

SAFFTY DATA SHEET

		SAFELT DALA SHEEL
SECTI	ON 1. IDENTIFICAT	ION
1.1. P	roduct Identifier(s)	
N	ame:	Carcinoembryonic Antigen Next Generation (CEA-Next Gen) AccuBind® ELISA Test System
	escription:	AccuBind® ELISA Microwells
-	ode:	4625-300
-	haracteristics:	Microplate Enzyme Immunoassay, Colorimetric
		the substance or mixture and uses advised against
		carcinoembryonic antigen concentration in human serum by a microplate enzyme immunoassay,
	olorimetric.	ly. Not for internal or external use in humans or animals.
	etails of the supplier of th	
	lanufacturer/Importer:	Manufacturer
	ame or commercial name:	
	egistered office:	100 North Pointe Drive, Lake Forest, California 92630, USA
	elephone number:	+1.949.951.2665
	ax number:	+1.949.951.3539
E	mail:	info@monobind.com
F	DA Established	
	egistration number:	2020726
	mergency telephone num	
+'	1.949.951.2665 (Hours: 8 a	m-5 pm PST, Monday-Friday)
N 2.2. La N 2.3. O	ON 2. HAZARD(S, lassification of the substa one abel elements one ther hazards one) IDENTIFICATION ince or mixture
SECTI		ION/INFORMATION ON INGREDIENTS
	ubstances and/or Mixture	
A	Il concentrations of potentia	Ily hazardous substances or mixtures are below the specific concentration limits and M-factors for
	azardous identification. As	preparations, the product components are not classified as hazardous. The following substance value and is listed with its concentration level. At this concentration level, the substance is not
		r definitions for all risk and hazards classifications.
3.1.1.	CEA-Next Gen Calibrato	
0	N/A	
3.1.2.	CEA-Next Gen Enzyme	Reagent
	N/A	
3.1.3.	Streptavidin Coated Plat	te
	N/A	
3.1.4.	Substrate A	
<u> </u>	N/A	
3.1.5.	Substrate B	
216	N/A Wash Solution Concenti	rato
3.1.6.	N/A	
3.1.7.	Stop Solution	

Chemical Name	Identification	Hazard Code Risk Phrase	Hazard Class Category Code	Hazard Statement	Concentration
Hydrochloric Acid	CAS: - EC: 231-595-7	C; R34 Xi; R37	Skin Corr. 1B STOT SE 3	H314 H335	< 5 %

SECTION 4. FIRST-AID MEASURES

4.1. Description of first aid measures

Immediately rinse with soap and plenty of water. Use personal protective working aids. General instructions: If inhaled: Transport the affected person into the open air. If there are respiratory complaints, oxygen must be administered. If irritation persists, seek medical advice. In case of skin contact: Wash contacted area with soap and water. Remove contaminated clothing. If irritation occurs, seek medical advice. In case of contact with eyes: Rinse with a stream of water for at least 15 minutes. Thorough rinsing must be ensured by opening the eyelids. If irritation occurs, seek medical advice. Do NOT induce vomiting. If conscious, rinse the mouth and administer a large amount of water to If ingested: dilute the substance. In the case of unconsciousness, never administer anything orally. If irritation occurs, seek medical advice.

Effective Date: 2019-09-17 Rev. 4

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	Mon	obind Inc.	
ISO	13485 0	Certified Co	ompany

- 4.2. Most important symptoms and effects, both acute and delayed No data available
- 4.3. Indication of any immediate medical attention and special treatment needed No data available

SECTION 5. FIRE-FIGHTING MEASURES

- 5.1. Extinguishing media
 - Carbon dioxide, dry powder, foam, water
- 5.2. Special hazards arising from the substance or mixture None
- 5.3. Advice for firefighters

Wear appropriate personal protective equipment and clothing. Wear self-contained breathing apparatus, if necessary.

SECTION 6. ACCIDENTAL RELEASE MEASURES

- 6.1. Personal precautions, protective equipment and emergency procedures
- Avoid contact with skin and eyes. Wear suitable personal protective clothing.
- 6.2. Environmental precautions
- Avoid penetration into sewerage systems, surface and ground water. Avoid soil pollution.
- 6.3. Methods and material for containment and cleaning up Cover with suitable absorbing material. After removing the substance, rinse the spot of spilling thoroughly with water and soap. Dispose of waste according to all federal, state, and local regulations.
- 6.4. Reference to other sections

See Section 8 for personal protective equipment. See Section 13 for appropriate disposal methods.

