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The impact, effectiveness and outcomes of targeted screening thresholds for programmatic latent tuberculosis infection testing in HIV

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Abstract

Background—Screening and treatment for latent tuberculosis infection (LTBI) are key for TB control. In the UK, the National Institute for Health and Care Excellence (NICE) and the British HIV Association (BHIVA) give conflicting guidance on which groups of people with HIV (PWH) should be screened, and previous national analysis demonstrated heterogeneity in how guidance is applied. There is an urgent need for a firmer clinical effectiveness evidence base on which to build screening policy.

Methods—We conducted a systematic, programmatic LTBI-screening intervention for all PWH receiving care in Leicester, UK. We compared yields (percentage IGRA positive) and number of tests required when applying the NICE and BHIVA testing strategies, as well as strategies targeting screening by TB incidence in patients' countries of birth.

Results—Of 1053 PWH tested, 118 were IGRA-positive (11.2%). Positivity was associated with higher TB incidence in country-of-birth [adjusted odds ratio, 50–149 cases compared with <50 cases/100 000: 11.6; 95% confidence interval (CI) 4.79–28.10]. There was high testing uptake

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Conflicts of interest

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(1053/1069, 98.5%). Appropriate chemoprophylaxis was commenced in 100 of 117 (85.5%) patients diagnosed with LTBI, of whom 96 of 100 (96.0%) completed treatment. Delivering targeted testing to PWH from countries with TB incidence greater than 150 per 100000 population or any sub-Saharan African country, would have correctly identified 89.8% of all LTBI cases while cutting tests required by 46.1% compared with NICE guidance, performing as well as BHIVA 2018 guidance.

Conclusion—Targeting screening to higher risk PWH increases yield and reduces the number requiring testing. Our proposed ‘PWH-LTBI streamlined guidance’ offers a simplified approach, with the potential to improve national LTBI-screening implementation.

Keywords

HIV; interferon gamma release assay; latent tuberculosis; screening; testing; tuberculosis

1 Introduction

In 2019, tuberculosis (TB) incidence in England rose for the first time in 9 years, from 4615 in 2018 to 4725 (2.4%) [1]. Incidence remains higher in the UK than most other countries in western and northern Europe [2] and its public health importance remains: until the SARS-CoV-2 pandemic, TB was the leading cause of death from an infectious disease among adults worldwide, with an estimated 10 million people falling ill with TB in 2019, a number that has been declining only very slowly in recent years [3].

There is now increasing focus on latent TB infection (LTBI) screening to reduce TB incidence in high-risk populations. Once latently infected, an individual is at highest risk of developing TB disease within the first 2 years, but can remain at risk for their lifetime [4]. As the global community looks to meet ambitious targets for reduction (90% reduction in TB incidence by 2035) and elimination of TB (<1 incident cases/1 000 000 per year) by 2050 [5], reducing the LTBI reservoir will be essential and is one of the WHO’s key performance indicators [6].

The WHO has published guidelines on groups at high risk to target for LTBI screening and treatment: people with HIV (PWH) are prime amongst these. HIV accelerates progression from LTBI to active TB from around a 10% lifetime risk to as high as 15% per year [7]. Although antiretroviral therapy (ART) reduces TB risk, it does not return to that of the HIV-negative population [8,9]. Screening and treatment (chemoprophylaxis) for LTBI reduce the risk of developing active TB, thereby preventing active TB with its attendant morbidity and mortality, transmission as well as additional costs for the health system [10].

Both 2019 European Centre for Disease Control (ECDC) guidance for use in the European Union and European Economic Area [11] and WHO guidelines for low-TB burden countries [12] recommend that all PWH should be targeted for LTBI screening. In the UK, national guidance is conflicting: the updated 2016 National Institute for Health and Care Excellence (NICE) guidance aligns with international recommendations to test all PWH [13]. By contrast, the British HIV Association (BHIVA) updated 2018 guidance recommends offering interferon gamma release assay (IGRA) testing to all PWH from high (> 150/100 000

population) or medium (40–150/100 000 population) TB incidence countries, and only screening those from low-TB-burden countries (<40/100 000 population) if additional risk factors for TB are present (listed in the guidance [14]).

This contradictory guidance may have contributed to the extreme heterogeneity in how LTBI testing for PWH is applied in the UK. Our national evaluation of practice highlighted that no widespread LTBI programmatic screening has been implemented in the UK in this population, with approximately half of HIV services offering no LTBI testing [15] despite LTBI screening and treatment being highly acceptable to this population [16]. Reasons for this heterogeneity in screening practice remain unclear but it likely represents a lack of confidence in existing guidelines and uncertainty as to which individuals should be offered LTBI screening. With patchy testing coverage, it is unsurprising that there are few previously published data on prospective, programmatic screening in low-TB-burden settings. Those which are available from cohort studies in low-incidence settings, including the UK, have included only a proportion of the active cohort being treated in that centre [17,18], contained estimated data [19], or had small sample sizes [17,20]. This highlights the need for a firmer clinical effectiveness evidence base on which to base national, and potentially international, screening policy.

