Pharmacopoeias included in the MHRA list, referred to in the definition of a homeopathic medicinal product.

In order to be a homeopathic medicinal product within the meaning of the Human Medicines Regulations, a product must fall within the following definition:

> ‘’Homeopathic medicinal product means a medicinal product prepared from homeopathic stocks in accordance with a homeopathic manufacturing procedure described by
>
> (a) the European Pharmacopoeia; or
>
> (b) in the absence of such a description in the European Pharmacopoeia,
>
> > (i) in relation to a certificate of registration or marketing authorisation for a national homeopathic product in force in Great Britain only, the British Pharmacopoeia, or in a pharmacopoeia used officially in a country that is included in a list published by the licensing authority for this purpose;
> >
> > (ii) in relation to a certificate of registration or marketing authorisation for a national homeopathic product in force in the whole United Kingdom or in Northern Ireland only, in the British Pharmacopoeia or in any pharmacopoeia used officially in an EEA State.’’

In line with this definition, homeopathic medicinal products marketed in Great Britain only must be prepared from homeopathic stocks manufactured in accordance with a homeopathic manufacturing procedure described in the European Pharmacopoeia or in the absence of such a description, in a pharmacopoeia from the following list:

- British Pharmacopoeia
- A pharmacopoeia used officially in an EEA State.

This list will be updated as new entries arise.

1 January 2021