

# Medicines and Healthcare products Regulatory Agency

CERTIFICATE NUMBER : UK GMP 33188 Insp GMP 33188/860751-0002[H]

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

### Part 1

Issued following an inspection in accordance with :  
Regulation 331A of The Human Medicines Regulations 2012 (SI 2012/1916)

The competent authority of United Kingdom confirms the following :

The Manufacturer : HYALURON INCORPORATED

Site address : HYALURON INCORPORATED, 99 SOUTH BEDFORD STREET, BURLINGTON, 01803, UNITED STATES

Other :

Voluntary inspection for investigational medicinal products

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 08/02/2010, it is considered that it complies with

- The principles and guidelines of Good Manufacturing Practice laid down in Regulation B17 of the Human Medicines Regulations 2012 (as amended)

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

#### Human Medicinal Products

#### 1. MANUFACTURING OPERATIONS

##### [ 1.1 ] Sterile Products

[ 1.1.1 ] Aseptically prepared (processing operations for the following dosage forms)

[ 1.1.1.2 ] Lyophilisates

[ 1.1.1.3 ] Semi-solids

[ 1.1.1.4 ] Small volume liquids

[ 1.1.2 ] Terminally Sterilised (processing operations for the following dosage forms)

[ 1.1.2.2 ] Semi-solids

[ 1.1.2.3 ] Small volume liquids

**[ 1.2 ] Non-sterile products**

[ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms)

[ 1.2.1.5 ] Liquids for external use

[ 1.2.1.6 ] Liquids for internal use

**[ 1.3 ] Biological medicinal products**

[ 1.3.1 ] Biological medicinal products

[ 1.3.1.2 ] Immunological products

[ 1.3.1.8 ] Other biological medicinal products

Monoclonal antibodies, Protein, RNA

**[ 1.4 ] Other products or manufacturing activity**

[ 1.4.2 ] Sterilisation of active substances/excipients/finished products:

[ 1.4.2.1 ] Filtration

[ 1.4.2.2 ] Dry heat

[ 1.4.2.3 ] Moist heat

**[ 1.5 ] Packaging**

[ 1.5.2 ] Secondary packaging

**[ 1.6 ] Quality control testing**

[ 1.6.2 ] Microbiological: non-sterility

[ 1.6.3 ] Chemical/Physical

[ 1.6.4 ] Biological

Any restrictions related to the scope of this certificate:

Building	Room	Line/equipment	QC Testing	Products
Suites in 99 South Bedford Street and 20 Blanchard Road				

05/07/2010	Name and signature of the authorised person of the Competent Authority of United Kingdom			
	Confidential			
	Medicines and Healthcare products Regulatory Agency			
	Tel : Confidential			