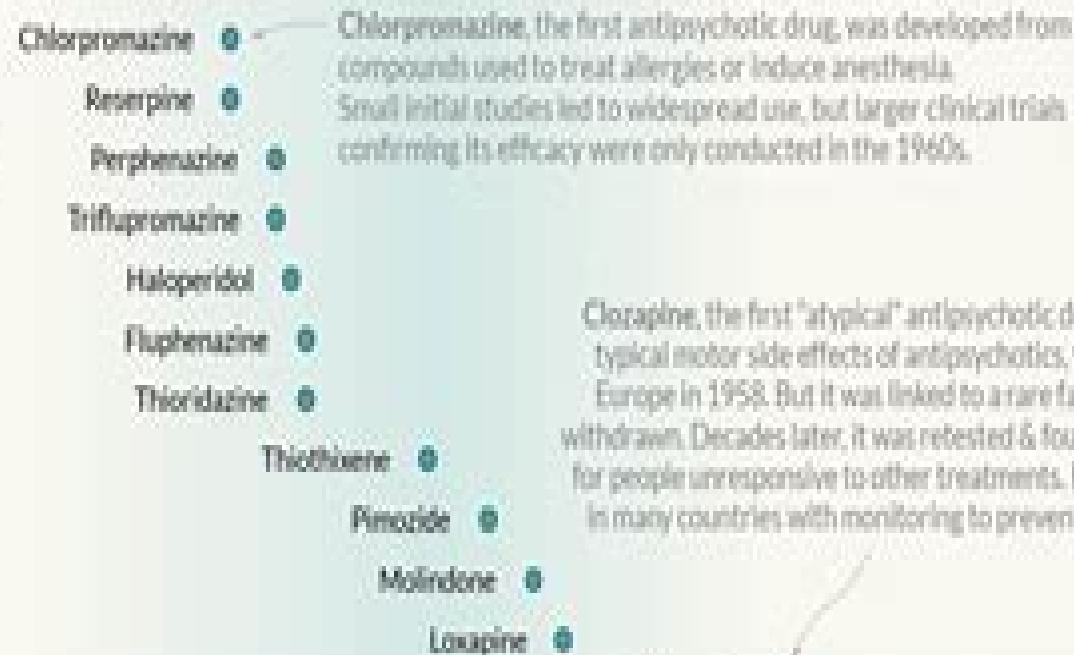


Timeline of the development of antipsychotic drugs

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

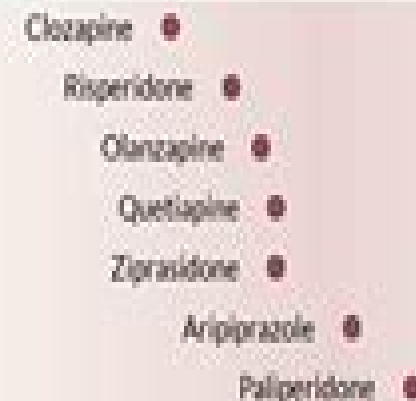
The first generation of antipsychotic drugs were initially called "major tranquilizers" for their sedative and calming effects, or "neuroleptics" for their side effects related to movement – such as spasms, stiffness, and tremors.



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 But it was linked to a rare fatal condition and withdrawn.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

Gang Liu



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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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Table of Contents Optimizing The Clinical Development Of Antipsychotic Drugs

1. Understanding the eBook Optimizing The Clinical Development Of Antipsychotic Drugs
 - The Rise of Digital Reading Optimizing The Clinical Development Of Antipsychotic Drugs
 - Advantages of eBooks Over Traditional Books
2. Identifying Optimizing The Clinical Development Of Antipsychotic Drugs
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Optimizing The Clinical Development Of Antipsychotic Drugs
 - User-Friendly Interface
4. Exploring eBook Recommendations from Optimizing The Clinical Development Of Antipsychotic Drugs
 - Personalized Recommendations
 - Optimizing The Clinical Development Of Antipsychotic Drugs User Reviews and Ratings
 - Optimizing The Clinical Development Of Antipsychotic Drugs and Bestseller Lists
5. Accessing Optimizing The Clinical Development Of Antipsychotic Drugs Free and Paid eBooks
 - Optimizing The Clinical Development Of Antipsychotic Drugs Public Domain eBooks

- Optimizing The Clinical Development Of Antipsychotic Drugs eBook Subscription Services
- Optimizing The Clinical Development Of Antipsychotic Drugs Budget-Friendly Options
- 6. Navigating Optimizing The Clinical Development Of Antipsychotic Drugs eBook Formats
 - ePub, PDF, MOBI, and More
 - Optimizing The Clinical Development Of Antipsychotic Drugs Compatibility with Devices
 - Optimizing The Clinical Development Of Antipsychotic Drugs Enhanced eBook Features
- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Optimizing The Clinical Development Of Antipsychotic Drugs
 - Highlighting and Note-Taking Optimizing The Clinical Development Of Antipsychotic Drugs
 - Interactive Elements Optimizing The Clinical Development Of Antipsychotic Drugs
- 8. Staying Engaged with Optimizing The Clinical Development Of Antipsychotic Drugs
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Optimizing The Clinical Development Of Antipsychotic Drugs
- 9. Balancing eBooks and Physical Books Optimizing The Clinical Development Of Antipsychotic Drugs
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection Optimizing The Clinical Development Of Antipsychotic Drugs
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Optimizing The Clinical Development Of Antipsychotic Drugs
 - Setting Reading Goals Optimizing The Clinical Development Of Antipsychotic Drugs
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Optimizing The Clinical Development Of Antipsychotic Drugs
 - Fact-Checking eBook Content of Optimizing The Clinical Development Of Antipsychotic Drugs
 - Distinguishing Credible Sources
- 13. Promoting Lifelong Learning
 - Utilizing eBooks for Skill Development
 - Exploring Educational eBooks

14. Embracing eBook Trends

- Integration of Multimedia Elements
- Interactive and Gamified eBooks

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