

**Section 1: COMPANY AND PRODUCT INFORMATION**

<b>1.1 Product Name:</b>	Inhibin A ELISA
<b>1.2 Product Code:</b>	AL-123 / AL-123-r
<b>1.3 Product Category:</b>	<b>GMDN N.A / EDMA N.A</b>
<b>1.4 Manufacturer:</b>	<b>Manufacturer:</b> Ansh Labs 445 Medical Center Blvd Webster, TX 77598 Ph: (281) 404-0260 <a href="mailto:techsupport@anshlab.com">techsupport@anshlab.com</a>
<b>1.5 Emergency telephone number:</b>	Emergency telephone (24 h): 911 Poison Control: 1-800-222-1222
<b>1.6 Relevant identified uses of the substance/mixture:</b>	For the quantitative measurement of Inhibin A. For <i>in vitro</i> professional laboratory use.
<b>uses advised against:</b>	Not for use in diagnostic procedures or therapeutic applications. Intended for laboratory professionals experienced in immunoassay techniques
<b>1.7 Kit content (name and label reference):</b>	

Component (Part No.)	Description	Hazard Information
CAL-123 (A-F)	Inhibin A Calibrators is in a serum-based matrix with $\leq 0.5\%$ ProClin™ 300 / Pro-Clean™ 400 and $\leq 0.09\%$ sodium azide	EUH208; H400–H410
CTR-123 (I-II)	Inhibin A Controls is in a serum-based matrix with $\leq 0.5\%$ ProClin™ 300 / Pro-Clean™ 400 and $\leq 0.09\%$ sodium azide	EUH208; H400–H410
BCC-123	Inhibin A Biotin Conjugate Concentrate with $\leq 0.5\%$ ProClin™ 300 / Pro-Clean™ 400	EUH208; H400–H410
ASB-123A	Inhibin A Assay Buffer A is in a non-ionic detergent with $\leq 0.5\%$ ProClin™ 300 / Pro-Clean™ 400 and $\leq 0.09\%$ sodium azide	EUH208; H400–H410
ASB-123B	Inhibin A Assay Buffer B with $\leq 0.5\%$ ProClin™ 300 / Pro-Clean™ 400	EUH208; H400–H410
CND-123	Inhibin A Biotin Conjugate Diluent with $\leq 0.5\%$ ProClin™ 300 / Pro-Clean™ 400	EUH208; H400–H410
SAR-123	Inhibin A Streptavidin Ready-to-Use (RTU) with $\leq 0.5\%$ ProClin™ 300 / Pro-Clean™ 400	EUH208; H400–H410

STP-100	Stopping Solution – 0.2 M sulfuric acid	Skin Corr. 1A (H314); Eye Dam. 1 (H318)
TMB-100	Substrate Solution – tetramethylbenzidine (TMB) in buffer	Eye Irrit. 2 (H319)
WSH-100	Wash Buffer Concentrate (25x) – phosphate-buffered saline solution with a nonionic detergent	EUH208
PLT-123	Microtiter plate coated with Inhibin A monoclonal antibody	Article – not hazardous

## Section 2: HAZARDS IDENTIFICATION

### 2.1 Classification of the substance or mixture:

#### According to Regulation (EC) No 1272/2008 (CLP) as amended:

This mixture is not classified as hazardous under the criteria of Regulation (EC) No 1272/2008 (CLP).

It contains trace concentrations of preservatives (isothiazolinones  $\leq 0.25\%$ , sodium azide  $\leq 0.1\%$ ) and dilute sulfuric acid ( $< 2\%$ ), all present below the concentration limits for classification.

The mixture does not meet the criteria for physical, health, or environmental hazard classification under CLP.

**Intended use:** For laboratory use by professional uses only. Use appropriate personal protective equipment while working with the reagents provided.

### 2.2 Label elements

#### Sulfuric acid:



H314 Causes severe skin burns and eye damage.

P280 Wear protective gloves, protective clothing and eye/ face protection.

P301+P330+P331 If swallowed: rinse mouth.

P303+P361+P353 If on skin (or hair): Rinse skin with water.

P305+P351+P338 If in eyes: Rinse cautiously with water for several minutes. Continue rinsing.

