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SAFETY DATA SHEET

IDENTIFICATION OF PRODUCT (SUBSTANCE) AND SUPPLIER

Product Name:	FastPack & FastPack IP Immunoassay Calibrators
Product Number:	25000002, 25000008, 25000014, 25000016, 25000024, 25000035,
	25000077
Intended Use:	In-Vitro Quantitative Testing
CAS No.	Mixture
Einecs No.	Mixture
Supplier's Name	Qualigen, Inc
Address:	2042 Corte del Nogal, Carlsbad, CA 92011
Phone Number:	(760) 918-9165
Emergency Phone Numbers:	(877) 709-2169

HAZARD IDENTIFICATION – HAZARDOUS COMPONENTS

The following information is furnished for the kits' hazardous constituents that require regulatory control or disclosure at the concentration found in the kit. Note that the information here is often based on data from the chemical raw material (LD50, exposure limits, etc.) The kit contains a significant diluted concentration in an aqueous solution yet the assessment below was made based on the hazards of its component parts at full concentration. This chemical is considered hazardous by the 2012 OSHA Hazard Communication Standard (29 CFR 1910.1200)

HMIS: Health-1, Flammability-0 Reactivity-0

Code letter and hazard designation of product:



Xi Irritant

Hazard-determining components of labeling:	Tris HCL
Hazard phrases:	H315: Causes skin irritation. H319: Causes serious eye irritation. H335: May cause respiratory irritation.
Precautionary phrases:	P264: Wash (hands and exposed skin) thoroughly after handling. P280: Wear protective gloves/protective clothing P501: Dispose of contents/container to: approved disposal facility.

3. COMPOSITION/INFORMATION ON INGREDIENTS – HAZARDOUS COMPONENTS

This vial should be handled only by qualified personnel trained in laboratory procedures and familiar with their potential hazards. Specific warnings are given in the instructions for use. The absence of a specific warning should not be interpreted as an indicator of safety.

Chemical Ingredient	CAS No.	Quantity in Calibrators
Tris HCL	1185-53-1	≤1.25%

4.	EMERGENCY FIRST AID MEASURES
General Information:	Symptoms of poisoning may even occur after several hours; therefore medical observation for at least 48 hours after the accident is recommended
Eye Contact:	Immediately flush eyes with plenty of tepid water for 15 minutes while
	separating eye lids with fingers. Remove contact lenses if worn. Obtain medical attention if needed or if symptoms, such as redness or irritation persist.
Skin Contact:	In case of contact, flush skin with copious amounts of cool water and remove
	contaminated clothing. Obtain medical attention if needed or if irritation or other symptoms develop.
Inhalation:	If inhaled, move from exposure area to fresh air. Seek medical attention if
	breathing becomes difficult or if cough or other symptoms develop.
Ingestion:	In case of ingestion, contact a poison control center or physician for instructions.
Notes to Physician:	According to OSHA Bloodborne Pathogens Standard (29 CFR 1910.1030),
	Universal Precautions apply. Persons handling human blood source samples should be offered hepatitis B vaccination prior to working with human source material.

FIRE FIGHTING MEASURES

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Flammable	Dilute aqueous solution not considered a fire hazard.
Properties:	
Suitable	Use extinguishing media appropriate for the surrounding fire.
Extinguishing Media:	
Unsuitable	Unknown
Extinguishing Media:	
Special Fire Fighting	Conventional fire fighting full protective equipment (with NIOSH-approved self-
Procedures:	contained breathing apparatus) and procedures appropriate for the surrounding fire should be sufficient.

ACCIDENTAL RELEASE MEASURES

Personal Precautions:	Wear Personal Protective Equipment (PPE) as indicated in Section 8. Avoid physical contact with material. Wash hands thoroughly after handling.
Environmental	This preparation may contain a small amount of sodium azide which can react
Precautions:	with copper, lead, brass or solder in plumbing systems and form potentially explosives metal azides. Follow proper disposal procedures.
Methods and	Absorb spills with inert material/sorbent. Decontaminate the spill site following
Materials for	standard procedures. Dispose of materials in accordance with all applicable
Containment and Clean-Up:	federal, state, local and provincial environmental regulations, per Section 13.

