reducing benzil using sodium borohydride

reducing benzil using sodium borohydride is a fundamental chemical reaction widely employed in organic synthesis for converting diketones into diols. This process involves the selective reduction of benzil, a 1,2-diketone, using sodium borohydride (NaBH4), a mild and versatile reducing agent. The reaction typically yields hydrobenzoin, a vicinal diol with important applications in pharmaceuticals and as a chiral building block. Understanding the mechanism, reaction conditions, and practical applications of this reduction is essential for chemists working in synthetic organic chemistry. This article will explore the chemistry behind reducing benzil using sodium borohydride, detail the reaction procedure, discuss factors influencing the reaction, and highlight its significance in laboratory and industrial settings.

- Chemistry of Benzil Reduction
- Role and Properties of Sodium Borohydride
- Reaction Mechanism of Benzil Reduction
- Experimental Procedure and Conditions
- Factors Affecting the Reduction Process
- Applications of Hydrobenzoin

Chemistry of Benzil Reduction

Benzil, chemically known as 1,2-diphenylethane-1,2-dione, is a diketone characterized by two adjacent carbonyl groups attached to phenyl rings. The reduction of benzil involves the conversion of these carbonyl groups into hydroxyl groups, resulting in the formation of hydrobenzoin. This reaction is a classic example of the selective reduction of diketones to vicinal diols, which can exist as different stereoisomers.

Structure and Properties of Benzil

Benzil is a crystalline solid with a distinctive yellow color, insoluble in water but soluble in organic solvents such as ethanol and acetone. Its two ketone groups are electrophilic centers susceptible to nucleophilic attack, which makes it an ideal substrate for reduction reactions. The proximity of

the two carbonyl groups affects the stereochemistry of the product formed during reduction.

Significance of Vicinal Diols

Vicinal diols like hydrobenzoin are valuable intermediates in organic synthesis. They serve as chiral auxiliaries, ligands in asymmetric synthesis, and precursors for various pharmaceuticals. The ability to selectively reduce benzil to hydrobenzoin allows for controlled synthesis of these important compounds.

Role and Properties of Sodium Borohydride

Sodium borohydride (NaBH4) is a widely used reducing agent in organic chemistry due to its mildness, selectivity, and ease of handling. It effectively reduces ketones and aldehydes to their corresponding alcohols without affecting many other functional groups.

Chemical Nature of Sodium Borohydride

NaBH4 is an inorganic compound composed of sodium cations and borohydride anions. It acts as a hydride donor, transferring hydride ions (H—) to electrophilic centers such as carbonyl carbons. It is stable in basic and neutral aqueous solutions but reacts violently with water under acidic conditions.

Advantages of Using NaBH4 for Benzil Reduction

Compared to other reducing agents like lithium aluminum hydride (LiAlH4), sodium borohydride offers several benefits:

- Selective reduction of ketones without affecting esters or carboxylic acids.
- Safer and easier to handle due to its stability in protic solvents such as ethanol or water.
- Operates under mild reaction conditions, reducing the risk of side reactions.

Reaction Mechanism of Benzil Reduction

The reduction of benzil by sodium borohydride proceeds through a nucleophilic addition mechanism, where hydride ions attack the electrophilic carbonyl carbons, followed by protonation to yield the diol.

Stepwise Mechanism

The mechanism involves several key steps:

- 1. **Hydride Transfer:** The borohydride ion donates a hydride ion to one of the carbonyl carbons in benzil, forming an alkoxide intermediate.
- 2. **Protonation:** The alkoxide intermediate is protonated by solvent molecules (usually water or alcohol), forming a hydroxy group.
- 3. **Second Hydride Attack:** A second hydride ion attacks the adjacent carbonyl group, repeating the process and leading to the formation of a vicinal diol.
- 4. **Final Protonation:** The remaining alkoxide is protonated, completing the reduction to hydrobenzoin.

Stereochemical Outcomes

The reduction can produce different stereoisomers of hydrobenzoin, including meso and racemic forms. Reaction conditions and solvents influence the stereoselectivity and yield of these isomers.

Experimental Procedure and Conditions

Conducting the reduction of benzil using sodium borohydride requires careful control of reaction parameters to ensure high yield and purity of hydrobenzoin.

