
New Drug Development A Regulatory Overview

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'Guide to Drug Development A Comprehensive Review

September 7th, 2008 - Written by one of the foremost authorities on clinical trials drug development and regulatory affairs Guide to Drug Development is a comprehensive review of the principles and activities involved in developing new drugs devices and other medical products'

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April 29th, 2018 - Since 2007 the US Food and Drug Administration FDA has issued a handful of special priority review vouchers which allow its recipient to expedite the review of any one of its new drug products'

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April 28th, 2018 - Who is this class for This class is for professionals and students interested in better understanding the process of drug development in the pharmaceutical and biotech industry as well as anyone who wants to learn how new drugs are developed after they received an IND'

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'Services Clinical Trials Regulatory Affairs

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zur Erlangung des Titels 'Master of Drug Regulatory Affairs?'

'Regulatory Focus? RAPS

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