Mouse IL-2 ELISA

Cat.n° 32670029U1

This ImmunoTools ELISA for mouse **Interleukin-2 (mIL-2)** contains appropriate reagents sufficient for processing of 1 microplate (1 x 96 wells; 100 μ l/well)

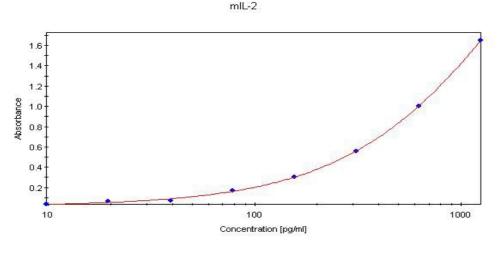
Typical standard curve range: 31 - 2000 pg/ml

Detection limit under optimal conditions: 26 pg/ml

Content	Working dilution	Storage
1 x vial 100 µl liquid anti-mIL-2 Capture-Antibody (red cap)	1:100	-20°C
1 x vial 100 µl liquid anti- mIL-2 Detection-Antibody (yellow cap)	1:100	-20°C
1 x vial 50 ng lyophilized rm IL-2 Standard (white cap)	customer specific	-20°C
1 x vial 10 μl Poly-HRP-Streptavidin (blue cap)	1:1000	-20°C

Additional material required:

96well-Microplates Wash-Buffer (e.g. PBS + 0.05% Tween20) Coating-Buffer (e.g. PBS) Blocking-Buffer / Reagent-Diluent (e.g. PBS + 2% BSA + 0.05% Tween20) Stop-Solution (e.g. 2 M H₂SO₄) TMB-Solution



4 Parameter (y = (A - D) / (1 + (x/C)^B) + D) A=4.4239 B=-1.0265 C=2099.2290 D=0.0168, R-Square = 0.9998

Note: All steps of incubation except <u>HRP-Streptavidin</u> and <u>TMB substrate</u> could be carried out overnight. Do not use solutions containing sodium azide, nor add sodium azide to the supplied reagents. Sodium azide inactivates the peroxidase.

Storage: Protect from light! Storage conditions are shown in the table above.

Reconstituted reagents should be stored at -20° C. Please prevent repeated freeze-thaw cycles. The performance of the unopened reagents is guaranteed until the expiration date shown on the label.

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Gladiolenweg 2, 26169 Friesoythe; Germany phone:+49-(0)4491-400997, fax:+49-(0)4491-400998, <u>info@immunotools.com</u> www.immunotools.com Spin down all vials before use!

Step	Incubation	Procedure		
Coating Capture- antibody	≥ OVERNIGHT at room temperature	Dilute <mark>capture-antibody</mark> 1:100 in COATING-BUFFER (100 µl capture-antibody in 10 ml COATING-BUFFER). Subsequently transfer 100 µl of this working-solution to each well and incubate.		
Remove capture-antibody completely by inverting the microplate and blotting it <i>vigorously</i> against clean paper towels.				
Blocking	≥ 1 Hour at room temperature	Add 300 µl BLOCKING-BUFFER to each well and incubate.		
Remove BLOCKING-BUFFER completely by inverting the microplate and blotting it <i>vigorously</i> against clean paper towels.				
Standard & Sample	≥ 2 Hours at room temperature	Dilute standard & samples in REAGENT-DILUENT and transfer 100 µl in the respective wells in duplicates. Standard: Make serial dilutions in REAGENT-DILUENT and begin with a high standard and end with blanks. The standard vial of this set contains 50 ng lyophilized standard . Reconstitute this in exactly 1 ml REAGENT-DILUENT (stock solution = 50 ng/ml) and choose a sufficient high standard working solution for your assay		
Wash 5x vigorously with WASHING-BUFFER and remove resting buffer completely by inverting the				
Detection- antibody	≥ 2 Hours at room temperature	ind blotting it vigorously against clean paper towels. Dilute detection-antibody 1:100 in REAGENT-DILUENT (100 μl detection-antibody in 10 ml REAGENT-DILUENT). Subsequently transfer 100 μl of this working-solution to each well and incubate.		
Wash 5x vi	Wash 5x vigorously with WASHING-BUFFER and remove resting buffer completely by inverting the microplate and blotting it vigorously against clean paper towels.			
Poly-HRP- Streptavidin	20-30 Min at room temperature	Dilute <mark>Poly-HRP-Streptavidin</mark> 1:1000 in REAGENT-DILUENT (10 μl in 10 ml REAGENT-DILUENT). Subsequently transfer 100 μl of this working-solution to each well and incubate.		
Wash 5x vi		SHING-BUFFER and remove resting buffer completely by inverting the ind blotting it vigorously against clean paper towels.		
Substrate solution	Up to 60 Min* at room temperature <u>in the dark</u>	Optionally warm the solution to room temperature before use. Add 100 µl of the SUBSTRATE-SOLUTION to each well and incubate. Control the development of the colour reaction continuously and stop at an appropriate time point.		
Stop solution	-	When the enzymatic colour reaction is sufficiently proceeded stop the reaction by adding of 50 μ l stop solution. Read the microplate at the substrate-depending wavelength. (e.g. 450 nm for TMB substrate) (if wavelength correction is available, set to 540 nm, 570 nm or 630 nm as reference)		

Warning: TMB-Solution is harmful if swallowed (R22). Keep out of reach of children (S2). Keep away from food, drink and animal feeding stuff (S13). Wear suitable protective clothing (S36). If swallowed, seek medical advice immediately and show this container or label (S46). Contact with acids liberates very toxic gas (R32). Azide compounds should be flushed with large volumes of water during disposal to avoid deposits in lead or copper plumbing where explosive conditions can develop.

This material is offered for **research only**. Not for use in diagnostic or therapeutic procedures.Not for use in human. For in vitro use only. ImmunoTools will not be held responsible for patent infringement or other violations that may occur with the use of our products.

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