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Description. Contents. This new 2017 edition of Rules and Guidance for Pharmaceutical Distributors (the Green Guide), provides you with a single source of guidance to, and legislation for, the distribution of medicines in Europe and UK. 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.

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The Rules and Guidance for Pharmaceutical Distributors aka "The Green Guide" is the authoritative resource for wholesalers and brokers. Manufacturers have their own guidance - Rules and Guidance for Pharmaceutical Manufacturers and Distributors (Orange Guide) 2017

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This compliance guidance is intended to assist companies that develop, manufacture, market, and sell pharmaceutical drugs or biological products (pharmaceutical manufacturers) in developing and implementing internal controls and procedures that promote adherence to applicable statutes, regulations, and requirements of the federal health care programs¹ and in evaluating and, as necessary, refining existing compliance programs.

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This title is an essential reference work for all those involved in the distribution of medicines in Europe. It reproduces relevant parts of Rules and Guidance for Pharmaceutical Manufacturers and Distributors (commonly known as the Orange Guide) specific to wholesale supply and distribution of medicines for human use.

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by pharmacyStudent on Nov 17, 2017. Book Reviews: Since the 2002 edition of Rules and Guidance for Pharmaceutical Manufacturers and Distributors (the “Orange Guide”) there have been many changes and additions to the detailed European Community guidelines on Good Manufacturing Practice (GMP). In addition, there is a new Directive dealing specifically with GMP and the Community code relating to medicinal products for human use (Council Directive 2001/83/EC) has itself been the subject of ...

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Rules and Guidance for Pharmaceutical Distributors 2007, Medicines and Healthcare Products Regulatory Agency, Jan 1, 2007, Medical, 69 pages. Compiled by the Medicines and Healthcare products Regulatory Agency (MHRA), this new publication provides guidance for distributors of medicines for human use in Europe.

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