



## BRIEF CONCLUSIVE REPORT

# IL-32 and its splice variants are associated with protection against *Mycobacterium tuberculosis* infection and skewing of Th1/Th17 cytokines

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### Abstract

Studies in IL-32 transgenic mice and in vitro suggest that IL-32 may have protective effects against *Mycobacterium tuberculosis*, but so far there are barely any studies in humans. We studied the role of IL-32 and its splice variants in tuberculosis (TB) in vivo and in vitro. Blood transcriptional analysis showed lower total IL-32 mRNA levels in pulmonary TB patients compared to patients with latent TB infection and healthy controls. Also, among Indonesian household contacts who were heavily exposed to an infectious TB patient, IL-32 mRNA levels were higher among those who remained uninfected compared to those who became infected with *M. tuberculosis*. In peripheral blood mononuclear cells from healthy donors, we found that IL-32 $\gamma$ , the most potent isoform, was down-regulated upon *M. tuberculosis* stimulation. This decrease in IL-32 $\gamma$  was mirrored by an increase of another splice variant, IL-32 $\beta$ . Also, a higher IL-32 $\gamma$ /IL-32 $\beta$  ratio correlated with IFN- $\gamma$  production, whereas a lower ratio correlated with production of IL-1Ra, IL-6, and IL-17. These data suggest that IL-32 contributes to protection against *M. tuberculosis* infection, and that this effect may depend on the relative abundance of different IL-32 isoforms.

### KEYWORDS

tuberculosis, *Mycobacterium tuberculosis*, immune response, interleukin-32, cytokines

## 1 | INTRODUCTION

Tuberculosis (TB) is an airborne infectious disease caused by *Mycobacterium tuberculosis*. TB remains a major public health problem, and approximately one-fourth of the world population is latently infected with *M. tuberculosis*.<sup>1</sup> Host factors may determine whether *M. tuberculosis* exposure results in infection, and whether infection progresses to disease. A comprehensive understanding of the immune response against *M. tuberculosis* is crucial for the development of preventive strategies.

Interleukin-32 (IL-32), which was previously called natural killer cell transcript 4, has been identified as an important player in innate and adaptive immune responses.<sup>2</sup> Although no receptor for IL-32 has been discovered so far, IL-32 acts as a pro-inflammatory cytokine<sup>3</sup> and an intracellular regulator of cytokine production, including tumor necrosis factor  $\alpha$  (TNF- $\alpha$ ).<sup>4</sup> IL-32 is abundantly expressed in T-cells and NK cells, but also in the lung, and alternative splicing of IL-32 mRNA results in at least 9 distinct isoforms,<sup>5</sup> of which not all functions are known yet.<sup>2</sup> The IL-32 $\gamma$  isoform is the most potent pro-inflammatory cytokine inducer,<sup>6</sup> which can splice into the most abundant isoform IL-32 $\beta$ .<sup>7,8</sup>

Abbreviations: IGRA, IFN-gamma release assay; TB, tuberculosis.

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Particular single nucleotide polymorphisms (SNPs) influence total IL-32 expression in different tissue types,<sup>9</sup> but polymorphisms influencing splicing are currently unknown. Although our knowledge of the role of IL-32 splice variants in health and disease has expanded, many aspects regarding their mechanism of action still remain unknown.

Recent studies have identified IL-32 as a modulating factor in the host response against *M. tuberculosis*. Stimulation of human PBMCs with heat-killed mycobacteria induced a strong production of IL-32, which is dependent on endogenous IFN- $\gamma$ .<sup>10</sup> In human monocyte-derived M $\phi$ s, addition of recombinant IL-32 increased killing of *M. tuberculosis* in vitro, and this effect was dependent on vitamin D.<sup>11</sup> Addition of IL-32 $\gamma$  induced caspase-3, caspase-1, and cathepsin-mediated apoptosis in THP-1 M $\phi$ s, and reduced the intracellular burden of *M. tuberculosis*.<sup>12</sup> Finally, transgenic mice expressing human IL-32 $\gamma$  in type II alveolar epithelial cells showed an 85% reduction in *M. tuberculosis* outgrowth in the lungs compared to control mice.<sup>13</sup> Together, these studies suggest a potential protective role for IL-32 upon *M. tuberculosis* infection.

