

IDK[®] Pancreatic Elastase ELISA

*For the in vitro determination of
human pancreatic elastase in stool
For laboratory professional use only*

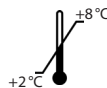
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IVD

Rx Only



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1. INTENDED USE

This Immundiagnostik AG assay is a quantitative test system intended for the measurement of pancreatic elastase in stool. Measurements obtained by this assay are to be used as an aid in the diagnosis of exocrine pancreatic insufficiency .

For *in vitro* diagnostic use only. For laboratory professional use only.

2. MATERIAL SUPPLIED

Art. no.	Label	Kit components	Quantity for cat. no.	
			K 6915. US	K 6915.20. US
K 6915	PLATE	Microtiter plate, pre-coated	12 x 8 wells	20 x 12 x 8 wells
K 0001.C.100	WASHBUF	Wash buffer concentrate, 10x	2 x 100 ml	40 x 100 ml
K 6915	CONJ	Conjugate concentrate, peroxidase-labelled (mouse anti pancreatic elastase)	1 x 200 µl	15 x 200 µl
K 6915	STD	Standards, lyophilised (see specification for concentrations)	4 x 5 vials	25 x 5 vials
K 6915	CTRL1	Control, lyophilised (see specification for range)	4 x 1 vial	25 x 1 vial
K 6915	CTRL2	Control, lyophilised (see specification for range)	4 x 1 vial	25 x 1 vial
K 0002.15	SUB	Substrate (tetramethylbenzidine), ready-to-use	1 x 15 ml	20 x 15 ml
K 0003.15	STOP	Stop solution, ready-to-use	1 x 15 ml	20 x 15 ml
K 6999.C.100	IDK Extract®	Extraction buffer concentrate IDK Extract® 2.5x	1 x 100 ml	--

For reorders of single components, use the catalogue number followed by the label as product number.

3. MATERIAL REQUIRED BUT NOT SUPPLIED

- Ultrapure water*
- Calibrated precision pipettors and 10–1 000 µl single-use tips
- Foil to cover the microtiter plate
- Horizontal microtiter plate shaker
- Multi-channel pipets or repeater pipets
- Vortex
- Standard single-use laboratory glass or plastic vials, cups, etc.
- Microtiter plate reader (required filters see chapter 7)

* Immundiagnostik AG recommends the use of ultrapure water (water type 1; ISO 3696), which is free of undissolved and colloidal ions and organic molecules (free of particles > 0.2 µm) with an electrical conductivity of 0.055 µS/cm at 25 °C (≥ 18.2 MΩcm).

4. STORAGE AND PREPARATION OF REAGENTS

- To run the assay more than once, ensure that reagents are stored at the conditions stated on the label. **Prepare only the appropriate amount necessary for each run.** The kit can be used up to 4 times within the expiry date stated on the label.
- Reagents with a volume less than **100 µl** should be centrifuged before use to avoid loss of volume.
- **Preparation of the wash buffer:** The **wash buffer concentrate (WASHBUF)** has to be diluted with ultrapure water **1:10** before use (100 ml WASHBUF + 900 ml ultrapure water), mix well. Crystals could occur due to high salt concentration in the concentrate. Before dilution, the crystals have to be redissolved at room temperature or in a water bath at 37 °C. The **WASHBUF** is stable at **2–8 °C** until the expiry date stated on the label. **Wash buffer** (1:10 diluted WASHBUF) can be stored in a closed flask at **2–8 °C for 1 month**.
- **Preparation of the extraction buffer:** The **extraction buffer concentrate IDK Extract®** has to be diluted with ultrapure water **1:2.5** before use (100 ml **IDK Extract®** + 150 ml ultrapure water), mix well. Crystals could occur due to high salt concentration in the concentrate. Before dilution, the crystals have to be redissolved at 37 °C in a water bath. The **IDK Extract®** is stable at **2–8 °C** until the expiry date stated on the label. Extraction buffer (1:2.5 diluted **IDK Extract®**) can be stored in a closed flask at **2–8 °C for 4 months**.
- The **lyophilised standards (STD)** and **controls (CTRL)** are stable at **2–8 °C** until the expiry date stated on the label. **Reconstitution** details are given in the **specification data sheet**. **Standards and controls** (reconstituted STD and CTRL) **are not stable and cannot be stored**.