We aimed to address this evidence gap by implementing a prospective screening programme for LTBI among PWH to understand levels of LTBI-testing uptake, prevalence of LTBI and levels of LTBI chemoprophylaxis uptake and completion for those testing positive, amongst this population. We also explored factors associated with LTBI in PWH, such as ethnicity and TB incidence in country-of-birth, to evaluate the performance of targeted screening strategies including the 2018 BHIVA [14] and NICE [13] guidance and formulate an alternative targeted testing strategy identifying groups of PWH to prioritize for testing, which optimizes testing yield (IGRA positivity rate amongst those tested) and efficiency (minimal IGRA tests required).

Methods

Study design and setting

We implemented a LTBI-screening programme in Leicester, UK, an ethnically diverse city with one of the highest TB incidence rates in the UK (40.5/100 000 general population in 2018 [21]). HIV prevalence is 3.96 of 1000 population aged 15–59 years, making Leicester one of 84 (out of 317) local authorities in England with ‘high-diagnosed prevalence’ (2/1000 population) [22]. Only inconsistent, patchy LTBI screening amongst PWH had been occurring in Leicester since the introduction of IGRA tests.

From 22 February 2014 onwards, we prospectively screened all remaining active HIV patients in Leicester for LTBI followed by treatment, irrespective of ethnicity, country-of-birth, age, sex or comorbidities, to assess acceptability and uptake of LTBI screening and treatment among PWH, IGRA positivity rate, LTBI treatment completion rate and correlates with IGRA positivity.

Study population and participants

We included all PWH who had sought care for HIV at University Hospitals Leicester (UHL) NHS Trust (which is the sole provider for HIV and TB care in Leicester city and Leicestershire) up until 30 June 2017. We excluded those who had had active TB or LTBI treated previously and those who had died, moved away from Leicester, been lost to follow-up or who had been screened for LTBI previously. Results from IGRA screening of the cohort, together with chemoprophylaxis uptake and completion data, were included until 30 June 2021 for the purposes of this analysis.

Ethics

No ethics approval was required as this was considered to be implementation of clinical care in line with national recommendations. Approval was given by the UHL Trust TB Board, the UHL HIV Department and the UHL Microbiology Department.

Screening and management

In our prospective screening study, we used Quanti-FERON-TB Gold In-Tube (QFN-GIT), with gradual switching to QuantiFERON-TB Gold Plus (QFT-Plus) between May 2016 and January 2017. Results were positive, negative or indeterminate, dependent on manufacturers' criteria. Indeterminate results were included in the denominator.

The majority of PWH with CD4⁺ cell counts at least 200 cells/μl received a single IGRA test. Those with CD4⁺ cell counts less than 200 cells/μl generally received two tests (T-SPOT. TB plus a QuantiFERON-TB test), in most cases performed on the same day (where this was not possible, dual results were included if tests were taken within 14 days). PWH taking two tests were classed IGRA positive if either test was positive.

The most recent previous CD4⁺ cell count to an IGRA test was used as the CD4⁺ cell count classification at the time of the test. Individuals who had CD4⁺ cell counts performed more than a year prior to the planned time of IGRA testing had testing withheld until a more recent CD4⁺ cell count became available.

Individuals with positive IGRA tests were recalled for chest radiography and further clinical assessment to exclude active TB [13]. We defined LTBI as PWH with a positive IGRA and normal chest radiography in the absence of any clinical features that would suggest active disease [23]. PWH diagnosed with LTBI were offered chemoprophylaxis, in most cases, 6 months of isoniazid, in accordance with UK guidelines [13], although individual clinicians made the final decision dependent on clinician and patient preference. Where active TB was diagnosed [23], treatment again followed UK guidelines [13].

Data acquisition

Date of birth and sex at birth were recorded, together with NHS number wherever available, to verify records. Ethnicity and country-of-birth were ascertained from electronic hospital, HIV records or paper hospital records, and ethnicity was coded according to the national NHS data dictionary [24]. Countries of birth were further classified into regions according to the World Bank Analytical Grouping [25]. We took TB incidence in country-of-birth (<50,

50–149, 150–249, 250–349 and 350 per 100 000 population) from WHO's Global Health Observatory and used figures available in March 2019 [26].

