



London, 21 February 2008
EMA/CHMP/86517/2008

**COMMITTEE FOR MEDICINAL PRODUCTS FOR HUMAN USE
POST-AUTHORISATION SUMMARY OF POSITIVE OPINION***
for
NEOCLARITYN

International Nonproprietary Name (INN): *desloratadine*

On 21 February 2008 the Committee for Medicinal Products for Human Use (CHMP) adopted a positive opinion** to recommend the variation to the terms of the marketing authorisation for the medicinal product Neoclarityn. The Marketing Authorisation Holder for this medicinal product is SP Europe.

The CHMP adopted a change to an indication as follows:

Neoclarityn is indicated for the relief of symptoms associated with:

- ~~chronic idiopathic~~ **urticaria** (see section 5.1)

Detailed conditions for the use of this product will be described in the updated Summary of Product Characteristics (SPC) which will be published in the revised European Public Assessment Report (EPAR) and will be available in all official European Union languages after the variation to the marketing authorisation has been granted by the European Commission.

For information, the full indications for Neoclarityn will be as follows***:

Neoclarityn is indicated for the relief of symptoms associated with:

- allergic rhinitis (see section 5.1)
- **urticaria (see section 5.1)**

* Summaries of positive opinion are published without prejudice to the Commission Decision, which will normally be issued within 44 days (Type II variations) and 67 days (Annex II applications) from adoption of the Opinion.

** Marketing Authorisation Holders may request a re-examination of any CHMP opinion, provided they notify the EMA in writing of their intention to request a re-examination within 15 days of receipt of the opinion.

*** The text in bold represents the new or the amended indication.