- **Preparation of the conjugate:** Before use, the **conjugate concentrate (CONJ)** has to be diluted **1:101** in wash buffer (100µl CONJ + 10ml wash buffer). The CONJ is stable at **2–8 °C** until the expiry date stated on the label. **Conjugate (1:101 diluted CONJ) is not stable and cannot be stored.**
- All other test reagents are ready-to-use. Test reagents are stable until the expiry date (see label) when stored at **2–8 °C**.

5. STORAGE AND PREPARATION OF SAMPLES

Sample stability and storage

According to literature, the stability of pancreatic elastase in **raw stool** is 3 days at room temperature [3], 3 days at 4–8 °C [1], and up to a year at -20 °C [1].

Stool extract is stable at room temperature (15–30 °C) for three days, at 2–8 °C as well as at -20 °C for seven days. Avoid more than one freeze-thaw cycle.

Extraction of the stool samples

Extraction buffer (1:2.5 diluted *IDK Extract*®) is used as a **sample extraction buffer**. We recommend the following sample preparation:

Stool Sample Application System (SAS) (Cat. No.: K 6998SAS.US)

For laboratory professional use only.

Stool sample tube – Instructions for use

Please note that the dilution factor of the final stool suspension depends on the amount of stool sample used and the volume of the buffer.

SAS with 1.5 ml sample extraction buffer:

Applied amount of stool: 15 mg

Buffer Volume: 1.5 ml

Dilution Factor: 1:100

Please follow the instructions for the preparation of stool samples using the SAS as follows:

- a) The raw stool sample has to be thawed. For particularly heterogeneous samples we recommend a mechanical homogenisation using an applicator, inoculation loop or similar device.

- b) Fill the **empty stool sample tube** with **1.5 ml sample extraction buffer** (1:2.5 diluted **IDK Extract®**) before using it with the sample. **Important:** Allow the sample extraction buffer to reach room temperature.
- c) Unscrew the tube (yellow part of cap) to open. Insert the yellow dipstick into the sample. The lower part of the dipstick has notches which need to be covered completely with stool after inserting it into the sample. Place dipstick back into the tube. When putting the stick back into the tube, excess material will be stripped off, leaving 15 mg of sample to be diluted. Screw tightly to close the tube.
- d) Vortex the tube well until no stool sample remains in the notches. **Important:** Please make sure that you have a maximally homogenous suspension after shaking. Especially with more solid samples, soaking the sample in the tube with sample extraction buffer for ~ 10 minutes improves the result.
- e) Allow sample to stand for ~10 minutes until sediment has settled. Floating material like shells of grains can be neglected.
- f) Carefully unscrew the complete cap of the tube including the blue ring plus the dipstick. Discard cap and dipstick. Make sure that the sediment will not be dispersed again.

Dilution I: 1:100

Dilution of samples

The supernatant of the sample preparation procedure (dilution I) is diluted **1:100 in wash buffer**. For this purpose, one of the two following dilution procedure variants can be used:

Variant A (recommended by Immundiagnostik AG):

- **100 µl** supernatant (dilution I) + **900 µl** wash buffer, mix well = **1:10 (dilution IIa)**
- **100 µl** dilution IIa + **900 µl** wash buffer, mix well = **1:10 (dilution IIIa)**.
This results in a final dilution of 1:10 000.

For analysis, pipet **100 µl** of **dilution IIIa** per well.

Variant B:

Alternatively, the 1:100 dilution can be done in one step. For example:

- **10 µl** supernatant (dilution I) + **990 µl** wash buffer, mix well = **1:100 (dilution IIb)**. This results in a final dilution of 1:10 000.