Statistical analysis

Continuous data were summarized with median and interquartile range (IQR) and categorical responses as proportions/percentages. Comparisons were made using Pearson's chi-square test (or Fisher's exact test, if appropriate). We assessed univariate associations of IGRA positivity with age at IGRA test, CD4⁺ cell count at IGRA test, year of HIV diagnosis, sex, ethnicity, UK birth status, region of birth, TB incidence level in country-of-birth and type of IGRA performed using logistic regression, reported as crude odds ratios (OR) and 95% confidence intervals (95% CIs). Region of birth and black ethnicity categories were collapsed because of small numbers for some regions/groups. Multivariable models were adjusted for age, sex and year of HIV diagnosis (selected *a priori*). On the basis of univariate associations, models were further adjusted for CD4⁺ cell count at IGRA test. However, as ethnicity, UK birth status and region of birth were closely linked to the TB incidence in country-of-birth, only TB incidence in country-of-birth was included in the multivariable logistic regression.

To evaluate the performance of NICE and BHIVA guidelines alongside other targeted testing strategies using different thresholds of TB incidence in country-of-birth, for each screening scenario we calculated number of PWH needing to be screened, LTBI yield and the proportion of all those IGRA-positives correctly identified. TB incidence thresholds evaluated were: more than 350 of 100 000, more than 250 of 100 000, more than 150 of 100 000, more than 50 of 100 000 and no threshold, that is, screening all PWH, the strategy recommended by NICE 2016 guidelines [13], ECDC guidelines for the EU/EEA [11] and WHO guidelines for low-TB-incidence countries [12]. We compared these strategies to 2018 BHIVA guidance, which recommends targeted IGRA testing for those born in low-TB-incidence countries (<40/100 000) for those with TB risk factors including recent exposure to a known TB case, injecting drug use and diabetes mellitus [14]. We, therefore, prospectively collected risk factor data for patients testing IGRA-positive and from a low incidence (<40/100 000) country from 2018 onwards, and retrospectively extracted data from medical notes for those testing IGRA-positive pre-2018. As BHIVA guidance changed part-way through our study, we did not prospectively collect data on risk factors for patients who tested IGRA-negative from low-incidence countries.

Finally, we compared these guidelines to our proposed 'PWH-LTBI streamlined guidance': targeting testing to PWH with country-of-birth TB incidence more than 150 cases per 100 000 population or any sub-Saharan African country. This alternative guidance was formulated to maximize yield while minimizing testing required, while streamlining guidance to be as simple and user-friendly for physicians as possible.

All data were analysed using Stata v15.1 (StataCorp, College Station, Texas, USA). All statistical tests were considered significant when *P* value 0.05 or less.

Results

Cohort description

Recruitment into the study is outlined in Fig. 1. Three hundred and twenty-three patients had already had active TB and were excluded from screening, as were 30 patients who had already been diagnosed with and treated for LTBI. Ten of 1069 patients eligible for screening (0.9%) remained untested at the end of follow-up and six (0.6%) patients declined screening, leaving 1053 who underwent LTBI screening.

Table 1 shows the demographic and HIV and LTBI testing-related characteristics of the screened population ($n = 1053$). Median age at IGRA testing was 42 years (IQR 36–49) and median CD4⁺ cell count was 530 cells/ μ l (IQR 380–700). Only 48 (4.6%) had a CD4⁺ cell count less than 200 cells/ μ l. The dominant ethnic groups were black African (498, 47.3%) and white (388, 36.8%). Sub-Saharan Africa and Europe and Central Asia were the most common regions of birth (51.1 and 40%, respectively).

Interferon gamma release assay-testing outcomes

IGRA results were available for all participants. Overall, 118 (11.2%) PWH had a positive IGRA result (Fig. 1 and Table 1), and two had indeterminate results (0.2%, further information in Table S1, Supplemental Digital Content, <http://links.lww.com/QAD/C621>).

All PWH with a positive IGRA test were diagnosed with LTBI apart from one individual who was found to have active TB disease during clinical/radiological assessment of a positive T-SPOT. TB test, performed at a CD4⁺ cell count of 340 cells/ μ l but with a detectable HIV viral load of 182 copies/ml.

Of the 117 PWH diagnosed with LTBI, 100 (85.5%) commenced LTBI chemoprophylaxis (Fig. 1). Nine of 117 (7.7%) declined; treatment was not advised by the treating physician in two of 117 (1.7%) cases; and four of 117 (3.4%) moved away before chemoprophylaxis could be given. Treatment is pending in two of 117 (1.7%) cases. Reasons behind declination were not well documented in patient notes. Ninety-eight of 100 (98%) of those initiating chemoprophylaxis had isoniazid monotherapy; the remaining two (2%) had combined rifampicin/ isoniazid.

