Hipro™ Cys-C Test (Rate Scattering Turbidimetric Method) Package Insert

【Product Name】

General Name：Cystatin C Test Kit (Rate Scattering Turbidimetric Method)

Trade Name： Cys-C Test

【Packing】

25 Tests/ Kit.

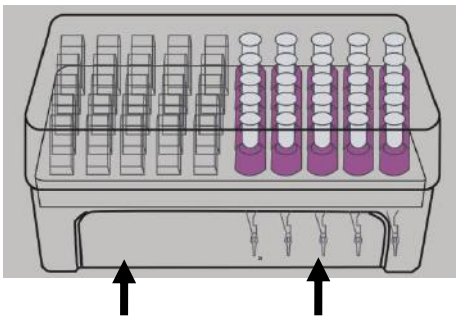
【Intended Use】

This product is used to determine the content of Cystatin C (CysC) in human serum or plasma.

【Principle】

The antibody of Cystatin C is coated on the latex surface. The CysC in the sample and the antibody become to immune complexes by Latex agglutination reaction. The immune complexes will produce the phenomenon of light scattering which is proportional to the intensity of scattered light and samples of CysC levels. Using specific protein analyzer to measure the intensity of scattered light, the concentration of CysC is determined by comparing the turbidity of samples to the standard concentration.

The kits contains all the reactive reagents.（figure 1）.



Reagent 1 Reagent 2

【Components】

	Components	Content
Reagent1 (R1)	Glycine buffer （pH7.4）	0.1mol/L
	Polyethylene glycol 6000	< 0.5%
	Sodium azide	0.1%
Reagent2 (R2)	Phosphate buffer（pH 8.0 ）	0.1mol/L
	Anti-Cystatin C antibody with latex	3.8ml/L

Do not mix different batches of reagents.

【Storage& Expire Date】

2 °C 8 °C Store at: 2 ~ 8℃.

Validity Period: 1 year.

Do not keep the kits at room temperature for long time. Restore the kits at 2-8℃ after use.

【Applicable Instrument】

HP-083/4-I specific protein analyzer, HP-083/4-II specific protein analyzer, HP-AFS/3 specific protein analyzer, HP-AFS/1 specific protein analyzer.

【Specimen】

Plasma (heparin anticoagulation, EDTA anticoagulant) or serum samples；Fasting blood collection and separation of serum as soon as possible; avoid hemolysis; store at 2～4℃ for 3 days.

【Procedure】

Preparation

The operation of specific protein analyzer please refer to the instruction. Start up the analyzer **30 minutes** before the test.

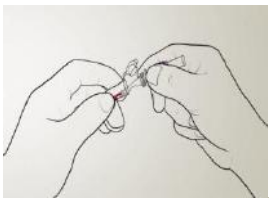
Attention：

HP-083/4-I 、 HP-083/4-II Specific protein analyzer: Insert the IC card into the slot, press the corresponding channel button to read the parameter information first.

Procedure

HP-083/4-I、HP-083/4-II Specific protein analyzer：

1、 Sample Preparation：



The test kit is equilibrated to room temperature, take samples by capillary in front of the sample collector, insert the sample collector into the cuvette.

Important: Due to the impact of evaporation, complete the test immediately once the capillary is full of samples. **Ensure the capillary full of samples**.

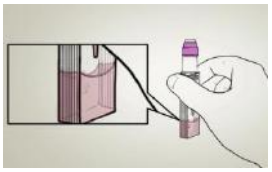
2. Sample Mixing：

Hold the middle of the cup on both sides, shake to mix

3. Reagent (R2) Injection：

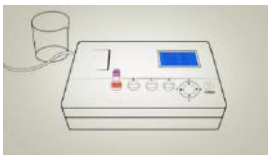
Inject the reagent R2 into the cuvette.

4. Reagents Mixing：



Attention：
Mixing Time:**3-5 seconds**

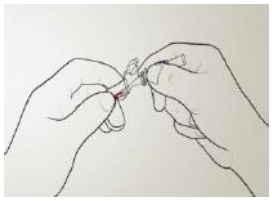
5. Test：



Insert into the corresponding test channel, The results will be displayed on the window and printed automatically

HP-AFS/1、HP-AFS/3 Specific protein analyzer:

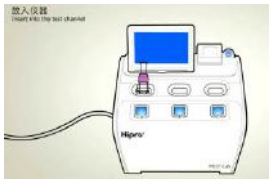
1. Sample Preparation:



The test kit is equilibrated to room temperature, take samples by capillary in front of the sample collector, insert the sample collector into the cuvette.

Important: Due to the impact of evaporation, complete the test immediately once the capillary is full of samples. **Ensure the capillary full of samples**.