For analysis, pipet **100 µl** of **dilution IIb** per well.

6. ASSAY PROCEDURE

Principle of the test

This ELISA is intended for the quantitative determination of pancreatic elastase in stool. In a first incubation step, the pancreatic elastase in the samples is bound to monoclonal antibodies, immobilised to the surface of the microtiter wells. To remove all unbound substances, a washing step is carried out. In a second incubation step, a peroxidase-labelled conjugate (mouse anti pancreatic elastase) is added which recognises specifically the bound pancreatic elastase. After another washing step to remove all unbound substances, the solid phase is incubated with the substrate, tetramethyl-benzidine (TMB), which reacts with the peroxidase. An acidic stop solution is added to stop the reaction. The colour changes from blue to yellow. The intensity of the yellow colour is directly proportional to the concentration of pancreatic elastase. A dose response curve of absorbance unit (optical density, OD at 450 nm) vs. concentration is generated using the values obtained from the standards. Pancreatic elastase, present in the patient samples, is determined directly from this curve.

Test procedure

Bring all **reagents and samples to room temperature** (15–30 °C) and mix well.

Mark the positions of standards/controls/samples on a protocol sheet.

Take as many microtiter strips as needed from the kit. Store unused strips together with the desiccant bag in the closed aluminium packaging at 2–8 °C. Strips are stable until expiry date stated on the label.

For automated ELISA processors, the given protocol may need to be adjusted according to the specific features of the respective automated platform. For further details please contact your supplier or Immundiagnostik AG.

We recommend to carry out the tests in duplicate.

1.	Add each 100 µl standards/controls/diluted samples into the respective wells.
2.	Cover the strips and incubate for 30 min at room temperature (15–30 °C) on a horizontal shaker* .
3.	Discard the content of each well and wash 5 times with 250 µl wash buffer . After the final washing step, remove residual wash buffer by firmly tapping the plate on absorbent paper.
4.	Add 100 µl conjugate (diluted CONJ) into each well.

5.	Cover the strips and incubate for 30 min at room temperature (15–30 °C) on a horizontal shaker* .
6.	Discard the content of each well and wash 5 times with 250 µl wash buffer . After the final washing step, remove residual wash buffer by firmly tapping the plate on absorbent paper.
7.	Add 100 µl substrate (SUB) into each well.
8.	Incubate for 10–20 min** at room temperature (15–30 °C) in the dark .
9.	Add 100 µl stop solution (STOP) into each well and mix well.
10.	Determine absorption immediately with an ELISA reader at 450 nm against 620 nm (or 690 nm) as a reference. If no reference wavelength is available, read only at 450 nm. If the extinction of the highest standard exceeds the range of the photometer, absorption must be measured immediately at 405 nm against 620 nm as a reference.

* We recommend shaking the strips at 550 rpm with an orbit of 2 mm.

** The intensity of the colour change is temperature sensitive. We recommend observing the colour change and stopping the reaction upon good differentiation.

7 RESULTS

The following algorithms can be used alternatively to calculate the results. We recommend using the “4 parameter algorithm”.

1. 4 parameter algorithm

It is recommended to use a linear ordinate for the optical density and a logarithmic abscissa for the concentration. When using a logarithmic abscissa, the zero standard must be specified with a value less than 1 (e.g. 0.001).

2. Point-to-point calculation

We recommend a linear ordinate for the optical density and a linear abscissa for the concentration.

3. Spline algorithm

We recommend a linear ordinate for the optical density and a linear abscissa for the concentration.

The plausibility of the duplicate values should be examined before the automatic evaluation of the results. If this option is not available with the programme used, the duplicate values should be evaluated manually.

Stool samples

The obtained results have to be multiplied by the **dilution factor of 10 000** to get the actual concentrations.

In case **another dilution factor** has been used, multiply the obtained result by the dilution factor used.

8. LIMITATIONS

Samples with concentrations above the measurement range (see definition below) can be further diluted and re-assayed. Please consider this higher dilution when calculating the results.

