

Revision date: 26-Mar-2014

Version: 2.0

Page 1 of 8

1. IDENTIFICATION OF THE SUBSTANCE/MIXTURE AND THE **COMPANY/UNDERTAKING**

Product Identifier

Material Name: New Bronz MG

Trade Name:	
Synonyms:	

New Bronz Newcastle (Kimber strain) - Infectious Bronchitis vaccine, Massachusetts (M41) and Holland serotypes + Mycoplasma gallasepticum inactivated vaccine Mixture

Chemical Family:

Relevant Identified Uses of the Substance or Mixture and Uses Advised Against Intended Use: Veterinary Vaccine

Details of the Supplier of the Safety Data Sheet

Zoetis Inc. 100 Campus Drive, P.O. Box 651 Florham Park, New Jersey 07932 (USA) Rocky Mountain Poison Control Center Phone: 1-866-531-8896 Product Support/Technical Services Phone: 1-800-366-5288

Emergency telephone number: CHEMTREC (24 hours): 1-800-424-9300 Contact E-Mail: VMIPSrecords@zoetis.com Zoetis Belgium S.A. **Mercuriusstraat 20** 1930 Zaventem Belgium

Emergency telephone number: International CHEMTREC (24 hours): +1-703-527-3887

2. HAZARDS IDENTIFICATION

Appearance:

Pale yellow to reddish-white opaque liquid **Classification of the Substance or Mixture** Not classified as hazardous

EU Classification:

Australian Hazard Classification

GHS - Classification

EU Indication of danger: Not classified

Label Elements

Signal Word: Not Classified Hazard Statements: Not classified in accordance with international standards for workplace safety.

Other Hazards Short Term:

This product is an oil-adjuvanted suspension. Oil-adjuvant containing products may cause severe vasospasm following accidental injection. In the event of accidental injection, an allergic reaction may occur. If an allergic reaction occurs, the worker should be removed to the nearest emergency room and the appropriate therapy instituted. May produce slight eve irritation. Non-Hazardous Substance. Non-Dangerous Goods.

(NOHSC):

Note:

This document has been prepared in accordance with standards for workplace safety, which requires the inclusion of all known hazards of the product or its ingredients regardless of the potential risk. The precautionary statements and warning included may not apply in all cases. Your needs may vary depending upon the potential for exposure in your workplace.

3. COMPOSITION/INFORMATION ON INGREDIENTS

Hazardous

In one die of		EU		0110	0/
Ingredient	CAS Number	EU EINECS/ELINCS	EU Classification	GHS Classification	%
		List		Classification	
Gentamicin	1403-66-3	215-765-8	Not Listed	Not Listed	##
Amphotericin B	1397-89-3	215-742-2	Not Listed	Not Listed	##

Ingredient	CAS Number	EU EINECS/ELINCS List	EU Classification	GHS Classification	%
Avian Infectious Bronchitis Massachusetts Serotype	NA	Not Listed	Not Listed	Not Listed	*
Avian Infectious Bronchitis Holland Serotype	NA	Not Listed	Not Listed	Not Listed	*
Newcastle, Kimber Strain	NA	Not Listed	Not Listed	Not Listed	*
Adjuvant - Oil Emulsion	NA	Not Listed	Not Listed	Not Listed	*
Mycoplasma Gallisepticum	Not Assigned	Not Listed	Not Listed	Not Listed	*

Additional Information:

Trace * Proprietary

Ingredient(s) indicated as hazardous have been assessed under standards for workplace safety.

4. FIRST AID MEASURES

Description of First Aid Measures Eye Contact:	Flush with water while holding eyelids open for at least 15 minutes. Seek medical attention immediately.
Skin Contact:	Remove contaminated clothing. Flush area with large amounts of water. Use soap. Seek medical attention.
Ingestion:	Never give anything by mouth to an unconscious person. Wash out mouth with water. Do not induce vomiting unless directed by medical personnel. Seek medical attention immediately.
Inhalation:	Remove to fresh air and keep patient at rest. Seek medical attention immediately.
Most Important Symptoms and Effe Symptoms and Effects of Exposure: Medical Conditions Aggravated by Exposure:	cts, Both Acute and Delayed For information on potential signs and symptoms of exposure, See Section 2 - Hazards Identification and/or Section 11 - Toxicological Information. None known
Indication of the Immediate Medical	Attention and Special Treatment Needed

Indication of the Immediate Medical Attention and Special Treatment Needed Notes to Physician: None

