ZEEL - silicon dioxide and comfrey root and arnica montana root and toxicodendron pubescens leaf and sulfur and sanguinaria canadensis root and solanum dulcamara stem and coenzyme a and nadide and sodium diethyl oxalacetate and .alpha.-lipoic acid and sus scrofa cartilage and sus scrofa embryo and sus scrofa umbilical cord and sus scrofa placenta and ointment
Biologische Heilmittel Heel

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

ACTIVE INGREDIENTS

Active ingredients: Each 50g of ointment contains: Silicea 6X 0.5g; Symphytum officinale 8X 0.375g; Arnica montana, radix 2X 0.15g; Rhus toxicodendron 2X 0.135g; Sulphur 6X 0.135g; Sanguinaria canadensis 2X 0.113g; Dulcamara 2X 0.038g; Coenzyme A 6X 0.005g; Nadidum 6X 0.005g; Natrum oxalaceticum 6X 0.005g; α- Lipoicum acidum 6X 0.005g; Cartilago suis 2X 0.0005g; Embryo suis 2X 0.0005g; Funiculus umbilicalis suis 2X 0.0005g; Placenta suis 2X 0.0005g.

PURPOSE

Minor joint pain and stiffness

KEEP OUT OF REACH OF CHILDREN

Keep this and all medicine out of the reach of children.

INDICATIONS AND USAGE

For the temporary relief of:

- Minor joint pain and stiffness

WARNINGS

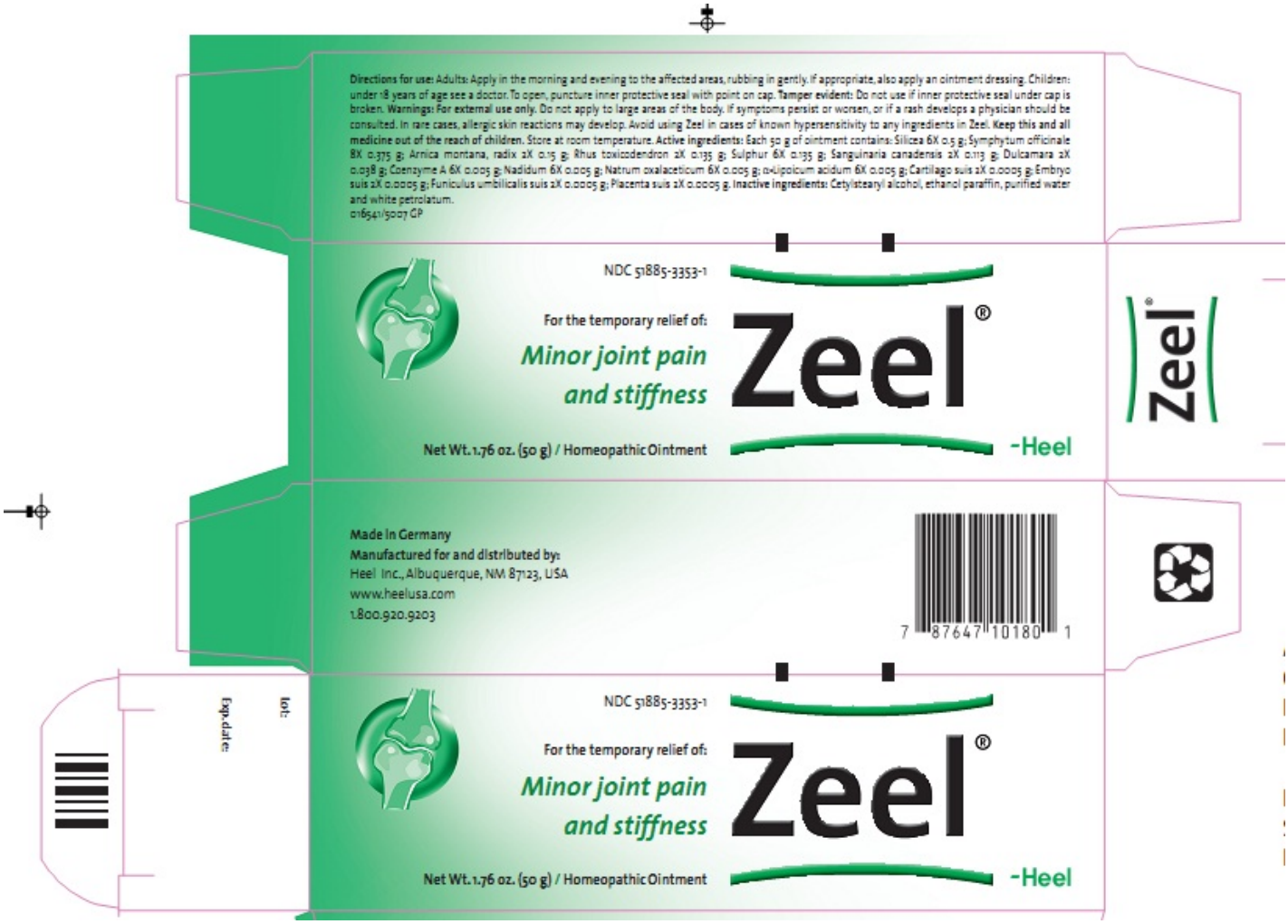
Warnings: For external use only. Do not apply to large areas of the body. If symptoms persist or worsen, or if a rash develops a physician should be consulted. In rare cases, allergic skin reactions may develop. Avoid using Zeel in cases of known hypersensitivity to any ingredients in Zeel.

