

Date of Issue: May 2012 Revision: Nov 2015

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

1.1 Product Identifier: 50-50 MiNT

1.2 Relevant uses of the substance or mixture and uses advised against:

Fertiliser

1.3 Manufacturer/Distributor: Vitax Limited

Owen Street Coalville LE67 3DE

Tel: 01530 510060 Fax: 01530 510299 Email: tech@vitax.co.uk

**1.4 Emergency Contact:** Tel: 01530 510060 (Office Hours)

### 2. HAZARDS IDENTIFICATION

**Primary Hazard** 

2.2 Label Elements

2.1 Classification Classification according to Regulation (EC) No 1272/2008 (EU-GHS/CLP)

Acute Tox. 4 H302 Harmful if swallowed.

Skin Irritation 2 H315 Causes skin irritation.

Eye Damage 1 H318 Causes serious eye damage

Aquatic Chronic 2 H411 Toxic to aquatic life with long lasting effects Causes serious eye damage. Harmful if swallowed. Causes skin irritation. **Vitax 50/50 MiNT** contains: Iron sulphate E.C. 231-753-5, Zinc suphate E.C.

231-793-3







Signal word: Danger

Hazard Statements: H302 Harmful if swallowed

H315 Causes skin irritation H318 Causes serious eye damage

H411 Toxic to aquatic life with long lasting effects

EUH208 Contains (reaction mass of: 5-chloro-2- methyl-4-isothiazolin-3-one [EC no. 247-500-7] and 2-methyl-2H -isothiazol-3- one [EC no. 220-239-6] (3:1).

May produce an allergic reaction.

**Precautionary Statements** P280 Wear protective gloves/protective clothing/eye protection/face protection.

P302 + P352 IF ON SKIN: Wash with plenty of soap and water.

P305 + P351 + P338 IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.

P310 Immediately call a POISON Center or doctor/physician. P362 Take off contaminated clothing and wash before reuse.

P391 Collect spillage

P501 Dispose of contents/container in accordance with local/national regulation

Mixture not classed as PBT or vPvB

### 3. COMPOSITION/INFORMATION ON INGREDIENTS

#### 3.2 Mixtures

2.3 Other Hazards

Chemical Name	CAS-No./ EINECS-No.	Annex Index or REACh number	Symbol(s)	R-phrase(s)	Concentration [%]
Iron sulphate	7720-78-7/ 231-753-5	Index number 026-003-00-7 REACh Number: 01 - 2119513203 – 57	According to 1272/2008: GHS07	According to 1272/2008:  Acute tox. 4, H302 Skin irrit. 2, H315 Eye Irrit. 2, H319  According to 67/548/EEC: R22, R36/38	10.0 – 19.9
Zinc sulphate	7733-02-0/ 231-793-3	Index No: 030-006-00-9 REACh No.: 01-2119474684-27	According to 1272/2008: GHS05 GHS07	According to 1272/2008: Acute Tox. 4 H302 Eye Damage 1 H318 Aquatic Acute 1 H400 Aquatic Chronic 1 H410	5.0 – 9.9



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			GHS09		
Manganese sulphate	10034-96-5/ 232-089-9	Index no.: 025-003-00-4 REACh registration no.: 01-2119456624-35	According to 1272/2008: GHS05 GHS08 GHS09	According to 1272/2008: Eye Damage 1, H318 STOT Rep. 2, H373 Aqu. Tox. chron. 2, H411	<2.0
Citric acid	77-92-9/ 201-069-1	REACh number: 01-2119457026-42	According to 1272/2008: GHS07	According to 1272/2008: Eye Irrit. 2, H319	<2.0
Boric acid	10043-35-3/ 233-139-2	Index No.: 005-007-00-2 REACh No: 01-2119486683-25	According to 1272/2008: GHS08	According to 1272/2008: Repr. 1B H360	<1.0
Reaction mass of: 5-chloro-2- methyl-4-isothiazolin-3-one [EC no. 247-500-7] and 2-methyl-2H - isothiazol-3- one [EC no. 220-239-6] (3:1)	55965-84-9/ 611-341-5	Index number: 613-167-00-5	According to 1272/2008: GHS05 GHS06 GHS07 GHS08	According to 1272/2008: Acute Tox. 3 * - H331 Acute Tox. 3 * - H311 Acute Tox. 3 * - H301 Skin Corr. 1B – H314 Skin Sens. 1 – H317 Aquatic Acute 1 – H400 Aquatic Chronic 1 – H410	<0.001

### 4. FIRST AID MEASURES

- 4.1 Description of first aid measures
- 4.1.1 Inhalation
- 4.1.2 Skin & Eye exposure

If symptoms arise remove from source of exposure to fresh air; seek medical attention if symptoms persist or develop

Skin: Drench immediately with water. Remove any contaminated clothing and launder before re-use. Obtain medical attention if symptoms develop or persist. Eyes: Immediately rinse with clean water for 15 minutes. Obtain medical attention IMMEDIATELY.