5. FIRE-FIGHTING MEASURES

Extinguishing Media:	Extinguish fires with CO2, extinguishing powder, foam, or water.	
Special Hazards Arising from the Sul Hazardous Combustion Products:	ostance or Mixture Formation of toxic gases is possible during heating or fire.	
Fire / Explosion Hazards:	Fine particles (such as dust and mists) may fuel fires/explosions.	
Advice for Fire-Fighters During all fire fighting activities, v	vear appropriate protective equipment, including self-contained breathing apparatus.	
6.	ACCIDENTAL RELEASE MEASURES	
Personal Precautions, Protective Equ Personnel involved in clean-up s	lipment and Emergency Procedures nould wear appropriate personal protective equipment (see Section 8). Minimize exposure.	
	abeled, sealed container for disposal. Care should be taken to avoid environmental release.	
Methods and Material for Containmer Measures for Cleaning / Collecting:	nt and Cleaning Up Contain the source of the spill if it is safe to do so. Wipe up with a damp cloth and place in container for disposal. Clean contaminated surface thoroughly.	
Additional Consideration for Large Spills:	Non-essential personnel should be evacuated from affected area. Report emergency situations immediately. Clean up operations should only be undertaken by trained personnel.	
	7. HANDLING AND STORAGE	
Precautions for Safe Handling Avoid breathing vapor or mist. Avoid contact with eyes, skin and clothing. Wash thoroughly after handling. Prevent environmental releases. Use appropriate personal protective equipment. Avoid accidental injection.		
Conditions for Safe Storage, Includir Storage Conditions:	g any Incompatibilities Store at room temperature in properly labeled containers. Keep away from heat, sparks and flames.	
Specific end use(s):	No data available	
8. EXPOSI	JRE CONTROLS / PERSONAL PROTECTION	

Control Parameters

Refer to available public information for specific member state Occupational Exposure Limits.

Gentamicin

Bulgaria OEL - TWA

0.1 mg/m³

The purpose of the Occupational Exposure Band (OEB) classification system is to separate substances into different Hazard categories when the available data are sufficient to do so, but inadequate to establish an Occupational Exposure Limit (OEL). The OEB given is based upon an analysis of all currently available data; as such, this value may be subject to revision when new information becomes available.

Gentamicin

Zoetis OEB

OEB 2 (control exposure to the range of 100ug/m³ to < 1000ug/m³)

8. EXPOSURE CONTROLS / PERSONAL PROTECTION

Exposure Controls	
Engineering Controls:	Engineering controls should be used as the primary means to control exposures. Use process containment, local exhaust ventilation, or other engineering controls to maintain airborne levels within the OEB range.
Personal Protective	Refer to applicable national standards and regulations in the selection and use of personal
Equipment:	protective equipment (PPE).
Hands:	Wear impervious gloves if skin contact is possible.
Eyes:	Safety glasses or goggles
Skin:	Use protective clothing (uniforms, lab coats, disposable coveralls, etc.) in both production and laboratory areas.
Respiratory protection:	If airborne exposures are within or exceed the Occupational Exposure Band (OEB) range, wear an appropriate respirator with a protection factor sufficient to control exposures to the bottom of the OEB range.

9. PHYSICAL AND CHEMICAL PROPERTIES

Physical State:	Liquid	Color:	Pale yellow to reddish- white
Odor:	Odorless	Odor Threshold:	No data available.
Molecular Formula:	Mixture	Molecular Weight:	Mixture
Solvent Solubility:	No data available		
Water Solubility:	Miscible		
pH:	No data available.		
Melting/Freezing Point (°C):	No data available		
Boiling Point (°C):	No data available.		
Partition Coefficient: (Method, pH, E No data available	ndpoint, Value)		
Decomposition Temperature (°C):	No data available.		
Evaporation Rate (Gram/s):	No data available		
Vapor Pressure (kPa):	No data available		
Vapor Density (g/ml):	No data available		
Relative Density:	No data available		
Viscosity:	No data available		
Flammablity:			
Autoignition Temperature (So	lid) (°C):	No data available	
Flammability (Solids):		No data available	
Flash Point (Liquid) (°C):		No data available	
Upper Explosive Limits (Liqui	d) (% by Vol.):	No data available	

10. STABILITY AND REACTIVITY

Reactivity: Chemical Stability: Possibility of Hazardous Reactions Oxidizing Properties: Conditions to Avoid: Incompatible Materials:

Lower Explosive Limits (Liquid) (% by Vol.):

No data available Stable under normal conditions of use.

No data available Fine particles (such as dust and mists) may fuel fires/explosions. As a precautionary measure, keep away from strong oxidizers

No data available

10. STABILITY AND REACTIVITY

Hazardous Decomposition No data available Products:

11. TOXICOLOGICAL INFORMATION

Information on Toxicological Effects General Information:

Toxicological properties of the formulation have not been fully investigated. The antigens included in this product are non-infectious. All have been prepared from killed or inactivated preparations of microorganisms. The information included in this section describes the potential hazards of the individual ingredients.

