

SAFETY DATA SHEET

(REACH regulation no. 1907/2006 and 453/2010)

### SECTION 1: IDENTIFICATION OF THE SUBSTANCE/MIXTURE AND OF THE COMPANY/UNDERTAKING

### 1.1 Product Identifier

Product name: Carporal 160 mg tablets **1.2 Relevant identified uses of the substance or mixture and uses advised against** Use: Non Steroidal Anti-Inflammatory Drug (NSAID) for veterinary use **1.3 Details of the supplier of the safety data sheet** Registered company name: Le Vet Beheer B.V. Address: Wilgenweg 7, 3421 TV Oudewater, The Netherlands Telephone: +31-(0)348-565858. Fax: +31-(0)348-565454 info@levetpharma.com www.levetpharma.com

1.4 Emergency telephone number:

+31-(0)6-10203326

**SECTION 2: HAZARDS IDENTIFICATION** 

#### **Classification of the Substance or Mixture**

### **GHS** – Classification

Reproductive Toxicity: Category 2

Specific target organ systemic toxicity (repeated exposure): Category 2

#### EU Classification:

EU Indication of danger: Harmful

EU Symbol: Xn EU Risk Phrases: R22 - Harmful if swallowed.

### Label Elements

Signal Word:	Warning
Hazard Statements:	H302 – Harmful if swallowed
	H361d - Suspected of damaging the unborn child
	H373 - May cause damage to organs through prolonged or repeated exposure
Precautionary Statements:	P201 – Obtain special instructions before use
	P202 – Do not handle until all safety precautions have been read and understood
	P280 – Wear protective gloves/protective clothing/eye protection/face protection
	P260 – Do not breathe dust/fume/gas/mist/vapors/spray
	P264 – Wash hands thoroughly after handling
	P270 – Do not eat, drink or smoke when using this product
	P308 + P313 – IF exposed or concerned: Get medical attention/advice
	P501 – Dispose of contents/container in accordance with all local and national
	regulations



Short Term:	Not an eye irritant ; Not a skin irritant (based on animal data) . There have been anecdotal reports that workers handling this material have experienced skin irritation and/or sensitivity reactions.
Long Term:	Repeat-dose studies in animals have shown a potential to cause adverse effects on gastrointestinal system, the developing fetus.
Known Clinical Effects:	Typical undesirable effects associated with NSAIDs are vomiting, soft faeces/diarrhea, faecal occult blood, loss of appetite and lethargy

### SECTION 3: COMPOSITION/INFORMATION ON INGREDIENTS

#### 3.1 Substances

No substances fulfill the criteria set forth in annex II section A of the REACH regulation (EC) no. 1907/2006

# 3.2 Mixtures

# Composition:

Ingredient	CAS-no.	Quantity
Carprofen	53716-49-7	160 mg
Lactose monohydrate	10039-26-6	Classified
Sodium starch glycolate	9063-38-1	Classified
Maize starch	9005-25-8	Classified
Purified talc	14807-96-6	Classified
Cellulose, powdered	9004-34-6	Classified
Pregelatinized maize starch	9057-07-2	Classified
Colloidal anhydrous silica	7631-86-8	Classified
Calcium behenate	3578-72-1	Classified
Yeast, deactivated	-	Classified
Chicken flavour	-	Classified

### SECTION 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	cts, Both Acute and Delayed
Symptoms and Effects	
of Exposure:	For information on potential signs and symptoms of exposure, See section 2 – Hazards Identification and/or Section 11 – Toxicological Information.
Medical Conditions	
Aggravated by Exposure:	None known

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

SECTION 5: FIREFIGHTING MEASU	IRES			
Extinguishing Media:	SMALL FIRE: Use DRY chemical powder. LARGE FIRE: Use water spray, fog or foam. Do not use water jet.			
Special hazards arising from the s	ubstance or mixture			
Hazardous Combustion				
Products:	May be combustible at high temperature. Products of combustion are carbon oxides (CO, CO2), nitrogen oxides (NO, NO2).			
Fire/Explosion Hazards:	Not applicable			
Advice for firefighters During all fire fighting ac apparatus.	tivities, wear appropriate protective equipment, including self-contained breathing			
SECTION 6: ACCIDENTAL RELEASE	MEASURES			
	Equipment and Emergency Procedures ean-up should wear appropriate personal protective equipment (See section 8). Minimize			
Environmental Precautions Place waste in an approp environmental release.	priately labeled, sealed container for disposal. Care should be taken to avoid			
Methods and Material for Contain Measures for Cleaning /				
Collecting:	Contain the source of spill if it is safe to do so. Collect spill with absorbent material. Clean spill area thoroughly.			
Additional Consideratio	n			
for Large Spills:				

### SECTION 7: HANDLING AND STORAGE

#### **Precautions for Safe Handling**

Avoid breathing vapor or mist. Avoid contact with eyes, skin and clothing. When handling, use appropriate personal protective equipment (see Section 8). Wash thoroughly after handling. Releases to the environment should be avoided. Review and implement appropriate technical and procedural waste water and waste disposal measures to prevent occupational exposure or environmental releases. Potential points of process emissions of this material to the atmosphere should be controlled with dust collectors, HEPA filtration systems or other equivalent controls.

### Conditions for Safe Storage, including any Incompatibilities

Storage Conditions: Store as directed by product packaging

### Specific end use(s)

No data available.

