

To

**The Principal**

St. Wilfreds Institute Of Pharmacy

Panvel

Dear Sir/Madam,

Subject: Intimation of research Project approval

We are pleased to inform you that the project proposal submitted by **Dr. Akshay Meshram**, Assistant Professor, Department of Pharmacology, titled "**Assessment Of Medicinal Plants' Anxiolytic, Antidepressant, Anticonvulsant, And Nootropic Properties**" has been approved and sanctioned to be proceeded and completed on the following lines:

S.NO	Particulars	Amount
1	Total Cost of the Proposed Project	Rs 6,00,000
2	Amount Sanction for the Project Completion	Rs 5,00,000
3	Amount Sanction after progress report submission	Rs 4,00,000
4	Principal Investigator	<b>Dr. Akshay Meshram</b> Assistant Professor Department of Pharmacology
5	Duration of the Project	2 Years

- The Principal investor should complete the project in the stipulated duration.
- The Project report with analysis should be submitted to the company with appropriate signatures from the university authorities.
- The Project sanctioned amount will be deposited to the university bank account only.
- The Principal investigator/any Project Assistant are not eligible for any additional Claims.
- A final copy should be hardbound and submitted with all necessary acknowledgements from the concerned authorities of the university

Authorized by:

Thanking You

  
Sanctioned Officer/seal

To

**The Principal**

St. Wilfreds Institute Of Pharmacy

Panvel

Dear Sir/Madam,

Subject: Intimation of research Project approval

We are pleased to inform you that the project proposal submitted by **Dr. Vanita Kanase**, Professor, Department of Pharmacology, titled “**Assessment of the Immunomodulatory Effects of a Novel Medicinal Herbs**” has been approved and sanctioned to be Proceeded and completed on the following lines:

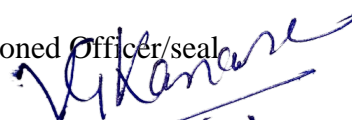
S.NO	Particulars	Amount
1	Total Cost of the Proposed Project	Rs 7,00,000
2	Amount Sanction for the Project Completion	Rs 6,00,000
3	Amount Sanction after progress report submission	Rs 5,00,000
4	Principal Investigator	<b>Dr. Vanita Kanase</b> Professor Department of Pharmacology
5	Duration of the Project	2 Years

- The Principal investor should complete the project in the stipulated duration.
- The Project report with analysis should be submitted to the company with appropriate signatures from the university authorities.
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To

**The Principal**

St. Wilfreds Institute Of Pharmacy

Panvel

Dear Sir/Madam,

Subject: Intimation of research Project approval

We are pleased to inform you that the project proposal submitted by **Dr. Deenanath Jhade**, Professor, Department of Pharmacognosy, titled **“Using in vitro in vivo and metabolomics approaches, a phytopharmaceutical and synergy based combination of Boerhavia diffusa L and tinospora cordifolia L is being developed for chronic kidney diseases”** has been approved and sanctioned to be Proceeded and completed on the following lines:

S.NO	Particulars	Amount
1	Total Cost of the Proposed Project	Rs 4,50,000
2	Amount Sanction for the Project Completion	Rs 3,00,000
3	Amount Sanction after progress report submission	Rs 2,50,000
4	Principal Investigator	<b>Dr. Deenanath Jhade</b> Professor Department of Pharmacognosy
5	Duration of the Project	2 Years

- The Principal investor should complete the project in the stipulated duration.
- The Project report with analysis should be submitted to the company with appropriate signatures from the university authorities.
- The Project sanctioned amount will be deposited to the university bank account only.
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Dear Sir/Madam,

Subject: Intimation of research Project approval

We are pleased to inform you that the project proposal submitted by **Dr. Sanchita Mandal**, Professor, Department of Pharmaceutics, titled **“Formulation and Evaluation of Novel Ophthalmic Drug Delivery System”** has been approved and sanctioned to be Proceeded and completed on the following lines:

S.NO	Particulars	Amount
1	Total Cost of the Proposed Project	Rs 3,00,000
2	Amount Sanction for the Project Completion	Rs 2,00,000
3	Amount Sanction after progress report submission	Rs 1,00,000
4	Principal Investigator	<b>Dr. Sanchita Mandal</b> Professor Department of Pharmaceutics
5	Duration of the Project	2 Years

- The Principal investor should complete the project in the stipulated duration.
- The Project report with analysis should be submitted to the company with appropriate signatures from the university authorities.
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Panvel

Dear Sir/Madam,

Subject: Intimation of research Project approval

We are pleased to inform you that the project proposal submitted by **Dr. Sanchita Mandal**, Professor, Department of Pharmaceutics, titled **“Formulation and Development of Effervescent system for Antihistamine Drugs”** has been approved and sanctioned to be Proceeded and completed on the following lines:

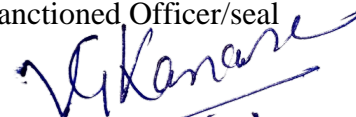
S.NO	Particulars	Amount
1	Total Cost of the Proposed Project	Rs 2,00,000
2	Amount Sanction for the Project Completion	Rs 1,00,000
3	Amount Sanction after progress report submission	Rs 1,00,000
4	Principal Investigator	<b>Dr. Sanchita Mandal</b> Professor Department of Pharmaceutics
5	Duration of the Project	2 Years

- The Principal investor should complete the project in the stipulated duration.
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To

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Dear Sir/Madam,

Subject: Intimation of research Project approval

We are pleased to inform you that the project proposal submitted by **Dr. Deenanath Jhade**, Professor, Department of Pharmacognosy, titled **“Development of a topical herbal formulation for wound healing from selected drug of Centella Asiatica”** has been approved and sanctioned to be Proceeded and completed on the following lines:

S.NO	Particulars	Amount
1	Total Cost of the Proposed Project	Rs 3,00,000
2	Amount Sanction for the Project Completion	Rs 2,00,000
3	Amount Sanction after progress report submission	Rs 1,00,000
4	Principal Investigator	<b>Dr. Deenanath Jhade</b> Professor Department of Pharmacognosy
5	Duration of the Project	2 Years

- The Principal investor should complete the project in the stipulated duration.
- The Project report with analysis should be submitted to the company with appropriate signatures from the university authorities.
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Near Shedung Toll Plaza, Old Mumbai Pune Highway, Panvel, Navi Mumbai.

Call : +91 897611122/33/44/66 E-mail : info@csmu.in | Website

**PRINCIPAL**  
ST. WILFRED'S INSTITUTE OF PHARMACY  
OLD MUMBAI PUNE HIGHWAY, NEAR SHEDUNG TOLL PLAZA,  
SHEDUNG, PANVEL- 410 206.

To

**The Principal**

St. Wilfreds Institute Of Pharmacy

Panvel

Dear Sir/Madam,

Subject: Intimation of research Project approval

We are pleased to inform you that the project proposal submitted by **Dr. Deenanath Jhade**, Professor, Department of Pharmacognosy, titled **“Assessment and Development of a herbal formulation for Acne treatment”** has been approved and sanctioned to be Proceeded and completed on the following lines:

S.NO	Particulars	Amount
1	Total Cost of the Proposed Project	Rs 2,50,000
2	Amount Sanction for the Project Completion	Rs 1,50,000
3	Amount Sanction after progress report submission	Rs 1,00,000
4	Principal Investigator	<b>Dr. Deenanath Jhade</b> Professor Department of Pharmacognosy
5	Duration of the Project	2 Years

- The Principal investor should complete the project in the stipulated duration.
- The Project report with analysis should be submitted to the company with appropriate signatures from the university authorities.
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# IDEAL COLLEGE OF PHARMACY

Affiliated to Mumbai University and Governed by Ideal Foundation

Pimplaj Road, Village Posheri, Taluka Wada, District Palghar- Maharashtra

Date: 10/07/20

To

## The Principal

St. Wilfreds Institute Of Pharmacy

Panvel

Dear Sir/Madam,

Subject: Intimation of research Project approval

We are pleased to inform you that the project proposal submitted by **Dr. Deenanath Jhade**, Assistant Professor, Department of Pharmacognosy, titled "**Comparative Pharmacognostical Photochemical in vitro Screening of Antidiabetic activities of Monochoria species**" has been approved and sanctioned to be proceeded and completed on the following lines:

S.NO	Particulars	Amount
1	Total Cost of the Proposed Project	Rs 2,00,000
2	Amount Sanction for the Project Completion	Rs 1,00,000
3	Amount Sanction after progress report submission	Rs 1,00,000
4	Principal Investigator	<b>Dr. Deenanath Jhade</b> Professor Department of Pharmacognosy
5	Duration of the Project	2 Years

- The Principal investor should complete the project in the stipulated duration.
- The Project report with analysis should be submitted to the company with appropriate signatures from the university authorities.
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# IDEAL COLLEGE OF PHARMACY

Affiliated to Mumbai University and Governed by Ideal Foundation  
Pimplaj Road, Village Posheri, Taluka Wada, District Palghar- Maharashtra

Date: 24/09/20

To

## The Principal

St.Wilfreds Institute Of Pharmacy

Panvel

Dear Sir/Madam,

Subject: Intimation of research Project approval

We are pleased to inform you that the project proposal submitted by **Dr. Sanchita Mandal**, Assistant Professor, Department of Pharmaceutics, titled "**Formulation and Evaluation of topical liposomal gel for management of acne**" has been approved and sanctioned to be proceeded and completed on the following lines:

S.NO	Particulars	Amount
1	Total Cost of the Proposed Project	Rs 2,00,000
2	Amount Sanction for the Project Completion	Rs 1,00,000
3	Amount Sanction after progress report submission	Rs 1,00,000
4	Principal Investigator	<b>Dr. Sanchita Mandal,</b> Professor Department of Pharmaceutics
5	Duration of the Project	2 Years

