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and The Directorate of Medical Education, Chennai.

B.Pharm Students Undertaking Project Work/ Field Work/ Internship For The Academic Year 2023 – 2024

S.NO	DESCRIPTION		
1	Certificate of Head of institution		
2	List of B.Pharm Students Undertaking Project Work/ Field Work/ Internship-HOI		
3	List of Students Undertaking Project Work/ Field Work/ Internship		



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Certificate Of Head Of Institution



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and The Directorate of Medical Education, Chennai.

Dr.C. Jothimanivannan., M. Pharm., Ph.D.

Principal

TO WHOMSOEVER IT MAY CONCERN

Number of students under taking **Project Work**/ Field Work/ Internship for the academic year 2023 - 2024 is 55.

The student participated in more than one activity has been counted as **one** only.

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PRINCIPAL, SS INSTITUTE OF PHARMACY, KUPPANUR (PO), SANKARI (TK). SALEM -637301



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Dr.C.Jothimanivannan., M.Pharm., Ph.D.

Principal

To whomsoever it may concern

This Is To Certify That The List Of **B.Pharm** Students_Under Taking **Project Work/** Field Work/ Internship For The Academic Year 2023 – 2024 Are Given Below.

S. No	Register No	Name Of The Student	Year	Project Work Topic	Guide	
entants provided the same states	561997227	PRADEEP .B	IV	PHARMACOGNOSTICAL,		
	561997234	SANTHOSHKUMAR.N	IV	PHYTOCHEMICAL &	Mr.T. SAMPATH KUMAR., M.PHARM	
1	561997236	SHALINI .M	IV	INVITRO ANTI- LICE ACTIVITY OF		
	561997238	SREEJITH .S	IV	ETHANOLIC EXTRACT		
	561997247	VIJAYAKUMAR .A	IV	OF CITRUS HYSTRIX		
2	561997235	SARANKUMAR.T	IV	IN-VITRO ACTIVITY OF		
-	561997237	SHANMUGAM.G	IV	ETHANOLIC EXTRACT		
of the latest and the	561997239	SRINIVASAN.M	IV	OF ALBIZIA AMARA IN	Mrs. M. VANITHA.,	
	561997240	SRITHER.R	IV	INDIAN ADULT EARTH	M.PHARM.	
	561997244	TAMILSELVAN.R	IV	WORMS (PHERETIMA POSTHUMA)		
3	561997208	DEEPASREE, M	IV	FORMULATION &		
	561997216	HEMAVATHI. D	IV	EVALUATION OF FAST	50	
	561997221	KUMANAN. N.V	IV	DISSOLVING ORAL THIN	Mrs.K.Sowndarya.,	
	561997226	PAVENDHARAN .S	IV	FILMS OF ATORVASTATIN CALCIUM	M.PHARM	
4	561997204	ARUN .S	IV	FORMULATION		
	561997209	DHANASEKARAN .A	IV	&EVALUATION OF	1	
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	561997215	GOKULNATH .M	IV	FUMARATE FLOATING	Mrs. M GOMATHI	
	561997217	KAMALESH .S	STEE STEE	TABLETS BY NON EFFERVESCENTRIA TECHNIQUE	M. Pharm	
5	561997241	SUBASHINI .M	S IV	PHYDOCHEMICAL & V	Dr.C.Jotnimaniyannan M.FHAR.A., PALD.	

