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EDQM - European Directorate for the Quality of Medicines

european pharmacopoeia 8th edition About the European Pharmacopoeia The European Pharmacopoeia (Ph. Eur.) defines requirements for the qualitative and quantitative composition of medicines, the tests to be carried out on medicines and on substances and materials used in their production.

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The European Pharmacopoeia (Ph. Eur.) 9th Edition has been obsolete since 1 January 2020. Consequently, the 9th Edition online and all previous versions, including the Ph. Eur. archives for 9th Edition clients, have no longer been accessible since 1 March 2020. The Ph. Eur. 10th Edition and its supplements are available to subscribers on the dedicated European Pharmacopoeia online platform.

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The European Medicines Agency (EMA) works with the European Directorate for the Quality of Medicines and HealthCare (EDQM), a directorate of the Council of Europe. The EDQM traces its origins and statutes to a European treaty promoting the elaboration of a common pharmacopoeia in Europe.

European Directorate for the Quality of Medicines and...

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