- The Principal investor should complete the project in the stipulated duration.
- The Project report with analysis should be submitted to the company with appropriate signatures from the university authorities.
- The Project sanctioned amount will be deposited to the university bank account only.
- The Principal investigator/any Project Assistant are not eligible for any additional Claims.
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# IDEAL COLLEGE OF PHARMACY

Affiliated to Mumbai University and Governed by Ideal Foundation

Pimplaj Road, Village Posheri, Taluka Wada, District Palghar- Maharashtra

Date: 18/07/19

To

## The Principal

St.Wilfreds Institute Of Pharmacy

Panvel

Dear Sir/Madam,

Subject: Intimation of research Project approval

We are pleased to inform you that the project proposal submitted by **Dr. Sanchita Mandal**, Professor, Department of Pharmaceutics, titled "**Formulation and Evaluation of Novel topical formulation based on surfactants for Antifungal treatment**" has been approved and sanctioned to be proceeded and completed on the following lines:

S.NO	Particulars	Amount
1	Total Cost of the Proposed Project	Rs 3,00,000
2	Amount Sanction for the Project Completion	Rs 2,00,000
3	Amount Sanction after progress report submission	Rs 1,00,000
4	Principal Investigator	<b>Dr. Sanchita Mandal</b> , Professor Department of Pharmaceutics
5	Duration of the Project	2 Years

- The Principal investor should complete the project in the stipulated duration.
- The Project report with analysis should be submitted to the company with appropriate signatures from the university authorities.
- The Project sanctioned amount will be deposited to the university bank account only.
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# IDEAL COLLEGE OF PHARMACY

Affiliated to Mumbai University and Governed by Ideal Foundation

Pimplaj Road, Village Posheri, Taluka Wada, District Palghar- Maharashtra

Date: 25/06/19

To

**The Principal**

St.Wilfreds Institute Of Pharmacy

Panvel

Dear Sir/Madam,

Subject: Intimation of research Project approval

We are pleased to inform you that the project proposal submitted by **Dr. Deenanath Jhade**, Professor, Department of Pharmacognosy, titled "**A Phytochemical and Pharmacological Pharmacognostic analysis of Ampelocissus Latifoliaroxb** " has been approved and sanctioned to be proceeded and completed on the following lines:

S.NO	Particulars	Amount
1	Total Cost of the Proposed Project	Rs 2,50,000
2	Amount Sanction for the Project Completion	Rs 1,50,000
3	Amount Sanction after progress report submission	Rs 1,00,000
4	Principal Investigator	<b>Dr. Deenanath Jhade,</b> Professor Department of Pharmaceutics
5	Duration of the Project	2 Years

- The Principal investor should complete the project in the stipulated duration.
- The Project report with analysis should be submitted to the company with appropriate signatures from the university authorities.
- The Project sanctioned amount will be deposited to the university bank account only.
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Thanking You

  
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# IDEAL COLLEGE OF PHARMACY

Affiliated to Mumbai University and Governed by Ideal Foundation

Pimplaj Road, Village Posheri, Taluka Wada, District Palghar- Maharashtra

Date: 27/07/18

To

**The Principal**

St.Wilfreds Institute Of Pharmacy

Panvel

Dear Sir/Madam,

Subject: Intimation of research Project approval

We are pleased to inform you that the project proposal submitted by **Dr. Uma Patil**, Professor, Department of Pharmaceutics, titled "**Development and Evaluation of Nano-ocular formulation for glaucoma**" has been approved and sanctioned to be proceeded and completed on the following lines:

S.NO	Particulars	Amount
1	Total Cost of the Proposed Project	Rs 2,00,000
2	Amount Sanction for the Project Completion	Rs 1,00,000
3	Amount Sanction after progress report submission	Rs 1,00,000
4	Principal Investigator	<b>Dr. Uma Patil,</b> Professor Department of Pharmaceutics
5	Duration of the Project	2 Years

- The Principal investor should complete the project in the stipulated duration.
- The Project report with analysis should be submitted to the company with appropriate signatures from the university authorities.
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- The Principal investigator/any Project Assistant are not eligible for any additional Claims.
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Authorized by:

Thanking You

  
Sanctioned Officer/seal

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# ST. WILFRED'S INSTITUTE OF PHARMACY

Affiliated to Mumbai University, Approved by AICTE- New Delhi, DTE Maharashtra  
Near Shedung Toll Plaza, Old Mumbai Pune Highway, Shedung, Panvel-410206

## PROJECT REPORT



*[Handwritten Signature]*  
**PRINCIPAL**  
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SHEDUNG, PANVEL- 410 206.

*[Handwritten Signature: V. Kanase]*  
**PRINCIPAL**  
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OLD MUMBAI PUNE HIGHWAY, NEAR SHEDUNG TOLL PLAZA,  
SHEDUNG, PANVEL- 410 206.