Although it has been shown that *M. tuberculosis* induces IL-32 production,<sup>10</sup> little is known about the different splice variants in the context of TB in primary cells. In addition, there are limited reports of studies performed in humans to support a role for IL-32 in vivo. We therefore examined whole blood IL-32 expression profiles in different TB phenotypes, and explored expression of IL-32 isoforms in response to *M. tuberculosis* in vitro.

## 2 | MATERIALS AND METHODS

Detailed material and methods can be found as Supplementary Information.

### 2.1 | Patient whole blood gene expression

We examined previously published whole blood gene expression data from patients with pulmonary TB, individuals with latent TB infection, and healthy controls from several cohorts from Africa and Europe.<sup>14–19</sup> The datasets were retrieved from the Gene Expression Omnibus (GEO) using the GSE identifiers GSE83456, GSE42826, GSE19491, GSE28623, GSE37250, and GSE34608. We also examined whole blood expression of pulmonary TB patients from the United Kingdom (GSE19491) during follow-up at 2 and 12 months into TB treatment.

In addition, we measured whole blood IL-32 expression among household contacts of patients with active pulmonary TB. A total of 44 household contacts of TB cases in an urban setting in Indonesia were recruited within 2 weeks of the index patient starting TB treatment. Using QuantiFERON-TB Gold, the presence of *M. tuberculosis* infection was tested at baseline and 3 months afterward. These IFN-gamma release assays (IGRAs) identified persistently negative contacts, who were exposed but remained uninfected, and converters<sup>20</sup>; individuals who were positive at baseline were not included.

### 2.2 | In vitro stimulation experiments

Clinical isolates were selected from a previous study<sup>21</sup> for in vitro stimulation of PBMCs of 8 healthy volunteers. Details can be found in the Supplementary Information. In short, 19 clinical isolates and the laboratory strain H37Rv were used at a 3  $\mu$ g/mL concentration to stimulate PBMCs ( $5 \times 10^5$  cells/well) in duplicate in a 96-well plate. The plates were incubated for 24 h (for TNF- $\alpha$ , IL-1 $\beta$ , IL-1Ra, IL-10, and IL-6 quantification) or 7 days (for IFN- $\gamma$ , IL-17, and IL-22 quantification) at 37°C in a 5% CO<sub>2</sub> environment. Cytokines in the supernatants were measured using commercial ELISA kits. RNA from stimulated PBMCs was used for quantitative PCR using different IL-32 primer sets and was corrected for expression of the housekeeping gene  $\beta$ 2-microglobulin (B2M).

