

7.2. Best Practice NIRF Ranking



National Institutional Ranking Framework Ministry of Education Government of India



India Rankings 2021: Pharmacy (Rank-band: 76-100)

Institution list in alphabetical order

Bat

AISSMS College of Pharmacy	Pune	Maharashtra
Arulmigu Kalasalingam College of Pharmacy	Srivilliputtur	Tamil Nadu
B. K. Mody Government Pharmacy College, Rajkot	Rajkot	Gujarat
CMR College of Pharmacy	Rangareddy	Telangana
College of Pharmacy, Madras Medical College	Chennal	Tamil Nadu
Dadasaheb Balpande College of Pharmacy	Nagpur	Maharashtra
Dr. B. C. Roy College of Pharmacy and Allied Health Sciences	Durgapur	West Bengal
Dr. Vishwanath Karad MIT World Peace University	Pune	Maharashtra
Galgotias University	Gautam Budh Nagar	Uttar Pradesh
Ganpat University	Ganpat Vidyanagar	Gujarat
Girijananda Chowdhury Institute of Pharmaceutical Science	Guwahati	Assam
Gokaraju Rangaraju College of Pharmacy	Hyderabad	Telangana
layoti Vidyapeeth Women's University	Jaipur	Rajasthan
KIET Group of Institutions	Ghaziabad	Uttar Pradesh
KLE College of Pharmacy	Bengaluru	Karnataka
KLE College of Pharmacy, Hubli	Hubballi	Karnataka
Maliba Pharmacy College	Tarsadi	Gujarat
Nirmala College of Pharmacy, Mangalagiri	Mangalagiri	Andbra Pradesh
Principal K.M. Kundnani College of Pharmacy	Mumbai	Maharashtra
Sanjivani College of Pharmaceutical Education and Research	Kopargaon	Maharashtra
Shree Warana Vibhag Shikshan Mandal's Talyasaheb Kore College of Pharmacy, Warananagar	Warananagar	Maharashtra
Shri Vishnu College of Pharmacy	Bhimavaram	Andhra Pradesh
Suresh Gyan Vihar University	Jaipur	Rajasthan
The Rashtrasant Tukadoji Maharaj Naggur University	Nagpur	Maharashtra
Vivekanand Education Society's College of Pharmacy	Mombai	Maharashtra

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National Institutional Ranking Framework Ministry of Education

Government of India



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India Rankings 2018: Pharmacy (Rank-band: 51-75)

Institution list in alphabetical order

Acharya & B M Reddy College of Pharmacy	Bengaluru	Karnataka
Acharya Nagarjuna University College of Pharmaceutical Sciences	Guntur	Andhra Pradesh
Adina Institute of Pharmaceutical Sciences	Sagar	Madhya Pradesh
AISSMS College of Pharmacy	Pune	Maharashtra
Al Shifa College of Pharmacy	Kozhikode	Kerala
B. K. Modi Government Pharmacy College	Rajkot	Gujarat
Bharati Vidyapeeth College of Pharmacy	Kolhapur	Maharashtra
Chalapathi Institute of Pharmaceutical Sciences	Guntur	Andhra Pradesh
Girijananda Chowdhury Institute of Pharmaceutical Science	Guwahati	Assam
Gupta College of Technological Sciences	Asansol	West Bengal
Karpagam College of Pharmacy	Colmbatore	Tamil Nadu
KMCH College of Pharmacy	Coimbatore	Tamil Nadu
KVSR Siddhartha College of Pharmaceutical Sciences	Vijayawada	Andhra Pradesh
Maharashtra Institute of Pharmacy	Pune	Maharashtra
Maharishi Markandeshwar	Ambala	Haryana
NSHM Knowledge Campus	Kolkata	West Bengal
S. E. T. 's College of Pharmacy	Dharwad	Karnataka
Samskruti College of Pharmacy	Hyderabad	Telangana
Sanjivani College of Pharmaceutical Education and Besearch	Kopargaon	Maharashtra
Shree Warana Vibhag Shikshan Mandal's Tatyasaheb Kore College of Pharmacy	Warananagar	Maharashtra
Sri Padmavathi Mahila Visva Vidyalayam	Tirupathi	Andhra Pradesh
Sri Ramakrishna Institute of Paramedical Sciences	Coimbatore	Tamil Nadu
Sri Venkateswara College of Pharmacy	Chittoor	Andhra Pradesh
The Rashtrasant Tukadoj/ Maharaj Nagour University	Nagpur	Maharashtra
Vivekanand Education Society's College of Pharmacy	Mumbai	Maharashtra

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India Rankings 2017: Pharmacy (Rank-band: 51-75)

Institution list in alphabetical order

Al Shifa College of Pharmacy	Perintalmanna	Kerala
B.K. Mody Govt. Pharmacy College	Rajkot	Gujarat
Bharati Vidyapeeth's College of Pharmacy	Navi Mumbai	Maharashtra
Chebrolu Hanumaiah Institute of Pharmaceutical Sciences	Guntur	Andhra Pradesh
Chitkara University	Rajpura	Punjab
Columbia Institute of Pharmacy	Raipur	Chhattisgarh
Dr. Samuel George Institute of Pharmaceutical Sciences	Markapur	Andhra Pradesh
Girijanunda Chowdhury Institute of Pharmaceutical Science	Guwahati	Assam
Gokaraju Rangaraju College of Pharmacy	Hyderabad	Telangana
Govt. College of Pharmacy	Aurangabad	Maharashtra
Guru Nanak Institute of Pharmaceutical Science & Technology	Kolkata	West Bengal
I.T.S College of Pharmacy	Ghaziabad	Uttar Pradesh
KMCH College of Pharmacy	Coimbatore	Tamif Nadu
Maharashtra Academy of Engineering & Educational Research's Maharashtra Institute of Pharmacy	Pune	Maharashtra
Maharishi Markandeshwar University	Ambala	Haryana
Ponda Education Society's Rajaram and Tarabai Bandekar College of Pharmacy	Ponda	Goa
Pt. Ravishankar Shukla University	Raipur	Chhattisgarh
S.E.T's College of Pharmacy	Dharwad	Karnataka
Shree Warana Vibhag Shikshan Mandai's Tatyasaheb Kore College of Pharmacy	Warananagar	Maharashtra
Sinhgad Technical Education Society's Sinhgad College of Pharmacy	Pune	Maharashtra
Srl Ramakrishna Institute of Paramedical Sciences	Colmbatore	Tamil Nadu
Srinivas College of Pharmacy	Mangalore	Karnataka
SSM College of Pharmacy	Jambat	Tamil Nadu
VELS Institute of Science Technology & Advanced Studies	Chennai	Tamil Nadu
Vivekanand Education Society's College of Pharmacy	Mumbai	Maharashtra

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Cell Culture Laboratory in the HEI



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Industrial grade cell culture laboratory equipped with air/ HEPA filter and full proof air-controlled system





Laminar Air Flow Unit

CO₂ Incubator

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Inverted Microscope

Cooling Centrifuge

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Common Facility Centre in the HEI





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Freeze Dryer



HPLC



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Uv-Visible Spectrophotometer



Brookfield Viscometer



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FTIR

Inverted Microscope

Incubation Facility Training availed to staff and students

CO₂ Extractor





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Research grants

Academic Year 2021-22

Sr. No.	Name of the research project/ endowment	Amount Sanctioned (INR in Lakhs)	Name of the Funding Agency	10/412
1.	Development and characterization of ribavirin-loaded nanoparticles for the treatment of cancer	0.1	Shivaji University Kolhapur	Page 9/113
	Total	0.1		

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BHARATI VIDYAPEETH

COLLEGE OF PHARMACY, KOLHAPUR

: Founder : Dr. Patangrao Kadam M.A., LL.B., Ph.D.

Courses: D.Pharm, B. Pharm, M. Pharm, Ph.D. DTE College Code No. - 6256 (Approved by A.I.C.T.E., P.C.I., New Delhi) hapur & Included in list under Sect:2(F) & 12(B) of UGC Act, 1956 Milated to Shivaji University, Kol B.Pharm. Course reaccredited by NBA, New Delhi

: Principal : Dr. H.N. MORE

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LEAD COLLEGE, SHIVAJI UNIVERSITY, KOLHAPUR, NIRF INDIA RANKING 2021:49 Near Chitranagari, Kolhapur - 416013 (MS) Tel. (0231) 2637286, 2638392, Fax : 2638833

Ref. No. : BV/CPK / 6/4 /2021 - 20 22

Date :01/02/2022

To

The Principals and Project guides. Pharmacy colleges under Lead College Research Sensitization Scheme, Shivaji University, Kolhapur.

Subject: Regarding research grant of maximum Rs. 10,000/- per project for 'Research Promotion Activity 2021-2022' under Lead College Research Sensitization

Sir/Madam

With reference to subject cited above, all the participated colleges in research promotion activity under lead college research sensitization scheme of Shivaji University hereby informed that research projects submitted by your college have been sanctioned from Shivaji University, Kolhapur for the research grant of maximum Rs. 10,000/- to meet the expenditure of recurring nature towards the project work.

The grant should be utilized for the sanctioned project work only. List of sanctioned projects is attached herewith.

Regards,

Principal Lead College

Bharati Vidyapeeth College Of Pharmacy, Kolhapur

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Research proposals submitted to lead college (Year 2021-2022)

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Sr. No.	Name of College	Research Title	Names of Students	Name of Advisor	
1	B. V. C. P. Kolhapur	Development and characterization of anti-diabetic and measteration regulation activity of Herbal formulation	/ Priyanka Mane / Monika Sabale /Mugdha Kambli	Mr. R.J. Jarag Mr. Sudhir Patil	
2	A. B. C. P. Sangli	In-vitro Antiproliferative and apoptotic inducing effect of plant extract on different cell line	tg effect of plant Sahif Bedmutha		
3	Shri santarupa Ghogaon	Development of new quality control method for Gul/ Jaggery	/ Rutuja Dhanawada / Arifa Naikawadi /Kashish Mulla /Tanuja Pawar / Tejal Shankar Veer	Dr. A.V. Belvotagi	
4	Ashokrao Mane Design, Development and in- Pethyadagson Vitro Autioxidant potential of Quercetin nanoparticles Akus		/ Sanjana Jadhav /Pratiksha Jadhav Akash Desai / Ankita Patil	Mrs. P. S. Sankpal	
5	SGMCP, Mahagaon	Design and Characterization of solid self Nano- Emulsifying Druy, delivery system of letrozole for Breusi Cancer	Shashikant Adsule Ashish Phutanc / Samrodhi Kadam / Shruti Mandakar	Dr. R.B. Kumbhar	
6	TKCP, Warnanagar	Development and Characterization of Ribavirin-loaded Nanoparticles for the Treatment of Cancer	/ Bhagyashri Thorat / Priti Barawade Sushant Todkar Shivprasad Patil	Dr. J. I. Disouza	
7	Vasentidevi Kodoli	Microwave assisted green synthesis antimicrobial activity of thiazolidine- 4-one derivatives	/ Namira Nadaf	Miss. Lalita Dahiwade	
<u>*</u>	Sarojani Kothapur	In vitro Evaluation of Antitubercular Activity of Coccinia grandis	/ Akanksha Gourkar / Utkarsha Ghatage / Sakshi Bhandari / Snehal Chavan	Ms. Preeti Patil- Vibhute	
9	Rajara:nbapu Kasegaon	Biological Evolution of some synthesized N- substituted 1,3,4 thiadiazole derivatives by using invitro model	/ Nutan Desai Anand Desai / Ashwarya Desahmukh Adesh Deshmukh	Dr. Sandeep Kane	

Secretary, Lead College Working Committee (Pharmacy)

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Shree Warana Vibhag Shikshan Mandal's

TATYASAHEB KORE COLLEGE OF PHARMACY

Warananagar, Tal: Panhala, Dist: Kolhapur, 416 113 (M.S.) Phone: (02328) 223501, Website: www.tkcpwarana.ac.in, Email: tkcp.pc@unishivaji.ac.in

NIRF RankBand: 2017 & 2018 - 51 to 75; 2021 - 75 to 100

Dr. John Disouza Principal

Hon. Dr. Vinayji V. Kore (Saavkar)

President

Thursday, March 17, 2022

Shivaji University, Sponsored Minor research project sanctioned under SUK lead college research sensitization scheme (Research Promotion Activity 2021-22)

Income & Expenditure Statement

Description: UG Minor Research Project

Advance Amount: Nil Cheque No.: Nil

Sr. No.	Income	Amo	ount	Sr. No.	Expenditure	Amount
1.	Gant Sanctioned by Shivaji University	10000	00	01	Chemicals	10100.00
2.	Actual Received		00	02		
3.	Amount to be receivable		10000			
4.	Additional Expenditure shared by College		100			
	Total		10100		Total =	10100.00

Sushant Phadnis & Co.

Chartered Accountant

PRINCIPAL

TATYASAHEB KORE COLLEGE OF PHARMACY
(DEGREE) WARANANAGAR, TAL. PANHALA.

Signature of the Finance Officer/ Account

IDIN: - 22122830AFGEDT1482

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Academic Year 2020-21

Sr. No.	Name of the research project/ endowment	Amount Sanctioned (INR)	Name of the Funding Agency	
1.	Induction of novice pharma academicians	3.32933	All India Council for Technical Education	Page 13/113
2.	Leveraging academic researchers on developing diagnostic kits, vaccines and drug product for improved therapy management against deadly viruses: Lesson learnt from COVID 19"	0.93	All India Council for Technical Education	
3.	Fostering pedagogy, research administration: Vital domains for effective professional academic career	2.79	All India Council for Technical Education	
	Total	7.04933		

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Annexure -I

SHORT TERM TRAINING PROGRAM

FEED BACK FORM

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1. AICTE File No. & Date of Offer Letter : 34-66/504/FDC/STTP/Policy-1/2019-20

2. Name of the Coordinator : Dr. John I. Disouza

3. Name and Address of the Institution : Shree Warana Vibhag Shikshan Mandal's Tatyasaheb Kore College of Pharmacy,

Warananagar, Tal- Panhala, Dist- Kolhapur,

M.S., 416 113.