Acute Toxicity: (Species, Route, End Point, Dose)

Amphotericin B

Oral LD50 Rat > 5000 mg/kg Para-periosteal LD50 Rat 1.6mg/kg Intraperitoneal LD50 > 5000mg/kg Rat Mouse Intravenous LD50 1.2mg/kg Mouse Intraperitoneal LD50 27.7mg/kg

Gentamicin

RatOralLD506600 mg/kgRatSubcutaneousLD50710mg/kgMouseIMLD50167 mg/kgRatIMLD50463 mg/kg

Irritation / Sensitization: (Study Type, Species, Severity)

Gentamicin

Eye Irritation Rabbit Non-irritating

Repeated Dose Toxicity: (Duration, Species, Route, Dose, End Point, Target Organ)

Amphotericin B

30 Day(s) Intravenous 37 mg/kg/day LOAEL Kidney Dog 2 Month(s) Dog Intravenous 16.5 mg/kg/day LOAEL Kidney Male reproductive system, Female reproductive system 13 Week(s) Rat Oral 2 mg/kg/day NOAEL Male reproductive system, Female reproductive system 13 Week(s) Dog Oral 1.6 mg/kg/day NOAEL

Reproduction & Development Toxicity: (Duration, Species, Route, Dose, End Point, Effect(s))

Amphotericin B

Embryo / Fetal Development Rat Oral 7.5 mg/kg/day NOAEL Not teratogenic, Fetotoxicity Embryo / Fetal Development Rabbit Oral 10 mg/kg/day NOAEL Not Teratogenic, Fetotoxicity

Gentamicin

Embryo / Fetal Development Rat Intramuscular 75 mg/kg/day LOAEL Developmental toxicity

Genetic Toxicity: (Study Type, Cell Type/Organism, Result)

11. TOXICOLOGICAL INFORMATION

Amphotericin B

Bacterial Mutagenicity (Ames)	Salmonella , E. coli	Negative	
In Vivo Micronucleus	Mouse	Negative		
In Vitro Chromosome A	berration	Chinese Hamster C	Ovary (CHO) cells	Negative

Carcinogen Status:

None of the components of this formulation are listed as a carcinogen by IARC, NTP or OSHA.

12. ECOLOGICAL INFORMATION

ntal properties have not been thoroughly investigated. Releases to the environment woided.
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13. DISPOSAL CONSIDERATIONS

Waste Treatment Methods:

Dispose of waste in accordance with all applicable laws and regulations. Member State specific and Community specific provisions must be considered. Considering the relevant known environmental and human health hazards of the material, review and implement appropriate technical and procedural waste water and waste disposal measures to prevent occupational exposure and environmental release. It is recommended that waste minimization be practiced. The best available technology should be utilized to prevent environmental releases. This may include destructive techniques for waste and wastewater.

14. TRANSPORT INFORMATION

The following refers to all modes of transportation unless specified below.

Not regulated for transport under USDOT, EUADR, IATA, or IMDG regulations.

15. REGULATORY INFORMATION

Safety, Health and Environmental Regulations/Legislation Specific for the Substance or Mixture

15. REGULATORY INFORMATION

Canada - WHMIS: Classifications WHMIS hazard class: None required

Gentamicin

CERCLA/SARA 313 Emission reporting California Proposition 65 Australia (AICS): Standard for the Uniform Scheduling for Drugs and Poisons: EU EINECS/ELINCS List	Not Listed Not Listed Present Schedule 4 215-765-8
EU EINEC3/ELINCS LIST	215-705-0
Avian Infectious Bronchitis Massachusetts Serotype CERCLA/SARA 313 Emission reporting California Proposition 65 EU EINECS/ELINCS List	Not Listed Not Listed Not Listed
Avian Infectious Bronchitis Holland Serotype CERCLA/SARA 313 Emission reporting California Proposition 65 EU EINECS/ELINCS List	Not Listed Not Listed Not Listed
Newcastle, Kimber Strain CERCLA/SARA 313 Emission reporting California Proposition 65 EU EINECS/ELINCS List	Not Listed Not Listed Not Listed
Adjuvant - Oil Emulsion CERCLA/SARA 313 Emission reporting California Proposition 65 EU EINECS/ELINCS List	Not Listed Not Listed Not Listed
Mycoplasma Gallisepticum CERCLA/SARA 313 Emission reporting California Proposition 65 EU EINECS/ELINCS List	Not Listed Not Listed Not Listed
Amphotericin B CERCLA/SARA 313 Emission reporting California Proposition 65 Australia (AICS): Standard for the Uniform Scheduling for Drugs and Poisons: EU EINECS/ELINCS List	Not Listed Not Listed Present Schedule 4 215-742-2
	210-142-2

16. OTHER INFORMATION

Data Sources:

The data contained in this MSDS may have been gathered from confidential internal sources, raw material suppliers, or from the published literature.

Reasons for Revision:	Updated Section 1 - Identification of the Substance/Preparation and the Company/Undertaking. Updated Section 2 - Hazard Identification. Updated Section 3 - Composition / Information on Ingredients. Updated Section 4 - First Aid Measures. Updated Section 6 - Accidental Release Measures. Updated Section 7 - Handling and Storage. Updated Section 8 - Exposure Controls / Personal Protection. Updated Section 9 - Physical and Chemical Properties. Updated Section 11 - Toxicology Information. Updated Section 12 - Ecological Information.
Prepared by:	Toxicology and Hazard Communication Zoetis Global Risk Management

Zoetis Inc. believes that the information contained in this Safety Data Sheet is accurate, and while it is provided in good faith, it is without warranty of any kind, expressed or implied. If data for a hazard are not included in this document there is no known information at this time.

End of Safety Data Sheet