

In accordance with Regulation (EC) No 2020/878 Revision date: 02/11/2022 Version number: 05

1. I	dentification of the	e substance or preparation and the company/ undertaking
1.1 <u>Product Identifier</u>		
Product Name: N	Mast C.E.M.O. Selec	tatab™
Product Code: N	MS60	
1.2 <u>Relevant identified uses of the substance</u>	e or mixture and use	es advised against
Recommended uses: 1	n vitro diagnostic pro	duct; Laboratory chemical
Uses advised against: N	No information availal	ble
1.3 Details of supplier of the product and s	<u>afety data sheet</u>	
1.4 Emergency Contact informatio	3 7277 (8am - 5pm GMT Monday to Friday)	
		2. Hazards identification
2.1 <u>Classification of the substance or mixtu</u> CLP Classification – Regulation (F		
CLA Classification – Regulation (E	Physical hazards:	Based on available data, there are no physical hazards
	Heath hazards:	Based on available data, there are no health hazards
Envir	onmental hazards:	Based on available data, there are no environmental hazards
2.2 Label elements		
	Pictogram:	None
	Signal word:	None
H	Hazard statements:	None
	ionary statements:	None
2.3 <u>Other hazards</u>	• • • • • • • • • • • • • • • • • • •	No information available



In accordance with Regulation (EC) No 2020/878 Revision date: 02/11/2022 Version number: 05

3. Composition/information on ingredients

Composition: A multi-component freeze-dried tablet containing amphotericin B in an inert carrier.

Hazardous ingredients:

Component	CAS-No.	EC-No.	Concentration	CLP Classification – 1272/2008/EC	Classification to – 67/548/EEC
Amphotericin B	1397-89-3	215-742-2	<2%	Skin Irritant Cat. 2; Eye Irritant Cat. 2; Specific target organ toxicity - single exposure Cat. 3. H315 - Causes skin irritation H319 - Causes serious eye irritation. H335 - May cause respiratory irritation	Xi – Irritant. R36/37/38 - May be irritating to eyes, respiratory system and skin.
Clindamycin	21462-39-5	244-398-6	<2%	None	None
Trimethoprim lactate	23256-42-0	245-533-1	<1%	None	None

4. First-aid measures

4.1 Description of First Aid measures

General advice:	Consult a physician. Show this safety data sheet to the doctor in attendance.
Eye contact: Rinse thoroughly with plenty of water for 10 to 15 minutes, also under the eyelids. Obtain medical attention if irr persists.	
Skin Contact:	Wash off skin thoroughly with soap and plenty of water. Obtain medical attention if irritation persists.
Ingestion:	Rinse mouth out with plenty of water. Obtain medical attention if symptoms occur.
Inhalation:	Move person to fresh air. Obtain medical attention immediately if symptoms occur.
	and offered which courts and deleged

4.2 Most important symptoms and effects, both acute and delayed

No information available.

4.3 Indicate any immediate medical attention and special treatment needed

No information available.

	5. Fire fighting measures		
5.1	Extinguishing medium		
	Suitable extinguishing media: Use water spray, CO2, foam or dry powder as the extinguisher medium.		
	Extinguishing media which must not be used for safety reasons: No information available		
5.2	2 Special hazards arising from the substance or mixture		
	Combustible material. Thermal decomposition may lead to release of irritating gases and vapours.		
5.3	Advice for firefighters		
	Wear suitable self contained breathing apparatus for fire fighting if necessary.		
5.4	Additional information		
	No data available.		



In accordance with Regulation (EC) No 2020/878 Revision date: 02/11/2022 Version number: 05

	6. Accidental release measures			
6.1	5.1 <u>Personal precautions, protective equipment and emergency procedures</u>			
		Ensure adequate ventilation.	Vear appropriate protective equipment. Avoid dust formation.	
6.2	6.2 <u>Environmental precautions</u>			
		Should not be released into the	e environment.	
6.3	6.3 <u>Methods and materials for containment and cleaning up</u>			
		Sweep up or vacuum up spilla	ge in suitable container for disposal. Avoid dust formation.	
6.4	6.4 <u>Further information</u>			
		No data available.		
	7. Handling and storage			
7.1	Precautions for safe handlin	ng		
	A	Avoid contact with eyes, skin an	nd clothing. Avoid ingestion and inhalation. Avoid dust formation.	
7.2	Conditions for safe storage,	including any incompatabilit	ies	
	Store at 2°C to 8°C. Keep tightly closed in the container provided. Protect from direct sunlight and moisture.			
7.3	Specific end use(s)			
	This product is for laboratory use only and should only be used by suitably trained laboratory personnel.			
		8. Exp	osure controls and personal protection	
8.1	Control parameters			
	Components with workspace	e control parameters: This pro	oduct has no occupational exposure limits.	
8.2	Exposure controls	1 1		
	Engineering controls:	No engineering protection rec	juired.	
	Personal protective measures:	Body protection:	Wear standard microbiology laboratory coat.	
	1	Eye/face protection:	Safety glasses with side shields conforming to EN 166.	
		Skin and hand protection:	Handle with suitable gloves conforming to EN 374. Product may be handled with forceps.	
		Respiratory protection:	If required use a nuisance type particle respirator type P1 EU EN 143.	
	General hygiene measures.	Handle in accordance with ac		

Handle in accordance with good laboratory practice. Wash hands before breaks and at the end of the working day. General hygiene measures:



In accordance with Regulation (EC) No 2020/878 Revision date: 02/11/2022 Version number: 05

9. Physical and Chemical properties

9.1 Information on basic physical and chemical properties

Physical appearance: Colour: Odour: Odour threshold: Melting point/freezing point: Initial boiling point/range Flammability (solid, gas): Explosive limits: Flash point: Auto-ignition temperature: Decomposition temperature: pH value: Partition coefficient (n-octanol/water): Vapour pressure: Vapour density: Relative density: Solubility in water: Solubility in other solvents: Viscosity:	Lyophilised tablet. White to off-white colour. No detectable odour. No data available. No data available.
Solubility in other solvents:	No data available.
Explosive properties Oxidising properties: Evaporation rate:	This product is not explosive in normal circumstances. No data available.

