MSDS
Material Safety Data Sheet

Product Name: Revamil Balm (15 g, 50 g)

Section I Manufacturer's Information

Manufacturer's Name and Address	
Bfactory Health Products B.V.	Emergency Telephone Number: +31(0)317 769 005
Remmerden 58	Number for Information: +31(0)317 769 005
3911 TZ Rhenen, The Netherlands	Date prepared: June 30, 2021

Section II Hazardous Identity Information

INCI code (common name)	**CAS number**	**Potential Health Effects**
Arachis hypogaea (purified peanut oil – nonallergenic –)	8002-03-7	Non-hazardous
Beeswax (cera alba)	8012-89-3	Non-hazardous
glyceryl oleate	67701-32-0	Not known
Mel (honey)	8028-66-8	Non-hazardous
Aqua (Purified water)	7732-18-5	Non-hazardous

Section III Physical/ Chemical Characteristics

Boiling point	Not known
Vapor pressure	Not known
Melting point	Not known
Evaporation rate	Not known
Solubility in water	Not soluble
Physical appearance	White cream

Section IV Health Hazard Data

Entry routes	**Eyes**: Flush eyes with copious amounts of water **Ingestion**: If person is conscious, dilute by drinking water. Do not induce vomiting. Contact a physician or poison control centre for instructions **Skin contact**: no measures needed **Special precautions**: no measures needed
Target Organs	Not known
Carcinogenity	This material is not known to have carcinogenic properties
Medical conditions aggravated by long-term exposure	Not known
Chronic effects	Not known
Sterility	Gamma-irradiated at 25 kGy

Section V Fire and Explosion Hazard data

Material is not inflammable

Section VI Reactivity data

Material is not reactive

Section VII Precautions for safe handling and use

Steps to be taken in case material is spilled or released	Not applicable
Waste disposal method	Not applicable
Precautions to be taken in Handling and Storing	Not considered necessary
Other precautions	Not considered necessary

Section VIII Control methods

Respiratory methods	None needed
Ventilation	Normal room ventilation
Protective equipment	Eyes: none needed Hands: none needed Other: none needed
Work/ hygienic practices	No unusual practices needed

Section IX Registration information

Registration	Medical device, class IIa
Notified Body	DEKRA Certification B.V., Notified body nr 0344 Arnhem, the Netherlands