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C) may and	561997243	SURYAPRAKASH .M	IV	ACTIVITY EVALUATION		
- Inches	561997245	THANGAPANDIYAMM	IV	OF ROOT		
Linguistici		AL.I		OF ROOT EXTRACT OF		
-	561997246	VEERASELVAN .V	IV	VITEX NEGUNDO LINN		
1	561997248	VIJAYAKUMAR .M	IV			
-	561997229	PRAMESHWARAN, A	IV	INVITRO EVALUATION		
-	561997230	PRIYANKA. P	IV	OF		
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6	561997242	SURESH. M	IV	OF ANTIFUNGAL	Mrs. M. VANITHA.,	
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	561997203	ARTHI. A	IV	ANALYTICAL METHOD		
	561997206	ARUNKUMAR. K	IV	DEVELOPMENT		
	561997207	AZHAGIRI, T. K	IV	VALIDATION FOR THE		
7	561997223	MANIKANDAN. S	IV	DETERMINATION OF	Mr.S.PRAVINKUMAR.	
, designation of the second	561997224	MUTHUKANNAN. S	īV	NICORANDIL BY UFIC IN ORAL SOLID DOSAGE FORMS	M.PHARM	
	561997202	ANUSUYA. N	IV	FORMULATION &		
	561997205	ARUNKUMAR. D	IV	EVALUATION OF HERBAL		
	561997213	ELAVARASAN. E	IV	CREAM BY USING	Mrs.C,KALAISELV	
8	561997218	KARTHICK, D	IV	BAUHINIA TOMENTOSA	M.Pharm., (Ph.D).,	
0	561997219	KARTHIKEYAN. R	IV	FOR EVALUATION OF ANTI INFLAMMATORY ACTIVITY	Associate Professor	
Λ	561997250	VINOTHKUMAR.R	IV	FORMULATION &		
2 %	560020523560	MYVIZHIKANNAN.R	IV	EVALUATION OF	Mrs. P.NANDHITHA	
9	560020523563	PRAKASH.V	IV	MATRIX TYPE	M. PHARM	
	560020523565	SUGAVARSHAN.C	IV	TRANSDERMAL	Assistant Professor	
	560020523566	VETRIVEL.R	IV	PATCHES OF IBUPROFEN		
	561997228	PRADEEP. T	IV	PHARMACOGNOSTICAL		
10	5600205235 58	GNANAVEL.S	IV	& PHARMACOLOGICAL ANALYSIS OF		
Political Politi	5600205235 59	GOKUL. K	IV	PHYTOCHEMICALS, WITH IN-VITRO	Mrs.V.DEEPABHARATH M.P.HARM	
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-	561997210	DHARUN. A	//e/ IV	IN-VITRO	Mr.M. PRAVEEM	
11	561997212	DINESH.T	* JV	ANTRELMINTIC	KUMAR	



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5600205235 64	SATHISHKUMAR. V	IV	(PHERETIMA POSTHUMA)	



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SALEM 637301





"Formulation and Evaluation of Fast Dissolving Oral Thin Films Of Atorvastatin Calcium"

A Dissertation submitted to

THE TAMILNADU Dr. M.G.R. MEDICAL UNIVERSITY

Chennai-600032

In partial fulfillment for the award of degree of BACHELOR OF PHARMACY

Submitted by

M.DEEPASREE

REG.NO: 561997208

D.HEMAVATHI

REG.NO: 561997216

N.V.KUMANAN

REG.NO: 561997221

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Under the Guidance of

Mrs.K. SOWNDARYA., M.PHARM.,

DEPARTMENT OF PHARMACEUTICS



SS INSTITUTE OF PHARMACY

SANKARI, SALEM (DT), TAMIL NADU - 637301

SEPTEMBER-2023 (Dec-2023)

PRINCIPAL, SS INSTITUTE OF PHARMACY, KUPPANUR (PO), SANKARI (TK).





CERTIFICATE

This is to certify that the Project entitled "Formulation and evaluation of fast dissolving oral film of atorvastatin calcium" submitted to The Tamilnadu Dr. M.G.R Medical University, Chennai, is a bonafide project work of

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in partial fulfillment of the degree of Bachelor of Pharmacy under the guidance of Mrs. K. SOWNDARYA, M.Pharm., Assistant Professor, Dept. of Pharmaceutics, SS Institute of Pharmacy, Sankari, Salem.

Dr. C. Jothimanivannan., M. Pharm., Ph.D.,

Principal,

SS Institute of Pharmacy,

Sankari, Salem - 637301.

Place:SS Institute of Pharmacy, Sankari- 637301.

Date: 30 · 12 · 20213 ·

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DECLARATION

The work presented in this dissertation entitled "Formulation and evaluation of fast dissolving oral film of atorvastatin calcium" was carried out by N.V.KUMANAN (REG.NO:561997221),

S.PAVENDHARAN(REG.NO:561997226),

D.HEMAVATHI(REG.NO:561997216), M.DEEPASREE (REG.NO:561997208) ,to The Tamilnadu Dr. M.G.R. Medical University, Chennai, in partial fulfillment for the degree of Bachelor of Pharmacy was carried out by the candidates at the department of pharmaceutics, under the supervision of Mrs. K. SOWNDARYA, M.Pharm., Assistant Professor, Dept. of Pharmaceutics, SS Institute of Pharmacy, Sankari, during the academic year 2022-23. We further declare that this work is original and has not been submitted in part or full for the award of any other degree or diploma in any other university.