# ST. WILFRED'S INSTITUTE OF PHARMACY

Affiliated to Mumbai University, Approved by AICTE- New Delhi, DTE Maharashtra  
Near Shedung Toll Plaza, Old Mumbai Pune Highway, Shedung, Panvel-410206

## **Project Title: A Phytochemical and Pharmacological Pharmacognostic Analysis of Ampelocissus Latifolia Roxb**

**Principal Investigator:** Dr. Deenanath Jhade, Professor, Department of Pharmacognosy

### **Funding Details:**

Total Cost of the Proposed Project: Rs. 2,50,000

Amount Sanctioned for Project Completion: Rs. 1,50,000

Amount Sanctioned after Progress Report Submission: Rs. 1,00,000

**Project Duration:** 2 Years

### **Objectives:**

Conduct a detailed phytochemical analysis of Ampelocissus Latifolia Roxb.

Evaluate the pharmacological potential of the plant, focusing on its anti-inflammatory, antioxidant, and antimicrobial activities.

Investigate the safety and tolerability of the plant extracts in animal models. Methodology:  
Collection and authentication of the plant material.

Extraction and fractionation of the plant material using various solvents.

Phytochemical screening and characterization of the plant extracts using chromatographic and spectroscopic techniques.

Evaluation of the anti-inflammatory, antioxidant, and antimicrobial activities of the plant extracts in vitro.

Safety and tolerability assessment of the plant extracts in animal models. Progress

Report:

Completed collection and authentication of the plant material. Extraction and

fractionation of the plant material are ongoing.

### **Next Steps:**

Complete extraction and fractionation of the plant material.

Perform phytochemical screening and characterization of the plant extracts.



*W. Kanase*  
**PRINCIPAL**  
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SHEDUNG, PANVEL- 410 206

*W. Kanase*  
**PRINCIPAL**  
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SHEDUNG, PANVEL- 410 206



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Affiliated to Mumbai University, Approved by AICTE- New Delhi, DTE Maharashtra  
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Conduct in vitro pharmacological activities assessment.

Evaluate the safety and tolerability of the plant extracts in animal models. Analyze the results and prepare the final project report.

Costing pattern:

Purchasing of medicinal Plants: 55000

Reagents and chemicals: 150000

Equipments : 100000

Documentation: 10000


Miscellaneous: 35000

## Conclusion:

The project is progressing as planned, with successful collection and authentication of the plant material. We expect to achieve the proposed objectives within the stipulated 2-year duration, leading to a comprehensive understanding of the phytochemical and pharmacological properties of *Ampelocissus Latifolia* Roxb.



  
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Near Shedung Toll Plaza, Old Mumbai Pune Highway, Shedung, Panvel-410206

## Project Title: Formulation and Development of Effervescent System for Antihistamine Drugs

**Principal Investigator:** Dr. Sanchita Mandal, Professor, Department of Pharmaceutics

### Funding Details:

Total Cost of the Proposed Project: Rs. 2,00,000

Amount Sanctioned for Project Completion: Rs. 1,00,000

Amount Sanctioned after Progress Report Submission: Rs. 1,00,000Project

Duration: 2 Years

### Objectives:

Develop an effervescent drug delivery system for antihistamine drugs to improve their bioavailability and patient compliance.

Evaluate the physicochemical properties, stability, and drug release profiles of the effervescent formulations.

Assess the in vivo performance and therapeutic efficacy of the effervescent antihistamine system in animal models.

Investigate the safety and tolerability of the effervescent system in animal models.

### Methodology:

Selection and optimization of effervescent system components, such as acids, bases, and excipients.

Formulation of effervescent antihistamine systems using optimized components.

Characterization of the formulations for drug content, effervescence time, pH, and stability.


In vitro drug release studies in simulated saliva and comparison with conventional oral dosage forms.

In vivo performance evaluation in animal models, including pharmacokinetic and pharmacodynamic studies.

Safety and tolerability assessment through acute and chronic toxicity studies in animal models.



  
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## Progress Report:

Optimized components for the effervescent drug delivery system.

Formulated effervescent antihistamine systems and characterized their physicochemical properties.

Completed in vitro drug release studies and compared the results with conventional dosage forms.

Costing pattern:

Purchasing of medicinal Plants: 55000

Reagents and chemicals: 150000

Equipments : 100000

Documentation: 10000

Miscellaneous: 35000

## Next Steps:

Perform in vivo studies in animal models to assess pharmacokinetics, pharmacodynamics, and therapeutic efficacy.

Conduct safety and tolerability studies in animal models to support the potential use of the effervescent system.


Finalize the formulation and prepare for scale-up and further preclinical and clinical evaluations.

## Conclusion:

The project has progressed well, with the successful development and initial characterization of an effervescent drug delivery system for antihistamine drugs. We anticipate completing the proposed objectives within the stipulated 2-year duration, leading to the development of an advanced oral formulation with improved bioavailability and patient compliance.