Samples with concentrations lower than the measurement range (see definition below) cannot be clearly quantified.

The upper limit of the measurement range can be calculated as:

highest concentration of the standard curve × sample dilution factor to be used

The lower limit of the measurement range can be calculated as:

LoB × sample dilution factor to be used

LoB see chapter "Performance Characteristics".

Liquid stools may lead to false pancreatic elastase results. In such cases, we recommend to also consider clinical symptoms and other diagnostic tests for the final diagnosis and/or to request another patient sample.

9. QUALITY CONTROL

Immundiagnostik AG recommends the use of external controls for internal quality control, if possible.

Control samples should be analysed with each run. Results, generated from the analysis of control samples, should be evaluated for acceptability using appropriate statistical methods. The results for the patient samples may not be valid if within the same assay one or more values of the quality control sample are outside the acceptable limits.

Reference range

We recommend each laboratory to establish its own reference concentration range.

10. PERFORMANCE CHARACTERISTICS

Accuracy – Precision

Repeatability (Intra-Assay); n = 42

The repeatability was assessed with 2 stool samples under **constant** parameters (same operator, instrument, day and kit lot).

Sample	Mean value [µg/ml]	CV [%]
1	187.92	5.7
2	308.06	4.0

Reproducibility (Inter-Assay); n = 20

The reproducibility was assessed with 2 stool samples under **varying** parameters (different operators, instruments, days and kit lots).

Sample	Mean value [µg/ml]	CV [%]
1	209.01	9.7
2	349.12	7.4

Accuracy – Trueness

The trueness states the closeness of the agreement between the result of a measurement and the true value of the measurand. Therefore, pancreatic elastase spikes with known concentrations were added to 3 different stool samples.

Sample [ng/ml]	Spike [ng/ml]	Expected [ng/ml]	Obtained [ng/ml]	Recovery [%]
23.69	4.0	27.69	27.05	97.70
	6.0	29.69	30.12	101.45
	8.0	31.69	32.59	102.82
	9.0	32.69	34.74	106.25

Sample [ng/ml]	Spike [ng/ml]	Expected [ng/ml]	Obtained [ng/ml]	Recovery [%]
68.63	4.0	72.63	74.16	102.12
	6.0	74.63	76.35	102.31
	8.0	76.63	76.32	99.60
	9.0	77.63	80.11	103.20
22.96	4.0	26.96	29.17	108.19
	6.0	28.96	29.71	102.58
	8.0	30.96	30.96	100.00

Analytical sensitivity

The following values have been estimated based on the concentrations of the standard without considering possibly used sample dilution factors.

Limit of blank, LoB	0.906 ng/ml
Limit of detection, LoD	1.884 ng/ml
Limit of quantitation, LoQ	2.055 ng/ml

Analytical specificity

The specificity of the antibody was tested by measuring the cross-reactivity against a range of compounds with structural similarity to pancreatic elastase. There was no cross-reactivity observed.

Substance tested	Concentration added	Concentration obtained [ng/ml]	Conclusion
Chymotrypsin	166.67 ng/ml	< 0.906	< LoB
PMN-elastase	10 µg/ml	< 0.906	< LoB
Pankreatic amylase	37 ng/ml	< 0.906	< LoB
Hemoglobin	500 ng/ml	< 0.906	< LoB
Pankreatic lipase	200 U/l	< 0.906	< LoB
Calprotectin	840 ng/ml	< 0.906	< LoB
α1-antitrypsin	90 µg/ml	< 0.906	< LoB
Pancreatin	80 mg/ml	< 0.906	< LoB

Linearity

The linearity states the ability of a method to provide results proportional to the concentration of analyte in the test sample within a given range. This was assessed according to CLSI guideline EP06-A with a serial dilution of 3 different stool-samples.

For pancreatic elastase in stool, the method has been demonstrated to be linear from 3.42 to 48.89 ng/ml, showing a non-linear behaviour of less than $\pm 20\%$ in this interval.