P310 Immediately seek physician assistance

#### Pro-Clin™ 300 / Pro-Clean 400:



H317 May cause an allergic skin reaction

P261 Avoid breathing vapors

P272 Contaminated work clothing should not be allowed out of the workplace

	P280 Wear protective gloves, protective clothing and eye/face protection P302+P352 If on skin: Wash with plenty of soap and water P333+P313 If skin irritation or rash occurs, seek medical assistance
<b>2.3 Hazards not otherwise classified (HNOC) or not covered by GHS:</b>	<ul style="list-style-type: none"> <li> <b>PBT / vPvB assessment:</b> The mixture does not contain any substances identified as PBT or vPvB according to Annex XIII of REACH.                     </li> <li> <b>Endocrine-disrupting properties:</b> None of the components are identified as having endocrine-disrupting properties under REACH or CLP.                     </li> <li> <b>Biological origin:</b> Some kit components may contain materials of human origin (e.g., serum-based buffers). These are tested and found non-reactive for known pathogens but should be handled using Biosafety Level 2 precautions per CDC/NIH <i>Biosafety in Microbiological and Biomedical Laboratories</i>, 5th Edition. The product also contains proteins of bovine or equine origin sourced from OIE-classified negligible BSE-risk herds in accordance with Reg. (EC) No 999/2001 and Commission Decision 2007/453/EC.                     </li> </ul> <p> <b>Other hazards:</b> Contains trace concentrations of isothiazolinones (&lt; 0.25 %) and sodium azide (&lt; 0.1 %) which are below classification thresholds but may cause sensitization in susceptible individuals. Under normal conditions of use, the mixture does not present any additional physical, health, or environmental hazards.                 </p>

### Section 3: COMPOSITION/INFORMATION ON INGREDIENTS

#### 3.1 Substances

<b>Stopping Solution</b>		<b>Hazard Classification of Pure Ingredients</b>			
Chemical Name	% by wt.	EU-67/548/EEC	EU 1272/2008 CLP/GHS	US OSHA	WHMIS
<b>Sulfuric Acid</b>  CAS # 7664-93-9 EINECS # 231-639-5 Index # 016-020-00-8	<1	C;R35	Eye Dam. 1 Skin Corr. 1A H314; H318	Water-Reactive Carcinogen Corrosive Highly Toxic	D1A; E

#### 3.2 Mixtures

This kit comprises buffered aqueous solutions and protein-based reagents containing the components listed below. Hazardous constituents are present at concentrations below classification thresholds under Regulation (EC) No 1272/2008 (CLP).

<u>Inhibin A Calibrators (A-F), Inhibin A Controls (I-II), Inhibin A Assay Buffer A, Inhibin A Assay Buffer B, Inhibin A Biotin Conjugate Concentrate, Inhibin A Biotin Conjugate Diluent, Inhibin A Streptavidin Conjugate Ready-to-Use</u>	<b>Hazard Classification of Pure Ingredients</b>
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Chemical Name	% by wt.	EU-67/548/EEC	EU 1272/2008 CLP/GHS	US OSHA	WHMIS
<b>ProClin™ 300<sup>2,8</sup> / Pro-Clean™ 400<sup>2,8</sup></b>  reaction mass of: 5-chloro-2-methyl-4-isothiazolin-3-one [EC# 247-500-7] and 2-methyl-4-isothiazolin-3-one [EC# 220-239-6](3:1)  <b>CAS # 55965-84-9</b> <b>EINECS # Not available</b> <b>Index # 613-167-00-5</b>	≤ 0.5	T;R23/24/25-34-43 N;R50/53	Acute Tox. Dermal 3 Acute Tox. Inhal. 3 Acute Tox. Oral 3 Aquatic Acute 1 Aquatic Longterm 1 Skin Corr. 1B Skin Sens. 1 H301; H311; H314; H317; H331; H400; H410	Corrosive Sensitizer Toxic	D1B; D2B; E
<b>Inhibin A Calibrators, Inhibin A Controls, Inhibin A Assay Buffer A</b>		<b>Hazard Classification of Pure Ingredients</b>			
Chemical Name	% by wt.	EU-67/548/EEC	EU 1272/2008 CLP/GHS	US OSHA	WHMIS
<b>Sodium Azide<sup>2,8</sup></b>  <b>CAS # 26628-22-8</b> <b>EINECS # 247-852-1</b> <b>Index # 011-004-00-7</b>	< 0.1	T+;R28-32 N;R50/53	T+;R28-32 N;R50/53 Acute Tox. Oral 2 Aquatic Acute 1 Aquatic Longterm 1 H300; H400; H410	Highly Toxic Irritant	D1A

### Section 4: FIRST AID MEASURES

#### 4.1 Description of first aid measures

General advice:	No special measures required. Consult a physician in case of complaints.
If inhaled:	If product is inhaled, move exposed individual to fresh air.
In case of skin contact:	In case of skin contact, flush with water for at least 15 minutes. Remove contaminated clothing and shoes. If pain or irritation occur, obtain medical attention.
In case of eye contact:	If product enters eyes, wash eyes gently under running water for 15 minutes or longer, making sure that the eyelids are held open. If pain or irritation occurs, obtain medical attention.
If swallowed:	If ingested, wash mouth out with water. Seek medical attention.

