

SECTION 1 : CHEMICAL PRODUCT AND COMPANY IDENTIFICATION

SECTION 1.1 : PRODUCT IDENTIFIER

PRODUCT NAME : ALBENDAZOLE Solution (10.0%)-All presentation Volumes

PRODUCT TYPE : MIXTURE REACH REGISTRATION : Not Available

SECTION 1.2 : RELEVANT IDENTIFIED USES, & USES ADVISED AGAINST

IDENTIFIED USES : VETERINARY; Oral Drench Anthelmintic with Ovicidal Properties for ruminants

SECTION 1.3 : DETAILS OF THE SUPPLIER OF THE SAFETY DATA SHEET

COMPANY NAME : Chanelle Pharmaceuticals Manufacturing Ltd., COMPANY ADDRESS : Dublin Road, Loughrea, Co. Galway, Rep. of Ireland

HOURS OF BUSINESS : 8.30 AM -5.00 PM (GMT)

SECTION 1.4 : EMERGENCY CONTACT INFORMATION

COMPANY TELEPHONE : +353-(0)-91-841-788 (Reception -during office hours)

COMPANY FAX : +353-(0)-91-841-303
COMPANY WEBSITE : http://www.chanellegroup.ie/

SECTION 2 : HAZARDS IDENTIFICATION

SECTION 2.1 : CLASSIFICATION OF THE SUBSTANCE/MIXTURE

CLASSIFICATION ACCORDING TO REGULATION 1272/2008/EC

Hazard Type: Category: H-Statement: Signal:

Reproductive Hazard Category 2
 Hazardous to the Aquatic Environment Category 3
 H402
 WARNING

SECTION 2.2 ; LABEL ELEMENTS

	Hazard Statements	Precautionary Statements					
	H361d: Suspected of Damaging the Unborn Child H402: Harmful to Aquatic Life	Prevention	Response	Storage	Disposal		
WARNING		P201: Obtain Special Instructions before Use	P308+P313: If Exposed or Concerned: Get Medical Advice/Attention	P405: Store Locked Up	P501: Dispose of Contents/Container in accordance with local/ regional/ national/ international regulations		
		P202: Do Not Handle until all Safety Precautions have been Read & Understood					
		P273: Avoid release to the environment					
		P280: Wear Protective Gloves/ Protective Clothing/ Eye Protection / Face Protection					

SECTION 2.3 : OTHER HAZARDS

- * Refer to Section 4 for First Aid Measures
- * Refer to Section 5 regarding Thermal Decomposition
- * Refer to Section 11 for Toxicological Data
- * Refer to Section 12 for Environmental Toxicity Data



ACCORDING TO REGULATION (EU) 2015/830 (AMENDING (EC) 1907/2006) & (EC) 1272/2008 CLP REGULATION

SECTION 3

: COMPOSITION/INFORMATION ON INGREDIENTS.

SECTION 3.1 : SUBSTANCE : NOT APPLICABLE

SECTION 3.2 : MIXTURE : INGREDIENTS ACCORDING TO 1272/2008/EC

COMPONENT	TYPE	CONC%	CAS	CLASSIFICATION	GHS & CLP LABEL (ingredient)
ALBENDAZOLE	API	10.0%	54965-21-8	Repr Tox Cat 2 WARNING Aquatic Acute Cat 1 WARNING Aquatic Chronic Cat 1 WARNING	H361: Suspected of Damaging Fertility or The Unborn Child H400: Very Toxic to Aquatic Life H400: Very Toxic to Aquatic Life with Long Lasting Effects
Citric Acid Monohydrate	Exciplent	0.50%	77-92-9	Eye Irrit. Cat 2 WARNING	H319: Causes Serious Eye Irritation
Methyl 4- Hydroxybenzoate	Excipient	0.20%	99-76-3	Aquatic Chronic Cat 3 WARNING	No Symbol H412: Harmful to Aquatic Life with Long Lasting Effects

Note: All other ingredients, not listed above, are either below reportable levels, or are not classified as hazardous according to GHS-CLP

SECTION 4 : FIRST AID MEASURES.

