Quick and Easy At Home Antigen Test Kit with 8 Minutes Result Time that use for testing Strep A

Our Product Introduction

Basic Information

- Place of Origin: china
- Brand Name: DVOT
- Certification: CE/ISO
- Model Number: Strep A Rapid Test Kit
- Minimum Order Quantity: 800 test
- Price: 0.9-1 USD
- Packaging Details: 800T/PER CTN Weight :16.6 KGS CTN SIZE :57*54*34CM
- Delivery Time: 15-20 work days
- Payment Terms: D/P, T/T, Western Union, MoneyGram
- Supply Ability: 1000000pcs

Product Specification

- Type: Rapid Test Kit
- Keyword: Antigen Rapid Test
- Result Time: 8 Minutes To Get
- Accuracy: 98%
- Test Time: 8 Minutes
- Certifications: CE ISO
- Time Of Use: Disposable
- Highlight: **ce clungene antigen rapid test**, **ce medical diagnostic test kits**, **disposable clungene antigen rapid test**

More Images

Product Description

Quick and Easy At Home Antigen Test Kit with 8 Minutes Result Time that use for testing Strep A

Product Description:

Our Influenza Test Kits are perfect for medical professionals looking for a quick and easy way to diagnose patients. With certifications such as CE and ISO, you can be sure that our test kits are of the highest quality.

Our Antigen Rapid Test Kits are perfect for those who are looking for a quick and easy way to test themselves for COVID19. With a simple to use design, you can get results in just 8 minutes.

GrOup A Streptococcal (StreD) infections are caused by Streptococcus,a bacterium GrOUD A responsible for a variety of health problems. These infections can range from mild skin infection or sore throat to severe,life-threatening conditions such as toxic shock syndrome (multi-organ failures) and necrotizing fasciitis (soft tissue disease), commonly Known as flesh eating disease.Most people are familiar with strep throat,which along with minor skin infection.is the most common form of the disease.Health experts estimate that more than 10 million mild infections (throat and skin) like these occur every year.

Technical Parameters:

Accuracy	98%
Certifications	CE, ISO
Time Of Use	Disposable
Test Time	8 Minutes
Result Time	8 Minutes To Get
Keyword	Antigen Rapid Test
Type	Rapid Test Kit

Applications:

The DOVT Strep A Rapid Test Kit is designed to be used in various medical diagnostic scenarios. One of the primary application occasions of this product is in the diagnosis of Streptococcal A infections. This kit can be used by healthcare professionals to quickly diagnose the presence of Streptococcal A bacteria in patients, allowing for timely treatment and management of the infection.

The DOVT Strep A Rapid Test Kit is also useful in emergency medical situations where a rapid diagnosis is required. The product can be used in hospitals, clinics, and other healthcare facilities to quickly diagnose a variety of medical conditions. The kit is easy to use and provides accurate results in a short amount of time, making it a valuable tool for healthcare professionals across a range of medical specialties.

In conclusion, the DOVT Strep A Rapid Test Kit is a reliable and accurate Antigen Rapid Test Kit that is suitable for use in a variety of medical diagnostic scenarios. It is a disposable product that is manufactured in China, and provides results within 8 minutes of testing time. This Rapid Test Kit is an important tool for the diagnosis of various medical conditions including Streptococcal A infections, COVID19 Infection, and emergency medical situations.

Support and Services:

Our Medical Diagnostic Test Kits are designed to provide accurate and reliable results for various medical conditions. We offer comprehensive technical support and services to ensure that our customers can use our products with confidence.

We are committed to providing the highest level of support and services to ensure that our customers are satisfied with our products. Contact us today to learn more about how we can assist you.

Advantages:

Conventional identification procedures for Group A Streptococcus from throat swabs involve the isolation and subsequent identification of viable pathogens by techniques that require 24 to 48 hours or longer. DVOT Strep A detects either viable or nonviable organisms directly from a throat swab,providing results within 8 minutes.

DVOT Strep A Rapid Test Kit offers a simple and highly sensitive screening assay to make a presump- tive diagnosis of Group A Strept coccal respiratory infection.

Pathogen and product description

Group A Streptococcal (Strep) infections are caused by Group A Streptococcus, a bacterium responsible for a variety of health problems. These infections can range from mild skin infection or sore throat to severe,life-threatening conditions such as toxic shock syndrome (multi-organ failures) and necrotizing fasciitis (soft tissue disease), commonly known as flesh eating disease. Most people are familiar with strep throat, which along with minor skin infection, is the most common form of the disease. Health experts estimate that more than 10 million mild infections (throat and skin) like these occur every year.