Of the 100 patients commencing chemoprophylaxis, 96 (96%) completed treatment to the satisfaction of the treating physician. One individual moved away and it was unclear whether chemoprophylaxis had been completed, and one defaulted from treatment. Only two of 100 (2%) had to stop treatment prematurely because of adverse drug effects.

Factors associated with interferon gamma release assay-positivity and latent tuberculosis infection

Non-UK born individuals were significantly more likely than UK-born individuals to be IGRA-positive (15.6 versus 2.8%, $P < 0.0001$). The majority of those testing positive were from sub-Saharan Africa (96/118, 81.4%), with the IGRA positivity rate for this group being 17.8%. Black African and South Asian patients had the highest IGRA positivity rates (both 18.1%). Patients from a country where TB incidence was more than 50 per 100 000

population had higher positivity rates: 17.3% (106/614) compared with 2.7% (12/439) for patients from low-TB-incidence countries (<50/100,000). Of the 12 from low-TB-incidence countries, only four (33.3%) had risk factors (Table S2, Supplemental Digital Content, <http://links.lww.com/QAD/C621>).

In univariable analysis, being born abroad, specifically in sub-Saharan Africa and the South Asia and East Asia and Pacific regions, and being of black African or South Asian ethnicities, were associated with positive IGRA (Table 2). TB incidence in country-of-birth was significant in both univariable and multivariable analysis with increasing likelihood of having a positive IGRA amongst individuals born in countries with TB incidence more than 50 per 100 000 population.

Yields by testing threshold

Table 3 outlines the outcome of PWH IGRA test screening in Leicester stratified by TB incidence in country-of-birth, as well as outcomes for other screening strategies including BHIVA 2018 [14] and NICE [13] guidance and our proposed alternative ‘PWH-LTBI streamlined guidance’ for targeted testing. As the incidence at which screening is instigated increases, fewer PWH are eligible to be screened and, consequently, the number of identified LTBI cases also decreases. The yield (IGRA positivity rate amongst those tested) does not correspondingly increase once above the 40 per 100 000 population on BHIVA 2018 incidence threshold because we did not observe a linear increase in IGRA positivity for PWH from countries with TB incidence in country-of-birth more than 40 per 100 000 population (Table 1).

The strategy we identified as optimizing yield and efficiency of testing (the ‘PWH-LTBI streamlined guidance’) involves testing all PWH with country-of-birth TB incidence greater than 150 per 100 000 plus all sub-Saharan African countries. Application of NICE [13] and international [11,12] guidance, that is, screening all PWH in our cohort, identifies 100% of IGRA-positive cases with yield 11.2%. Applying BHIVA 2018 [14] guidance or our proposed ‘PWH-LTBI streamlined guidance’ both reduce the number of patients eligible for screening (to 622, 59.1% and 568, 53.9%, respectively). These screening strategies produce yields of 17.7 and 18.7%. Both yields are significantly higher than NICE [13] guidelines (proposed guidance v NICE, $P < 0.0001$; BHIVA guidance v NICE, $P = 0.0002$). BHIVA 2018 [14] guidance misses marginally fewer infections than in our proposed strategy (percentage IGRA positives correctly identified 93.2% versus 89.8%). There was no statistically significant difference in any of the outcomes shown in Table 3 between BHIVA 2018 [14] and the ‘PWH-LTBI streamlined guidance’ ($P = 0.66$).

Discussion

Our study describes a large prospective, systematic LTBI-screening programme implemented among PWH in a low-TB-incidence country and is the first to report chemoprophylaxis treatment uptake and completion rates. Overall, 11.1% (117/1053) of screened patients had LTBI, confirming that there is significant potential to reduce incident TB rates amongst PWH in the UK. TB incidence in this Leicester cohort is extremely high: of the 2158 patients ever treated for HIV in Leicester, 325 (15%) have had active

TB, with 100 of these (31%) having incident TB occurring more than 3 months after HIV diagnosis [15,27]. Therefore, it is imperative that the burden of LTBI amongst PWH is addressed to prevent incidence of active infection. Our study showed high acceptance of LTBI testing among PWH, with high chemoprophylaxis uptake and completion for IGRA-positive patients. It is, therefore, feasible to achieve high levels of retention at each stage of the cascade of care.

Our assessment of the outcomes of IGRA screening at different incidence thresholds and using different testing guidelines showed that an alternative to current NICE [13] and BHIVA [14] guidelines, the 'PWH-LTBI streamlined guidance', performed statistically significantly as well as BHIVA guidelines in reducing number of IGRA tests performed and increasing yield of LTBI identified. Additionally, it offered a simpler, more streamlined approach to testing than BHIVA guidance, without the need to consult a complex set of TB risk factors to determine test eligibility that may constitute a barrier to effective implementation. 89.8% of IGRA positive cases could have been identified by restricting screening to those from countries with TB more than 150 per 100 000 population or any sub-Saharan African country. This strategy led to a significantly higher yield (LTBI positivity rate) in those tested than if all patients were screened, as is currently proposed in the ECDC, WHO and 2016 NICE guidelines [11–13].