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4.1.3 Ingestion

Do not induce vomiting. Wash out mouth with water and give water to drink.

Obtain medical attention if symptoms develop or persist.

4.2 Most important symptoms and effects, both acute and delayed

Causes serious eye damage.

4.3 Indication of any immediate medical attention and special treatment needed.

Information not available.

5. FIRE-FIGHTING MEASURES

**5.1 Extinguishing media**Use foam, carbon dioxide, dry powder, sand. The mixture is not classified as

flammable as such extinguishing media should also be chosen as appropriate for

surrounding materials.

5.2 Special Hazards arising from the substance or mixture

Possible irritant fumes arising from combustion.

**5.3 Advice for fire-fighters** Cool down containers/equipment exposed to heat with a water spray. Contain

spread of extinguishing fluids (these fluids may be hazardous for the

environment). Wear complete protective clothing and self-contained breathing

apparatus.

#### 6. ACCIDENTAL RELEASE MEASURES

6.1 Personal precautions, protective equipment and emergency procedures

The following precautions are considered to be good practice when using any chemicals irrespective of their classification unless otherwise specified.

Ensure adequate ventilation. Use personal protective equipment, Gloves, Eye

protection, Suitable respirator if dust is generated during handling

**6.2 Environmental Precautions** Do not allow to enter storm drains or water courses. If this product enters a water

course or a sewer (including via contaminated soil & vegetation) in large quantities contact local water authority and inform the Environment Agency

6.3 Methods and material for containment and cleaning up

Use soil, sand or other absorbent material. Contact specialist waste disposal

contractor.

**6.4 Reference to other sections** See also section 8

7. HANDLING AND STORAGE

**7.1 Precaution for safe handling** Avoid contact with skin and eyes. Wash hands thoroughly after handling. Do not

eat, drink or smoke when using this product

7.2 Conditions for safe storage, including any incompatibilities

Store in a cool dry atmosphere, in original labelled containers. Refer to

manufacturer for maximum safe stacking height. Keep away from heat sources,

combustible materials and strong oxidising agents.

**7.3 Specific end use(s)** No Information available

#### 8. EXPOSURE CONTROLS/PERSONAL PROTECTION

### **8.1 Control Parameters**

Workplace exposure Limits as defined by UK HSE in document EH40/2005 where available:

Substance	CAS number	Workplace Exposure Limit				Comments
		Long-term exposure limit (8-hr TWA reference period)		Short-term exposure limit (15 minute reference period)		The Carc, Sen and Sk notations are not exhaustive. Notations have been applied
		ppm	mg.m <sup>-3</sup>	ppm	mg.m <sup>-3</sup>	to the substances identified in IOELV Directives*
Manganese and its inorganic compounds (as Mn)		-	0.5	-	-	-
Iron salts (as Fe)		-	1.0	-	2.0	-

<sup>\*</sup>IOELV - Indicative Occupational Exposure Limit Values (IOLEV).

**8.2 Exposure controls** 

The following precautions are considered to be good practice when using any chemicals irrespective of their classification unless otherwise specified.

Primary Hazard considered as handling of concentrate. Gloves: to BS EN374 of gauntlet type in Natural Rubber or PVC (not Nitrile) recommended for acid

resistance. Clothing: Coveralls/apron to BS EN465/466/467



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#### 9. PHYSICAL AND CHEMICAL PROPERTIES

9.1 Information on basic physical and chemical properties

Appearance; Brown liquid

Odour: No Information available Odour threshold; No Information available

pH;  $1.7 \pm 0.5$ 

Melting point/freezing; No Information available Initial boiling point and boiling range; No Information available Flash point; No Information available **Evaporation rate;** No Information available

Flammability (solid, gas); Mixture is not classed as flammable Upper /lower flammability or explosive limits; Mixture is not classed as explosive

Vapour Pressure: Negligible

Vapour density: No Information available

Specific Gravity;  $1.28 \pm 0.2$ 

Solubility (ies); No Information available Partition coefficient: n-octanol/water; No Information available **Auto ignition temperature:** No Information available **Decomposition temperature:** No Information available

9.2 Other Information No other relevant information available

#### 10. STABILITY AND REACTIVITY

10.1 Reactivity Unknown.

10.2 Chemical Stability Stable under normal conditions 10.3 Possibility of hazardous reactions No Information available. 10.4 Conditions to avoid Extremes of temperature

10.5 Incompatible materials None known

10.6 Hazardous decomposition products

Possible irritant fumes.