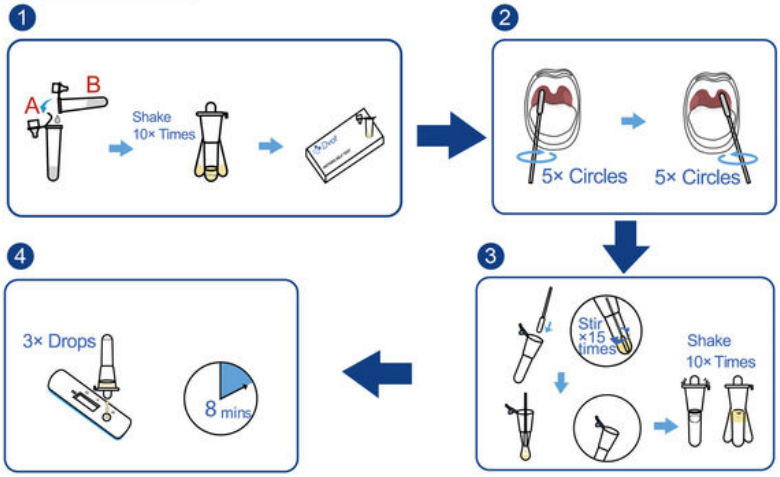
Conventional identification procedures for Group A Streptococcus from throat swabs involve the isolation and subsequent identification of viable pathogens by techniques that require 24 to 48 hours or longer. DVOT Strep A detects either viable or nonviable organisms directly from a throat swab, providing results within 8 minutes.

DVOT Strep A Rapid Test Kit offers a simple and highly sensitive screening assay to make a presump- tive diagnosis of Group A Strept coccal respiratory infection.

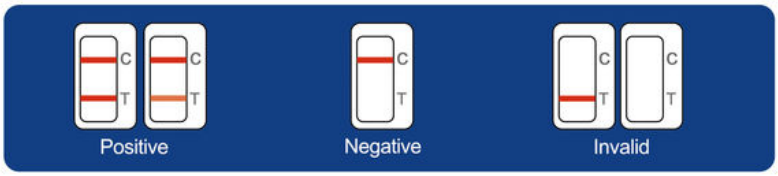
EXCELLENT CLINICAL COMPARISON RESULT

Items	Sensitivity	Specificity
Strep A Rapid Test Kit	92.5%	98.11%

How to test?



Result Reading



The packing details

CB	
QTY :	800PCS(40Boxs)
G.W :	15.5KGS
MEAS :	57x54x36cm

CERTIFICATE CERTIFICATE CERTIFICATE

NOTIFICATION OF REGISTRATION

Spanish Medicines and Health Products Administration

This is to certify that, according to the European Council Directive 98/79/EC, The manufacturer has completed the registration of the listed products:

MANUFACTURER: Feng Chun Yuan Medical Equipment (Shenzhen) Co., Ltd
ADDRESS: Room 1304&1306, No.48, Xinyu Road, Xiangshan Community, Xinqiao Street, Baoan District, Shenzhen, China 518000
SRN#: CN-MF-000002326
Mail: market@fcy-medical.com

The manufacturer has provided EU authorized representative with all the appropriate declaration according to the European Council Directive 98/79/EC including the Declaration of Conformity confirming that its in vitro diagnostic medical device, as stipulated here below, is fulfilling the essential requirements of the European Council Directive 98/79/EC.

IVD Devices: Strep A Rapid Test Kit
Classification: Others

Where the manufacturer affix the CE mark to the device listed they must ensure that all the essential requirements of European Council Directive 98/79/EC are met. The notification of abovementioned device has been completed by the European Authorized Representative in Spain. The Spain Competent Authority is notifed of the manufacture's device and has allocated registration. The registration number is RPS/1389/2024 .

CE

RIOMAVIX

Issue date: 23/MAY/2024
Cert. No.: RPS/1389/2024

Riomavix S.L. (ES-AR-000001202)
Calle de Almansa 55, 1D, Madrid 28039 Spain
Email:info@jiameiconsulting.com

DEPARTAMENTO DE
PRODUCTOS SANITARIOS

N/REF: PS/RPS/1389/2024

O F I C I O

Comunicación: RPS/1389/2024
Nº AEMPS: 24-02001
Fecha: 23/05/2024
Asunto: **Anotación de la comunicación en el Registro de Responsables de la puesta en el mercado de Productos Sanitarios**

RIOMAVIX SL
Calle de Almansa 55,1D
28039 - Madrid
MADRID
Madrid, Comunidad de

Con fecha **23/05/2024** ha sido **registrada** en la aplicación de Registro de Responsables de la puesta de mercado de Productos Sanitarios (RPS) de la Agencia Española de Medicamentos y Productos Sanitarios (AEMPS) la comunicación presentada por **RIOMAVIX SL**, con la siguiente información:

1. Número de identificación asignado en el registro

RPS/1389/2024

2. Responsable de la puesta en el mercado de los productos sanitarios

Empresa **RIOMAVIX SL**
Calle de Almansa 55,1D
28039 - Madrid (MADRID)
Madrid, Comunidad de

En calidad de **Representante**

3. Legislación que declara cumplir:

DIV - Directiva 98/79/EC.

4. Página(s) adicional(es) de productos sanitarios incluidos en esta comunicación.

REGISTRO DE RESPONSABLES DE LA PUESTA EN EL MERCADO DE PRODUCTOS SANITARIOS
DEPARTAMENTO DE PRODUCTOS SANITARIOS

Nota.- Esta notificación no tiene el carácter de una autorización sanitaria de comercialización, ni entraña un juicio sobre la conformidad del producto con la legislación vigente.Únicamente avala el cumplimiento del Registro de Responsables según el artículo 9 del RD 1662/2000 por el que se regulan los Productos Sanitarios para Diagnóstico in vitro.

Agencia Española de Medicamentos y Productos Sanitarios (AEMPS) CSV: Q3WKLE99AD
Fecha de la firma: 23/05/2024
Puede comprobar la autenticidad del documento en la sede de la AEMPS:https://localizador.aemps.es

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