Extremely few patients declined IGRA testing (0.6%), although a higher proportion of these IGRA-positive individuals declined chemoprophylaxis (7.7%). Over 85% of IGRA-positive individuals started chemoprophylaxis, comparing favourably with rates of 17–87% from elsewhere in the UK and other low-TB-incidence countries [18,20,28]. Ninety-six percent successfully completed treatment, and adverse drug effects from chemoprophylaxis led to cessation of therapy in only two of 100 (2%) cases, supporting previous evidence showing that chemoprophylaxis regimens, and particularly isoniazid monotherapy regimens, are well tolerated in PWH [10,29]. Although there was high retention at each stage of the cascade of care, small drop-outs at each stage still led to 14.5% of IGRA-positive cases not being treated. Further research to identify barriers and facilitators to improve uptake are required in order to avert reactivation to active TB cases as far as possible.

We were fortunate to have all data available on the country of birth for patients in our cohort, which made analysis straightforward. Encouraging the recording of country-of-birth without stigma or discrimination is helpful in health systems, so that any targeted testing based on country-of-birth can be implemented effectively UK-wide.

Our work has several limitations. Most notable of these is generating testing eligibility estimates according to the 2018 BHIVA guidance testing criteria, which recommends offering IGRA testing to PWH from low-TB-burden countries (<40/100 000 population) only if additional risk factors for TB are present (see details in Table 3 footnotes) [14]. TB risk factor data were not collected prospectively for IGRA tests performed pre-2018 (date of BHIVA guidance [14] publication). Information on risk factors was retrieved from medical records only for IGRA-positive cases from low-TB-burden countries. Therefore, our estimate of IGRA eligibility under BHIVA guidance [14] is likely to be an underestimate.

This would make our proposed 'PWH-LTBI streamlined guidance' even more efficient than BHIVA 2018 guidance [14] in reducing IGRA tests required.

Secondly, we included indeterminate results in the denominator, which will lead to an under-estimation of the overall IGRA positivity rate. As there were only two cases of indeterminate results, however, this effect will be marginal. A further limitation was that we used country-specific TB incidence data available at a single time-point in our analysis, rather than using incidence estimates corresponding to year of entry to the UK for non-UK-born PWH. TB incidence may have changed in individual countries over time. However, date of UK entry was incomplete in our dataset and may not be routinely available, and an accessible, risk-based testing approach requires a simplified approach.

LTBI prevalence was moderately high at 11.1% for the whole cohort compared with 7–10% in other settings [18–20]. IGRA positivity for PWH from low-TB-incidence countries was comparable: 3.1% among PWH born in countries with TB incidence less than 30 per 100 000 in London [20] compared with 2.7% for those from less than 50 per 100 000 in our study. Key to the performance of screening criteria dependent on TB incidence in country-of-birth is LTBI prevalence amongst those from countries below the determined threshold. It is reassuring to observe a similar prevalence from a contrasting UK region, but more evidence on IGRA positivity rates by TB incidence in country-of-birth for other PWH populations in the UK would be useful to validate our proposed testing strategy and determine the generalizability of the results.

Our proposed 'PWH-LTBI streamlined guidance' performed statistically significantly as well as BHIVA 2018 guidance in terms of yield, number screened and proportion of latent infections identified. The next step is to undertake a full cost-effectiveness analysis of this and other LTBI testing strategies for PWH, both for the Leicester cohort and more generally across the UK. This would bring together the costs of the intervention, not only in terms of IGRA tests and chemoprophylaxis but also costs saved by averting cases of active TB and associated health benefits of reducing active TB morbidity and mortality, under a single framework, to inform formation of the next round of UK guidance. A previous cost-effectiveness analysis of LTBI screening among PWH based in London found that a targeted approach to screening was more cost-effective than universal testing but at the expense of missing some cases [30,31]. We now have the empirical data to inform new health economic analyses with realistic assumptions regarding IGRA positivity rates by risk group, chemoprophylaxis uptake and treatment.