### 11. TOXICOLOGICAL INFORMATION

#### 11.1 Information on toxicological effects

The mixture has not been assessed for toxicological effects, the mixture classification is given in section 2 based on individual component contents.

Individual component hazards if any are given in section 3

Toxicological information on individual components where available:

Iron sulphate:

Dose descriptor:

Toxicological information

Acute toxicity The overall pattern of oral toxicity for iron salts is that they are harmful if

> swallowed. The human oral lethal dose is approximately 1000 mg/kg and 500-2000 mg/kg in rats. Toxic effects may, however, be produced by much lower doses especially when administered systemically. There is limited evidence that inhaled soluble iron salts are tolerated by rats plus limited evidence that inhaled soluble iron salts do not impair lung function and the dermal lethal dose would be greater than 2000 mg/kg. The dermal limit dose of Ferrous Chloride in rats is greater than 2,000 mg/kg (>881 mg Fe/kg) and thus should be used to compare against Ferrous Sulphate. This suggests little potential for systemic toxicity in

humans after dermal contact. Oral - LD50S 300-2000 mg/kg bw

Dermal - LD50S >2000 mg/kg bw

Inhalation-No data

Skin corrosion / irritation Ferrous Sulphate is skin irritant based on (2:1 animals majority) in rabbit test and

is an eye irritant. Read across from Ferrous Sulphate and Ferric Chloride,

indicates that solutions have the same or a lower classification than the solid and that classification based on pH would be overly cautious. On this basis an irritant classification, Skin Irritation Cat 2. H315: Causes skin irritation should be applied to solutions based on rules for mixtures. This classification therefore applies for solutions of concentration > 10%. Ferrous Sulphate should not be seen as

corrosive just as an irritant.

Results are available for a GLP-compliant guideline study (Johnson, 2003), which Eye damage / irritation

> showed that a 25% solution of Ferrous Sulphate Heptahydrate caused no more than mild redness and chemosis after instillation into the rabbit eye. The predicted



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Respiratory / Skin sensitisation

Germ cell mutagenicity

Reproductive toxicity

classification based on reading across of several iron salts would be a classification between no classification and causes serious eye damage however due to the lack of test data and low pH (<2) a precautionary approach should be taken with classification as Eye Damage 2.

Ferrous Sulphate Heptahydrate has been tested in a guideline, GLP, Local Lymph Node Assay (Stitzinger, 2010: reliability 1). In this test Ferrous Sulphate gave a clear negative result and is therefore not considered a skin sensitiser. Results of a reliable LLNA test were clearly negative for Ferrous Sulphate Heptahydrate. There are a few case studies in which human subjects showed signs of sensitisation to iron; however overall these data are poor and do not provide convincing evidence of a positive reaction in humans. There is also poor evidence in animal studies of sensitisation as a result of exposure to iron. The widespread exposure of iron and its role in biological processes, together with the extensive use of dietary supplements suggest that sensitisation is not a concern.

With regard to their mutagenic properties, iron salts have been extensively tested in microbial and mammalian systems in vitro, and in mammalian and insect tests in vivo. There are inconsistencies in the in vitro findings, with a small number of studies returning positive results. This has been attributed to DNA damage following reduction of Fe(II) to Fe(II) with free radical or superoxide formation and subsequent redox recycling. This contrasts with the consistently negative results obtained in vivo where, presumably, more efficient control mechanisms exist that protect the body from iron-induced oxidative damage. It is concluded that iron salts are not genotoxic.

Due to its potential pro-oxidant effects, there has been extensive research into possible links between iron and cancer development. These include many clinical investigations into the effects of oral (dietary) iron salts in humans and links to cancer. Although iron has been implicated in the development of cancers at various sites because of its role as a pro-oxidant, the UK Scientific Advisory Committee on Nutrition concluded that there is not enough evidence to reach conclusions for any specific links (EVM, 2003).

