Biomarker-guided tuberculosis preventive therapy (CORTIS): a randomised controlled trial

Thomas J Scriba*, Andrew Fiore-Gartland*, Adam Penn-Nicholson, Humphrey Mulenga, Stanley Kimbung Mbandi, Bhavesh Borate, Simon C Mendelsohn, Katie Hadley, Chris Hikuam, Masooda Kaskar, Munyaradzi Musvosvi, Nicole Bilek, Steven Self, Tom Sumner, Richard G White, Mzwandile Erasmus, Lungisa Jaxa, Rodney Raphela, Craig Innes, William Brumskine, Andriëtte Hiemstra, Stephanus T Malherbe, Razia Hassan-Moosa, Michèle Tameris, Gerhard Walzl, Kogieuleum Naidoo, Gavin Churchyard, Mark Hatherill, and the CORTIS-01 Study Team

Summary

Background Targeted preventive therapy for individuals at highest risk of incident tuberculosis might impact the epidemic by interrupting transmission. We tested performance of a transcriptional signature of tuberculosis (RISK11) and efficacy of signature-guided preventive therapy in parallel, using a hybrid three-group study design.

Methods Adult volunteers aged 18–59 years were recruited at five geographically distinct communities in South Africa. Whole blood was sampled for RISK11 by quantitative RT-PCR assay from eligible volunteers without HIV, recent previous tuberculosis (ie, <3 years before screening), or comorbidities at screening. RISK11-positive participants were block randomised (1:2: block size 15) to once-weekly, directly-observed, open-label isoniazid and rifapentine for 12 weeks (ie, RISK11 positive and 3HP positive), or no treatment (ie, RISK11 positive and 3HP negative). A subset of eligible RISK11-negative volunteers were randomly assigned to no treatment (ie, RISK11 negative and 3HP negative). Diagnostic discrimination of prevalent tuberculosis was tested in all participants at baseline. Thereafter, prognostic discrimination of incident tuberculosis was tested in the untreated RISK11-positive versus RISK11-negative groups, and treatment efficacy in the 3HP-treated versus untreated RISK11-positive groups, during active surveillance through 15 months. The primary endpoint was microbiologically confirmed pulmonary tuberculosis. The primary outcome measures were risk ratio [RR] for tuberculosis of RISK11-positive to RISK11-negative participants, and treatment efficacy. This trial is registered with ClinicalTrials.gov, NCT02735590.

Findings 20207 volunteers were screened, and 2923 participants were enrolled, including RISK11-positive participants randomly assigned to 3HP (n=375) or no 3HP (n=764), and 1784 RISK11-negative participants. Cumulative probability of prevalent or incident tuberculosis disease was 0.066 (95% CI 0.049 to 0.084) in RISK11-positive (3HP negative) participants and 0.018 (0.011 to 0.025) in RISK11-negative participants (RR 3.69, 95% CI 2.25–6.05) over 15 months. Tuberculosis prevalence was 47 (4.1%) of 1139 versus 14 (0.78%) of 1984 in RISK11-positive compared with RISK11-negative participants, respectively (diagnostic RR 5.13, 95% CI 2.93 to 9.43). Tuberculosis incidence over 15 months was 2.09 (95% CI 1.97 to 3.19) vs 0.80 (0.30 to 1.30) per 100 person years in RISK11-positive (3HP-negative) participants compared with RISK11-negative participants (cumulative incidence ratio 2.6, 95% CI 1.2 to 5.9). Serious adverse events related to 3HP included one hospitalisation for seizures (unintentional isoniazid overdose) and one death of unknown cause (possibly temporally related). Tuberculosis incidence over 15 months was 1.94 (95% CI 0.35 to 3.50) versus 2.09 (95% CI 0.97 to 3.19) per 100 person-years in 3HP-treated RISK11-positive participants compared with untreated RISK11-positive participants (efficacy 7-0%, 95% CI -145 to 65).

Interpretation The RISK11 signature discriminated between individuals with prevalent tuberculosis, or progression to incident tuberculosis, and individuals who remained healthy, but provision of 3HP to signature-positive individuals after exclusion of baseline disease did not reduce progression to tuberculosis over 15 months.

Funding Bill and Melinda Gates Foundation, South African Medical Research Council.

Copyright © 2021 The Author(s). Published by Elsevier Ltd. This is an Open Access article under the CC BY 4.0 license.
as a screening (triage) test to identify undiagnosed tuberculosis disease would allow earlier curative treatment. WHO and FIND have developed target product profiles (TPP) for triage tests for tuberculosis (optimal and minimum sensitivity of >95% and >90%, and specificity of >80% and >70%, respectively), and incipient tuberculosis tests (minimum sensitivity and specificity of 75% and 75%, and optimal sensitivity and specificity of 90% and 90%, respectively).1

We previously developed a 16-gene transcriptomic signature by whole blood RNA sequencing for identification of individuals at high risk of developing tuberculosis (the Zak16 signature).2 Measurement of Zak16 was adapted to quantitative RT-PCR (RT-qPCR), and predictive ability for incident tuberculosis was validated in an independent longitudinal cohort of household contacts of tuberculosis patients.3 We reduced this signature to 11 genes (RISK11) with equivalent performance,4 to allow testing in 96-well PCR format. Zak16 and RISK11 also did well as non-sputum screening tests for prevalent, active tuberculosis in case-control studies, measured by RNA sequencing, microarray,5 or microfluidic RT-qPCR.6,7 In a 2020 systematic review and patient-level pooled meta-analysis of 17 transcriptomic signatures for prognosis of incident tuberculosis, Zak16 was among eight signatures that achieved a positive predictive value above the WHO TPP benchmark for incident tuberculosis tests.8 Since case-control studies might overestimate performance characteristics, testing in unselected populations is needed.

We report on a randomised controlled trial (CORTIS; NCT02735590), which prospectively measured diagnostic and prognostic performance of RISK11 for triage of prevalent and prediction of incident tuberculosis in South African adults; and, in parallel, estimated efficacy of short-course TPT to avert incident disease in RISK11-positive individuals.