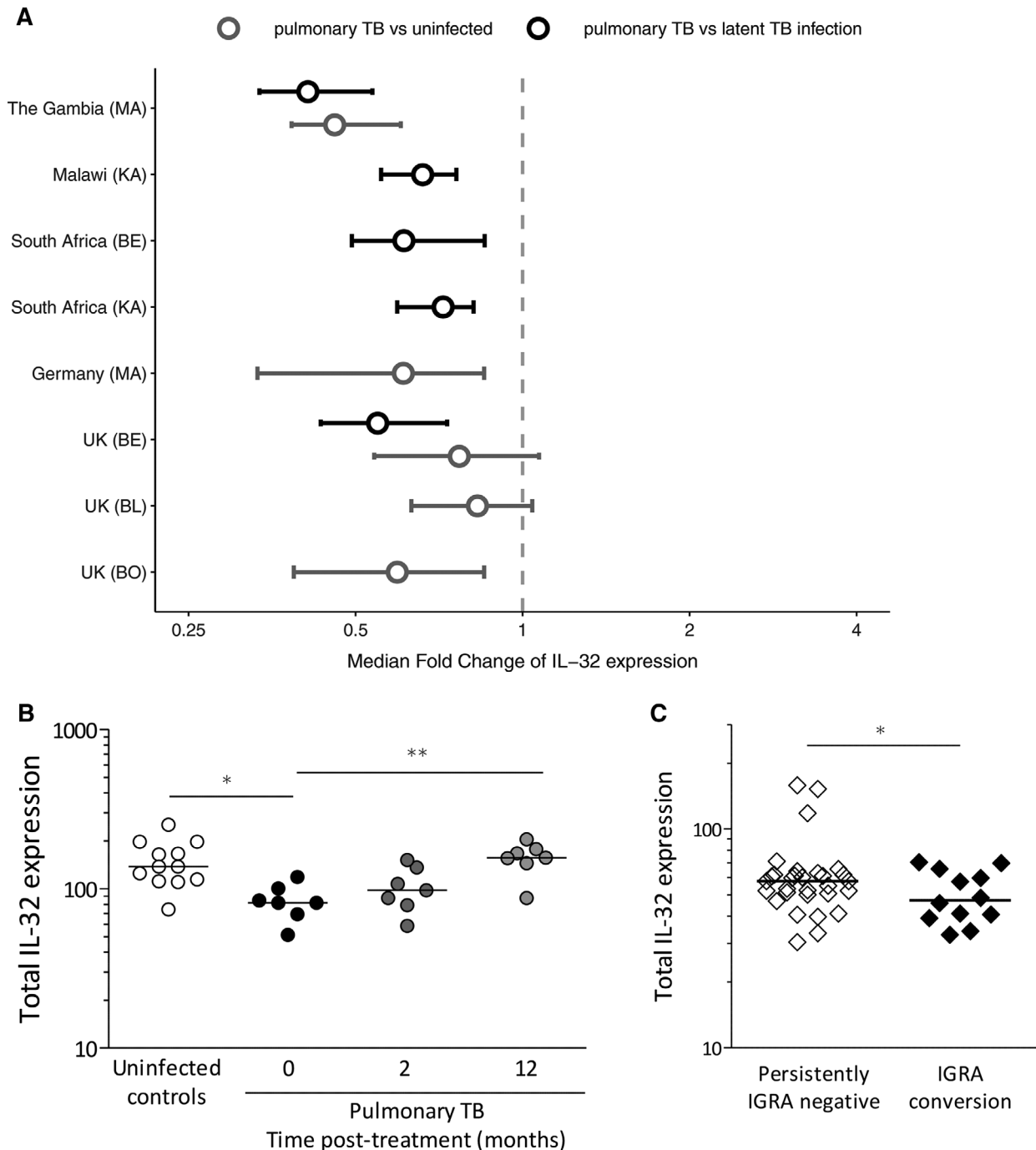
### 2.3 | Data analysis and statistics

All analyses were performed using GraphPad Prism version 5.3 or in R 3.2.4 using RStudio. Statistical analyses of the whole blood microarray data were performed using Kruskal–Wallis tests, including the post hoc Dunn's multiple comparison tests, and Mann–Whitney *U*-tests. Differential gene expression analysis for IL-32 in the TB contact cohort was performed using the R package DESeq2.<sup>22</sup> Multiple univariate linear regression analyses assessed the relation between the IL-32 $\beta$ /IL-32 $\gamma$  ratio and cytokine levels. Cytokine levels were positively skewed and therefore log-transformed. *P*-values were corrected for multiple testing using Bonferroni correction, and *P*-values of 0.05 or less were considered statistically significant.

