

2000

USP 24

THE UNITED STATES PHARMACOPEIA

NF 19

THE NATIONAL FORMULARY

By authority of the United States Pharmacopeial Convention, Inc., meeting at Washington, D.C., March 9–12, 1995. Prepared by the Committee of Revision and published by the Board of Trustees

Official from January 1, 2000

Vol. 1

UNITED STATES PHARMACOPEIAL CONVENTION, INC.
12601 Twinbrook Parkway, Rockville, MD 20852

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ISSN 0195-7996

ISBN 1-889788-03-1

Printed by National Publishing, Philadelphia, PA

Continuous Revision of the USP and the NF

The need for continuously refining specifications and updating standards is a natural consequence of the introduction of new drugs and the accelerated growth of knowledge in the pharmaceutical sciences.

In order to keep the *United States Pharmacopeia* and the *National Formulary* abreast of these developments and to maintain the official standards accordingly, the main volume is revised regularly by *Supplements*. *Interim Revision Announcements* are issued between *Supplements* as necessary.

Interim Revision Announcements pertaining to this volume of *USP* and *NF* are published in the journal, *Pharmaceutical Forum*. Reprints of *Interim Revision Announcements* are available on order. A single copy of each *Interim Revision Announcement* is available without charge for each subscription to *USP-NF*. Each request should be sent to USPC at the address shown below. Corrections and revisions included in *Interim Revision Announcements* are incorporated in the next regular *Supplement*.

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- Interested parties submit comments regarding compendial revisions.
- ↓
- Scientific staff liaison sends comments to Subcommittee for review/approval.
- ↓
- Revision proposal appears in *Pharmaceutical Forum* for public comment.
- ↓
- Comments on *PF* are reviewed by Subcommittee.
- ↓
- Subcommittee approves proposal for official adoption.
- ↓
- DSD Executive Committee approves official adoption.
- ↓
- Board of Trustees approves publication [fiduciary responsibility].

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 Herman Jaggy, Karlsruhe, Germany
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 Raymond N. Johnson, Rouses Point, NY
 Jose C. Joseph, North Chicago, IL
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 Toshio Konomi, Osaka, Japan
 Sabine Kopp-Kubel, Geneva, Switzerland
 Santosh Kottayil, Mundelein, IL
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 Johannes Kramer, Eschborn/Taunus, Germany
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 Rod Lenoble, Boulder, CO
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 Hetty Lima, Lincolnshire, IL
 Amy Little, Cary, NC
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 Alan R. Rogers, London, UK
 Martina Rogge, Ingelheim am Rhine, Germany
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 Wayne B. Rose, Mendota Heights, MN
 Michael M. Rosen, New York, NY
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