SUMMARY OF PRODUCT CHARACTERISTICS

1. NAME OF THE MEDICINAL PRODUCT
VERRUPAN, lotion for local application

2. QUALITATIVE AND QUANTITATIVE COMPOSITION
Cedar tincture ..... 15 g
Salicylic acid ..... 3 g
Lactic acid ..... 3 g
For 100 g of lotion.
For the complete list of excipients, see section 6.1.

3. PHARMACEUTICAL FORM
Solution for local application.

4. CLINICAL PARTICULARS

4.1 Therapeutic indications
Treatment of plantar warts, vulgar hand warts and mosaic warts.

4.2 Posology and method of administration
Two to three applications per day.
In the evening before the last application, gently rub the surface of the wart with a cardboard file or pumice stone, avoiding bleeding, so as to remove the rough surface of the wart.
If bleeding occurs, stop treatment for three days.
Treatment may continue for several weeks or even months.

4.3 Contraindications
Allergy to salicylic acid.
Do not apply to mucous membranes.

4.4 Special warnings and precautions for use
Warnings
Do not apply to mucous membranes.
Precautions for use
When applying, do not overflow onto healthy skin.

4.5 Interactions with other medicinal products and other forms of interactions
Not applicable.

4.6 Pregnancy and breast-feeding
Not applicable.

4.7 Effects on ability to drive and use machines
Not applicable.

4.8 Undesirable effects
Possible skin irritation if applied to healthy skin or mucous membranes.

4.9 Overdose
Not applicable.

5. PHARMACOLOGICAL PROPERTIES

5.1 Pharmacodynamic properties
Pharmacotherapeutic class : KERATOLYTIC, WART REMOVER, CORN REMOVER. (D. DERMATOLOGY)

5.2 Pharmacokinetic properties
Not documented.

5.3 Preclinical safety data
Not documented.

6. PHARMACEUTICAL PARTICULARS

6.1 List of excipients
Elastic collodion.

6.2 Incompatibilities
Not applicable.

6.3 Shelf life
3 years.

6.4 Special precautions for storage
This product is very volatile and flammable, it should be kept away from all sources of heat and flame.
Close the bottle tightly after use.

6.5 Nature and contents of container
Vial (glass) of 3 g. Box of 2 vials.

6.6 Special precautions for disposal and handling
No particular requirements.

7. MARKETING AUTHORISATION HOLDER
PHARMA DEVELOPPEMENT SAS
CHEMIN DE MARCY
58800 CORBIGNY
FRANCE

8 . MARKETING AUTHORISATION NUMBER (S)
- 3400932950072 : 2 glass vials of 3 g.

9. DATE OF FIRST AUTHORISATION / RENEWAL OF THE AUTHORISATION
23/05/1979

10. DATE OF REVISION OF THE TEXT
12/09/2013

11. DOSIMETRY
Not relevant.

12. INSTRUCTIONS FOR PREPARATION OF RADIOPHARMACEUTICALS
Not relevant.

PRESCRIPTION AND SUPPLY CONDITIONS
This medicinal product is available without a prescription.