

SUMMARY OF PRODUCT CHARACTERISTICS

1. NAME OF THE MEDICINAL PRODUCT

Immunokind

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

1 tablet (100 mg) contains:

Active substances:

Calcium carbonicum Hahnemanni trit. D6	20.00 mg
Calcium fluoratum trit. D6	20.00 mg
Calcium phosphoricum trit. D6	20.00 mg
Sulfur jodatum trit. D12	20.00 mg

Excipient with known effect: lactose monohydrate (used as carrier for the homeopathic triturations)
For the full list of excipients, see section 6.1.

3. PHARMACEUTICAL FORM

Tablet

4. CLINICAL PARTICULARS

4.1 Therapeutic indications

To increase resistance to recurring upper respiratory tract infections in infants and children of early age.

4.2 Posology and method of administration

Posology

Children up to 6 years of age: 1 tablet is administered 3 times per day.

Method of administration

Tablets should be allowed to dissolve slowly in the mouth. If necessary, tablets can be dissolved in a little bit of water.

An interval of at least half an hour to meals should be kept.

4.3 Contraindications

Hypersensitivity to one of the active substances or to any of the excipients.

4.4 Special warnings and precautions for use

This medicinal product contains lactose. Patients with rare hereditary problems of galactose intolerance, the Lapp lactase deficiency or glucose-galactose malabsorption should not take this medicinal product.

A reassessment of the therapy is necessary

- if the child is poorly responding to Immunokind,
- if additional complaints arise.

Parents are advised to consult a doctor if their child is suffering from a thyroid disease before starting the treatment with Immunokind.

Parents are advised not to use Immunokind for long-term treatment without consultation with a doctor.

4.5 Interaction with other medicinal products and other forms of interaction

Interactions with other medicinal products have not been observed. The use of this homeopathic medicinal product does not exclude the use of other medicinal products.

Parents are advised to consult a doctor if their child is taking, has recently taken or might take any other medicinal products.

4.6 Pregnancy and lactation

This medicinal product is meant for children.

4.7 Effects on ability to drive and use machines

This medicinal product is meant for children.

4.8 Undesirable effects

No side effects are known up to present.

If any side effects are noticed, appropriate measures have to be taken.

Note: Existing complaints may temporarily become worse when homeopathic medicines are taken (initial homeopathic aggravation). In such cases, patients are advised to discontinue the medication and consult a doctor.

4.9 Overdose

No case of overdose has been reported.

5. PHARMACOLOGICAL PROPERTIES

Pharmacotherapeutic group: Homeopathic medicinal product

Homeopathic therapy differs markedly from conventional medicine with regard to the understanding of disease and the use of medicinal substances. Homeopathic remedies stimulate the organism's own defence mechanisms and thereby activate its self-healing power. The response of the organism to a homeopathic remedy is highly individual. For these reasons dose-specific pharmacological effects, pharmacodynamic or pharmacokinetic data can not be determined.

5.1 Pharmacodynamic properties

Not applicable.

5.2 Pharmacokinetic properties

Not applicable.

5.3 Preclinical safety data

Not applicable. Immunokind as a homeopathic pharmaceutical have not gone through preclinical testing.

6. PHARMACEUTICAL PROPERTIES

6.1 List of excipients

Cellulose microcrystalline, magnesium stearate, talc.

6.2 Incompatibilities

Not applicable.

6.3 Shelf life

5 years.

Do not use after the expiration date indicated on the package

6.4 Special precautions for storage

Store at a temperature not exceeding 25 ° C. Store out of reach and sight of children.

6.5 Nature and contents of container

150 tablets in bottle; 1 bottle and PIL in carton box.

6.6 Special precautions for disposal

No special requirements.

7. MANUFACTURER

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8. MARKETING AUTHORISATION HOLDER

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9. MARKETING AUTHORISATION NUMBER(S)

[To be completed]

10. DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

19/05/2015

11. DATE OF REVISION OF THE TEXT

October 2023