

**MATERIAL SAFETY DATA SHEET**Rev. 00  
27/03/06**IMUNO-RÁPIDO DENGUE****COMPANY AND PRODUCT IDENTIFICATION**

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Product Name	Trade name: IMUNO-RÁPIDO DENGUE IGG/IGM Code: 626010 – 10 determinations Code: 626020 – 20 determinations Code: 626025 – 25 determinations Code: 626040 – 40 determinations Code: 626080 – 80determinations

**PRODUCT INFORMATION**

Description	Kit contains: - Test Device - Diluent solution
Product Use	In Vitro Diagnostic only
Hazard Components	Test device is composed of plastic and test strip coated, Diluent solution contains Sodium azide (0.1%) as preservative.

**HAZARD IDENTIFICATION**

The components of this product are non-toxic when handled according to Good Manufacturing Practices. Do not open the test device.


Sodium azide 0.1% may react with lead and cooper plumbing to form highly explosive metal azides. Upon disposal of liquids, flush with large volumes of water to prevent azide buildup.

**FIRST AID MEASURES**

Contact with eyes	Wash with plenty of water. Seek medical advice.
Contact with skin	Wash with plenty of water.
Inhalation	Exposure to fresh air immediately. Seek medical advice.
Ingestion	Seek medical advice.

**FIRE FIGHTING MEASURES**

Non-inflamable

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<b>MEASURES OF CONTROL – LEAKING/SPILLAGE</b>	
Isolate and ventilate the area. Remove with absorbent material. Clean up area and wash with plenty of water.	

<b>HANDLING AND STORAGE</b>	
Handling	Avoid contact with eyes, skin and mouth. Wash hands after handling.
Storage	Keep at 2 to 30°C

<b>EXPOSURE CONTROL AND PERSONAL PROTECTION</b>	
Preventive Measure	Avoid ingestion, inhalation and contact with eyes and skin. Protection for eyes: yes Protection for hands: wear gloves; wash hands after handling.

<b>PHYSICAL AND CHEMICAL PROPERTIES</b>	
Appearance	Reagents in liquid Plastic device
Others	None relevant data for safety

<b>STABILITY AND REACTIVITY</b>	
Stability	The product is stable when used appropriately.
Avoid	Exposure to heat can damage the product. Do not freeze.

<b>DISPOSAL CONSIDERATIONS</b>	
Disposal in accordance with local regulations.	

<b>REGULATORY INFORMATION</b>	
Product is manufactured according to Good Manufacturing Practices for In Vitro Diagnostic kits.	

<b>OTHER INFORMATION</b>	
All products can contain unknown risks and they should be handled with care. It is the duty of the user to comply with all local regulations.	