This kit employs a sandwich enzyme immunoassay technique to quantify Human IL-17F in biological samples.

Human IL-17F ELISA Kit
Sandwich ELISA
Catalog EL-213-91217

INTRODUCTION

IL-17F is an inflammatory cytokine most closely associated with the Th17 lineage of T-helper cells, although it is also produced by CD8+ T cells, NK cells, mast cells, and other innate lymphoid populations. It signals largely as a heterodimer with IL-17A through a shared IL-17RA/IL-17RC receptor complex, though it can also act as a homodimer via IL-17RC alone, triggering NF-kB and MAPK signaling that drives production of neutrophil-recruiting chemokines, antimicrobial peptides, and matrix-degrading enzymes. Through this pathway it recruits and activates neutrophils to help clear extracellular bacteria and fungi at epithelial surfaces, and it further protects mucosal barriers by prompting epithelial cells to produce defensin peptides. Beyond direct antimicrobial defense, IL-17F has been reported to restrain new blood vessel growth in endothelial cells even as it stimulates their output of other inflammatory mediators, and dysregulated IL-17F/IL-17A activity is linked to chronic inflammatory conditions including psoriasis and rheumatoid arthritis.

PRINCIPLE OF THE ASSAY

This kit uses a sandwich enzyme immunoassay format in which the microtiter plate wells are pre-coated with a capture antibody specific to Human IL-17F. Standards or samples are added to the appropriate wells along with a biotin-conjugated detection antibody that also recognizes Human IL-17F. Avidin conjugated to horseradish peroxidase (HRP) is then added to each well and allowed to incubate, binding to the biotin on the detection antibody. Upon addition of TMB substrate solution, a colorimetric signal develops exclusively in wells where Human IL-17F, the biotin-conjugated antibody, and avidin-HRP are all present. The enzymatic reaction is stopped by the addition of a sulphuric acid solution, and the resulting color is measured spectrophotometrically at 450 nm (± 10 nm). Sample concentrations are determined by comparing the absorbance readings of each sample against a standard curve generated from the provided standards.

MATERIALS AND STORAGE

Store kit components at -20°C unless specified otherwise. DO NOT USE past kit expiration date. Some vials contain a small amount of reagents. Spin tubes on pulse setting prior to opening.

Each kit includes:	Units
Pre-Coated Microplate	12 strips x 8 wells
Standard (Lyophilized)	2 vials
Biotinylated Antibody (100×)	120 µL
Streptavidin-HRP (100×)	120 µL
Standard/Sample Diluent Buffer	20 mL
Biotinylated Antibody Diluent	12 mL
HRP Diluent	12 mL
Wash Buffer (25×)	20 mL
TMB Substrate Solution	10 mL
Stop Reagent	6 mL
Plate Covers	2 pieces
Do not mix or substitute reagents with those from other lots.	

MATERIALS AND INSTRUMENTS REQUIRED BUT NOT SUPPLIED

- Precision pipettes calibrated to deliver 5-1000µL
- Multi-channel pipette calibrated to deliver 50-200µL
- Plate shaker
- Disposable tips
- Vortex-Mixer
- Distilled or de-ionized water
- Microplate reader capable of reading 450nm with background subtraction at 620nm

SAFETY PRECAUTIONS

- The test protocol must be followed strictly.
- All reagents containing human material should be handled as if potentially infectious. Operators should wear gloves and protective

clothing when handling any patient sera or serum based products.

- The kit reagents contain antimicrobial agents, acid and 3,3’,5,5’-tetramethylbenzidine. Avoid contact with the skin and eyes. Rinse immediately with plenty of water if any contact occurs.
- Any liquid that has been brought into contact with potentially infectious material has to be discarded in a container with a disinfectant. Disposal must be performed in accordance with local regulations.
- Disposal must be performed in accordance with local regulations.
- Only trained laboratory personnel should execute this test.

This kit employs a sandwich enzyme immunoassay technique to quantify Human IL-17F in biological samples.

Human IL-17F ELISA Kit
Sandwich ELISA
Catalog EL-213-91217

PREPARATION OF REAGENTS

Prepare only the appropriate amount of required reagent on the day of use. Store all reagents as per instructions stated on the label.

