

Food and Drug Administration
[Docket No. FDA-2015-N-0540]

Homeopathic Product Regulation: Evaluating the Food and Drug Administration's Regulatory Framework After a Quarter-Century; Public Hearing

Presenter: Dr Peter Fisher MB,BChir, FRCP, FFHom

Presentation Outline

- 1) Background / Credentials
 - a. Current positions:
 - i. Clinical director and director of research, Royal London Hospital for Integrated Medicine
 - ii. Fellow, Royal College of Physicians and of the Faculty of Homeopathy
 - b. Accreditation, board certification, and degree
 - i. Homeopathy
 - ii. Rheumatology
 - c. Affiliations
 - i. National Institute for Health and Care Excellence's external advisory panel
 - ii. WHO Expert Advisory Panel on Traditional and Complementary Medicine
- 2) Consumer/patient attitudes toward homeopathy
 - a. Patients report benefit
 - i. Patient outcomes surveys show condition improvement has impact on daily lives
 - b. Growth and use in Europe
 - i. UK = 10% growth
 - ii. France = 98% of retail pharmacies stock homeopathy products; >50% of consumers report use
 - iii. Germany = 30-40% of consumers report use
- 3) Healthcare provider attitudes toward homeopathy
 - a. Favorable, especially in insurance-based systems
- 4) Data sources
 - a. CORE-HOM database
 - i. Searchable by diagnosis, trial design, and medication use
 - ii. Comparative effectiveness research
- 5) Consumer labeling
 - a. Education on homeopathic vs herbal
 - b. Distinctions on label & in-store supporting materials