#### 4.2 Most important symptoms and effects, both acute and delayed:

To the best of our knowledge, the chemical, physical and toxicological properties have not been thoroughly investigated.

#### 4.3 Indication of any immediate medical attention and special treatment needed

No data available.

### Section 5: FIREFIGHTING MEASURES

**5.1 Flammable Properties:**

Nonflammable solution.

**5.2 Extinguishing media:**

Chemical or water fire extinguisher.

**5.3 Special hazards arising from the substance or mixture:**

No special hazards determined.

**5.4 Advise for Firefighters**

Wear self-contained breathing apparatus for firefighting, if necessary.

**5.5 NFPA Rating****Health:** 2**Flammability:** 0**Reactivity:** 1**Section 6: ACCIDENTAL RELEASE MEASURES****6.1 Personal precautions, protective equipment and emergency procedures:**

Use appropriate personal protective equipment (Wear rubber gloves, safety goggles, impermeable shoe covers and long laboratory coat).

**6.2 Spill and Leak Procedures:**

Absorb spilled material with an appropriate inert, non-flammable absorbent and dispose according to local regulations.

**6.3 Environmental precautions:**

Contain the spill to the smallest area possible. Do not let product enter drains. Discharge into the environment must be avoided.

**6.4 Methods and material for containment and cleaning up:**

Absorb with inert absorbent material and dispose of a waste (see section 13).

**6.5 Reference to other sections:**

For disposal see section 13.

**Section 7: HANDLING AND STORAGE****7.1 Precautions for safe handling:**

Wear suitable personal protective equipment. Take care not to splash spill or splatter reagents. Do not eat, drink, smoke or apply cosmetics in laboratory areas. Do not pipette samples or reagents by mouth.

**7.2 Recommended Storage and Conditions:**

Keep away from incompatible material (see Section 10).  
To maintain efficacy, store according to the instructions in the product labelling

**7.3 Specific end use(s):**

This product is intended for laboratory use by professional users only.

**Section 8: EXPOSURE CONTROLS / PERSONAL PROTECTION**
**8.1 Control parameters:**

Substance	EU IOELV / OEL (TWA 8 h)	OSHA PEL (8 h TWA)	ACGIH TLV	NIOSH REL	DFG MAK	Notes
Sulfuric acid (aerosol, thoracic fraction)	0.05 mg/m <sup>3</sup> (TWA, Directive 2000/39/EC)	1 mg/m <sup>3</sup>	0.2 mg/m <sup>3</sup> (thoracic fraction)	1 mg/m <sup>3</sup>	0.1 mg/m <sup>3</sup>	Irritant threshold limit
Sodium azide (as NaN <sub>3</sub> )	None established	None established	0.1 mg/m <sup>3</sup> ceiling	0.1 mg/m <sup>3</sup> ceiling	None	Applies only for aerosols/powder
Reaction mass of CMIT/MIT (3:1)	None established	None established	None	None	None	Trace preservative (<0.25%) – EUH208 applies

**8.2 Exposure controls**

- Engineering controls: Use only in a well-ventilated laboratory. Handle open reagents under local exhaust ventilation or a chemical fume hood to avoid aerosol generation.
- Eye/face protection: Safety glasses with side-shields or chemical goggles conforming to EN 166 / ANSI Z87.1.
- Hand protection: Disposable nitrile, neoprene, latex or rubber gloves (EN 374). Replace immediately if contaminated or torn.
- Body protection: Laboratory coat or equivalent protective garment.
- Respiratory protection: Not normally required under intended use. If airborne concentrations may exceed limits, use an approved respirator (NIOSH N95 / EU P2) after evaluation by a qualified safety professional.
- Environmental controls: Avoid release to drains or waterways. Flush waste solutions with excess water to prevent metal-azide accumulation in plumbing.
- Hygiene measures: Follow good laboratory practice. Do not eat, drink, or smoke while handling reagents. Wash hands thoroughly after use.