Internal Examiner 30/12/23

Convener of Examination

Place: SS Institute of pharmacy, Sankari

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SUMMARY AND CONCLUSION

- OTFs have emerged as a revolutionary trend and most pharmaceutical companies in this field continue their research and development activities to adapt their drugs in various categories to this technology
- ➤ The present study indicates a good orally fast dissolving films containing atorvastatin calcium for systemic delivery with an added advantage of faster drug action.
- ➤ Finally, it is concluded that the drug release from the fast dissolving film was increased by using the increased concentration of superdisintegrant, thus assisting in faster disintegration in the buccal cavity.
- As the drug having low solubility, fast disintegration may leads to more drug availability for dissolution, resulting in faster absorption in systemic circulation increased systemic availability of drug may leads to quick onset of action.
- > The formulation F4 was found to be best formulation which releases 98.95 % of the drug within 15 min and disintegration time is 168sec which was significantly high when compared to other formulation.

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PHYTOCHEMICAL AND ANTIMICROMICROBIAL ACTIVITY EVALUATION OF ROOT EXTRACT OF VITEX NEGUNDO Linn

A Dissertation submitted to

THE TAMILNADU Dr. M.G.R. MEDICAL UNIVERSITY

Chennai-600032

In partial fulfillment for the award of degree of

BACHELOR OF PHARMACY

Submitted by

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M. SURIYAPRAKASH REG.NO:561997243
I.THANGAPANDIYAMMAL REG.NO:561997245
V.VEERASELVAN REG.NO:561997246
M.VIJAYKUMAR REG.NO:561997248

Under the Guidance of

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This is to certify that the Project entitled "PHYTOCHEMICAL AND ANTIMICROBIAL ACTIVITY EVALUVATION OF ROOT EXTRACT OF VITEX NEGUNDO LINN" submitted to the Tamilnadu Dr. M.G.R Medical University, Chennai, is a bonafide

project work of

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in partial fulfillment of the degree of Bachelor of Pharmacy under the guidance of Dr. C.

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DECLARATION

presented work in this dissertation entitled "phytochemical The Pharmacological Antimicrobial activity of root extract of vitex negundo linn"was Carried M.SUBASHINI (561997241), V.VEERASELVAN (561997246) by out I.THANGAMPANDIYAMMAL (561997245), M.SURIYAPRAKESH (561997243)

M.VIJAYKUMAR (561997248),,to the Tamilnadu Dr.MGR.medical university, Chennai, in partial fulfillment for the degree of Bachelor of Pharmacy was carried out by the department of pharmacology, under the supervision candidate the Dr. C. Jothimanivannan, M.Pharm., Ph.D., Department of Pharmaceutical Chemistry SS Institute of Pharmacy, Sankari, during the academic year Aug 2023 (March 2023). We further declare that this work is original and has not been submitted in part or full for the award of any other degree in any other university.

Convener of Examination

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V.VEERASELVAN

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CHAPTER- VIII

8. SUMMARY AND CONCLUSION

Antimicrobial properties of medicinal plants are being increasingly reported from different parts of the world.

The World Health Organization (WHO) estimates the plant extract or active constituents are used as folk medicine in traditional therapies of 80% of the world population.

In the present work the aqueous extract obtained from Vitex negundo linn roots showed strong activity against the selected microbial strains.

The result of phytochemical in the present investigation showed that the plant contains effective constituents like alkaloid, glycosides, flavonoids, tannins, amino acids, proteins and reducing sugar.

In this study different concentration of aqueous root extract 25µl, 50µl, 75µl, 100µl. Were used against the different microbial strains shows increase in zone of inhibition was observe on increasing the concentration of aqueous root extract.

Among the different microbial strains a significant inhibitory action was observed against Staphylococcus (G+ve), E.Coli (G-ve) and a minimal inhibitory action against Streptococcus (G-ve). However further studies are needed to better evaluate the potential effectiveness of the crude extract, attempt is worth to isolate a broad spectrum antimicrobial agents.

The present result will form the basis for selection investigation in the potential discovery of new natural bio-active compound.