  
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Affiliated to Mumbai University, Approved by AICTE- New Delhi, DTE Maharashtra  
Near Shedung Toll Plaza, Old Mumbai Pune Highway, Shedung, Panvel-410206

## Project Title: Assessment and Development of a Herbal Formulation for Acne Treatment

**Principal Investigator:** Dr. Deenanath Jhade, Professor, Department of Pharmacognosy

### Funding Details:

Total Cost of the Proposed Project: Rs. 2,50,000

Amount Sanctioned for Project Completion: Rs. 1,50,000

Amount Sanctioned after Progress Report Submission: Rs. 1,00,000

Duration: 2 Years

### Objectives:

Assess the anti-acne potential of selected medicinal plants and their bioactive compounds.

Develop a herbal formulation for acne treatment using the most promising plant extracts.

Evaluate the physicochemical properties, stability, and drug release profiles of the formulation.

Assess the in vitro and in vivo anti-acne efficacy of the herbal formulation. Investigate the safety and tolerability of the herbal formulation in animal models.

### Methodology:

Screening of medicinal plants for their anti-acne potential using in vitro assays, such as antibacterial and anti-inflammatory activity.

Selection of the most promising plant extracts for formulation development. Formulation of a herbal gel or cream using the selected plant extracts.

Characterization of the formulation for drug content, consistency, pH, and stability. In vitro drug release studies using artificial membranes.

In vitro and in vivo anti-acne efficacy assessment using suitable models, such as antibacterial and anti-inflammatory activity studies, as well as animal models.



*Deenanath Jhade*  
**PRINCIPAL**  
ST WILFRED'S INSTITUTE OF PHARMACY  
OLD MUMBAI PUNE HIGHWAY, NEAR SHEDUNG TOLL PLAZA,  
SHEDUNG, PANVEL-410206

*V. Kanase*  
**PRINCIPAL**  
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SHEDUNG, PANVEL-410206





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Safety and tolerability studies, including skin irritation and histopathological examinations in animal models.

## Progress Report:

Completed screening and selection of medicinal plants with promising anti-acne potential.

Developed a herbal gel formulation using the selected plant extracts.

Ongoing characterization of the formulation for physicochemical properties and stability.

## Next Steps:

Complete characterization of the formulation. Conduct in vitro drug release studies.

Assess the in vitro and in vivo anti-acne efficacy of the herbal formulation. Perform safety and tolerability studies in animal models.


Finalize the formulation and prepare for scale-up and further preclinical and clinical evaluations.

## Conclusion:

The project is progressing well, with the successful screening of medicinal plants for anti-acne potential and the development of a herbal gel formulation. We anticipate completing the proposed objectives within the stipulated 2-year duration, leading to the development of an effective and safe herbal remedy for acne treatment.



  
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## **Project Title: Assessment of the Immunomodulatory Effects of Novel Medicinal Herbs**

Principal Investigator: Dr. Vanita Kanase, Professor, Department of Pharmacology

### **Funding Details:**

Total Cost of the Proposed Project:Rs.7,00,000

Amount Sanctioned for Project Completion:Rs.6,00,000

Amount Sanctioned after Progress Report Submission:Rs.5,00,000

Project Duration:2Years

### **Objectives:**

Evaluate the immunomodulatory effects of selected novel medicinal herbs.

Identify bioactive compounds responsible for these therapeutic effects.

Investigate the molecular mechanisms underlying the observed immunomodulatory activities.

Assess the potential for developing novel phyto pharmaceuticals with immunomodulatory properties.

### **Methodology:**

Selection of novel medicinal herbs based on traditional usage and existing scientific literature.

Extraction and characterization of bioactive compounds from these elected plants.

In vitro assessment of immunomodulatory activities using relevant cellular model and biochemical assays.


In vivo evaluation of the immunomodulatory effects of these medicinal herbs using animal models.

Meta bolomics analysis to identify key metabolites and pathways associated with the observed immunomodulatory activities.

### **Progress Report:**

Selected three novel medicinal herbs with potential immunomodulatory properties.

Complete dextraction and characterization of bioactive compounds from these plants.

  
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In vitro assessment of immunomodulatory activities using relevant cellular model and biochemical assays is ongoing.

## Next Steps:

Complete the in vitro evaluation of immunomodulatory activities.

Conduct in vivo studies in animal models to assess the immunomodulatory effects of these medicinal herbs.

Perform metabolomics analysis to identify key metabolites and pathways involved in the observed immunomodulatory activities.


## Conclusion:

The project is progressing as planned, and we anticipate completing the proposed objectives within the stipulated 2-year duration. The findings from this study will

Contribute to the development of novel phyto pharmaceuticals with immunomodulatory properties, addressing the growing need for safe and effective therapies in this area.



