

# SAFETY DATA SHEET

SAFELT DATA SHEET							
SECTION 1. IDENTIFICATION							
1.1. Pro	duct Identifier(s)						
Nar	me:	Progesterone AccuLite® CLIA Test System					
Des	scription:	AccuLite® CLIA Microwells					
Coc	de:	4875-300					
	aracteristics:	Microplate Enzyme Immunoassay, Chemiluminescence					
		the substance or mixture and uses advised against					
Quantitative determination of progesterone concentration in human serum or plasma by a microplate enzyme immunoassay, Chemiluminescence.							
		y. Not for internal or external use in humans or animals.					
1.3. Det	ails of the supplier of the						
	nufacturer/Importer:	Manufacturer					
	me or commercial name:						
	gistered office:	100 North Pointe Drive, Lake Forest, California 92630, USA					
	ephone number:	+1.949.951.2665					
	number:	+1.949.951.3539					
Ema		info@monobind.com					
	A Established						
	gistration number:	2020726					
	ergency telephone num						
+1.9	949.951.2665 (Hours: 8 ai	m-5 pm PST, Monday-Friday)					
Nor	ssification of the substance bel elements	IDENTIFICATION nce or mixture					
2.3. Oth	er hazards						
Nor	ne						
SECTIO		ION/INFORMATION ON INGREDIENTS					
	ostances and/or Mixture						
		Ily hazardous substances or mixtures are below the specific concentration limits and M-factors for					
		preparations, the product components are not classified as hazardous.					
3.1.1.	Progesterone Calibrator	s (A-G)					
	N/A						
3.1.2.	Progesterone Tracer Re	agent					
	N/A						
3.1.3.	Progesterone Biotin Rea	agent					
	N/A						
3.1.4.	Light Reaction Wells						
o	N/A						
3.1.5.	Signal Reagent						
	N/A						
3.1.6.	Wash Solution Concentr	ate					
	N/A						

## SECTION 4. FIRST-AID MEASURES

# 4.1. Description of first aid measures

Booonprion of mot and	
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
	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

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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.

### 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

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.

### 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. None
- 8.2.4. Thermal hazards:

#### PHYSICAL AND CHEMICAL PROPERTIES **SECTION 9.**

### 9.1. Information on basic physical and chemical properties

9.1.1. Appearance:

Physical state (at )	20 °C)
Liquid:	Calibrators, Tracer Reagent, Biotin Reagent, Wash Solution Concentrate, Signal Solution
Solid:	Microtiter strips
Colour	
Straw:	Calibrators
Yellow:	Tracer Reagent, Biotin Reagent
Clear:	Signal, Wash
Odour:	Odourless
Odour threshold:	Not applicable
pH value:	Stop solution: < 3
	Calibrators: $7.4 \pm 0.2$
	Tracer Reagent: 7.3 ± 0.2
	Biotin: $7.3 \pm 0.2$
	Microtiter strips: $7.5 \pm 0.2$
	Wash Solution Concentrate: 8.8 ± 0.2
	Substrate Reagent A: 9.0 ± 0.2
	Substrate Reagent B: 5.0 ± 0.2
Melting point/freez	ing point: Not determined
Initial boiling point	/ boiling range: Not determined
Flash point:	Not applicable
	Solid: Colour Straw: Yellow: Clear: Odour: Odour threshold: pH value: Melting point/freez Initial boiling point/

- 9.1.8. Evaporation rate: Not determined
- Not flammable 9.1.9. Flammability (solid, gas):

- 9.1.10. Upper/lower flammability or explosive limits:
- 9.1.11. Vapour pressure: Not determined
- 9.1.12. Vapour density: Not determined
- Relative density: 9.1.13. Not determined
- 9.1.14. Solubility: Water soluble
- 9.1.15. Partition coefficient: n-octanol/water: Not determined
- 9.1.16. Auto-ignition temperature: Not applicable
- 9.1.17. Decomposition temperature: Not determined
- 9.1.18. Viscosity: Not determined
- 9.1.19. Explosive properties: None Not determined
- Oxidising properties: 9.1.20.
- 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
- 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
- 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

Not applicable

- 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:
  - No known health effects If contact with skin:
  - 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

#### **ECOLOGICAL INFORMATION SECTION 12.**

### 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

#### SECTION 13. **DISPOSAL CONSIDERATIONS**

### 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.

#### SECTION 14. TRANSPORT INFORMATION 14.1.UN number

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#### MSDS 4875-300

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

TSCA All components in product preparations are lifted on the US Toxic Substances Control Act inventory of 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 2 (2019-Sep-17): Updated to include component pH value details

Revision 1 (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 0 (2013-MAY-01): 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								
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APPROVE	DBY: Frallap	DEPT: Administration	EFFECTIVE DATE: 2019-SEP-17					
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