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The British Pharmacopoeia 2026 edition

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It includes approximately 4,000 monographs, which are legally enforced by the Human Medicines Regulations 2012. Medicinal products or active pharmaceutical ingredients sold or supplied in the UK must comply with the relevant monograph.

All monographs and requirements of the European Pharmacopoeia (Ph. Eur.) are reproduced in the BP, making the BP a convenient and fully comprehensive set of standards that can be used across Europe and internationally.

New for the BP 2026 edition

- 19 new BP monographs and 34 new monographs reproduced from the 11th edition of the Ph. Eur., as amended by issues 11.6 to 11.8
- Two new appendices, four supplementary chapters, and six new infrared reference spectra have been added to the collection
- 12 new BPCRS have been produced and referenced
- 131 amended BP monographs
- All monographs from the Ph. Eur. 11th edition and Ph. Eur. issues up to 11.8
- The Ph. Eur. 12th edition, comprising issues 12.1, 12.2 and 12.3, included as an in-year online and download update

To order, call **+44 (0)1603 696860** or email **pharmahealth@tso.co.uk**

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British Pharmacopoeia

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Price and format options

Single-user online licence = £950

A single-user licence grants access to one designated user and offers a flexible solution – ideal for individuals or smaller organisations. A licence provides access from 1 August 2025 to 31 March 2027 (a total of 20 months).

Optional add-ons include:

- BP Archive (from BP 2014 onwards): £200 per licence
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The six-volume print set remains an essential reference for professionals who require reliable, quick, and uninterrupted access to official standards.

The complete package will no longer be available. Instead, new bespoke packages are being introduced to allow customers greater flexibility at a discounted rate. Distributor and reseller discounts will not apply to these, but you can offer something similar by creating your own bespoke packages.

Title Information

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The British Pharmacopoeia (BP) Commission

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British Pharmacopoeia

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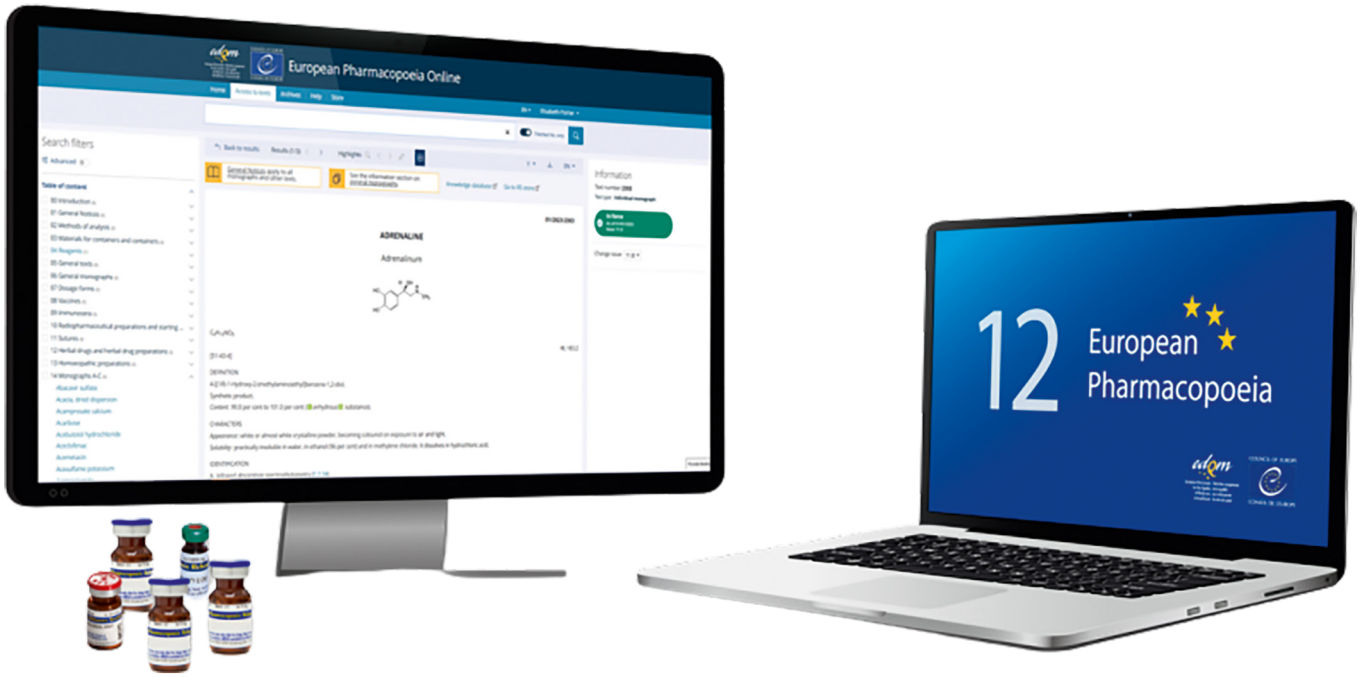

