

KRIBIOLISA® Anti-Panitumumab (VECTIBIX™) ELISA

REF : KBI2082

Ver 1.2

RUO

Enzyme Immunoassay for the Quantitative Determination of Antibodies to Panitumumab in human serum and plasma

RUO	For Research Use Only	REF	Catalog Number
	Store At	LOT	Batch Code
	Manufactured By		Biological Risk
	Expiry Date		Consult Operating Instructions

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REF KBI2082  96 tests

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Introduction:

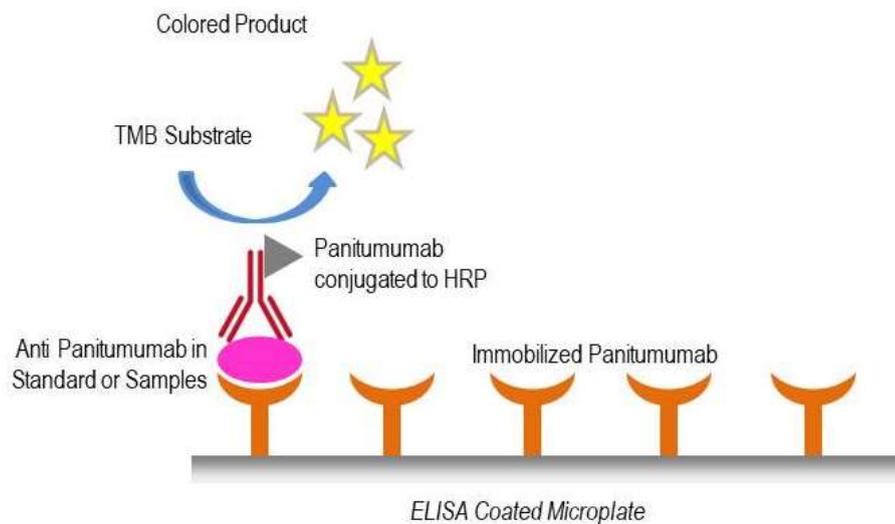
Panitumumab is a recombinant humanized IgG1 monoclonal antibody directed against the human lymphocyte $\alpha 4\beta 7$ integrin, a key mediator of gastrointestinal inflammation. It is used in the treatment of moderate to severe active ulcerative colitis and Crohn's disease for patients who have had an inadequate response with, lost response to, or were intolerant to inhibitors of tumor necrosis factor-alpha (TNF-alpha) or other conventional therapies. By blocking its primary target, $\alpha 4\beta 7$ integrin, Panitumumab reduces inflammation in the gut.

Intended Use:

The KRIBIOLISA® Anti-Panitumumab (VECTIBIX™) ELISA is used as an analytical tool for quantitative determination of antibodies to Panitumumab (VECTIBIX™) in human serum and plasma.

Principle:

The method employs the quantitative sandwich enzyme immunoassay technique. Panitumumab is pre-coated onto microwells. Samples and standards are pipetted into microwells and antibodies to human Panitumumab present in the sample are bound by the capture antibody. Then, a HRP (horseradish peroxidase) conjugated Panitumumab is pipetted and incubated. After washing microwells in order to remove any non-specific binding, the ready to use substrate solution 3,3', 5,5' Tetra Methyl Benzidine (TMB) is added to microwells and color develops proportionally to the amount of Anti-Panitumumab in the sample. Color development is then stopped by addition of stop solution. Absorbance is measured at 450 nm.



Materials Provided:

Part	Description	Qty
Panitumumab Coated Microtiter Plate	96 well polystyrene microplate (12 strips of 8 wells) coated with Panitumumab.	1 x 96 wells
Anti- Panitumumab Standard	Recombinant Anti- Panitumumab in a buffered protein base and preservatives 0.02% Methylisothiazolinone and 0.02% bromonitrodioxane – (lyophilized, concentrated 1 ug/ml)	2 vials
Panitumumab:HRP conjugate Concentrated	Panitumumab conjugated to HRP concentrated (1 mg/ml)	2 vials
Detection Diluent	Buffered protein base with protein stabilizer and preservatives 0.02% Methylisothiazolinone and 0.02% bromonitrodioxane	12 ml
(1X) Standard Diluent	Buffered protein base with protein stabilizer and 1:1000 human serum and preservatives 0.02% Methylisothiazolinone and 0.02% bromonitrodioxane	10 ml

