

## Yersinia IgG ELISA

**Enzyme immunoassay** based on microtiter plate for the detection and quantitative determination of human IgG antibodies against **Yersinia enterocolitica** in serum and plasma

**Cat. No.: ILE-YER01**

Storage: 2-8°C  
For in-vitro diagnostic use only

### Symbols and Translations

Symbol	English	French	German	Italian	Spanish	Greek
<b>CAL</b>	Calibrator	Etalon	Kalibrator	Calibratore	Calibrador	Πρότυπο Διάλυμα
<b>CONJ</b>	Conjugate	Conjugué	Konjugat	Coniugato	Conjugado	Διάλυμα Συμπλόκου
<b>CONC</b>	Concentrate (<n>-fold)	Concentré (<n> fois)	Konzentrat (<n>-fach)	Concentrato (<n>-volte)	Concentrado (<n>-veces)	Συμπύκνωση (<n> φορές)
<b>SAMP DIL</b>	Sample Diluent	Diluant échantillon	Proben-verdüner	Diluente del campione	Diluyente de muestra	Διάλυμα Αραίωσης Δειγμάτων
<b>STOP</b>	Stop Solution	Solution d'arrêt	Stopp-Lösung	Soluzione d'arresto	Solución de parada	Διάλυμα Αναστολής
<b>SUBS</b>	Substrate	Substrat	Substrat	Substrato	Sustrato	Διάλυμα Υποστρώματος
<b>MT PLATE</b>	Microtiter plate	Microplaque	Mikrotiterplatte	Piastre	Placa microtiter	Μικρόπλακα
<b>WASH BUF</b>	Wash buffer	Tampon de lavage	Waschpuffer	Soluzione di lavaggio	Tampón de lavado	Πλύστικό Διάλυμα

### 1. Intended Use

The IMMUNOLAB Yersinia IgG Antibody ELISA Test Kit has been designed for the detection and the quantitative determination of specific IgG antibodies against Yersinia enterocolitica in serum and plasma. Further applications in other body fluids are possible and can be requested from the Technical Service of IMMUNOLAB. This assay is intended for in-

vitro diagnostic use only. Laboratory results can never be the only base of a medical report. The patient history and further tests have additionally to be taken into account.

## **2. General Information**

The pathogenic germs *Yersinia pestis*, *Y. pseudotuberculosis*, *Y. enterocolitica* and *Y. ruckeri* belong to the genus *Yersinia* as a member of the enterobacteriaceae family. All the species of medical importance occur facultatively inside the cells, which leads to the characteristic inflammation of the lymphatic tissue in the course of an illness. *Yersinia enterocolitica* is taken up orally, and the symptoms in a patient are terminal ileitis as well as diarrhoea. It is difficult to make a separation from appendicitis by differential diagnosis. In the course of a retarded immunological reaction, extraintestinal manifestations like erythema nodosum, uveitis, and arthritis can appear. It has been claimed that the background for a reactive arthritis caused by *Yersinia* consists in the local synthesis of antibodies in the joints (synovial fluid). *Yersinia* can be characterized without problems by standardized bacterial tests, after isolation in a pure culture. Following a human disease, above all the serotypes O3, O8 and O9 are found. Crossreactivities to *Brucella* are described and have to be taken into account by the differential diagnosis. The classical serological detection method has always been the Widal test. Further increasingly popular methods are the HAT, the KBR test as well as the ELISA. The enzyme immunoassay is characterized on the one hand side by a high sensitivity, but also by the possibility to differentiate between IgG and IgA/IgM antibodies. The immunoglobulin classes IgA and IgM should be referred to as a criterium of interpretation for an active process, when an arthritic disease is suspected. Because there exists also an immunological similarity with the thyrotropin receptor, the test can also be employed for the confirmation of the autoimmune Graves' disease. As a confirmatory test with the possibility of an identification of separate bacterial proteins as well as for the exclusion of cross-reactivities, the Western Blot method has proved successful.