SECTION 4.1: Description of First Aid Measures (First Aid Section Refers to the Finished Product Mixture)

<u>General:</u> If unconscious, place person on their side in the recovery position; contact medical aid; give access to this SDS.

<u>Inhalation:</u> If coughing, dizziness, or irritation develops remove to Fresh Air; Keep Patient Warm; Seek Medical Assistance

Skin Contact: Wash affected area with mild soap & water; if irritation develops contact physician.

Eye Contact: Rinse open eye with water for several minutes; Remove Contact Lenses if present. Seek Medical Advice

Ingestion:
Rinse Mouth with water; immediately contact medical physician/assistance showing the pack information or this SDS; Keep patient in resting position; Allow patient (if conscious) to drink water; Never administer anything to patient in an unconscious state; DO

NOT INDUCE VOMITING unless directed to do so by a medical physician; if patient loses consciousness, place in recovery position.

SECTION 4.2: Most Important Symptoms and Effects (Acute & Delayed): This product contains ingredients which are known to induce

Gastrointestinal Disturbance and/or Allergic Reaction

SECTION 4.3: Indication of Any Immediate Medical Attention & Special Treatment Needed

Notes to Physician: Treat Symptomatically. No Specific Antidote.

SECTION 5

: FIREFIGHTING MEASURES.

SECTION 5.1:

Suitable Extinguishing Media: Unsuitable Extinguishing Media: EXTINGUISHING MEDIA

Dry Chemical Powder; Carbon Dioxide; Foam; Water Spray; Full Water Jet deemed unsuitable as it may spread the flame.

SECTION 5.2:

SPECIAL HAZARDS (arising from substance/mixture)

Hazardous Thermal Decomposition Products: Possibility of Carbon Oxides (COx), Nitrogen Oxides (NOx) & Sulphur Oxides (SOx).

SECTION 5.3:

ADVICE FOR FIREFIGHTERS

Wear self-contained respiratory protection; Wear Protective Clothing; Evacuate Area

ADDITIONAL INFORMATION

Collect contaminated Fire Fighting Media, Not to be discharged to drain.

Exposure to fire may cause container to rupture or if heated under confinement



SECTION 6

: ACCIDENTAL RELEASE MEASURES.

SECTION 6.1: Personal Precautions, Protective Equipment & Emergency Procedures

For Non-Emergency Personnel: No action should be taken without appropriate training, PPE, or if danger of personal risk exists.

Do not walk through, touch or attempt to contain spilled material without PPE.

Refer to Section 8 for PPE requirements. No Smoking.
Unnecessary personnel should be removed to safe area.

For Emergency Personnel: Refer to section 8 for required PPE to contain spillage.

Ensure proper containment for disposal as detailed in Section 13. Affected Area may then be cleaned with Water and Detergent.

SECTION 6.2: ENVIRONMENTAL PRECAUTIONS

Do NOT discharge spilled material to soil, sewers, waterways, surface or ground waters.

Contact relevant authorities if contamination occurs.

SECTION 6.3: MATERIALS/METHODS (for containment and clean-up)

Approach release from up-wind. Absorb spilled material with a suitable absorptive agent (E.g. Spill Kit/ Diatomaceous Earth). Label collected material appropriately and store for disposal

Referring to Section 13 for more information

SECTION 7

: HANDLING & STORAGE.

SECTION 7.1: PRECAUTIONS FOR SAFE HANDLING: (Refer to section 4 for First Aid Protocols)

Avoid contact with skin and eyes. Avoid breathing mist/vapours. DO NOT eat, drink or smoke during use of product.

Wash Hands after use. Keep container tightly closed when not in use.

