Blood and Transplant
Effective date: 22/09/2023

According to EC Regulations 1907/2006 (REACH), 1272/2008 (CLP) & 2015/830

Safety Data Sheet

1 SECTION 1: Identification of the substance/mixture and company

1.1 Product identifier

Human antisera reagent.

Product Code	Product Name	UDI-DI
PN061	AB serum	5055232400376
PN062	AB Serum for use in antibody Quantification	5055232400475

CAS No. : Mixtures EC No. : Mixtures

1.2 Relevant identified uses of the substance or mixture and uses advised against

Identified uses: Human antisera reagent for in-vitro diagnostic use only.

Uses advised against: Anything other than the above

1.3 Details of the supplier of the safety data sheet

NHSBT Reagents, 14 Estuary Banks, Speke, Liverpool, L24 8RB. United Kingdom,

Telephone: 0151 268 7157

Fax: 0151 268 7156

Email: reagents@nhsbt.nhs.uk

1.4 Emergency telephone number

NHSBT Reagents customer services 0151 268 7157 (Monday to Friday 9am to 5pm. Calls will

be forwarded to an answering machine outside of these hours) Further information available from: www.blood.co.uk/reagents

Languages spoken: English

2 **SECTION 2:** Hazards identification

2.1 Classification of the substance or mixture

Not classified as hazardous for supply/use according to Regulation (EC) 1272/2008 (CLP).

2.2 Label elements

No Hazard Pictogram(s), Signal Word(s), Hazard Statement(s) or Precautionary Statement(s) have been assigned according to Regulation (EC) 1272/2008 (CLP).

2.3 Other hazards

Sodium azide is very toxic by inhalation, in contact with skin and if swallowed.

Contact with acid liberates very toxic gas.

Heating may cause explosion.

Readily absorbed through skin.

Avoid contact with metals, may react with lead and copper plumbing to form highly explosive metal azides.

However, the concentrations used within this reagent are not associated with any hazards

The donations used in this product are of human origin. They have been tested and found negative for the mandatory microbiological tests required by the UK guidelines for blood transfusion services during donation testing. No known test methods can offer assurances that products derived from human blood will not transmit infectious diseases. This device should be treated as clinical material for its use and disposal.

Blood and Transplant
Effective date: 22/09/2023

According to EC Regulations 1907/2006 (REACH), 1272/2008 (CLP) & 2015/830

3 **SECTION 3: Composition/information on ingredients**

3.1 Substances

N/A

3.2 Mixtures

Donated human plasma which has been pooled.

Component	Approximate Concentration	CAS number	EC number	Hazard
	(%)			
Sodium azide	<0.1	26628-22- 8	247-852-1	Acute Tox. 2; H300: Fatal if swallowed
		0		Aquatic Acute 1;
				H400: Very toxic to aquatic life
				Aquatic Chronic 1;
				H410: Very toxic to
				aquatic life with long
				lasting effects
K ₂ EDTA	<1	25102-12-	2001-94-7	Acute Tox. 4; H332:
		9		harmful if inhaled.
				STOT RE 2; H373:
				May cause damage
				to organs through
				prolonged or
N. EDTA		04.00.0	000 570 0	repeated exposure.
Na ₄ EDTA	<1	64-02-8	200-573-9	Acute Tox. 4; H302:
				Harmful if
				swallowed
				Eye Dam. 1; H318:
				causes serious eye
				damage

None of the products at the concentration used within this reagent are associated with any hazards.

4 SECTION 4: First aid measures

4.1 Description of first aid measures

Eye contact: Flush eyes with water for at least 15 minutes while holding eyelids open. Remove contact lenses, if present and easy to do. Continue rinsing. If eye irritation persists, get medical advice/attention.

Skin contact: Wash affected skin with soap and water. Remove contaminated clothing and wash clothing before reuse. If irritation (redness, rash, blistering) develops, get medical attention.

Ingestion: Rinse mouth. Give plenty of water to drink. Do not give anything by mouth to an unconscious person. Get medical advice/attention if you feel unwell.

Inhalation: Not applicable.

4.2 Most important symptoms and effects, both acute and delayed All known important symptoms are described in Section 3.

Blood and Transplant
Effective date: 22/09/2023

According to EC Regulations 1907/2006 (REACH), 1272/2008 (CLP) & 2015/830

4.3 Indication of any immediate medical attention and special treatment needed No special treatment indicated. Treat Symptomatically

5 SECTION 5: Firefighting measures

5.1 Extinguishing media

Non-Flammable. Use extinguishing media appropriate to the surrounding conditions.