Sample	Dilution	Expected [ng/ml]	Obtained [ng/ml]	Recovery [%]
A	1:40 000	48.89	48.89	100.00
	1:80 000	24.44	28.52	116.69
	1:160 000	12.22	12.69	103.84
	1:320 000	6.11	6.86	112.24
	1:640 000	3.06	3.42	111.92
B	1:80 000	40.10	40.10	100.00
	1:160 000	20.05	22.85	113.97
	1:320 000	10.03	10.95	109.22
	1:640 000	5.01	5.34	106.49
C	1:40 000	40.68	40.68	100.00
	1:80 000	20.34	23.54	115.71
	1:160 000	10.17	11.71	115.10
	1:320 000	5.09	5.73	112.76

11. PRECAUTIONS

- Human materials used in kit components were tested and found to be negative for HIV, Hepatitis B and Hepatitis C. However, for safety reasons, all kit components should be treated as potentially infectious.
- Kit reagents contain sodium azide or ProClin as bactericides. Sodium azide or ProClin are hazardous to health and the environment. Substrates for enzymatic colour reactions may also cause skin and/or respiratory irritation. Any contact with the substances must be avoided. Further safety information can be found in the safety data sheet, which is available from Immundiagnostik AG on request.

- The 10x Wash buffer concentrate (WASHBUF) contains surfactants which may cause severe eye irritation in case of eye contact.

Warning: Causes serious eye irritation. **IF IN EYES:** Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. If eye irritation persists: get medical Advice/attention.

- The stop solution consists of diluted sulphuric acid, a strong acid. Although diluted, it still must be handled with care. It can cause burns and should be handled with gloves, eye protection, and appropriate protective clothing. Any spill should be wiped up immediately with copious quantities of water. Do not breath vapour and avoid inhalation.

12. TECHNICAL HINTS

- Do not interchange different lot numbers of any kit component within the same assay. Furthermore we recommend not assembling wells of different microtiter plates for analysis, even if they are of the same batch.
- Control samples should be analysed with each run.
- Reagents should not be used beyond the expiration date stated on kit label.
- Substrate solution should remain colourless until use.
- To ensure accurate results, proper adhesion of plate sealers during incubation steps is necessary.
- Avoid foaming when mixing reagents.
- Do not mix plugs and caps from different reagents.
- The assay should always be performed according to the enclosed manual.

13. GENERAL NOTES ON THE TEST AND TEST PROCEDURE

- The guidelines for medical laboratories should be followed.
- **IDK®** and **IDK Extract®** are trademarks of Immundiagnostik AG.
- Incubation time, incubation temperature and pipetting volumes of the components are defined by the producer. Any variation of the test procedure, which is not coordinated with the producer, may influence the results of the test. Immundiagnostik AG can therefore not be held responsible for any damage resulting from incorrect use.














- Warranty claims and complaints regarding deficiencies must be logged within 14 days after receipt of the product. The product should be send to Immunodiagnostik AG along with a written complaint.

14. REFERENCES

General literatur

1. Nandhakumar, N. & Green, M.R., 2010. Interpretations: How to use faecal elastase testing. Archives of disease in childhood. *Education and practice edition*, **95**(4), pp.119–23.
2. Löser, C., Möllgaard, A. & Fölsch, U.R., 1996. Faecal elastase 1: a novel, highly sensitive, and specific tubeless pancreatic function test. *Gut*, **39**(4), pp.580–6.
3. Stein, J. et al., 1996. Immunoreactive elastase I: clinical evaluation of a new noninvasive test of pancreatic function. *Clinical chemistry*, **42**(2), pp.222–6.

Used symbols:

	Temperature limitation		Catalogue number
	In Vitro Diagnostic Medical Device		To be used with
	Manufacturer		Contains sufficient for <n> tests
	Lot number		Use by
	Attention		Consult instructions for use
	Consult specification data sheet		Irritant
	Prescription use only		

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