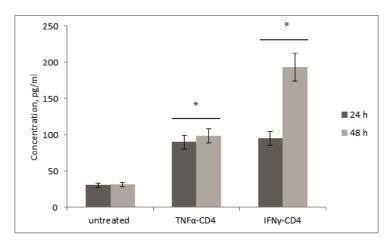
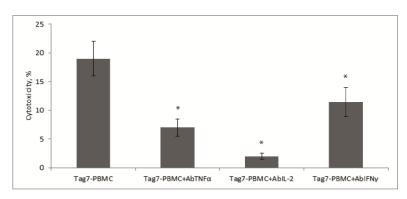


Supplementary Fig. S1. Changes in cytotoxic activity of different lymphocyte subpopulations induced in PBMC incubated with Tag7 (10-9M) for different periods of time. Lymphocyte subpopulations were isolated by magnetic cell separation and tested for cytotoxicity against K562 cells. Data are presented as the mean \pm SD of 3 independent experiments. Differences from the control in all cases are significant at * p < 0.03, ** p < 0.05 (Student t test).



Supplementary Fig. S2. Secretion of IL-2 by CD3⁺CD4⁺ - lymphocytes incubated with TNF α or IFN γ for 24 and 48 hours. CD3⁺CD4⁺ - lymphocytes were isolated by magnetic cell separation from PBMC on 0 day and treated with TNF α or IFN γ . Conditioned medium from untreated lymphocytes was used as a control. The medium was sampled every 24 h to determine IL-2 level by ELISA. Data are presented as the mean ± SD of 3 independent experiments. Differences from the control in all cases are significant at * p < 0.05 (2-way ANOVA).



Supplementary Fig. S3. Cytotoxicity Tag7-PBMC in presense specific antibodies to TNF α , IFN γ , IL-2. Each antibody (1:1000) was added to PBMC before 1h to Tag7 (10-9M). After 6 days of incubation PBMC were tested for cytotoxicity against K562 cells. Data are presented as the mean ± SD of 3 independent experiments. Differences from the control in all cases are significant at * p < 0.05 (Student t test).