This large, prospective screening cohort showed that PWH from high-TB-burden countries are at the highest risk of having LTBI but also that programmatic LTBI screening is achievable and can lead to impressive outcomes in terms of chemoprophylaxis completion. We now recommend that a full cost-effectiveness analysis is undertaken in order to produce the most user-friendly, evidence-based guidelines for screening in the UK and other low-TB-incidence settings, to enable consistent implementation.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgements

Data sharing

The patient cohort was extracted under Caldicott Guardian approval for a specific purpose and as part of our undertaking with them, we are not to further routinely share the data presented here. The data are held in an institutional repository and interested parties, with appropriate approvals, can apply for data access through the Corresponding author. Reasonable requests will be assessed on a case-by-case basis in discussion with the Caldicott Guardian.

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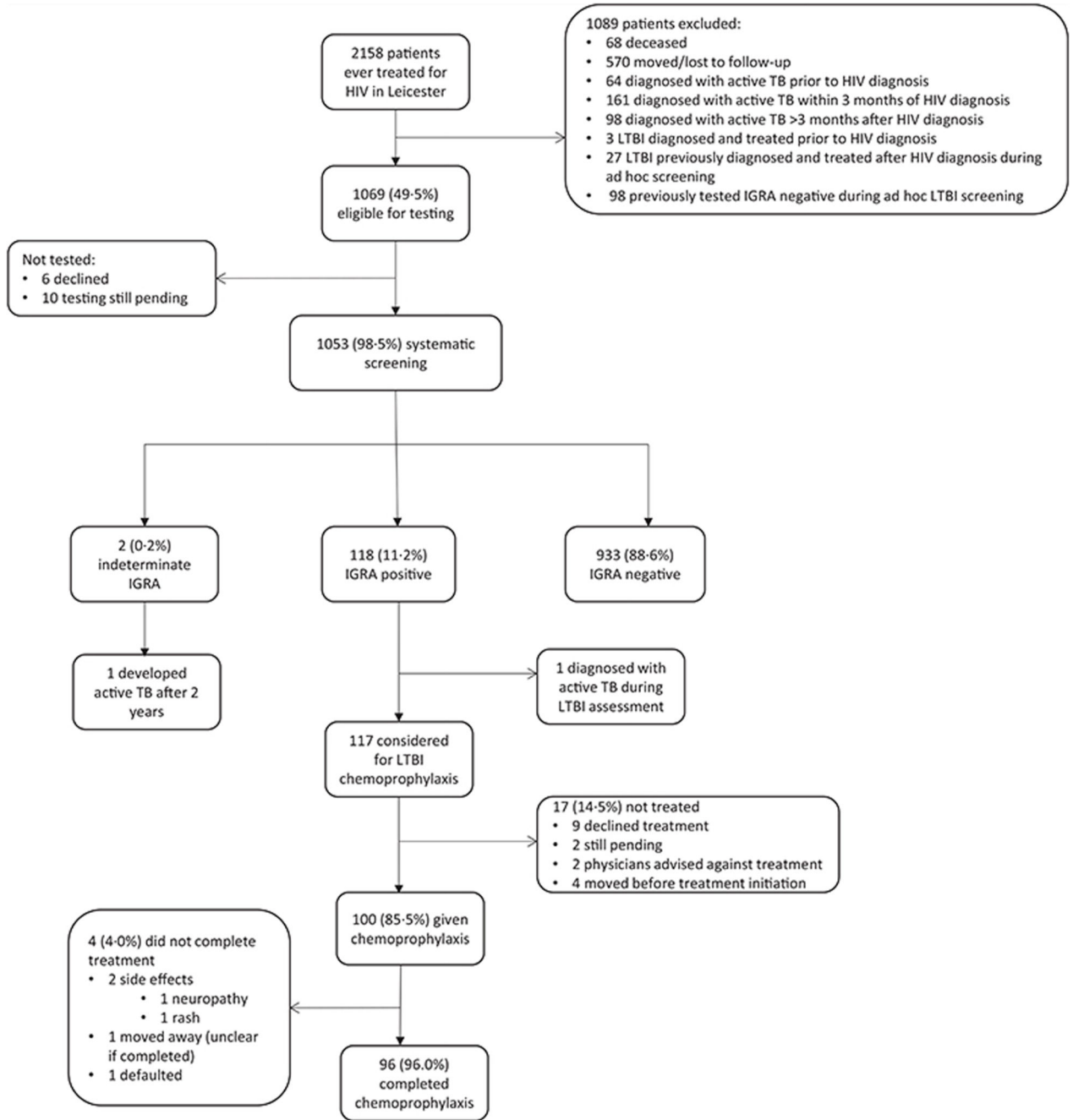


Fig. 1. Latent tuberculosis infection screening and treatment cascade of care. IGRA, interferon gamma release assay; LTBI, latent TB infection.

Table 1
Description of total cohort and those testing interferon gamma release assay-positive.

