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Food and Drug Administration vs European Medicines Agency: Review times and clinical evidence on novel drugs at the time of approval

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British Journal of Clinical Pharmacology, 2020, 86, 170-174.

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10	Food and Drug Administration vs European Medicines Agency: Review times and clinical evidence on novel drugs at the time of approval. <i>British Journal of Clinical Pharmacology</i> , 2020 , 86, 170-174	3.8	6
9	Approval of Cancer Drugs With Uncertain Therapeutic Value: A Comparison of Regulatory Decisions in Europe and the United States. <i>Milbank Quarterly</i> , 2020 , 98, 1219-1256	3.9	10
8	Evolving Landscape of New Drug Approval in Japan and Lags from International Birth Dates: Retrospective Regulatory Analysis. <i>Clinical Pharmacology and Therapeutics</i> , 2021 , 109, 1265-1273	6.1	5
7	Transparency in European Medicines Agency and US Food and Drug Administration Decision Making: Is It Possible to Identify the Rationale for Divergences in Approved Indication From Public Assessment Reports?. <i>Clinical Therapeutics</i> , 2021 , 43, 888-905	3.5	
6	Current challenges in applying gene-driven therapies in clinical lung cancer practice. <i>World Journal of Clinical Oncology</i> , 2021 , 12, 656-663	2.5	0
5	Recent Clinical Trials on Natural Products and Traditional Chinese Medicine Combating the COVID-19. <i>Indian Journal of Microbiology</i> , 2020 , 61, 1-6	3.7	7
4	A detailed analysis of expedited regulatory review time of marketing authorization applications for new anticancer drugs in the US and EU.. <i>Clinical and Translational Science</i> , 2022 ,	4.9	0
3	Transferability of real-world data across borders for regulatory and health technology assessment decision-making. 9,		0
2	Regional disparity in first-in-class anticancer drug development in the US, EU, and Japan. 2023 ,		0
1	The role of drug regulatory authorities and health technology assessment agencies in shaping incentives for antibiotic R&D: a qualitative study. 2023 , 16,		0