**Methods**

**Study design**

This randomised controlled trial used a hybrid treatment selection, three-group study design to evaluate efficacy of the intervention and, in parallel, performance of the biomarker used to allocate that intervention (figure 1).2 The
coprimary aims were to test, over 15 months, whether RISK11 status differentiates between people with and without cumulative prevalent or incident tuberculosis; and whether preventive therapy (weekly high-dose isoniazid and rifapentine for 12 weeks [3HP]) reduces tuberculosis incidence among RISK11-positive people compared with active surveillance only. RISK11 performance to detect prevalent tuberculosis was evaluated in all groups at baseline. RISK11 performance to predict incident tuberculosis was evaluated in the untreated RISK11-positive and RISK11-negative groups, and treatment efficacy was estimated from the 3HP treated and untreated RISK11-positive groups, after omitting participants with baseline tuberculosis. Efficiency of the hybrid study design was maximised by using the RISK11-positive and 3HP-negative group to evaluate both biomarker performance and treatment efficacy.

Routine implementation of TPT requires that patients are screened to exclude prevalent tuberculosis before TPT is provided to prevent incident disease. Therefore, because risk for prevalent and incident tuberculosis by RISK11 status must be understood before efficacy of preventive therapy against incident tuberculosis in RISK11-positive people can be interpreted, these secondary analyses are presented before the primary analysis of treatment efficacy.

The trial protocol (appendix p 21) was approved by the South African Health Products Regulatory Agency (20160305) by the Institutional Human Ethics Committees of participating sites; and was registered on ClinicalTrials.gov (NCT02735590). All participants provided written informed consent in their language of choice.

**Participants**

Adult volunteers living in tuberculosis-endemic communities in South Africa were recruited at five sites (South African Tuberculosis Vaccine Initiative, Worcester; Immunology Research Group, Stellenbosch University, Ravensmead; Aurum Institute, Klerksdorp and Rustenburg; and Centre for the AIDS Programme of Research in South Africa, Durban). Community-based recruitment was by word-of-mouth, house-to-house visits, and liaison with non-governmental organisations. Recruitment did not target groups at high risk of tuberculosis, such as household contacts. Eligible participants were aged between 18 years and 59 years, HIV-negative, without a history of tuberculosis disease in the last 3 years or preselected comorbidities (appendix p 8).

**Screening**

Venous blood was collected in PAXgene RNA tubes from all potentially eligible people at screening, frozen at −20°C, and shipped weekly to the South African Tuberculosis Vaccine Initiative Immunology Laboratory for RISK11 testing by qRT-PCR assay. Participants with a RISK11 score of at least 60% were classified a priori as RISK11 positive and less than 60% as RISK11 negative. This 60% threshold was the optimal point at which sensitivity and specificity for prognosis of incident tuberculosis were balanced in case-control studies.† Samples with failed reference primer-probe reactions, with marked deviation in internal positive control sample from historical runs, or more than 30% failed interferon-stimulated genes primer-probe reactions (see quality control criteria and analysis script in Bitbucket instance) were classified as indeterminate. Per-participant qualitative results (RISK11 positive and RISK11 negative) were provided to the Triclinium Clinical Development (TCD) Data Centre for participant randomisation (see appendix pp 2–3).

**Randomisation and masking**

Assignment to study group was managed by an unmasked randomisation team from the TCD Data Centre, based on RISK11 status. RISK11-positive volunteers were randomly assigned in blocks to ensure concurrent enrolment. †Groups randomly assigned in blocks to ensure concurrent enrolment.

---

**Figure 1: Study design**

The prevalence of RISK11 positivity was not precisely known in the study population; therefore, the number of individuals to be screened and the randomisation of RISK11-negative participants to enrolment was monitored and adjusted adaptively to ensure concurrent enrolment of the target number of RISK11-positive and RISK11-negative participants, per protocol specifications. The study used a three-group design to evaluate efficacy of the intervention and, in parallel, performance of the biomarker used to allocate that intervention. Diagnostic performance for differentiation of prevalent tuberculosis was tested in all three groups at baseline; prognostic performance for differentiation of incident tuberculosis over 15 months was tested in the two untreated groups (untreated RISK11 positive and untreated RISK11 negative), and treatment efficacy of 3HP over 15 months was tested in the two RISK11-positive groups (treated and untreated RISK11 positive). *Participants evaluated for eligibility at screening and enrolment. †Groups randomly assigned in blocks to ensure concurrent enrolment.
assigned (1:2; block size 15) to either open-label 3HP (3HP-positive group), or active tuberculosis surveillance without 3HP (3HP-negative group), in accordance with a randomisation schedule generated using SAS, version 9.4. RISK11-negative volunteers were concurrently randomly assigned either to active tuberculosis surveillance (3HP-negative group) or to non-participation, to enrich the study population for RISK11-positive participants.

Study group allocation was revealed to site staff after enrolment of eligible participants within 28 days of screening. The 3HP-negative group was double-blinded to RISK11 status, but unblinded to treatment allocation; the 3HP-positive group was unblinded to both RISK11-positive status and treatment allocation (figure 1). RISK11-positive participants randomly assigned to active surveillance did not receive a placebo, to maintain blinding of participants and study team members to RISK11 status.

Procedures
Target enrolment was maximally 3200 participants (1500 RISK11 positive and 1700 RISK11 negative). The randomisation ratio for RISK11-negative volunteers was adapted to ensure concurrent enrolment of the recruitment target. Adaptations occurred at intervals, informed by 3-monthly operational monitoring reports based on enrolment rate, RISK11-positive and RISK11-negative prevalence, and tuberculosis case accrual blinded to RISK11 status (appendix p 7).

Enrolment procedures included phlebotomy for IFNγ release assays (QuantiFERON TB Gold-Plus, Qiagen), tuberculosis symptom screen (a positive tuberculosis symptom screen included one or more symptoms of persistent unexplained cough, fever, night sweats, weight loss, or any haemoptysis), and collection of two spontaneous expectorated sputum samples for Xpert MTB/RIF assay (Cepheid) from all sputum-productive participants, regardless of symptoms.

All participants attended up to seven study visits, including four study site visits at months 3, 6, 12, and 15 (end of study), and three telephonic contact or field visits at months 1, 2, and 9. HIV testing was repeated at months 6 and 12.

Participants in the 3HP group received open-label, high-dose isoniazid (15 mg/kg; maximum dose 900 mg) with pyridoxine supplementation (25 mg) and rifapentine based on body weight (>32–50 kg, 750 mg; >50 kg, 900 mg), given weekly as directly observed oral doses, ideally with food, over 12 weeks. Completion of 3HP treatment was defined as receipt of 11 doses within 16 weeks (appendix p 3).