9.2 Other information

No data available.

10. Stability and reactivity

10.1 <u>Reactivity</u>

No known on information available.

10.2 Chemical stability

Stable under normal conditions.

10.3 Possibility of hazardous reactions

No data available.

10.4 Conditions to avoid

Hygroscopic. Avoid exposure to sunlight.

10.5 Incompatible materials

None known.

10.6 Hazardous decomposition products

None under normal use conditions.

11. Toxicological information

11.1 Information on toxicological effects

Overall product information: This product does not present an acute toxicity based on known or supplied information.



In accordance with Regulation (EC) No 2020/878 Revision date: 02/11/2022 Version number: 05

Acute toxicity:	For amphotericin B - LD ₅₀ Oral - rat - >5,000 mg/kg For further information see RTECS: BU2625000	
	For clindamycin - LD_{50} Oral - rat - 2,193 mg/kg, LD_{50} Intramuscular - rat - 273 mg/kg LD_{50} Intraperitoneal - rat - 745 mg/kg, LD_{50} Subcutaneous - rat - 2,618 mg/kg For further information see RTECS: no data available.	
	For trimethoprim – no data available. For further information see RTECS: no data available.	
Skin corrosion/irritation:	No data available.	
Serious eye damage/ eye irritation:	No data available.	
Respiratory or skin sensitisation:	May cause allergic respiratory and skin reactions.	
Germ cell mutagenicity:	No data available.	
Carcinogenicity:	IARC: No component of this product is present at levels greater than or equal to 0.1% is identified as probable, possible or confirmed human carcinogen by IARC.	
Reproductive toxicity:	No data available.	
Specific target organ toxicity – single exposure:	No data available.	
Specific target organ toxicity - repeated exposure:	No data available.	
Aspiration hazard:	No data available.	
Potential health effects:	Inhalation - May be harmful if inhaled. May cause respiratory tract irritation. Ingestion - May be harmful if swallowed. Skin - May be harmful if absorbed through skin. May cause skin irritation. Eyes - May cause eye irritation.	
	To the best of our knowledge, the chemical, physical, and toxicological properties have not been thoroughly investigated.	

11.2 <u>Toxicological data for the components</u>

Assess endocrine disrupting properties for human health. This product does not contain any known or suspected endocrine disruptors.

12. Ecological information

12.1 Toxicity

This product is not hazardous to the environment or that are not degradable in waste water treatment plants.

12.2 Persistence and degradability

Expected to be biodegradable.

12.3 Bioaccumulative potential

No data available.

12.4 Mobility in soil

No data available.

12.5 Results of PBT and vPvB assessment

None known.

12.6 Endocrine disrupting properties: requiring you to give product information on adverse effects on the environment caused by endocrine-disrupting properties

This product does not contain any known or suspected endocrine disruptors.



In accordance with Regulation (EC) No 2020/878 Revision date: 02/11/2022 Version number: 05

12.7 Other adverse effects

No data available

	13. Disposal considerations
Disposal considerations	 Dispose of in accordance with local and national regulations. Dispose of contaminated waste, e.g. used plates, according to local microbiological rules.
	14. Transport information
14.1 <u>UN Number</u>	
ADR/RID: IMDG: IATA:	Not dangerous for shipping
14.2 <u>UN proper shipping name</u>	
ADR/RID: IMDG: IATA:	Not dangerous for shipping
14.3 <u>Transport hazard class(es)</u>	
ADR/RID: IMDG: IATA:	Not dangerous for shipping
14.4 Packaging group	
ADR/RID: IMDG: IATA:	Not dangerous for shipping
14.5 Environmental hazards	
ADR/RID: IMDG: IATA:	None known.
14.6 Special precautions for user	
	No data available
14.7 Maritime transport in bulk a	according to IMO instruments No data available
	15. Regulatory information
15.1 Safety, health and environm	ental regulations/legislation specific for the substance or mixture
	No data available.
15.2 Chemical safety assessment	

15.2 Chemical safety assessment

Chemical safety assessment reports are not required for mixtures/IVD products.

16. Other information		
Original origination date:	17/05/2001	
Reason for change to document:	Updated in accordance with Regulation (EC) No 2020/878	
	The above information is believed to be correct but does not purport to be all inclusive and shall be used as a guide only. The information in this document is based on the present state of our knowledge and is applicable to the product with regard to appropriate safety precautions. It does not represent any guarantee of the properties of the product and is designed only as a guide for safe handling, use, processing, storage, transportation, disposal and release and is not to be considered a warranty or quality specification. Mast Group Ltd. shall not be held liable for any damage resulting from handling or from contact with the above product. See our website at www.mast.grp.com and/or the reverse side of our invoice for additional terms and conditions of sale.	