

In accordance with Regulation (EC) No 2020/878 Revision date: 03/11/2022 Version number: 03

1. Identification of the substance or preparation and the company/ undertaking		
1.1 Product Identifier		
Product Name: Mast Legionella	Mast Legionella Selectavial [™] (MWY)	
Product Code: SV36		
1.2 Relevant identified uses of the substance or mixtu	re and uses advised against	
Recommended uses: In vitro diagnostic	In vitro diagnostic product; Laboratory chemical	
Uses advised against: No information a	No information available	
1.3 Details of supplier of the product and safety data s	heet	
Manufacturer/supplier: Mast Group Ltd., Mast House, Derby Road, Bootle, Merseyside, UK. L20 1EA Telephone: +44 (0) 151 933 7277 Email: uksales@mastgrp.com Feldstraße 20 23858 Reinfeld Germany Telephone: +49 4533 20 07 34 Email: mast@mast-diagnostica.de Web: www.mast-group.com 1.4 Emergency Contact information Web : 493 7277 (8am - 5pm GMT Monday to Friday) EU contact: Telephone: +44 (0) 151 933 7277 (8am - 5pm GMT Monday to Friday) 2. Hazards identification		
2.1 <u>Classification of the substance or mixture</u>		
CLP Classification – Regulation (EC) No. 1272/200	8: All hazardous components present below a level that requires hazard labelling	
Physical hazard	s: Based on available data, there are no physical hazards	
Heath hazard	s: This product may cause an allergic skin reaction.	
Environmental hazard	s: Based on available data, there are no environmental hazards	
2.2 Label elements		
Pictogram	n: None	
Signal wor	d: None	
Hazard statement	s: None	
Precautionary statement	s: None	
2.3 <u>Other hazards</u>	No information available	



In accordance with Regulation (EC) No 2020/878 Revision date: 03/11/2022 Version number: 03

3. Composition/information on ingredients

Composition: A multi-component freeze-dried tablet containing polymyxin B, vancomycin, amphotericin B, bromothymol blue, bromocresol purple and glycine in an inert carrier.

Hazardous ingredients:

Component	CAS-No.	EC-No.	Concentration	CLP Classification – 1272/2008/EC	Classification to – 67/548/EEC
Vancomycin hydrochloride	1404-93-9	-	0.03%	Skin sensitization Cat. 1: H317 - may cause an allergic skin reaction	Xi, irritant. R43 – may cause sensitisation by skin contact
Polymyxin B	1405-20-5	215-774-7	<0.3%	Acute Toxicity Oral Cat. 4: H302 - harmful if swallowed.	Xn, harmful. R22 - harmful if swallowed
Amphotericin B	1397-89-3	215-742-2	<0.3%	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 – irritating to eyes, respiratory system and skin.
Bromocresol purple	115-40-2	204-087-7	0.3%	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 – irritating to eyes, respiratory system and skin.

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 irritation 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.

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 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.



In accordance with Regulation (EC) No 2020/878 Revision date: 03/11/2022 Version number: 03

5.4 Additional information

No data available.

6. Accidental release measures

6.1 <u>Personal precautions, protective equipment and emergency procedures</u>

Ensure adequate ventilation. Wear appropriate protective equipment. Avoid dust formation.

6.2 Environmental precautions

Should not be released into the environment.

6.3 Methods and materials for containment and cleaning up

Sweep up or vacuum up spillage in suitable container for disposal. Avoid dust formation.

6.4 Further information

No data available.

7. Handling and storage

7.1 Precautions for safe handling

Avoid contact with eyes, skin and clothing. Avoid ingestion and inhalation. Avoid dust formation.

7.2 Conditions for safe storage, including any incompatabilities

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. Exposure controls and personal protection

8.1 Control parameters

Components with workspace control parameters: This product has no occupational exposure limits.

8.2 Exposure controls

Engineering controls:	No engineering protection required.	
Personal protective measures:	Body protection:	Wear standard microbiology laboratory coat.
	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 good laboratory practice. Wash hands before breaks and at the end of the working day.	



In accordance with Regulation (EC) No 2020/878 Revision date: 03/11/2022 Version number: 03

9. Physical and Chemical properties

9.1 Information on basic physical and chemical properties

Physical state:	Lyophilised tablet.
Colour:	White to off-white colour.
Odour:	Odourless.
Melting point/freezing point:	No data available.
Initial boiling point/range:	No data available.
Flammability:	No data available.
Lower and upper explosion limit:	No data available.
Flash point:	No data available.
Auto-ignition temperature:	No data available.
Decomposition temperature:	No data available.
pH	No data available.
Kinetic viscosity:	No data available.
Solubility in water/other solvents:	No data available.
Partition coefficient (n-octanol/water):	No data available.
Vapour pressure:	No data available.
Density/Relative density:	No data available.
Relative vapour density:	No data available.
Decomposition temperature	No data available.
Particle characteristics:	No data available.

9.2 Other information

No data available.

10. Stability and reactivity

10.1 Reactivity

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.

 Acute toxicity:
 For vancomycin - LD₅₀ Oral - rat - >10,000 mg/kg. Also see RTECS: YW4380000

 For polymyxin B - LD₅₀ Oral - mouse - 790 mg/kg; LD₅₀ Intraperitoneal - mouse - 20.5 mg/kg; LD₅₀ Intraperitoneal - mouse - 20.5 mg/kg; LD₅₀ Intravenous - mouse - 5.4 mg/kg. Also see RTECS: TR1150000

 For amphetericin B - LD₅₀ Oral - rat - >5.000 mg/kg. Also see RTECS: RU2625000

For amphotericin B - LD_{50} Oral - rat - >5,000 mg/kg. Also see RTECS: BU2625000



In accordance with Regulation (EC) No 2020/878 Revision date: 03/11/2022 Version number: 03

Skin corrosion/irritation:	No data available.
Serious eye damage/ eye irritation:	No data available.
Respiratory or skin sensitisation:	May cause allergic respiratory reaction.
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. Causes respiratory tract irritation. Ingestion May be harmful if swallowed. Skin May be harmful if absorbed through skin. Causes skin irritation. Eyes Causes serious eye irritation.

11.2 Toxicological data for the components

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

12. Ecological information

12.1 Toxicity

No data available.

12.2 Persistence and degradability

No data available.

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.

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.



In accordance with Regulation (EC) No 2020/878 Revision date: 03/11/2022 Version number: 03

14. Transport information

14.1 UN Number

ADR/RID: IMDG: IATA: Not dangerous for shipping

14.2 UN proper shipping name

ADR/RID: IMDG: IATA: Not dangerous for shipping

14.3 Transport hazard class(es)

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 <u>Maritime transport in bulk according to IMO instruments</u> No data available

15. Regulatory information

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

No data available.

15.2 Chemical safety assessment

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

16. Other information		
Original origination date:	23/04/2009	
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.	