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FORMULATION AND EVALUATION OF MATRIX TYPE TRANSDERMAL PATCHES OF IBUPROFEN

Dissertation submitted to

THE TAMILNADU DR. M.G.R. MEDICAL UNIVERSITY,

CHENNAI-32

In partial fulfillment for the award of degree of BACHELOR OF PHARMACY

Submitted By

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MAY 2024(MARCH 2024)





CERTIFICATE

This is to certify that the dissertation entitled "FORMULATION AND EVALUATION OF MATRIX TYPE TRANSDERMAL PATCHES OF IBUPROFEN" is the bonafide work carried out by Mr.VINOTHKUMAR R, Mr.MYVIZHIKANNAN R, Mr.PRAKASH V, Mr.SUGAVARSAN C, Mr.VETRIVEL R, under the guidance of Mrs.P.Nandhitha, M.PHARM., Assistant Professor, Department of Pharmaceutics, SS Institute of Pharmacy, Sankari, Salem in partial fulfillment of requirements for the award of degree of Bachelor of Pharmacy.

This is forwarded to The Tamilnadu Dr. M.G.R. Medical University, Chennai.

Prof. Dr.C. Jothimanivannan, M.Pharm, Ph.D.,

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Date: 19/06/24

Place: Sankari

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CERTIFICATE

This is to certify that the dissertation entitled "FORMULATION AND EVALUATION OF MATRIX TYPE TRANSDERMAL PATCHES OF IBUPROFEN" is the bonafide work carried out Mr.VINOTH KUMAR R, Mr.MYVIZHIKANNAN R, Mr.PRAKASH V, Mr.SUGAVARSAN C, Mr.VETRIVEL R, under my guidance and supervision. This is forwarded to The Tamilnadu Dr. M.G.R. Medical University, Chennai in a partial fulfillment of requirements for the award of degree of Bachelor of Pharmacy.

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DECLARATION

We hereby declare that this dissertation entitled "FORMULATION AND EVALUATION OF MATRIX TYPE TRANSDERMAL PATCHES OF IBUPROFEN" has been carried out by us under the Guidance and supervision of Mrs.P.Nandhitha, M.Pharm, Assistant professor, SS Institute of Pharmacy, Sankari. In a partial fulfillment of requirements for the Bachelor of Pharmacy Degree Course. We further declare that we have not submitted this dissertation previously for the award of any degree or diploma.

Mr. VINOTH KUMAR.R

(561997250)

R.mysizlikannan Mr.MYVIZHIKANNAN. R (560020523560)

v .Frakash Mr. PRAKASH. V (560020523563)

Mr. VETRIVEL. R (560020523566) Mr. SUGAVARSAN.C (560020523565)

Date: 19/06/24

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Signature of the Internal Examiner

Date of Examination 19 06 24.

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CHAPTER XI

SUMMARY AND CONCLUSION

In the present work an attempt has been made to formulate and evaluate the transdermal patches of Ibuprofen using various types of polymers (Methylcellulose, Hydroxy propyl methyl cellulose, & Polyvinyl pyrolidone).

The results of compatibility studies by Fourier transform infrared spectroscopy and showed no interaction between the drug and polymers.

The polymers Hydroxypropyl methyl cellulose & Polyvinyl pyrolidone in 1:2 ratioused for the formulation of transdermal patches showed good film forming property when compared to other polymer ratios.

The patches prepared by using Polyvinyl pyrolidone were thin, flexible, smooth and transparent whereas the patches prepared by without using Polyvinyl pyrolidone were thin, flexible, smooth and opaque.

The weight variation test showed less variation in weight and suggesting uniform distribution of drug and polymer.

The thickness of the transdermal patches showed less variation almong all the PRINCIPAL.

polymers with different ratios.

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Polymer patches with methyl cellulose, HPMC & PVP of ratio 3:7:7, & HPMC & pVP of ratio 2:1 &1:1 showed good flexibility and folding endurance properties when compared to other patches.

All the formulations showed 100% flatness which indicating that all patches could maintain a smooth surface when applied on to the skin.

The moisture content in the patches were found to low and the formulations containing polymers are hydrophillic polymer so it should be carefully packed along with dessicant.

The drug content analysis showed minimum variations suggesting uniform distribution of drug. .In vitro release studies showed that drug release of the formulations containing 2:1 ratio of HPMC & PVP (F₆) shows best results.