Outcomes
The two primary outcome measures were RISK11-positive to RISK11-negative risk ratio (RR) for prevalent or incident tuberculosis disease, and 3HP treatment efficacy through 15 months. Diagnostic performance for prevalent tuberculosis was evaluated on the presence of tuberculosis at the enrolment visit within the ITT cohort. Participants in the 3HP group had each dose directly observed by study staff, and attended clinic for evaluation of solicited adverse events and possible adverse events of special interest at weeks 1–11. Solicited adverse events included gastrointestinal signs and symptoms suggestive of hepatotoxicity, such as nausea, vomiting, and jaundice. Possible hypersensitivity reactions, including influenza-like illness, were reported as adverse events of special interest. Safety events meeting the definition for a serious adverse event, whether deemed related or unrelated to study drug, were reported for both all participants through end of study.

Statistical analysis
The intention-to-treat (ITT) cohort included all enrolled participants who completed investigation for tuberculosis endpoints at baseline. The modified intention-to-treat (mITT) cohort included all participants in the ITT population who completed at least one post-baseline tuberculosis endpoint investigation and omitted participants with endpoint-defined tuberculosis disease at baseline (prevalent tuberculosis).

The diagnostic RR was estimated as a proportion of participants with tuberculosis disease among RISK11-positive divided by RISK11-negative participants. Prognostic performance for incident tuberculosis after enrolment was evaluated among 3HP-negative participants within the mITT cohort (ie, excluding participants with tuberculosis at baseline). Prognostic RR was estimated as the cumulative incidence through 15 months for RISK11-positive divided by RISK11-negative participants. The primary RR was an estimate of the probability of having prevalent tuberculosis or developing incident tuberculosis among RISK11-positive divided by RISK11-negative participants; this combined probability of prevalent and incident tuberculosis was computed for each group as the probability of prevalent tuberculosis plus the probability of incident tuberculosis through 15 months (conditioned on not having prevalent tuberculosis). Efficacy of 3HP preventive therapy to reduce the rate of incident tuberculosis disease compared with the untreated RISK11-positive participants was also evaluated in the mITT population. Treatment efficacy was estimated as one minus the cumulative incidence of RISK11-positive and 3HP-positive participants divided by RISK11-positive and 3HP-negative participants through 15 months. A per-protocol analysis of treatment efficacy was done and excluded RISK11-positive and 3HP-positive participants that received less than 11 of the 12 weekly doses of 3HP.

For statistical efficiency, a random subset of RISK11-negative participants were enrolled, therefore creating an ITT cohort artificially enriched with RISK11-positive participants. Therefore, unless stated otherwise, all analyses were adjusted so that results reflected the screened population. Secondary performance metrics
Articles

**Performance of RISK11 and IGRA for prevalent and incident tuberculosis**

Data are risk ratio (95% CI), % (95% CI), AUC (95% CI), or NNS or NNT (95% CI). ITT=intention to treat. mITT=modified intention to treat. IGRA=interferon γ release assay. AUC=area under the receiver operating characteristic curve. PPV=positive predictive value. NPV=negative predictive value. NNS=number needed to screen. NNT=number needed to treat. *Computed over 15-month prognostic window. Performance of RISK11 and IGRA for incident tuberculosis over 6-month and 12-month prognostic windows is in the appendix (p 16). †RISK11 score threshold at 60% or 26%. ‡AUC is computed across all score thresholds and characteristic curve. PPV=positive predictive value. NPV=negative predictive value. NNS=number needed to screen. NNT=number needed to treat. *Computed over 15-month prognostic window. Performance of RISK11 and IGRA for prevalent and incident tuberculosis based on secondary endpoint (≥1 sample+) is in the appendix (p 15).

| Race or ethnicity | Total (n=2923) | RISK11 positive and 3HP positive (n=375) | RISK11 positive and 3HP negative (n=764) | RISK11 negative (n=1784) | RISK11 positive vs RISK11 negative, p value* |
|-------------------|---------------|------------------------------------------|-----------------------------------------|--------------------------|-----------------------------------|
| Mixed race        | 968 (33·1%)   | 148 (39·5%)                              | 285 (37·3%)                            | 523 (30·0%)              | <0·001                            |
| Asian             | 4 (0·1%)      | 0                                        | 1 (0·1%)                               | 3 (0·2%)                 | 0·567                             |
| Black African     | 3947 (66·6%)  | 227 (60·5%)                              | 477 (62·4%)                            | 1243 (69·7%)             | <0·001                            |
| White             | 4 (0·1%)      | 0                                        | 1 (0·1%)                               | 3 (0·2%)                 | 0·567                             |
| Body-mass index, kg/m² | 24·6 (7·7)   | 24·1 (6·3)                              | 24·4 (6·1)                             | 24·9 (8·6)               | 0·354                             |
| Smoking           | 1478 (50·6%)  | 203 (54·1%)                              | 391 (51·2%)                            | 884 (49·6%)              | 0·170                             |
| Interferon γ release assay positive | 1895 (64·8%) | 250 (66·7%)                             | 528 (69·1%)                            | 1117 (62·6%)             | <0·001                            |
| Follow-up, months | 15 (9·4–15·0) | 15 (11·1–15·0)                           | 15 (9·1–15·0)                         | 13·6 (9·1–15·0)          | 0·056                             |
| Prevalent tuberculosis, n, (probability, 95% CI)¶ | 51 (1·1%) 0·77–1·6)† | 47 (4·1%, 3·1–5·4)§ | 47 (4·1%, 2·5–4·5)§ | 10 (0·7%, 0·4–7·3) NA | NA                                |
| Incident tuberculosis, n (cases per 100 person years, 95% CI)† | 24 (1·05, 0·59–1·5)§ | 6 (1·5, 0·35–3·5)§ | 14 (2·0, 0·97–3·19) | 10 (0·8, 0·30–1·30) NA | NA                           |
| Cumulative tuberculosis, n (probability, 95% CI)] | 95 (0·022, 0·016–0·028)† | 61 (0·066, 0·049–0·084)¶ | 61 (0·066, 0·049–0·084)‡ | 24 (0·018, 0·011–0·026) NA | NA                                         |