Typical Reaction Setup

The procedure generally involves dissolving benzil in an appropriate solvent, usually ethanol or methanol, followed by the gradual addition of sodium borohydride while stirring. The reaction is often performed at room temperature or under mild heating to accelerate the process.

Step-by-Step Protocol

- Dissolve a measured amount of benzil in ethanol under stirring.
- Slowly add sodium borohydride powder or solution to the benzil solution.
- Maintain stirring and monitor the reaction progress using thin-layer chromatography (TLC) or other analytical methods.
- After completion, quench the reaction by adding water or dilute acid to deactivate excess NaBH4.
- Isolate the hydrobenzoin product by filtration or extraction, followed by purification through recrystallization or chromatography.

Factors Affecting the Reduction Process

Several variables influence the efficiency and outcome of reducing benzil using sodium borohydride. Understanding these factors is critical for optimizing the reaction.

Solvent Effects

The choice of solvent plays a vital role in solubility, reaction rate, and selectivity. Protic solvents such as ethanol or methanol facilitate protonation steps and stabilize intermediates, enhancing reaction speed and yield.

Temperature

Elevated temperatures can accelerate the reduction but may also increase side reactions or decrease selectivity. Mild heating is often sufficient, while excessively high temperatures are typically avoided.

Stoichiometry of Reagents

Using stoichiometric or slight excess amounts of sodium borohydride ensures complete reduction of benzil. Insufficient hydride can lead to incomplete conversion, while excessive amounts may complicate purification.

Reaction Time

The duration of the reaction affects product formation. Prolonged reaction times may improve yield but also risk over-reduction or byproduct formation.

Purity of Starting Materials

Impurities in benzil or sodium borohydride can hinder the reaction or affect product quality. Using high-purity reagents is recommended for optimal results.

Applications of Hydrobenzoin

Hydrobenzoin, the product of benzil reduction, is a valuable compound with diverse applications in organic synthesis and industry.

Use in Chiral Synthesis

Hydrobenzoin is widely used as a chiral auxiliary and ligand in asymmetric synthesis. Its stereocenters enable enantioselective reactions, producing optically active compounds essential in pharmaceuticals.

Pharmaceutical Intermediates

The vicinal diol functionality of hydrobenzoin facilitates the synthesis of various drugs and biologically active molecules. It serves as a precursor for compounds with antifungal, antibacterial, and anticancer properties.

Material Science and Catalysis

Hydrobenzoin derivatives find applications in the preparation of catalysts and materials with specialized properties. Its chemical versatility allows modification to tailor specific functionalities.

Frequently Asked Questions

What is the product of reducing benzil using sodium borohydride?

The reduction of benzil with sodium borohydride typically yields benzoin, which is an α -hydroxy ketone formed by the selective reduction of one of the

Why is sodium borohydride used for the reduction of benzil?

Sodium borohydride is used because it is a mild and selective reducing agent that can reduce ketones to alcohols without affecting other sensitive functional groups, making it suitable for selectively reducing benzil to benzoin.

What is the mechanism of benzil reduction by sodium borohydride?

Sodium borohydride delivers hydride ions (H-) to one of the carbonyl carbons in benzil, converting it to an alcohol group, resulting in benzoin. The reaction proceeds via nucleophilic attack of hydride on the electrophilic carbonyl carbon.

Can sodium borohydride reduce both carbonyl groups in benzil to diols?

Under typical conditions, sodium borohydride selectively reduces one carbonyl group in benzil to produce benzoin. However, with excess reagent and prolonged reaction time, both carbonyl groups can be reduced to give hydrobenzoin, a diol.

What solvents are commonly used for the reduction of benzil with sodium borohydride?

Common solvents include methanol, ethanol, or other protic solvents because they dissolve sodium borohydride and facilitate the hydride transfer to benzil.

How does the reaction temperature affect the reduction of benzil by sodium borohydride?

Lower temperatures help control the selectivity and rate of reduction, typically favoring the formation of benzoin. Higher temperatures can increase reaction rates but may lead to over-reduction to the diol.

Is the reduction of benzil by sodium borohydride stereoselective?