Hence it was concluded that solvent casting method was useful for successful development of matrix type transdermal patches of Ibuprofen. The controlled release of drug from the transdermal patches suggested that the frequency of administration can be reduced.

The transdermal patches can improve the bioavailability of the Ibuprofen by avoiding hepatic first pass metabolism. The present investigation suggested that the matrix type transdermal patches of Ibuprofen could be explored for the management of the pair.

Inflammatory response.

SS INSTITUTE OF PHARMACY, KUPPANUR (PO), SANKARI (TK). SALEM -637301





"ANALYTICAL METHOD DEVELOPMENT AND VALIDATION FOR THE DETERMINATION OF NICORANDIL BY UFLC IN ORAL SOLID DOSAGE FORMS"

A Dissertation submitted to

THE TAMILNADU Dr. M.G.R. MEDICAL UNIVERSITY

Chennai-600032

In partial fulfillment for the award of degree of

BACHELOR OF PHARMACY

Submitted by

A.ARTHI REG.NO: 561997203

K.ARUNKUMAR REG.NO: 561997206

T.K.AZHAGIRI REG.NO: 561997207

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S.MUTHUKANNAN REG.NO: 561997224

Under the Guidance of

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SANKARI, SALEM (DT), TAMIL NADU - 637381 SEPTEMBER-2023 (Dec-2023)





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This is to certify that the Project entitled "ANALYTICAL METHOD DEVELOPMENT AND VALIDATION FOR THE DETERMINATION OF NICORANDIL BY UFLC IN ORAL SOLID DOSAGE FORMS" submitted to The Tamilnadu Dr. M.G.R Medical University, Chennai, is a bonafide project work of

A.ARTHI REG.NO: 561997203

K.ARUNKUMAR REG.NO: 561997206

T.K.AZHAGIRI REG.NO: 561997207

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in partial fulfillment of the degree of Bachelor of Pharmacy under the guidance of Mr.S.PRAVINKUMAR, M.Pharm., Assistant Professor, Dept. of Pharmaceutical Analysis, SS Institute of Pharmacy, Sankari, Salem.

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Principal,

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Date: 30.12.2023

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CERTIFICATE

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in partial fulfillment of the Degree of Bachelor of Pharmacy under my guidance and supervision in the Department of Pharmaceutical Analysis, SS Institute of Pharmacy, Sankari, Salem.

Mr. S.PRAVINKUMAR., M.Pharm.,

S. P. Jumas

Supervisor/guide, Asst. Professor,

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The work presented in this dissertation entitled "ANALYTICAL METHOD DEVELOPMENT AND VALIDATION FOR THE DETERMINATION OF NICORANDIL BY UFLC IN ORAL SOLID DOSAGE FORMS" was carried out by A.ARTHI (REG.NO:561997203), K.ARUNKUMAR (REG.NO:561997206), T.K.AZHAGIRI (REG.NO:561997207), S.MANIKANDAN (REG.NO:561997223) S.MUTHUKANNAN (REG.NO:561997224), to The Tamilnadu Dr. M.G.R. Medical University, Chennai, in partial fulfillment for the degree of Bachelor of Pharmacy was carried out by the candidates at the department of pharmaceutical analysis, under the supervision of Mr. S.PRAVINKUMAR, M.Pharm., Assistant Professor, Dept. of Pharmaceutical Analysis, SS Institute of Pharmacy, Sankari, during the academic year 2022-23. We further declare that this work is original and has not been submitted in part or full for the award of any other degree or diploma in any other university.

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ANALYTICAL MEHOD DEVELOPMENT AND VALIDATION FOR THE DETERMINATION OF NICORANDIL BY UFLC IN ORAL SOLID DOSAGE FORMS

7. SUMMARY AND CONCLUSION

Parameters		Nicorandil	LIMIT	
Linear Regression equa		y = 0.0000x-5.2442 ($r^2 = 0.9993$)	r² not less than 0.99	
Assa	ıy			
(% mean	assay)	100.78%	98%-102%	
Specificity		Complies	No interference of any peak	
Method precision %RSD		0.51 %	RSD NMT 2.0%	
Intermediate precision day-01 %RSD		0.66%	RSD NMT 2.0%	
Intermediate precision day-02 %RSD Accuracy %		0.71%	RSD NMT 2.0%	
		99.18% to 100.91%	98-102%	
	FM	0.82%	%RSD NMT 2.0	
	FP	0.48%		
Robustness	WM	0.48%		
	WP	0.68%		

Conclusion:

A simple, Accurate, precise method was developed for the simultaneous estimation of the Nicorandil in solid dosage form. Retention time of Nicorandil was found 2.173min. %RSD of the Nicorandil found 0.13%. %Assay was obtained as 100.78% for Nicorandil. Regression equation of Nicorandil is y=0.0000x-5.2442 & r² =0.9993. Retention times were decreased and that run time was decreased, so the method developed was simple and economical that can be adopted in regular Quality control test in Industries.