Results from recent guideline oral screening studies performed on Ferrous Chloride and Ferrous Sulphate gave NOAELs for reproductive and developmental effects of >500 mg/kg body weight/day or >1000 mg/kg body weight/day (no adverse effects were observed), respectively. These findings are considered to be relevant to Ferric as well as Ferrous salts, as oxidation of Ferrous to Ferric occurs in the low pH of stomach before ingested iron is absorbed into the body. In humans, iron supplementation of about 5.8 to 11.7 mg/kg bw/day (for a 60kg individual) is routinely prescribed throughout pregnancy with no adverse effects on pregnancy outcome. Evidence of adverse effects on male testes has only been observed at acutely toxic, overload doses, at which some of the experimental animals died.

Oral - LD50S >1000 mg/kg bw day

Dermal - No data Inhalation - No data

No human data is available for Ferrous Sulphate and repeated dose toxicity and even though effects are shown in some animal studies the overall conclusion is that no classification should be assigned for all endpoints oral, inhalation and dermal. NOAEL 49 days –100mg/kg Ferrous Sulphate Heptahydrate, result = no

effect.

**Aspiration hazard** No data, not an aspiration hazard.

Zinc sulphate:

Dose descriptor:

Repeated dose toxicity

Toxic Dose 1 - LD 50 862 - 4429 mg/kg (oral rat)

Acute Toxicity (Dermal LD50) > 2000 mg/kg Rat

Health Warnings INHALATION. Prolonged inhalation of high concentrations may damage respiratory system. SKIN CONTACT. Acts as a defatting agent on skin. May

cause cracking of skin, and eczema. Prolonged or repeated exposure may cause severe irritation. EYE CONTACT. May cause severe irritation to eyes.

INGESTION. The product causes irritation of mucous membranes and may cause

abdominal discomfort if swallowed. Skin Eyes Respiratory system, lungs

Target Organs



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Manganese sulphate:

HEALTH WARNINGS INHALATION. Prolonged inhalation of high concentrations may damage

respiratory system. SKIN CONTACT. Acts as a defatting agent on skin. May cause cracking of skin, and eczema. Prolonged or repeated exposure may cause

severe irritation. EYE CONTACT. May cause severe irritation to eyes.

INGESTION. The product causes irritation of mucous membranes and may cause

abdominal discomfort if swallowed.

TARGET ORGANS Skin Eyes Respiratory system, lungs

Citric acid

Acute toxicity

LD/LC50 Oral 3000mg/Kg (rat)

**Primary irritant effect:** 

On the skin: No irritant effect
On the eye: Irritating effect

Sensitisation: No sensitising effects known

Additional toxicological information: The product shows the following dangers according to the calculation method of

the General EU Classification Guidelines for Preparations as issued in the latest

version: Irritan

Routes of exposure: The substance can be absorbed into the body by inhalation (of solution mist and

dust) and by ingestion.

**Boric acid** 

Acute toxicity:

Acute Toxicity (Oral LD50) > 2000 mg/kg Rat

Test method(s): OECD 401.

Acute Toxicity (Dermal LD50) > 2000 mg/kg Rabbit

Test method(s): FIFRA (40 CFR 163)

Acute Toxicity (Inhalation LC50) > 2.03 mg/l (dust/mist) Rat 4 hours

Test method(s): OECD 403. Skin Corrosion/Irritation:

Dose 500 mg 24 hr Rabbit

Primary dermal irritation index (PDI) 0.1 (72hr)

Erythema\eschar score Very slight erythema -barely perceptible (1).

Oedema score No oedema (0).
Test method(s): FIFRA (40 CFR 163) Not classified.
Serious eye damage/irritation: Not classified.
Test method(s): equivalent or similar to OECD 405.

Respiratory or skin sensitisation:

Skin sensitisation Buehler test: Guinea Pig

Test method(s): OECD 406. Not Sensitising.

Germ cell mutagenicity:

Genotoxicity - In Vitro Negative.
Gene Mutation: Test method(s): OECD 471.

Genotoxicity - In Vivo Negative.

Chromosome aberration: Test method(s): equivalent or similar to OECD 474.

Carcinogenicity: Not classified.

Carcinogenicity NOEL>1150 mg/kg Oral Mouse

Test method(s): equivalent or similar to OECD 451.

Reproductive Toxicity:

Reproductive Toxicity – Fertility Known reproductive toxicant based on animal evidence.

