

# AGGRECETIN REAGENT (100mg) SAFETY DATA SHEET (SDS)

*Prepared in accordance with:*

- OSHA Hazard Communication Standard (29 CFR 1910.1200, HazCom 2012)
- Regulation (EC) No 1272/2008 (CLP)
- Regulation (EC) No 1907/2006 (REACH)
- UK REACH (post-Brexit)
- GHS (Globally Harmonized System of Classification and Labelling of Chemicals)
- TSCA (U.S. Toxic Substances Control Act) requirements, where applicable
- California Proposition 65 (where applicable)

*This SDS also supports compliance with:*

- Regulation (EU) 2017/746 (IVDR)
- ISO 13485:2016 Quality Management Systems
- FDA Quality Management System Regulation (21 CFR Part 820)

## SECTION 1: Identification of the Substance and Supplier

Product Name: AggRecetin Reagent (Ristocetin)

Catalog Number:

**REF** 101241

Product Contents:

- AggRecetin Reagent, 1 vial (100 mg)

Relevant Identified Use:

**IVD** In vitro diagnostic reagent used in light transmission platelet aggregation testing.

**AN ISO 13485 REGISTERED COMPANY**

155 Gibraltar Road, Horsham, PA 19044 USA

Telephone Worldwide: +1 215-441-4000 / Telephone USA: +1-800-257-3282 / Fax Worldwide: +1 215-443-8820

Website: [www.biodatacorp.com](http://www.biodatacorp.com) / Email: [customer.service@biodatacorp.com](mailto:customer.service@biodatacorp.com)

Manufacturer / Supplier: Bio/Data Corporation  
Address: 155 Gibraltar Road, Horsham, PA 19044 USA  
Telephone: +1 215-441-4000  
Website: [www.biodatacorp.com](http://www.biodatacorp.com)  
Email: customer.service@biodatacorp.com

Emergency Contact: +1 215-441-4000 (Available Monday–Friday, 8:30 AM – 5:00 PM EST)

## SECTION 2: Hazard Identification

Classification (EC 1272/2008):

Not classified as hazardous.

Label Elements (CLP/GHS):

No hazard pictogram, signal word, hazard, or precautionary statements required.

Other Hazards:

Not considered hazardous under IVDR or CLP regulations. Use according to standard laboratory and professional practice.

OSHA Classification (29 CFR 1910.1200 / HazCom 2012):

This product is not classified as hazardous under OSHA Hazard Communication Standard (HazCom 2012).

WHMIS Classification (Canada, 2015):

Not classified as hazardous under WHMIS 2015.

## SECTION 3: Composition/Information on Ingredients

Substance	CAS No.	EC No.	Concentration	Classification
Ristocetin A Sulphate	11140-99-1	215-770-5	100 mg per vial	Not classified

## SECTION 4: First Aid Measures

General Advice: Not expected to pose a hazard. Use standard laboratory precautions.

Inhalation: Not an expected route of exposure.

Skin Contact: Wash with water. Seek medical attention if irritation persists.

Eye Contact: Rinse cautiously with water. Seek medical attention if needed.

Ingestion: Rinse mouth. Seek medical attention if symptoms occur.

## **SECTION 5: Fire-Fighting Measures**

Suitable Extinguishing Media: Water spray, CO<sub>2</sub>, dry chemical, or foam.

Special Hazards: Non-flammable.

Protective Equipment: Standard chemical fire PPE.

## **SECTION 6: Accidental Release Measures**

Personal Precautions: Wear gloves, eye protection.

Cleanup Methods: Absorb with inert material. Dispose in accordance with local regulations.

Environmental Precautions: Avoid runoff and prevent discharge into waterways.

## **SECTION 7: Handling and Storage**

Handling:

Use per standard laboratory practice. Avoid contamination. Reconstitute the reagent using an appropriate diluent such as AggRecetin Diluent or TRIS Buffered Saline (as specified in the IFU).

Storage Conditions:

No temp protection required during shipment.

Upon receipt, store at 2 – 8°C in original packaging.

Once reconstituted, the reagent is stable for 7 days in tightly capped vial at 2 – 8°C.

Avoid light and repeated freeze-thaw cycles.

## **SECTION 8: Exposure Controls / Personal Protection**

Occupational Exposure Limits:

No occupational exposure limits are established for the components.

Engineering Controls:

Use in a well-ventilated area or within a biosafety cabinet as appropriate.

Personal Protective Equipment (PPE):

Eye/Face Protection: Safety goggles or face shield

Skin Protection: Laboratory gloves (e.g., nitrile or latex)

Body Protection: Standard laboratory coat

Respiratory Protection: Not required under normal use conditions

Hygiene Measures:

Wash hands thoroughly after handling.

Avoid ingestion, inhalation, or contact with eyes and skin.