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Page 110 of 112





FORMULATION AND EVALUATION OF HERBAL CREAM BY USING BAUHINIA TOMENTOSA FOR EVALUATION OF ANTI INFLAMMATORY ACTIVTY

THE TAMILNADU DR. M.G.R. MEDICAL UNIVERSITY

Chennai-600032

In partial fulfilment for the award of degree of

BACHELOR OF PHARMACY

Submitted by

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D. ARUNKUMAR REG.NO: 561997205

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CHAPTER X

10. SUMMARY AND CONCLUSION

10.1 SUMMARY AND CONCLUSION

The topic of our research work is "FORMULATION AND EVALUATION OF HERBAL CREAM BY USING BAUHINIA TOMENTOSA FOR EVALUATION OF ANTI INFLAMMATORY ACTIVITY"

Herbal medicines are the oldest form of health care known to mankind. Though there are several allopathic medicines available, many people still rely on nature which has blessed human kind with abundant medicinal plants for prevention and cure of disease.

The risk of inflammation is still increasing world-wide. Many synthetic drugs are commonly used to treat the different types of inflammation conditions. Even though there is an availability of different synthetic drugs to treat inflammation, an exploration of herbal medicine is essential to meet out the cure for the same. Herbs that are having potential anti-inflammation activity are used to further develop novel drugs nowadays. Among those, two particular plants especially medicinal herbs are

The plants have flavanoids which may be responsible for their potential anti-inflammatory properties as mentioned in the traditional claim. In the present research work, an unique combination of active extracts of the herbs are chosen to formulate a herbal CREAM for the treatment of inflammation. Thus, the study is an attempt to evaluate the anti- inflammation activity of herbal CREAM containing active extracts of Bauhinia Tomentosa. The review of literature showed that the formulation of herbal CREAM containing combined active extracts of Bauhinia Tomentosa and evaluation of anti- inflammation activity has not been reported so far

The project entitled "FORMULATION AND EVALUATION OF HERBAL CREAM BY USING BAUHINIA TOMENTOSA FOR EVALUATION OF ANTI INFLAMMATORY ACTIVITY" has been achieved with the following results.

The above findings helps to standardize both morphology and anatomical characters. The Physicochemical parameters with Asharalues and Extractive papers in passion

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"Pharmacognostical, Phytochemical and Invitro Anti-Lice activity of Ethanolic extract of Citrus hystrix"

A Dissertation submitted to

THE TAMILNADU Dr. M.G.R. MEDICAL UNIVERSITY

Chennai-600032

In partial fulfillment for the award of degree of

BACHELOR OF PHARMACY

Submitted by

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Summary

The project on the antilice activity of Citrus hystrix involved comprehensive investigations, including pharmacognostical studies, phytochemistry studies, and in vitro anti-lice studies.

The pharmacognostical studies provided valuable insights into the botanical characteristics and identification of Citrus hystrix.

Phytochemistry studies revealed the presence of bioactive compounds in the plant, contributing to its medicinal properties.

The in vitro anti-lice studies involved the introduction of lice across different concentrations, demonstrating notable activity.

Conclusion:

In conclusion, the collective findings of the project underscore the potential of Citrus hystrix as a promising candidate for anti-lice treatments.

The integration of pharmacognostical and phytochemistry studies enriched our understanding of the plant's composition.

The in vitro anti-lice studies further affirmed its efficacy, albeit with variations in activity levels across concentrations.

This research lays the foundation for future exploration and development of Citrus hystrixbased formulations for combating lice infestations.