The reduction can produce a mixture of stereoisomers of benzoin or hydrobenzoin, as the hydride can attack from either face of the planar carbonyl group, leading to racemic mixtures.

What safety precautions should be taken when using sodium borohydride for benzil reduction?

Sodium borohydride should be handled with care as it reacts violently with water and acids, releasing hydrogen gas. Use appropriate protective equipment, work in a well-ventilated area, and avoid contact with moisture.

How can the progress of benzil reduction by sodium borohydride be monitored?

The reaction progress can be monitored by thin-layer chromatography (TLC), observing the disappearance of benzil and appearance of benzoin, or by spectroscopic methods such as NMR and IR spectroscopy.

What are common side reactions during the reduction of benzil with sodium borohydride?

Side reactions may include over-reduction to hydrobenzoin, decomposition of sodium borohydride in protic solvents, or formation of byproducts due to impurities or improper reaction conditions.

Additional Resources

1. Selective Reduction of Benzil with Sodium Borohydride: Mechanisms and Applications

This book offers an in-depth exploration of the chemical processes involved in the selective reduction of benzil using sodium borohydride. It covers the reaction mechanisms, conditions affecting selectivity, and practical applications in organic synthesis. Researchers and students will find detailed experimental procedures and case studies that highlight the nuances of this reduction.

- 2. Organic Synthesis Techniques: Sodium Borohydride Reductions of α -Diketones Focusing on the broader category of α -diketones, this book details the use of sodium borohydride as a reducing agent, with benzil reduction as a key example. It discusses reaction kinetics, solvent effects, and product characterization. The text is ideal for chemists looking to optimize reduction protocols in laboratory settings.
- 3. Practical Guide to Sodium Borohydride in Laboratory Reductions
 This practical manual emphasizes the safe and efficient use of sodium
 borohydride in various reductions, including benzil to hydrobenzoin. It
 provides step-by-step instructions, safety precautions, and troubleshooting
 tips. The book is particularly useful for undergraduate and graduate students
 performing organic reductions.
- 4. Advanced Organic Chemistry: Reductions of Carbonyl Compounds
 A comprehensive textbook that covers the reduction of diverse carbonyl

compounds, with a dedicated chapter on benzil reduction using sodium borohydride. The book explains stereochemical outcomes and compares different reducing agents. It is a valuable resource for advanced students and professional chemists.

5. Green Chemistry Approaches in Carbonyl Reductions: Sodium Borohydride Case Studies

This book explores environmentally friendly methods for reducing carbonyl compounds, highlighting sodium borohydride's role in sustainable chemistry. It includes case studies on benzil reduction, emphasizing solvent choice and waste minimization. Ideal for researchers interested in green synthetic methodologies.

- 6. Mechanistic Insights into Sodium Borohydride Reductions of α -Diketones Delving into the reaction pathways, this text offers detailed mechanistic analyses of sodium borohydride reducing benzil and related compounds. It integrates spectroscopic data and computational studies to explain product formation. The book is suited for chemists focused on reaction mechanism research.
- 7. Laboratory Manual: Reduction of Benzil to Hydrobenzoin
 Designed as a hands-on guide, this manual provides detailed experimental
 protocols for reducing benzil to hydrobenzoin using sodium borohydride. It
 includes data sheets, expected results, and tips for optimizing yield and
 purity. Perfect for teaching laboratories and practical courses in organic
 chemistry.
- 8. Sodium Borohydride in Industrial Organic Synthesis
 This book addresses the scale-up and industrial application of sodium
 borohydride reductions, featuring benzil as a model substrate. Topics include
 process optimization, cost analysis, and safety considerations. Industrial
 chemists and process engineers will find valuable insights into large-scale
 operations.
- 9. Chiral Diol Synthesis via Sodium Borohydride Reduction of Benzil Focusing on the synthesis of chiral diols, this book discusses the stereoselective reduction of benzil using sodium borohydride and related catalysts. It covers asymmetric synthesis techniques and applications in pharmaceuticals. Researchers involved in chiral molecule synthesis will benefit from its detailed methodologies.

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Reducing Benzil Using Sodium Borohydride

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Ebook Chapter Outline:

Introduction: The significance of reduction reactions in organic chemistry, introducing benzil and its reduction products. Brief overview of sodium borohydride as a reducing agent.