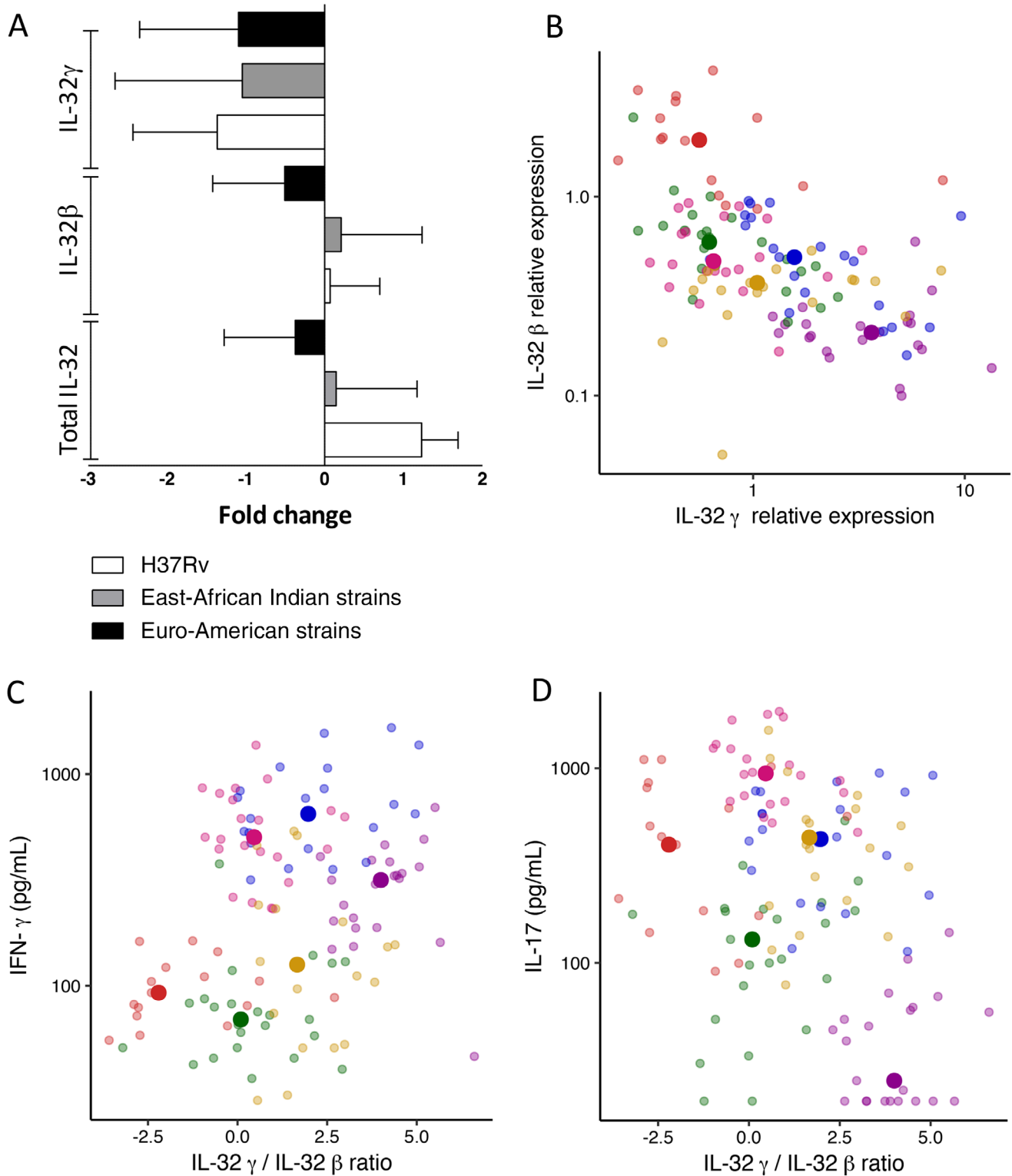
## 3 | RESULTS AND DISCUSSION

### 3.1 | IL-32 expression levels differ between TB phenotypes

We first compared expression of total IL-32 in pulmonary TB patients with individuals with latent TB infection and healthy controls using publicly available data. IL-32 mRNA expression levels were lower in pulmonary TB patients compared to healthy controls and individuals with latent TB infection, and this was consistently observed in several cohorts (Fig. 1A). During the course of TB treatment, levels IL-32 restored to normal (Fig. 1B). To examine a possible protective role of IL-32 in primary *M. tuberculosis* infection, whole blood transcription levels were compared between 32 TB household contacts that remained IGRA-negative upon heavy exposure to an infectious TB patient (so-called “early clearers<sup>20</sup>”) and 12 household contacts who became infected with *M. tuberculosis*. Total IL-32 mRNA levels were higher in the persistently IGRA-negative group compared to those who developed latent TB infection ( $P = 1.8 \times 10^{-2}$ ; Fig. 1C). The difference in IL-32 expression was not caused by a difference in lymphocyte count: 2.64 (interquartile range [IQR], 2.08–3.06)  $\times 10^9$ /L in individuals who remained IGRA negative vs. 2.56 (IQR 2.13–3.13) in patients who converted to a positive IGRA ( $P = 0.37$ ).