Variable	Total screened cohort [<i>n</i> (%)]		Proportion IGRA-positive [<i>x/n</i> (%)]	
Total	1053	(100)	118/1053	(11.2)
Median age at IGRA test, years (IQR)	42	(36–49)	42	(38–48)
Year of HIV diagnosis				
1985–1989	8	(0.8)	2/8	(25.0)
1990–1999	68	(6.5)	3/68	(4.4)
2000–2009	604	(57.4)	74/604	(12.3)
2010–2017 ^a	373	(35.4)	39/373	(10.5)
Year of IGRA test				
2014	393	(37.3)	43/393	(10.9)
2015	358	(34.0)	47/358	(13.1)
2016	147	(14.0)	18/147	(12.2)
2017	125	(11.9)	8/125	(6.4)
2018	24	(2.3)	2/24	(8.3)
2019	3	(0.3)	0/3	(0.0)
220	1	(0.1)	0/1	(0.0)
2021 ^b	2	(0.2)	0/2	(0.0)
Sex				
Male	597	(56.7)	53/597	(8.9)
Female	456	(43.3)	65/456	(14.3)
CD4 ⁺ cell count at IGRA testing (cells/μl)				
Median (IQR)	530	(380–700)	545	(440–720)
Range	10–2260		90–1350	
200	48	(4.6)	2/48	(4.2)
201–350	181	(17.2)	12/181	(6.6)
351–500	250	(23.7)	36/250	(14.4)
>500	574	(54.5)	68/574	(11.8)
Ethnicity				
Black African	498	(47.3)	90/498	(18.1)
South Asian	94	(8.9)	17/94	(18.1)
White	388	(36.8)	9/388	(2.3)
Mixed	16	(1.5)	0/16	(0.0)
Black Caribbean	14	(1.3)	0/14	(0.0)
Black other	10	(0.9)	0/10	(0.0)
Other	32	(3.0)	2/32	(6.3)
Unknown	1	(0.1)	0/1	(0.0)
UK birth status				
UK born	361	(34.3)	10/361	(2.8)
Non-UK born	692	(65.7)	108/692	(15.6)

Variable	Total screened cohort [<i>n</i> (%)]		Proportion IGRA-positive [<i>x/n</i> (%)]	
Region of birth				
Sub-Saharan Africa	538	(51.1)	96/538	(17.8)
South Asia	50	(4.7)	8/50	(16)
Europe and Central Asia	421	(40.0)	12/421	(2.9)
East Asia and Pacific	22	(2.1)	2/22	(9.1)
Latin America and Caribbean	13	(1.2)	0/13	(0.0)
Middle East and North Africa	5	(0.5)	0/5	(0.0)
North America	4	(0.4)	0/4	(0.0)
TB incidence in country-of-birth				
<50/100 000 population	439	(41.7)	12/439	(2.7)
50–149/100 000 population	58	(5.5)	12/58	(20.7)
150–249/100 000 population	427	(40.6)	74/427	(17.3)
250–349/100 000 population	63	(6.0)	11/63	(17.5)
350/100 000 population	66	(6.3)	9/66	(13.6)
Type of IGRA performed				
QuantiFERON-TB test ^c only	1013	(96.2)	115/1013	(11.4)
QuantiFERON-TB tests ^c and T- SPOT. <i>TB</i>	25	(2.4)	2/25	(8)
T-SPOT. <i>TB</i> only	15	(1.4)	1/15	(6.7)

IGRA, interferon gamma release assay; IQR, interquartile range.

^a Individuals were included up and including to 30 June 2017.

^b 30 June 2021 was used as the cut-off for following up patients for IGRA testing.

^c QuantiFERON-TB GIT or QuantiFERON-TB Plus.

Table 2
Univariate and multivariable logistic regression for having a positive interferon gamma release assay test at latent tuberculosis infection screening.

