# QuickVue Influenza A+B Test Safety Data Sheet

## Section 1 - Product and Company Identification

### 1.1 Manufacturer Information

Quidel Corporation  
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Web: quidel.com  
E-mail: qehs@quidel.com

### 1.2 Product Information

- **Product Name:** QuickVue Influenza A+B Test (Catalog #: 20183, 20189, 20305, 20183IN)  
- **Recommended Use:** The QuickVue Influenza A+B test allows for the rapid, qualitative detection of influenza type A and type B antigens directly from nasal swab and nasopharyngeal swab specimens. The test is intended for use as an aid in the rapid differential diagnosis of acute influenza type A and type B viral infections. This test is intended for professional and laboratory use only.

### Components:

- Kit is composed of individually pouched test cassettes, extraction tubes, extraction reagent solution, disposable droppers, sterile nasal swabs, Influenza Type A positive control swab (swab is coated with non-infectious recombinant influenza A antigen), Influenza Type B positive control swab (swab is coated with non-infectious recombinant influenza B antigen) and negative control swab (swab is coated with formalin-inactivated, non-infectious *Streptococcus C* antigen).

## Section 2 – Hazards Identification

### 2.1 Emergency Overview

Significant health effects are not anticipated from routine use of this kit when following the precautions listed within the kit specific Package Insert, Universal Precautions and general safety laboratory practices. None of the components listed within this kit are considered hazardous as defined by the Occupational Safety and Health Administration (OSHA), the Canadian Workplace Materials Information System (WHMIS), or the European Union (EU) Directives.

When working with this kit we always recommend that employees wear appropriate personal protective equipment (PPE), including gloves, a lab coat and eye protection, and follow good laboratory hygiene practices to avoid accidental exposure to workplace materials. Universal Precautions should be followed when working with any potentially infectious material.

### SAFETY DATA SHEET (SDS) NOT REQUIRED FOR THIS TEST KIT

- Contains non-hazardous quantities of proprietary ingredients according to OSHA (29 CFR 1910.1200).
- Not a hazardous mixture according to Regulation (EC) No. 1272/2008.
- No chemicals need to be disclosed according to the applicable regulations for the components of this kit.

## Section 3 – Composition / Information on Ingredients

### Table

<table>
<thead>
<tr>
<th>Component</th>
<th>Chemical Name</th>
<th>CAS #</th>
<th>EINECS #</th>
<th>Concentration (%)</th>
<th>Volume</th>
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Other Information

Every effort has been made to adhere to the hazard criteria and content requirements of the U.S. OSHA Hazard Communication Standard, European Communities Safety Data Sheets Directive, Canadian Controlled Products Regulations, UK Chemical Hazard information and Packaging Regulations, and UN Globally Harmonized System of Classification and Labeling of Chemicals.

Prepared By: Quidel Corporation, EHS Department  
Supercedes: February 21, 2017  
Revisions: Updated catalog numbers; removed nasal aspirate and nasal wash as specimen types.

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