Safeguarding public health

MHRA

Mr R Holiday
CLF DISTRIBUTION LIMITED
UNIT 16
PARKERS CLOSE
DOWNTON BUSINESS CENTRE
SALISBURY
WILTSHIRE
SP5 3RB
UNITED KINGDOM

Medicines and Healthcare products Regulatory Agency
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An executive agency of the Department of Health

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MEDICINES AND HEALTHCARE PRODUCTS REGULATORY AGENCY
On behalf of the Licensing Authority under The Medicines Act 1968 as amended

Wholesale Dealer's (General Sales List Products) Licence

SECTION 1

1. This licence is granted in accordance with section 8 (3) and (3A) of The Medicines Act 1968 as amended and is subject to the provisions of The Medicines Act 1968 as amended and 1971 and The Medicines for Human Use (Manufacturing, Wholesale Dealing and Miscellaneous Amendments) Regulations 2005 (SI 2005/2789). By virtue of section 47 of the Act, when new standard provisions are made by regulation such provisions shall be deemed to be incorporated in existing licences as from the end of the three months from the date on which the regulations come into operation but it is provided that at any time before the end of that period the holder may apply to the Licensing Authority to direct that the new provisions should not be incorporated or will apply subject to any exemption or modifications specified in the application.

2. It authorises distribution by way of wholesale dealing within the EEA of licensed medicinal products on a general sales list by the licence holder named and storage of such products only on the premises specified.

3. The licence holder must provide and maintain such personnel, equipment and facilities as are necessary to avoid the deterioration of the medicinal products. If any change of premises is proposed prior approval must be sought from the Licensing Authority. Any proposals to make structural alterations to the premises must also be notified to the Licensing Authority. If the business should change hands, the company or person taking over the business will need to obtain their own licence, if they do not already have one, before commencing wholesale dealing in medicinal products.

Attention is drawn to the structure of this licence (as detailed on page 2 of Section 1) and to its completeness in accordance with that structure. This is of particular relevance where the holder of the licence is using it as evidence to a third party in support of claims to carry out those operations and activities to which this licence applies on premises and using personnel covered by this licence.

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SECTION 1 (continued)

1. A medicinal product may be placed on the market in the United Kingdom only in accordance with a marketing authorisation or pursuant to an exemption from the requirements to hold a marketing authorisation granted under the provisions of the Medicines for Human Use (Marketing Authorisations) Regulations 1994 as amended (SI 1994/3144) or by relevant European Community provisions ("an exempt medicinal product"). Wholesale dealers may only distribute such products. This licence only authorises the distribution of medicinal products on a general sales list with a marketing authorisation or product licence.

2. If a licence holder intends to import licensed medicinal products from outside the EEA an application for a manufacturer's licence that authorises import must be made and a licence granted for that purpose. Such a licence requires the holder to have available at all times a Qualified Person who must be named on the licence.

3. If a licence holder intends to import exempt medicinal products into the UK, an application for a manufacturer's licence that authorises import must also be made and a licence granted for that purpose.

4. If a licence holder intends to carry out any manufacture and/or assembly process (e.g. packing, filling or labelling) of medicinal products, an application for a manufacturer's licence must be made and a licence granted for that purpose.

5. A licence may be suspended if any fees are not paid in full as they fall due.

6. The Medicines and Healthcare Products Regulatory Agency (MHRA) acts on behalf of the Licensing Authority established under The Medicines Act 1968 as amended.

7. Further information and specified guidelines may be obtained from the MHRA website (www.mhra.gov.uk).

8. Licence Structure

This Licence is divided into three sections.

(a) Section 1 (this section) identifies the licence holder and holds the authorising name for the issue of the licence. This section would not usually be replaced during routine variations of the licence unless the licence holder details are varied.

(b) Section 2 lists variations to the licence. A replacement section 2 will be issued each time the licence is varied.

(c) Section 3 contains the details relating to each site named on the licence. Where there is more than one site there will be more than one part to Section 3. When a variation is made to the details of a site named in Section 3 the relevant part of Section 3 will be replaced.

(d) The licence holder is required to attach to his licence any replacement pages issued by the Licensing Authority and to mark or destroy superseded pages as to render them invalid.

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SECTION 1 (continued)

9. Licence Holder

(a) WDL Number: WDL 20891 has been granted to –

NAME:	MR R HOLIDAY
COMPANY:	CLF DISTRIBUTION LIMITED
TRADING AS:	
ADDRESS:	UNIT 16, PARKERS CLOSE, DOWNTON BUSINESS CENTRE, SALISBURY, WILTSHIRE, SP5 3RB, UNITED KINGDOM

(b) This licence permits the holder of the licence to distribute by way of wholesale dealing within the EEA medicinal products, on a General Sales List to be stored at the named premises and using the named personnel in accordance with Section 3 of this licence.

(c) This authorisation will continue to remain in force from the date of issue by the Licensing Authority unless suspended, revoked or varied as to the period of its validity or relinquished by the licence holder.

(d) Date granted - 30/07/2008

(e) Authorised by -

Name: Sean Kaiser

(A person authorised to approve on behalf of the Secretary of State for Health.)

Date: 30/07/2008

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SECTION 2

VARIATION HISTORY

This page will be amended if the licence is varied.

Date	Variation Detail
28/01/2004	Initial Application
08/12/2004	Variation to change address.
30/07/2008	Re-issue new format licence.

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SECTION 3 - SITE INFORMATION

The premises –

SITE NAME:	CLF DISTRIBUTION
ADDRESS:	UNIT 16, PARKERS CLOSE, DOWNTON BUSINESS CENTRE, SALISBURY, WILTSHIRE, SP5 3RB, UNITED KINGDOM
MHRA SITE NUMBER:	91848

is named on WDL number: WDL 20891 and licensed for the following:

1. Those operations as specified.
2. Those descriptions of products or classes of product as specified.
3. The personnel named to carry out the roles as specified.

Any restrictions or clarifying remarks related to the scope of these Wholesaling operations:

SECTION 3 – SITE INFORMATION (continued)

USE OF PRODUCTS AT SITE	
Administration to human beings	Yes
SITE TYPES	
Distribution only	No
Storage and Handling (picking of goods)	Yes
Procurement/administration only (no storage)	No

PRODUCT CLASSES	
Non-sterile liquids (including solutions, syrups and suspensions)	Not Licensed
Semi-solid non-sterile dosage forms (including non- sterile creams and ointments)	Not Licensed
Solid non-sterile dosage forms (including tablets, capsules, suppositories and powders)	Licensed
Other non-sterile products (as specified)	Not Licensed

SECTION 3 – SITE INFORMATION (continued)

Personnel

Person Number	Responsible Person
	Name
120850	Mr M Watson

Key to Role:

RP –Responsible Person