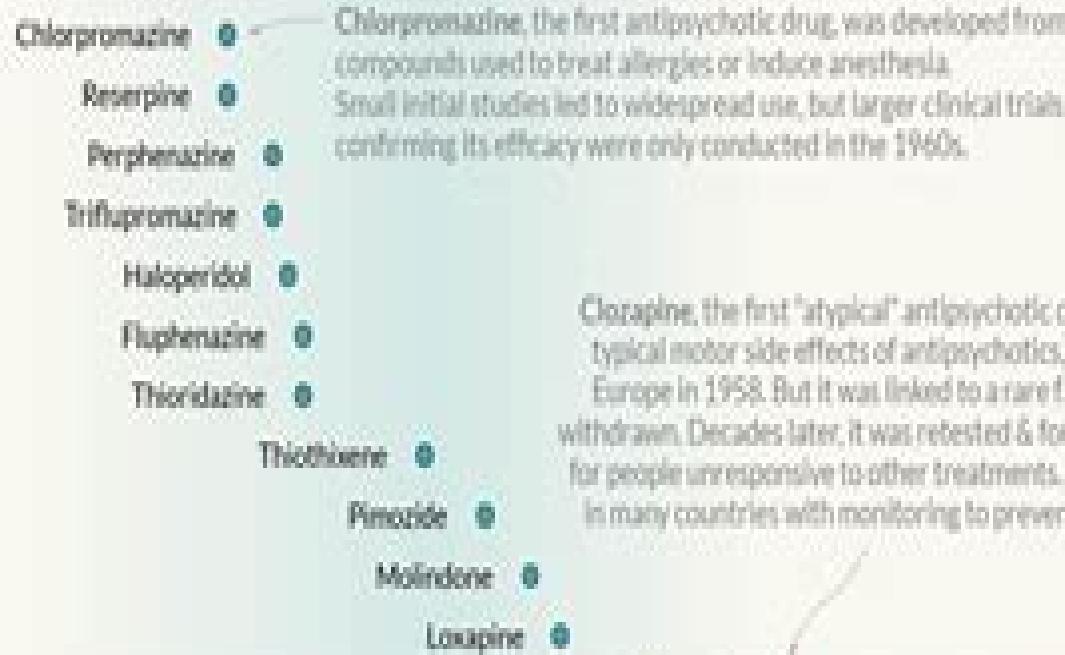


Timeline of the development of antipsychotic drugs

The year when each antipsychotic drug was first licensed in the United States. This is shown alongside key changes in drug regulation and approval by the Food and Drug Administration (FDA).

"Conventional" antipsychotics

The first generation of antipsychotic drugs were initially called "major tranquilizers" for their sedative and calming effects, or "neuroleptics" for their side effects related to movement - such as spasms, stiffness, and tremors.



"Atypical" antipsychotics

Following clozapine, many more atypical antipsychotic drugs were developed. They generally have lower risks of side effects of movement and sexual dysfunction, but also generally carry greater risks of weight-gain, diabetes, and heart complications.

Clozapine, the first "atypical" antipsychotic drug, which lacked typical motor side effects of antipsychotics, was developed in Europe in 1958. But it was linked to a rare fatal condition and withdrawn. Decades later, it was retested & found very effective for people unresponsive to other treatments. It was introduced in many countries with monitoring to prevent potentially fatal side effects.

Clozapine

Risperidone

Olanzapine

Quetiapine

Zimelidine

Aripiprazole

Paliperidone

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Gang Liu

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Dose Optimization in Drug Development Rajesh Krishna, 2006-05-01 This reference provides a concise overview of the key principles in dose selection and optimization and demonstrates applicability to recent successful new drug applications. Compiling key issues and current research on safety, efficacy and clinical pharmacology and PK/PD, this volume critically highlights the multidisciplinary nature of drug dev.

Appropriate Dose Selection - How to Optimize Clinical Drug Development J. Venitz, W. Sittner, 2007-03-06 Optimal dose individualization has become more important in improving clinical efficacy and safety given the variability in drug response e.g. due to concurrent illnesses or co-medication. Therefore, the role of optimal dose finding in early clinical drug development so as to maximize successful clinical use is emphasized. The continued use of biomarkers based on the known pharmacology of the drug and/or biology of the underlying disease along with exposure-response evaluation throughout all phases of drug development can quantitatively integrate clinical pharmacology knowledge, provide early proof of concept and help in rational dose selection and rational drug product labeling for clinical use.

Translational Medicine in CNS Drug Development George G. Nomikos, Douglas E. Feltner, 2019-06-18 Translational Medicine in CNS Drug Development Volume 29 is the first book of its kind to offer a comprehensive overview of the latest developments in translational medicine and biomarker techniques. With extensive coverage on all aspects of biomarkers and personalized medicine and numerous chapters devoted to the best strategies for developing drugs that target specific disorders, this book presents an essential reference for researchers in neuroscience and pharmacology who need the most up-to-date techniques for the successful development of drugs to treat central nervous system disorders. Despite increases in the number of individuals suffering from CNS related disorders, the development and approval of drugs for their treatment have been hampered by inefficiencies in advancing compounds from preclinical discovery to the clinic. However, in the past decades, game changing strides have been made in our understanding of the pathophysiology of CNS disorders and the relationship of drug exposure in plasma and CNS to pharmacodynamic measures in both animals and humans. Includes comprehensive coverage of biomarker tools and the role of personalized medicine in CNS drug development. Discusses strategies for drug development for a full range of CNS indications with particular attention to neuropsychiatric and neurocognitive disorders. Includes chapters written by international experts from industry.

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