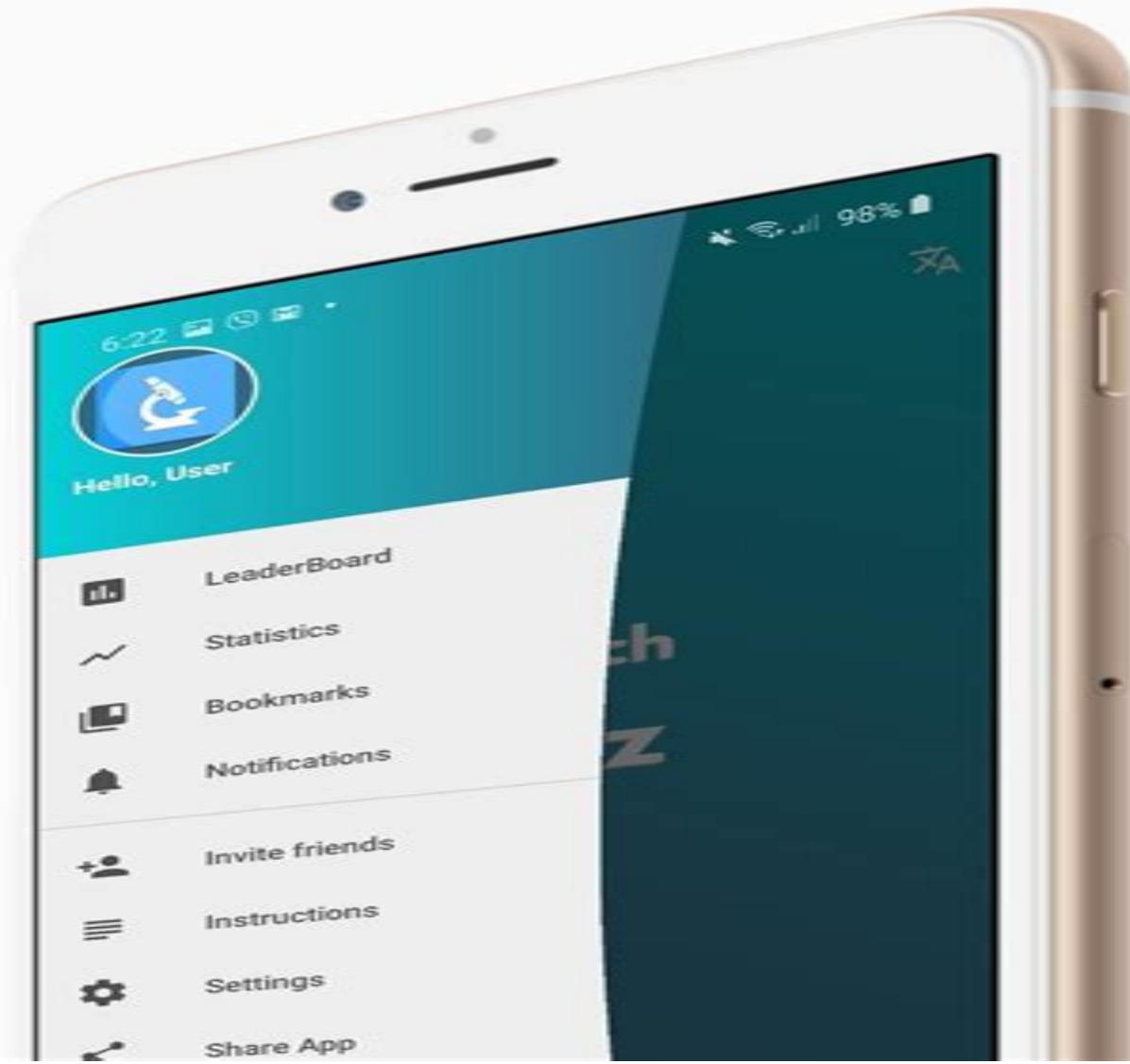


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the thalidomide tragedy of the mid 20th century In the past decades a lot of progress has been made in the application and evolution of regulatory legislative procedures for the benefit of patients and public health but also in driving and enabling innovation Medicines development is a global endeavor and exchange of experience and knowledge between regulatory agencies working under different jurisdictions is not only necessary but seen increasingly as essential There are several factors playing a role in this process Patients are increasingly well informed about their disease existing treatments and novel developments on the horizon and share information with others across regions Scientific progress is facing a rapid development with impressive achievements in medicine pharmacology basic science and technical disciplines With this we also face several novel challenges inherent with the possibilities that technology can provide e g whole genome sequencing AI etc Regulatory bodies are facing new challenges with decisions to be made faster for the sake of not delaying the availability and access to newly developed treatments The COVID 19 crisis has only reiterated the need for collaboration coherence and solidarity on a global scale With this in mind we as guest editors are opening a research topic which is intended to invite experts from around the globe to contribute their views on the regulation of medicines today and in the future More specifically we expect manuscripts related to the following topics Summary of the key aspects of the 3 main regulatory frameworks globally USA Europe and Japan commonalities and differences Future changes to EU legislation case study of drivers for change Global trends in regulatory science that impact legislation drug development and patient access The importance of regulatory collaboration and harmonization and the role of not for profit organizations Emerging regulatory frameworks Africa Latin America and the ROW Reliance regulation pros and cons is this the direction of travel Novel regulatory procedures case studies of innovation in regulatory practice Future proofing the regulatory framework and the role of horizon scanning Scientific advice and other regulatory support tools Clinical trials centralized versus decentralized Digital revolution impact of AI and ML Drug device combinations Specific regulatory pathways for innovative medicines Possibilities for regulatory convergence Type of articles expected but not limited to are Original articles Reviews

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Digital Remote Patient Monitoring in Neurodegenerative Diseases Graham Jones,Amit Khanna ,Diane Stephenson,2026-01-06 The development of sensitive miniature electronic digital health sensors has stimulated growth of a multi billion dollar consumer products for the health tech industry Devices like wrist and finger worn wearables smartphones specialized peripherals and home based IoMT devices are capable of utilizing these technologies coupled with associated algorithms to accurately measure vital signs cognitive functions and fine motor movements This now opens the opportunity for measuring subtle changes in chronic degenerative diseases using a suite of digital tools with the potential to detect a change within a reasonable time during the early course of the disease These slowly progressing conditions manifest clinical symptoms and functional impairment over a period of several years Thus deploying technologies to facilitate early diagnosis characterization and monitoring disease progression longitudinally might enable informed