## **3. Principle of the Test**

The IMMUNOLAB *Yersinia enterocolitica* IgG antibody test kit is based on the principle of the enzyme immunoassay (EIA). *Yersinia* antigen is bound on the surface of the microtiter strips. Diluted patient serum or ready-to-use standards are pipetted into the wells of the microtiter plate. A binding between the IgG antibodies of the serum and the immobilized *Yersinia* antigen takes place. After a one hour incubation at room temperature, the plate is rinsed with diluted wash solution, in order to remove unbound material. Then ready-to-use anti-human-IgG peroxidase conjugate is added and incubated for 30 minutes. After a further washing step, the substrate (TMB) solution is pipetted and incubated for 20 minutes, inducing the development of a blue dye in the wells. The color development is terminated by the addition of a stop solution, which changes the color from blue to yellow. The resulting dye is measured spectrophotometrically at the wavelength of 450 nm. The concentration of the IgG antibodies is directly proportional to the intensity of the color.

## **4. Limitations, Precautions and General Comments**

- Only for in-vitro use! Do not ingest or swallow! The usual laboratory safety precautions as well as the prohibition of eating, drinking and smoking in the lab have to be followed.

- All sera and plasma or buffers based upon, have been tested respective to HBsAg, HIV and HCV with recognized methods and were found negative. Nevertheless precautions like the use of latex gloves have to be taken.
- Serum and reagent spills have to be wiped off with a disinfecting solution (e.g. sodium hypochlorite, 5%) and have to be disposed of properly.
- All reagents have to be brought to room temperature (18 to 25 °C) before performing the test.
- Before pipetting all reagents should be mixed thoroughly by gentle tilting or swinging. Vigorous shaking with formation of foam should be avoided.
- It is important to pipet with constant intervals, so that all the wells of the microtiter plate have the same conditions.
- When removing reagents out of the bottles, care has to be taken that the stoppers are not contaminated. Further a possible mix-up has to be avoided. The content of the bottles is usually sensitive to oxidation, so that they should be opened only for a short time.
- In order to avoid a carry-over or a cross-contamination, separate disposable pipet tips have to be used.
- No reagents from different kit lots have to be used, they should not be mixed among one another.
- All reagents have to be used within the expiry period.
- In accordance with a Good Laboratory Practice (GLP) or following ISO9001 all laboratory devices employed should be regularly checked regarding the accuracy and precision. This refers amongst others to microliter pipets and washing or reading (ELISA-Reader) instrumentation.
- The contact of certain reagents, above all the stopping solution and the substrate with skin, eye and mucosa has to be avoided, because possible irritations and acid burns could arise, and there exists a danger of intoxication.

## 5. Reagents Provided

Store kit components at 2-8°C and do not use after the expiry date on the box outer label. Before use, all components should be allowed to warm up to ambient temperature (18-25°C). After use, the plate should be resealed, the bottle caps replaced and tightened and the kit stored at 2-8°C. The opened kit should be used within three months.

Components	Volume / Qty.
Yersinia antigen coated microtiter strips	12
Calibrator A (Negative Control)	2 mL
Calibrator B (Cut-Off Standard)	2 mL
Calibrator C (Weak Positive Control)	2 mL
Calibrator D (Positive Control)	2 mL
Enzyme Conjugate	15 mL
Substrate	15 mL
Stop Solution	15 mL
Sample Diluent	60 mL
Washing Buffer (10×)	60 mL
Plastic foils	2
Plastic bag	1

### **5.1. Microtiter Strips**

12 strips with 8 breakable wells each, coated with a *Yersinia enterocolitica* antigen (*Yersinia* outer proteins (YOPs), purified). Ready-to-use.

### **5.2. Calibrator A (Negative Control)**

2 mL, protein solution diluted with PBS, contains no IgG antibodies against *Yersinia*. Addition of 0.01 % methylisothiazolone and 0.01 % bromonitrodioxane. Ready-to-use.

