

Annual Report 2021

Reporting Desk for New Drugs

What is the Reporting Desk for New Drugs?

The Reporting Desk for New Drugs is a large-scale national surveillance study in the Netherlands, which is coordinated by the Trimbos Institute. Through this surveillance study, data are collected about new psychoactive substances in the Netherlands. The Reporting Desk for New Drugs reports on which new psychoactive substances had been discovered to be present in the Netherlands, and if available, in which quantities these new psychoactive substances have been detected on the illicit drug market during the last year.



Organization: The Reporting Desk for New Drugs assembles, analyzes, and reports on data collected by a variety of organizations in the Netherlands about the production, trade, and consumption of new psychoactive substances. The Customs Laboratory of the Netherlands and the Netherlands Forensic Institute provide data about seized new psychoactive substances that have been sent to their laboratories for analysis. The Drugs Information and Monitoring System (DIM5) provides data about new psychoactive substances that have been detected in consumer samples submitted for laboratory analysis at a drug checking service¹. The Monitor Drug-related Incidents (MDI) and the Dutch Poisons Information Centre (DPIC) share data about adverse health-related events related to the use of new psychoactive substances. Additional information about new psychoactive substances being used in the Netherlands is also collected from pre-selected online discussion boards about drugs.

Application: The annual report from the Reporting Desk for New Drugs is used by the Ministry of Health, Welfare and Sports in the Netherlands and the Coordination point for Assessment and Monitoring new drugs (CAAM²) to assess the distribution and possible risks associated with the sale, transport, and use of new psychoactive substances in the Netherlands.

What are new psychoactive substances?

The European Monitoring Centre for Drugs and Drug Addiction (EMCDDA) defines new psychoactive substances as "substances of abuse, either in a pure form or a preparation, that are not controlled by the 1961 Single Convention on Narcotic Drugs or the 1971 Convention on Psychotropic Substances, but which may pose a public health threat"³. The Reporting Desk for New Drugs focuses on substances that have been produced in and introduced to the illicit drug market for their psychoactive properties since the early 2000s.

A few substances, such as 2C-B, GHB/GBL, DMT, and ketamine, have also been included in the Reporting Desk for New Drugs. These substances are being monitored by the European Monitoring Centre for Drugs and Drug Addiction under the framework of the European Joint Action on new synthetic drugs.

New psychoactive substance groups

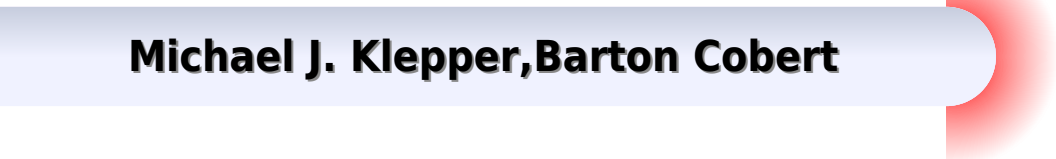
New psychoactive substances are usually classified according to their chemical structure or pharmacological properties. These classifications include:

- Synthetic cannabinoids
- Cathinones
- Phenethylamines
- Tryptamines
- Piperazines
- Other new psychoactive substances:
 - arylcyclohexylamines
 - synthetic opioids (i.e. fentanyl derivatives)
 - new benzodiazepines

Currently, there is no nationwide surveillance system for new psychoactive substances in the Netherlands, which means that underreporting of all new psychoactive substances is likely. However, the information from the Reporting Desk for New Drugs gives an indication of the situation regarding new psychoactive substances being present in the Netherlands on the illicit drug market.

New Drugs Annual Update

Michael J. Klepper, Barton Cobert



New Drugs Annual Update:

New Drugs Bruce H. Woolley, 1979 *New Drugs* Bruce H. Woolley, Edward S. Brady, 1976 **Merck's Annual Report of Recent Advances in Pharmaceutical Chemistry and Therapeutics**, 1912 *Annual Report* National Cancer Institute (U.S.). Division of Cancer Treatment, 1976 **New Drug Approval Process** Richard A. Guarino, Richard Guarino, 2016-04-19

The thoroughly revised Fifth Edition of *New Drug Approval Process* supplies readers with the latest global changes that affect pharmaceutical product approval and influence how new products are researched and marketed. Updated chapters include advances in international regulatory requirements including ICH guidelines and harmonization a step by step

Federal Register, 2013-05 Current Catalog National Library of Medicine (U.S.), 1983 First multi year cumulation covers six years 1965-70 *Guidebook for Drug Regulatory Submissions* Sandy Weinberg, 2009-03-11 Destined to become every regulatory director's essential desktop companion Professionals working to submit major documents to the Food and Drug Administration FDA are guaranteed to encounter numerous unexpected and daunting hurdles *Guidebook for Drug Regulatory Submissions* offers a readable and clearly written road map for effective submission of documents for required regulatory reviews during drug development Demystifying this complex high stakes process author and nationally recognized drug regulation expert Sandy Weinberg presents professionals with authoritative tips tools and advice including suggestions for preparation checklists for submission an FDA evaluation tool for review and copies of relevant FDA guidelines As well vital information is provided on the most common types of submissions including Meeting Requests Orphan Drug Applications Investigatory New Drug Applications INDAs New Drug Applications NDAs 505 b 2 NDAs Abbreviated New Drug Applications ANDAs Annual Report This reference also explores the pressures affecting the industry and the general public as well as how these pressures will change the general nature and specific aspects of the submissions process over the near future In addition retired Canadian trade consul and regulatory consultant Carl Rockburne guest authors a chapter comparing the FDA process to the four other major regulatory environments of Canada the European Union Japan and Australia *Guidebook for Drug Regulatory Submissions* is more than a useful guide it is an essential tool to be kept on the desk of every regulatory director submissions manager vice president of Regulatory Affairs and Food and Drug Administration reviewer responsible for the process of drug regulatory submissions

Drug Safety Data Michael J. Klepper, Barton Cobert, 2010-09-15 *Drug Safety Data How to Analyze Summarize and Interpret to Determine Risk* was selected for The First Clinical Research Bookshelf Essential reading for clinical research professionals by the Journal of Clinical Research Best Practices *Drug Safety Data How to Analyze Summarize and Interpret to Determine Risk* provides drug safety pharmacovigilance professionals pharmaceutical and clinical research scientists statisticians programmers medical writers and technicians with an accessible practical framework for the analysis summary and interpretation of drug safety data The only guide of its kind *Drug Safety Data How to Analyze Summarize and Interpret to Determine Risk* is an invaluable

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- Chapter 1: Introduction to New Drugs Annual Update
- Chapter 2: Essential Elements of New Drugs Annual Update
- Chapter 3: New Drugs Annual Update in Everyday Life
- Chapter 4: New Drugs Annual Update in Specific Contexts
- Chapter 5: Conclusion

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6. In chapter 5, the author will draw a conclusion about New Drugs Annual Update. The final chapter will summarize the key points that have been discussed throughout the book.

The book is crafted in an easy-to-understand language and is complemented by engaging illustrations. This book is highly recommended for anyone seeking to gain a comprehensive understanding of New Drugs Annual Update.

https://pinsupreme.com/results/Resources/index.jsp/Mothertalk_Life_Stories_Of_Mary_Kiyoshi_Kiyooka.pdf

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