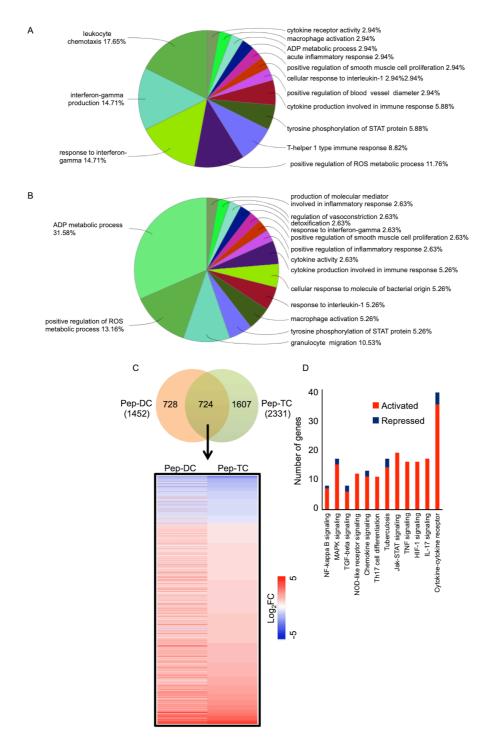
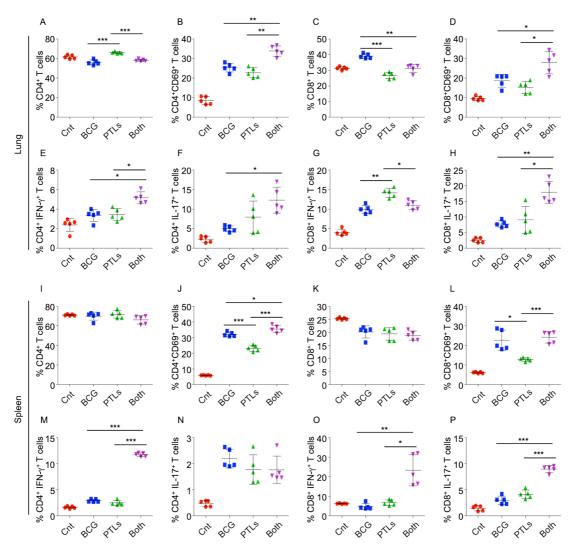
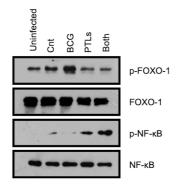
1 Supplementary Materials:



Supplementary Figure 1: *M.tb* peptides induce upregulation of genes that confer protection against TB in both DCs and T cells. (A) Pie diagram indicating the GO biological processes upregulated in DCs pulsed with peptide pool as compared to unstimulated DCs. (B) Pie chart indicating the percentage of genes belonging to different GO biological processes in stimulated T cells as compared to unstimulated T cells. (C) Heatmap depicting the expression profile of the genes common between the DCs and T cells. (D) KEGG pathway analysis of the shared genes representing the significantly affected pathways. Un-DC: unstimulated DCs, Pep-DC: DCs pulsed with peptide pool, Un-TC: T cells co-



Supplementary Figure 2: Pre-challenge immune responses in the lungs and the spleen of vaccinated mice. (**A-H**) T lymphocytes were isolated from the lungs of all experimental groups and stained with 7AAD, anti-CD3, anti-CD4, anti-CD8, anti-CD69, anti-IFN-γ and anti-IL-17 antibodies. Percentage of CD4⁺T cells (**A**), CD4⁺CD69⁺T cells (**B**), CD8⁺T cells (**C**) and CD8⁺CD69⁺T cells (**D**) in the lungs of unvaccinated (Cnt) and vaccinated animals. (**E-H**) Expression of IFN-γ and IL-17 on CD4⁺ and CD8⁺ T cells in the lungs. (**I-P**) T lymphocytes isolated from the spleen of all experimental groups were stained with 7AAD, anti-CD3, anti-CD4, anti-CD8, anti-CD69, anti-IFN-γ and anti-IL-17 antibodies. Percentage of CD4⁺ T cells (**I**), CD4⁺CD69⁺ T cells (**J**), CD8⁺ T cells (**K**) and CD8⁺CD69⁺ T cells (**L**) in the spleen of different groups. (**M-P**) Expression of IFN-γ and IL-17 on CD4⁺ and CD8⁺ T cells in the spleen. One-way ANOVA followed by multiple tukey tests was performed for statistical analysis. Data is representative of two independent experiments (n=5 mice/group). *p<0.05, **p<0.005, ***p<0.005.



 Supplementary Figure 3: Enhanced NF-kB and FOXO-1 activation in the spleen of infected mice co-immunized with BCG-PTLs. Cell lysates prepared from the splenocytes of infected mice from all the experimental groups were used to analyze the phosphorylation status of FOXO-1 and NF-kB by immunoblotting. Data is representative of two independent experiments.

Supplementary Table 1: List of Peptides used in the study

S.			Protein	Rv
No.	Peptide Sequence	Position	name	number
		10aa -		
P1	AWGRRLMIGTAAAVVLPG	27aa	Ag85B	Rv1886c
		19aa - 36		
P2	TAAAVVLPGLVGLAGGAA	aa	Ag85B	Rv1886c
		91aa -		
P3	WDINTPAFEWYYQSGLSI	108aa	Ag85B	Rv1886c
		29aa -		
P4	LDEGKQSLTKLAAAW	43aa	ESAT6	Rv3875
		33aa -		
P5	KQSLTKLAAAWGGSG	47aa	ESAT6	Rv3875
		37aa -		
P6	TKLAAAWGGSGSEAY	51aa	ESAT6	Rv3875
		72aa -		
P7	LARTISEAGQAMASTEGNVTGMFA	95aa	ESAT6	Rv3875
		61aa-		
P8	AVAASNNPELTTLTAALSGQLNPQV	85aa	Mpt70	Rv2875
		76aa-		
P9	ALSGQLNPQVNLVDTLNSGQYTVFA	100aa	Mpt70	Rv2875
		106aa-		
P10	FSKLPASTIDELKTNSSLLTSILTY	130aa	Mpt70	Rv2875
		166aa-		
P11	GNADVVCGGVSTANATVYMIDSVLM	190aa	Mpt70	Rv2875
		181aa-		
P12	TVYMIDSVLMPPA	193aa	Mpt70	Rv2875
		278-		
P13	AVDAADKVLGYRNWL	292aa	GlcB	Rv1837
		284-		
P14	KVLGYRNWLGLNKGD	298aa	GlcB	Rv1837

- 101 Supplementary Table 2: List of seven antigenic peptides used for further studies.
- Experiments from A-C were repeated with the pool of seven peptides.

103

Peptides	Sequences
P-1	AWGRRLMIGTAAAVVLPG
P-2	TAAVVLPGLVGLAGGAA
P-3	WDINTPAFEWYYQSGLSI
P-4	KQSLTKLAAAWGGSG
P-5	TKLAAAWGGSGSEAY
P-6	LDEGKQSLTKLAAAW
P-7	LARTISEAGQAMASTEGNVTGMEA