Data are n (%), mean (SD), or median (IQR), unless stated otherwise. Secondary tuberculosis endpoints by group are in the appendix (p 13). NA=not applicable. *For continuous data, p values from Wilcoxon rank sum test. For categorical data, p values from Fischer’s exact test. †Overall rate estimates are weighted combinations of the enrolled participants to reflect the screened population. ‡Overall incidence and cumulative tuberculosis excludes RISK11-positive and 3HP-positive incident cases. §Prevalent tuberculosis among RISK11 positive was assessed by combining the 3HP-negative and 3HP-positive groups. Value is repeated for RISK11 positive and 3HP-positive, and RISK11 positive and 3HP-negative. ¶Probability of observing prevalent or incident tuberculosis over 15 months. ||Probability of prevalent or incident tuberculosis not estimated for RISK11-positive and 3HP-positive because it would combine data from before and after 3HP treatment and is therefore potentially misleading. **Estimate for RISK11 positive includes 3HP-positive prevalent cases and RISK11 positive and 3HP positive, and RISK11 positive and 3HP negative. *Computed over 15-month prognostic window. Performance of RISK11 and IGRA for prevalent and incident tuberculosis based on secondary endpoint (≥1 sample+) is in the appendix (p 15).

| Biomarker prevalence | 9·2% (9·2 to 9·2) | 25·8% (24·1 to 27·4) | 63·4% (61·3 to 65·4) | 9·0% (9·0 to 9·0) | 25·3% (23·7 to 26·9) | 63·2% (61·1 to 65·3) | 0·170 |
| Sensitivity          | 34·9% (23·7 to 52·2) | 72·1% (54·5 to 90·2) | 88·7% (77·1 to 96·4) | 25·0% (12·7 to 45·9) | 47·5% (25·9 to 75·0) | 83·2% (61·9 to 100·0) | 0·567 |
| Specificity          | 91·0% (90·9 to 91·1) | 74·3% (73·1 to 76·4) | 36·9% (34·9 to 39·0) | 91·4% (91·0 to 91·2) | 74·9% (73·7 to 75·5) | 37·0% (34·9 to 39·1) | 0·567 |
| PPV§               | 4·1% (3·0 to 5·4) | 3·1% (2·0 to 4·3) | 1·5% (1·0 to 2·2) | 1·9% (0·9 to 3·0) | 1·3% (0·6 to 2·1) | 0·9% (0·5 to 1·4) | 0·567 |
| NPS or NNT|| | 29·9 (21·8 to 46·6) | 37·8 (25·5 to 65·7) | 83·1 (53·2 to 179·8) | 75·1 (40·4 to 277·5) | 123·8 (47·2 to 834·1) | 168· (44·0 to 1059) | 0·567 |

such as sensitivity and specificity were estimated using standard formulas with binary endpoints; a percentile bootstrap with 20000 samples was used to estimate 95% CIs. For descriptive analyses (tables 1, 2), rank-based Wilcoxon rank-sum tests were used to compare continuous variables in RISK11-positive versus RISK11-negative
The study was designed to have 90% power to reject the null hypothesis of a RISK11-positive and RISK11-negative cumulative risk ratio less than 2 with one-sided alpha of 0.025. For treatment efficacy there was 80% power to reject the null hypothesis of efficacy less than 20%, with one-sided alpha of 0.05 and under the alternative design hypothesis that efficacy was 80%. To compute statistical power for these aims, a stochastic simulation of the trial was constructed, based on which we expected to observe 33 tuberculosis disease endpoints among 1500 RISK11-positive participants and seven tuberculosis disease endpoints among 1700 RISK11-negative participants (appendix p 6).

Role of the funding source
The trial was funded by the Bill & Melinda Gates Foundation (OPP1116632, OPP1137034) and the Strategic Health Innovation Partnerships Unit of the South African Medical Research Council, with funds received from the South African Department of Science and Technology. The Gates Foundation contributed to the study design. The regulatory sponsor was the University of Cape Town. Rifapentine (PRIFTIN) was donated by the manufacturer (Sanofi), who had no role in the design, implementation, analysis, or reporting of the trial. All authors had access to all the data reported in the study. The corresponding author had final responsibility for the decision to submit for publication.

Results
Between Sept 20, 2016, and Oct 19, 2018, 20207 volunteers consented to participation; 16248 met inclusion criteria at screening and 15777 with a RISK11 result were potentially eligible for enrolment (figure 2). Of the 20207 adults assessed for eligibility, common reasons for exclusion included HIV infection (1246 [6.1%]) and comorbid conditions (1369 [6.8%]; appendix p 8).

2923 eligible participants were enrolled after randomisation (1784 [61.0%] RISK11 negative and 1139 [39.0%] RISK11 positive; table 1). Participants were enrolled at five geographically diverse sites across South Africa (appendix pp 9–11). Among participants with a RISK11 result, 1434 (9.3%) of 15494 were RISK11 positive, with the proportion ranging from 6.2% to 13.0% across the five sites. RISK11-positive participants were randomly assigned either to receive treatment (375 [32.9%] RISK11 positive and 3HP positive) or undergo observation without treatment (764 [67.1%] RISK11 positive and 3HP negative).

There were no significant differences in smoking history, family history of tuberculosis, or febrile illness between RISKII-positive and RISKII-negative participants (table 1). A higher proportion of RISKII-positive (75 [9.8%] of 764) than RISKII-negative (18 [6.7%] of 1784) participants reported previous tuberculosis
disease (p=0·003). Compared with RISK11 negative participants, a greater proportion of RISK11-positive participants were female and mixed race (p<0·001; table 1).

Median duration of follow-up for incident tuberculosis was 13·9 months (IQR 9·0–15·0) and 1879 (66%) of 2500 3HP-negative mITT participants attended at least six scheduled visits. 1416 (49%) of 2838 participants were followed up for 15 months and 2160 (75%) of 2838 participants were followed up for at least 9 months. 585 (21%) of 2838 participants did not complete the study because of withdrawal, death, or loss to follow-up (figure 2; table 1).

Among 91 participants with tuberculosis who reached the primary endpoint, 61 were diagnosed with prevalent tuberculosis at baseline (47 RISK11 positive and 14 RISK11 negative; table 1; appendix p 13). Prevalence of tuberculosis in RISK11-positive and RISK11-negative participants was 4·1% (95% CI 3·1–5·4) and 0·78% (0·5–1·3), respectively (figure 3). Thereafter, 24 participants in the untreated group were diagnosed with incident tuberculosis disease (14 RISK11 positive and ten RISK11 negative; table 1; appendix p 13) with overall incidence of 1·05 cases (95% CI 0·59–1·5) per 100 person-years.