- 5.2 Special hazards arising from the substance or mixture None known.
- 5.3 Advice for firefighters

Fight fire with normal precautions from a reasonable distance.

6 SECTION 6: Accidental release measures

- 6.1 Personal precautions, protective equipment and emergency procedures Use personal protective equipment during removal of spillages.
- 6.2 Environmental precautions

Control spillage in accordance with local regulations.

6.3 Methods and material for containment and cleaning up

Spray spillage with appropriate virucidal detergent and absorb in suitable inert material, wash spill site after material pick up is complete.

6.4 Reference to other sections
See sections 8 and 13 for disposal information

7 SECTION 7: Handling and storage

7.1 Precautions for safe handling

When handling product use personal protective equipment as required. Wash hands thoroughly after handling. Contaminated clothing should be thoroughly cleaned. Protect from contamination.

- 7.2 Conditions for safe storage, including any incompatibilities

 Store the reagent at +2°C to +8°C in the original container/packaging.
- 7.3 Specific end use(s) See section 1.2

8 SECTION 8: Exposure controls/personal protection

8.1 Control parameters

None of the products at the concentration used within this reagent are associated with any hazards. No monitoring procedures required

8.2 Exposure controls

When handling product use personal protective equipment as required. Avoid contact. Keep good hygiene and housekeeping measures.

9 **SECTION 9: Physical and chemical properties**

- 9.1 Information on basic physical and chemical properties
 - (a) Appearance: Human serum diluted in a clear liquid
 - (b) Odour: Odourless
 - (c) Odour threshold: Not established

INF111/2 – Safety Data Sheet NHSBT

AB Serum

According to EC Regulations 1907/2006 (REACH), 1272/2008 (CLP) & 2015/830

Blood and Transplant
Effective date: 22/09/2023

- (d) pH: Not established
- (e) Melting point/freezing point: Not established
- (f) Initial boiling point and boiling range: Not established
- (g) Flash point: Not established
- (h) Evaporation rate: Not established
- (i) Flammability (solid, gas): Not established
- (j) Upper/lower flammability or explosive limits: Not applicable
- (k) Vapour pressure: Not established
- (I) Vapour density: Not established
- (m) Relative density: Not established
- (n) Solubility(ies): Miscible with water
- (o) Partition coefficient: n-octanol/water: Not established
- (p) Auto-ignition temperature: Not established
- (q) Decomposition temperature: Not established
- (r) Viscosity: Not established
- (s) Explosive properties Not explosive
- (t) Oxidising properties: Not oxidising
- 9.2 Other information

No further information available

10 SECTION 10: Stability and reactivity

10.1 Reactivity

Stable under normal conditions

10.2 Chemical stability

Stable for stated expiry date of the product when stored between +2°C and +8°C.

Do not freeze

Protect from extremes of temperature.

Do not use if reagent exhibits turbidity or if a precipitate, gel or particles are present.

10.3 Possibility of hazardous reactions

None under normal processing

10.4 Conditions to avoid

Keep away from heat and direct sunlight.

10.5 Incompatible materials

There are no known materials that could react with this product to produce a hazardous situation.

10.6 Hazardous decomposition products

There are no known hazardous decomposition products produced, but as the product is derived from human blood it should be treated as potentially infectious and be disposed of appropriately.

11 **SECTION 11:** Toxicological information

11.1 Information on toxicological effects

(a) acute toxicity;

Ingestion: based on available data, the classification criteria are not met for this mixture.

- (b) skin corrosion/irritation; based on available data, the classification criteria are not met for this mixture.
- (c) serious eye damage/irritation; based on available data, the classification criteria are not met for this mixture.

NHS
Blood and Transplant
Effective date: 22/09/2023

According to EC Regulations 1907/2006 (REACH), 1272/2008 (CLP) & 2015/830

- (d) respiratory or skin sensitisation; based on available data, the classification criteria are not met for this mixture.
- (e) germ cell mutagenicity; based on available data, the classification criteria are not met for this mixture.
- (f) carcinogenicity; based on available data, the classification criteria are not met for this mixture.
- (g) reproductive toxicity; based on available data, the classification criteria are not met for this mixture.
- (h) STOT-single exposure; based on available data, the classification criteria are not met for this mixture.
- (i) STOT-repeated exposure; based on available data, the classification criteria are not met for this mixture.
- (j) aspiration hazard: based on available data, the classification criteria are not met for this mixture.

