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reference work for all those involved in the manufacture and distribution of medicines in Europe. It is compiled by the UK drug regulatory body, MHRA, and brings together the European and UK guidance documents and information on legislation relating to the manufacture and distribution of medicines for human use.

Rules and Guidance for Pharmaceutical Manufacturers and ... Full form of MHRA is Medicines and Healthcare products Regulatory Agency. This agency is of United Kingdom (UK). This agency is responsible for MHRA audits throughout the world. The companies those comply their GMP regulations can export their pharmaceutical products to UK. The GMP guidelines of MHRA are known as Orange Guide.

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All the GMP regulation are given in this guide that is to be followed in pharmaceuticals according to MHRA guidelines.

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The Orange Guide Rules and Guidance for Pharmaceutical ...

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First published in 1971 the original Orange Guide contained British Good Manufacturing Practice and was entitled " Guide to Good Pharmaceutical Manufacturing Practice ". Not much more than 30 pages in length this voluntary guide was an aid to manufacturers to

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