Quidel Molecular Influenza A+B Control Set

Intended Use
The Quidel Molecular Influenza A+B Control Set is intended to be used as full process controls in molecular testing. These unassayed Control set for in vitro diagnostic use consists of a mixture of highly purified, inactivated strains of Influenza A/New Caledonia/20/99 (H1N1), Influenza B/Florida/04/06 and an influenza RNA-free matrix.

INSTRUCTIONS FOR USE
The Quidel Molecular Influenza A+B Controls should be used at the same volume as unknown samples in a validated molecular test methodology.

Materials Provided
SKU # M106
Control Kit (2 vials) – Store at 2° to 8°C

<table>
<thead>
<tr>
<th>#</th>
<th>Component</th>
<th>Quantity</th>
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</thead>
<tbody>
<tr>
<td>⊞CONTROL▶</td>
<td>Influenza A+B Positive Control Part M5029</td>
<td>1 vial/kit 2 mL</td>
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<tr>
<td>⊞CONTROL▼</td>
<td>Negative Control Part M5031</td>
<td>1 vial/kit 2 mL</td>
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Warnings and Precautions
• Treat all specimens, samples and controls as potentially infectious. Follow universal precautions when handling samples, this kit, and its contents.
• Avoid contact with skin and eyes.
• Wear suitable protective clothing, gloves, eye, and face protection when using this kit.
• Use micropipettes with an aerosol barrier or positive displacement tips for all procedures.
• Avoid microbial and cross contamination of the kit reagents. Follow Good Laboratory Procedures.
• Controls contain 0.09% Sodium Azide. Do not empty these controls into drains. Dispose of the containers in a safe way.
• Proper sample collection, storage, and transport are essential for correct results.
• Store assay reagents as indicated on their individual labels.
• MSDS is available upon request or can be accessed on the product website.

Storage and Handling of Kit Reagents
• Store the kit at 2° to 8°C until the expiration date listed on the labels.

QUALITY CONTROL
• The controls included in the Quidel Molecular Influenza A+B Control Set are unassayed controls. Control ranges may vary based on testing method. A specific range should be determined for each desired testing method. Each laboratory should establish their own Quality Control ranges and frequency of QC testing based on applicable local laws, regulations and standard good laboratory practice.
Limitations

- For *in vitro* diagnostic use.
- This control has been studied using a molecular testing methodology; it is not intended for use with other methodologies.

Customer and Technical Support

To place an order or for technical support, please contact a Quidel Representative at (800) 874-1517 (toll-free in the U.S.A.) or (858) 552-1100, Monday through Friday, between 8:00 a.m. and 5:00 p.m., Eastern Time. Orders may also be placed by fax at (740) 592-9820. For e-mail support contact customer service@dhiusa.com or technical_services@dhiusa.com. For services outside the U.S., please contact your local distributor. Additional information about Quidel, our products, and our distributors can be found on our website quidel.com.

References
