






# Helicobacter Pylori Antibody IgG GENLISA™ ELISA

**REF** : KBD931

Ver 2.0


**IVD**

Enzyme Immunoassay for the Quantitative Determination of  
Helicobacter Pylori Antibody IgG in human serum and plasma.

<b>IVD</b>	For In-vitro Diagnostic Use	<b>REF</b>	Catalog Number
	Store At	<b>LOT</b>	Batch Code
	Manufactured By		Biological Risk
	Expiry Date		Consult Operating Instructions

*For In-vitro Diagnostic Use only. Purchase does not include or carry the right to resell or transfer this product either as a stand-alone product or as a component of another product. Any use of this product other than the permitted use without the express written authorization of KRISHGEN Pudgala LLP is strictly prohibited.*

**REF** KBD931

 96 tests

**Krishgen Pudgala LLP** Unit Nos#318/319, Shah & Nahar, Off Dr E Moses Road, Worli, Mumbai 400018. India.  
Tel: +91-22-49198700 | email: [sales@krishgenpudgala.com](mailto:sales@krishgenpudgala.com)

**Introduction:**

Helicobacter pylori (H.pylori) is a gram negative bacterium which is located at gastric mucosa. The organism is present in 95-98% of patients with duodenal ulcer and 60-90% of patients with gastric ulcers. Estimation of infection rate by various diagnostic tests including bacteriological, histo-logical and serological tests have revealed that 90% of symptomatic patients are affected and 50% of old age adults (> 50 years) is only colonized by bacteria lifelong without any clinical symptoms. Studies also have been demonstrated that removal of the organism by antimicrobial therapy is correlated with the resolution of symptoms and cure of diseases. Patients who present clinical symptoms relating to the gastrointestinal tract can be diagnosed for H. pylori infection by two methods:

- 1) Invasive techniques include biopsy followed by culture or histological examination of biopsy specimen or direct detection of urease activity.
- 2) Non-invasive techniques include urea breath tests and serological methods.

All of the tests performed on biopsy samples are subject to errors related to sampling and interference of bacterial contamination. Helicobacter pylori infection stimulates humoral immune response and provokes specific antibodies like IgG, IgM, and IgA. H. pylori IgG ELISA test is an accurate and simple technique to determine colonization of bacteria and ELISA test is technique of choice for detection of IgG response.

**Intended Use:**

The Helicobacter Pylori Antibody IgG GENLISA™ ELISA is intended for the quantitative determination of Helicobacter Pylori Antibody IgG in human serum and plasma.

**Principle:**

The Helicobacter Pylori Antibody IgG GENLISA™ ELISA is an indirect enzyme linked immunosorbent assay for quantitative determination of IgG antibody present in the human serum and plasma. The Microtiter wells are pre-coated with specific H.pylori antigens. Samples and Controls are pipetted into microwells and Helicobacter Pylori Antibody IgG present in sample binds to the antigen coated on the wells. Anti-human-IgG:Enzyme Conjugate antibody is pipetted and incubated to form an immune complex. After washing microwells in order to remove any non-specific binding, the substrate solution is added to microwells and color develops proportionally to the amount of The Helicobacter Pylori Antibody IgG present in the sample. Color development is then stopped by addition of stop solution. Absorbance is measured at 450 nm.

**Materials Provided:**

1. H.pylori antigen Microtiter Coated Plate (8x12 wells) - 1 no
2. Standards (1.5 ml/vial) - 0, 5, 10, 20, 50, 100 U/ml
3. Sample diluent – 2 X 50 ml
4. Low Control Serum – 1.5 ml/vial
5. High Control Serum – 1.5 ml/vial
6. Anti-human-IgG:Enzyme Conjugate - 12 ml
7. (20X) Wash Buffer – 50 ml
8. TMB Substrate - 12 ml
9. Stop Solution - 12 ml
10. Instruction Manual

**Materials to be provided by the End-User:**

1. Microtiter Plate Reader able to measure absorbance at 450 nm.
2. Adjustable pipettes and multichannel pipette to measure volumes ranging from 25 ul to 1000 ul
3. Deionized (DI) water
4. Wash bottle or automated microplate washer
5. Graph paper or software for data analysis
6. Timer
7. Absorbent Paper

**Handling/Storage:**

1. Store main kit components at recommended storage temperature indicated on the component label.
2. Before using, bring all components to room temperature (18-25°C). Upon assay completion return all components to appropriate storage conditions.
3. The Substrate is light-sensitive and should be protected from direct sunlight or UV sources.