4. Title of the Faculty Development Programme: Induction of Novice Pharma Academicians

5. Dates : Phase I:- 22/11/2021 to 27/11/2021

Phase II:- 29/11/2021 to 04/12/2021

Phase III:- 06/12/2021 to 11/12/2021

6. Venue : By online mode at SWVSM's TKCP College

Warananagar

7. Total No. of participants proposed and actually attended

Proposed 40 Attended 146

8. No. and date of the offer letter

Letter No.	Date
34-66/504/FDC/STTP/Policy-1/2019-20	10/08/2020

9. Total amount sanctioned : Rs. 332933/

10. No. and date of Sanction letter:

Letter No. Date Grant Released

34-66/504/FDC/STTP/Policy-1/2019-20 10/08/2020 332933/-

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Name of the Coo	rdinator Dr.	John I. Disouza	Academicians	(1		Pag
Sanction No. and Date	Grant Sanctioned	Details of expenditure incurred Item wise	Amount Rs. (in each head)	No. of Participants	Duration of the Programme (with dates)	
1	6	1. Honorarium for	15000.00	16	2	
5		Coordinator 2. Honorarium to Experts	216000.00	100	Phase I:- 22/11/2021 to 27/11/2021	
34- 66/504/FDC/ST1 P/Policy-1/2019- 20	332933/-	3. Payment to lab attendant engaged during the lab practices	9000.00	146	Phase II:- 29/11/2021 to 04/12/2021	
10/08/2020	3	4. Miscellaneous charges	28533.00		Phase III;- 06/12/2021 to	
	198	Total Grant received	268533.00 332933.00		11/12/2021	
1	1	Total excess amount with interest returned to AICTE (Unspent Amount Rs. 64400 + Interest Rs. 7869)	72269,00	~	>	
	7	Balance to be Receive	d 00.00			
See Miles	Congress of Constant	Rog. No. 20137 PH. Williams 1200-01-11-120-0004 Mar. Suthent Phadams Countant	TATVASAITE MA	John T. Dissuzz PRINGIPAL I III INSTANCIAR, TAL I LIP. PDE 415113, M.	HARMACY MANUALA	

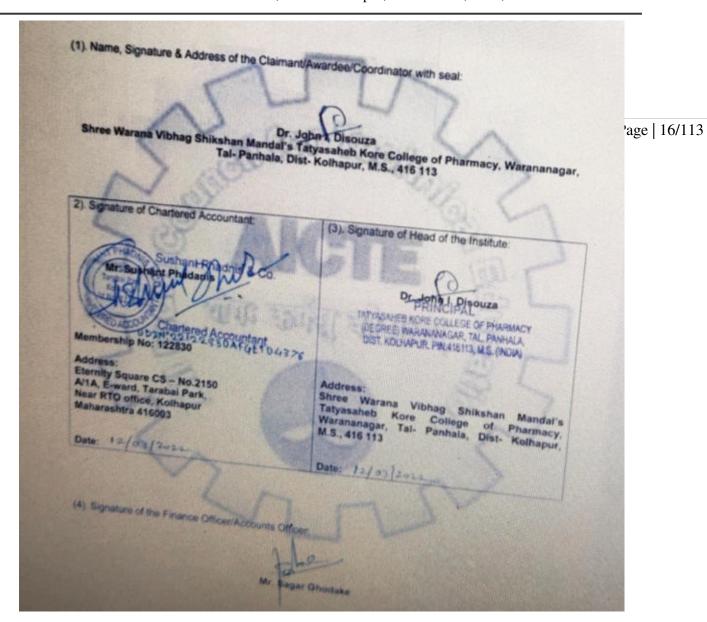
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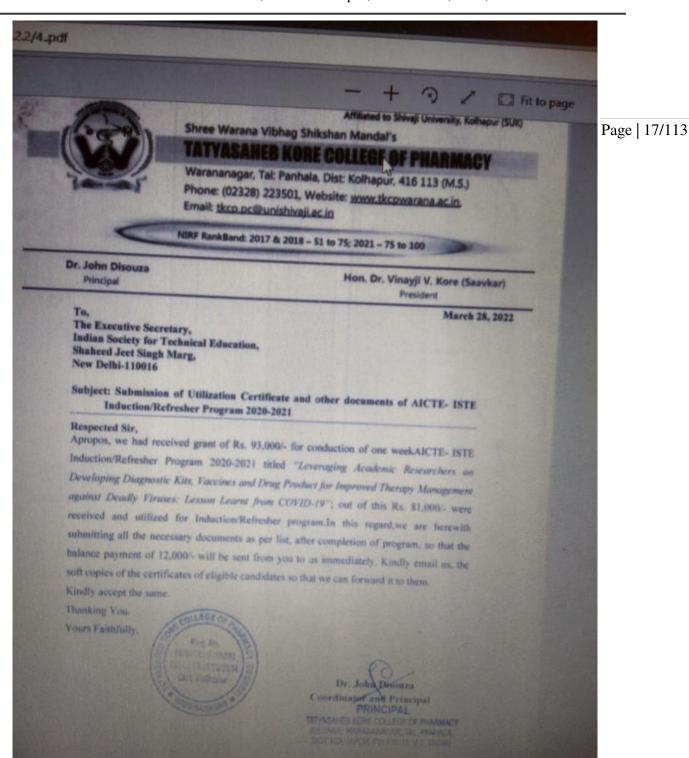
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Warananagar

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	UTILIZTION CERTIFICATE
Name and Address of Institute File No Name of Coordinator	Grant was sanction: AICTE-ISTE Induction/Refresher Program 2021-22 1 Sieve Waruna Vibbag Shikshan Mandal's Tatyasaheb Kore College of Pharmacy, Warananagar Tai: Panhaia Dist: Kolhapur 416113 1 ISTE/AICTE-ISTE Induction-Refresher Program/2021-22 Nov 5, 2021 1 Dr. John I. Disouza
Dates of the Programme Title of the Programme	: 13 th December to 18 th December 2021 : Leveraging Academic Researchers on Developing Diagnostic Kits, Vaccines and Drug Product for Improved Therapy Management against Deadly Viruses: Lesson Learnt from COVID-19
SLNo. ISTE Sanction Order/Letter No. & Date under which grant was sanctioned	Amount
ISTE/AICTE-ISTE Induction-Refresher Program/2021-22 Nov 5, 2021	WARANANAGAR as per letter mentioned in the margin. Rs.0.00/- on account of unspent balance of previous year. Rs. 0.00 on account of other income/receipts, a sum of Rs. 93010 has been utilized for the purpose for which it was sanctioned and the balance of Rs. 0.00 remained unutilized at
Kinds of checks exercised Statement of Income and Expending Receipt and Payment account, bank Hills Venchen/receipts etc. Mr. Sugar Ghodke	the conditions on which the grant-m-aid was satisficed have been duly fulfilled and that I see that the money was actually utilized for the purpose for which it wassanctioned.

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AICTE-ISTE INDUCTION/REFRESHER PROGRAMMES – 2018-19



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ISTE/AICTE-ISTE Orientation/Refresher Programme/2018-19

March 1, 2021.

Dear Sir/Madam,

Sub: AICTE-ISTE Sponsored Induction/Refresher Programs - regarding

Ref.: 1. Our letter dated October 19, 2020.

2. Your consent letter by email to conduct the program.

Sanction is hereby accorded to the institute to conduct three programs in Online Format on the subject titled Fostering Professional Performance during the following dates:

> Phase 1 : 19/03/2021 to 25/03/2021 Phase II : 26/03/2021 to 01/04/2021 Phase III : 02/04/2021 to 08/04/2021

The headwise breakup of each online programme will be:

	Total for each program	Rs.93,000.00
d.	Miscellaneous charge	Rs.10,000.00
c.	Provision for payment to lab attendant engaged during lab practices	Rs.3,000.00
b.	Honorarium to Experts	Rs.75,000.00
a.	Honorarium for Coordinator	Rs.5,000.00

You are requested to take steps to conduct the program subject to the following guidelines (as laid down by the AICTE and ISTE):

- > The program duration will be six days.
- The total budget sanctioned for online program is Rs.93,000/- for one week. Expenses should not exceed the prescribed budget provisions.
- Coordinator may use any available software (Google Meet/WebEx/MS Team/ Go to Webinar etc.) for smooth conduction of online FDP and also he/she may explore any other available software.
- Minimum two sessions on inauguration day after inauguration and minimum two sessions before Valedictory function. Institutions/Coordinator will ensure minimum three sessions for remaining four days and one session from that may be utilized for feedback and assessment). Each Session should be of minimum one & half hours.

SHAHEED JEET SINGH MARG, NEAR KATWARIA SARA!, OPP. SANSKRIT VIDYAPEETH, NEW DELHI – 110 016
Phone: 011-26513542, 26963431; email: <u>istedha@isteonline.org</u>; website: www.isteonline.in

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Warananagar, Tal: Panhala, Dist: Kolhapur, 416 113 (M.S.)
Phone: (02328) 223501, 223526, Fax: 223501; Website: www.tkepwarana.ac.ir
Email: tkcp.pc@unishivaji.ac.in

MIRF 2017 & 2018: 51 to 75 BankBand

Dr. John Disouza Principal

Hon. Dr. Vinayji V. Kore (Saavkar) President

Date: 09/08/2021

AICTE-ISTE Sponsored Induction/Refresher Program on "Fostering Pedagogies, Research, Administration: Vitals Domains of Effective Professional Academic Career"

UTILIZATION CERTIFICATE

Certificate that the grant of Rs. 93000/- (Rupees Ninety Three Thousand only) for each program from AICTE-ISTE to conduct Induction/Refresher Program in online format in three phases titled "Fostering Pedagogies, Research, Administration: Vitals Domains of Effective Professional Academic Career", vide reference no. ISTE/AICTE-ISTE Orientation/Refreshers Programme/2018-19 dated 01/03/2021 for all three phases. The amount sanctioned for the conduction of each program and the utilization for the purpose for which it was sanctioned, in association with the terms and conditions laid down by AICTE-ISTE is mentioned as below.

Sr. No.	Date of Program	Sanctioned Amount	Heiliand Amount
- 1	19/03/2021 to 25/03/2021	93000.00	
2	26/03/2021 to 01/04/2021		87890.00
	02/04/2021 to 08/04/2021	75000.00	84720.00
-		93000.00	86870.00
	Total Amount	279000.00	259480.00

The total amount received from AICTE-ISTE is Rs. 150000.00 (Rs. One Lakh Fifty Thousand Only) out of Rs. 279000.00 (Rs. Two Lakh Seventy Nine Thousand Only) for all three phases. The actual expenditure of all three phases are Rs. 259480.00 (Rs. Two Lakh Fifty Nine Thousand Four Hundred Eighty Only). The balance to be received from AICTE-ISTE is Rs. 109480.00 (Rs. One Lakh Nine Thousand Four Hundred Eighty only)

Signature of Coordinator

Signature of Principal
PRINCIPAL
MAKER KORE COLLEGE OF PHARM

TATYASAHEB KORE COLLEGE OF PHARMACY (DEGREE) WARANANAGAR, TAL. PANNALA, DIST. KOLHAPUR, PINM16113, M.S. (INDIA) Signature of Auditor (Govt. Internal Aud./ Chartered Acc.)

Sushant Phadnis & C

Chartered Accountant

UDIN-2H22830AAAA9K

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Academic Year 2019-20

Sr. No.	Name of the research project/ endowment	Amount Sanctioned (INR)	Name of the Funding Agency	
1.	Development of docetaxel nanoparticles: Effect of metabolism inhibition on its anticancer activity	0.1	Shivaji University - Kolhapur	Page 21/113
2.	Modernization and development of microbiology and cell culture laboratory for advanced academic research	14.885	All India Council for Technical Education	
3.	Supercritical fluid extraction of medicinal plants and screening of their extracts for pharmacological activity using BIOPAC	14.985	All India Council for Technical Education	
	Total	29.97		

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SHIVAJI UNIVERSITY, KOLHAPUR

RESEARCH SENSITIZATION SCHEME FOR COLLEGE STUDENTS

Application for the financial support to research project

(To be submitted to the lead college)

1	Name of the College/Institute	***	Shree Warana Vibhag Shikshan Mandal's Tatyasaheb Kore College of Pharmacy, Warananagar Tal: Panhala, Dist: Kolhapur 416 113 (M. S.), Phone: (O) +912328 223501, (R) +912328 224349
2	Name of the Students		Ms. Rasika Amne (Final Year B. Pharm)
	(Up to 4)	:	2. Ms. Grishma Patil (Final Year B. Pharm)
			3. Mr. Nachiket Banne (Final Year B. Pharm)
3	Title of Project	:	Development of Docetaxel Nanoparticles: Effect of Metabolism Inhibition of its Anticancer Activity
4	Area of Research Project/Subject	:	Pharmaceutical Sciences & Technology
5	Details of the Research Project	:	Annexure – 1
6.	Financial Requirements (Up to rupees 10000/-)	**	11000/-
	Chemicals/ Consumables	**	10,500/-
	Travelling	:	500/-

Name and signature of the project advisor

Dr. A. S. Manjappa

05/07/MS/PHARM 2004/042/11/05/2004 Name and signature of students

1. Ms. Rasika Amne RA mme

2. Ms. Grishma Patil Goi shreet

3. Mr. Nachiket Banne Borne

TATYASAHER KORE COLLEGE OF PHARMACY (DEGREE) WARANANAGAR, TAL PANHALA, DIST, KOLHAPUR, PIN-416113, M.S. (INDIA)

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Shree Warana Vibhag Shikshan Mandal's

TATYASAHEB KORE COLLEGE OF PHARMACY

Warananagar, Tal: Panhala, Dist: Kolhapur, 416 113 (M.S.) Phone: (02328) 223501, 223526, Fax: 223501; Website: www.tkcpwarana.ac.in Email: tkep.pe@unishivaji.ac.in

NIRF 2017 & 2018: 51 to 75 RankBand

Dr. John Disouza Principal

Hou. Dr. Vinayji V. Kore (Saavkar) President

Date:

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MODERNISATION AND REMOVAL OF OBSOLENSCENCE RURAL (MODROB-RURAL): 2019-20 "Supercritical Fluid Extraction of Medicianal Plants and Screening of their Extracts for Pharmacological Activities using BIOPAC"

UTILIZATION CERTIFICATE

Certificate that the grant of Rs. 1498500/- (Rupees Fourteen Lakhs Ninety Eight Thousand Five Hundred Only) under MODERNISATION AND REMOVAL OF OBSOLENSCENCE RURAL (MODROB -RURAL)scheme from AICTE New Delhi has been sanctioned for the project entitled "Supercritical Fluid Extraction of Medicianal Plants and Screening of their Extracts for Pharmacological Activities using BIOPAC", vide order/letter no. F.No. 84-7/RIFD/MODROB/Rural/Policy-1/2019-20 Dated 16 May 2019. The amount sanctioned for the above said project has been utilized, and the utilization for the purpose for which it was sanctioned in association with the terms and conditions laid down by AICTE New Delhi is mentioned as below;

AICTE Sanction Order/Letter No. &	Total Sanctioned Amount (Rs.)		Total Utilized Amount (Rs.)		Unspent Balance
Date under which the amount was sanctioned	Non Recurring (Rs.)	Recurring (Rs.)	Non Recurring (Rs.)	Recurring (Rs.)	
F.No. 84- 7/RIFD/MODROB/Rural/Policy- 1/2019-20 Dated 16 May 2019.	Rs. 1273725/- (Rupees Twelve Lakhs Seventy Three Thousand Seven Hundred Twenty Five Only)	Rs.224775/- (Rupees Two Lakh Twenty Four Thousand Seven Hundred Seventy Five Only)	Rs. 1460000/- (Rupees Fourteen Lakhs SixtyThousand Only)	Rs. 298660/- (Rupees Two Lakhs Ninety Eight Thousand Six Hundred Sixty Only)	Zero Only

The total amount received from AICTE New Delhi is Rs. 1198800/-(Rupees Eleven Lakhs Ninety Eight Thousand Eight Hundred Only) out of sanctioned amount of Rs. 1498500/-(Rupees Fourteen Lakhs Ninety Eight Thousand Five Hundred Only). The actual expenditure incurred is Rs. 1758660/-(Rupees Seventeen Lakhs Fifty Eight Thousand Six Hundred Sixty Only). The balance amount to be received from AICTE is Rs.299700/- (Rs. Two Lakhs Ninety Nine Thousand Seven Hundred Only).

Signature of Coordinator Dr. John I. Disouza (DEGREE) WARANANAGAR, TAL DIST. KOLHAPUR. PIR.416 N.

RINCIPAL

ATYASAHER KORE COLLEGE OF PHARMAC (DEGREE) WARANANAGAR, TAL PANHALA Charter BETACCOURT PRISA 16113, M.S. (NDIA)

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Tatyasaheb Kore College of Pharmacy, Warananagar

Tal: - Panhala, Dist.: - Kolhapur, Maharashtra, India, Pin 416 113

Approved by PCL AICTE New Delhi Recognized by Govt of Maharashtra Affiliated to Shivaji University, Kolhapur (SUK)

Shree Warana Vibhag Shikshan Mandal's

TATYASAHEB KORE COLLEGE OF PHARMACY

Warananagar, Tal: Panhala, Dist: Kolhapur, 416 113 (M.S.)
Phone: (02328) 223501, 223526, Fax: 223501; Website: www.tkcpwarana.ac.in
Email: tkcp.pc@unishivaji.ac.in

NIRF 2017 & 2018: 51 to 75 RankBand

Dr. John Disouza Principal

Hon. Dr. Vinayji V. Kore (Saavkar) President Page | 24/113

Annexure-1

MODERNISATION AND REMOVAL OF OBSOLENSCENCE RURAL (MODROB - RURAL): 2019-20

STATEMENT OF EXPENDITURE

AICTE File No.