Figure 3: RISK11 detection of combined prevalent and incident tuberculosis and diagnostic performance
(A) Prevalence of tuberculosis in RISK11-positive (47 cases,) and RISK11-negative (14 cases, red bar) participants at trial enrolment. Error bars depict 95% CI. Cumulative incidence probability of tuberculosis in RISK11-positive (14 cases, blue line) or RISK11-negative (10 cases, red line) mITT participants during follow-up. Shaded areas represent 95% CI. (B) Ratio of RISK11-positive versus RISK11-negative cumulative incidence probability of observing prevalent or incident tuberculosis disease, in the ITT population of the observation group. (C) RISK11 signature scores (each dot represents a participant) measured at screening in trial participants, stratified on tuberculosis diagnosis. Boxes depict IQR, midline represents the median, and whiskers indicate range among enrolled participants. (D) RISK11 signature scores measured at screening in prevalent tuberculosis cases with or without any tuberculosis symptoms, in incident tuberculosis cases or those who did not have a tuberculosis diagnosis. The enrolled population, not the screened population, is represented in (C) and (D), because a large fraction of RISK11-negative participants were not enrolled by design. (E) ROC curves depicting RISK11 diagnostic performance for prevalent tuberculosis in the ITT population, for prevalent tuberculosis among individuals with no symptoms of tuberculosis (asymptomatic) and among individuals with at least one symptom consistent with tuberculosis disease (symptomatic). Shaded areas represent the 95% CI. The grey and black dots indicate the minimum and optimal criteria, respectively, set out in the WHO target product profile for a triage test. The empty dot indicates the criteria set out in the WHO target product profile for a confirmatory diagnostic test. TRP=true positive rate. ITT=intention to treat. FPR=false positive rate. AUC=area under the receiver operating characteristic curve.
In the primary analysis of biomarker performance for cumulative tuberculosis, cumulative probability of observing prevalent or incident tuberculosis disease in the ITT population was 0·066 (95% CI 0·049–0·084) in RISK11-positive participants and 0·018 (0·011–0·025) in RISK11-negative participants, with a ratio of 3·69 (95% CI 2·25–6·05) over 15 months (figure 3).

A wide range of RISK11 scores was observed, irrespective of tuberculosis outcome (figure 3). Among enrolled participants, those who remained tuberculosis free (controls) had significantly lower RISK11 scores (24·2%, IQR 8·2–75·3) than those with prevalent or incident tuberculosis disease (76·7%, 36·6–94·4; Wilcoxon rank-sum test p<0·0001; figure 3).

In the secondary analysis of biomarker performance for prevalent tuberculosis, using the prespecified RISK11 test threshold (60%) there was 5·13-times (95% CI 3·01–10·69) increased risk of prevalent tuberculosis disease at baseline in RISK11-positive versus RISK11-negative participants, with sensitivity of 35% (95% CI 24–52) and specificity of 91% (95% CI 91–91; table 2). The receiver operating characteristic (ROC) analysis (figure 3) showed that a RISK11 threshold of 26% provided sensitivity of 72% (95% CI 54–90) and specificity of 75% (95% CI 73–76; table 2); with area under the diagnostic ROC curve (AUC) of 0·77 (95% CI 0·68–0·86). These performance estimates did not meet the minimum criteria for a tuberculosis triage test (table 2).

In the secondary analysis of biomarker performance for prevalent tuberculosis, using the prespecified RISK11 test threshold (60%) there was 5·13-times (95% CI 3·01–10·69) increased risk of prevalent tuberculosis disease at baseline in RISK11-positive versus RISK11-negative participants, with sensitivity of 35% (95% CI 24–52) and specificity of 91% (95% CI 91–91; table 2). The receiver operating characteristic (ROC) analysis (figure 3) showed that a RISK11 threshold of 26% provided sensitivity of 72% (95% CI 54–90) and specificity of 75% (95% CI 73–76; table 2); with area under the diagnostic ROC curve (AUC) of 0·77 (95% CI 0·68–0·86). These performance estimates did not meet the minimum criteria for a tuberculosis triage test (table 2).

In the primary analysis of biomarker performance for cumulative tuberculosis, cumulative probability of observing prevalent or incident tuberculosis disease in the ITT population was 0·066 (95% CI 0·049–0·084) in RISK11-positive participants and 0·018 (0·011–0·025) in RISK11-negative participants, with a ratio of 3·69 (95% CI 2·25–6·05) over 15 months (figure 3).
between asymptomatic controls and asymptomatic prevalent tuberculosis cases with an AUC of 0·75 (95% CI 0·66–0·84; figure 3).

In the secondary analysis of biomarker performance for incident tuberculosis, annualised incidence was 2·1 versus 0·8 per 100 person-years among the RISK11-positive versus RISK11-negative participants, respectively, which was equivalent to a 0·026 (95% CI 0·01–0·04) versus 0·010 (0·004–0·02) cumulative incident probability of developing tuberculosis disease over 15 months, respectively (figure 3). No incident tuberculosis cases were detected in RISK11-negative participants until 8–7 months (figure 3). Tuberculosis incidence through 15 months among RISK11-positive participants was 2·61 (95% CI 1·15–5·94) times higher than RISK11-negative participants (table 2); and the RISK11 signature discriminated between incident tuberculosis cases and controls with AUC of 0·63 (95% CI 0·47–0·80; figure 4). Over this period, at the predefined 60% threshold, RISK11 showed very low sensitivity of 25% (95% CI 12–46) with specificity of 91% (95% CI 91–91). At an exploratory RISK11 threshold of 26%, which provided 75% specificity, sensitivity was 47% (95% CI 26–74). By comparison, the prognostic sensitivity of IFNγ release assay release assay over 15 months was 83% (62–100), but more than 60% of the population was IFNγ release assay positive (specificity 37%, 95% CI 35–40; table 2).

RISK11 prognostic performance was highly dependent on time to disease. Instantaneous RISK11 performance, estimated from 6-month sliding windows, showed prognostic discrimination was high (AUC >0·80) for approximately 9 months, before waning towards 0·58 between months 9 and 15 (figure 4; appendix p 16). Prognostic performance of RISK11 for incident tuberculosis within 6 months (AUC 0·95, 95% CI 0·92–1·0) exceeded the optimal TPP for an incipient tuberculosis test (appendix p 16), and for tuberculosis disease within 12 months (0·80, 0·65–0·94; figure 4) approached the minimum TPP (appendix p 16), but over a 15-month period did not meet minimum criteria for a prognostic tuberculosis test (table 2).

