

Annual Report 2022

Reporting Desk for New Drugs

What is the Reporting Desk for New Drugs?

The Reporting Desk for New Drugs is a 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 were recently detected 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 (DIMS) 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 (CAM²) to assess the distribution and possible risks associated with the sale, transport, and use of new psychoactive substances in the Netherlands.



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 or have been re-introduced after being absent on the illicit drug market for several decades.

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
- Indole alkaloids (Tryptamines)
- Arylcyclohexylamines
- Synthetic opioids
- New benzodiazepines
- Other new psychoactive substances

Highlights 2022

In the Netherlands, a total of 122 different new psychoactive substances were reported in 2022 to the Reporting Desk for New Drugs by the participating laboratories; 14 of these new psychoactive substances were detected on the illicit drug market in the Netherlands for the first time. Seven of these substances were reported by Netherlands Forensic Institute, two by both the Netherlands Forensic Institute and DIMS, two by DIMS only and three by the Customs Laboratory.

- Netherlands is an important transit country for various New Psychoactive Substances (NPS). A small portion of them is intended for the Dutch consumer market. A notable example of this are synthetic cannabinoids, which are typically primarily reported by the Customs Laboratory of the Netherlands. However, in 2022, they only observed that large quantities of JWH-210 were imported.
- Regarding cathinones, it was particularly noteworthy that the Customs Laboratory of the Netherlands detected record amounts of the substance 3-CMC. This substance was not frequently submitted to DIMS but was found in the majority of samples submitted as 3-MMC.
- What's also striking is that, in the Netherlands in 2022, there is minimal reporting of the presence of synthetic opioids, unlike other European countries. O-desmethyiltramadol is the substance most commonly reported by some of the participating partners.

New Drugs Annual Update

Bruce H. Woolley, Edward S. Brady



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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.), 1985 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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Table of Contents New Drugs Annual Update

1. Understanding the eBook New Drugs Annual Update
 - The Rise of Digital Reading New Drugs Annual Update
 - Advantages of eBooks Over Traditional Books
2. Identifying New Drugs Annual Update
 - 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 New Drugs Annual Update
 - User-Friendly Interface
4. Exploring eBook Recommendations from New Drugs Annual Update
 - Personalized Recommendations
 - New Drugs Annual Update User Reviews and Ratings
 - New Drugs Annual Update and Bestseller Lists
5. Accessing New Drugs Annual Update Free and Paid eBooks
 - New Drugs Annual Update Public Domain eBooks
 - New Drugs Annual Update eBook Subscription Services

- New Drugs Annual Update Budget-Friendly Options
- 6. Navigating New Drugs Annual Update eBook Formats
 - ePub, PDF, MOBI, and More
 - New Drugs Annual Update Compatibility with Devices
 - New Drugs Annual Update Enhanced eBook Features
- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of New Drugs Annual Update
 - Highlighting and Note-Taking New Drugs Annual Update
 - Interactive Elements New Drugs Annual Update
- 8. Staying Engaged with New Drugs Annual Update
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers New Drugs Annual Update
- 9. Balancing eBooks and Physical Books New Drugs Annual Update
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection New Drugs Annual Update
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine New Drugs Annual Update
 - Setting Reading Goals New Drugs Annual Update
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of New Drugs Annual Update
 - Fact-Checking eBook Content of New Drugs Annual Update
 - 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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