FINISHED PRODUCT:

<u>For Persons administering the Product:</u> Keep all medicines away from children & pets; Children are particularly at risk from product and may prove harmful. Pets can be overly susceptible to Toxicity therefore access & exposure to this product should be restricted. In all cases of accidental exposure through ingestion, contact medical physician immediately. Avoid

contact with Skin & Eyes.

SECTION 7.2:

<u>CONDITIONS FOR SAFE STORAGE including incompatibilities;</u> Keep all medicines stored away from children, pets and animals; Finished Product does not require any special storage conditions though it is recommended to store securely,

under 30°C, dry, & well ventilated area. Avoid storing in direct sunlight.

SECTION 7.3:

SPECIFIC END USE(s); Use ONLY as detailed in Section 1.2.

SECTION 8

: EXPOSURE CONTROLS / PERSONAL PROTECTION.

SECTION 8.1:

EXPOSURE CONTROLS: None specified; No existing Data regarding OEL for finished product; Appropriate engineering controls should be in place where individuals are exposed to dust,

Vapours; mist; airborne particles; to ensure worker exposure are within/below any recommended limits.

ENVIRONMENTAL EXPOSURE CONTROLS: None specified; Emissions from ventilation or work processes should meet

requirements of environmental protection legislation.

SECTION 8.2:

CONTROL PARAMETERS; Ensure good ventilation and/or exhaust measures within the workplace.

List of PPE is as follows: Finished Product itself requires no Respiratory Protection under Normal Conditions of Use and a well ventilated environment. A (minimum) Grade Mask (EN 143) in enclosed areas recommended especially if working with bulk volumes and as a back-up to engineering controls. Gloves are required (EN 374) when handling bulk product for extended periods, wash hands thoroughly after handling product. Eye protection normally required when handling the finished product, protective eyewear (EN 166) is recommended when working with large or bulk volumes where the incidence of exposure is increased.

Respiratory Protection	Eye Protection	Hand Protection	General Protection
	(m)		
Quarter Mask Grade as per EN143	Safety Glasses Grade as per EN166	Protective Gloves Grade as per EN374	General Protective Garb and Safety Footwear appropriate to task

ACCORDING TO REGULATION (EU) 2015/830 (AMENDING (EC) 1907/2006) & (EC) 1272/2008 CLP REGULATION

SECTION 9 : PHYSICAL & CHEMICAL PROPERTIES.

SECTION 9.1: **INFORMATION ON BASIC PHYSICAL & CHEMICAL PROPERTIES**

	ACTIVE PHARMACEUTICAL	FINISHED PRODUCT
APPEARANCE: Form/Colour	White/Off-White-Yellow Powder	Pale Blue Free Flow Suspension
Odour	Mild Odour	Characteristic Scent
Odour Threshold	Not Available	Not Available
.pH	Not Available	5.0 (+/- 0.5)
Melting/Freezing Point	Melting : 208-210°C	Not Available
Boiling Point (Range)	Not Available	Not Available
Flash Point	Not Available	Not Available
Evaporation Rate	Not Available	Not Available
Flammability	Not Available	Not Available
Upper/Lower Explosive Limits	Not listed as Explosive Hazard	Not Available
Vapour Pressure	Not Available	Not Available
Vapour Density	Not Available	Not Available
Relative Density	Not Available	Not Available
Solubility	Soluble: Formic Acid (Anh) or Methylene Chloride Insoluble: Water or Alcohol	Not Available
Partition Coefficient n-octanol/water	Not Available	Not Available
Auto-Ignition Temperature	Not Available	Not Available
Decomposition Temperature	Not Available	Not Available
Viscosity	Not Available	Not Available
Explosive properties	Not Available	Not Available
Oxidising Properties	Not Available	Not Available

SECTION 9.2 : OTHER INFORMATION; No other information available; Refer to Section 10

SECTION 10 : STABILITY & REACTIVITY DATA.