F.No. 84-7/RIFD/MODROB/Rural/Policy-1/2019-20 Dated 16

May 2019

Title of Project

Supercritical Fluid Extraction of Medicianal Plants and

Screening of their Extracts for Pharmacological Activities using

BIOPAC

Name of the Coordinator

Dr. John I. Disouza

Tatyasaheb Kore College of Pharmacy, Warananagar

Sanction No. and Date	Total Grant Sanctioned	Details of Expenditure Incurred Item wise	Amount Rs. (In each head)
F.No. 84- 7/RIFD/MODROB/Rural/Policy- 1/2019-20	Rs. 1498500/-	CO ₂ Extractor (Supercritical Fluid Extractor)	1460000/-
Dated 16 May 2019.		Make: Amar Equipments, Mumbai	
		Recurring expenditure includes chemicals, Glasswares & other consumables.	298660/-
		Total Expenditure	1758660/-
		Grant Released	1198800/
		Grant Remaining	299700/
		Funds utilized from Institute	260160/

Signature of Coordinator Dr. John I. Disouza

Signature of Principal

TATYASAHEB KORE COLLEGE OF PHARM (DEGREE) WARANANAGAR, TAL. PANHAL DIST. KOLHAPUR. PIN:416113, M.S. MNU As Per Report of even da

Sushant Phadnis Chartered Accountant

UDIN: 21122830AAAATQ5927

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Tatyasaheb Kore College of Pharmacy, Warananagar

Tal: - Panhala, Dist.: - Kolhapur, Maharashtra, India, Pin 416 113

Dated: 15 May 2019 F.No. \$4-7/R/FD/MODROB/Rural/Policy-1/2019-20 All India Council for Technical Education (A Statutory body under Ministry of HRD, Govt. of India) Nelson Mandela Marg, Vasant Kunj, New Delhi-110070 Website: www.aicte-india.org MODROB-RURAL - Sanction Letter The Drawing and Disbursing Officer, All India Council for Technical Education, Nelson Mandela Marg. Vasant Kunj. New Delhi - 110070 Sub: Release of a sum of Rs.1198800 /- (Rupees Eleven Lakh Ninety Eight Thousand Eight Hundred Only), being the Grant-in-Aid under the scheme Modernization and Removal of Obsolescence Rural (MODROB-Rural) for the year 2019-20 payable during the current financial year 2019-20. rem year 2019-20- reg. With reference to the proposal submitted by the institute, this is to convey that the sanction of the Council for payment of Rs. 1498500/- (Rupees Fourteen Lakh Ninety Eight Thousand Five Hundred Only) as Grant-in-Aid under the Modernization and Removal of Obsolescence Rural (MODROB-Rural) scheme, as per details given below: Director/ Principal/ Registrar Name and address of the SHREE WARANA VIBHAG SHIKSHAN MANDAL'S TATYASAHEB KORE Beneficiary Institution: COLLEGE OF PHARMACY, WARANANAGAR, TATYASAHEB KORE COLLEGE OF PHARMACY, WARANANAGAR, AT POST: WARANANAGAR, TAL: PANHALA, DIST: KOLHAPUR 4116113 MAHARASHATRA STATE, Maharashtra Supercritical Fluid Extraction of medicinal plants and screening of their Title of proposal: extracts for pharmacological activities using BIOPAC. Dr. JOHN DISOUZA Name of Coordinator: 2 years Duration of the project: Total: Non-Recurring (85%): Recurring (15%): Total Grant-In-aid Sanctioned: Rs. 1498500/-Rs.1273725/-Rs.224775/-Recurring (15%): 1st Installment Non-Recurring (85%) Amount to be released during Rs.1018980/-Rs.179820/-Rs. 1198800 /the year 2019-20: Major Head 601.18(a) Gen. (Plan Head) Sanctioned grant-in-aid debitable to: The amount of the Grant shall be drawn by the Drawing and Disbursing Officer, All India Council for Technical Education on the Grant-in-Aid bill and shall be disbursed to and credited to the account of Director/Principal/ Registrar of the Institute through RTGS/PFMS. This Grant-in-Aid is being released in conformity with the terms & conditions as well as norms of the scheme as already communicated, and also being communicated in this letter. The instructions/guidelines to be followed by University/Instituted Release of funds

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Tatyasaheb Kore College of Pharmacy, Warananagar

Tal: - Panhala, Dist.: - Kolhapur, Maharashtra, India, Pin 416 113

Approved by PCI, AICTE New Delhi Recognized by Govt. of Maharashtra

Affiliated to Shivaji University, Kolhapur (SUK)

Page | 26/113

Shree Warana Vibhag Shikshan Mandal's

TATYASAHEB KORE COLLEGE OF PHARMACY

Warananagar, Tal: Panhala, Dist: Kolhapur, 416 113 (M.S.) Phone: (02328) 223501, Website: www.tkcpwarana.ac.in, Email: tkcp.pc@unishivaji.ac.in

NIRF RankBand: 2017 & 2018 - 51 to 75: 2021 - 75 to 100

Dr. John Disouza

Principal

Hon. Dr. Vinayji V. Kore (Saavkar) President

TKCP/1227/22-23

Monday, July 25, 2022

To, Prof. Dileep N. Malkhede The Advisor (RIFD) AICTE, New Delhi

Subject: Submission of Utilization Certificate for Grant Received under MODROB-Rural Scheme 2019-20 (F.No. 84-7/RIFD/MODROB/Rural/Policy-1/2019-20, Dated 16 May 2019)

Respected Sir,

Apropos, the grant of Rs. 1498500/- (Rupees Fourteen Lakhs Ninety Eight Thousand Five Hundred Only) under MODROB-RURAL scheme from AICTE New Delhi has been sanctioned for the project entitled "Supercritical Fluid Extraction of Medicianal Plants and Screening of their Extracts for Pharmacological Activities using BIOPAC", vide order/letter no. F.No. 84-7/RIFD/MODROB/Rural/Policy-1/2019-20 Dated 16 May 2019. The amount sanctioned for the above said project has been utilized for the purpose for which it was sanctioned in association with the terms and conditions laid down by AICTE New Delhi.

We are herewith forwarding the Utilization Certificate (UC) along with other Mandatory Documents for your kind consideration and further reimbursement of the grant balance amount.

We kindly request you to reimburse the grant balance amount.

Thanking You.

Yours Faithfully,

Project Coordinator Dr. John I. Disouza

TATYASAHEB KORE COLLEGE OF PHARMACY

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Academic Year 2018-19

Sr. No.	Name of the research project/ endowment	Amount Sanctioned (INR)	Name of the Funding Agency
1.	Development of multifunctional liposomal drug delivery system targeting multiple myeloma and associated bone damage	0.25	Shivaji University Kolhapur
2.	Development of nanoparticles co- loaded with docetaxel and ritonavir: role of ritonavir in enhancement of docetaxel anti- tumor efficacy	0.175	Shivaji University Kolhapur
3.	Two days workshop for teachers on fostering creativity and innovation in science education	1.03	Vigyan Prasar
4.	Self -assembled mixed micelles composed of drug-polymer conjugates: Improved docetaxel efficacy against cytochrome P-450 3A4 expressing tumors	0.1	Shivaji University Kolhapur
	Total	1.55	

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NAAC "A" Grade

Shree Warana Vibhag Shikshan Mandal's

Tatyasaheb Kore College of Pharmacy, Warananagar

Tal: - Panhala, Dist.: - Kolhapur, Maharashtra, India, Pin 416 113

SHIVAJI UNIVERSITY, KOLHAPUR-416 004 MAHARASHTRA Colleges and University Development Section

PHONE :EPABX-2609000, 2609145

D FAX:0091-231-2691533 & 0091-231-2692333

Website : www.unishivaji.ac.in E-mail: stats@unishiyaji.ac.in शिवाजी विद्यापीठ, कोल्हापूर -४१६००४ महाराष्ट्र (महाविद्यालये व विद्यापीठ विकास विभाग)

दुरध्वनी: (ईपीएबीएक्स) २६०९०००, २६०९१४५

ं फॅक्स: ००९१-२३१-२६९१५३३,२६९२३३३,२६९३२९४

Ref No.: SU/C&U.D.Section/54/857

20 FEB 2020

Page | 28/113

Dr. Manjappa Arehalli S., Tatyasaheb Kore College of Pharmacy Waranngar, Dist: Kolhapur.

Sub.:- Grants Release order under Research Grants to Collge Teachers 2017-2018. Sir/Madam,

With reference to above mentioned subject, I am directed to inform you that, the University authorities have approved your research proposal entitled "Development of Multifuctional Liposomal Drug Delivery System : Targeting Multiple Myeloma and Associated Bone Damage" under Research Grants to Collge Teachers 2017-2018.

- As per project guidelines, total grants of ₹.25000/- has been sanctioned to your research project and out of grant ₹.12500/- sending herewith as a first installment vide cheque bearing number 143791, dt. 12-02-2020.
- · The second installment of remaining grants will be released to you after compliance of the project.
- The total period of the project will be for two years and under no circumstances it will be extended further. The effective date of start of the project should be the date on which grant is issue of day to the Principal Investigator.
- · Submit the workdone report/ project completion report within a prescribed period alongwith the bills duly completed viz. "Accession No.", "Rates are reasonable and paid by me" with P.I signature.

Thanking you,

Encl.: As above.

Deputy Registrar,

Shivaji University, Kolhapur.

Lours faithfully,

The Principal, Tatyasaheb Kore College of Pharmacy Waranngar, Dist: Kolhapur

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Approved by Checked by Authorized by



PHONE :EPABX-2609000, 2609145 □ FAX :0091-231-2691533 & 0091-231-2692333 Website: www.unishivaji.ac.in E-mail: stats@unishivaji.ac.in

SHIVAJI UNIVERSITY, KOLHAPUR-416 004 MAHARASHTRA Colleges and University Development Section

शिवाजी विद्यापीठ, कोल्हापूर -४१६००४ महाराष्ट्र (महाविद्यालये व विद्यापीठ विकास विभाग)

दुरध्वनी: (ईपीएबीएक्स) २६०९०००, २६०९१४५

कंक्सः ००९१-२३१-२६९१५३३,२६९२३३३,२६९३२९४

Ref No.: SU/C&U.D.Section/53/1312

Estd: 1962

NAAC "A" Grade

2 7 MAR 2019

Page | 29/113

To, Dr. Disouza John I., Tatyasaheb Kore College of Pharmacy Waranangar, Dist: Kolhapur.

Sub. :- Grants Release order under Research Grants to Collge Teachers 2017-2018. Sir/Madam,

With reference to above mentioned subject, I am directed to inform you that, the University authorities have approved your research proposal entitled "Development of Nanoparticles Co-Loaded with Docetaxel and Ritonavir" under Research Grants to Collge

- As per project guidelines, total grants of ₹.17500/- has been sanctioned to your research project and out of grant ₹.8750/- sending herewith as a first installment vide cheque bearing number 338534, dt. 24/3/2019.
- The second installment of remaining grants will be released to you after compliance of
- The total period of the project will be for two years and under no circumstances it will be extended further.
- Submit the workdone report/ project completion report within a prescribed period alongwith the bills duly completed viz. "Accession No.", "Rates are reasonable and paid by me" with P.I signature.

Thanking you,

Yours faithfully,

50 huts

Assistant Registrar, Shivaji University, Kolhapur.

Encl.: As above.

Copy to;

The Principal, Tatyasaheb Kore College of Pharmacy Waranangar, Dist: Kolhapur

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Tatyasaheb Kore College of Pharmacy, Warananagar

Tal: - Panhala, Dist.: - Kolhapur, Maharashtra, India, Pin 416 113

विज्ञान प्रसार

-50, इंस्टीट्यूशनल एरिया, सेक्टर-62 नोएडा 201 309 (उ.प्र.)



A-50, Institutional Area, Sector-6; Noida 201 309 (U.P.

VP/982/Science Hands on activity/2018 / 18

04-04-2018

Page | 30/113

Sub: Two days workshop for teachers on Fostering Creativity and Innovation in Science Education.

Dear Sir,

As per your proposal dated 14 March 2018, an amount of Rs. 1,03,000 (One lakh three thou ctioned from Vigyan Prasar for Two days workshop for teachers on fostering creativity and innovation in science education.

The expenditure heads are as mentioned below:-

SN	Particulars	
1,	TA to resource person	Amount in Rs
2.	Accommodation to resource persons	Rs .50,000
	Honorarium to Resource persons outside as well as local	Rs.10,000
	Material cost for the experiments	Rs .8,000
	Miscellaneous	Rs.30,000
	Total	Rs .5,000
		Rs 1.03.000

Terms and Conditions:-

- 1. Warana Science Innovation Activity Center will organised the workshop on 26-27 April 2018 and center will invite participants and will do all logistic arrangements for both participants and resource persons whereas Vigyan Prasar will provide resource persons for different sessions.
- 2. Warana Science Innovation Activity Center is requested to submit SE and UCs in original within 30 days
- 3. Attendance sheet of all Resource Persons and Coordinators in original need to be submitted with SE and
- TA will be paid on actual basis.
- 5. Vigyan Prasar is releasing 80 % of total budget Rs 82,400/- of the budget as an advance and remaining 20% i.e Rs 20,600/-will be paid after receipt of SE and UCs.
- 6. Warana Science Innovation Activity Center will mention specifically in the SE and UC about the
- 7. Since this is a joint activity of Vigyan Prasar and Warana Science Innovation Activity Center, so Vigyan Prasar name and logo will be mentioned in all publicity materials.

Kindly send your acceptance for the same above.

Thanking You

Yours Sincerely,

(Kapil Kr Tripathi) Scientist E

Dr John L D'Souza Principal Coordinator Warana Science Innovation Activity Center Shree Warana Vibhag Shikshan Mandal's Panhala, District Kolhapur - 416113 (MS)

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Tatyasaheb Kore College of Pharmacy, Warananagar

Tal: - Panhala, Dist.: - Kolhapur, Maharashtra, India, Pin 416 113

Project Aided by -Rajiv Gandhi Science & Technology Commission, Govt. of Maharashtra

Shree Warana Vibhag Shikshan Mandal's

Warana Science & Innovation Activity Centre

Warananagar, Taluka: Panhala, Dist: Kolhapur, 416 113 (M.S.), Phone: (02328) 223501, Fax: 223501 Email: waranasiac@gmail.com Project by RGSTC M.S. RGSTC M.S. RGSTC M.S.

Page | 31/113

Receipt payment A/C for two days Science Teachers workshop on 'Fostering Creativity & Innovation in Science Education' on 26^{th} & 27^{th} April, 2018.

Sr. No.	Receipt	Amount in	Sr. No	Payments	Amount in
1	Received fromVigyan Prasar	82,400.00	1	TA for resource persons	55640.00
2	Receivable fromVigyan Prasar	20,600.00	2	Accommodation to resource person	14809.00
3	Registration Fees	10,200.00	3	Honorarium to Resource person outside as well as local	8000.00
4	WSIAC Contribution	48402.00	4	Material cost and expenses	32179.00
		7.	5	Miscellaneous	14517.00
			6	TA & DA for Teachers	22512.00
			7	Food Lunch, Breakfast, Tea	13945.00
	Total	1,61,602.00		Total	1,61,602.00

Utilization Certificate

Certified that (Workshop Expenditure) of Rs. 161602.00 (Rs. One Lack Sixty One Thousand Six Hundred Two Only) (Sanctioned by Vigyan Prasar & Warana Science and Innovation Activity Centre) for the conduction of Science Teachers workshop organized on 26th & 27th April, 2018 has been incurred by observing scrupulously all the rules of Vigyan Prasar & Warana Science and Innovation Activity Centre.

Certified that, from total expenditure Rs. 161602.00 (Rs. One Lack Sixty One Thousand Six Hundred Two Only) amount Rs. 20600 (Rs. Twenty Thousand Six Hundred Only) is receivable from Vigyan Prasar.

