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Drug Approval Pathways: Deciding Parameters Involve in Approvals Drugs in United States & Europe

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Abstract:

Developing a new drug requires incredible amount of research work in chemistry, producing, controls, preclinical and clinical preliminaries. Drug analysts in administrative organizations around the globe bear the obligation of assessing whether the exploration information bolster the security, adequacy and quality control of a new drug product to serve the general public health. In this present exertion, study communicates the drug endorsement pathway and administrative necessities as indicated by United States Food and Drug Administration (USFDA) and European Medical Agency (EMA).

Key words: *Endorsement Pathway, USFDA, EMA.*

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