Variable		Observation (%)	Unadjusted OR (univariate analysis)	P value	Adjusted OR (multivariable analysis)	P value
Age at IGRA test (years)			1.01 (0.992–1.028)	0.26	1.022 (1.0–1.05)	0.05
Year of HIV diagnosis			0.99 (0.96–1.02)	0.53	1.03 (0.99–1.07)	0.23
Sex	Male	53/118 (44.9)	1		1	
	Female	65/118 (55.1)	1.71 (1.16–2.51)	0.01	0.95 (0.62–1.45)	0.79
CD4 ⁺ cell count at IGRA test (cells/ μ l)	<200	2/118 (1.7)	1		1	
	200–349	12/118 (10.2)	1.63 (0.36–7.56)	0.53	1.67 (0.35–7.97)	0.52
	350–499	36/118 (30.5)	3.87 (0.90–16.65)	0.008	4.39 (0.99–19.53)	0.05
	500	68/118 (57.6)	3.09 (0.73–13.02)	0.12	3.92 (0.89–17.12)	0.07
Ethnicity	Black ^a	90/118 (76.3)	1		–	–
	South Asian	17/118 (14.4)	1.06 (0.60–1.88)	0.84	–	–
	White	9/118 (7.6)	0.11 (0.06–0.23)	<0.0001	–	–
	Mixed/other	2/118 (1.7)	0.20 (0.05–0.86)	0.03	–	–
UK birth status	Non-UK born	108/118 (91.5)	1		–	–
	UK born	10/118 (8.5)	0.15 (0.08–0.30)	<0.0001	–	–
World Bank region of birth	Europe and Central Asia,	12/118 (10.2)	1		–	–
	North America and Latin America and Caribbean and Middle East and North Africa ^b					
	South Asia and East Asia and Pacific ^c	10/118 (8.5)	5.79 (2.40–13.97)	<0.0001	–	–
	Sub-Saharan Africa	96/118 (81.4)	7.80 (4.22–14.42)	<0.0001	–	–
	Less than 50 of 100 000 population	12/118 (10.2)	1		1	
	50–149/100 000 population	12/118 (10.2)	9.28 (3.94–21.85)	<0.0001	11.6 (4.79–28.10)	<0.0001
TB incidence in country-of-birth [26] ^f	150–249/100 000 population	74/118 (62.7)	7.46 (3.99–13.95)	<0.0001	8.26 (4.27–15.98)	<0.0001
	250–349/100 000 population	11/118 (9.3)	7.53 (3.16–17.92)	<0.0001	8.13 (3.33–19.86)	<0.0001
	350/100000 population	9/118 (7.6)	5.62 (2.27–13.92)	<0.0001	6.16 (2.42–15.67)	<0.0001

IGRA, interferon gamma release assay; OR, odds ratio.

^a All were black African; none were black Caribbean or black other.

^b All were from Europe and Central Asia; none were from Latin America and Caribbean, North America or the Middle East and North Africa.

^cEight of 10 were from South Asia; 2/11 were from East Asia and Pacific region.

Table 3
Yield and percentage of interferon gamma release assay-positive results obtained by implementing latent tuberculosis infection screening at different tuberculosis incidence thresholds.

Threshold: TB incidence in country-of-birth	Number screened (%)	Number IGRA positive	Yield (% IGRA positive of those tested)	Percent of all IGRA positives correctly identified
>350/100 000	66 (6.3)	9	13.6%	7.6%
>250/100 000	129 (12.3)	20	15.5%	16.9%
>150/100 000	556 (52.8)	94	16.9%	79.7%
>150/100 000 plus all sub-Saharan African countries: the proposed 'PWH-LTBI streamlined guidance'	568 (53.9)	106	18.7%	89.8%
>50/100 000	614 (58.3)	106	17.3%	89.8%
40/100 000 plus risk factors: BHIVA 2018 guidelines ^a [14]	622 ^b (59.1)	110	17.7%	93.2%
Screen all PWH:* 2016 NICE guidelines [13], ECDC guidelines for the EU/EEA [11], WHO guidelines for low tuberculosis burden countries [12]	1053 (100)	118	11.2%	100%

BHIVA, British HIV Association; ECDC, European Centre for Disease Prevention and Control; EEA, European Economic Area; EU, European Union; IGRA, interferon gamma release assay; NICE, National Institute for Health and Care Excellence. All included guidelines mention dual use of IGRA/Mantoux testing in some, or all PWH. We have assumed in this table that IGRA is as effective at diagnosing LTBI as Mantoux.

^aRecommends screening all those from high (>150/100 000 population) or medium (40-150/100 000 population) TB incidence countries; only screening those from low-TB-burden countries (<40/100 000 population) if additional risk factors for TB are present: CD4⁺ cell count less than 200 cells/μl; recent exposure to a known TB case; diabetes mellitus; stage 4/5 chronic kidney disease; receipt of chemotherapy for malignancy; immunosuppression following transplantation; biological disease modifiers for inflammatory conditions; prolonged duration of high-dose corticosteroids (prednisolone 20 mg o.d., or equivalent, for ≥2 months); travel to or periods of time spent in medium-incidence or high-incidence countries; history of working in medical settings in countries with medium or high TB incidence; injecting drug use (detailed in Table 6.1 of guidance [14]).

^bThis figure is an underestimate (includes all patients from countries where TB at least 40/100 000 population; plus four IGRA-positive patients from countries where TB incidence less than 40/100 000 for whom BHIVA cited additional risk factors were evident, but does not include patients with negative IGRA results from countries where TB incidence less than 40 of 100 000 because BHIVA-cited risk factors were not collected prospectively).