Place: Warananagar

Date: Thursday, May 24, 2018

PHADA Sushant Phadnis & 20

Chartered Accountant

Al DUAL CAMERA Shot on realme 3 Principal Coordinator
Warana Scio 49 & Tol-00 Dicco
Activity Centre Warananagar,

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Research, review, book chapters, and books publications

Year	2021-22	2020-21	2019-20	2018-19	2017-18
Number	8	13	9	11	4

Page | 32/113

Papers published in WoS/ Scopus journals with impact factor

Sr. No.	Name of Journal	Impact Factor
1.	Journal of Controlled Release	11.46
2.	Drug Disccovert Todr	8.36
3.	International Journal of Pharmaceutics	6.5
4.	Microchemical Journal	5.304
5.	Chemico-Biological Interactions	5.168
6.	European Journal of Pharmacology	5.195
7.	Journal of Drug Delivery Science and Technology	5.062
8.	AAPS PharmSciTech	4.026
9.	Drug Development and Industrial Pharmacy	3.7
10.	Journal of Pharmaceutical Innovation	2.53
11.	Current Nanoscience	1.53
12.	Brazilian Journal of Pharmaceutical Sciences	1.24
13.	Journal of Research in Pharmacy	0.88

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	TVA 641	O66. 111. 1
Sr.	Title of the paper	Official link
No.		
1.	Drug repurposing: An emerging strategy in	https://doi.org/10.1016/j.ejphar.2022.1750
	alleviating skin cancer	<u>31</u> Pag
2.	D-a-tocopheryl polyethylene glycol succinate:	https://doi.org/10.1016/j.onano.2022.1000
	A review of multifarious applications in	<u>36</u>
	nanomedicines	
3.	Carbohydrate anchored lipid nanoparticles	https://doi.org/10.1016/j.ijpharm.2022.121
		<u>681</u>
4.	Emulgel for improved topical delivery of	https://doi.org/10.1016/j.pharma.2021.05.
	tretinoin: Formulation design and	<u>004</u>
	c0haracterization	
5.	Recent advances in developing polymeric	https://doi.org/10.1016/j.drudis.2022.02.0
	micelles for treating cancer: Breakthroughs and	<u>05</u>
	bottlenecks in their clinical translation	
6.	Development of topical nanogel as a promising	http://dx.doi.org/10.52711/2231-
	delivery of NSAID's tenoxicam using natural	<u>5713.2022.00048</u>
	permeation enhancer essential oil (euckolyptus)	
7.	Inhalation delivery of repurposed drugs for lung	https://doi.org/10.1016/j.jconrel.2021.11.0
	cancer: Approaches, benefits and challenges	<u>15</u>
8.	Design, development, in silico and in vitro	https://doi.org/10.1016/j.jddst.2021.10268
	characterization of docetaxel-loaded	<u>5</u>
	TPGS/Pluronic F 108 mixed micelles for	
	improved cancer treatment	
9.	Carbohydrates-based diagnosis, prophylaxis	https://doi.org/10.1016/j.carpta.2021.1000
	and treatment of infectious diseases: Special	<u>52</u>
	emphasis on COVID-19	
10.	Lyophilization: principle, methods, and	http://www.dap.sciencearchives.org/
	applications	
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11.	Pharmacosomes: An approach to improve	http://dx.doi.org/10.52711/0974-
	biopharmaceutical properties of drugs basic	360x.2021.00779
	considerations in development	
12.	Microneedles: An advanced approach for	http://dx.doi.org/10.5958/2231-
	transdermal delivery of biologics	5691.2021.00010.1
13.	Drug delivery nanocarriers and recent advances	https://doi.org/10.1186/s43046-021-
	ventured to improve therapeutic efficacy	00059-3
	against osteosarcoma: an overview	
14.	Antidiabetic and antihyperlipidemic effects of	https://doi.org/10.1016/j.jtcme.2020.08.00
	Argyreiapierreana and Mateleadenticulata:	<u>1</u>
	Higher activity of the micellar nanoformulation	
	over the crude extract	
15.	Formulation, evaluation and optimization of	https://www.researchgate.net/publication/
	sustain release matrix tablet ofdiltiazem HCL	350707364 Formulation Evaluation and
	by using hydrophilic natural polymers	Optimization of Sustain Release Matri
		x Tablet of Diltiazem HCL by Using
		Hydrophilic Natural Polymers
16.	A review on current nutraceuticals in the	https://www.hortijournal.com/article/view
	management of osteoarthritis	<u>/56/3-1-11</u>
17.	A remarkable in vitro cytotoxic, cell cycle	https://doi.org/10.1007/s13346-020-
	arresting and proapoptotic characteristics of	00752-1
	low-dose mixed micellar simvastatin combined	
	with alendronate sodium	
18.	Design and development of nifedipine	10.36648/0975-9344.12.4.154
	extendedrelease tablet double rotary bi- layered	
	compression machine	
19.	Development of lipid-drug conjugate	10.2174/1573413716666200319130830
	nanoparticles for hydrophilic and lipophilic	

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	drug: a comparative ex vivo gut and Caco-2 cell		
	permeability study		
20.	Podophyllotoxin-polyacrylic acid conjugate	https://doi.org/10.1186/s43046-020-	
	micelles: improved anticancer efficacy against	00053-1	
	multidrug-resistant breast cancer	Page	: 35
21.	Development and validation of RP-HPLC	https://doi.org/10.1016/j.pharma.2020.07.	
	method for simultaneous estimation of	004	
	docetaxel and ritonavir in PLGA nanoparticles		
22.	Nanoparticulate combination of drugs for the	https://doi.org/10.1186/s43046-021-	
	treatment ofosteosarcoma: A review	00059-3	
23.	Complete genome sequence of Lactobacillus	https://doi.org/10.1128/mra.01199-19	
	plantarum strain JDARSH, isolated from sheep		
	milk		
24.	Accelerated stability study of arsenazo iii used	10.13040/IJPSR.0975-8232.12(3).1615-	
	for detection of calcium from biological system	<u>23</u>	
	through uv-spectrophotometer, biochemistry		
	analyzer, pheter, HPLC and HPTLC		
25.	Design, development and assessment of herbal	https://globalresearchonline.net/journalco	
	lipstick from natural pigments	ntents/v61-1/10.pdf	
26.	Ameliorated in vitro anticancer efficacy of	https://doi.org/10.1186/s43094-019-0013-	
	methotrexate d-α-Tocopheryl polyethylene	<u>x</u>	
	glycol 1000 succinate ester against breast		
	cancer cells		
27.	Biological activities of Cassia occidentalis	10.20959/wjpr20199-15430	
	Linn: a systematic review		
28.	Comparative studies of various adsorbent	https://www.researchgate.net/publication/	
	carriers for enhancing dissolution profile of	335676307_COMPARATIVE_STUDIES	
	ketoprofen	OF VARIOUS ADSORBENT CARRI	
		ERS FOR ENHANCING DISSOLUTI	
		ON PROFILE OF KETOPROFEN	

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29.	Studies on solubility enhancement of	https://www.researchgate.net/publication/
	telmisartan by adsorption method	343163641_STUDIES_ON_SOLUBILIT
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	7-epidocetaxel: in vitro and in vivo results	<u>62461</u>
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	acid-cholesterol biomolecular scaffold as	5d76a6b2289bbeb16b818f5d2182/1?pq-
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Drug repurposing: An emerging strategy in alleviating skin cancer

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D-a-tocopheryl polyethylene glycol succinate: A review of multifarious applications in nanomedicines

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Popat S. Kumbhar ^a ♀ ☒, Sameer Nadaf ^b, Arehalli S. Manjappa ^a ♀ ☒, Niraj Kumar Jha ^c, Sunita S. Shinde ^a, Swapnil S. Chopade ^a, Amol S. Shete ^d, John I. Disouza ^a, Unnam Sambamoorthy ^e, Sanapala A. Kumar ^f

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Recent advances in developing polymeric micelles for treating cancer: Breakthroughs and bottlenecks in their clinical translation

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PMID: 35158056 DOI: 10.1016/j.drudis.2022.02.005

Abstract

Polymeric micelles (PMs) have been explored pre-clinically for the delivery of chemotherapeutics to

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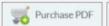
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DOI: 10.52711/2231-5713.2022.00048

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ABSTRACT

Tenoxicam (TNX) is an effective non-steroidal anti-inflammatory drug (NSAIDs) used to treat rheumatoid arthritis. Like other NSAIDs, tenoxicam has the disadvantage of being linked to gastrointestinal side effects. Furthermore, this drug is having weak transdermal penetration, preventing transdermal administration. The goal of this study was to develop a TNX nanogel with a smaller particle size to improve the anti-inflammatory drug's bioavailability and assess its potential in rheumatoid arthritis. The modified emulsification-diffusion method is used to develop nanosized dispersion of TNX using noveon polycarbophil AA-1 as a gelling agent. Moreover, essential oils increase skin penetration by interacting with the stratum corneum (SC). They were found to be successful in increasing skin penetration of both lipophilic and hydrophilic drugs. The rheology, particle size, drug content, % drug release, and in-vitro diffusion study of prepared TNX nanogel were performed. Based on the rheological features of the formulations it was found to be substantial, with the particle size of 125.05nm and zeta potential -8.47mV, drug content of 97.05%, % drug release 97.40% drug diffusion of 97.42%, and pH of 6.2. Tenoxicam nanogel was prepared by using noveon

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Review article

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Inhalation delivery of repurposed drugs for lung cancer: Approaches, benefits and challenges

Popat Kumbhar ^a, Arehalli Manjappa ^a, Rohit Shah ^b, Niraj Kumar Jha ^c, Sachin Kumar Singh ^d, Kamal Dua ^{e f} ♀ ☒ , John Disouza ^a ♀ ☒ , Vandana Patravale ^g

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Design, development, in silico and in vitro characterization of Docetaxel-loaded TPGS/Pluronic F 108 mixed micelles for improved cancer treatment

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Carbohydrates-based diagnosis, prophylaxis and treatment of infectious diseases: Special emphasis on COVID-19



Popat S. Kumbhar^a, Anjali K. Pandya^b, Arehalli S. Manjappa^a, John I. Disouza^a, Vandana B. Patravale b,

ARTICLE INFO

Keywords: COVID-19 SARS-CoV-2 Carbohydrates Diagnosis Nanomedicines

ABSTRACT

COVID-19 pandemic is taking a dangerous turn due to unavailability of approved and effective vaccines and therapy. Currently available diagnostic techniques are time-consuming, expensive, and maybe impacted by the mutations produced in the virus. Therefore, investigation of novel, rapid, and economic diagnosis techniques, prophylactic vaccines and targeted efficacious drug delivery systems as treatment strategy is imperative. Carbo hydrates are essential biomolecules which also act as markers in the realization of immune systems. Moreover, they exhibit antiviral, antimicrobial, and antifungal properties. Carbohydrate based vaccines and therapeutics including stimuli sensitive systems can be developed successfully and used effectively to fight COVID-19. Thus, carbohydrate-based diagnostic, prophylactic and therapeutic alternatives could be promising to defeat COVID-19 propitiously. Morphology of SARS-CoV-2 and its relevance in devising combat strategies has been discussed. Carbohydrate-based approaches for tackling infectious diseases and their importance in the design of various diagnostic and treatment modalities have been reviewed.

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REVIEW **Open Access**

Drug delivery nanocarriers and recent advances ventured to improve therapeutic efficacy against osteosarcoma: an overview



Page | 46/113

Sujit Arun Desai^{1,2*}, Arehalli Manjappa¹ and Preeti Khulbe¹

Abstract

Background: Osteosarcoma (OS) is one of the key cancers affecting the bone tissues, primarily occurred in children and adolescence. Recently, chemotherapy followed by surgery and then post-operative adjuvant chemotherapy is widely used for the treatment of OS. However, the lack of selectivity and sensitivity to tumor cells, the development of multi-drug resistance (MDR), and dangerous side effects have restricted the use of chemotherapeutics.

Main body: There is an unmet need for novel drug delivery strategies for effective treatment and management of OS. Advances in nanotechnology have led to momentous progress in the design of tumor-targeted drug delivery nanocarriers (NCs) as well as functionalized smart NCs to achieve targeting and to treat OS effectively. The present review summarizes the drug delivery challenges in OS, and how organic nanoparticulate approaches are useful in overcoming barriers will be explained. The present review describes the various organic nanoparticulate approaches such as conventional nanocarriers, stimuli-responsive NCs, and ligand-based active targeting strategies tested against OS. The drug conjugates prepared with copolymer and ligand having bone affinity, and advanced promising approaches such as gene therapy, gene-directed enzyme prodrug therapy, and T cell therapy tested against OS along with their reported limitations are also briefed in this review.

Conclusion: The nanoparticulate drugs, drug conjugates, and advanced therapies such as gene therapy, and T cell therapy have promising and potential application in the effective treatment of OS. However, many of the above approaches are still at the preclinical stage, and there is a long transitional period before their clinical application.

Keywords: Osteosarcoma, Nanocarriers, Stimuli-responsive nanocarriers, Active targeting, Gene therapy, T cell

Background

Of the many bone cancers, osteosarcoma (OS) is the most general prime malignant bone tumor accounting for 60% [1]. Both children and adults between 10 and 20 years of age are affected by OS. OS is a complex unbalanced karyotype tumor having some chromosomal aberrations. Although a variety of genetic factors has been

correlated with OS, the specific cause of the OS is not known. Pain is one of the frequent symptoms of OS.

Recently, chemotherapy followed by surgery and then post-operative adjuvant chemotherapy is the widely used conventional strategies for OS treatment. However, the clinical applications of most of the chemotherapeutics have been limited due to lack of selectivity and sensitivity to tumor cells, toxicity towards normal cells, multidrug resistance (MDR), poor pharmacokinetic performance and, etc. [2, 3]. Furthermore, lower blood flow to the bone also acts as a barrier (blood-bone marrow bar-rier) in the delivery of anti-tumor therapeutics to the bone [4]. Therefore, there is an unmet need to develop

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Antidiabetic and antihyperlipidemic effects of Argyreia pierreana and Matelea denticulata: Higher activity of the micellar nanoformulation over the crude extract



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ARTICLEINFO

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HFD-STZ Type 2 diabetes Argyreia pierreana Matelea denticulata Ethanolic extracts

ABSTRACT

Background and aim: Herbal medicine combined with nanotechnology is widely proposed to improve the oral bioavailability, reduce the required dose and side effects, and improve the pharmacological efficacy of extracts. Thus, this study evaluated the in vivo antidiabetic and antihyperlipidemic activities of ethanolic leaf extracts of Argyreia pierreana (AP) and Matelea denticulata (MP) plants in comparison with their micellar nanoformulations.

Materials and methods: The mixed micelles (MMs) loaded with crude extracts (CEs) of AP and MD (AP-MMs and MD-MMs) were prepared using a film dispersion technique. Type 2 diabetes was induced in rats using high-fat diet (HFD) and low-dose (35 mg/kg) streptozotocin (STZ) injection. The pharmacological actions of CEs, AP-MMs and MD-MMs were determined in type 2 diabetic Sprague-Dawley rats. Results: Oral treatments with low-dose AP-MMs and MD-MMs having a mean particle size of 163 ± 10 nm and 145 ± 8 nm respectively, resulted in significantly decreased fasting blood glucose level and increased serum insulin, glucokinase levels, and normalized the elevated levels of hemoglobin A1C and glucose-6-phosphatase. Both extracts significantly decreased serum total cholesterol, triglycerides, and low-density lipoprotein, as well as elevated high-density lipoprotein levels. Additionally, improvements in antioxidant enzymes (superoxide dismutase, catalase, glutathione peroxidase) and malondialdehyde levels were evidenced clearly in tested vital organs (brain, heart, liver).

Conclusion: This is the first report of the antidiabetic and antihyperlipidemic activities of ethanolic leaf extracts of AP and MP plants. Our findings indicate the potential utility of nanotechnology in improving the oral therapeutic efficacy of herbal extracts.