In the primary analysis of treatment efficacy among RISK11-positive participants, tuberculosis incidence in the 3HP-positive and 3HP-negative groups was 1·94 cases per RISK11-positive participants, tuberculosis incidence in the 15-month period did not meet minimum criteria for a approach the minimum TPP (appendix p 16), but over a 6-month horizon. While risk-targeted 3HP did not prevent tuberculosis disease over 15 months, there was some evidence of transient efficacy through 9 months among fully adherent participants.

Community-based recruitment of ambulant volunteers was not focused on individuals with known risk factors for tuberculosis, such as household contact. Individuals with recent previous history of tuberculosis (>3 years before screening) and HIV infection were excluded. Nevertheless, more than 1% of study volunteers had previously undiagnosed, microbiologically confirmed
tuberculosis at enrolment, based on spontaneous expectorated rather than induced sputum samples. More than 80% of these baseline tuberculosis cases did not have any symptom compatible with tuberculosis disease and would not have been detected by a tuberculosis screening strategy that requires symptoms as the entry point to investigation. This finding is consistent with 46–79% prevalence (median 50%) of subclinical tuberculosis reported in prevalence surveys.15-18 It is not known whether subclinical tuberculosis would have progressed to symptomatic disease, spontaneously halted, or even reversed if left untreated.15,18 nor whether subclinical disease directly contributes to M tuberculosis transmission.15 Further research is needed to determine the importance of detection, treatment, and prevention of subclinical disease for global tuberculosis control.

Although very few prevalent tuberculosis cases were symptomatic, RISK11 performance for discrimination of symptomatic prevalent tuberculosis cases from symptomatic controls exceeded the optimal TPP criteria for a triage test, while not meeting the stringent criteria for a confirmatory diagnostic test.5 A 2020 prospective observational study among symptomatic individuals who self-presented to a tuberculosis clinic assessed diagnostic accuracy of 27 transcriptomic signatures for discrimination between prevalent tuberculosis cases and controls.16 The 16-gene Zak signature, from which RISK11 was derived, did not meet the minimum WHO criteria for a triage test, but four of the 27 signatures met these criteria, suggesting that another signature might perform as well as or better than RISK11. RISK11 discrimination of asymptomatic prevalent tuberculosis cases from asymptomatic controls was modest and did not meet the minimal criteria for a triage test. These findings suggest that host blood biomarker development must consider that subclinical tuberculosis might be characterised by more heterogeneous or less pronounced peripheral blood inflammatory responses than symptomatic tuberculosis, or both, which affects RNA signature performance. This finding is consistent with lower inflammatory profiles, observed by blood transcriptomic and proteomic analyses, during the subclinical phases of tuberculosis progression compared with subsequent symptomatic, active disease.18-20 The modest performance characteristics for asymptomatic prevalent tuberculosis would limit applicability of RISK11 for biomarker-targeted, mass community screening in the South African setting, where the majority of baseline tuberculosis cases were subclinical. A host blood biomarker with good sensitivity and specificity for symptomatic tuberculosis might be most useful in countries with low to medium tuberculosis burden, where individuals with compatible symptoms might not otherwise be investigated for tuberculosis. Future diagnostic studies in symptomatic patients seeking health care should also include patients with extrapulmonary and culture-negative tuberculosis, particularly in HIV-infected cohorts, and the full spectrum of differential diagnoses that commonly mimic tuberculosis.

Prognostic performance of RISK11 for risk of progression to incident tuberculosis was poor through 15 months and did not meet the minimum TPP criteria for an incipient tuberculosis test through this prognostic horizon.7 However, good prognostic performance was observed for incident tuberculosis within 12 months of testing and performance exceeded the optimal TPP criteria for tuberculosis occurring within 6 months of testing. The positive predictive value for incident disease (1-9 vs 0-9 for RISK11 compared with IFNγ release assay, respectively) was computed from the observed tuberculosis incidence rate, which was lower than that typically used in estimations (2%).16 It is not possible to determine whether the late incident tuberculosis cases occurring among RISK11-negative participants were due to reactivation or new M tuberculosis infection. We infer that a prognostic test with optimal short-term performance might be useful in identifying people who would benefit from an efficacious intervention in a low-incidence setting, in which the timing of M tuberculosis exposure is often known and subsequent exposure is unlikely.

The 3HP regimen was previously shown to be as effective as 9 months of isoniazid in preventing tuberculosis among individuals with known exposure or positive TST.21 Ethical equipoise of the intervention and control groups in this trial was based on the fact that RISK11 had been validated in selected case-control studies, which can overestimate biomarker performance, and thus risk for tuberculosis among RISK11-positive individuals needed to be tested prospectively in the field. Furthermore, although TPT is given commonly to IFNγ release assay-positive and tuberculin skin test-positive individuals, and individuals with known household exposure to a tuberculosis patient, the vast majority do not progress to tuberculosis disease if left untreated. The risk–benefit balance of 3HP preventive therapy for RISK11-positive individuals was unknown. However, we found no evidence that 3HP treatment reduced the rate of incident tuberculosis over 15 months in RISK11-positive individuals, who might be further advanced along the spectrum of tuberculosis pathogenesis.3 Interpretation of the results is limited by the wide CIs. It is also notable that no tuberculosis cases were observed in fully adherent participants through 9 months. This finding is consistent with the possibility that 3HP was sufficient to temporarily halt, but insufficient to sterilise, incipient tuberculosis, resulting in reactivation. A RISK11-targeted preventive therapy strategy for high tuberculosis transmission settings like South Africa might require a more potent therapeutic regimen than 3HP. 3HP might have been sufficient to sterilise incipient tuberculosis disease, but not to protect against tuberculosis disease resulting from reinfection after completion of the treatment course. Although it is
not possible to distinguish between reactivation and reinfection tuberculosis cases in this study, we suggest that the limited time available for reinfection and subsequent progression to disease after completing 3HP makes the latter possibility less likely.