**FIGURE 1** IL-32 expression in different tuberculosis phenotypes. (A) Whole blood IL-32 gene expression data from healthy controls, latently infected individuals, and pulmonary TB patients are presented. The data were obtained from publicly available datasets published by Maertzdorf et al. (MA), Kaforou et al. (KA), Berry et al. (BE), Bloom et al. (BO), and Blankley et al. (BL). The hollow circles represent the median fold changes of IL-32 expression in tuberculosis patients compared to healthy controls (gray) or latently infected individuals (black), and the lines represent the 95% confidence intervals, determined by bootstrap resampling. (B) Whole blood IL-32 expression from pulmonary TB patients ( $N = 7$ ) at 0, 2, and 12 months after start of treatment is presented. Data were previously published by Berry et al., and medians are indicated. Statistical analyses of the IL-32 mRNA levels were performed using Kruskal–Wallis tests, including the post hoc Dunn’s multiple comparison tests, and Mann–Whitney  $U$ -tests. \* $P < 0.05$ ; \*\* $P < 0.01$ ; \*\*\* $P < 0.001$ . (C) IL-32 gene expression measured by RNA sequencing in whole blood of household contacts who remained uninfected (persistently IGRA negative,  $N = 32$ ) and contacts who became infected after exposure (IGRA conversion,  $N = 12$ ). Differential gene expression analysis of IL-32 was performed using the R package DESeq2 with RStudio in R 3.2.4 ( $P = 0.018$ ). Individual data points and medians are presented



**FIGURE 2** IL-32 expression in relation to cytokine production after stimulation with *M. tuberculosis*. (A) Total IL-32, IL-32 $\beta$ , and IL-32 $\gamma$  gene expression in PBMCs from 8 different donors stimulated with *M. tuberculosis* for 24 h. The East-Asian Indian strains group shows the median of 15 strains, and the Euro-American strains group shows the median of 4 strains compared to the unstimulated control. Data are presented as median  $\pm$  interquartile range (IQR). (B–D) PBMCs from 6 different donors were stimulated with 20 *M. tuberculosis* isolates for 24 h or 7 days. Each color represents a donor, and each dim point shows 1 individual data point. The brighter point shows the median value of 1 donor for all 20 *M. tuberculosis* isolates. (B) The relative expression of IL-32 $\gamma$  is plotted against the relative expression of IL-32 $\beta$  in PBMCs stimulated for 24 h with *M. tuberculosis* isolates. The levels of IL-32 $\beta$  and IL-32 $\gamma$  were measured using quantitative PCR and normalized against the housekeeping gene  $\beta$ 2-microglobulin (B2M) and the relative expression was calculated as  $2^{(-\Delta Ct)}$ . (C–D) Concentrations of IFN- $\gamma$  (C) and IL-17 (D) measured by ELISA after 7 days are plotted against IL-32 $\gamma$ /IL-32 $\beta$  ratio

**TABLE 1** Multiple linear regression models assessing the variability in cytokine production explained by IL-32 corrected for stimulus

	Ratio IL-32 $\gamma$ /IL-32 $\beta$	
	Estimate	P-value
TNF- $\alpha$	0.0377	0.1673
IL-1 $\beta$	-0.0372	0.0629
IL-1Ra	-0.0568	0.0010*
IL-6	-0.0506	0.0018*
IL-10	0.0285	0.1957
IFN- $\gamma$	0.143	0.0018*
IL-17	-0.213	0.0003*
IL-22	0.0614	0.1550

\*Significance at the Bonferroni-corrected level for the number of cytokines tested ( $\alpha = 0.05/8 = 0.00625$ ).