DOSAGE AND ADMINISTRATION

Direction for use: Adults: Apply in the morning and evening to the effected areas, rubbing in gently. If appropriate, also apply an ointment dressing. Children under 18 years of age see a doctor.

INACTIVE INGREDIENTS

Inactive ingredients: Cetylstearyl alcohol, ethanol, paraffin, purified water and white petrolatum.



Zeel 50g Carton.jpg

ZEEL

silicon dioxide and comfrey root and arnica montana root and toxicodendron pubescens leaf and sulfur and sanguinaria canadensis root and solanum dulcamara stem and coenzyme a and nadide and sodium diethyl oxalacetate and .alpha.-lipoic acid and sus scrofa cartilage and sus scrofa embryo and sus scrofa umbilical cord and sus scrofa placenta and ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51885-3353
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	6 [hp_X] in 50 g
COMFREY ROOT (UNII: M9VVZ08EKQ) (COMFREY ROOT - UNII:M9VVZ08EKQ)	COMFREY ROOT	8 [hp_X] in 50 g
ARNICA MONTANA ROOT (UNII: MUE8Y11327) (ARNICA MONTANA ROOT - UNII:MUE8Y11327)	ARNICA MONTANA ROOT	2 [hp_X] in 50 g

TOXICODENDRON PUBESCENS LEAF (UNII: 6IO182RP7A) (TOXICODENDRON PUBESCENS LEAF - UNII:6IO182RP7A)	TOXICODENDRON PUBESCENS LEAF	2 [hp_X] in 50 g
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	6 [hp_X] in 50 g
SANGUINARIA CANADENSIS ROOT (UNII: N9288CD508) (SANGUINARIA CANADENSIS ROOT - UNII:N9288CD508)	SANGUINARIA CANADENSIS ROOT	2 [hp_X] in 50 g
SOLANUM DULCAMARA STEM (UNII: IR986LE7DF) (SOLANUM DULCAMARA STEM - UNII:IR986LE7DF)	SOLANUM DULCAMARA STEM	2 [hp_X] in 50 g
COENZYME A (UNII: SAA04E81UX) (COENZYME A - UNII:SAA04E81UX)	COENZYME A	6 [hp_X] in 50 g
NADIDE (UNII: 0U46U6E8UK) (NADIDE - UNII:0U46U6E8UK)	NADIDE	6 [hp_X] in 50 g
SODIUM DIETHYL OXALACETATE (UNII: 6CA025Y4FG) (SODIUM - UNII:9NEZ333N27)	SODIUM DIETHYL OXALACETATE	6 [hp_X] in 50 g
.ALPHA.-LIPOIC ACID (UNII: 73Y7P0K73Y) (.ALPHA.-LIPOIC ACID - UNII:73Y7P0K73Y)	.ALPHA.-LIPOIC ACID	6 [hp_X] in 50 g
SUS SCROFA CARTILAGE (UNII: 73ECW5WG2F) (SUS SCROFA CARTILAGE - UNII:73ECW5WG2F)	SUS SCROFA CARTILAGE	2 [hp_X] in 50 g
SUS SCROFA EMBRYO (UNII: 9928MC12VO) (SUS SCROFA EMBRYO - UNII:9928MC12VO)	SUS SCROFA EMBRYO	2 [hp_X] in 50 g
SUS SCROFA UMBILICAL CORD (UNII: 118OYG6W3H) (SUS SCROFA UMBILICAL CORD - UNII:118OYG6W3H)	SUS SCROFA UMBILICAL CORD	2 [hp_X] in 50 g
SUS SCROFA PLACENTA (UNII: C8CV8867O8) (SUS SCROFA PLACENTA - UNII:C8CV8867O8)	SUS SCROFA PLACENTA	2 [hp_X] in 50 g

Inactive Ingredients

Ingredient Name	Strength
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	
PETROLATUM (UNII: 4T6H12BN9U)	
PARAFFIN (UNII: I9O0E3H2ZE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51885-3353-1	1 in 1 CARTON		
1		50 g in 1 TUBE		
2	NDC:51885-3353-3	1 in 1 CARTON		
2		50 g in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/20/2011	

Labeler - Biologische Heilmittel Heel (315635359)

Revised: 2/2012 Biologische Heilmittel Heel