



**STATEMENT ON GOOD MANUFACTURING PRACTICES FOR
PHARMACEUTICAL STARTING MATERIALS (API)**

This one-page certificate conforms to the format recommended
by the World Health Organization

*On the basis of a voluntary inspection carried out by the Danish Medicines Agency on:
September 30th 2010*

we certify that the site indicated on this certificate complies with Good Manufacturing Practices for active pharmaceutical ingredients according to The Rules governing Medicinal Products in the European Union, Volume 4, Part II (ICH Q7) for the dosage forms, categories and activities listed in Table 1.

Name and address of site:

**Akzo Nobel Salt A/S
Hadsundvej 17
DK-9550 Mariager**

Manufacturer's authorisation number (DKMA company no.):

257120

Table 1

Product / Product category:

**Sanal P
Sanal P+
Sanal SQ**

Dosage form(s):

N.a.

Activity(ies):

**Manufacture
Quality control**

The responsibility for the purity and quality of the individual batches of the active pharmaceutical ingredients manufactured through the process lies with the manufacturer.

This certificate remains valid for three years from the date of last inspection.

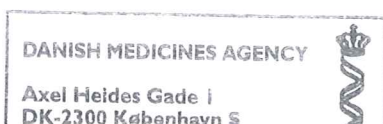
This certificate becomes invalid if the activities and/or categories certified are changed or if the site is no longer considered to be in compliance with GMP.

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H5-205
CGMP-API
Chem/Bio

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