  
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## **Project Title: Comparative Pharmacognostical Photochemical in vitro screening of Ant diabetic Activities of Monochoria Species**

**Principal Investigator:** Dr. Deenanath Jhade, Professor, Department of Pharmacognosy

### **Funding Details:**

Total Cost of the Proposed Project: Rs.2,00,000

Amount Sanctioned for Project Completion:Rs.1,00,000

Amount Sanctioned after Progress Report Submission: Rs.1,00,000

Project Duration: 2Years

### **Objectives:**

Compare the pharmacognostical and phytochemical profiles of different Monochoria species.

Screen the anti diabetic potential of various Monochoria species using in vitro assays, such as alpha-amylase and alpha-glucosidase inhibition.

Identify the most active Monochoria species and isolate the bioactive compounds responsible for antidiabetic activity.

Evaluate the safety and tolerability of the selected Monochoria species and their bioactive compounds.

### **Methodology:**

Collection and authentication of various Monochoria species. Pharmacognostical and phytochemical characterization of the selected species.

In vitro antidiabetic activity screening using enzyme inhibition assays (alpha-amylase and alpha-glucosidase).

Isolation and identification of bioactive compounds from the most active Monochoria species.


Safety and tolerability assessment of the selected species and their bioactive compounds using in vitro and in vivo models.

### **Progress Report:**

Completed collection and authentication of different Monochoria species.



  
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Pharmacognostical and phytochemical characterization of the selected species is ongoing.

## Next Steps:

Complete pharmacognostical and phytochemical characterization of the selected species.

Conduct in vitro antidiabetic activity screening using enzyme inhibition assays.

Identify the most active Monochoria species and isolate their bioactive compounds.

Perform safety and tolerability studies for these lectedspecies and bioactive compounds.

Analyze the results and prepare the final project report.

## Conclusion:

The project is progressing as planned, with successful collection and authentication of different Monochoria species. We anticipate completing the proposed objectives within the stipulated 2-year duration, leading to the identification of potential antidiabetic agents from Monochoria species for further research and development.



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## Project Title: Comparative Pharmacognostical Photochemical in vitro Screening of Antidiabetic Activities of Monochoria Species

Principal Investigator: Dr. Deenanath Jhade, Professor, Department of Pharmacognosy Funding

Details:

Total Cost of the Proposed Project: Rs. 2,00,000

Amount Sanctioned for Project Completion: Rs. 1,00,000

Amount Sanctioned after Progress Report Submission: Rs. 1,00,000 Project

Duration: 2 Years

### Objectives:

Compare the pharmacognostical and phytochemical profiles of different Monochoria species.

Screen the antidiabetic potential of various Monochoria species using in vitro assays, such as -  
amylase and -glucosidase inhibition.

Identify the most active Monochoria species and isolate the bioactive compounds responsible for  
antidiabetic activity.

Evaluate the safety and tolerability of the selected Monochoria species and their bioactive  
compounds.

Methodology:

Collection and authentication of various Monochoria species. Pharmacognostical and  
phytochemical characterization of the selected species.

In vitro antidiabetic activity screening using enzyme inhibition assays ( -amylase and -  
glucosidase).

Isolation and identification of bioactive compounds from the most active Monochoria species.

Safety and tolerability assessment of the selected species and their bioactive compounds  
using in vitro and in vivo models.

Progress Report:

Collected and authenticated different Monochoria species.



*Deenanath Jhade*  
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Completed pharmacognostical and phytochemical characterization of the selected species.  
Ongoing in vitro antidiabetic activity screening using enzyme inhibition assays. Next Steps:

Complete the in vitro antidiabetic activity screening.

Identify the most active Monochoria species and isolate their bioactive compounds.


Conduct safety and tolerability studies for the selected species and bioactive compounds.

Analyze the results and prepare the final project report. Conclusion:

The project is on track, with successful collection, authentication, and characterization of different Monochoria species. We anticipate completing the proposed objectives within the stipulated 2-year duration, leading to the identification of potential antidiabetic agents from Monochoria species.



  
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## **Project Title: Assessment of Medicinal Plants' Anxiolytic, Antidepressant, Anticonvulsant, and Nootropic Properties**

**Principal Investigator:** Dr. Akshay Meshram, Assistant Professor, Department of Pharmacology

### **Funding Details:**

Total Cost of the Proposed Project: Rs. 6,00,000

Amount Sanctioned for Project Completion: Rs. 5,00,000

Amount Sanctioned after Progress Report Submission: Rs. 4,00,000 Project

Duration: 2 Years

### **Objectives:**

Evaluate the anxiolytic, antidepressant, anticonvulsant, and nootropic properties of selected medicinal plants.

Identify bioactive compounds responsible for these therapeutic effects.

Investigate the molecular mechanisms underlying the observed pharmacological activities.

Assess the potential for developing novel phytopharmaceuticals targeting mental health conditions.

### **Methodology:**

Selection of medicinal plants based on traditional usage and existing scientific literature.