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Formulation, Evaluation and Optimization of Sustain Release Matrix Tablet of Diltiazem HCL by Using Hydrophilic Natural Polymers

Page | 48/113

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ABSTRACT

Article Info Volume 6, Issue 1 Page Number: 16-29 Diltiazem HCl is a Calcium channel blocker which is used as anti-anginal and Class IV anti-arrhythmic drug. It is a drug of choice for stable and unstable angina pectoris, myocardial infarction, coronary artery spasm, cardiac arrhythmia, PSVT and hypertension. In this study, sustained release matrix tablets of Diltiazem HCl were prepared by wet granulation method. The formulation of each Diltiazem HCl sustained release matrix tablets is composed of two selected polymers i.e. chitosan and xanthan gum in alone or in combination. The other excipients used were lactose monohydrate for its diluent property, PVP K-30 as a binder and magnesium stearate and talc for lubrication. The weight of tablet was adjusted to 200 mg and each tablet contained 90 mg Diltiazem HCl. Total 9 batches (F1-F9) were prepared.Batch F1, F2 and F3 containing a single polymer i.e. xanthan gum in concentration of 15, 20 and 25% of total weight of the tablet. Batch F4, F5 and F6 containing a single polymer i.e. chitosan in concentration of 20, 30 and 40% of total weight of the tablet Batch F7, F8 and F9 containing combination of both polymers i.e.

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A review on current nutraceuticals in the management of osteoarthritis

Sharad Kamble, Amol Patil, Sunita Shinde and Hrithik Ankush

Abstract

Osteoarthritis (OA) is a progressive degenerative joint syndrome that has a major impact on joint function and quality of life. Nutraceuticals and nutritional supplement derived from herbs have long been used in traditional remedy and there is considerable evidence that nutraceuticals may play an important role in irritation and joint demolition in OA. We review the biological effects of some medicinal fruits and herbs like pomegranate, green tea, cat's claw, devil's claw, ginger, Indian olibaum, turmeric and ananas. So in an attempt to understand the essential molecular targets involved in irritation and the joint destruction process and to summarize their toxicities and efficacy for OA management. So far there is insufficient reliable evidence on the effectiveness of ginger, turmeric and ananas. Pomegranate and green tea only have preclinical evidence of efficacy due to the bee deficient in of clinical data. In vivo and clinical studies are required to understand their targets and efficacy in OA. There is strong clinical evidence of the efficacy of devil's claw in relieving pain. However, high-

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> Drug Deliv Transl Res. 2020 Aug;10(4):1122-1135. doi: 10.1007/s13346-020-00752-1.

A remarkable in vitro cytotoxic, cell cycle arresting and proapoptotic characteristics of low-dose mixed micellar simvastatin combined with alendronate sodium

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PMID: 32221848 DOI: 10.1007/s13346-020-00752-1

Abstract

The objective of the present study was to screen the effect of increased simvastatin (SVS) solubility, through mixed micelles as a model approach on in vitro entirencer efficacy in combination with

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Design and Development of Nifedipine Extended Release Tablet Double Rotary Bi- Layered Compression Machine

Abstract

The aim of the present work was to Design and development of Nifedipine extended release tablet 90 mg by double rotary bi layered compression machine that can provide continuous drug release for period of 24 hours. The granules prepared using polymers such as polyethylene glycol 4000, HPMC and cellulose acetate etc. the osmotic pump mechanism was used, after that mechanical drilling machine was used for drilling with respect in size. Prior to compression, the prepared granules were evaluated for flow and compression characteristics. The principle shows two compartments was present such as the drug layer and push layer, after some time push layer goes contact with aqueous medium then swelling of push layer and suspend drug particle and flow through the delivery orifice. Prepared Nifedipine extended release tablet was evaluated for in vitro drug release study. The prepared Nifedipine extended release tablet 90 mg showed good mechanical properties (hardness and friability) as well as good in vitro dissolution profile showing the release of constant drug for 24 hours. The increase in binder has retarding effect leading to the decrease in the dissolution. The Change in granulation time with respect to increase in parameters

Keywords: Nifedine; Extended release tablet; Infrared Spectroscopy; Push pull layer

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Development of Lipid-Drug Conjugate Nanoparticles for Hydrophilic and Lipophilic Drug: A Comparative Ex vivo Gut and Caco-2 Cell Permeability Study



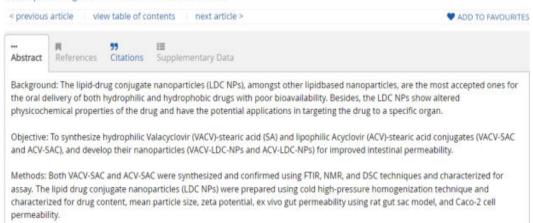
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Authors: Kumbhar, Popat S.; Manjappa, Arehalli Sidramappa; Shete, Abhijeet Dilip; Disouza, John Intru

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Journal of the Egyptian National Cancer Institute

RESEARCH

Podophyllotoxin-polyacrylic acid conjugate micelles: improved anticancer efficacy against multidrug-resistant breast cancer



Page | 53/113

Popat S. Kumbhar, Asmita M. Sakate, Onkar B. Patil, Arehalli S. Manjappa and John I. Disouza

Background: Podophyllotoxin (PPT) is a naturally occurring compound obtained from the roots of Podophyllum species, indicated for a variety of malignant tumors such as breast, lung, and liver tumors. This toxic polyphenol (PPT) exhibited significant activity against P-glycoprotein (P-gp) mediated multidrug-resistant (MDB) cancer cells. thowever, extremely poor water solubility, a narrow therapeutic window, and high toxicity have greatly restricted the clinical uses of PPT. Therefore, the present research was aimed to synthesize the water-soluble ester prodrug of PPT with polyacrylic acid (PAAI, a water-soluble polymer by Steglich esterification reaction, and to screen it for assay, solubility, in vitro hemolysis, in vitro release, and in vitro anticancer activity.

Assults: The Fourier transform infrared (FTIR) and nuclear magnetic resonance (NMR) spectroscopy results revealed the successful synthesis of podophyllotoxin-polyacrylic acid conjugate (FPC). The assay and saturation solubility of the prodrug is found to be 64.01 ± 4.5% and 1.39 ± 0.05 mg/ml. (FPT equivalent) respectively. The PPC showed CMC (critical micelle concentration) of 0.430 mg/ml. In distilled water at room temperature. The PPC micelles showed a mean particle size of 2.15 ± 1.1 nm with polydispersity index (PDTI of 0.199 ± 0.006. Further, the transmission electron microscope (TEM) results confirmed the self-assembling character of PPC into micelles. The PPC caused significantly (ess hemolysis (1.8.6 ± 2.9%) than plain PPT solution. Also, it demonstrated significantly (p < 0.01) includes a within a contraction between the self-assembling characters. 0.01) higher in vitro cytotoxicity against both sensitive as well as resistance human breast cancer cells (MCF-7 and

Conclusion: The obtained study results clearly revealed the notable in vitro anticancer activity of FPT followin esterification with PAA. However, further in vivo studies are needed to ascertain its efficacy against a variety of

Keywords: Padophyllotoxin prodrug, PAA, Hemolysis, In vitro release, Cytotoxicity

Chemotherapy is the most preferred among the avail-able treatment strategies and has been proven to be

effective in clinics. But, the multidrug resistance (MDR) Cancer or malignancy is a heterogeneous disease characterized by abnormal cell mitosis, and is a serious health
concern around the world. Cancer predominance and
mortality are expanding year by year and creating a
heavier burden globally [1]. division [3, 4]. In earlier research papers, it is reported that PPT is capable to kill effectively the MDR (P-gp me-diated) cancer cells and therefore used to treat a variety of MDR tumors efficiently [5-8]. However, the clinical applications of PPT are significantly restricted due to its



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Original article

Development and validation of RP-HPLC method for simultaneous estimation of docetaxel and ritonavir in PLGA nanoparticles Développement et validation de la méthode RP-HPLC pour la détermination simultanée du docétaxel et du ritonavir dans des nanoparticules polymériques

P.S. Kumbhar ^a, Sonali K. Diwate ^{a b}, U.G. Mali ^a, T.U. Shinde ^a, J.I. Disouza ^a Q 🖾,
A.S. Manjappa ^a Q 🖾

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Academic year 2019-20

NANOPARTICULATE COMBINATION OF DRUGS FOR THE TREATMENT OF OSTEOSARCOMA: A REVIEW

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SGVU JOURNAL OF PHARMACEUTICAL RESEARCH & EDUCATION, VOLUME 5 ISSUE 1, 2020

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ABSTRACT

Malignancy speaks to a gathering of heterogeneous ailments described by uncontrolled development and spread of abnormal cells, at last leading to death. Nanomedicine assumes a huge job in the advancement of nanodrugs, nanodevices, drug delivery systems as well as nanocarriers. A portion of the significant issues in the treatment of cancer are multidrug resistance (MDR), restricted helpful window and undesired symptoms of accessible anticancer drug and the constraints of anticancer drugs. A few nano systems being used for recognition, determination and treatment, for example, theranostic bearers, liposomes, carbon nanotubes, quantum spots, polymeric micelles, dendrimers and metallic nanoparticles. Nonetheless, non-biodegradable nanoparticles cause high tissue aggregation and prompts harmfulness. MDR is viewed as a significant obstruction to disease treatment because of metastatic tumors that create protection from chemotherapy. MDR adds to the disappointment of chemotherapies in different diseases, including bosom, ovarian, lung,

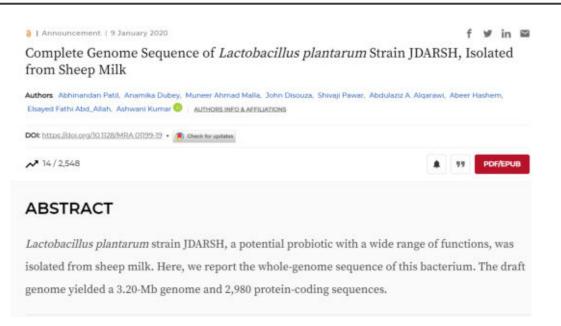
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ACCELERATED STABILITY STUDY OF ARSENAZO III USED FOR DETECTION OF CALCIUM FROM BIOLOGICAL SYSTEM THROUGH UV-SPECTROPHOTOMETER, BIOCHEMISTRY ANALYZER, PH METER, HPLC AND HPTLC

J. R. Kamble, A. S. Sherikar and J. I. Disouza

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Keywords:

In-vitro, linearity, Arsenazo III, Biochemistry analyzer, Shelf life and retention time

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ABSTRACT: Objective: Stability of In-vitro Diagnostics (IVDs) reagents was performed to check its quality standards, performance, and linearity. IVDs like Calcium reagent containing Arsenazo III were studied by Accelerated stability studies by considering temperature, pH, and light parameters. Materials and Methods: Stability data were obtained by using different instruments like UV spectrophotometer, Biochemistry analyzer, pH meter, HPLC, and HPTLC. This gives information about the degradation that occurred during storage, transportation, etc. Results: Calcium reagent containing Arsenazo III degrade 90.81% when placed at 42 °C by UV spectrophotometer analysis. The stability conditions' effect on actual serum concentration was measured by taking reagent performance on a biochemistry analyzer. The linearity of reagents decreases at 42 °C and at normal temperature, linearity does not change. HPLC spectra gave degradation of reagent, which was analyzed by its retention time, peak height, and % area. Arsenazo III produces 91.25% remains undecomposed in 3 months when exposed to light. The shelf life of the calcium reagent was found to be 85.36. HPTLC spectra gave degradation of Arsenazo III, which was analyzed by its retention time, peak height. The reagent, during its stability studies, shows a slight change in its pH. Conclusion: From HPLC and HPTLC analysis, it is confirmed that the degradation occurred in Arsenazo III after exposed to an rated etability etudy

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Design, Development and Assessment of Herbal Lipstick from Natural Pigments

*Mahanthesh M.C., Manjappa A.S., Shinde M.V., Sherikar A.S., Disouza J.I., Namrata B.U, Kranti K.R., Ajija W.C.
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The natural pigment or colorant in the cosmetics, are inconceivable in demand since, historical time till day. The colors that add to the shade of the lipstick, are unsafe to people on utilization can bring about sensitivity, sickness, dermatitis, and drying of the lips. The usage of natural dyes and pigments, increase more importance in food and textile industries because of their non toxic and eco friendly characteristics. Coloring pigments is obtained from petals of Rosa rubiginosa, flower of Bougainvillea spectabilis. Beta vulgaris (Beetroot) and flower of Crocus sativus.(F1 to F4) and were evaluated on the parameters such as melting point, breaking point, force of application, surface anomalies, aging stability, solubility, pH, skin irritation and perfume stability etc... The results are shown in tab.4 and prepared lipstick in fig.2 F1 to F4. The prepared lipstick formulations F1, F2 and F4 showed ideal properties like shining, spreading and smoothness of lips after application. Further studies through a detailed clinical trial may be suggested to ensure safety of these formulations. Hence from present investigation it was concluded that, formulated herbal lipstick having minimal and no side effects and thus showing maximum local effect on lips.

Keywords: Herbal cosmetics, lipstick, natural pigments, formulation evaluation

INTRODUCTION

ith the beginning of the civilization, Herbal cosmetic also known as "natural cosmetics"., peoples (men and women) had the magnetic dip towards impressing others with their looks was reported¹and there area number of wide range of herbal cosmetics products to satisfy your beauty regime, is very safe for the skin. The human beings have been using herbs for different purpose like food, medicine, beatifying with the advancement of science & technology was studied².

The phenomenon of herbals, nowadays becoming a full fledged, encircling both health and beauty care. The lips perhaps constitute the most sensitive part of our body and it is also very close to the nose and mouth. The great demand in both developing and developed countries⁴, the demand of herbal medicines is increasing rapidly due to their lack of side effects was reported⁵. Page | 58/113

In another study, Natural pigment or color in biological system is one, that is synthesized and accumulated in, or extracted from living cells and natural dyes may be defined as chemicals which are obtained from vegetable and animal sources without chemical processing. The applied colour should be fast to sunlight, water washing and to action of mild acid and alkali. The different natural colorants are obtained from following categories is shown in the table 16-9.

The taking into consideration the importance of natural products, the present work was aimed at formulating and evaluating lipsticks containing only natural ingredients. The ingredients included in the study, extracts of the

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Future Journal of Pharmaceutical Sciences

RESEARCH Open Access

Ameliorated in vitro anticancer efficacy of methotrexate D-α-Tocopheryl polyethylene glycol 1000 succinate ester against breast cancer cells



Page | 59/113

Arehalli S. Manjappa", Popat S. Kumbhar, Rohini Kasabe, Sonali K. Diwate and John I. Disouza

Abstract

Background: Methotrexate (MTX), a folate anti-metabolite, has been used widely in the treatment of plenty of malignancies. However, the clinical use is limited because of its poor water solubility (BCS class II drug), nonspecific distribution, drug resistance, short circulation half-life, and toxicity. The objective of the present research was to synthesize the ester prodrug of MTX with p-q-Tocopheryl polyethylene glycol 1000 succinate (TPGS) and characterize for in vitro anticancer efficacy.

Results: The FTIR and NMR results revealed the successful synthesis of the prodrug. The assay and saturation solubility of the prodrug is found to be $23 \pm 2.5\%$ and 6.7 ± 1.3 mg/mL (MTX equivalent) respectively. The CMC of the prodrug in distilled water at room temperature is found to be 36.9 ± 2.6 µg/mL. The prepared prodrug micelles showed a mean particle size of 166 ± 10 nm (PDI, 0.325 ± 0.09). Further, the TEM results confirmed the self-assembling character of the prodrug into micelles with a nearly spherical shape. The prodrug caused the significantly (p < 0.01) less hemolysis ($16.8 \pm 1.5\%$) when compared to plain MTX solution and significantly higher (p < 0.01) in vitro cytotoxicity, cell cycle arresting, and apoptosis against human breast cancer cells (MCF-7 and MDA-MB-231).

Conclusion: Our study results revealed the remarkable in vitro anticancer activity of MTX following its esterification with TPGS. However, further, in vivo studies are needed to prove its efficacy against different cancers.