### 3.2 | Induction of IL-32 isoforms after *M. tuberculosis* stimulation in vitro

In order to explore which IL-32 isoforms are induced in TB, PBMCs from healthy volunteers were stimulated with 20 different *M. tuberculosis* clinical isolates, and after 24 h of stimulation, the mRNA levels of total IL-32 were determined by RT-PCR, as well as the isoforms primarily induced by *M. tuberculosis*<sup>8</sup>: IL-32 $\beta$  and IL-32 $\gamma$ . Different clinical isolates either up-regulated or down-regulated the expression of total IL-32 mRNA (Fig. 2 and Supplementary Fig. S1A). The isoform IL-32 $\beta$  followed the pattern of total IL-32 (Fig. 2 and Supplementary Fig. S1B). It has been shown before that genetically diverse mycobacterial strains vary widely in their induction of cytokines<sup>21,23</sup> and that these differences are observed between *M. tuberculosis* lineages<sup>24</sup> and even within lineages.<sup>25</sup> Here, we also observe differences in the expression of total IL-32 and IL-32 $\beta$  induced by different clinical isolates, suggesting that the host-microorganism relationship can determine the expression of different IL-32 isoforms. Nevertheless, all clinical isolates induced down-regulation of IL-32 $\gamma$  (Fig. 2 and Supplementary Fig. S1C). These in vitro experiments reveal that *M. tuberculosis* stimulation leads to active splicing of the most potent IL-32 isoform, IL-32 $\gamma$ , into other isoforms, including IL-32 $\beta$ .

### 3.3 | Association between IL-32 mRNA levels and cytokine profile

In human PBMCs, IL-32 $\beta$  and IL-32 $\gamma$  expression showed a strong inverse relationship ( $P = 6.6 \times 10^{-10}$ ) in a linear regression model corrected for stimulus, as visualized in Fig. 2B. We explored the immunological consequences by correlating IL-32 expression with *M. tuberculosis*-induced cytokines in PBMCs. The ratio between IL-32 $\gamma$  and IL-32 $\beta$  was predictive of cytokine production in a linear regression model with stimulus as covariate after correction for multiple testing (Table 1). The IL-32 $\gamma$ /IL-32 $\beta$  ratio correlated positively with IFN- $\gamma$  (Fig. 2C) and negatively with IL-6, IL-1Ra, and IL-17 (Fig. 2D). Together with previously published literature, these data suggest that different IL-32 isoforms have very distinct effects and regulation patterns. This

provides a strong rationale for measuring distinct IL-32 isoforms in future laboratory and clinical studies.

IFN- $\gamma$  has been repeatedly shown to be crucial in the host defense against TB.<sup>26,27</sup> In contrast, excessive IL-17 is thought to play a role in immunopathology through neutrophil recruitment. A balance between Th1 and Th17 responses is critical to control bacterial growth and to limit immunopathology.<sup>28</sup> Here, the IL-32 $\gamma$ /IL-32 $\beta$  ratio was inversely correlated with IL-17, and positively correlated to IFN- $\gamma$ . This suggests that splicing of IL-32 $\gamma$  may influence the balance between Th1 and Th17, and therefore play a role in steering the immune response in TB. This could indicate that IL-32 $\gamma$  contributes to control of *M. tuberculosis* infection and reduction of Th17-mediated lung damage.

## 4 | CONCLUDING REMARKS

Previous studies have suggested a protective effect of IL-32 against *M. tuberculosis*. We analyzed multiple TB cohorts to further explore these hypotheses in vivo. We observed decreased levels of total IL-32 mRNA in blood from pulmonary TB patients compared to individuals with latent TB infection and healthy controls. Additionally, in TB contacts in Indonesia, heavily exposed TB contacts who remained uninfected showed higher total IL-32 expression compared to those who became latently infected, suggesting a possible role in innate protection against *M. tuberculosis* infection. In PBMCs from healthy individuals, a higher ratio of IL-32 $\gamma$ /IL-32 $\beta$  expression correlated with higher IFN- $\gamma$  and a lower ratio with higher IL-17 cytokine responses. This is the first study to identify an association between IL-32 and resistance against *M. tuberculosis* infection in an in vivo setting in humans, and of IL-32 splice variants with specific *M. tuberculosis* cytokine profiles. More study is needed to unravel the exact mechanism of action of IL-32 and its splice variants in TB. Of note, IL-32 is also known to play a role in Leishmaniasis and viral infections, which might imply a role in host response against intracellular pathogens.<sup>29</sup>

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## DISCLOSURE

The authors declare no conflict of interest.

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## SUPPORTING INFORMATION

Additional information may be found online in the Supporting Information section at the end of the article.

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