Extraction and characterization of bioactive compounds from the selected plants.

In vitro assessment of anxiolytic, antidepressant, anticonvulsant, and nootropic activities using relevant cellular models and biochemical assays.

In vivo evaluation of these therapeutic effects in animal models of anxiety, depression, epilepsy, and cognitive impairment.

Metabolomics analysis to identify key metabolites and pathways associated with the observed pharmacological activities.



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## Progress Report:

Selected five medicinal plants with potential anxiolytic, antidepressant, anticonvulsant, and nootropic properties.

Completed extraction and characterization of bioactive compounds from these plants.

In vitro assessment of therapeutic activities using relevant cellular models and biochemical assays is ongoing.

## Next Steps:

Complete the in vitro evaluation of anxiolytic, antidepressant, anticonvulsant, and nootropic activities.

Conduct in vivo studies in animal models of anxiety, depression, epilepsy, and cognitive impairment.

Perform metabolomics analysis to identify key metabolites and pathways involved in the observed pharmacological activities.

## Conclusion:

The project is progressing as planned, and we anticipate completing the proposed objectives within the stipulated 2-year duration. The findings from this study will

Contribute to the development of novel phytopharmaceuticals targeting mental health conditions, thus addressing the growing need for safe and effective therapies in this area.



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## Project Title: Development and Evaluation of Nano-ocular

### Formulation for Glaucoma

Principal Investigator: Dr. Uma Patil, Professor, Department of Pharmaceutics

Funding Details:

Total Cost of the Proposed Project: Rs. 2,00,000

Amount Sanctioned for Project Completion: Rs. 1,00,000

Amount Sanctioned after Progress Report Submission: Rs. 1,00,000Project

Duration: 2 Years

### Objectives:

Develop a nano-ocular formulation for the effective delivery of antiglaucoma drugs.

Evaluate the physicochemical properties, stability, and drug release profiles of the formulation.

Assess the in vitro and in vivo efficacy of the nano-ocular formulation in reducing intraocular pressure.

Investigate the safety and tolerability of the nano-ocular formulation in animal models.

### Methodology:

Selection and optimization of nanocarrier systems for the formulation. Preparation of the nano-ocular formulation with the chosen antiglaucoma drug. Physicochemical characterization and stability studies of the formulation.

In vitro drug release studies using appropriate methods.

In vivo efficacy assessment in animal models with induced glaucoma, including intraocular pressure measurement and histopathological analysis.

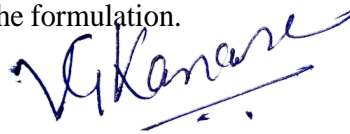
Safety and tolerability evaluation through ocular irritation and systemic toxicity studies in animal models.

Progress Report:

Completed selection and optimization of nanocarrier systems for the formulation.



  
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Formulation development and preliminary physicochemical characterization are ongoing.

Next Steps:

Complete physicochemical characterization and stability studies. Conduct in vitro drug release studies.

Evaluate in vivo efficacy and safety of the nano-ocular formulation. Analyze the results and prepare the final project report.

## Conclusion:

The project is progressing well, with successful selection and optimization of nanocarrier systems for the formulation. We expect to achieve the proposed objectives within the stipulated 2-year duration, contributing to the development of an effective and safe nano-ocular formulation for improved glaucoma treatment.



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## **Project Title: Formulation and Evaluation of Novel Topical Formulation Based on Surfactants for Antifungal Treatment**

Principal Investigator: Dr. Sanchita Mandal, Professor, Department of Pharmaceutics Funding

Details:

Total Cost of the Proposed Project: Rs. 3,00,000

Amount Sanctioned for Project Completion: Rs. 2,00,000

Amount Sanctioned after Progress Report Submission: Rs. 1,00,000 Project

Duration: 2 Years

### **Objectives:**

Develop a novel topical formulation using surfactants as a delivery system for antifungal drugs.

Evaluate the physicochemical properties of the formulation, including stability, viscosity, and drug content.

Investigate the in vitro antifungal activity of the formulation against various fungal strains.

Conduct in vivo studies to assess the efficacy and safety of the formulation in an appropriate animal model.

### **Methodology:**

Selection and optimization of surfactants for the formulation. Preparation of the antifungal drug-loaded surfactant-based formulation.

Physicochemical characterization of the formulation, including stability, viscosity, and drug content determination.

In vitro antifungal activity assessment against fungal strains using broth dilution and/or agar diffusion methods.

In vivo efficacy and safety evaluation of the formulation using an appropriate animal model.

### **Progress Report:**

Completed selection and optimization of surfactants for the formulation.



*Sanchita Mandal*  
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Formulation development and preliminary physicochemical characterization are ongoing.

## Next Steps:

Complete the physicochemical characterization of the formulation. Conduct

in vitro antifungal activity studies.

Perform in vivo efficacy and safety assessment. Analyze the

results and prepare the final project report.

