Status: This is the original version (as it was originally made).

SCHEDULE 1

STANDARD PROVISIONS WHICH MAY BE INCORPORATED IN A MANUFACTURER'S LICENCE RELATING TO THE MANUFACTURE AND ASSEMBLY OF RELEVANT MEDICINAL PRODUCTS

2. The manufacturer's licence holder may use a contract laboratory pursuant to Article 11(2) of Commission Directive 2003/94/EC if operated by a person approved by the licensing authority.