SECTION 7. HANDLING AND STORAGE

- 7.1. Precautions for safe handling
 - Avoid spills. Avoid contact with skin, eyes and clothing. Use suitable protective means to work with the substance. Use in a well-ventilated area. Follow good manufacturing practices when using product. Do not drink, smoke, or eat in work areas.

7.2. Conditions for safe storage, including any incompatibilities

7.2.1. Kit and unopened components:

- Store at temperatures between + 2 and + 8 °C in a dry and dark place until expiration date.
- 7.2.1. Opened components:
- Opened reagents are stable for sixty (60) days when stored at 2-8 °C.
- 7.2.2. For prepared reagents (see product insert):
 - Diluted wash buffer should be stored at room temperature (2-30 °C) for up to 60 days.

Working substrate solution should be stored at 2-8 °C and is stable for one (1) year.

7.3. Specific end uses

Product procedure should be performed by a skilled individual or trained professional for in vitro diagnostic use only.

EXPOSURE CONTROL/PERSONAL PROTECTION **SECTION 8.**

8.1. Control parameters

No substances with occupational exposure limits.

8.2. Exposure controls

8.	.2.1.	Eye/face protection:	Safety glasses or goggles with side shields recommended
8.	.2.2.	Skin protection:	Compatible protective gloves recommended. Wash hands after properly removing and disposing of gloves.
		Other skin protection:	Laboratory coats are recommended.
8.	.2.3.	Respiratory protection:	No respiratory protection is required. Use product in rooms enabling good ventilation. If local exhaustion is necessary, general (forced) exhaustion is recommended.
8.	.2.4.	Thermal hazards:	None

SECTION 9. PHYSICAL AND CHEMICAL PROPERTIES

9.1. Information on basic physical and chemical properties

Annearance 9.1.1.

5.1.1.	Appearance.	
	Physical state (at 2	20 °C)
	Liquid:	Calibrators, Enzyme Reagent, Wash Solution Concentrate, Substrate Solutions, Stop Solution
	Solid:	Microtiter strips
	Colour	
	Straw:	Calibrators
	Red:	Enzyme Reagent
	Clear:	Stop, Substrates, Wash
9.1.2.	Odour:	Odourless
9.1.3.	Odour threshold:	Not applicable
9.1.4.	pH value:	Stop solution: < 3
		Calibrators: 7.4 \pm 0.2
		Enzyme: 7.3 ± 0.2
		Streptavidin Wells: 7.5 ± 0.2
		Wash Solution Concentrate: 8.8 ± 0.2
		Substrate Reagent A: 3.2 ± 0.2
		Substrate Reagent B: 5.0 ± 0.2



- 9.1.5. Melting point/freezing point: Not determined
- 9.1.6. Initial boiling point/ boiling range: Not determined
- 9.1.7. Flash point: Not applicable
- 9.1.8. Evaporation rate: Not determined
- 9.1.9. Flammability (solid, gas): Not flammable
- 9.1.10. Upper/lower flammability or explosive limits: Not applicable
- 9.1.11. Vapour pressure: Not determined
- 9.1.12. Vapour density: Not determined
- Not determined 9.1.13. Relative density:
- 9.1.14. Solubility: Water soluble
- Partition coefficient: n-octanol/water: Not determined 9.1.15.
- Not applicable 9.1.16. Auto-ignition temperature:
- 9.1.17. Decomposition temperature: Not determined Not determined
- 9.1.18. Viscosity: None
- 9.1.19. Explosive properties:
- 9.1.20. Oxidising properties: Not determined
- 9.2. Other information

None

SECTION 10. STABILITY AND REACTIVITY

- 10.1.Reactivity
 - No known reactivity hazards associated with product

10.2.Chemical stability

- Stable under recommended storage conditions
- 10.3.Possibility of hazardous reactions
- No hazardous polymerization
- 10.4.Conditions to avoid
- Excessive heat and light
- 10.5.Incompatible materials
- Acids
- 10.6.Hazardous decomposition products Not determined

TOXICOLOGICAL INFORMATION: SECTION 11.