The study was subject to a number of limitations. The hybrid design required an open-label treatment group,² with no placebo for RISK11-positive participants, so that risk for tuberculosis could also be evaluated in the control group while concealing RISK11 status from participants, site staff, and tuberculosis endpoint laboratory staff for analysis of biomarker performance. Tuberculosis case accrual during the latter stages of follow-up suggested no evidence of ascertainment bias during treatment. Extensive study simulations were done to inform the size of each group for the coprimary analyses. However, due to lower than expected prevalence of RISK11-positive status in the screened population (9.3% vs 15%), enrolment of 1139 RISK11-positive individuals required 2 years, compared with planned enrolment of 1500 RISK11-positive individuals over 1 year. As a result, fewer than expected incident tuberculosis outcomes were observed (24 vs 40 primary endpoints), which might have reduced power for both treatment efficacy and RISK11 performance analyses. Not all participants in the treatment group while concealing RISK11 status from participants, overall and in 4.9% due to an adverse event.²¹ The loss to follow-up in this trial (14.4%) was higher than expected and might reflect the challenges of retaining the study participants, in the absence of traditional tuberculosis risk factors for which surveillance is routine. However, loss to follow-up occurred predominantly after the treatment period and did not seem to have been biased by treatment factors. We note that median duration of participation was 13.9 months and thus loss to follow-up did not have major unforeseen effect on statistical power. Strengths of the study include enrolment in geographically distinct sites across South Africa that were representative of populations with different rates of IFNγ release assay positivity and tuberculosis disease, which were congruent with local rates of RISK11 positivity, although the sample size was not sufficient for analysis of signature performance or 3HP efficacy at site level. The findings broadly reflect the South African tuberculosis epidemic, but might not be directly applicable to countries with much lower tuberculosis incidence.

It is not yet known whether other parsimonious tuberculosis signatures, developed and validated like RISK11 in carefully curated case-control studies, will exhibit similar performance characteristics when prospectively tested in the field, where undiagnosed subclinical tuberculosis might pose a challenge to diagnostic performance. Head-to-head analyses of several transcriptomic tuberculosis signatures with promising diagnostic performance are currently underway on CORTIS samples and novel near-patient testing platforms are in development, which might bring cost-effective, community-based tuberculosis biomarker screening closer to implementation in the field. However, although we have shown that a strategy of biomarker-guided tuberculosis preventive therapy is feasible, the optimal preventive therapy regimen for use in such a strategy remains elusive.

**Contributors**

TJS, AF-G, AP-N, SS, GW, KN, GC, and MH designed the study. CI, WB, AH, STM, RH-M, MT, GW, KN, and GC recruited and evaluated participants. TJS, AF-N, SKM, SCM, KH, MM, NB, ME, LJ, and RR collected and analysed laboratory data. AF-G, and BB accessed and verified the data. TJS, AF-G, AP-N, HM, BB, TS, RGW, and MH analysed data and interpreted results. KH, CH, and MK provided operational support. TJS, AF-G, HM, and MH drafted the Article. All authors had full access to the data, and reviewed, revised and gave final approval of the Article before submission.

**The CORTIS-01 Study Team**

Kesenogile Baepanye, Thobiso Baepanye, Ken Clarke, Marelize Collignon, Audrey Dlamini, Candice Eyre, Tebogo Feni, Moogo Fikizolo, Phinda Galane, Thelma Goliath, Ali Gagat, Shirley Malefo-Grootboom, Elia Janse van Rensburg, Bonita Janse van Rensburg, Sophy Kekana, Marietjie Zietsman, Adrianne Kock, Israel Kiemene, Anesca Lakhi, Nonhumbuso Langa, Hilda Ledwaba, Marillyn Luphoko, Ines Macalewa, Dorah Mabe, Nkosinathi Mahuza, Molly Majola, Mantai Makhetha, Mpho Makoayane, Blossom Makhubalo, Vernon Malay, Juana Market, Selv Mathiego, Nontsikelelo Mhpa, Tsiamo Mnontsa, Sylvester Modipa, Samuel Mopati, Palesa Mowewgo, Primrose Motlaga, Dorothy Muller, Grace Nchwe, Maryna Nel, Lindwe Nhlangulela, Bantu Buhlobo, Ntiamo Ntiamo, Lawrence Ntshabu, Nomusa Sanyaka, Letlhogonolo Seabela, Pearl Selepe, Melissa Senne, MG Serake, Maria Thlapi, Vincent Tshikovhi, Lebogang Tswaile, Amanda van Aswegen, Lungile Mbata, Constance Takavamanya, Pedro Pinho. Jindal Mdulu, Marthinette Taljazed, Nadyene Slabbert, Sharfuddin Sayed, Tanya Nielson, Melissa Senne, Ni Ni Sein, Lungile Mbata (The Aurum Institute); Dhineshree Govender, Tilagavathy Chinappa, Mbilangi Ignatia Zulu, Nonhle Bridgette Maphanga, Senzo Ralph Hlati, Goodness Khanyisile Gumede, Thandwe Yonne Shezi, Jahlulwile Lethabo Maphanga, Zandile Patrick Jai, Thololani Cwele, Nonhlanhla Zanele Elsie Gwamanda, Celaphiwe Dlamini, Zibuyile Phindile Penle Sing, Ntombonzuko Gloria Ntanjana, Sphelele Simo Nzimande, Siyabonga Mthatha, Bhavna Maharaj, Atika Moosa, Cara-Mia Corris (Centre for the AIDS Programme of Research in South Africa); Fazlin Kafar, Marwou De Kock, Hennie Geldenhuys, Angelique Kany Kany Luabeey, Justin Shenje, Natalja Botes, Susan Rosouw, Hadin Africa, Borgani Samdent, Samdonta Braf, Sonia Struers, Alida Carstens, Ruwiyda Jansen, Simbharashesi Malwe, Runaze Herling, Ashley Veldsman, Lebohang Makhele, Marcia Steyn, Sivuyile Buhlungu, Margareth Erasmus, Ilse Davids, Patsiswa Plaatjie, Alessandro Companie, Frances Ratangere, Helen Veldsman, Christel Petersen, Charmaine Abrahams, Miriam Moses, Xoliswa Kelepu, Yolande Gregg, Leticia Swanepoel, Nomitho Magasa, Non punelo Cetrywao, Lauren Maxtavie, Handbooka Valley, Elizabeth Filander, Nambitha Ngxaka, Angelique Mouton, Fajwa Opperman, Elna Van Rooyen, Petrus Tyambetyu (South African Tuberculosis Disease Division, Fred Hutchinson Cancer Research Center)
Declaration of interests

TJS has a patent of the RISK11 signature pending and reports grants from the Bill and Melinda Gates Foundation and South African Medical Research Council. KN reports grants from CAPRISA. GW reports a patent (PCT/IB2019/052043) pending to University of Cape Town, Stellenbosch University, Max-Planck-Gesellschaft Zur Foerderung Der Wissenschaften EV. Seattle Children’s Hospital Doing Business as Seattle Children’s Research Institute, and UK Research and Innovation. GC reports grants from the Bill & Melinda Gates Foundation. AP-N has a patent of the RISK11 signature pending and reports grants from the Bill & Melinda Gates Foundation and the South African Medical Research Council. MH reports grants from the Bill & Melinda Gates Foundation and the South African Medical Research Council.

