Medicines and Healthcare products Regulatory Agency

CERTIFICATE NUMBER : UK API 10082 Insp GMP 10082/20306-0009

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 : FUERST DAY LAWSON LIMITED

Site address : FUERST DAY LAWSON LIMITED, GOOSES FOOT INDUSTRIAL ESTATE, KINGSTONE, HEREFORD, HR2 9HY, UNITED KINGDOM

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 12/08/2025 , 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

Human Medicinal Products

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

- [2000008199] CAPSICUM TINCTURE
- [4000010248] CASCARA EXTRACT
- [2000013194] SQUILL OXYMEL
- [2000011380] COAL TAR SOLUTION
- [2000007466] SQUILL TINCTURE

3. MANUFACTURING OPERATIONS - ACTIVE SUBSTANCES

CAPSICUM TINCTURE

3.2	Processing Activities of Active Substance from Natural Sources
	3.2.1 Plant Source Extraction
3.5	General Finishing Steps
	3.5.1 Physical Processing Steps
	Mixing, blending and filtration
	3.5.2 Primary Packaging
3.6	Quality Control Testing
	3.6.1 Physical / Chemical testing
	3.6.2 Microbiological testing (excluding sterility testing)

CASCARA EXTRACT

3.2	Processing Activities of Active Substance from Natural Sources
	3.2.1 Plant Source Extraction
3.5	General Finishing Steps
	3.5.1 Physical Processing Steps
	Mixing, blending and filtration
	3.5.2 Primary Packaging
3.6	Quality Control Testing
	3.6.1 Physical / Chemical testing
	3.6.2 Microbiological testing (excluding sterility testing)

SQUILL OXYMEL

3.2	Processing Activities of Active Substance from Natural Sources
	3.2.1 Plant Source Extraction
3.5	General Finishing Steps
	3.5.1 Physical Processing Steps
	Mixing, blending and filtration
	3.5.2 Primary Packaging
3.6	Quality Control Testing
	3.6.1 Physical / Chemical testing
	3.6.2 Microbiological testing (excluding sterility testing)

COAL TAR SOLUTION

3.2	Processing Activities of Active Substance from Natural Sources

	3.2.4 Mineral Source Extraction
3.5	General Finishing Steps
	3.5.1 Physical Processing Steps Mixing, blending and filtration
	3.5.2 Primary Packaging
3.6	Quality Control Testing
	3.6.1 Physical / Chemical testing
	3.6.2 Microbiological testing (excluding sterility testing)

SQUILL TINCTURE

3.2	Processing Activities of Active Substance from Natural Sources
	3.2.1 Plant Source Extraction
3.5	General Finishing Steps
	3.5.1 Physical Processing Steps Mixing, blending and filtration
	3.5.2 Primary Packaging
3.6	Quality Control Testing
	3.6.1 Physical / Chemical testing
	3.6.2 Microbiological testing (excluding sterility testing)

Restrictions or Remarks

This certificate is issued based on a desk-based assessment of GMP compliance information provided by the manufacturer. This certificate should be used in combination with the relevant authorisation/registration. A risk-based site inspection programme remains in force.

12/08/2025	Name and signature of the authorised person of the Competent Authority of United Kingdom Confidential Medicines and Healthcare products Regulatory Agency Tel : Confidential