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norms, standards and guidelines to promote quality assurance and quality control is an integral part of WHO's Constitution and a unique responsibility. It has been endorsed and supported through numerous World Health Assembly resolutions, and more recently in those on the Revised Drug Strategy.

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The WHO Expert Committee on Specifications for Pharmaceutical Preparations was established in the very first World Health Assembly in 1948 to provide advice to WHO and its Member States. Global Regions

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## Pharmaceutical ... Forty Third Meeting Who

This booklet describes the basic working procedures of the World Health Organization (WHO) Expert Committees, and goes on to address specifically the WHO Expert Committee on Specifications for Pharmaceutical Preparations, which has provided formal WHO advice on medicines testing and quality assurance to Member States for more than half a century.

## ~~How does it work? WHO Expert Committee on Specifications ...~~

Geneva, 14-18 October 2019  
New agenda. Follow link below to access a list of items representing the work

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that is currently underway within the Quality Assurance team. Members of the WHO Expert Advisory Panel on The International Pharmacopoeia and Pharmaceutical Preparations serving the WHO Expert Committee on Specifications for Pharmaceutical Preparations and the WHO INN Expert Group.

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Groups serve without remuneration in their personal capacities rather than as representatives of governments or other bodies; their views do not necessarily reflect the decisions or the stated policy of WHO.

## ~~54th report of the WHO Expert Committee on Specifications ...~~

The WHO Expert Committee on Specifications for Pharmaceutical Preparations met in Geneva from 25 to 29 October 2004. Dr Hans V. Hogerzeil, Director ad interim, Essential Drugs and Medicines Policy (EDM), welcomed the Committee members and other participants on behalf of the Director-General, Dr LEE



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Forty-fifth report of the WHO Expert Committee on specifications for pharmaceutical preparations. (WHO technical report series ; no. 961) 1. Pharmaceutical preparations — standards. 2. Technology, Pharmaceutical — standards. 3. Drug industry — legislation. 4. Quality control. I. World Health Organization. II. Series.

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WHO Expert Committee on Specifications for  
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2009 Members Professor Saleh A. Bawazir, Head of  
Drug Sector and Vice-President, Saudi Food and Drug  
Authority (SFDA), Riyadh, Saudi Arabia Mr Jean-Michel

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The WHO Expert Committee on Specifications for Pharmaceutical Preparations met in Geneva from 31 May to 4 June 1999. The meeting was opened on behalf of the Director-General by Dr M. Scholtz, Executive Director of Health Technology and Pharmaceuticals, who stressed that it was of the utmost importance that WHO should v igor-

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standards and guidelines for the quality assurance of medicines. Standards are developed by the Committee through worldwide consultation and an international

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