## Conclusion:

The project is progressing as planned, with successful selection and optimization of surfactants for the formulation. We expect to achieve the proposed objectives within the stipulated 2-year duration, leading to the development of a safe and effective surfactant-based antifungal topical formulation for improved treatment outcomes.



  
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## **Project Title: Development of a Topical Herbal Formulation for Wound Healing from Selected Drug of Centella Asiatica**

**Principal Investigator:** Dr. Deenanath Jhade, Professor, Department of Pharmacognosy

### **Funding Details:**

Total Cost of the Proposed Project: Rs. 3,00,000

Amount Sanctioned for Project Completion: Rs. 2,00,000

Amount Sanctioned after Progress Report Submission: Rs. 1,00,000 Project

Duration: 2 Years

### **Objectives:**

Develop a topical herbal formulation from Centella Asiatica for wound healing. Evaluate the physicochemical properties and stability of the developed formulation. Assess the in vitro drug release and in vivo wound healing efficacy in animal models. Investigate the safety and tolerability of the topical herbal formulation.

### **Methodology:**

Extraction and characterization of bioactive compounds from Centella Asiatica.

Formulation of the herbal topical preparation, such as a gel or ointment.

Characterization of the formulation for drug content, consistency, pH, and stability. In vitro drug release studies using artificial membranes.

In vivo wound healing efficacy evaluation in animal models, including measurement of wound closure rate, re-epithelialization, and collagen deposition.

Safety and tolerability assessment through skin irritation and histopathological studies in animal models.

### **Progress Report:**

Completed extraction and characterization of bioactive compounds from Centella Asiatica.

Developed a herbal gel formulation and characterized its physicochemical properties. In vitro drug release studies using artificial membranes are ongoing.



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## Next Steps:

Complete the in vitro drug release studies.

Conduct in vivo studies in animal models to evaluate wound healing efficacy. Perform safety and tolerability studies in animal models.

Finalize the formulation and prepare for scale-up and further preclinical and clinical evaluations.

## Conclusion:

The project is progressing well, with the successful development of a topical herbal gel formulation from Centella Asiatica. We anticipate completing the proposed objectives within the stipulated 2-year duration, leading to the development of an effective and safe herbal remedy for wound healing.



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**Project Title:** Formulation and Evaluation of Novel Ophthalmic Drug Delivery System

**Principal Investigator:** Dr. Sanchita Mandal, Professor, Department Of Pharmaceutics

## Funding Details:

Total Cost of the Proposed Project:Rs.3,00,000

Amount Sanctioned for Project Completion:Rs.2,00,000

Amount Sanctioned after Progress Report Submission: Rs. 1,00,000

**Project Duration:** 2Years

## Objectives:

- 1.Develop a novel ophthalmic drug delivery system for improved bioavailability and therapeutic efficacy.
- 2.Evaluate the physicochemical properties and stability of the developed formulations.
- 3.Assess the in vitro drug release and in vivo performance of the formulations in animal models.
- 4.Investigate the safety and tolerability of the novel ophthalmic drug delivery system.

## Methodology:

- 1.Selection and optimization of drug delivery system components, such as polymers, surfactants, and stabilizers.
- 2.Formulation of drug-loaded ophthalmic drug delivery systems using optimized components.
- 3.Characterization of the formulations for particle size, zeta potential, drug loading, and encapsulation efficiency.
- 4.Invivo drug release studies in simulated tear fluid and Comparison with conventional ophthalmic formulations.
- 5.In vivo performance evaluation in animal models, including pharmacokinetic and pharmacodynamic studies.

- 1.Selection and optimization of drug delivery system components, such as polymers, surfactants, and stabilizers.

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2. Formulation of drug-loaded ophthalmic drug delivery systems using optimized components.
3. Characterization of the formulations for particlesize, zeta potential, drug loading, and encapsulation efficiency.
4. In vitro drug release studies in simulated tear fluid and Comparison with conventional ophthalmic formulations.
5. In vivo performance evaluation in animal models, including pharmacokinetic and pharmacodynamic studies.

## PROGRESS REPORT:

1. Optimized components for the novel ophthalmic drug delivery system.
2. Formulated drug-loaded ophthalmic drug delivery systems and characterized their physicochemical properties.
3. Completed invitro drug release studies and compared the results with conventional formulations.

## NEXT STEP:


1. Perform in vivo studies in animal models to assess pharmacokinetics, pharmacodynamics, and therapeutic efficacy.
2. Conduct safety and tolerability studies to support the potential use of the novel ophthalmic drug delivery system.
3. Finalize the formulation and prepare for scale-up and further preclinical and clinical evaluations.

## Conclusion:

The project that progressed well, with the successful development and initial characterization of a novel ophthalmic drug delivery system. We anticipate completing the proposed objectives within the stipulated 2- year duration, leading to the development of an advanced ophthalmic formulation with improved bioavailability and therapeutic efficacy



  
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