Keywords: Methotrexate prodrug, TPGS, Cytotoxicity, Cell cycle analysis and apoptosis

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Review Article

ISSN 2277-7105

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BIOLOGICAL ACTIVITIES OF CASSIA OCCIDENTALIS LINN: A SYSTEMATIC REVIEW

Mahanthesh M. C. 4, Manjappa A. S.2, Sherikar A. S.3, Disouza J. L.4 and Shinde M. V.5

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ABSTRACT

Cassia occidentalis L. (Kasamardah), Negro coffee, Family leguminose, is an erect, perennial plant and have traditional practice, as well as wide Phytochemicals and having diverse biological activities, known to possess antiallergic, antibacterial, antidote for poison, blood purifier, antifungal, antidiabetic, anti-inflammatory, antimutagenic, psoriasis, melanoblast cell line leprosy and hepatoprotective activity. Chemicals including achrosin, aloeemodin, cassia occidentanol I, cassia occidentanol II, emodin, anthraquinones, anthrones, apigenin, aurantiobtusin, campesterol, cassiollin, chryso-obtusin, chrysophanic acid, chrysophin, chrysophanol, chrysopriol. The presented review

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Research Article EJPMR

COMPARATIVE STUDIES OF VARIOUS ADSORBENT CARRIERS FOR ENHANCING DISSOLUTION PROFILE OF KETOPROFEN

Page | 61/113

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ABSTRACT

In recent times, a large number of studies concerning the improvement of the dissolution rate of poorly watersoluble drugs is focused on the application of various porous materials as the drug carriers. These materials have attracted the attention of researchers owing to their outstanding properties such as large surface area, high pore volumes, microporosity and possibility of surface functionalization. In the present study, the biopharmaceutical performance of porous adsorbents as a carrier for the poorly water soluble drug Ketoprofen was investigated. Ketoprofen loaded different adsorbents with high specific surface area were used like Neusilin, Sylysia, Fujicalin and Aerosil, and it was done by solvent evaporation method. It was noticed that porous structure is responsible for an amorphous state of the drug and thus the improvement of its dissolution rate. From this research work it can be concluded that although the porous carrier particles help to enhance dissolution rate, including stability studies.

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RESEARCH PAPERS

STUDIES ON SOLUBILITY ENHANCEMENT OF TELMISARTAN BY ADSORPTION METHOD

Page | 62/113

By

SHARAD KAMBLE *

SUNITA SHINDE **

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Date Received: 15/07/2019 Date Revised: 29/09/2019

Date Accepted: 26/11/2019

ARSTRACT

In this study precipitated form of Telmisartan was prepared by using Solvent evaporation technique and Spray drying technique. For this the author used adsorbents like Sylysia, Neusilin, and diluents like Lactose Monohydrate and Avice I. All these prepared batch es were screened by performing Drug content, saturation solubility and dissolution study of the prepared batches. This obtained optimized batch, was further characterized by using the dissolution test, Differential Scanning Calorimetry (DSC), X-Ray powder Diffractometer (XRD), and USP Dissolution test apparatus.

Keywords: Telmisartan, Solvent Evaporation Method, Spray Drying Technique, Adsorbent Carrier like Neusllin, and Sylysia, Diluents Like Lactose Monohydrate and Avicel, Dissolution Rate.

INTRODUCTION

Low aqueous solubility is the major problem encountered with formulation development of new chemical entities as well as for the generic development. More than 40% of NCEs (new chemical entities) developed in pharmaceutical industries are practically insoluble in water. Solubility is a major challenge for formulation scientist. Solubility occurs under dynamic equilibrium, which means

drug is considered highly soluble, when the highest dose Saturated solutions of ionic compounds of relatively low strength is soluble in 250mL or less of aqueous media over the pH range of 1 to 7.5. Solubility is sometimes described by solubility constants. This is a case of equilibrium process. It describes the balance between dissolved ions from salt and undissolved salt. Similar to other equilibrium constants, temperature affects the numerical value of solubility

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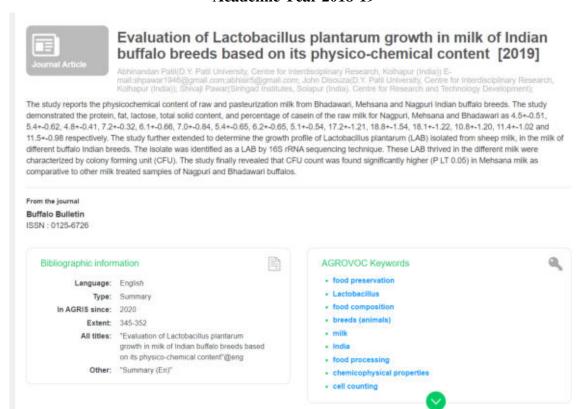
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Original Article

Page | 65/113

DESIGN AND EVALUATION OF GUANFACINE EXTENDED RELEASE FORMULATION *SANJEEVANI DESAI1, DURGACHARAN BHAGWAT2, SUNITA SHINDE1, JOHN DISOUZA1

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Received: 31 Oct 2018, Revised and Accepted: 30 Jan 2019

Objective: The present study was aimed to develop of the Guardacine Hydrochloride Extended-release tablets for the treatment of Attention Deficit Hyperactivity Disorder (ADHD). The dorage regimes of Guardacine Hydrochloride is 4 mg at every 6 h. The concentration of Guardacine in plasma is fluctuating. Hence, to control the plasma fluctuation and to avoid toxicity problem, Guardacine Hydrochloride was chosen as a drug with an aim to develop an extended release system for 20 to 24 h.

Methods. The design of the system was based on the use of pH-dependent polymer (Hydroxypropyl Methyl Celluloss), pH-independent polymer (Eudragit L 100-55), along with microenvironment medifiers such as organic acid (Pamaric acid) were used in the formulation. Drug-exciptent compatibility was studied by FTIR Before compression, the grazules were evaluated for precompression parameters such as halk density, tapped density, an angle of repore, compressibility index and Hausner's ratio. After compression, evaluation tests of tablets such as general appearance, hardness, thickness, weight variety fitability, content uniformity, in vitro release studies and stability studies were performed.

Results: Out of 9 formulations, the drug release was found to be within the isnovator formulation F9. The stability study of formulation F9 revealed there was no significant change in physical and chemical properties of drug stored at 40 °C/75 % RH, 30 °C/65 % RH, 25 °C/60 % RH for 2 mo.

Couclusion: Optimized formulation batch F9 showed highest F2 value which indicates similarity with innovator product. The study indicates that Guanfacine Hydrochioride Extended-release tables was

Keywords: Extended-release, Solubility, pH-dependent polymer, In vitro study

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(Research Article)



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SIMVASTATIN LOADED NANO MIXED MICELLES: AN APPROACH TO TREAT HORMONE DEPENDENT CARCINOMAS

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Keywords:

Simvastatin, Single copolymer micelles, Mixed micelles, In-vitro hemolysis, MTT assay

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ABSTRACT: The main objective of the present research was to develop mixed micelles using two biocompatible copolymers, D-a-tocopheryl polyethyleneglycol 1000 succinate (TPGS) and poloxamer 188 (P188) to improve the aqueous solubility and targeting efficacy of Simvastatin (SMV) against a variety of hormone-dependent cancers. A solvent evaporation technique prepared the plain/single copolymer micelles (SCMs) and mixed micelles (MMs). The prepared SCMs and MMs were characterized for critical micelle concentration (CMC), SMV content, particle size by dynamic light scattering (DLS), surface morphology by transmission electron microscopy (TEM), in-vitro SMV release and hemolysis. The SCMs and MMs showed mean particle size of 98 ± 5 nm and 129 \pm 6 nm, respectively. SCMs showed SMV loading of 79.7 \pm 5.6% while MMs exhibited improved SMV loading of 94.5 ± 6.5. The developed MMs system showed significantly lower CMC (3.5 fold less) than SCMs revealing their higher in-vivo stability. Moreover, SCMs and MMs exhibited zero order release profile, lower hemolytic behavior (<5% of hemolysis), when compared to plain SMV solution. The in-vitro cytotoxicity assay was conducted on MCF-7 (human breast cancer) cell line. Cytotoxicity studies revealed significantly improved antitumor activity of MMs when compared to SCMs and plain SMV after both incubation time points (24 and 48 h). In conclusion, the developed

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Polymeric Mixed Micelles: Improving the Anticancer Efficacy of Single-Copolymer Micelles

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Arehalli S. Manjappa,^{a,*} Popat S. Kumbhar,^a Ajit B. Patil,^a John I. Disouza,^a & Vandana B. Patravale^b

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ABSTRACT: Mixed micelles self-assembled from two or more dissimilar block copolymers provide a direct and convenient approach to improved drug delivery. The present review is focused on mixed micelles (prepared from block copolymers only) for various drug delivery applications along with their merits over single-copolymer micelles. Presented are the physi-

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Small Ruminant Research

Volume 170, January 2019, Pages 19-25



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Shelf life stability of encapsulated lactic acid bacteria isolated from sheep milk thrived in different milk as natural media

Abhinandan Patil a, John Disouza a, Shivaji Pawar a b 🙎 🖾

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Future Journal of Pharmaceutical Sciences

RESEARCH

Open Access

In vitro free radical scavenging and antidiabetic activity of aqueous and ethanolic leaf extracts: a comparative evaluation of Argyreia pierreana and Matelea denticulata



Page | 69/113

Venkatalah Gudise^{1*} G. Bimalendu Chowdhury² and Arehalli S. Manjappa³

Abstract

Background: Oxidation is believed to play a vital role in the pathogenesis of diabetes melitus by lipid peroxidation; DNA and protein damage leads to the development of vascular complications like coronary heart disease, stroke, neuropathy, retinopathy, and nephropathy. The herbal preparations are complementary and alternative medicines to allopathic drugs which are believed to cause adverse events. Therefore, the current study was aimed to identify the novel plants, which belong to the genera Arganeia (Aggreia pierreana (API)) and Matelea (Matelea denticulata (MDI)), and assess the aqueous and ethanolic leaf extracts for in vitro antioxidant and antidiabetic potential by DPPH, OH; superoxide, and glucose uptake and gene expression (GLUT-4 and PPARy) studies using the L-6 cell line respectively.

Results: The preliminary scrutiny revealed the presence of polyphenols, flavonoids, terpenoids, steroids, tunnins, alkaloids, and glycosides. The total phenolic and flavonoid contents of ethanolic extracts were found higher than those of aqueous extracts. The ethanolic extracts exhibited the superior antioxidant capacity when compared with aqueous extracts. However, the ethanolic extract of MD was shown superlative glucose uptake activity (72.54%) over control (0.037%) and GLUT-4 and PPARy gene expressions (1.17 and 1.20) in term of folds respectively over cell control (1.00).

Conclusion: The ethanolic leaf extracts of both plants showed significant in vitro antioxidant and antidiabetic activities compare to aqueous extracts. The Matelea denticulata ethanolic leaf extract exhibited superior activity. This superior activity might be due to their higher phenolic and flavonoid content. However, further approaches are needed to define these activities

Keywords: Argyreia pierreana, Matelea denticulata, Antiradical activity, Antidiabetic activity, GLUT-4 and PPARy

Background

Traditional herbal medicines have shaped the basis of human health care, and further research will improve global health [1, 2]. Presently, about 80% of the world population (according to WHO) uses herbal drugs for some aspects of primary health care. Globally, the use of medicinal plants predates antibiotics and other

contemporary drugs [3, 4]. In addition, many culinary herbs and spices were tested for their biological activ ities in Alzheimer's disease management and other chronic diseases [5, 6].

The natural antioxidant defence mechanism, in all human and other aerobic organisms, prevents the oxidative damage. Since the natural antioxidant defence mechanism is inadequate on its own, the nutritional consumption of antioxidants is suggested [7, 8]. Currently, synthetic antioxidants are replaced by natural antioxidants as the former are reported to have carcinogenic properties. Plants are the



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spondence venkephumaDdgmat.com timent of Pharmacology, SSJ College of Pharmacy, Vattinagulipally, et, Hyderabad, Telanganis State 10001f, India of author information is available at the end of the article

Drug Dev Ind Pharm. 2019 Mar;45(3):474-484. doi: 10.1080/03639045.2018.1562461.
Epub 2019 Jan 1.

Unravelling the anticancer efficacy of 10-oxo-7epidocetaxel: in vitro and in vivo results

Page | 70/113

Arehalli S Manjappa 1 2 3, Rayasa S Ramachandra Murthy 2

Affiliations - collapse

Affiliations

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- 3 c Advanced Centre for Treatment Research and Education in Cancer, Tata Memorial Centre, Kharghar, India.

PMID: 30599774 DOI: 10.1080/03639045.2018.1562461

Abstract

Purpose: To prepare 7-epidocetaxel (7ED) and 10-oxo-7-epidocetaxel (10-O-7ED) formulations as like marketed Taxotere® (TXT) injection and to screen them for in vitro and in vivo anticancer efficacy including their in vivo toxicity behavior.

Methods: The 7ED and 10-O-7ED formulations were screened for in vitro anti-proliferative, anti-metastatic and cell cycle arresting behaviors. Further, in vivo acute toxicity of TXT injection containing 10% of 7ED and 10-O-7ED separately and the therapeutic study of 10-O-7ED alone were studied in B16F10 experimental metastasis mouse model.

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Research Article

Design and development of aliphatic amino acid-cholesterol biomolecular scaffold as anticancer conjugates.

Mayuresh Shinde1*, Shitalkumar Patil2, Manish Bhatia3, Dhanashri Patil4, Sanjay Mishra4

¹Department of Pharmaceutical Chemistry, Tatyasaheb Kore College of Pharmacy, Warananagar, 416113, Maharashtra, India.

Ashokrao Mane College of Pharmacy, Peth Vadgaon, 416112, Maharashtra, India.

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Received 22 March 2019; received in revised form 18 June 2019; accepted 20 June 2019

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ARSTRACT

We have developed lipoprotein macromolecular motif to target multiple type of cancerous cells. These scaffold moieties linked with anticancer agents for targeting release at specific site. Biomolecular network increases cellular penetration, specificity and efficacy. Molecular motifs containing these agents are readily degradable by enzymatic cleavage. Structural functionalities of these modified molecules generate response against cancerous cells. Lipids and protein conjugates improve drug delivery towards target tissues. Bioactive lipoprotein exerts inhibitory effect for progressing tumor tissues. Lipid-protein bioconjugates interact with tumor tissue proteins selectively for reducing toxicity of antitumor agents. Complexation of cholesterol with bioactive aliphatic amino acid yields complex scaffold possessing anticancer activity. Reaction was conducted using dicyclohexyl carbodiimide (DCC) and 4-dimethylamino pyridine (DMAP) in pyridine solvent. Developed conjugates were characterized by using TLC, IR, NMR and HRMS studies. Conjugates were screened for anticancer activity by using MTT assay for human lung cancer (A549), liver hepatocellular carcinoma (HepG2), Human colon cancer (HT-29), Breast carcinoma (MCF-7), Glioblastoma cell lines (U87 MG).All molecular motifs exhibited remarkable antitumor activity against specified cell lines. Non-toxicity towards normal mouse fibroblast (L-929) is the promising feature of synthetic biomolecular scaffold which indicates

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Original Contribution | Published: 03 November 2018

D-Gluconic acid—based methotrexate prodrug—loaded mixed micelles composed of MDR reversing copolymer: in vitro and in vivo results

Popat S. Kumbhar, Swapnil Birange, Mahesh Atavale, John I, Disouza & Arehalli S. Manjappa 🖾

Colloid and Polymer Science 296, 1971-1981 (2018) Cite this article

364 Accesses 11 Citations Metrics

Abstract

The main aim of the present research was to synthesize carbohydrate (D-gluconic acid, DGA) prodrug of methotrexate (MTX) to improve the aqueous solubility and to develop mixed micelles (MMs) composed of D-α-tocopheryl poly (ethylene glycol) 1000 succinate (TPGS) as an MDR reversing copolymer and poloxamer 407 (P-407) to deliver the MTX prodrug to tumor tissue via enhanced permeability and retention (EPR) mechanism. MTX-DGA conjugate (MDGAC) was synthesized using Steglich esterification reaction. The MDGAC-loaded TPGS and P-407 MMs (MDGAC-TP MMs) were prepared by solvent evaporation technique. MDGAC-TP MMs showed low critical micelle concentration, high drug loading, sustained release profile, lower hemolytic behavior, higher in vitro cytotoxicity against the human carcinoma cell lines KB and MDR KBv, and significantly reduced in vivo toxicity. Therefore, the developed MDGAC-TP MMs could be a promising and effective approach for

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CODEN: IJRSFP (USA)

International Journal of Recent Scientific Research Vol. 9, Issue, 12(C), pp. 29951-29957, December, 2018

International Journal of Recent Scientific Rezearch

age | 73/113

DOI: 10.24327/IJRSR

Research Article

FORMULATION DEVELOPMENT AND EVALUATION OF ANTI-INFLAMMATORY POTENTIAL OF TOPICAL TENOXICAM NANOGEL ON ANIMAL MODEL

Chopade Swapnil¹., Khabade Sheeba¹., Patil Ajit³ and Powar Sayali²

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 Department of Pharmaceutical Chemistry, Tatyasaheb Kore College of Pharmacy, Warananagar,
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 Department of Pharmacology, Tatyasaheb Kore College of Pharmacy, Warananagar, Tal- Panhala,
 Dist- Kolhapur, 416113, M.S., INDIA

DOI: http://dx.doi.org/10.24327/ijrsr.2018.0912.2967

ARTICLE INFO

ABSTRACT

Article History: Received 6th September, 2018 Background: The present study is to increase the transport of tenoxicam through transdermal route, and also to present it as a possible replacement for the oral NSAID therapy for rheumatoid arthritis.