Part	Description	Qty
(1X) Sample Diluent	Buffered protein base with protein stabilizer and preservatives 0.02% Methylisothiazolinone and 0.02% bromonitrodioxane	2 x 50 ml
(20X) Wash Buffer	20-fold concentrated solution of buffered surfactant with preservative thiomersol < 0.01%. May turn yellow over time.	25 ml
TMB Substrate	Stabilized Chromogen	12 ml
Stop Solution	0.73M Phosphoric Acid	12 ml
Instruction Manual		1 no

Materials to be provided by the End-User:

1. Microtiter Plate Reader able to measure absorbance at 450 nm.
2. Adjustable pipettes and multichannel pipettor 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. It is advisable to aliquot and stores the Panitumumab:HRP Conjugate concentrated at -20°C upon receipt. Rest of the kit components should be stored at 2-8°C. Immediately discard any excess Working Panitumumab:HRP Conjugate after running your assay.
2. All the reagents and wash solutions should be used within 12 months from manufacturing date.
3. Before using, bring all components to room temperature (18-25°C). Upon assay completion ensure all components of the kit are returned to appropriate storage conditions.
4. 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.
2. For Research Use Only.



Sample Preparation and Storage:

Blood is taken by venipuncture. Serum is separated after clotting by centrifugation. Plasma can be used, too. Lipaemic, hemolytic or contaminated samples should not be run. Repeated freezing and thawing should be avoided. If samples are to be used for several assays, initially aliquot samples and keep at - 20°C.

Preparation Before Use:

Allow samples to reach room temperature prior to assay. Take care to agitate patient samples gently in order to ensure homogeneity.

Test Sample preparation:

- Serum Samples have to be diluted 1:1000 (v/v), e.g. for 1:1000 (1 ul sample + 999 ul sample diluent) prior to assay. The samples may be kept at 2 - 8°C for up to three days. Long-term storage requires -20°C.
- Plasma Samples have to be diluted 1:1000 (v/v), e.g. for 1:1000 (1 ul sample + 999 ul sample diluent) prior to assay. The samples may be kept at 2 - 8°C for up to three days. Long-term storage requires -20°C.

Reagent Preparation (all reagents should be diluted immediately prior to use):

1. Label any aliquots made with the kit Lot No and Expiration date and store it at appropriate conditions mentioned.
2. Bring all reagents to Room temperature before use.

- To make Wash Buffer (1X); dilute 25 ml of 20X Wash Buffer in 475 ml of DI water.
- Standard Preparation:** Reconstitute the lyophilized standard in 1ml of Standard diluent (1X) to get a concentration of 1000ng/ml. Keep the vial for 15 mins with gentle agitation before making further dilutions. Dilute 640 ul of Reconstituted **Standard (1000 ng/ml)** with 360 ul of Standard Diluent to generate a **640 ng/ml Standard Solution**. Prepare further **Standards** by serially diluting the Standard Solution as per the below table. Use the Standard Diluent as the Zero Standard (Standard No.0)

Standard Concentration	Standard Vial	Dilution Particulars
1000 ng/ml	Lyophilized Standard	Lyophilized Standard provided in the Kit + 1 ml of Standard Diluent
640 ng/ml	Standard No.7	640 ul Reconstituted Standard (1 ug/ml) + 360 ul Standard Diluent
320 ng/ml	Standard No.6	500 ul Standard No.7 + 500 ul Standard Diluent
160 ng/ml	Standard No.5	500 ul Standard No.6 + 500 ul Standard Diluent
80 ng/ml	Standard No.4	500 ul Standard No.5 + 500 ul Standard Diluent
40 ng/ml	Standard No.3	500 ul Standard No.4 + 500 ul Standard Diluent
20 ng/ml	Standard No.2	500 ul Standard No.3 + 500 ul Standard Diluent
10 ng/ml	Standard No.1	500 ul Standard No.2 + 500 ul Standard Diluent
0 ng/ml	Standard No.0	Only Standard Diluent

Use the Standards as soon as possible upon reconstitution. Discard balance standard after use.

5. Working Panitumumab:HRP Conjugate – Refer to the Reagent Preparation sheet attached with the IFU and COA (enclosed in the kit).