European Pharmacopoeia (Ph. Eur.) 12th Edition

The European Pharmacopoeia (Ph. Eur.) is the primary source of official quality standards for medicines and their ingredients in Europe. Ph. Eur. standards provide a legal and scientific basis for the quality control of a product throughout its life cycle, supporting the pharmaceutical industry and healthcare systems.

The European Directorate for the Quality of Medicines and Healthcare (EDQM) has published the latest edition of the European Pharmacopoeia (Ph. Eur.) as an online licence only. The 12th edition of the European Pharmacopoeia (Ph. Eur.) marks the beginning of a new, online era in European pharmacopoeial standards. Consult the Ph. Eur. texts on a redesigned, user-friendly platform launched alongside the 12th Edition.

The new Ph. Eur. platform features:

- access to all texts currently in force and texts published since volume 11.0
- a text-status traffic light system (green for “in force”, orange for “not yet in force” and red for “no longer in force”)
- in-text menus to navigate between different versions
- improved filter functions, predefined searches and updated result displays
- access to the archives and to complementary information in the Knowledge database

Publication model

The former three-year cycle comprising one edition and eight supplements has been replaced by an annual edition made up of three issues, numbered .1 to .3 (for example, the 12th Edition will consist of Issues 12.1, 12.2 and 12.3). Each of these issues will contain the new and revised texts adopted at one of the three European Pharmacopoeia Commission (EPC) sessions held annually. What will not change is the implementation schedule, which remains January, April and July, respectively, of the year following publication of a text. A detailed calendar presenting the EPC sessions and adoption and implementation dates will be published in due time.

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Product	Format	Subscription no.	Price
European Pharmacopoeia 12th edition	Electronic	7704067	£550

British National Formulary: 89 (March-September 2025)

The British National Formulary (BNF) is the first choice for concise medicines information. Trusted by healthcare professionals across the world to support confident decision-making at the point of care.

The new edition (BNF 85) provides up-to-date guidance on prescribing, dispensing, and administering medicines, plus legal and professional guidelines. Access to the latest edition of the BNF is vital for healthcare professionals, as it reflects current best practice as well as legal and professional guidelines relating to the uses of medicines.

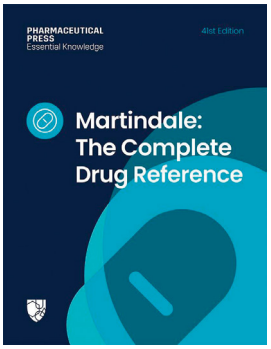
Author: British Medical Association, Royal Pharmaceutical Society of Great Britain
Publisher: BMJ Publishing Group Ltd and Royal Pharmaceutical Society
ISBN: 9780857114877 **Price: £79.00**
Format: Paperback 210 x 148mm **Extent: 1984pp**

BNF for Children (BNFC) 2024-2025

The BNF for Children (BNFC) 2022-2023 provides essential practical information to all healthcare professionals involved in the prescribing, dispensing, monitoring and administration of medicines to children. It addresses a significant knowledge gap in many areas of paediatric practice by providing practical information on the use of medicines in children of all ages from birth to adolescence.

Recommendations in the BNFC have been constructed on the basis of authoritative sources, emerging evidence and best practice guidelines. The content has been carefully validated by a network of paediatric experts and the process is overseen by a paediatric formulary committee.