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IN-VITRO ACTIVITY OF ETHONOLIC EXTRACT OF ALBIZIA AMARA IN INDIAN ADULT EARTH WORMS (PHERETIMA POSTHUMA)

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8. DISCUSSION

Albizia Amara worldwide growing plant exhibits potent diabetic, anti inflammatory, anti oxidant and antihypertensive activity, anti oxidant (Tamaki Hajime b, et al., 2008), Anti ulcer (Thamotharan G et al., 2012). In the present study, we have tested the anti-hyperglycemic potency of Albizia Amara ethanolic extract in Alloxan induced diabetic rats as well as anti-hyperlipidemic activity. It is suggested that the anti-hyperglycemic and anti-hyperlipidemic activities of the plant is due to the active principles of plant source. The phytochemical screening of the ethanolic leaf extract shown that isolation and identification of four flavanoids glycosides such as Rutin, apigenin 6-neohesperidose, kaempferol 3-robinobioside and kaempferol 3-rutinoside (Rakesh kumar gupta et al., 2012) and alkaloids, carbohydrates, glycosides, tannins, and triterpens. (Wollenweber et al., 1988).

Phytochemical screening of the crude extracts revealed the presence of flavonoids and polyphenolic compound as one of the major chemical constituents. Polyphenolic compounds shown activity; chemically tannins are polyphenolic compounds. (Bate-Smith E.C. 1962) Some synthetic phenolic s e.g., niclosamide, oxyclozanide and bithionol are shown to interfere with energy generation in helminth parasites by uncoupling oxidative phosphorylation. (Martin R.J. 1997).

It is possible that tannins contained in the extracts of *Albizia Amara* produced similar effects. Another possible effect of tannins is that they can bind to free protein in the gastrointestinal tract of host animal or glycoprotein on the cuticle of the parasite and cause death. (Mali R.G.et al., 2008)

Ethanolic leaves extracts of the Albizia Amara showed better activity in dose

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PHARMACOGNOSTICAL AND PHARMACOLOGICAL ANALYSIS OF PHYTOCHEMICALS, WITH IN-VITRO EVALUATION OF DIURETIC ACTIVITIES ON BAMBUSA VULGARIS LEAVES

A Dissertation submitted to

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CONCLUSIONS

Pharmacognostical studies

The pharmacognostical studies of the leaves Bambusa vulgaris was carried out to confirm its unique features. This study reveals the following,

- ➤ The macroscopical studies showed the characteristics features of Bambusa vulgaris leaves are green colored, smooth surface, faint characteristics, astringent taste and other morphological characters has been well established. Microscopic investigations are the diagnostic features of the plant and its powder analysis helps to distinguish this particular plant (leaves) in whole form and in crude powder form, Microscoical analysis of Bambusa vulgaris leaves showed the pre Physiochemical constants were determined for the leaves of Bambusa vulgaris such as Ash values, Extractive values, loss on drying. This helps in confirming the identity and purity of this plant and to detect adulterants and its nature.
- > The qualitative analysis were carried out to identify inorganic elements present in the plant.

Phytochemical Studies:

- ➤ The phytochemical studies on the leaves of *Bambusa vulgaris* were carried out to bring its importance as valuable medicinal plant.
- The leaves were extracted with ethanol using soxhlet apparatus and the percentage yield was found to be 3.076% W/W
- The plant powder and extracted were subjected to preliminary phytochemical analysis. The preliminary screening analysis reveals the presence of glycosides, saponins, terpenoids, tannins and phenols, steroids, fixed oils and fats flavoneids.

> TLC analysis of all the extracts was carried out to identify the presence of phytoconstituents.

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IN-VITRO ANTHELMINTIC ACTIVITY OF VARIOUS EXTRACT OF ABUTILON INDICUM IN INDIAN ADULT EARTHWORMS (PHERETIMA POSTHUMA)

A Dissertation submitted to

THE TAMILNADU Dr. M.G.R. MEDICAL UNIVERSITY
Chennai-600032

In partial fulfillment for the award of degree of BACHELOR OF PHARMACY

Submitted by

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Under the Guidance of

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SS INSTITUTE OF PHARMACY SANKARI, SALEM (DT), TAMIL NADU - 637301 MARCH-2024 (MAY-2024)

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This is to certify that the Project entitled *IN-VITRO ANTHELMINTIC ACTIVITY OF VARIOUS EXTRACT OF ABUTILON INDICUM IN INDIAN ADULT EARTHWORMS (PHERETIMA POSTHUMA) ** submitted to The Tamilnadu Dr. M.G.R Medical University, Chennai, is a bonafide project work of

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CONCLUSION

The plant Abutilon indicum proves the Anthelmintic activity in the seeds, Malvaceae family. From the results, the ethanolic extract has potent Anthelmintic activity on Pherethima posthuma., compared with the standard albendazole drug. When the easy availability of the plant, it is suggested to conducting further Invivo studies.