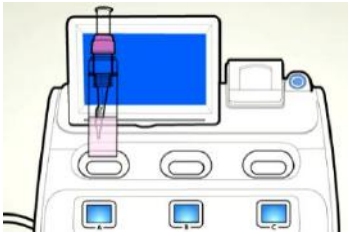
2. Test：



Insert the R2 reagent into R1 cuvette, insert the R1 cuvette into the test channel, the test will be done automatically, The results will be displayed on the window and printed.

Attention:

In step 2, insert the cuvette into test channel directly and the two-dimensional code on the cuvette face to screen (As figure). **Do not mix the sample and press the piston.**



Calibration

This product can be traced back toERM-DA471/IFCC.The calibration values for the different lots of the kits are stored on the calibration IC card or the two-dimensional code on the cuvette. Before test the new lot of kits, read the calibration card parameters first. Or the instrument automatically scan the two-dimensional code on the cup to obtain the corresponding calibration curve during testing.

Quality control

3- level calibration system guarantee the results' reliability for each lot of test kits, including the instrument calibration, remote reagent calibration and the third party calibration.

The third party calibration applicable for:

1. The daily indoor quality control test.
2. New lots of reagent.
3. New operator training.
4. The results can not match the clinical symptoms.
5. The first use of the reagent.

If still can not be calibrated, contact the manufacture for further technical support.

【Reference Value】

Normal reference range: ＜1.16mg/L

Recommended that each laboratory establish its own reference range

【Interpretation】

The test results of this reagent are only for clinical reference. the clinical diagnosis and treatment of patients should be considered in combination with their symptoms / signs, medical history, other laboratory tests and treatment responses.

【Limitation】

Hemoglobin>5g/L, triglyceride>23mmol/L, bilirubi＞684μmol/L will affect the test result.

【Performance】

1.Linearity range:0.2mg/L～9mg/L

2. Detection limit: ≤0.12mg/L

The limit of detection means the lowest detectable analyte level that can distinguish the concentration. Calculate based on the minimum standard above the two standard deviation of the data (Blank table, 1+2SD, within-run precision, n=20).

3. Precision

Test the control material by Cystatin C Test Kit 2 times per day for 20 days (n=80) according to EP5-A2 of CLSI. The data as below:

a)

HP-083/4-II Specific Protein Analyzer					
Sample	Mean	Within-Run		Between-Run	
	mg/L	S.D.	%C.V.	S.D.	%C.V.
Control 2	0.71	0.04	5.9	0.04	5.8
Control 3	3.19	0.12	3.9	0.13	4.1

b)

HP-AFS/3 Specific Protein Analyzer					
Sample	Mean	Within-Run		Between-Run	
	mg/L	S.D.	%C.V.	S.D.	%C.V.
Control 2	0.71	0.05	6.5	0.04	6.2
Control 3	3.19	0.11	3.5	0.13	4.1

4. Methodology comparison

Compared to Hitachi 7060 Cys-C TIA(x) by test the same serum sample, the relative data as below:

HP-AFS/3 Specific Protein Analyzer				
Site No.	Sample Type	No.of Assays	Regression Line	Coefficient correlation
1	Serum	50	Y= 1.03X+00.06	0.98

The concentration of sample is about 0mg/L-7mg/L

【Precaution】

Attention:

Only for in vitro diagnostic.

Only for professional use.

All samples and reactive wastes are treated as sources of infection.

Do not use the kits beyond shelf life.

Do not mix different batches of reagents.

Warning :

To avoid error, do not forced to take out the cuvette from the device. Follow the device operation manual strictly, If the problem cannot be solved, contact the manufacturer for further technical support.

【SYMBOLS USED】

Symbol	Usage
	Use By
LOT	Batch Code
	Manufacturer
	Keep Away from Sunlight
2 °C 8 °C	Temperature Limitation
IVD	In Vitro Diagnostic Medical Device
EC REP	Authorized Representative in the European Community
CE	CE Mark
	Consult Instruction for use
	warning

【References】

Michael G，Shlipak MD，Mark J，et a1．Cystatin C and mortality in elderly persons with heart failure．Journal of the American College of Cardiology，2005，45：268-271．

【Manufacturer】

Shijiazhuang Hipro Biotechnology Co.,ltd

No.3 Building,Block C,Fangyi Tech. Park, No. 313Zhujiangdadao Street,Shijiazhuang, 050000 China.

EC REP Lotus NL B.V.

Koningin Julianaplein 10, 1e Verd, 2595AA, The Hague, Netherlands

R＆D CENTER: 3938 Trust Way, Hayward, CA94545 USA.

【Approval date and date of revision】

Approval date：Sept 9,2015

Date of Revision：May 6,2016

Date of Revision：May 1, 2017

Date of Revision：Jan 1, 2021