**Health Hazard Warnings:**

1. Reagents that contain preservatives may be harmful if ingested, inhaled or absorbed through the skin. Refer to the MSDS online for details.
2. To reduce the likelihood of blood-borne transmission of infectious agents, handle all serum and/or plasma in accordance with NCCLS regulations.

**Specimen Collection and Handling:**

**Serum-** Coagulate at room temperature for 10-20 minutes; centrifuge for 20-min at 2000-3000 rpm. Remove the supernatant. If precipitation appears, re-centrifuge.

**Plasma-** Use EDTA or citrate plasma as an anticoagulant, mix for 10-20 minutes; centrifuge for 15-min at 2000-3000 rpm. Remove the supernatant carefully. If precipitation appears, re-centrifuge.

**Sample Dilution:**

To make 1:51 Dilutions, dilute 10 ul Sample + 500ul Sample Diluent. Mix well.

**Reagent Preparation:**

1. Allow all components to reach RT (Room Temperature) prior to use in the assay.
2. Wash Buffer (1X) Dilution: To make Wash Buffer (1X), add 50ml of Wash Buffer (20X) to 450ml of DI water. This is the working solution.

**Test Procedure:**

1. All reagents should be allowed to reach room temperature before use.
2. Add **100 ul Standard, Controls, Diluted Sample** in appropriate wells.
3. Seal the plate and Incubate at room temperature for 30 minutes.
4. Aspirate and wash plate 5 times with **(1x) Wash Buffer** and blot residual buffer by firmly tapping plate upside down on absorbent paper. Wipe of any liquid from the bottom outside of the microtiter wells as any residue can interfere in the reading step. All the washes should be performed similarly.
5. Add **100 ul of Anti-human-IgG:Enzyme Conjugate** to each well except blank well.
6. Incubate at room temperature for 30 minutes.
7. Repeat the Wash Step as mentioned in step 4.
8. Add **100 ul of TMB Substrate** into each well.
9. Incubate at room temperature for 15 minutes.
10. Add **100 ul of Stop Solution**. Read result with an ELISA reader at 450 nm within 15 minutes of stopping the reaction.

**Interpretation of Results:**

Determine the Mean Absorbance for each set of duplicate or triplicate Standards and Samples. Using graph paper, plot the average value (absorbance 450 nm) of each standard on the Y-axis versus the corresponding concentration of the standards on the X-axis. Draw the best fit curve through the standard points. To determine the unknown Helicobacter Pylori Antibody IgG concentrations, find the unknown's Mean Absorbance value on the Y-axis and draw a horizontal line to the standard curve. At the point of intersection, draw a vertical line to the X-axis and read the Helicobacter Pylori Antibody IgG Concentration. If samples were diluted, multiply by the appropriate dilution factor.

Software which is able to generate a cubic spline curve-fit, 4-PL or a polynomial curve (2nd order) is best recommended for automated results.

**Note:**

- If the sample reads more than 100 U/ml, then dilute with Standard 0. The obtained result should be multiplied by the dilution factor.