7.	HANDLING AND STORAGE
Handling:	Follow universal/standard precautions when handling this material. See Section 8. Minimize contact and contamination of personal clothing and skin. Wash
Storage:	hands thoroughly after handling. Store the kit components as specified in the product instructions/package insert provided with the test kit or the instrument operation manual.

8. EXPOSURE CONTROL/PERSONAL PROTECTION MEASURES

The following personal protective equipment (PPE) is recommended to prevent blood or other potentially infectious or hazardous materials from reaching the user's work or street cloths, skin, mouth, mucous membranes and eyes, under normal conditions of use and for the time during which the protective equipment is utilized:

Ventilation:	Adequate lab ventilation is required. It is recommended that users handle potentially infectious human source material/patient samples in a dedicated location.
Eye Protection:	Wear appropriate protective chemical safety glasses.
Protective Gloves:	Wear chemical resistant protective gloves
Protective Clothing:	Wear a labcoat, clinic jacket, gown, apron and/or smock.
Other:	All personal protective equipment should be removed before leaving the work area and placed in an appropriate designated area or container for storage, processing, decontamination or disposal.

PHYSICAL AND CHEMICAL PROPERTIES

Appearance:Clear to yellowish liquid.Fire Hazard:NoneFlash Point:NoneAuto Igniting:None

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STABILITY AND REACTIVITY

Chemical Stability: Incompatible Materials: Hazardous Decomposition Products: Possibility of Hazardous Reactions: Stable under normal conditions Strong Acids Variable Oxides under fire conditions only. Can react if in contact with – Strong acids (releases toxic gas)

11.	TOXICOLOGICAL INFORMATION
Acute Health Effects:	Harmful in contact with skin and if swallowed. May cause headache, nausea and vomiting.
Biohazard Potential:	Consists of $\geq 1\%$ human and/or animal protein. Human source material used was tested and found negative for HIV 1/2, HBsAg, HCV, HIV-1(NAT), HCV(NAT) and RPR by FDA approved methods.
Carcinogenicity:	No component of this product present at levels greater than or equal to 0.1% is identified as probable, possible or confirmed human carcinogen.
Reproductive Effects:	No information available

ECOLOGICAL INFORMATION

Toxicity **Bioaccumulative Potential Persistence and Degradability Other Adverse Effects**

No information available No information available No information available None known

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Methods

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DISPOSAL CONSIDERATIONS

Waste Treatment Disposal should be in accordance with local, state or national legislation. Do NOT dispose of via domestic waste. Ensure that all packaging is disposed of safely.

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TRANSPORT INFORMATION

Basic Shipping Description:

Not classified as dangerous for transport via land, sea, or air.

15.

REGULATORY INFORMATION

US Federal Regulations:	This preparation is a component of an FDA-regulated in vitro diagnostic device.
US State Regulations:	This product does not contain components whose ingredients are listed under California Proposition 65.

OTHER INFORMATION

This test kit should be handled only by qualified personnel trained in laboratory procedures and familiar with their potential hazards.

Specific warnings are given in the instructions for use. The absence of a specific warning should not be interpreted as an indication of safety.

Contact for general information:

Qualigen, Inc. 2042 Corte del Nogal, Suite B Carlsbad, CA 92011 USA Phone : (760) 918-9165 (8 am to 5 pm PST) (877) 709-2169 Qualigeninc.com

Disclaimer:

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This document was developed from information obtained from reputable sources, but does not purport to be all-inclusive. The data contained herein, which is based on our present knowledge and is intended for information only, shall not constitute a guarantee for any specific product features and shall not establish a legally valid contractual relationship. Regulatory requirements are subject to change and vary from one location to another; thus, it is the buyer's responsibility to ensure that its activities comply with national, regional and local laws and regulations. In no event shall Qualigen, Inc. be liable for any claims, losses, or damages of any individual or for lost profits or any special, indirect, incidental, consequential or exemplary damages, howsoever arising, even if Qualigen, Inc. has been advised of the possibility of such damages.