Objective: The present investigation was to develop a tenoxicam panogel with reduced particle size

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Research Article

age | 74/113

DEVELOPMENT AND VALIDATION OF A SIMPLE UV SPECTROPHOTOMETRIC AND FLUOROMETRIC METHOD FOR THE DETERMINATION OF VALACYCLOVIR HYDROCHLORIDE BOTH IN BULK AND MARKETED DOSAGE FORM

PS. Kumbhar*, AC. Rukade, PS. Sawant, AT. Gaikwad, AA. Patil, CM. Jamkhandi, MV. Shinde, AS. Manjappa and JI. Disouza

Tatyasaheb Kore College of Pharmacy, Warananagar, Dist: Kolhapur, Maharashtra, India.

ABSTRACT

Introduction: Several analytical methods such as high performance liquid chromatography (HPLC), Uv-spectrophotometry and colorimetry have been reported for quantitative estimation of Valacyclovir hydrochloride in bulk and pharmaceutical formulations. The aim of this study was to develop simple, easily accessible and economic UV spectrophotometric and newer fluorometric methods. Methods: A simple, rapid, specific and cost effective spectrophotometric method using different solvents like methanol (Method A), ethanol (Method B), water (Method C) and phosphate buffer of pH 7.4 (Method D) and fluorometric method using solvents such as methanol (Method A), water (Method B) and 0.1N HCl (Method C) has been developed to determine the Valacyclovir hydrochloride content in bulk and pharmaceutical dosage formulations. Results: The calibration graph are linear and obeys beer's law in the concentration range of 2-20 μg/mL for all four spectrophotometric methods with a correlation coefficient (r²) of 0.998, 0.996, 0.999 and 0.997, respectively while the calibration graph are linear in the concentration range of 1-10 μg/mL for all three fluorometric methods with a correlation coefficient (r²) of 0.998, 0.999 and 0.999, respectively. The accuracy and precision of the methods were evaluated based on the intra-day and inter-day variations. The accuracy of the methods was further confirmed by standard addition procedure. The other characteristics such as limit of detection (LOD) and limit of quantification (LOQ) values are also reported. Conclusion: The obtained results proved that the developed methods can be employed for the routine analysis of

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/ Vol. 12 No. 01 (2018): ASIAN JOURNAL OF PHARMACEUTICS SUPPLEMENTARY ISSUE ORIGINAL ARTICLES

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Quality by Design-based Formulation and Evaluation of Fast Dissolving Tablet of Aspirin

PDF

Published: May 13, 2018

DOI: https://doi.org/10.22377/aj

p.v12i01.2046

Dr. S. R. Desai

Abstract

Aim: The focus of the current study was to develop fast dissolving tablet (FDT) of aspirin using quality by design (QbD) approach. QbD was applied for better understanding the process and to enhance design space, using quality target product profile, critical quality attributes, and risk assessment. The aim of the project is to achieve early onset of aspirin by FDT. Materials and Methods: FDT of aspirin was developed by 32 factorial using BoxâC"Behnken design. In factorial design we have selected two variables povidone and crospovidone at three levels. The response surface plots were generated. Ultraviolet (UV), Fourier-transform infrared, differential scanning calorimeter (DSC), and X-ray diffraction (XRD) analysis have been done for pre-formulation and post-formulation evaluations. The tablets were prepared by direct compression method. Results and Discussions: The l'smax was confirmed at 275 nm by UV spectroscopy. In compatibility study IR, it was observed that the drug was in pure form and there were no major

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Article

Mixed Micelles as Nano Polymer Therapeutics of Docetaxel: Increased In vitro Cytotoxicity and Decreased In vivo Toxicity

Page | 76/113

June 2017 · <u>Current Drug Delivery</u> 15(4) DOI:10.2174/1567201814666170621113637

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International Journal of Current Pharmaceutical Research

ISSN: 0975-7066

Int J Curr Pharm Res, Vol 10, Issue 1, 13-19

Original Article

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DEVELOPMENT OF SPECTROPHOTOMETRIC AND FLUOROMETRIC METHODS FOR ESTIMATION OF DARUNAVIR USING QBD APPROACH

R. D. GODAMBE, J. L. DISOUZA, C. M. JAMKHANDI*, P. S. KUMBHAR

Tatyasaheb Kore College of Pharmacy Warananagar, Dist: Kolhapor (MS) India Email: cmjamakhandi@gmail.com

Received: 22 Oct 2017, Revised and Accepted: 12 Dec 2017

ABSTRACT

Objective: The main objective of the present study is to develop newer simple, precise spectrophotometric and fluoremetric methods of estimation for Darumavir using coupling agent 0-pthaladehyde.

Methods: The experimental work was designed for both spectroscopic and fluorometric method development and validation. The method is based on formation complex of Darunavir with 0-pthaladehyde. QbD approach was applied by varying different parameters. These parameters were designed into Ishikawa diagram.

Results: The complex Darunavir-Pirhalaidehyde in methanol with 0.1 N HCl showed linearity for both spectrophotometric and fluorometric methods. The calibration curve by spectrophotometry is linear in concentration range of 2-22 µg/ml with regression coefficient (R²) = 0.998 at 355 mm and for fluorometry it is linear in concentration range of 0.5-5.0 ng/ml with regression coefficient (R²) = 0.999. This method was found to be rugged and orbivation in different verting criteria with % RSD less than 2. The limit of detection and limit of quantification was found to be 0.2 µg/ml and 0.8 µg/ml for a spectrophotometric method and 0.1 µg/ml ml 0.4 x µg/ml for fluorometric method and 0.1 µg/ml ml 0.8 x µg/ml for fluorometric method and 0.1 µg/ml ml 0.8 x µg/ml for fluorometric method and 0.1 µg/ml ml 0.8 x µg/ml for fluorometric method and 0.1 µg/ml ml 0.8 x µg/ml for fluorometric method and 0.1 µg/ml ml 0.8 x µg/ml for fluorometric method and 0.1 µg/ml ml 0.8 x µg/ml for fluorometric method and 0.8 x µg/ml fluorometric method

Conclusion: Both methods were found to be precise with % RSD of less than 2. The % recovery of the spectrophotometric and fluorometric methods was found to be 101.04 %, 90.15 % respectively. In this way, the results of all validation parameter were within the limits as per international Conference on Harmonization guideline.

Keywords: Spectrophotometry, Fluorometry, Darumavir, Condensation reaction

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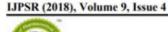
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Patil et al., IJPSR, 2018; Vol. 9(4): 1594-1599.

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Received on 21 June, 2017; received in revised form, 05 September, 2017; accepted, 17 September, 2017; published 01 April, 2018

GRANULES OF UNISTRAIN LACTOBACILLUS AS NUTRACEUTICAL ANTIOXIDANT AGENT

Abhinandan Patil "1, Shivaji Pawar 1,2 and John Disouza1

D. Y. Patil University ¹, Centre for Interdisciplinary Research, Kolhapur - 416006, Maharashtra India. Centre for Research and Technology Development ², Sinhgad Institutes, Solapur - 413255, Maharashtra, India.

Keywords:

Probiotics, Lactobacillus acidophilus, Antioxidant

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E-mail: abhisir5@gmail.com

ABSTRACT: The present study was conducted with the aim to prepare probiotics Lactobacillus acidophilus (L. acidophilus) granules which are stable at room temperature. Lactobacillus acidophilus 2285 probiotics was obtained from the N.C.I.M (National Collection of Industrial Microorganism), Pune. The formation of the semi-solid mass occurred after the further incubation at 34 °C from range (33 °C to 37 °C) in an incubator kept for the less than 24 hour time duration. This mass was homogenized and converted into granule formulation. The viability of the granule formulation was achieved with a maximum viable cell count after 24 hours of incubation in de Man, Rogosa, and Sharpe (M.R.S) agar media. Spray dried and tray dried powder of the probiotics is used for granulation, these drying methods served as a cheap alternative to the expensive freeze-drying procedure. The selected strain of L. acidophilus NCIM 2285 assessed for antioxidant activity. The antioxidant activity of L. acidophilus was demonstrated by invitro test using 2, 2--diphenyl-1-picrylhydrazyl free radical scavenging assay. The results showed that intact cells and cell-free extract of two formulations exhibited obviously higher antioxidative activity in scavenging DPPH radical than standard L. rhamnose GG, which was shown to have an

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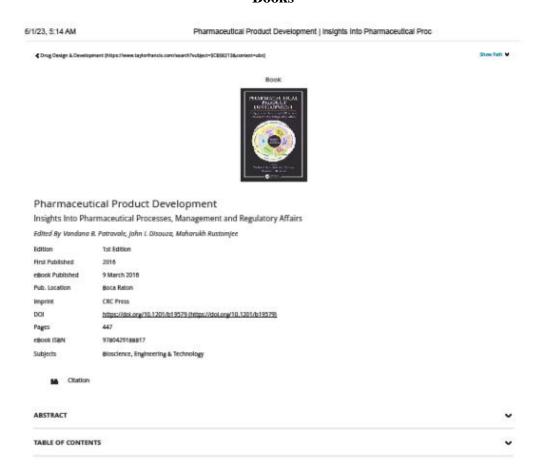
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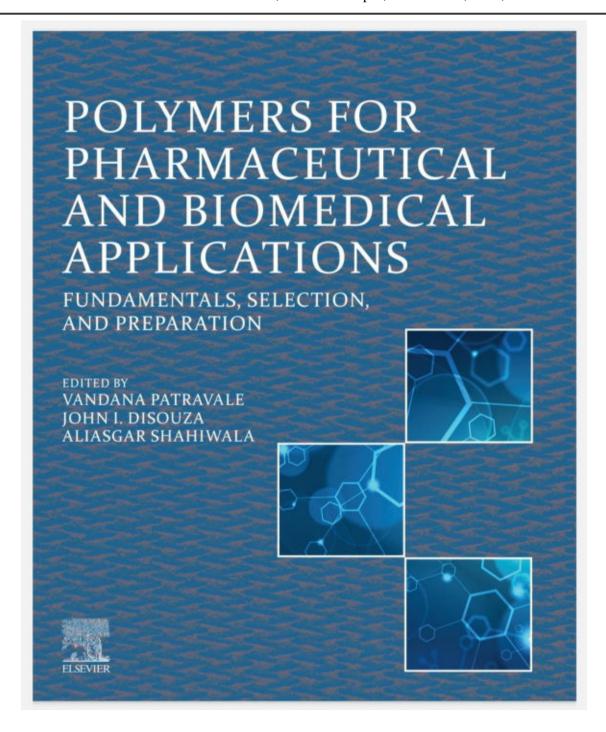
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Number of workshops/seminars/conference including programs conducted on Research Methodology, Intellectual Property Rights (IPR) and entrepreneurship during the last five years

Year	2021-22	2020-21	2019-20	2018-19	2017-18	
Number	06	04	01		01	Page 8

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Academic Year 2021-22

"Nanotechnology in Healthcare: Opportunities and Challenges



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AICTE Sponsored One Week Online Short-Term Training Program (STTP) on Induction of Novice Pharma Academicians



Manuscript writing under AICTE Sponsored One Week Online Short Term Training Program (STTP) on Induction of Novice Pharma Academicians

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AICTE-ISTE Sponsored One Week Online Induction/Refresher Program on Leveraging Academic Researchers on Developing Diagnostic Kits, Vaccines and Drug Product for Improved Therapy Management against Deadly Viruses: Lesson Learnt from COVID-19"

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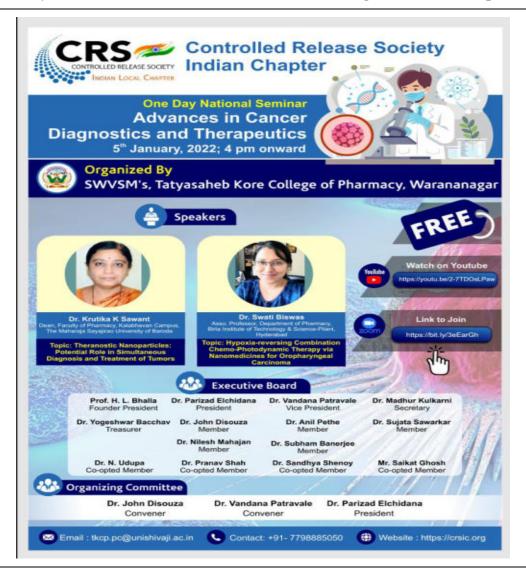
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One day National Seminar on Advances in Cancer Diagnostic and Therapeutics



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Hands-on-Training: Development and stabilization methods for Nanoparticulate Drugs



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Academic Year 2020-21 Oral Delivery of Biologics



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Manufacturing, Characterization, and Applications of Monoclonal Antibodies

Greetings from Controlled Release Society-Indian Chapter (CRS-IC).

The mission of CRS-IC is to promote education, create awareness and to encourage scientific research towards the creation of intellectual wealth in the area of drug delivery systems in India. In an attempt to fulfil our mission, we have planned A National Level Workshop on "Manufacturing, Characterization and Applications of Monoclonal Antibodies" on Saturday, 10th October, 2020.

The recent years have seen phenomenal growth in the field of research and commercialization of monoclonal antibody based therapeutics. With the right confluence of Pharmacy and Biotechnology, this field is set to progress by leaps and bounds. The workshop is hence designed to provide insight into the development, manufacturing, and applications of monoclonal antibody based formulations.

Please find attached the flyer of the workshop and the link for registration.

https://docs.google.com/forms/d/e/1FAlpQLSe1UEQG_iutNm9KCYI9oteZoEGW4hBPRfo-aoqLxukdTWAbg/viewform?usp=pp_url

E-certificates shall be provided to the participants upon successful completion of the workshop.