**8.3 Personal protective equipment:**

- This mixture is supplied as a small-volume in vitro diagnostic reagent kit for professional laboratory use only.
- REACH exposure scenarios: Not required; quantities are below registration thresholds.
- Regulatory alignment: Information in this SDS meets or exceeds requirements of EU REACH / CLP (2020/878), UK REACH, U.S. OSHA HazCom 2012 (29 CFR 1910.1200), ACGIH TLVs, NIOSH RELs, DFG MAKs, and GHS-aligned systems in Switzerland, Turkey, GCC, and China.
- Typical laboratory precautions: Avoid splashes and aerosols; open vials slowly; maintain secondary containment for liquids. Engineering and PPE measures above are adequate for all listed jurisdictions.
- Intended end use: Diagnostic or research use testing only; not for consumer or industrial manufacturing use. Product is exempt from OSHA workplace labeling when supplied in sealed diagnostic kits.

**Section 9: PHYSICAL AND CHEMICAL PROPERTIES**
**9.1 Information on basic physical and chemical properties:**

Property	a) Typical value / description	b) Property	c) Typical value / description
Appearance	Clear to slightly opalescent aqueous liquids (component-dependent)	Appearance	Clear to slightly opalescent aqueous liquids (component-dependent)
Color	Colorless to pale yellow (component-dependent)	Color	Colorless to pale yellow (component-dependent)
Odor	Odorless or faint preservative odor	Odor	Odorless or faint preservative odor
Odor threshold	Not determined	Odor threshold	Not determined
pH	6.0–7.2 (buffers); <2 for stopping solution	pH	6.0–7.2 (buffers); <2 for stopping solution
Melting/freezing point	≈ 0 °C (aqueous mixtures)	Melting/freezing point	≈ 0 °C (aqueous mixtures)
Initial boiling point and range	≈ 100 °C	Initial boiling point and range	≈ 100 °C
Flash point	Not applicable (non-flammable aqueous mixtures)	Flash point	Not applicable (non-flammable aqueous mixtures)
Evaporation rate	Comparable to water	Evaporation rate	Comparable to water
Flammability (solid, gas)	Not applicable	Flammability (solid, gas)	Not applicable
Upper/lower flammability or explosive limits	Not applicable	Upper/lower flammability or explosive limits	Not applicable

Component	d) Appearance	e) Odor	f) pH
Inhibin A Calibrators (A-F)	Lyophilized, white	Odorless	7.2
Inhibin A Controls (I-II)	Lyophilized, white	Odorless	7.2
Inhibin A Ab. Plate	Plastic, clear plate	Odorless	N/A
Inhibin A Biotin Conjugate Concentrate	Liquid, clear	Odorless	7.2
Inhibin A Assay Buffer A	Liquid, clear	Odorless	7.2
Inhibin A Assay Buffer B	Liquid, clear	Odorless	6.0
Inhibin A Biotin Conjugate Diluent	Liquid, colorless	Odorless	7.2
Inhibin A Streptavidin Conjugate Ready-to-Use	Liquid, yellow	Odorless	6.2
TMB Solution	Liquid, colorless	Odorless	4.0
Stopping Solution	Liquid, colorless	Odorless	1.2
Wash Concentrate A	Liquid, colorless	Odorless	7.2

**For all components**

g) odor threshold	no data available
h) melting point / freezing point	no data available
i) initial boiling point and boiling range	no data available
j) flash point	no data available
k) evaporation rate	no data available
l) flammability (solid, gas)	no data available
m) upper/lower flammability or explosive limits	no data available
n) vapor pressure	no data available
o) vapor density	no data available
p) relative density	no data available
q) solubility(ies)	no data available
r) partition coefficient: n-octanol / water;	no data available

**9.2 Other information:**

No additional physicochemical information relevant to safe use.

**Section 10: STABILITY AND REACTIVITY**

- 10.1 Reactivity: Stable under normal laboratory conditions.
- 10.2 Chemical stability: Stable at 2–8 °C within stated shelf life.
- 10.3 Possibility of hazardous reactions: None known under normal use; avoid mixing with incompatible materials.
- 10.4 Conditions to avoid: Freezing, excessive heat, drying out; aerosol generation.
- 10.5 Incompatible materials: Strong acids and bases; strong oxidising agents; reactive metals (may form metal azides with azide-containing residues).
- 10.6 Hazardous decomposition products: Thermal decomposition may release oxides of sulfur, nitrogen, and carbon.

