BÜHLMANN fPELA® turbo

Warning

Immuno turbidimetric fecal ELASTASE Assay

Fast

Flexible

Efficient fecal testing

BÜHLMANN

Fecal Pancreatic Elastase results within 10 minutes

- Optimize your workflow by automation
- Integrate fecal samples into routine
- Measuring Range from 10 to 5000 µg/g

Simplify and improve fecal sample preparation with CALEX[®] Cap

- High quality for fast and efficient preparation
- Two fecal markers with one preparation Fecal calprotectin / pancreatic elastase

Highest correlation to manual reference method

- Continue with the established cut-off
- Specific to human enzymatic isoforms

BÜHLMANN fPELA® turbo

Fecal pancreatic elastase result within 10 minutes

BÜHLMANN fPELA® turbo, the turbidimetric immunoassay, is a flexible solution to be applied on most clinical chemistry analyzers.

The technology is a milestone in automation of pancreatic elastase quantification. It allows very rapid and flexible random access use, as well as being the ideal solution for high throughput applications in the routine laboratory. The fPELA turbo assay reduces the hands-on time dramatically and allows reporting pancreatic elastase results from human stool samples within shortest time.

Fecal pancreatic elastase in PEI

Pancreatic exocrine insufficiency (PEI), is a condition in which people are unable to adequately digest fats, carbohydrates and proteins due to a lack of digestive enzymes being produced from the pancreas. This results in nutrient malabsorption and malnutrition with severe consequences in the quality of life.

The determination of pancreatic Elastase levels is the most commonly employed indirect test for exocrine pancreatic function. The concentration of the enzyme in feces is five times higher than that in the pancreatic juice. It reflects the level of pancreatic output and correlates also with the output of other pancreatic enzymes such as lipase, amylase, and trypsin^{1,2}).

Lévy, Gastroenterol Clin Biol. 2006;30(6-7)
 Van de Vijver, J Pediatr Gastroenterol Nutr. 2011;53(1)

Application notes available	
Roche cobas c501 / c502	Siemens Atellica
Roche cobas c701 / c702	Beckman AU Series
Roche cobas Pro c503	Mindray BS-240Pro
Roche cobas Pure c303	Thermo Fisher Indiko
Abbott Alinity	BioSystems BA200
Abbott Architect c	The Binding Site Optilite

PRODUCT DESCRIPTION	
Method	Particle-enhanced turbidimetric immunoassay (PETIA)
Sample Type	Fecal extract
Kit Format	2 reagents (R1/ R2) Calibrators and controls provided separately Reagent set lasts for ≥100 tests
Sample Preparation	CALEX® Cap extracts ready to use without dilution
Reagent on board	Stable for 3 months
Calibration	Stable for 30 days
Calibration range	0-500 µg/g
Measuring range	10-5000 µg/g
Sample volume	~10 µL centrifuged fecal Extract (1:500)
Time to result	~10 min CALEX® Extraction ~20 min

Combination fecal calprotectin and pancreatic elastase

PEI is usually associated with other medical conditions, including cystic fibrosis, chronic pancreatitis, pancreatic cancer, diabetes, gastrointestinal surgery, coeliac disease, irritable bowel syndrome, or inflammatory bowel disease.

A significant amount of laboratory requests combines the quantification of fecal calprotectin and pancreatic elastase. The CALEX® Cap preparation device prefilled with a unique extraction buffer allows using the same fecal extract for quantification of both analytes at the same time. This synergy allows an additional significant reduction and streamlining of work load for fecal testing in the modern automated laboratory.

The analyzers mentioned in the left table are registered trademarks held by the corresponding manufactures.

Simplify and improve fecal preparation with CALEX® Cap

The CALEX[®] Cap is a unique device for the fast and efficient quantitative preparation of calprotectin in stool specimen.

The prefilled tubes are ready to use. Three simple steps are required for preparation:



The resulting extract (1:500) is ready to use in the turbidimetric assay. The CALEX[®] Cap preparation highly correlates with the manual weighing method.

Specificity and correlation to the manual reference method

The BÜHLMANN fPELA® turbo is based on polyclonal antibodies specific for the relevant human isoforms of the enzyme.

The assay is <u>not</u> affected by PERT (pancreatic enzyme replacement therapy).

Method comparison: A total of 130 stool samples from normal donors and PEI patients spanning the assay range were tested by the BÜHLMANN fPELA® turbo assay and a commercial Elastase-1 monoclonal assay. Results are summarized in the table below. Applying a cut-off at 200 µg/g, an overall agreement of 93.1% was found.

BÜHLMANN fPELA® turbo	Elastase 1 ELISA				
	>200 µg/g	100-200 µg/g	<100 µg/g	Total	
>200 µg/g	37	9	2	48	
100-200 µg/g	5	22	9	36	
<100 µg/g	1	8	37	46	
Total	43	39	48	130	



BÜHLMANN Laboratories AG Baselstrasse 55 4124 Schönenbuch Switzerland Phone +41 61 487 12 12 Orders logistics@buhlmannlabs.ch info@buhlmannlabs.ch www.buhlmannlabs.ch

BÜHLMANN fPELA® turbo Ordering Codes:

 Reagent Kit (~100 tests)
 B-KPELA-RSET
 R1 27 mL, R2 5.1 mL

 Calibrator Kit
 B-KPELA-CASET
 6 levels, 1 mL each; Ready to use

 Control Kit
 B-KPELA-CONSET
 3 x 2 levels, 1 mL each; Ready to use

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 $\mathsf{B}\ddot{U}\mathsf{HLMANN}$ fPELA* and CALEX* are registered trademarks of $\mathsf{B}\ddot{U}\mathsf{HLMANN}$ in many countries.

Parts of the CALEX® Cap are patent protected by: EP2833795(B1); US9752967(B2); AU2016203121(B2); CA2997598(C); JP6307132(B2); KR10-1875862(B1)