Mechanism of Benzil Reduction: A detailed step-by-step explanation of the reaction mechanism, including relevant electron movements and intermediate formation. Discussion of stereochemistry. Experimental Procedure: A comprehensive guide to performing the reduction, including detailed instructions on reagent preparation, reaction setup, workup, and purification techniques (recrystallization). Safety precautions are emphasized.

Characterization of the Product: Techniques for confirming the identity and purity of the reduced product (benzoin). Discussion of melting point determination, IR spectroscopy, and NMR spectroscopy.

Yield Calculation and Analysis: Explanation of yield calculation, potential sources of error, and methods for improving the reaction yield.

Applications of Benzil Reduction: Exploring the applications of benzoin and its derivatives in various fields, including organic synthesis and material science.

Conclusion: Summary of the key findings, highlighting the importance of the reaction and its practical applications.

Reducing Benzil Using Sodium Borohydride: A Comprehensive Guide

Introduction: The Power of Reduction in Organic Synthesis

Reduction reactions are fundamental transformations in organic chemistry, enabling the conversion of functional groups with higher oxidation states to those with lower oxidation states. These reactions are crucial for the synthesis of a vast array of organic molecules, including pharmaceuticals, polymers, and natural products. One such reaction involves the reduction of benzil, a diketone, to benzoin, an α -hydroxyketone. This transformation is a classic example of a nucleophilic addition reaction, frequently used in undergraduate organic chemistry laboratories to illustrate the reducing power of sodium borohydride (NaBH4). Benzil's reduction to benzoin is particularly significant because benzoin itself serves as a valuable intermediate in the synthesis of many other compounds. Understanding this reaction offers a practical understanding of reaction mechanisms, purification techniques, and spectroscopic characterization methods vital to organic chemistry.

Mechanism of Benzil Reduction: A Step-by-Step Analysis

Sodium borohydride ($NaBH_4$) is a mild reducing agent commonly used to reduce ketones and aldehydes. Its selectivity makes it a valuable tool in organic synthesis, as it avoids reducing other functional groups that might be present in a molecule. The reduction of benzil to benzoin proceeds via a nucleophilic addition mechanism.

- 1. Hydride Attack: The hydride ion (H^-) , a powerful nucleophile, from NaBH4 attacks the electrophilic carbonyl carbon of benzil. This attack leads to the formation of an alkoxide intermediate. The reaction is often carried out in a protic solvent like methanol or ethanol, which helps to stabilize the alkoxide.
- 2. Protonation: The alkoxide intermediate is then protonated by the solvent (methanol or ethanol) or water, resulting in the formation of a hydroxyl group (-OH).
- 3. Second Hydride Attack (Less Common): While a single hydride attack from NaBH4 is sufficient for the majority of benzil reduction, under certain conditions a second hydride attack can occur on the other carbonyl group, followed by a second protonation step leading to a diol. However, this side reaction is usually minimized by controlling the stoichiometry and reaction conditions.
- 4. Stereochemistry: The reduction of benzil can produce both diastereomers (three and erythro benzoin), depending on the reaction conditions. However, the stereochemical outcome is often not highly controlled using sodium borohydride reduction alone.

Experimental Procedure: A Detailed Guide to Benzil Reduction

Materials:

Benzil (precisely weighed amount)
Sodium borohydride (NaBH4)
Methanol (or ethanol)
Ice bath
Erlenmeyer flask
Filter paper
Buchner funnel
Recrystallization solvent (e.g., ethanol/water)

Procedure:

- 1. Dissolution: Dissolve the weighed benzil in a suitable amount of methanol in an Erlenmeyer flask. An ice bath helps to control the exothermic nature of the reaction.
- 2. Addition of NaBH₄: Slowly add the NaBH₄ to the benzil solution, ensuring that the temperature remains below 25°C. Stir the mixture continuously. The reaction is exothermic and may cause foaming.
- 3. Reaction Completion: Allow the reaction to proceed for a specific time (generally 30-60 minutes)

with constant stirring. Monitor the reaction using TLC (thin-layer chromatography) if possible.