**Section 11: TOXICOLOGICAL INFORMATION**

**11.1 Information on hazard classes as defined in Regulation (EC) No 1272/2008**

- Acute toxicity (oral/dermal/inhalation): Not classified for the mixture. Sodium azide and isothiazolinones are present below classification thresholds.

- Skin corrosion/irritation: Not classified; prolonged or repeated contact may cause transient irritation.
- Serious eye damage/irritation: Not classified; splashes may cause transient irritation. The stopping solution is corrosive at component level but below classification threshold at mixture level.
- Respiratory or skin sensitization: Contains isothiazolinones (CMIT/MIT) which may cause skin sensitization in susceptible individuals (EUH208).
- Germ cell mutagenicity: No data indicating mutagenic effects for the mixture.
- Carcinogenicity: No components are classified as carcinogenic by IARC, NTP, EU CLP.
- Reproductive toxicity: No data indicating reproductive effects for the mixture.
- STOT – single exposure: Not classified.
- STOT – repeated exposure: Not classified.
- Aspiration hazard: Not applicable (aqueous mixtures of low viscosity and low hydrocarbon content).

### 11.2 Information on other hazards

No evidence of endocrine disrupting properties is available for this mixture. The product is intended for professional laboratory use; exposure is expected to be minimal under recommended handling conditions.

## Section 12: ECOLOGICAL INFORMATION

**12.1 Toxicity:** This mixture is not classified as hazardous to the environment. Contains very small quantities of sodium azide and isothiazolinones which are toxic to aquatic organisms at higher concentrations. Due to the low volume and dilution used in diagnostic testing, environmental exposure is minimal.

**12.2 Persistence and degradability:** Components are primarily biodegradable. Small amounts of preservatives may persist in aquatic environments.

**12.3 Bioaccumulative potential:** No data available for the mixture. Sodium azide has low bioaccumulation potential.

**12.4 Mobility in soil:** Expected to be mobile in soil due to water solubility of components.

**12.5 Results of PBT and vPvB assessment:** The mixture does not contain any substances identified as PBT or vPvB according to Annex XIII of REACH.

**12.6 Endocrine disrupting properties:** None known or expected based on available data.

**12.7 Other adverse effects:** Avoid uncontrolled release to the environment. Flush small quantities with excess water.

## Section 13: DISPOSAL CONSIDERATIONS

### 13.1 Waste treatment methods:

Dispose of contents and containers in accordance with local, regional, national, and international regulations. Small quantities may be diluted with large volumes of water and discharged to drain after neutralization. Do not dispose of sodium azide solutions via metal plumbing due to potential formation of explosive metal azides. Contaminated packaging: Rinse thoroughly before recycling or disposal as laboratory waste. Follow all applicable regulations under EU Directive 2008/98/EC, U.S. EPA (40 CFR Part 261), and other local environmental authorities.

**Section 14: TRANSPORT INFORMATION**

Transportation of this product is not regulated under ICAO, IMDG, US DOT, European ADR or Canadian TDG, because it is in a very small quantity, the product benefits from a total exemption from the ADR regulation.

**14.1 UN Number:**

Not applicable (mixture not classified as dangerous for transport).

**14.2 UN proper shipping name:**

Not applicable.

**14.3 Transport hazard class(es):**

Not regulated as hazardous under ADR/RID/IMDG/IATA/ICAO or U.S. DOT.

**14.4 Packing group:**

Not applicable.

**14.5 Environmental hazards:**

Not classified as environmentally hazardous for transport.

**14.6 Special precautions for user:**

Handle as laboratory reagent; avoid leaks or spills. Ensure containers remain tightly closed during transport.

**14.7 Transport in bulk according to Annex II of MARPOL 73/78 and the IBC Code:**

This product is not intended for bulk transport under the terms of Annex II of MARPOL or the International Bulk Chemical (IBC) Code.

The mixture is supplied exclusively in sealed primary containers as part of small-volume in vitro diagnostic reagent kits, not as bulk liquid cargo.

Therefore, no ship type assignment or pollution category applies.

**Section 15: REGULATORY INFORMATION****15.1 Safety, health and environmental regulations/legislation specific for the substance or mixture****European Union / EEA**

- Prepared in accordance with Regulation (EC) No 1907/2006 (REACH) as amended by Regulation (EU) 2020/878, and classified under Regulation (EC) No 1272/2008 (CLP).