**Performance Characteristics:****Sensitivity**

The analytical sensitivity of **Helicobacter Pylori Antibody IgG** kit was calculated and determined to be **97%**

**Specificity**

The analytical specificity of **Helicobacter Pylori Antibody IgG** kit was calculated and determined to be **95%**

**Precision:****Intra-Assay precision:**

The within assay variability is shown below:

No.	No. of Tests Performed	Means U/ml	SD U/ml	CV%
1	24	6.5	0.3	4.6
2	24	18	0.9	5.0
3	24	45	2.0	4.4

**Inter-Assay precision:**

The between assay variability is shown below:

No.	No. of Tests Performed	Means U/ml	SD U/ml	CV%
1	10	8.7	0.71	8.1
2	10	18.8	1.76	9.4
3	10	43.5	2.90	6.7

\*Each test has been run in duplicate

**Safety Precautions:**

- **This kit is For In-vitro Diagnostic Use only.** Follow the working instructions carefully.
- The expiration dates stated on the kit are to be observed. The same relates to the stability stated for reagents
- Do not use or mix reagents from different lots.
- Do not use reagents from other manufacturers.
- Avoid time shift during pipetting of reagents.
- All reagents should be kept in the original shipping container.
- Some of the reagents contain small amount of sodium azide (< 0.1 % w/w) as preservative. They must not be swallowed or allowed to come into contact with skin or mucosa.



- Source materials maybe derived from human body fluids or organs used in the preparation of this kit were tested and found negative for HBsAg and HIV as well as for HCV antibodies. However, no known test guarantees the absence of such viral agents. Therefore, handle all components and all patient samples as if potentially hazardous.
- Since the kit contains potentially hazardous materials, the following precautions should be observed
  - Do not smoke, eat or drink while handling kit material
  - Always use protective gloves
  - Never pipette material by mouth
  - Wipe up spills promptly, washing the affected surface thoroughly with a decontaminant.
- In any case GLP should be applied with all general and individual regulations to the use of this kit.

## LIMITED WARRANTY

Krishgen Pudgala LLP does not warrant against damages or defects arising in shipping or handling, or out of accident or improper or abnormal use of the product; against defects in products or components not manufactured by Krishgen Pudgala LLP, or against damages resulting from such non-Krishgen Pudgala LLP made products or components. Krishgen Pudgala LLP passes on to customer the warranty it received (if any) from the maker thereof of such non-Krishgen made products or components. This warranty also does not apply to product to which changes or modifications have been made or attempted by persons other than pursuant to written authorization by Krishgen Pudgala LLP.

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Krishgen Pudgala LLP, 2023.

## THANK YOU FOR USING KRISHGEN PRODUCT!



**Krishgen Pudgala LLP**

Unit No.1/2, Om Sainath Commercial Complex,  
Off Mankoli-Anjur Phata Road. Village Dapode, Bhiwandi 421302.



**KinesisDx, Lyoner Strasse 14, Frankfurt, Germany**

### Regulatory Status:






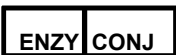







CE Marked	Europe
FDA registered	USA
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\* Under CDSCO Registration, please note

## SCHEMATIC ASSAY PROCEDURE

1	All reagents should be allowed to reach room temperature before use.
2	Add <b>100 ul Control, Diluted Sample</b> in appropriate wells.
3	Seal the plate and <b>Incubate at 37°C for 30 minutes.</b>
4	Aspirate and wash plate 5 times with (1x) Wash Buffer and blot residual buffer by firmly tapping plate upside down on absorbent paper. Wipe of any liquid from the bottom outside of the microtiter wells as any residue can interfere in the reading step. All the washes should be performed similarly.
5	Add <b>100 ul of Anti-human-IgG:Enzyme Conjugate</b> to each well except blank well.
6	<b>Incubate at 37°C for 30 minutes.</b>
7	Repeat the Wash Step as mentioned in step 4.
8	Add <b>100 ul of TMB Substrate</b> into each well.
9	<b>Incubate at 37°C for 10 minutes.</b>
10	Add <b>100 ul of Stop Solution.</b> Read result with an ELISA reader at 450 nm within 15 minutes stopping the reaction.

## SYMBOLS KEY

	Microtiter Plate (8x12 wells)
	Standards
	Sample Diluent
	Low Control Serum
	High Control Serum
	Ani-human-IgG:Enzyme Conjugate
	(20X) Wash Buffer
	TMB Substrate
	Stop Solution
	Consult Instructions for Use
	Catalog Number
	Expiration Date
	Storage Temperature