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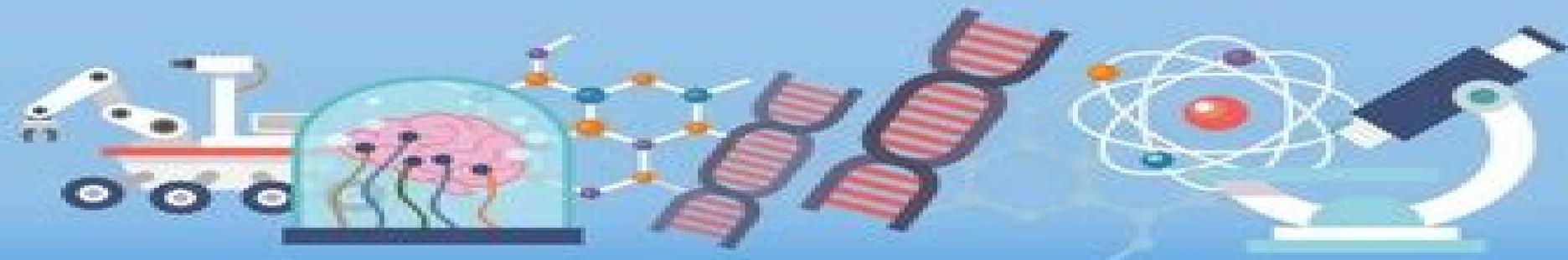
Project Details

Proposed Title: Creation and synthesis of bifunctional degrader libraries for the discovery of possible cancer targets.

Background of the Project

The success of the target discovery of cancer drug is highly dependent on the determination of protein targets which can be modulated using small molecules, thus producing anti-cancer effects. The purpose of this project is to possibly apply the use of synthetic organic chemistry, combined with medicinal chemistry to the creation and synthesis of a library of different compounds which can promote protein degradation. After, these will be used in identifying protein targets which will then be used in the discovery of new cancer drugs.

Bifunctional degraders are a newly discovered molecule class wherein an E3 ubiquitin ligase binding compound is connected to a warhead which binds its therapeutic target. In contrast with other conventional smaller molecule drugs which bind and blocks the action of their protein targets, these bifunctional degraders work by catalyzing the formulation of a complex in between an E3 ligase, as well as a target protein, thus allowing for ubiquitination, thus resulting to the irreversible and rapid degradation of the potential target.



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Table of Contents Cancer Research Proposal Sample

1. Understanding the eBook Cancer Research Proposal Sample
 - The Rise of Digital Reading Cancer Research Proposal Sample
 - Advantages of eBooks Over Traditional Books
2. Identifying Cancer Research Proposal Sample
 - 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 Cancer Research Proposal Sample
 - User-Friendly Interface
4. Exploring eBook Recommendations from Cancer Research Proposal Sample
 - Personalized Recommendations
 - Cancer Research Proposal Sample User Reviews and Ratings
 - Cancer Research Proposal Sample and Bestseller Lists
5. Accessing Cancer Research Proposal Sample Free and Paid eBooks
 - Cancer Research Proposal Sample Public Domain eBooks
 - Cancer Research Proposal Sample eBook Subscription Services
 - Cancer Research Proposal Sample Budget-Friendly Options
6. Navigating Cancer Research Proposal Sample eBook Formats

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