

## Instruction for DS-Ref Ref. Filling Solution

#### [Product Name]

DS-Ref Ref. Filling Solution

### [Package Specification]

10mL/bottle REF:42046001

#### [Intended Use]

Used as filling solution for reference electrode.

#### Test Principle

Not Applicable

### [Main Composition]

Saturated KCI solution and Preservative.

#### 【Storage and Transportation】

Stable for 2 years when stored at 2-30 °C in the shade. The products should be transported with outer package.

#### [ Applicable Instruments ]

Electrolyte Analyzers

#### [Sample Requirements]

Not Applicable

#### Test Methods

Fill Ref. filling solution into reference electrode. For more Instrument operations, please refer to electrolyte analyzer operation manual.

## 【Reference Range】

Not Applicable

### 【Interpretation of Test Results】

Not Applicable

#### [Calibration and QC]

Not Applicable

### 【Limitations of Testing Methods】

Not Applicable

## [Performance Indicator]

Not Applicable

## [Precaution]

- 1. Please seal it properly after using, to avoid long-time exposure in the air.
- 2. Avoid skin contact .Wear gloves when operating.
- If the reagent is carelessly spattered onto the skin, eyes, etc., flush with clean water immediately, and if mistakenly swallowed, seek medical advice.
- 4. The best volume of solution for normal operation is 2/3 level of the intra-cavity.
- 5. Do not use the solution when package is damaged.

### 【Icon Illustration】

Label	Meaning
M	Date of manufacture
IVD	In vitro diagnostic medical device
***	Manufacturer
\$€	Biological risks
LOT	Batch code
1	Temperature limit
	Use-by date

EC REP	Authorized representative in the European Community
REF	Catalogue number
C€	CE Marking
₽	Volume
	Main component

#### Training information

Please refer to the service manual.

#### 【Help information】

If you need help please contact after-sales.

### 【Trouble shooting】

Please contact after-sales.

# [References]

Not Applicable.

## [Manufacturer]



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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.

# 【Instruction Approved and Revised Date】

Approved date: November 6<sup>th</sup> 2015 Revised date: October 20<sup>th</sup> 2017



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