LÄKEMEDELSVERKET
MEDICAL PRODUCTS AGENCY

Tillstånd

Datum: 2009-12-09 Dnr: 24:2009/515006

Tillstånd att tillverka naturläkemedel

Tillståndet är utformat enligt principerna i Community Basic Format for Manufacturers Authorisation.

Tillståndets nummer	24:2009/515006. Ersätter 24:2009/512856
Tillståndsinnehavarens namn	**Vitamex Production AB**
Organisationsnummer	556582-2953
Tillverkningsställets adress	Bergslagsgatan 9 Box 715
Tillståndsinnehavarens postadress	610 16 Norrköping
Tillståndets omfattning	Produktion Icke sterila produkter Fasta beredningsformer - Tabletter - Brustabletter Endast förpackning: - Mixturer - Kapslar - Import och frisläppning av läkemedel från tredje land
Begränsningar	-
Författningsrum	LVFS 2004:7
Sakkunnig person, titel och namn	**Fil.mag. Charlotta Gromulski**
Tillståndets giltighetstid	**2009-12-09 – 2014-10-22.**

På Läkemedelsverkets vägnar

Tor Gråberg
Chefsinspektör

Erik Larsson
Läkemedelsinspektör

Postadress/Postal address: P.O. Box 26, SE-751 03 Uppsala, SWEDEN
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Date: December 9, 2009 Dnr: 24:2009/522235

Certificate[1)]

Manufacturer: Vitamex Production AB
Address: Bergslagsgatan 9
SE-610 16 Norrköping
Sweden

It is certified that:
The manufacturer is registered as a producer of natural remedies in Sweden, and that the production has been and is being carried out under competent management, all in accordance with the Medicinal Products Act (1992:859) governing the importation, manufacture and distribution of such products in Sweden.
Manufacturing licence: 24:2009/515006 valid until 2014-10-22.

The manufacturing plant is subject to inspections at suitable intervals.
The last inspection was performed 2009-10-20-22.

The manufacturer conforms to requirements for good practices in the manufacture and quality control, as recommended by the World Health Organization and European Directive 2003/94/EC in respect of products to be sold or distributed within the country of origin or to be exported[2)].

This certificate reflects the status of the manufacturing site at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more then three years have elapsed since the date of the inspection, after which time the issuing authority should be consulted. The authenticity of this certificate may be verified with the issuing authority.

On duty

Tor Gråberg
Chief Pharmaceutical Inspector

Certification fee 475 SEK

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1. WHA 28.65 (see official Records No. 226, Annex 12, Part 1).
2. The requirements for good practices in the manufacture and quality control of drugs mentioned in the certificate refer to the text adopted by the Twenty-eighth World Health Assembly in its resolution WHA 28.65

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