The plants activity is comparable to that of standard anthelmintic drug(albendazole), making it a potential natural alternative for treating intestinal worm infestations.

Further studies are needed to isolate and characterize the specific bioactive compounds responsible for the anthelmintic activity and to evaluate the plants safety and efficacy in human clinical trials.

Overall, the studies suggest that Abulilon indicum has promising anthelmintics properties, making it a valuable resource for the development of natural and sustainable treatment for intestinal worm infection.

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FORMULATION AND EVALUATION OF QUETIAPINE FUMARATE FLOATING TABLETS BY NON EFFERVESCENT TECHNIQUE

Dissertation submitted to

THE TAMILNADU DR. M.G.R. MEDICAL UNIVERSITY,

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This is to certify that the dissertation entitled "FORMULATION AND EVALUATION OF QUETIAPINE FUMARATE FLOATING TABLETS BY NON EFFERVESCENT TECHNIQUE" is the bonafide work carried out by Mr.Arun S, Mr.Dhanasekaran A, Mr.Gokul prasath.M, Mr.Gokulnath M, Mr.Kamalesh.S under the guidance of Mrs.M.Gomathi, M.PHARM., Associate Professor, Department of Pharmaceutics, SS Institute of Pharmacy, Sankari, Salem in partial fulfillment of requirements for the award of degree of Bachelor of Pharmacy.

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CHAPTER XII

SUMMARY AND CONCLUSION

- Gastro retentive Floating drug delivery system offers a simple and practical approach to achieve increased gastric residence time (GRT)
- Quetiapine Fumarate is preferentially absorbed in the proximal small intestine (narrow absorption window), the drug displays oral bioavailability problems in conventional dosage forms.
- The objective of the present investigation was to develop gastroretentive floating matrix tablet of quetiapine fumarate to achieve controlled release and improve the absorption, by retaining the drug in stomach for a prolonged period of time.
- FT-IR and DSC studies indicated that were was no interaction between drug and excipients.
- > Floating matrix tablets were prepared by Non-effervescent technique using different concentrations of gel forming hydrophilic polymers.
- > The formulated tablets were analysed for precompression & post compression parameters, in vitro drug release, etc.
- The precompression parameter of all the formulations were within the required limit that was suitable for formulation of the tablets.
- > The postcompression parameters such as hardness, friability, uniformity in weight, & drug content of all the formulated tablets were within the acceptable limits.

> The floating lag time and duration of buoyancy were found to be satisfied

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INVITRO EVALUATION OF ANTIFUNGAL POTENCY OF THE PLANT EXTRACTS OF DIFFERENT PARTS OF CELOSIA ARGENTEA LINN AGAINST CANDIDA SPECIES

A Dissertation submitted to

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In partial fulfillment for the award of degree of

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CHAPTER X 10. SUMMARY AND CONCLUSION

The dried plant of Celosia argentea belonging to the family Amaranthaceae has been examined to gain an insight of its phytochemical and pharmacological behaviors.

The phytochemical investigation of ethanolic extract of Celosia argentea plant showed the presence of Flavonoids, Alkaloids, Saponins, Steroids, Triterpenoides, Glycosides, Tannins.

The ethanolic extract of dried Celosia argentea plant was also tested for its in vitro antifungal activity against the candida aspergillus, the extract also showed significant anti-fungal activity.

In conclusion, ethanolic extracts of Celosia argentea showed significant antifungal activity against the fungal species candida aspergillus by Invitro studies.

The findings of the present study concluded that the ethanolic extracts of Celosia argentea can effectively control the growth of fungi by invitro studies.

These plant extracts showed excellent inhibitory activity against the fungal species, Candida aspergillus.

Among these ethanolic extracts possessed superior antifungal activity extract of Celosia argentea

The present study justified the claimed used of plants in the traditional system of medicines to treat various fungal infections caused by the Candida species.

However, further studies are needed to better evaluate the potential effortiveness of the crude extracts as the Antifungal agents.

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