people with presumptive TB?
  - How to integrate with NPOC tests?



# Future Work Needed



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- Documenting pooling processes in lab
  - Who makes decisions, what resources are needed, how to lab techs work with pooled testing?
- Both NPOC and pooled testing saw drop-offs in performance with very low and trace Ultra semiquantitative results
- What does this mean for different screening populations?
- For ACF? Primary care level? How do yield patient characteristics matter for bacillary load?



# The Start4All Partnership



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For more information, see our online Acknowledgements page:  
[www.lstmed.ac.uk/start4all/acknowledgements](http://www.lstmed.ac.uk/start4all/acknowledgements)

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