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### Warnings

Reagents contain sodium azide. Under acidic conditions sodium azide yields hydrazoic acid, which is extremely toxic. Azide compounds should be diluted with running water before discarding. These precautions are recommended to avoid deposits in plumbing where explosive conditions may develop.

# 1. Description

This product is for research use only.

Components

1 mL monoclonal Rapid Cytokine Inspector (RCI) Anti-Cytokine Antibodies, human.

Conjugate	Clone/ Isotype	Order no.
RCI Anti-IFN-γ-PE	45-15/ mouse IgG1	130-097-600
RCI Anti-IFN-γ-APC	45-15/ mouse lgG1	130-097-596
RCI Anti-IL-2-PE	N7.48A/ mouse lgG1	130-099-450
RCI Anti-IL-17A-PE	CZ8-23G1/ mouse lgG1	130-097-599
RCI Anti-IL-17A-APC	CZ8-23G1/ mouse lgG1	130-099-448
RCI Anti-TNF-α- APC	cA2/ human lgG1	130-099-481

# Rapid Cytokine Inspector Anti-Cytokine antibodies

# human

Capacity 100 tests or up to 10<sup>8</sup> total cells

Product format Antibodies are supplied in buffer containing

stabilizer and 0.05% sodium azide.

Storage Store protected from light at 2-8 °C. Do not

freeze. The expiration date is indicated on the

vial label.

#### 1.1 Background information

The Rapid Cytokine Inspector Anti-Cytokine antibodies, human are optimized for use in combination with Rapid Cytokine Inspector Kits, human, to enable the fast intracellular staining of cytokines produced by activated T cells. The performance of the assay in 96-well cell culture plates enables the analysis of multiple samples in parallel.

### 1.2 Applications

For use in combination with Rapid Cytokine Inspector (RCI) Kits:

	RCI (CD4/CD8 T Cell) Kit (# 130-097-343)	RCI (CD4 T Cell) Kit (# 130-097-341)	RCI (CD8 T Cell) Kit (# 130-097-342)
IFN-γ	RCI Anti-IFN-γ-PE	RCI Anti-IFN-γ-PE	RCI Anti-IFN-γ-PE RCI Anti-IFN-γ-APC
IL-2	RCI Anti-IL-2-PE	RCI Anti-IL-2-PE	RCI Anti-IL-2-PE
IL-17	RCI Anti-IL-17A-PE	RCI Anti-IL-17A-PE	RCI Anti-IL-17A-PE RCI Anti-IL-17A-APC
TNF-α			RCI Anti-TNF-α-APC

# 2. Protocol overview

Stimulation	Plate $10^6$ cells (100 $\mu$ L) per well. Add antigen or control. After 2 hours of cultivation, add brefeldin A and continue cultivation for additional 4 hours.		
Cell surface staining	Pipette off 80 μL cell culture supernatant. Add 50 μL staining mix containing RCI T Cell Detection Cocktail and RCI Anti-Cytokine antibodies. Shake plate for 2 min and incubate for 8 min at RT.	Ī	
Fixation	Add 25 µL Inside Fix. Shake plate for 2 min and incubate for 18 min at RT.	45 minutes	
Permeabilization and intracellular staining	Add 25 µL Inside Perm. Shake plate for 2 min and incubate for 8 min at RT.	tes	
Washing step	Add 100 µL of buffer. Centrifuge plate for 5 min at 300×g. Remove supernatant and resuspend cells in 200 µL of buffer.		
Flow cytometric analysis	Proceed to sample aquisition and analysis.		

For a detailed protocol, refer to the data sheet of the respective Rapid Cytokine Inspector Kit.

Refer to **www.miltenyibiotec.com** for all data sheets and protocols. Miltenyi Biotec provides technical support worldwide. Visit www.miltenyibiotec.com for local Miltenyi Biotec Technical Support contact information.

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