- 4. Work-up: Carefully add dilute hydrochloric acid or acetic acid to quench any excess NaBH₄. The solution will become slightly acidic.
- 5. Filtration: Filter the resulting mixture using vacuum filtration to remove any insoluble impurities.
- 6. Recrystallization: Recrystallize the crude benzoin from a suitable solvent mixture (e.g., ethanol/water) to obtain pure crystals.
- 7. Drying: Dry the purified crystals and weigh them to determine the yield.

Safety Precautions: Sodium borohydride reacts vigorously with water, so careful addition and quenching are crucial. Always wear appropriate safety glasses and gloves. Dispose of waste materials properly according to the safety guidelines.

Characterization of the Product: Confirming Benzoin Identity and Purity

The identity and purity of the synthesized benzoin can be confirmed through various techniques:

Melting Point Determination: The melting point of the recrystallized benzoin can be compared to the literature value (137°C) to assess its purity.

Infrared (IR) Spectroscopy: IR spectroscopy provides information about the functional groups present in the molecule. The presence of a broad hydroxyl (O-H) stretch and a carbonyl (C=O) stretch confirms the formation of benzoin.

Nuclear Magnetic Resonance (NMR) Spectroscopy: ¹H NMR and ¹³C NMR spectroscopy provide detailed information about the structure of the molecule. The chemical shifts and coupling patterns in the NMR spectra help confirm the identity and purity of benzoin.

Yield Calculation and Analysis: Optimizing the Reaction

The percentage yield of the reaction is calculated using the following formula:

Percentage Yield = (Actual Yield / Theoretical Yield) x 100%

The actual yield is the weight of the purified benzoin obtained, while the theoretical yield is calculated based on the stoichiometry of the reaction and the initial amount of benzil used. Low yields can result from incomplete reaction, loss of product during work-up, or inefficient purification. Careful attention to experimental details and optimization of reaction conditions (solvent, temperature, reaction time) are crucial for maximizing the yield.

Applications of Benzil Reduction: Beyond the Laboratory

Benzoin, the product of benzil reduction, has various applications in:

Organic Synthesis: Benzoin serves as a crucial building block in the synthesis of other organic compounds, including pharmaceuticals and natural products.

Material Science: Benzoin and its derivatives find use in the preparation of polymeric materials and advanced materials.

Conclusion: A Valuable Synthetic Transformation

The reduction of benzil using sodium borohydride is a simple yet powerful reaction with significant pedagogical and practical value. It allows for a detailed exploration of reaction mechanisms, purification techniques, and spectroscopic characterization methods in organic chemistry. Understanding this reaction provides a strong foundation for further exploration of reduction reactions and their importance in the synthesis of a wide range of organic compounds. The applications of benzoin, the product of this reaction, further highlight its importance in various scientific and industrial fields.

FAQs

- 1. What are the safety precautions when handling sodium borohydride? Sodium borohydride reacts exothermically with water and acids. Always wear appropriate personal protective equipment (PPE), including gloves and safety glasses. Add the reagent slowly and carefully.
- 2. What is the role of the solvent in the benzil reduction? The solvent, usually methanol or ethanol, acts as both a solvent for the reactants and a proton source for the alkoxide intermediate.
- 3. Why is recrystallization necessary? Recrystallization is a purification technique used to remove impurities from the crude benzoin, resulting in a higher purity product with a sharper melting point.
- 4. What spectroscopic techniques are used to characterize the product? IR, ¹H NMR, and ¹³C NMR spectroscopy are commonly used to confirm the structure and purity of the benzoin.
- 5. What factors can affect the yield of the reaction? Incomplete reaction, loss of product during work-up, and inefficient purification can all affect the yield.
- 6. Can other reducing agents be used to reduce benzil? Yes, other reducing agents, such as lithium aluminum hydride (LiAlH₄), can reduce benzil. However, LiAlH₄ is a more powerful reducing agent and may reduce other functional groups as well.

- 7. What are the differences between threo and erythro benzoin? Threo and erythro benzoin are diastereomers, differing in the stereochemistry of the hydroxyl and ketone groups.
- 8. How can I monitor the progress of the reaction? Thin-layer chromatography (TLC) can be used to monitor the reaction progress and determine when the reaction is complete.
- 9. What are some common errors to avoid during the experiment? Common errors include adding NaBH₄ too quickly, insufficient stirring, and using impure starting materials.