SECTION 10.1 : REACTIVITY : Considered Stable

SECTION 10.2 : CHEMICAL STABILITY : Stable under Normal Conditions

SECTION 10.3 : POSSIBILITY OF HAZARDOUS REACTIONS : None Listed or Known SECTION 10.4 : No Specific Data (Refer to Section 7.2) Avoid Extremes of Light & Heat

: CONDITIONS TO AVOID **SECTION 10.5** : INCOMPATIBLE MATERIALS : No Information or NONE Known

SECTION 10.6 : HAZARDOUS DECOMPOSITION PRODUCTS : Normal Conditions NONE Known: Refer to Section 5 (fire conditions)

SECTION 11

: TOXICOLOGICAL INFORMATION

SECTION 11.1

: INFORMATION ON TOXICOLOGICAL EFFECTS

11.1.1	: 1	\cu	te	Tox	ici	ty

Ingredient	LD50	Species	Result
ALBENDAZOLE	Oral	Rat	1,300.00 to 2,400.00 mg/kg
Citric Acid Monohydrate	Oral	Rat	11,700.00 mg/kg
lethyl 4- Hydroxybenzoate	Oral	Rat	2.100.00 mg/kg

Supplementary Information: The Information provided in this Table is for individual components for which Acute Toxicity Data is available at 100% concentration of that specific ingredient. Each component listed above is present in the Finished Product at the concentrations listed in SECTION 3.2.

For the purposes of this SDS, the Acute Toxicity Estimate (ATE) for the Finished Product is estimated using the Rat LDSO data (most potent) and is estimated at >5,000.00mg/kg by calculation (actual 12,771.00 mg/kg) and accordingly, classification of the Finished Product within Section 2 is based on the GHS-CLP guidance based on this calculated figure.

Ingredients not listed in Section 3.2 are classified as non-hazardous according to Regulation EC 1272/2008 and are exempt from inclusion in the ATE calculation.

11.1.2	: Skin Irritation/Corrosion Data	: No Available Data for Finished Product: Specific Ingredients May Cause Skin Irritation.
11.1.3	: Eye Irritation/Damage Data	: No Available Data for Finished Product: Specific Ingredients May Cause Eye Irritation.
11.1.4	: Respiratory &/or Skin Sensitisation Data	: No Available Data for Finished Product: No Specific Data for Ingredients
11.1.5	: Germ Cell Mutagenicity	: Finished Product Not Classified. Contains no ingredient classified as Mutagenic
<u>11.1.6</u>	: Carcinogenicity	: Finished Product Not Classified. Contains no ingredient classified as Carcinogenic
11.1.7	: Reproductive Toxicity	: Note: ALBENDAZOLE, the Active Ingredient, is classified as Reproductive Hazard Cat 2 Positive in certain animal studies. Finished Product is classified in Section 2, in this regard as the Final Concentration of ALBENDAZOLE within the Finished Product is between ≥9.5% & ≤10.5%
11.1.8	: STOT- Single Exposure	: No Available Data for Finished Product. NOT CLASSIFIED
11.1.9	: STOT- Repeated Exposure	: No Available Data for Finished Product. NOT CLASSIFIED
11.1.10	: Aspiration Hazard	: No Available Data for Finished Product. NOT CLASSIFIED



SECTION 12.5

SECTION 12.6

SECTION 15.2

ACCORDING TO REGULATION (EU) 2015/830 (AMENDING (EC) 1907/2006) & (EC) 1272/2008 CLP REGULATION

SECTION 12 : ECOLOGICAL INFORMATION.

SECTION 12.1	: TOXICITY	: No Available Data for Finished Pro	duct:
Ingredient	LC50	Species Species	Result
ALBENDAZOL	E Crustacean	Water Flea (48Hr)	0.025 mg/L
SECTION 12.2	: PERSISTANCE AND DEGRADEA	BILITY : No Available Data	
SECTION 12.3	: BIOACCUMULATIVE POTENTIA	L : For Albendazole: Log Pow is 2.7 at	: 20°C
SECTION 12.4	: MOBILITY IN SOIL	: No Available Data	

: No Available Data

: No Available Data: Active Ingredient considered environmentally toxic

SECTION 13 : DISPOSAL CONSIDERATIONS.