Author: British Medical Association, Royal Pharmaceutical Society of Great Britain
Publisher: BMJ Publishing Group Ltd and Royal Pharmaceutical Society
ISBN: 9780857114792 **Price: £79.00**
Format: Paperback 210 x 148mm **Extent: 1408pp**



Martindale: The Complete Drug Reference 41st edition

The Complete Drug Reference provides unbiased and evaluated information on drugs and medicines in use around the world.

It is prepared by an experienced team of pharmacists and life scientists who use their professional expertise to select the most clinically relevant and appropriate information from reliable published sources, to provide an unbiased and evaluated digest of the literature.

It contains:

- Over 6,300 drug monographs
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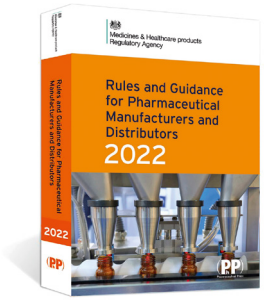
- Nearly 700 treatment reviews, with references from the published literature
- Information to help you identify medicines, the local equivalent and the manufacturer
- Herbals, diagnostic agents, radiopharmaceuticals, pharmaceutical excipients, toxins and poisons

Author: Royal Pharmaceutical Society
Publisher: Pharmaceutical Press
ISBN: 9780857114846 **Price: £650**
Format: Hardback 276 x 219mm **Extent: 4800pp**

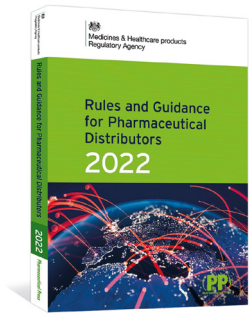


Rules and Guidance for Pharmaceutical Manufacturers and Distributors 2022 The Orange Guide

This is the tenth edition of Rules and Guidance for Pharmaceutical Manufacturers and Distributors, compiled by MHRA. Commonly known as the Orange Guide, it remains an essential reference for all manufacturers and distributors of medicines in Europe. It provides a single authoritative source of European and UK guidance, information and legislation relating to the manufacture and distribution of human medicines.

The new 2017 edition has been updated to incorporate changes and additions made to the detailed European Community guidelines on Good Manufacturing Practice (GMP) and the revised EU Guidelines on Good Distribution Practice (GDP), including Annexes 15 and 16.

Author: MHRA
Publisher: Pharmaceutical Press
ISBN: 9780857114396 Price: £82
Format: Paperback 244 x 172mm Extent: 1140pp

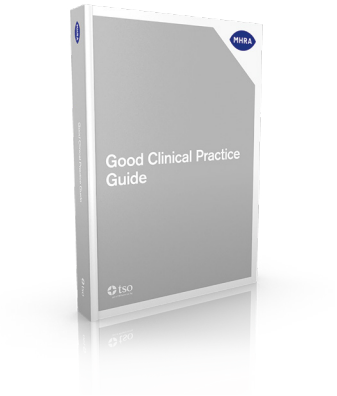


Rules and Guidance for Pharmaceutical Distributors 2022 The Green Guide

This new 2017 edition provides you with a single source of guidance to, and legislation for, the distribution of medicines in Europe and the UK. This tenth edition has been updated to incorporate the revised EU Guidelines on Good Distribution Practice.

The Green Guide reproduces all the elements of the new Rules and Guidance for Pharmaceutical Manufacturers and Distributors 2017 (the Orange Guide) that are relevant to distributors. So if you're involved in the wholesale supply, distribution and brokering of medicines for human use and the distribution of active substances, this is the one-stop guide you need.

Author: MHRA
Publisher: Pharmaceutical Press
ISBN: 9780857114419 Price: £72
Format: Paperback 244 x 172mm Extent: 400pp



Good Clinical Practice Guide

This detailed and authoritative guide covers the legislation, guidance and good practice that relates to the conduct of clinical trials of medicinal products for human use in the UK. It references European and international standards so will also be relevant to organisations conducting trials across Europe and beyond.

Written and produced by the MHRA, this is the only guide on Good Clinical Practice available within Europe which has been produced by a regulatory agency.

Author: MHRA
Publisher: Pharmaceutical Press
ISBN: 9780117081079 Price: £45
Format: Paperback 244 x 172mm Extent: 542pp

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