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The past several decades have been a time of rapid globalization in the development, manufacture, marketing, and distribution of medical products and technologies. Increasingly, research on the safety and effectiveness of new drugs is being conducted in

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such standards could improve the efficiency and clarity of the drug development and evaluation process and, ultimately, promote and enhance product quality and the public health. To explore the need and prospects for greater international regulatory

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harmonization for drug development, the IOM Forum on Drug Discovery, Development, and Translation hosted a workshop on February 13-14, 2013.

Discussions at the workshop helped identify principles, potential approaches, and strategies to

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advance the development or evolution of more harmonized regulatory standards. This document summarizes the workshop.

This exciting new text illustrates and advances the argument that

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policy coordination. Examining and analysing both positive and negative examples of this independence, this text is a valuable resource on the topic of the internal workings of IOs, providing the richest and most focused textbook so far dealing

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with the capacity of IOs for independent action in international politics. It is essential reading for all students of international organizations.

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