Medicines and Healthcare products Regulatory Agency

CERTIFICATE NUMBER: UK API 44519 Insp GMP 44519/12076689-0001 [V]

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER (1),(2)

Part 1

Issued following an inspection in accordance with:

Regulation 5 of the current Veterinary Medicines Regulations

The competent authority of United Kingdom confirms the following:

The Manufacturer: NANJING KING FRIEND BIOCHEMICAL PHARMACEUTICAL COMPANY LIMITED

Site address: NANJING KING FRIEND BIOCHEMICAL PHARMACEUTICAL COMPANY LIMITED, MA-010-1, NANJING HIGH & amp; NEW TECHNOLOGY DEVELOPMENT ZONE, NANJING, CN-210 061, CHINA

Is an active substance manufacturer that has been inspected in accordance with Regulation 327 of The Human Medicines Regulations 2012 (SI 2012/1916).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 24/09/2018, it is considered that it complies with

• The principles of GMP for active substances referred to in Regulation B17 and C17 of the Human Medicines Regulations 2012 (SI 2012/1916)

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 than three years have elapsed since the date of that inspection. However, this period of validity may be reduced or extended using regulatory risk management principles by an entry in the Restrictions or Clarifying remarks field. This certificate is valid only when presented with all pages and both Parts 1 and 2. The authenticity of this certificate may be verified in MHRA-GMDP. If it does not appear, please contact the issuing authority.

- (1) Guidance on the interpretation of this template can be found in the Help menu of MHRA-GMDP database.
- (2) These requirements fulfil the GMP recommendations of WHO.

Part 2

Veterinary Medicinal Products

Manufacture of active substance. Names of substances subject to inspection :

- [2000007085] ENOXAPARIN SODIUM
- 3. MANUFACTURING OPERATIONS ACTIVE SUBSTANCES

ENOXAPARIN SODIUM

3.1 Manufacture of Active Substance by Chemical Synthesis

3 1 1	Manufacture	Of Active	Substance	Intermediates

3.1.2 Manufacture Of Crude Active Substance

3.1.3 Salt Formation/Purification steps (eg. Crystallisation)

Oxidation, filtration, precipitation, dehydration

3.5 General Finishing Steps

3.5.1 Physical Processing Steps

Drying, milling

3.5.2 Primary Packaging

3.5.3 Secondary Packaging

3.6 Quality Control Testing

3.6.1 Physical / Chemical testing

Any restrictions related to the scope of this certificate:

Building	Room Line/equipment QC Testing	Products
LMWH Workshop 3 and associated storage	IPC testing and testing of intermediates	Enoxaparin Sodium
areas and utilities	only at this site.	only

06/11/2018	Name and signature of the authorised person of the Competent Authority of United Kingdom		
	Confidential		
	Medicines and Healthcare products Regulatory Agency		
	Tel : Confidential		