Instruction for High Sensitive C-Reactive Protein (HS-CRP) QC

【Product Name】

High Sensitive C-Reactive Protein (HS-CRP) QC

【Package Specification】

Applicable for PA50&PA54		
REF No.	Level	Specification
32081019	QC(N)	0.2ml/bottle × 2
32081018	QC(H)	0.2ml/bottle × 2

Applicable for PA120		
REF No.	Level	Specification
32081081	QC(N)	0.2ml/bottle × 2
32081080	QC(H)	0.2ml/bottle × 2

【Intended Use】

HS-CRP QC is designed for quality control of High Sensitive C-Reactive Protein (HS-CRP) detection by in vitro immune diagnostic system.

【Precaution】

This product is only used for in vitro diagnosis. Dispose the reagents according to the usual laboratory precautions.

Absorb the required amount for each use, and the remaining samples should not be returned to the original bottle.

If the quality control product is contaminated, the stability of the components will be decreased, it should be stop using and a new quality control product should be selected.

【Main Composition】

HS-CRP antigen

【Storage and Stability】

The sealed detection kit can be stored at 2-8℃ for 12 months. Do not freeze. Once opened, the reagents stored at 2-8℃ are stable for 30 days.

【Instructions for use】

The quality control product is liquid ready-to-use type. The usage is as follows:

1. Remove bottles from the refrigerator(2-8℃) and allow to warm to room temperature for 5-10 minutes before mixing.
2. Gently invert at least 3 times before sampling, but avoid bubbles.
3. Open the bottle carefully to avoid ejecting the contents.
4. After sampling, cover the bottle cap as soon as possible and store it in a sealed condition of 2 ~ 8℃ in time.

For specific usage, please refer to the corresponding kit instruction.

【Assignment】

Assigned the quality control product in the company laboratory, using kit and specific protein analyzer, the methodology was Nephelometry.

The assignment result is shown on the QC bottle label.

【Icon Illustration】

	Date of manufacture		Manufacturer
IVD	In vitro diagnostic medical device		Volume
	Biological risks	REF	Catalogue number
LOT	Batch code		Main component
	Use-by date		Temperature limit
EC REP	Authorized representative in the European Community	CE	CE Marking

【Manufacturer】

Genrui Biotech Inc.

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【Medical Device Manufacturing Enterprise Permit No.】

Guangdong SFDA (State Food and Drug Administration) authorized Medical Device Manufacturing Permit No. 20041046.

【Guarantee and Technical Support】

If invalid message repeats or need technical support, please contact Genrui Customer Service and Support Center.

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