Data sharing

Deidentified RISK11 signature scores and primary tuberculosis endpoint data from all participants will be available with publication. The dataset is deposited in Zivahub (https://doi.org/10.25375/uct.13573337v1), an open access data repository hosted by the University of Cape Town’s institutional data repository powered by Figshare for Institutions.

Acknowledgments

The trial was funded by the Bill & Melinda Gates Foundation and the South African Medical Research Council. KN reports grants from CAPRISA. GW reports a patent of the RISK11 signature pending and reports grants from the Bill and Melinda Gates Foundation and South African Department of Science and Technology. Rifapentine was donated by the manufacturer (Sanofi).

References

1 Behr MA, Edelstein PH, Ramakrishnan L. Is Mycobacterium tuberculosis infection life long? BMJ 2019; 367:15770.
2 Fiore-Gartland A, Carpp LN, Naikou K, et al. Considerations for biomarker-targeted intervention strategies for tuberculosis disease prevention. Tuberculosis 2018; 109: 61–68.
3 Petrucchioli E, Scriba TJ, Petrone L, et al. Correlates of tuberculosis risk: predictive biomarkers for progression to active tuberculosis. Eur Respir J 2016; 48: 1751–63.
4 Sumner T, Scriba TJ, Penn-Nicholson A, Hatherill M, White RG. Potential population level impact on tuberculosis incidence of using an mRNA expression signature correlate-of-risk test to target tuberculosis preventive therapy. Sci Rep 2019; 9: 11126–10.
5 WHO. High-priority target product profiles for new tuberculosis diagnostics: report of a consensus meeting. Geneva: World Health Organization, 2014. https://apps.who.int/iris/bitstream/handle/10665/135617/WHO-HTM_TB_2014_18_eng.pdf [accessed Jan 7, 2021].
6 WHO. FIND. Development of a Target Product Profile (TPP) and a framework for evaluation for a test for predicting progression from tuberculosis infection to active disease. Geneva: World Health Organization, 2017. https://apps.who.int/iris/bitstream/handle/10665/259176/WHO-HTM-TB-2017.18-eng.pdf [accessed Jan 7, 2021].
7 Penn-Nicholson A, Scriba TJ, Hatherill M, White RG, Sumner T. A novel blood test for tuberculosis prevention and treatment. S Afr Med J 2016; 107: 4–5.
8 Zak DE, Penn-Nicholson A, Scriba TJ, et al. A blood RNA signature for tuberculosis disease risk: a prospective cohort study. Lancet 2016; 387: 2312–22.
9 Darabe F, Mhando SK, Naidoo K, et al. Detection of tuberculosis recurrence, diagnosis and treatment response by a blood transcriptomic risk signature in HIVinfected persons on antiretroviral therapy. Front Microbiol 2019; 10: 1441.
10 Darabe F, Mhando SK, Thompson EG, et al. Diagnostic performance of an optimized transcriptomic signature of risk of tuberculosis in cryopreserved peripheral blood mononuclear cells. Tuberculosis 2018; 108: 124–26.
11 Gupta RK, Turner CT, Venturini C, et al. Concise whole blood transcriptional signatures for incipient tuberculosis: a systematic review and patient-level pooled meta-analysis. Lancet Respir Med 2020; 8: 395–406.
12 Onozaki I, Law I, Sismanidis C, Zignol M, Glazion P, Floyd K. National tuberculosis prevalence surveys in Asia, 1990–2012: an overview of results and lessons learned. Trop Med Int Health 2015; 20: 1126–45.
13 Gunasekara K, Cohen T, Gao W, Ayles H, Godfrey-Faussett P, Claassens M. Smoking and HIV associated with subclinical tuberculosis: analysis of a population-based prevalence survey. Int J Tuberc Lung Dis 2020; 24: 340–46.
14 Frascella B, Richards AS, Sossen B, et al. Subclinical tuberculosis disease—a review and analysis of prevalence surveys to inform definitions, burden, associations and screening methodology. Clin Infect Dis 2020; published online Sept 16. https://doi.org/10.1093/cid/ciaa1402.
15 Drain PK, Bajena KL, Dowdy D, et al. Incipient and subclinical tuberculosis: a clinical review of early stages and progression of infection. Clin Microbiol Rev 2018; 31: e00218–18.
16 Tiemersma EW, van der Werf MJ, Borgdorff MW, Williams BG, Nagelkerke NJ. Natural history of tuberculosis: duration and fatality of untreated pulmonary tuberculosis in HIV negative patients: a systematic review. PLoS One 2011; 6:e17601.
17 Dowdy DW, Basu S, Andrews JR. Is passive diagnosis enough? The impact of subclinical disease on diagnostic strategies for tuberculosis. Am J Respir Crit Care Med 2013; 187: 541–51.
18 Turner CT, Gupta RK, Tsaliik L, et al. Blood transcriptional biomarkers for active pulmonary tuberculosis in a high-burden setting: a prospective, observational, diagnostic accuracy study. Lancet Respir Med 2020; 8: 407–19.
19 Scriba TJ, Penn-Nicholson A, Shankar S, et al. Sequential inflammatory processes define human progression from M. tuberculosis infection to tuberculosis disease. PLoS Pathog 2017; 13:e1006687.
20 Moreira-Texeira I, Tâbhone O, Graham CM, et al. Mouse transcriptome reveals potential signatures of protection and pathogenesis in human tuberculosis. Nat Immunol 2020; 21: 464–76.
21 Sterling TR, Villarino ME, Borisov AS, et al. Three months of rifapentine and isoniazid for latent tuberculosis infection. N Engl J Med 2011; 365: 2355–66.

www.thelancet.com/infection  Published online January 25, 2021  https://doi.org/10.1016/S1473-3099(20)30914-2