Three-generation study: LOAEL 58.5 mg/kg Oral Rat P

The units are expressed in 'mg/µg' of: Boron. Test method(s): Toxicology and Applied Pharmacology 23: 351 - 364.

Reproductive Toxicity – Development May damage the unborn child. Developmental toxicity: NOAEC 21.8 mg/kg Oral Rabbit

The units are expressed in 'mg/µg' of: Boron. Test method(s): equivalent or similar to OECD 414.

Specific target organ toxicity - single exposure:

STOT - Single exposure No information available.

Specific target organ toxicity - repeated exposure: Not classified.

STOT - Repeated exposure LOAEL 58.5 mg/kg Oral Rat

Test method(s): Toxicology and Applied Pharmacology 23: 351 - 364.

Aspiration hazard: No data available.



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Inhalation Dust may irritate throat and respiratory system and cause coughing.

Ingestion Irritating. May cause nausea, stomach pain and vomiting.

Skin contact May cause skin irritation/eczema.

Eye contact Irritating and may cause redness and pain.

reaction mass of: 5-chloro-2- methyl-4-isothiazolin-3-one [EC no. 247-500-7] and 2-methyl-2H -isothiazol-3- one

[EC no. 220-239-6] (3:1)

Oral LD50: 67 mg/Kg (rat)
Dermal LD50: >140 mg/Kg (rat)

Inhalation LC50/4hr 0.17 mg/l (rat) Aerosol THR 48/971458

Primary irritant effect:

On the skin: Caustic effect on skin and mucous membranes

On the eye: Strong caustic effect

Sensitization: Sensitization possible by skin contact

12. ECOLOGICAL INFORMATION

**12.1 Toxicity** Mixture classified as toxic to aquatic life with long lasting effects.

**12.2 Persistence and degradability 12.3 Bioaccumulative potential 12.4 Mobility in soil**Information not available.
Information not available.

12.5 Results of PBT and vPvB Not classified

**12.6 Other adverse effects** Information not available

13. DISPOSAL CONSIDERATIONS

**13.1 Waste Treatment Methods**Use only licensed waste disposal companies for unwanted chemical. Do not re-use

empty containers for any purpose.

14. TRANSPORT INFORMATION

**14.1 UN number:** UN3082

**14.2 UN proper shipping name:** Environmentally hazardous preparation, liquid N.O.S. (contains: Iron sulphate

E.C. 231-753-5, Zinc suphate E.C. 231-793-3)

**14.3 Transport hazard** 9 **14.4 Packing group:** III

**14.5 Environmental hazards:** Product is classified as toxic to aquatic life with long lasting effects.

**14.6 Special precautions for user:** No information available

14.7 Transport in bulk according to Annex II of MARPOL 73/78 and the IBC code

Applicable for Maritime bulk transport only. Check with carrier.

## 15. REGULATORY INFORMATION

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

This substance is classified and labelled in accordance with Directive 1999/45/EC regulation (EC) 1272/2008 and the EC Fertiliser Regulations 2003, Regulation (EC) No 1907/2006 of the European Parliament and of the Council of 18 December 2006 concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH), establishing a European Chemicals Agency, amending Directive 1999/45/EC and repealing Council Regulation (EC) No 793/93 and Commission Regulation (EC) No 1488/94 as well as Council Directive 76/769/EEC and Commission Directives 91/155/EEC, 93/67/EEC, 93/105/EC and 2000/21/EC, including amendments. Regulation (EC)

15.2 Chemical Safety Assessment CSA not undertaken for this mixture

## 16. OTHER INFORMATION

### Hazard Information not otherwise listed in full elsewhere:

H301: Toxic if swallowed

H311: Toxic in contact with skin.

H314: Causes severe skin burns and eye damage

H317: May cause an allergic skin reaction.

H319: Causes serious eye irritation.

H331: Toxic if swallowed or if inhaled

H360FD: May damage fertility. May damage the unborn child.

H373: May cause damage to organs through prolonged or repeated exposure

H400: Toxic to aquatic life.



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**SDS** information:

H410: Very toxic to aquatic life with long lasting effects.

This Safety data sheet is compiled using data submitted for raw materials and practical experience. This product is intended for professional users only. This Safety Data Sheet is prepared in compliance with Directive 1999/45/EC, regulation 1272/2008 and Annex I of the REACH regulation 453/2010. The information given herein is, to the best of our knowledge, correct and is presented in good faith but no warranty, expressed or implied is given.