SECTION 8: EXPOSURE CONTROLS,	
Control Parameters	
Occupational	
Exposure Limits:	No data available.
Exposure controls	
Engineering Controls:	Engineering controls should be used as the primary means to control exposures. General room ventilation is adequate, unless the process generates dust, mist or fumes.
Personal Protective	
Equipment:	Refer to applicable national standards and regulations in the selection and use of personal protective equipment (PPE).
Hands:	None
Eyes:	None
Skin:	None

## SECTION 9: PHYSICAL AND CHEMICAL PROPERTIES

Physical State:	Round and convex flavoured tablet with a cross-shaped break line on one side.			
Odor:	Odourless			
Colour:	Light brown with brown s	pots		
Molecular formula:	Mixture			
Molecular weight:	Mixture			
Solvent Solubility:	Not applicable			
Water Solubility:	Not applicable			
pH:	Not applicable			
Melting/Freezing Point:	No data available			
Boiling Point (°C):	No data available			
Partition Coefficient:	No data available			
Evaporation Rate:	No data available			
Vapor Pressure:	No data available			
Vapor Density:	No data available			
Viscosity:	No data available			
Flammability:				
Autoignition Ten	nperature (Solid):	No data available		
Flammability (So	olids):	No data available		
Flash Point (Liquid):		No data available		
Upper Explosive Limits (Liquid):		No data available		
Lower Explosive Limits (Liquid):		No data available		

### SECTION 10: STABILITY AND REACTIVITY

#### Reactivity

No data available

### Chemical stability

Stable under normal conditions of use.

#### Possibility of hazardous reactions

No data available

### **Conditions to avoid**

Fine particles (such as dust and mists) may fuel fires/explosions.

#### Incompatible materials

As a precautionary measure, keep away from strong oxidizers.

### Hazardous decomposition products

No data available

### SECTION 11: TOXICOLOGICAL INFORMATION

### Information on Toxicological Effects

**General Information:** 

Toxicological properties of the formulation have not been investigated. The information in this section describes the potential hazards of the individual ingredients and the formulation. Routes of exposure: ingestion, eye contact, skin contact.

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

# Carprofen

Carprofen

Mouse	Oral	LD <sub>50</sub>	282 mg/kg
Rat	Oral	LD <sub>50</sub>	149 mg/kg
Rat (M/F)	SC	LD <sub>50</sub>	230/190 mg/kg
Rat (M/F)	IP	LD <sub>50</sub>	140/110 mg/kg

#### Irritation/sensitization (Study type, species, severity)

Carprof	en		
Eye	Irritation	Rabbit	Non-irritating
Skin	Irritation	Rabbit	Non-irritating
Skin	Sensitization – GPMT	Guinea Pig	Negative
	Antigenicity- Delayed skin reaction	Guinea Pig	No effect

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

13 Week(s)	Rat	Oral	5 mg/kg/day	NOAEL	Gastrointestinal System
13 Week(s)	Dog	Oral	5 mg/kg/day	NOAEL	None identified

### Reproduction & Developmental Toxicity: (Study Type, Species, Route, Dose, End Point, Effect(s))

#### Carprofen **Reproductive & Fertility** Rat 20 mg/kg/day NOAEL Fetotoxicity, Maternal toxicity Embryo / Fetal Development Rat 20 mg/kg/day NOAEL Not Teratogenic Prenatal & Postnatal Development Mouse 40 mg/kg/day NOAEL Not Teratogenic Prenatal & Postnatal Development Rabbit Oral 6 mg/kg/day NOAEL Embryotoxicity, Early embryonic development

### Genetic Toxicity: (Study Type, Cell Type/Organism, Result) Carprofen Bacterial Mutagenicity (Ames) Salmonella Negative

Bacterial Mutagenicity (Ames)	Sumonenu	negative
In Vivo Micronucleus	Mouse	Negative

### Carcinogenicity: (Duration, Species, Route, Dose, End Point, Effect(s))

Carprofen				
2 Year(s) Rat	Oral	10 mg/kg/day	NOAEL	Not carcinogenic, Gastrointestinal system
2 Year(s) Dog	Oral	25 mg/kg/day	NOAEL	Not carcinogenic, No effects at maximum dose

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

### SECTION 12: ECOLOGICAL INFORMATION

Environmental Overview:	Environmental properties have not been investigated. Releases to the environment should be avoided.
Toxicity:	No data available.
Persistence and Degradability:	No data available
Bio-accumulative Potential:	No data available
Mobility in Soil:	No data available

### SECTION 13: DISPOSAL CONSIDERATIONS

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

#### SECTION 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.

### SECTION 15: REGULATORY INFORMATION

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

Particular provisions:Veterinary medicinal product, under the scope of 2001/82/EC as amendedChemical safety assessment:No data available

# SECTION 16: OTHER INFORMATION

### Text of R phrases and GHS Classification abbreviations mentioned in Section 3

Acute toxicity, oral-Cat.3; H301 - Toxic if swallowed

Specific target organ toxicity, repeated exposure-Cat.2; H373 - May cause damage to organs through prolonged or repeated exposure Reproductive toxicity-Cat.2; H361d - Suspected of damaging the unborn child

T - Toxic Toxic to Reproduction: Category 3 Xn - Harmful

R25 - Toxic if swallowed.R63 - Possible risk of harm to the unborn child.R48/22 - Harmful: danger of serious damage to health by prolonged exposure if swallowed.

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