- 11.1.Information on toxicological effects
- 11.1.1. Acute toxicity: Not determined
- 11.1.2. Skin corrosion/irritation: Not determined
- 11.1.3. Serious eye damage/irritation: Not determined
- 11.1.4. Respiratory or skin sensitisation: Not determined
- 11.1.5. Germ cell mutagenicity: Not determined No component of this product present at levels $\geq 0.1\%$ is identified as probable, possible or
- 11.1.6. Carcinogenicity:
- confirmed human carcinogen by NTP (National Toxicology Program), IARC (International
- Agency for Research on Cancer), or OSHA (Occupational Safety & Health Administration)
- 11.1.7. Reproductive toxicity: Not determined
- STOT-single exposure: Not determined 11.1.8. Not determined
- 11.1.9. STOT-repeated exposure: 11.1.10. Aspiration hazard: Not determined
- 11.1.11. Information on likely routes of exposure:
- If ingested: No known health effects No known health effects
- If inhaled:
- If contact with skin: No known health effects
- No known health effects If contact with eyes:

11.1.12. Symptoms related to the physical, chemical, and toxicological characteristics: None after short or long-term exposure

SECTION 12. ECOLOGICAL INFORMATION

12.1.Toxicity

- Not determined.
- 12.2.Persistence and degradability
- Not determined
- 12.3.Bioaccumulative potential
- Not determined
- 12.4.Mobility in soil
- Not determined
- 12.5.Results of PBT and vPvB assessment Not determined
- 12.6.Other adverse affects
- Not determined

DISPOSAL CONSIDERATIONS SECTION 13. 13.1.Waste treatment methods

Effective Date: 2019-09-17 Rev. 3



All waste disposals must be carried out in accordance with federal, state, and local legislation and administrative regulations. A licensed professional waste disposal service should be utilized to dispose of material and packaging.

TRANSPORT INFORMATION SECTION 14

14.1.UN number

- Not available 14.2.UN proper shipping name
- Not available

14.3.Transport hazard class(es)

- Not available
- 14.4.Packing group Not available

14.5.Environmental hazards

Overland transport (ADR/RID):	None
Water transport (ADN/IMDG):	None
Air transport (ICAO/IATA):	None

14.6.Special precautions for user

None

14.7.Transport in bulk according to Annex II of MARPOL73/78 and the IBC Code Not applicable

SECTION 15. REGULATORY INFORMATION

15.1.Safety, health and environmental regulations/legislation specific for the substance or mixture

SARA Reporting Requirements: None

All components in product preparations are lifted on the US Toxic Substances Control Act inventory of TSCA chemicals or are exempt from listing.

This safety data sheet has been prepared to comply with the requirements of Annex II, European Community Regulation No. 1907/2006 REACH (Registration, Evaluation, Authorisation and Restriction of Chemicals) and OSHA (Occupational Safety & Health Administration) 1910.1200, Appendix D.

15.2.Chemical safety assessment

None

SECTION 16. OTHER INFORMATION

Revision 3 (2019-Sep-17): Updated to include component pH value details

Revision 2 (2015-MAY-05): updated to comply with requirements of Annex II, European Community Regulation No. 1907/2006 (REACH) and OSHA 1910.1200, Appendix D Revision 1 (2010-DEC-01): updated to 16 point format

Revision 0 (2005-DEC-22): Initial creation

Hazard Statements		Hazard Class and Category Codes	
H314	Causes severe skin burns and eye damage	Skin Corr.	Skin Corrosion/Irritation
H335	May cause respiratory irritation	STOT SE 3	Specific Target Organ toxicity - Single Exposure
Hazard	Codes	Risk Phrases	
С	Corrosive	R34	Causes burns
Xi	Irritant	R37	Irritating to respiratory system

The material safety data sheet contains data necessary to ensure safety and health and environmental protection in working with chemical substances. This product is a chemical substance and can be solely used by persons with chemical education at their own risk. Monobind kits are designed for biomedical research. The manufacturer has no responsibility for damage caused by unsuitable use and by disrespecting the enclosed working instructions. The above-stated information cannot be considered as complete and must be understood to be only a methodical instruction.

	DOCUMENT HIS	TORY			
PREPARED BY: APPROVED BY: REVISION: 3	DEPT: Records Administration	VERIFIED BY: EFFECTIVE DAT DCO: 1361	AShatola E: 2019-SEP-17	DEPT: QA	

