

SAFETY DATA SHEET

SALETT DATA SHEET							
SECTION 1. IDENTIFICAT	ION						
1.1. Product Identifier(s)							
Name:	High Sensitivity C-Reactive Protein (hsCRP) AccuLite® CLIA Test System						
Description:	AccuLite® CLIA Microwells						
Code:	3175-300						
Characteristics:	Microplate Enzyme Immunoassay, Chemiluminescence						
1.2. Relevant identified uses of the substance or mixture and uses advised against							
Quantitative determination of C-Reactive protein concentration in human serum or plasma by a microplate enzyme							
immunoassay, Chemiluminescence.							
For in vitro diagnostic use only. Not for internal or external use in humans or animals.							
1.3. Details of the supplier of th							
Manufacturer/Importer:	Manufacturer Manakird Inc.						
Name or commercial name:							
Registered office:	100 North Pointe Drive, Lake Forest, California 92630, USA						
Telephone number: Fax number:	+1.949.951.2665 +1.949.951.3539						
Email:	+1.949.951.5559 info@monobind.com						
FDA Established							
Registration number:	2020726						
1.4. Emergency telephone num							
	im-5 pm PST, Monday-Friday)						
 2.1. Classification of the substa None 2.2. Label elements None 2.3. Other hazards None 	ince or mixture						
 3.1. Substances and/or Mixture All concentrations of potentia hazardous identification. As p 3.1.1. hsCRP Calibrators (A-F N/A 	ally hazardous substances or mixtures are below the specific concentration limits and M-factors for preparations, the product components are not classified as hazardous.						
N/A							
3.1.3. Serum Diluent N/A							
3.1.4. Light Reaction Wells N/A							
3.1.5. Signal A N/A							
3.1.6. Signal B N/A							
3.1.7. Wash Solution Concent N/A	rate						
SECTION 4. FIRST-AID MEASURES							
4.1. Description of first aid mea							
General instructions: Immediately rinse with soap and plenty of water. Use personal protective working aids.							

 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.

 If ingested:
 Do NOT induce vomiting. If conscious, rinse the mouth and administer a large amount of water to

ed: Do NOT induce vomiting. If conscious, rinse the mouth and administer a large amount of water to dilute the substance. In the case of unconsciousness, never administer anything orally. If irritation occurs, seek medical advice.

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.

ACCIDENTAL RELEASE MEASURES **SECTION 6.**

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.

HANDLING AND STORAGE SECTION 7.

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

Kit and unopened components: 7.2.1.

Store at temperatures between + 2 and + 8 °C in a dry and dark place until expiration date. Avoid extended exposure to heat and light.

- 7.2.1. Opened components:
- 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 Signal solution should be stored at 2-8 °C and is stable for thirty-six (36) hours. Diluted serum diluent solution should be stored at 2-8 °C and is stable for up to 60 days.

7.3. Specific end uses

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

SECTION 8. EXPOSURE CONTROL/PERSONAL PROTECTION

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

9.1.1. Appearance

9.1.2. 9.1.3. 9.1.4.

Appearance.				
Physical state (at 20 °C)				
Liquid:	Calibrators, Tracer Reagent, Wash Solution Concentrate, Signal Solutions			
Solid:	Microtiter strips			
Colour				
Straw:	Calibrators			
Green:	Tracer Reagent			
Orange:	Diluent			
Clear:	Signals, Wash			
Odour:	Odourless			
Odour threshold:	Not applicable			
pH value:	Stop solution: < 3			
	Calibrators: 7.4 ± 0.2			
	Tracer Reagent: 7.3 ± 0.2			
	Microtiter strips: 7.5 ± 0.2			
	Wash Solution Concentrate: 8.8 ± 0.2			
	Substrate Reagent A: 9.0 ± 0.2			
	Substrate Reagent B: 5.0 ± 0.2			
Melting point/freezing point: Not determined				

- 9.1.5. Melting point/freezing point: Not determined
- 9.1.6. Initial boiling point/ boiling range: Not determined

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- 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
- 9.1.13. Relative density: Not determined
- 9.1.14. Solubility: Water soluble
- Partition coefficient: n-octanol/water: 9.1.15. Not determined Auto-ignition temperature: Not applicable 9.1.16.
- Decomposition temperature: Not determined
- 9.1.17. 9.1.18. Not determined Viscosity:
- None
- 9.1.19. Explosive properties: Oxidising properties: 9.1.20. 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

SECTION 11. **TOXICOLOGICAL INFORMATION:**

- 11.1.Information on toxicological effects
- 11.1.1. Acute toxicity: Not determined
- 11 1 2 Skin corrosion/irritation: Not determined
- Serious eye damage/irritation: Not determined 11.1.3.
- 11.1.4. Respiratory or skin sensitisation: Not determined
- 11.1.5. Germ cell mutagenicity: Not determined 11.1.6.
 - Carcinogenicity: No component of this product present at levels $\geq 0.1\%$ is identified as probable, possible or 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: 11.1.8. Not determined
- 11.1.9. STOT-repeated exposure: Not determined
- 11.1.10. Aspiration hazard: Not determined
- 11.1.11. Information on likely routes of exposure: If ingested: No known health effects
- If inhaled: No known health effects
- If contact with skin: No known health effects
- If contact with eyes: No known health effects

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

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.

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SECTION 14. 14.1.UN number	TRANSPORT IN	FORMATION				
Not available						
14.2.UN proper shipping name						
Not available						
14.3.Transport hazard class(es)						
Not available						
14.4.Packing group	0					
Not available						
14.5.Environmental hazards						
	port (ADR/RID):	None				
Water transpor	t (ADN/IMDG):	None				
Air transport (IC		None				
14.6.Special precan None	utions for user					
14.7.Transport in bulk according to Annex II of MARPOL73/78 and the IBC Code						
Not applicable	J					
SECTION 15.	REGULATORY	INFORMATION				
15.1.Safety, health and environmental regulations/legislation specific for the substance or mixture SARA Reporting Requirements: None						
TSCA		in product preparations are lifted on the US Toxic Substances Control Act inventory of 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 (2011-SEP-22): updated to 16 point format Revision 0 (2005-DEC-22): Initial creation

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 HISTORY							
PREPARED BY: APPROVED BY: Frollaps	_DEPT: Records Administration _DEPT: Administration	VERIFIED BY: AShatok EFFECTIVE DATE: 2019-SEP-17	DEPT: QA				
REVISION: 3		DCO: 1361					

