



Parainfluenza 1 DFA
REAGENT

**A replacement component for Quidel Product 01-010000.v2 D³ Ultra™ DFA
Respiratory Virus Screening & ID Kit.**

FOR *IN VITRO* DIAGNOSTIC USE



INTENDED USE

Parainfluenza 1 DFA Reagent is a replacement component for Quidel Product 01-010000.v2 D³ Ultra DFA Respiratory Virus Screening & ID Kit.

MATERIALS PROVIDED

Quidel Product 01-013502.v2 contains:

Parainfluenza 1 DFA Reagent **2 mL**

One dropper bottle containing 2 mL of fluorescein labeled murine monoclonal antibodies directed against antigens produced by parainfluenza virus type 1. The buffered, stabilized, aqueous solution contains Evans Blue as a counter-stain and 0.1% sodium azide as preservative.

STORAGE AND HANDLING

Reagent will retain its full potency through the expiration date when stored at 2°C to 8°C. Store in the dark.

WARNINGS AND PRECAUTIONS

- For *in vitro* diagnostic use.
- Use undiluted.
- Refer to 01-010000.v2 Package Insert for complete directions.
- Sodium azide is included in this reagent at 0.1%.
 - Aqueous solutions of sodium azide, when mixed with acids, may liberate toxic gas.
 - Sodium azide may react with lead and copper plumbing to form highly explosive metal azides.
 - Avoid disposal of these solutions down sanitary or industrial plumbing systems.
 - Avoid release to the environment.
- Evans Blue is a potential carcinogen. If skin contact occurs, flush with water immediately.
- Testing should be performed in an area with adequate ventilation.
- Dispose of containers and unused contents in accordance with Federal, State and Local regulatory requirements.
- Wear suitable protective clothing, gloves, and eye/face protection when handling the contents of this kit.
- Wash hands thoroughly after handling.
- For additional information on hazard symbols, safety, handling and disposal of the components within this kit, please refer to the Safety Data Sheet (SDS) located at quidel.com.

WARRANTY STATEMENT

These products are warranted to perform as described in their labeling and the Quidel literature when used in accordance with their instructions. THERE ARE NO WARRANTIES WHICH EXTEND BEYOND THIS EXPRESS WARRANTY AND QUIDEL DISCLAIMS ANY IMPLIED WARRANTY OF MERCHANTABILITY OR WARRANTY OF FITNESS FOR PARTICULAR PURPOSE. Quidel sole obligation and purchaser's exclusive remedy for breach of this warranty shall be at the option of Quidel to repair or replace the products.

ASSISTANCE

To place an order or for technical support, please contact a Quidel Representative at 800.874.1517 (in the U.S.) or 858.552.1100 (outside the U.S.), Monday through Friday, from 8:00 a.m. to 5:00 p.m., Eastern Time. Orders may also be placed by fax at 740.592.9820. For e-mail support contact customerservice@quidel.com or technicalsupport@quidel.com.

For services outside the U.S.A., please contact your local distributor. Additional information about Quidel, our products, and our distributors can be found on our website quidel.com.

REF 01-013502.v2 – Parainfluenza 1 DFA Reagent

IVD



EC REP

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GLOSSARY

REF

Catalogue number



CE mark of conformity

EC REP

Authorized Representative
in the European Community

LOT

Batch code



Use by



Manufacturer



Temperature limitation



Intended use



Consult e-labeling
instructions for use



Do not reuse

IVD

For *In Vitro* diagnostic use