### **5.3. Calibrator B (Cut-Off Standard)**

2 mL human serum diluted with PBS, contains a low concentration of IgG antibodies against *Yersinia*. Addition of 0.01 % methylisothiazolone and 0.01 % bromonitrodioxane. Ready-to-use.

### **5.4. Calibrator C (Weak Positive Control)**

2 mL, human serum diluted with PBS, contains a medium concentration of IgG antibodies against *Yersinia*. Addition of 0.01 % methylisothiazolone and 0.01 % bromonitrodioxane. Ready-to-use.

### **5.5. Calibrator D (Positive Control)**

2 mL, human serum diluted with PBS, contains a high concentration of IgG antibodies against *Yersinia*. Addition of 0.01 % methylisothiazolone and 0.01 % bromonitrodioxane. Ready-to-use.

### **5.6. Enzyme Conjugate**

15 mL, anti-human-IgG-HRP (rabbit), in protein-containing buffer solution. Addition of 0.01 % methylisothiazolone and 0.01 % bromonitrodioxane and 5 mg/L Proclin™. Ready-to-use.

### **5.7. Substrate**

15 mL, TMB (tetramethylbenzidine). Ready-to-use.

### **5.8. Stop Solution**

15 mL, 0.5 M sulfuric acid. Ready-to-use.

### **5.9. Sample Diluent**

60 mL, PBS/BSA buffer. Addition of 0.095 % sodium azide. Ready-to-use.

### **5.10. Washing Buffer**

60 mL, PBS + Tween 20, 10x concentrate. Final concentration: dilute 1+9 with distilled water. If during the cold storage crystals precipitate, the concentrate should be warmed up at 37°C for 15 minutes.

### **5.11. Plastic Foils**

2 pieces to cover the microtiter strips during the incubation.

### **5.12. Plastic Bag**

Resealable, for the dry storage of non-used strips.

## 6. Materials Required but not Provided

- 5 µL-, 100 µL- and 500 µL micro- and multichannel pipets
- Microtiter Plate Reader (450 nm)
- Microtiter Plate Washer
- Reagent tubes for the serum dilution
- Bidistilled water

## 7. Specimen Collection and Handling

Principally serum or plasma (EDTA, heparin) can be used for the determination. Serum is separated from the blood, which is aseptically drawn by venipuncture, after clotting and centrifugation. The serum or plasma samples can be stored refrigerated (2-8°C) for up to 48 hours, for a longer storage they should be kept at -20 °C. The samples should not be frozen and thawed repeatedly. Lipemic, hemolytic or bacterially contaminated samples can cause false positive or false negative results. For the performance of the test the samples (not the standards) have to be diluted 1:101 with ready-to-use sample diluent (e.g. 5 µL serum + 500 µL sample diluent).

## 8. Assay Procedure

### 8.1. Preparation of Reagents

**Washing Solution:** dilute before use 1+9 with distilled water. If during the cold storage crystals precipitate, the concentrate should be warmed up at 37°C for 15 minutes.

- Strict adherence to the protocol is advised for reliable performance. Any changes or modifications are the responsibility of the user.
- All reagents and samples must be brought to room temperature before use, but should not be left at this temperature longer than necessary.
- Standards and samples should be assayed in duplicates.
- A standard curve should be established with each assay.
- Return the unused microtiter strips to the plastic bag and store them dry at 2-8°C.