Related Articles:

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- 2. Stereochemistry of Reduction Reactions: A detailed explanation of stereochemical aspects of reduction reactions and how to control stereoselectivity.
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applications of nanomaterials from renewable resources; hydrogels and its nanocomposites from renewable resources: preparation of chitin-based nanocomposite materials through gelation with ionic liquid; starch-based bionanocomposites; biorenewable nanofiber and nanocrystal; investigation of wear characteristics of dental composite reinforced with rice husk-derived nanosilica filler particles; performance of regenerated cellulose/vermiculite nanocomposites fabricated via ionic liquid; preparation, structure, properties, and interactions of the PVA/cellulose composites; green composites with cellulose nanoreinforcements; biomass composites from bamboo-based micro/nanofibers; synthesis and medicinal properties of polycarbonates and resins from renewable sources; nanostructured polymer composites with modified carbon nanotubes; organic-inorganic nanocomposites derived from polysaccharides; natural polymer-based nanocomposites; cellulose whisker-based green polymer composites; poly (lactic acid) nanocomposites reinforced with different additives; nanocrystalline cellulose; halloysite-based bionanocomposites; nanostructurated composites based on biodegradable polymers and silver nanoparticles; starch-based biomaterials and nanocomposites; green nanocomposites based on PLA and natural organic fillers; and chitin and chitosan-based nanocomposites.

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drugs, this book fills the gap that exists in the literature of drug synthesis. It provides the kind of information that will be of interest to those who work, or plan to begin work, in the areas of biologically active compounds and the synthesis of medicinal drugs. This book presents the synthesis of various groups of drugs in an order similar to that traditionally presented in a pharmacology curriculum. This was done with a very specific goal in mind - to harmonize the chemical aspects with the pharmacology curriculum in a manner useful to chemists. Practically every chapter begins with an accepted brief definition and description of a particular group of drugs, proposes their classification, and briefly explains the present model of their action. This is followed by a detailed discussion of methods for their synthesis. Of the thousands of drugs existing on the pharmaceutical market, the book mainly covers generic drugs that are included in the WHO's Essential List of Drugs. For practically all of the 700+ drugs described in the book, references (around 2350) to the methods of their synthesis are given along with the most widespread synonyms. Synthesis of Essential Drugs is an excellent handbook for chemists, biochemists, medicinal chemists, pharmacists, pharmacologists, scientists, professionals, students, university libraries, researchers, medical doctors and students, and professionals working in medicinal chemistry. * Provides a brief description of methods of synthesis, activity and implementation of all drug types* Includes synonyms* Includes over 2300 references

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Stéphane Caron, 2020-02-05 This book is a hands-on guide for the organic chemist. Focusing on the most reliable and useful reactions, the chapter authors provide the information necessary for a chemist to strategically plan a synthesis, as well as repeat the procedures in the laboratory.

Consolidates all the key advances/concepts in one book, covering the most important reactions in organic chemistry, including substitutions, additions, eliminations, rearrangements, oxidations, reductions Highlights the most important reactions, addressing basic principles, advantages/disadvantages of the methodology, mechanism, and techniques for achieving laboratory success Features new content on recent advances in CH activation, photoredox and electrochemistry, continuous chemistry, and application of biocatalysis in synthesis Revamps chapters to include new and additional examples of chemistry that have been demonstrated at a practical scale

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thorough overview of the synthetic routes used to access specific classesof therapeutic agents, this volume covers approximately 220 new non-proprietary drug entities introduced since the publication of Volume 6. Many of these compounds represent novel structural types firstidentified by sophisticated new cell-based assays. Specifically, a significant number of new antineoplastic and antiviral agents are covered. As in the previous volumes, materials are organized by chemical class and syntheses originate with available starting materials. Organized to make the information accessible, this resource covers disease state, rationale for method of drug therapy, and the biological activities of each compound and preparation. The Organic Chemistry of Drug Synthesis, Volume 7 is a hands-on reference for medicinal and organic chemists, and a great resource for graduate and advanced undergraduate students in organic and medicinal chemistry.

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