Procedural Notes:

- In order to achieve good assay reproducibility and sensitivity, proper washing of the plates to remove excess un-reacted reagents is essential.
- High Dose Hook Effect may be observed in samples with very high concentrations of Anti-Panitumumab. High Dose Hook Effect is due to excess of antibody for very high concentrations of Anti-Panitumumab present in the sample. High Dose Hook effect is most likely encountered from samples early in the purification process. If Hook Effect is possible, the samples to be assayed should be diluted with a compatible diluent. Thus if the Anti-Panitumumab concentration of the undiluted sample is less than the diluted sample, this may be indicative of the Hook Effect.
- Avoid assay of Samples containing sodium azide (NaN₃), as it could destroy the HRP activity resulting in under-estimation of the amount of Anti-Panitumumab.
- It is recommended that all Standards and Samples be assayed in duplicates.
- Maintain a repetitive timing sequence from well to well for all the steps to ensure that the incubation timings are same for each well.
- If the Substrate has a distinct blue color prior to use it may have been contaminated and use of such substrate can lead to compromisation of the sensitivity of the assay.
- The plates should be read within 30 minutes after adding the Stop Solution.
- Make a work list in order to identify the location of Standards and Samples.

Assay Procedure:

- It is strongly recommended that all Standards and Samples be run in duplicates or triplicates. A standard curve is required for each assay. All steps must be performed at 37°C.
- Add **100 ul of prepared Standards or diluted Samples** into the respective wells.
- Cover the plate and incubate for 60 minutes at 37°C.
- Aspirate and wash plate 4 times with **Wash Buffer (1X)** 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.
- Add **100 ul of Working Panitumumab:HRP Conjugate** into each well.

KRIBIOLISA® Anti-Panitumumab (VECTIBIX™) ELISA

6. Cover the plate and incubate for 60 minutes at 37°C
7. Aspirate and wash plate 4 times with **Wash Buffer (1X)** same as in step 4.
8. Add **100 ul** of **TMB Substrate** in each well.
9. Incubate the plate at 37°C for 30 minutes in dark. **DO NOT SHAKE** or else it may result in higher backgrounds and worse precision. Positive wells should turn bluish in color.
10. Pipette out **100 ul** of **Stop Solution**. Wells should turn from blue to yellow in color.
11. Read the absorbance at 450 nm with a microplate within 10-15 minutes after addition of Stop solution.

Calculation of Results:

Determine the Mean Absorbance for each set of duplicate or triplicate Standards and Samples. Using Semi-Log 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 Anti-Panitumumab 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 Anti-Panitumumab Concentration. If samples were diluted, multiply by the appropriate dilution factor. Software which is able to generate a cubic spline curve-fit or 4PL is best recommended for automated results.

Note:

It is recommended to repeat the assay at a different dilution factor in the following cases:

- If the sample absorbance value is below the first standard.
- If the absorbance value is equivalent or higher than the 640 ng/ml standard.

Quality Control:

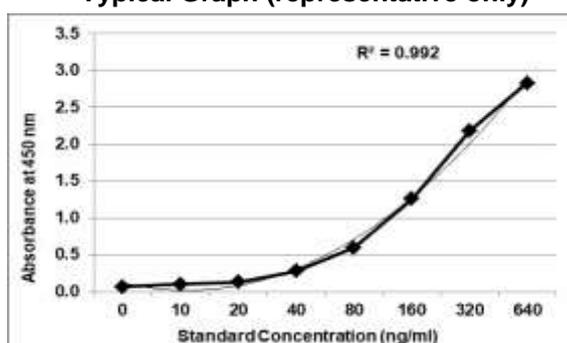
It is recommended that for each laboratory assay appropriate quality control samples in each run to be used to ensure that all reagents and procedures are correct.

Typical Data (representative only)

(for reference only, please refer to the CoA for lot specific data)

Standards (ng/ml)	Abs A	Abs B	Mean Abs	Interpolated Concentration	% Interpolated Concentration against Actual Concentration
0	0.073	0.065	0.069	---	---
10	0.099	0.109	0.104	11.9	118.8
20	0.146	0.111	0.128	17.7	88.4
40	0.298	0.261	0.279	42.3	105.8
80	0.666	0.523	0.595	80.4	100.5
160	1.397	1.125	1.261	158.0	98.7
320	2.365	2.009	2.187	323.6	101.1
640	2.818	2.844	2.831	635.3	99.3

Typical Graph (representative only)



Abs = absorbance at 450nm

Performance Characteristics of the Kit:

This kit has been validated as per EMA/FDA guidelines in line with ICH Code for Harmonization of Biological Assays.