: OTHER ADVERSE EFFECTS

: RESULTS OF PBT & vPvB ASSESSMENT

<u>SECTION 13.1</u> : WASTE TREATMENT METHODS; It is recommended that the product be considered hazardous waste and disposed of in accordance with Local Authority Requirements & Regulations. Waste Products should not be disposed of untreated to drain.

SECTION 14 : TRANSPORT INFORMATION.

Please Note: Section 14 applies to the Finished Product

SECTION 14.1 : UN NUMBER : UN3082

SECTION 14.2 : UN PROPER SHIPPING NAME : Environmentally Hazardous Substance, Liquid, n.o.s. (Albendazole)

 SECTION 14.3
 : TRANSPORT HAZARD CLASS (ES)
 : 9

 SECTION 14.4
 : PACKING GROUP
 : III

SECTION 14.5 : ENVIRONMENTAL HAZARDS : No (Refer to Section 12)

SECTION 14.6 : SPECIAL PRECAUTIONS FOR USER : No Data
SECTION 14.7 : TRANSPORT IN BULK : Not Applicable

SECTION 15 : OTHER REGULATORY INFORMATION.

SECTION 15.1 : SAFETY, HEALTH & ENVIRONMENTAL REGULATIONS/ LEGISLATION

(Specific for the substance/mixture) : Data Not Available : CHEMICAL SAFETY ASSESSMENT : Data Not Available

SECTION 16 : OTHER INFORMATION.

Document History : Version 1 (Created 22-July-2016)

Revised Sections : N/A

Prepared by : Materials Safety Data Officer, Chanelle Pharmaceuticals Ltd.

Abbreviations Legend : Listed in alphabetical order.

*	ADR	: "Accord-Dangereuses-Route" Dangerous Goods by Road; European Agree	IBC	; Intermediate Bulk Container
	ATE	: Acute Toxicity Estimate	IMDG	: International Maritime code for Dangerous Goods
	API	: Active Pharmaceutical Ingredient	LCSO	: Lethal Concentration 50 %
	BCF	: Blo-concentration Factor	LD50	: Lethal Dose, 50%
	CAS	: Chemical Abstracts Service	LQ	: Limited Quantity (stated in ADR-LQ)
	EINECS	: European Inventory of Existing Chemical Substances	MARPOL 73/78	: International Convention for the Prevention of Pollution from Ships
	Em5	: Emergency Schedule (Code of Emergency Labelling)	NOEC	: No Observable Effect Concentration
	GHS	: Globally Harmonised System (of Classification and Labelling of Chemicals)	N.O.S	: "Not Otherwise Specified"
	IATA	: International Air Transport Association	UN	: United Nations

IMPORTANT NOTICE: All Information contained herein is accurate to the best of our knowledge, however the above named supplier, nor any of its subsidiaries nor affiliates assumes any liability whatsoever for the accuracy, completeness, or absence of the Information contained herein. All materials contained by this document may present their own hazards and should be used with caution. Neither the Chanelle group, subsidiaries, nor affiliates guarantee that the Information contained by this SDS is complete, nor do they provide any guarantee, contract or liability that other hazards exist. Final determination of the suitability of any material or information supplied rests solely with the user. No guarantee or warranty of any kind is offered either express or implied with respect to the information and commendations contained herein. We accept no responsibility and disclaim all liability for any harmful effects that may be caused by (Incorrect) use, handling, purchase, resale or exposure to our product. Customers and users of our product must comply with all applicable health and safety laws, regulations and orders. In particular they are under an obligation to carry out a risk-assessment for the particular work place and to take adequate risk management measures in accordance with the national implementation legislation of EU directive 89/391/98 — 24.

Prepared by:

Date of Approval

22-07-20/6

Material Safety Data Officer

END OF SDS