decisions on treatment care and early interventions They can also provide key data to underscore the efficacy of therapeutics designed to prevent delay or modify the course of the disease addressing an unmet need in this area and generating data of relevance to the regulatory authorities and Payors The need for use of such technologies is becoming acute as the rising prevalence of diseases such as Alzheimer s Parkinson s and multiple sclerosis represent an enormous societal economic and healthcare burden globally In this Research Topic we will highlight recent developments in remote patient monitoring derived from experts in the academy pharmaceutical industry government laboratories and research foundations Examples range from routine measurements of blood pressure heart rate variability and sleep cycles to more complex interrogation of acoustic oculomotor and ballistographic signals which can provide insight to disease states to enable early intervention Additionally specialized digital programs can give insight to cognitive state and function If captured longitudinally they can provide informative data on disease progression and classification A benefit of digital monitoring in the early stages of disease lies in the potential to explore disease preventing approaches during limited time periods of a clinical study and offer long term monitoring via registries and electronic health records Based on the considerable promise of these digital technologies we are delighted to showcase cutting edge advances and emerging directions in the field through this exciting collection which we cordially welcome authors to consider submitting to Potential

topics could include but are not restricted to the following subject areas Use of digital technologies to facilitate early diagnosis characterization and monitoring of neurodegenerative disease Data driven approaches to interrogate the efficacy of symptomatic and disease modifying therapies Development of standards and metrics to guide registration validation and regulatory approaches Future remote monitoring environments from clinical studies to IoMT at home Ethical considerations around the capture storage transmission and ownership of patient captured data Clinical and real world evaluation of sensors and sensing technologies in neurodegenerative disease Progress in human factor engineering in the design of patient friendly remote monitoring devices Passive versus active monitoring the impact of patient friction burden on longitudinal data capture Strategies to align and translate patient voice into clinical meaningfulness measures **MIT's Technology Review** ,1998 **The Software Encyclopedia 2000** Bowker Editorial Staff,2000-05 **Ontario Medical Review** ,1996

Digital Review of Asia Pacific 2007/2008 Idrc,2008-01-07 The Digital Review of Asia Pacific provides an overview of how information and communication technology ICT is being diffused throughout the Asia Pacific region to facilitate socio economic development This third annual review provides an analytical overview of the state of ICT4D in the Asia Pacific region It covers 31 countries and economies including for the first time North Korea Each country is dealt within a separate chapter which attempts to provide comprehensive coverage of the various aspects of ICT4D in the concerned country at the time of writing in 2006 The chapters have been written by a team of authors representing different sectors such as government academia industry and civil society The American Biology Teacher ,2000 **Proceedings of the Sawtooth Software Conference** ,2005 Harvard Business Review ,2001 Review Copy Larry E. Long,Sara Long,2000-12

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