

INFORMATION SOCIETY IN PHARMACEUTICALS PHARMACEUTICALS POLICY AND LAW 4

Jan 20, 2021



[Information Society In Pharmaceuticals Pharmaceuticals Policy And Law 4](#)

Information Society in Pharmaceuticals (Pharmaceuticals Policy and Law, 4) | J. L. Valverde | ISBN: 9781586031787 | Kostenloser Versand für alle Bücher mit Versand und Verkauf durch Amazon.

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Acknowledgement Acknowledgement J.L. Valverde 2001-01-01 00:00:00 Pharmaceuticals Policy and Law 4 (2001) 3â 5 IOS Press This monograph of Pharmaceuticals, Policy and Law is focused on the issues concerning the Information Society in Pharmaceuticals. We have had the very valuable collaboration of Esteban Gonzalez-Juarros, from the Pharmaceuticals and Cosmetics Unit within the European Commission.

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A pharmaceutical corporation that wishes to meet its obligations to the developing world or to some chosen part of it should ensure the following formal legal obligations: (1) its products meet international standards for quality, efficacy, and safety, (2) the products are accessible, particularly in terms of price, (3) its research programs take account of third-world needs, (4) the ...

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In the dozen years between 2001 and 2012, at least 38 states had passed some type of state pharmaceutical assistance law. 2 However, the number of operating programs is on the decline, due to Medicare Part D benefits and expanded discounts available under the Affordable Care Act, starting in 2011. Most use state funds to help pay for a portion of the cost of pharmaceuticals for eligible ...

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Here we propose an approach to identifying key information across the pharmaceutical system that can help policy-makers and other stakeholders strengthen accountability and we illustrate its application. In developing this approach, we have drawn on the experience of two international programmes that focused on transparency, accountability and anticorruption practices in the pharmaceutical system.

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PBS Information Management Section, Pharmaceutical Policy Branch (2016), 'Expenditure and prescriptions twelve months to 30 June 2016', Canberra: Department of Health. Philippon , D. J. and Braithwaite , J. (2008), ' Health system organization and governance in Canada and Australia: a comparison of historical developments, recent policy changes and future implications ', Healthcare ...

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However, the new medicine law, which included several progressive, but controversial, pricing policy components, such as generic substitution and parallel importation, was challenged in court by the research-based local and international pharmaceutical industry and delayed for three years. For various political reasons, some of the provinces remained skeptical and hesitated to become partners ...

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@inproceedings{Kanavos2011ThePD, title={The pharmaceutical distribution chain in the European Union: structure and impact on pharmaceutical prices}, author={Panos Kanavos and Willemien Schurer and Sabine Vogler}, year={2011} } figure 3.1 table 3.1 figure 3.2 figure 3.3 figure 3.4 table 4.1 table 4.2 ...

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Tort claims related to products such as pharmaceuticals are authorized by state law, either common law or statutory. Federal law has no statutory provisions authorizing similar causes of action, and there is no general common law . Thus, particularly in personal injury tort claims, the absence of a state law remedy means the absence of any remedy. In Ledbetter, Merck's argument was in essence ...

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Corruption in the health sector can hurt health outcomes. Improving good governance can in turn help prevent health-related corruption. We understand good governance as having the following characteristics: it is consensus-oriented, accountable, transparent, responsive, equitable and inclusive, effective and efficient, follows the rule of law, is participatory and should in theory be less ...

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Pursuant to Chapter 4 of the Medicinal Products Act, a pharmaceutical product will, in connection with being granted marketing authorisation, be classified either as a prescription-only or a non-prescription pharmaceutical product. The MPA will decide the classification for the pharmaceutical product depending on its intended use and characteristics. A prescription-only pharmaceutical product must be subject to either the prescription or requisition regulation in order to reach the patient ...

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1.4 About our Operating Policies and Procedures. The operating policies and procedures contained in this document are intended to provide guidance on the way in which PHARMAC carries out its statutory role and functions. Supporting information about PHARMAC's operations is available on our website however this information does not constitute part of the operating policies and procedures. 1.5 ...

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