### 8.2. Assay Steps

1. Prepare a sufficient amount of microtiter wells for the standards, controls and samples in duplicate as well as for a substrate blank.
2. Pipet 100 µL each of the **diluted** (1:101) samples and the **ready-to-use** standards and controls respectively into the wells. Leave one well empty for the substrate blank.
3. Cover plate with the enclosed foil and incubate at room temperature for 60 minutes.
4. Empty the wells of the plate (dump or aspirate) and add 300 µL of diluted washing solution. This procedure is repeated totally three times. Rests of the washing buffer are afterwards removed by gentle tapping of the microtiter plate on a tissue cloth.
5. Pipet 100 µL each of ready-to-use conjugate into the wells. Leave one well empty for the substrate blank.
6. Cover plate with the enclosed foil and incubate at room temperature for 30 minutes.
7. Empty the wells of the plate (dump or aspirate) and add 300 µL of diluted washing solution. This procedure is repeated totally three times. Rests of the washing buffer are afterwards removed by gentle tapping of the microtiter plate on a tissue cloth.

8. Pipet 100  $\mu$ L each of the ready-to-use substrate into the wells. This time also the substrate blank is pipetted.
9. Cover plate with the enclosed foil and incubate at room temperature for 20 minutes in the dark (e.g. drawer).
10. To terminate the substrate reaction, pipet 100  $\mu$ L each of the ready-to-use stop solution into them wells. Pipet also the substrate blank.
11. After thorough mixing and wiping the bottom of the plate, perform the reading of the absorption at 450 nm (optionally reference wavelength of 620 nm). The color is stable for at least 60 minutes.

## 9. Evaluation

The mean values for the measured absorptions are calculated after subtraction of the substrate blank value. The difference between the single values should not exceed 10%.

### Example

	OD Value	corrected OD	Mean OD Value
Substrate Blank	0.018		
Negative Control	0.030 / 0.026	0.012 / 0.008	0.010
Cut-Off Standard	0.583 / 0.611	0.565 / 0.593	0.579
Weak Positive Control	0.856 / 0.879	0.838 / 0.861	0.850
Positive Control	1.961 / 1.921	1.943 / 1.903	1.923

The above table contains only an example, which was achieved under arbitrary temperature and environmental conditions. The described data constitute consequently **no reference values** which have to be found in other laboratories in the same way.

### 9.1. Qualitative Evaluation

The calculated absorptions for the patient sera, as mentioned above, are compared with the value for the cut-off standard. If the value of the sample is higher, there is a positive result. For a value below the cut-off standard, there is a negative result. It seems reasonable to define a range of +/-20 % around the value of the cut-off as a grey zone. In such a case the repetition of the test with the same serum or with a new sample of the same patient, taken after 2-4 weeks, is recommended. Both samples should be measured in parallel in the same run. The positive control must show at least the double absorption compared with the cut-off standard.

### 9.2. Quantitative Evaluation

The ready-to-use standards and controls of the Yersinia antibody kit are defined and expressed in arbitrary units (U/mL). This results in an exact and reproducible quantitative evaluation. Consequently for a given patient follow-up controls become possible. The values for controls and standards in units are printed on the labels of the vials. For a quantitative evaluation the absorptions of the standards and controls are graphically drawn against their concentrations. From the resulting reference curve the concentration values for each patient sample can then be extracted in relation to their absorptions. It is also possible to use automatic computer programs.



## 10. Assay Characteristics

Yersinia ELISA	IgG	IgA	IgM
Intra-Assay-Precision	5.0 %	10.3 %	10.3 %
Inter-Assay-Precision	5.2 – 11.9 %	5.3 – 10.0 %	5.3 – 10.0 %
Inter-Lot-Precision	3.9 – 12.8 %	5.2 – 13.0 %	5.2 – 13.0 %
Analytical Sensitivity	1.92 U/mL	1.02 U/mL	1.02 U/mL
Recovery	87 – 100 %	88 – 107 %	74 – 98 %
Linearity	70 – 113 %	65 – 116 %	66 – 114 %
Cross-Reactivity	No cross-reactivity to <i>Helicobacter pylori</i>		
Interferences	No interferences to bilirubin up to 0.3 mg/mL, hemoglobin up to 8.0 mg/mL and triglycerides up to 5.0 mg/mL		
Clinical Specificity	92 %	92 %	92 %
Clinical Sensitivity	100 %	100 %	100 %

GENTIA