See section 3 for individual component hazard classification.

12 SECTION 12: Ecological Information

12.1 Toxicity

Based on available data, the classification criteria are not met for this mixture. (see section 3 for individual component hazard classification)

12.2 Persistence and degradability

Not established. Predicted to be unlikely.

12.3 Bio accumulative potential

Not established. Predicted to be unlikely.

12.4 Mobility in soil

This product has high mobility in soil. Miscible with water.

12.5 Results of PBT and vPvB assessment

Not classified as PBT or vPvB. None of the substances in this product fulfil the criteria for being regarded as a PBT or vPvB substance.

12.6 Other adverse effects

None known.

13 SECTION 13: Disposal Considerations

13.1 Waste treatment methods

The product and any contaminated packaging should be disposed in accordance with local state or national legislation.

14 SECTION 14: Transport Information

Not classified according to the United Nations 'Recommendations on the Transport of Dangerous Goods.'

14.1 UN number

Not assigned ADR/RID, IMDG or IATA/ICAO numbers.

14.2 UN proper shipping name

Not assigned ADR/RID, IMDG or IATA/ICAO numbers.

14.3 Transport hazard class(es)

Not assigned ADR/RID, IMDG or IATA/ICAO numbers.

Blood and Transplant
Effective date: 22/09/2023

According to EC Regulations 1907/2006 (REACH), 1272/2008 (CLP) & 2015/830

14.4 Packing group

Not assigned ADR/RID, IMDG or IATA/ICAO numbers.

14.5 Environmental hazards

Not assigned ADR/RID, IMDG or IATA/ICAO numbers.

14.6 Special precautions for user

See section 2

14.7 Transport in bulk according to Annex II of MARPOL and the IBC Code Not Applicable

15 SECTION 15: Regulatory information

- 15.1 Safety, health and environmental regulations/legislation specific for the substance or mixture This product does not contain substances subject to EU authorisation or restriction of use. This safety datasheet was prepared in accordance with the requirements of EC regulation 1907/2006 (REACH) (Article 32, Annex II) and 1272/2008 (CLP) & 2015/830
- 15.2 Chemical safety assessment None.

16 SECTION 16: Other information

Amendments from the previous version of this SDS are in purple text

	LEGEND		
ADR	Agreement concerning the International Carriage of Dangerous Goods by Road		
CAS	Chemical abstracts service		
CLP	Classification, labelling and packaging of substances and mixtures		
EC	European Commission		
EDTA	Ethylenediaminetetraacetic acid		
IATA	International Air Transport Association		
IBC	International Code for the Construction and Equipment of Ships carrying Dangerous Chemicals in Bulk		
ICAO	International Civil Aviation Organization		
IMDG	International Maritime Dangerous Goods		
LC50	Lethal Concentration to 50% of a test population		
LD50	Lethal Dose to 50% of a test population (Median Lethal Dose)		
MARPOL	International Convention for the Prevention of Pollution from Ships		
OECD	Organisation for Economic Cooperation and Development		
PBT	Persistent, Bio accumulative and Toxic		
RID	Regulations concerning the International Carriage of Dangerous Goods by Rail		
SDS	Safety data sheet		
STOT	Specific target organ toxicity		
vPvB	Very persistent and very bio accumulative		

Blood and Transplant
Effective date: 22/09/2023

According to EC Regulations 1907/2006 (REACH), 1272/2008 (CLP) & 2015/830

Disclaimers

Customers are urged to ensure that the product is entirely suitable for their own purpose. It is the customers' responsibility to ensure that a suitable and sufficient assessment of the risks created by the use of the product is undertaken. The use of the reagent and the interpretation of results must be conducted by professionally trained and qualified personnel in accordance with the requirements of the country where the reagent is in use.

Information contained in this publication or as otherwise supplied to Users is believed to be accurate and is given in good faith, but it is for the Users to satisfy themselves of the suitability of the product for their own particular purpose. NHSBT reagents gives no warranty as to the fitness of the product for any particular purpose and any implied warranty or condition (statutory or otherwise) is excluded except to the extent that exclusion is prevented by law. NHSBT reagents accepts no liability for loss or damage (other than that arising from death or personal injury caused by defective product, if proved), resulting from reliance on this information. Freedom under Patents, Copyright and Designs cannot be assumed.

Training advice

Consideration should be given to the work procedures involved and the potential extent of exposure as they may determine whether a higher level of protection is required.

Date of First Issue 27/08/2008