- Waste management per Directive 2008/98/EC.
- Animal-derived components comply with Regulation (EC) No 999/2001 and Commission Decision 2007/453/EC.
- This product is for research use only (RUO) and is not subject to EU IVDR (2017/746) or CE marking requirements.

**United Kingdom**

- Complies with UK REACH and GB CLP Regulations (The REACH etc. (Amendment etc.) (EU Exit) Regulations 2019).

**United States**

- Complies with the OSHA Hazard Communication Standard (29 CFR 1910.1200) (HazCom 2012).
- Components are listed or exempt on the U.S. TSCA Inventory.
- Waste disposal is subject to EPA Resource Conservation and Recovery Act (40 CFR Parts 260–273).
- Product contains no infectious or biohazardous material and is not regulated as an infectious substance under the U.S. DOT Hazardous Materials Regulations (49 CFR 173).

**Canada**

- Complies with Hazardous Products Regulations (SOR/2015-17) aligned with WHMIS 2015 (GHS Rev. 5).
- Components are listed or exempt under the Domestic Substances List (DSL) or Non-Domestic Substances List (NDSL).

**Other jurisdictions**

- Switzerland – ChemO Art. 54 (accepts EU SDS format)
- Turkey – KKDİK Reg. 30105 and SEA Reg. 28848 (REACH/CLP equivalent)
- Gulf Cooperation Council (GCC) / Saudi Arabia – Implementing GHS Rev. 6; English SDS accepted
- China (PRC) – GB/T 16483-2008 and GB 30000 series
- Japan – ISHL and GHS (JIS Z 7253:2019)
- Australia – WHS Regulations and SDS Code of Practice 2011; listed or exempt on AICS
- Korea – K-REACH and KECI inventory compliant

**Label Elements (EU CLP)**

- No hazard pictogram or signal word required
- EUH208 – Contains CMIT/MIT (3:1) and sodium azide. May produce an allergic reaction.

**Authorizations and Restrictions**

- None of the substances in this mixture are subject to REACH Annex XIV (authorization) or Annex XVII (restriction).
- Substances of Very High Concern (SVHC)
- No SVHC present  $\geq 0.1$  % (w/w).

**Chemical Inventories**

All intentionally added components are listed or exempt on the following national or regional inventories: EU REACH, UK REACH, US TSCA, Canada DSL/NDSL, Australia AICS, China IECSC, Japan ENCS, Korea KECI, Philippines PICCS, and New Zealand NZIoC.

**Bovine/Equine Material Compliance**

All animal-derived materials originate from OIE-classified negligible BSE-risk herds and comply with Regulation (EC) No 999/2001.

**15.2 Chemical safety assessment**

A chemical safety assessment has not been performed for this mixture. The product is supplied in small volumes for professional in vitro diagnostic use only, and potential exposure is minimal under normal laboratory conditions.

**Section 16: OTHER INFORMATION**

*This information is based on our present knowledge. However, this shall not constitute a guarantee for any specific product features and shall not establish a legally valid contractual relationship.*

**16.1 Abbreviations and acronyms**

ADR – European Agreement concerning the International Carriage of Dangerous Goods by Road

ACGIH – American Conference of Governmental Industrial Hygienists

BSE – Bovine Spongiform Encephalopathy

CAS – Chemical Abstracts Service

CLP – Classification, Labelling and Packaging Regulation

DNEL – Derived No-Effect Level

DOT – U.S. Department of Transportation

GHS – Globally Harmonised System

IATA – International Air Transport Association

IMDG – International Maritime Dangerous Goods Code

NIOSH – U.S. National Institute for Occupational Safety and Health

OEL – Occupational Exposure Limit

OSHA – U.S. Occupational Safety and Health Administration

PBT – Persistent, Bioaccumulative and Toxic

PNEC – Predicted No Effect Concentration

REACH – Registration, Evaluation, Authorisation and Restriction of Chemicals

SVHC – Substance of Very High Concern

TSE – Transmissible Spongiform Encephalopathy

vPvB – Very Persistent and Very Bioaccumulative

**16.2 Training advice**

Product intended for professional use only. Laboratory personnel should receive training in safe handling of diagnostic reagents and chemical hygiene.

**16.3 Further information**

The information herein is believed to be correct at the time of publication and is provided in good faith. It describes safety requirements but does not constitute a product specification or guarantee.

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