

Special Issue

Drug Discovery, Development and Regulatory Affairs

Message from the Guest Editors

The discovery and development processes, inclusive of regulatory affairs, manufacturing, and post-market operations, are cornerstones in the commercialization of innovative pharmaceutical and biotechnology products to address unmet clinical needs. Effective drug discovery processes provide a continuous pipeline of candidates for drug development, which in turn generates approvable compounds. Novel approaches across the fields of drug discovery, drug development, and regulatory affairs are of paramount interest to multiple healthcare industry stakeholders, including patients, biopharmaceutical manufacturers, clinicians, provider institutions, and payers. This Special Issue, “Drug Discovery, Drug Development and Regulatory Affairs”, will focus on novel approaches (original research articles) as well as reviews of current practices surrounding the continuum of taking products from the beaker to the bedside. It will present innovative research involving aspects of drug discovery, drug development, regulatory science, commercialization, and market access for both small-molecule pharmaceuticals, as well as biotechnology products.

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Deadline for manuscript submissions

closed (31 May 2019)



International Journal of Molecular Sciences

an Open Access Journal
by MDPI

Impact Factor 4.9
CiteScore 9.0
Indexed in PubMed



mdpi.com/si/20849

*International Journal of
Molecular Sciences*
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