1. **Wash Buffer (1X) Preparation: Dilute the 25X wash buffer concentrate 1:25 with double-distilled Water.**
2. **Detection Reagent (1X) Preparation: Dilute the biotinylated antibody and streptavidin-HRP concentrates 100-fold with Biotinylated Antibody Diluent and HRP Diluent, respectively, immediately before use.**
3. **Preparation of Calibrators: Prepare calibrators by serial dilution from a 500 down to 7.82 (plus a 0 blank), per the source manual.**

Sol'n ID	Source	Source Vol (µL)	Cal. Diluent (µL)	Final Vol (µL)	Final Conc. (ng/mL)
Std 1	Stock Standard	500	500	1000	500
Std 2	Std 1	500	500	1000	250
Std 3	Std 2	500	500	1000	125
Std 4	Std 3	500	500	1000	62.5
Std 5	Std 4	500	500	1000	31.25
Std 6	Std 5	500	500	1000	15.63
Std 7	Std 6	500	500	1000	7.82
Blank	Std 7	500	500	1000	0

SPECIMEN STORAGE

This kit is compatible with EDTA-plasma, heparin-plasma and serum samples. Samples can be stored at or below -20°C for up to 1 year.

ASSAY PROCEDURE

1. After the kit is equilibrated at room temperature, add 100 µL of Standard Working Buffer (gradually diluted according to the instructions) or 100 µL of sample to each well, and incubate at 37°C for 80 minutes.

2. Discard the liquid in the plate, add 200 µL 1× Wash Buffer to each well, and wash the plate 3 times. After pat it dry against clean absorbent paper, add 100 µL Biotinylated Antibody Working Solution (1×) to each well, incubate at 37°C for 50 minutes.

3. Discard the liquid in the plate, add 200 µL 1× Wash Buffer to each well, and wash the plate 3 times. After pat it dry against clean absorbent paper, add 100 µL 1× Streptavidin-HRP Working Solution to each well, incubate at 37°C for 50 minutes.

4. Discard the liquid in the plate, add 200 µL 1× Wash Buffer to each well, and wash the plate 5 times. After pat it dry against clean absorbent paper, add 90 µL TMB Substrate Solution to each well, incubate at 37°C for 20 minutes in the dark.

5. Add 50 µL Stop Solution to each well, shake plate on a plate shaker for 1 minute to mix. Record the OD at 450 nm immediately, calculation of the results.

CALCULATIONS AND RESULTS

1. Construct a standard curve by plotting the absorbance obtained from each standard against concentration. Use a 4 or 5 parameter curve fit. Alternatively a log-log curve fit may be used.
2. The concentration of the unknowns can be read directly from this standard curve using the absorbance value for each sample.

3. Any sample undiluted or diluted still reading greater than the highest standard should be diluted appropriately and retested. If the samples have been diluted, the concentration determined from the standard curve must be multiplied by the dilution factor.

PERFORMANCE CHARACTERISTICS

Precision: Intra-assay CV < 8%. Inter-assay CV < 10%.

Detection Limit: 2.8 pg/mL

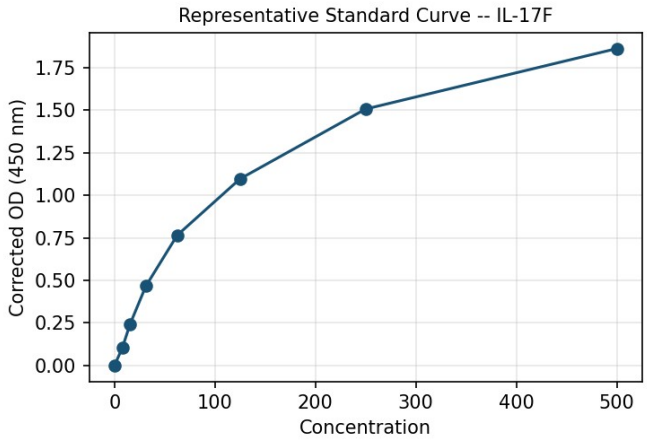
Recovery: Serum (n = 5): 81-97% (avg 89%); EDTA plasma (n = 5): 84-99% (avg 91%); Heparin plasma (n = 5): 79-91% (avg 85%)

Specificity: This assay has high sensitivity and excellent specificity for detection of Human IL17F. No significant cross-reactivity or interference between Human IL17F and analogues was observed.

This kit employs a sandwich enzyme immunoassay technique to quantify Human IL-17F in biological samples.

Human IL-17F ELISA Kit
Sandwich ELISA
Catalog EL-213-91217

SAMPLE STANDARD CURVE



Representative standard curve shown for illustration only. This curve should not be used to calculate unknown sample concentrations -- run a new standard curve with each assay.

ORDERING INFORMATION

Please visit *www.affinityimmuno.com* to order this product. Visa, Mastercard, AMEX and PayPal are accepted in our online store.

Your order will be processed immediately and you will be notified with a delivery timeframe.