Specificity:

The standard / calibrator antibodies used in the kit are monoclonal antibodies, anti-idiotypic and specific for Anti-Panitumumab.

Sensitivity:

Limit Of Detection: It is defined as the lowest detectable concentration corresponding to a signal of Mean of '0' standard plus 2* SD.

10 replicates of '0' standards were evaluated and the LOD was found to be less than 8 ng/ml.

Linearity:

The kit has been validated for linearity of Anti- Panitumumab in the claimed matrix.

Precision:

Precision is defined as the percent coefficient of variation (%CV) i.e. standard deviation divided by the mean and multiplied by 100. Assay precision was determined by both intra (n=5 assays) and inter assay (n=5 assays) reproducibility on two pools with low (10ng/ml), medium (160ng/ml) and high (640ng/ml) concentrations. While actual precision may vary from laboratory to laboratory and technician to technician, it is recommended that all operators achieve precision below these design goals before reporting results.

Pool	Intra Assay %CV	Inter Assay %CV
Low	<10%	<10%
Medium	<5%	<5%
High	<5%	<5%

Safety Precautions:

- **This kit is For Research 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/v) 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.

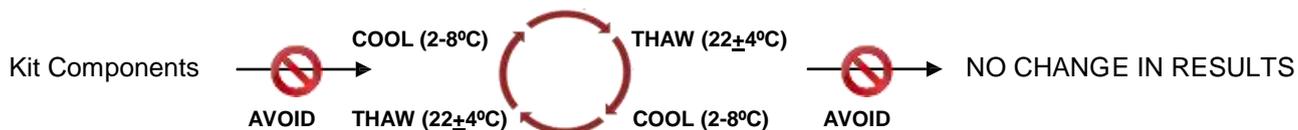


SCHMATIC ASSAY PROCEDURE

1. Remove all components, 30 minutes before adding into the assay plate.



2. Avoid repeated cool-thaw of the components as there will be a loss of activity and this can affect the results.



3. Pipette **100 ul prepared Standards / diluted Samples** into the respective wells.

4. Cover plate and incubate for **60 min** at 37°C.

5. Aspirate and wash wells 4 times with **Wash Buffer (1X)**

6. Pipette **100 ul Working Panitumumab:HRP** into each well.

7. Cover plate and incubate for **60 min** at 37°C

8. Aspirate and wash wells 4 times with **Wash Buffer (1X)**

9. Pipette **100 ul TMB Substrate** into each well.

10. Cover plate and incubate for **30 min** at 37°C.

11. Pipette **100 ul Stop Solution** into each well.

12. Read absorbance at 450 nm with a microplate reader within **15 min** of stopping reaction.

Typical Example of a Work List

Well #	Contents	Absorbance at 450nm	Mean Absorbance	ng/ml Anti-Panitumumab (VECTIBIX™) equivalent
1A	zero std			
2A	zero std			
1B	10 ng/ml			
2B	10 ng/ml			
1C	20 ng/ml			
2C	20 ng/ml			
1D	40 ng/ml			
2D	40 ng/ml			
1E	80 ng/ml			
2E	80 ng/ml			
1F	160 ng/ml			
2F	160 ng/ml			
1G	320 ng/ml			
2G	320 ng/ml			
1H	640 ng/ml			
2H	640 ng/ml			
3A	Sample			
4A	Sample			
3B	Sample			
4B	Sample			

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SYMBOLS KEY

MTP	Panitumumab Coated Microtiter Plate (12x8 wells)
STD	Anti-Panitumumab Standard, Lyophilized
HRP CONJ	Panitumumab:HRP conjugate Concentrated
DETN DIL	Detection Diluent
1X SAMP DIL	(1X) Sample Diluent
1X STD DIL	(1X) Standard Diluent
20X WASH BUF	(20X) Wash Buffer
SUB TMB	TMB Substrate
SOLN STOP	Stop Solution
	Consult Instructions for Use
REF	Catalog Number
	Expiration Date
	Storage Temperature