## **SECTION 9: Physical and Chemical Properties**

Physical State:

- Reagent: Powder
- Reconstituted Reagent: Liquid

Appearance:

- Reagent: White to yellow powder
- Reconstituted Reagent: Clear, colorless to slightly amber solution

Odor: Odorless

pH (reconstituted): 5.0 – 6.0

Melting/Freezing Point: Not applicable

Boiling Point: Not applicable

Flash Point: Not applicable

Solubility: Soluble in water

Vapor Pressure: Not applicable

Specific Gravity: Similar to water

## **SECTION 10: Stability and Reactivity**

Reactivity: No known reactivity hazards under normal conditions.

Chemical Stability: Stable under recommended storage and handling conditions.

Possibility of Hazardous Reactions: No hazardous polymerization or decomposition expected.

Conditions to Avoid: Excessive heat, light exposure, or repeated freeze-thaw cycles.

Incompatible Materials: Strong oxidizing agents

Hazardous Decomposition Products: None under normal use

## **SECTION 11: Toxicological Information**

Acute Toxicity: Not classified as acutely toxic; very low systemic toxicity expected at use concentrations.

Skin Corrosion/Irritation: Not expected to be irritating

Serious Eye Damage/Irritation: May cause mild mechanical irritation

Sensitization: No data available; not expected

Carcinogenicity / Mutagenicity / Reproductive Toxicity: Not classified based on available data

STOT – Single/Repeated Exposure: No data indicating specific target organ toxicity

## **SECTION 12: Ecological Information**

Ecotoxicity: Not expected to pose a significant environmental hazard

Persistence and Degradability: Expected to be biodegradable

Bioaccumulation Potential: Not expected to bioaccumulate

Mobility in Soil: High water solubility; likely to move with water

Other Adverse Effects: None known

## **SECTION 13: Disposal Considerations**

Waste Treatment Methods:

Dispose of unused reagent and contaminated packaging in accordance with local, regional, or national regulations.

Do not dispose of via sewage or regular waste.

Follow procedures for laboratory chemical disposal.

## **SECTION 14: Transport Information**

UN Number: Not regulated

Proper Shipping Name: Not regulated

Transport Hazard Class: Not applicable

Packing Group: Not applicable

Environmental Hazards: None

Special Precautions: None

ADR/RID/IMDG/IATA Status: Not regulated for ground, air, or sea

## **SECTION 15: Regulatory Information**

EU IVDR: CE-marked under Regulation (EU) 2017/746.

EU REACH: Exempt per Article 2(5).

EU CLP: Not classified as hazardous under Regulation (EC) 1272/2008.

UK REACH: Exempt; UK Responsible Person required for distribution within the United Kingdom.

U.S. FDA / OSHA: Exempt from OSHA Hazard Communication Standard (29 CFR 1910.1200(b)(6)(viii)); complies with FDA Quality Management System Regulation (21 CFR Part 820).

U.S. TSCA (EPA): This product is exempt from TSCA inventory requirements as it is manufactured and distributed solely for FDA-regulated in vitro diagnostic use.

ISO / UKAS: This SDS supports documentation requirements for ISO 15189 and ISO 17025 accredited laboratories. UKAS accreditation does not apply directly to SDS content.

California Proposition 65: This product contains no chemicals known to the State of California to cause cancer, birth defects, or other reproductive harm.

## **SECTION 16: Other Information**

Preparation Date: April 2023

Document Number: SDS-101241 Rev AA

Revision Level and Date: Revision AA, December 2025

Prepared by: Regulatory Affairs & Quality Assurance Department

Bio/Data Corporation

Email: [customer.service@biodatacorp.com](mailto:customer.service@biodatacorp.com)

Telephone: +1 215-441-4000

### QMS Cross-Reference:

This SDS is supported by internal, controlled documentation including Material Specifications (MS), Process Specifications (PS), Formulation Procedures (FP), and the Instruction for Use (IFU) for the product. Refer to the current versions of these documents maintained under Bio/Data Corporation's Quality Management System.

DISCLAIMER: THIS PRODUCT IS INTENDED FOR IN VITRO DIAGNOSTIC USE ONLY. NOT FOR INJECTION OR INGESTION. THE INFORMATION PROVIDED IN THIS SAFETY DATA SHEET IS BELIEVED TO BE ACCURATE AND RELIABLE AS OF THE DATE OF PUBLICATION. HOWEVER, NO WARRANTY, EXPRESS OR IMPLIED, IS MADE REGARDING ITS COMPLETENESS OR ACCURACY IN SPECIFIC APPLICATIONS. THE END USER IS SOLELY RESPONSIBLE FOR ENSURING THAT THE PRODUCT IS USED IN ACCORDANCE WITH ALL APPLICABLE LOCAL LAWS, REGULATIONS, AND INDUSTRY GUIDELINES.