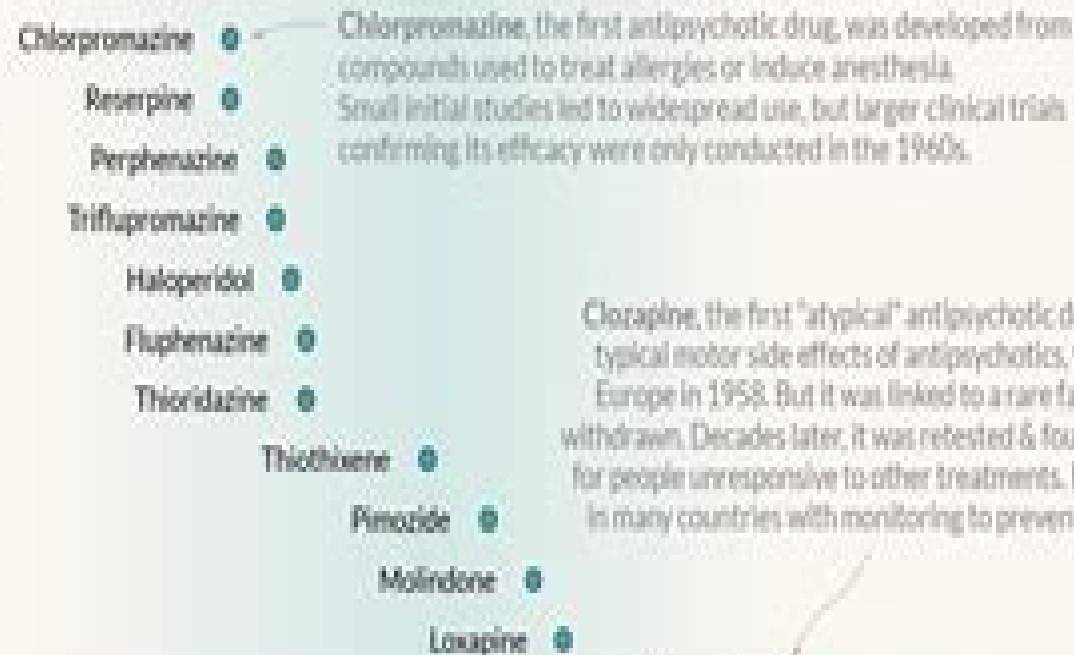


Timeline of the development of antipsychotic drugs

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

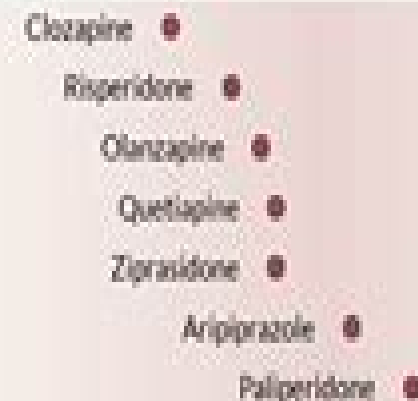
The first generation of antipsychotic drugs were initially called "major tranquilizers" for their sedative and calming effects, or "neuroleptics" for their side effects related to movement – such as spasms, stiffness, and tremors.



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 But it was linked to a rare fatal condition and withdrawn.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.

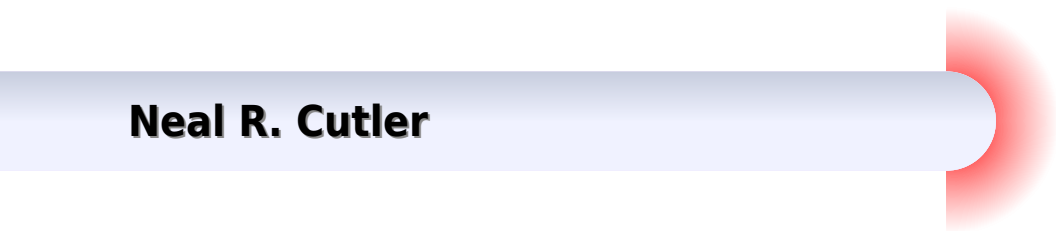
"Atypical" antipsychotics

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.



Optimizing The Clinical Development Of Antipsychotic Drugs

Neal R. Cutler



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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**

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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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Accelerating CNS Drug Development Neal R. Cutler, 1998-04-15 There has been a remarkable increase in the number of available drugs for disorders of the CNS in recent years and there are many more novel compounds in the pipeline But most of these compounds will be in development for many years Although it is vital that trials are thorough there are ways of accelerating development so that these vital drugs are available sooner Based on evidence from real examples this book examines options on the design of trials and questions the need for prolonged testing in some areas It is therefore an essential text for all those involved in the development of CNS drugs **Clinical Handbook of Psychotropic Drugs**

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