CONTROLLED RELEASE SOCIETY INDIAN CHAPTER

ORGANIZES A NATIONAL LEVEL WORKSHOP ON

Manufacturing, Characterization & Applications of Moneclonal Antibodies

On Saturday, 10th October 2020 between 6.30 PM- 9.30 PM

Time	Speaker Details	Topic
6.30 PM- 7.30 PM	Dr. K. Rajeshwari Founder & Managing Director, Bioklone Biotech Pvt. Ltd., Chennai, India	Making of Human Monoclonal Antibodies
7.30 PM – 8.30 PM	Dr. Suneet Shukla Senior Pharmacologist US FDA	Basics of Monoclonal Antibodies Drug Development
8.30 PM- 9.30 PM	Dr. Sachin Dubey Deputy Director- Formulation & Analytical Development, Ichnos Sciences SA, Switzerland	Role of Antibodies Based Therapeutics in the Modern Healthcare System

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Advances in Technology and Business Potential of New Drug Delivery Systems



THIS IS TO HONOUR

Mr. Andy De

WITH A BEAUTIFUL TREE PLANTATION

Therrivy you for participating on the 18th Assertational e-Symposium on

Advances in Technology and Business Potential of New Drug Delivery Systems

from 15th So 27th Sebruary 2021 as an Invited Speaker

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AICTE-ISTE Sponsored One Week Online Induction/Refresher Programmes on Fostering Pedagogy, Research Administration: Vital Domains for Effective Professional Academic Career



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Academic Year 2019-20 Industrial Pharmacy-III



Department of Pharmaceutical Sciences and Technology Nathatal Parekh Marg, Matunga (East), Mumbai – 400019

Under the Scheme of

Pandit Madan Mohan Malaviya National Mission on Teachers and Teaching (PMMMNMTT) (MHRD, Govt. of India)

Based on RUSA, Government of Maharashtra vision of Skill Development, Faculty Development Centre (FDC) called Centre for Education in Pharmaceutical Sciences, Technology and Management (CEPSTM) has been established at Institute of Chemical Technology, Mumbai under the scheme of Pandit Madan Mohan Malaviya National Mission on Teachers and Teaching (PMMMNMTT) by MHRD, Government of India.

The main objective of this centre is to serve as the nodal agency for training pharmacy, teachers in the field of industrial pharmacy, clinical data management and pharmaceutical management with respect to syllabus designed by RUSA, for the entire India.

Chief - Patron

Hon. Dr. Shri. Vinayji V. Kore (Savkar) President, Warana Industrial & Educational Complex, Warananagar

Hon. Dr. Vasanti Rasam

Administrative Officer, Shree Warana Vibhag Shikshan Mandal, Warananagar

Organizing Committee

Conveners

Dr. John Disouza, Principal, TKCP Prof. Vikas Telvekar, ICT, Mumbai

Coordinators

Mr. Kiran Patil (7798384959)

Dr. Arehalli Manjappa (9552826871)

Committee Members

Dr. Mahantesh Mattad

Mrs. Sunita Shinde Mrs. Shalaka Patki

Mr. Popat Kumbhar

Mr. Pratik Maske

Mrs. Sayali Powar Mr. Swapnil Chopade

Mr. Pritesh Lole



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9"-19" December, 2019

und huncimes



Venue
Shree Watsna Vibhag Shiksha



Taluka Panhala, District Kolhapur, Maharashtra, Pin Code: 416113

Contact

Dr. John I Disouza: 07798885050 Prof. Vikas N. Telvekar: 09869539929

No registration fees and free accommodation.



http://bit.ly/FDP-ICT-TKCP

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Academic Year 2017-18

Pharma Entrepreneurs Conclave 2017



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One Day International Conference

on

Pharmaceutical Research: Fundamentals and Advanced Trends

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Participation and Research Awards of Students in Conferences, Workshops, etc

Sr. No.	Name	Class	Level	Year	Award	Rank	
1.	Onkar Patil	M. Pharmacy	National Level	2020	Best Poster	1 st Prize	Page 96/113
2.	Onkar Patil	Poster Presentation	International Level	2020	Best Poster	1 st Prize	1 4 g c 7 0/ 1 1 c
3.	Rajeshwari Patil	Poster Presentation	International Level	2020	Best Poster	1 st Prize	
4.	Tejaswini Jadhav	Poster Presentation	National Level	2020	Best Poster	1 st Prize	
5.	Dipika Gaikwad	Poster Presentation	International Level	2020	Best Poster	Consolation Prize	
6.	Apurva Chougule	Poster Presentation	International Level	2020	Best Poster	Consolation Prize	
7.	Amruta Mhatugade	National Level	Plan of Ideal Community Pharmacy	2020		2 nd Prize	
8.	Deepali Patil	National Level	Techno Pharma Model Presentation	2020		2 nd Prize	
9.	Pradyumana Magdum	National Level	Techno Pharma Model Presentation	2020		2 nd Prize	
10.	Rohini Kulkarni	National Level	Techno Pharma Model Presentation	2020		2 nd Prize	
11.	Somesh Waghmode	State Level	Intercollegiate Competition in Microbiology	2020		1 st Prize	
12.	Girish Parle	State Level	Intercollegiate Competition in Microbiology	2020		1 st Prize	
13.	RutujaRhatwal	District Level	Avishkar	2020		1 st Prize	
14.	Rajnandini Patil	District Level	Avishkar	2020		3 nd Prize	

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Participation and Research Awards of Faculty in Conferences, Workshops, etc

Sr. No	Name of Faculty	Type of Event	Title	Yea r	Award	Rank	
1.	John Disouza	Internationa l Level	AD Scientific Index	2018	Involved in Scientist List		Page 97/113
1.	Mrs. S. S. Shinde	Conference	Nyctanthus Arbor- Tristis: As a Natural Colorant in Drug Products	2018	Best Poster	1 st Prize	
2.	Mr. P. P. Maske	Conference	Synthesis & Antimicrobial Activities of Some Novel Mercaptobenzimidazol e Derivatives.	2018	Best Poster	1 st Prize	
3.	Mrs. S. D. Gaikwad	Conference	Rational Design & Development of Novel Cadherin Inhibitors as Anticancer Agents.	2018	Best Poster	2 nd Prize	
4.	Miss. M. S. Shete	Conference	Design, Development & Characterization of Curcumin Loaded Nanoemulsion.	2018	Best Paper	2 nd Prize	
5.	Mr. K. S. Patil	Conference	A study of Imapact of Medical Advertisment on Public Health	2018	Best Poster	2 nd Prize	
6.	Mr. P. S. Kumbha r	Conference	TPGS Prodrug of Methotrexate: Improved In-vitro Anticancer Efficacy against MDA-MB 231 MDR Breast Cancer Cells	2018	Best Poster	2 nd Prize	
7.	Mr. K. S. Patil	National Level e- Poster Competitio	"Development of Pharma Educational App in COVID-19 Pandemic"	2019	Special Appreciatio n	Special Appreciatio n	

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	-				
	n on "COVID- 19 Pandemic"				
8. Mr. K S. Pati		Design, development, and characterization of Docetaxel-loaded TPGS/ Pluronic F 108 mixed micelles for improved cancer treatment	2019	Best Poster	3 rd Prize
Dr. Joh Disouz		AD Scientific Index	2020	Involved in Scientist List	Involved in Scientist List
Mr. 10. Kiran S Patil	Avishkar S.	Design, Development and Characterization of Stable Vacuum Foam Dried Docetaxel-Loaded Mixed Micelles for Improved Cancer Treatment	2021	Best Poster	First Prize
Mr. Popat S 11. Kumbh		Fabrication and Characterization of ribavirin-loaded liposomes for cancer treatment	2021	Best Poster	First Prize

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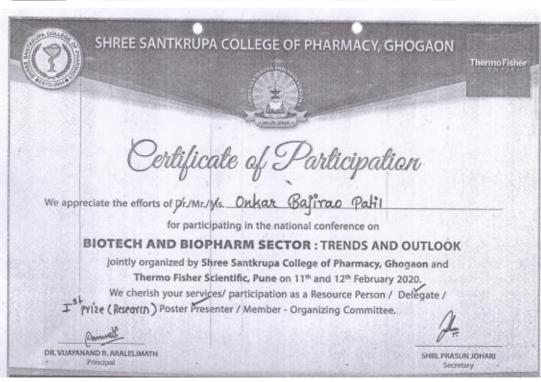
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(No: MMC/Accre.cert/MED-0207/2013 CPD Code: MMC/MAC/2020/F-014575] International Conference on 'Cancer Biology: Basic Science to Translational Research' (CBTR-2020) held on 17"-18" January 2020. Organized by Department of Stem Cell and Regenerative Medicine D. Y. Patil Medical College ent Units Of D. Y. Patil Education Society (Deemed to be University), Kolhapur Re-accredited by NAAC with "A" grade This is to certify that Mr. / Ms. / Mrs. / Dr. / Prof. Onkar Patil UHIQU€ has awarded First / Secontd / Taird prize in Oral / Poster presentation in CBTR 2020. Maharashtra Medical Council (MMC) has granted 04 (FOUR) CPO Credit Points for the Delegates).

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(No: MMC/Accre.cert/MED-0207/2013 CPD Code: MMC/MAC/2020/F-014575) Cancer Biology: Basic Science to Translational Research' (CBTR-2020) held on 17°-18° January 2020. Kolhapur Cancer Centre Organized by Department of Stem Cell and Regenerative Medicine and D. Y. Patil Medical College Constituent Units Of D. Y. Patil Education Society (Deemed to be University), Kolhapur Re-accredited by NAAC with "A" grade This is to certify that Mr./ Ms./Mrs./Dr./Prof. Apurva R. Chouque has Participated in Oral/ Poster presentation in CBTR-2020. rashtra Medical Council (MMC) has granted 04 (FOUR) CPD Credit Points for the Delegates). Dr. Salprasad Kavatheka MMC Observer MMC/MAO-01302/2017

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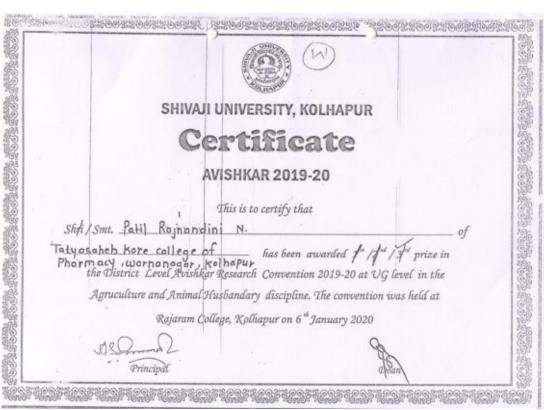


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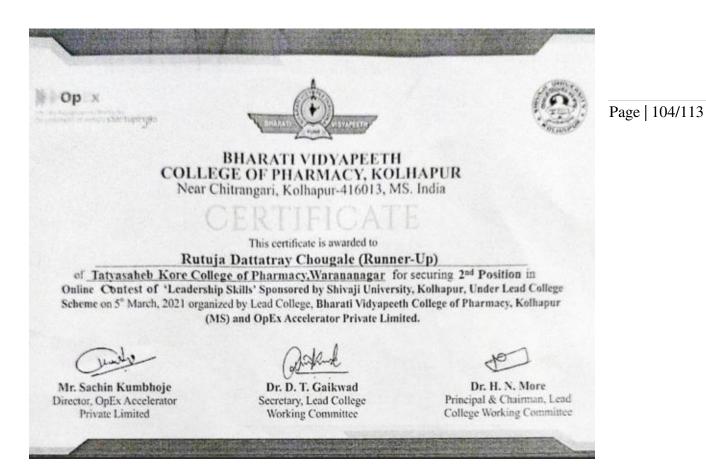
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Patents

Patent Application No.	Status of Patent (Published/ Granted)	Inventor/s Name	Patent	Applicant/s Name	Patent Published Date / Granted Date Page 108/113
202221024815A	Published	Dr. A. S. Manjappa	Combination drug therapy	Dr. A. S. Manjappa	13/05/2022
202121023742	Published	Mr. Kiran Shivaji Patil	Transdermal ethosome composition of ranozaline	Mr. Kiran Shivaji Patil	
202021038512	Published	Mr. P. S. Kumbhar	A novel bike friendly bright helmet with different safety features	Mr. P. S. Kumbhar	18/09/2020
201921009581 A	Published	Dr. A. S. Manjappa	Microparticles containing montelukast for inhalation therapy	Dr. A. S.Manjappa	19/04/2019
1943/MUM/2015	Published	Dr. J. I. Disouza	A novel herbal extract with anticancer activity	Dr. J. I. Disouza	28/04/2017
2021/MUM/2008A	Published	Dr. A, S. Sherikar	Synthesis of phenyl nitrate derivatives of free carboxylic acid group containing NSAIDS as cyclooxygenase inhibitor for antiflammatory, analgesic and smooth muscle relaxant activity	Dr. A. S. Sherikar	02/04/2010

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(12) PATENT APPLICATION PUBLICATION

(19) INDIA

(51) International

(86) International Application No

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classification

(22) Date of filing of Application :27/04/2022

(21) Application No.202221024815 A

(43) Publication Date: 13/05/2022

(54) Title of the invention : COMBINATION DRUG THERAPY FOR ANTICANCER

:A61K0031145000. A61K0031496000.

A61K0045060000, A61K0031337000,

A61K0031498500

NA

-NA

71)Name of Applicant :

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Page | 109/113

2)Dr.Shashikant C. Dhawale

3)Dr. Arehalli Sidramappa Manjappa

Name of Applicant : NA Address of Applicant : NA (72)Name of Inventor :

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ramardad@rediffmail.com Mob no.9503756634. --

2)Dr. Shashikant C. Dhawale Address of Applicant :Dr. Shashikant C. Dhawale ,58, Veer Sawarkar Nagar, Wadi Bk, Purna Road Nanded. 431605 MS India. Email id-shashiprathmesh@gmail.com Mob no

9970700030.

3)Dr. Arehalli Sidramappa Manjappa

Address of Applicant :Department of Pharmaceutics, Tatyasaheb Kore College of Pharmacy, Warananagar -416113 Tal: PangalaDistrict: Kolhapur Maharashtra Email Id: nanju_as82@yahoo.co.in Mob No: 9552826871;8956647419 ----

COMBINATION DRUG THERAPY FOR ANTICANCER Abstract The present invention states that the combination therapy useful for treatment of oncological disorders. Further invention relates to Ketoconazole; Disulfiram; and Tadalafil having 1:1:1 molar ratio respectively. Further embodiment of present invention relates to Ketoconazole; Disulfiram; Tadalafil cocktail in combination with

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Application	on Details
APPLICATION NUMBER	202121023742
APPLICATION TYPE	ORDINARY APPLICATION
DATE OF FILING	28/05/2021
APPLICANT NAME	1 . Ms. Hemalata Suhasrao Dol 2 . Dr. Ashok Ananda Hajare 3 . Dr. Trupti Ashok Powar 4 . Mr. Kiran Shivaji Patil
FITLE OF INVENTION	TRANSDERMAL ETHOSOME COMPOSITION OF RANOLAZINE
FIELD OF INVENTION	CHEMICAL
E-MAIL (As Per Record)	hemalatadol@gmail.com
ADDITIONAL-EMAIL (As Per Record)	hemalatadol@gmail.com
E-MAIL (UPDATED Online)	
PRIORITY DATE	

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	Age to see	100
Applica	ation Details	
APPLICATION NUMBER	202021038512	'age
APPLICATION TYPE	ORDINARY APPLICATION	
DATE OF FILING	07/09/2020	-
APPLICANT NAME	1 . BHOJILING EKNATH KUMBHAR 2 . POPAT SONAPPA KUMBHAR	
	3. SUHAS SANJAY KUMBHAR 4. NILESH ARUN KUMBHAR 5. RAVINDRA SOPAN KUMBHAR	
TITLE OF INVENTION	A NOVEL BIKE-FRIENDLY BRIGHT HELMET WITH DIFFERENT SAFETY FEATURES.	-
FIELD OF INVENTION	TEXTILE	
E-MAIL (As Per Record)		
Record)	pskumbhar1.tkcp@gmail.com	
E-MAIL (UPDATED Online)		
PRIORITY DATE		
REQUEST FOR EXAMINATION DATE	07/09/2020	1
PUBLICATION DATE (U/S	18/09/2020	

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(12) PATENT APPLICATION PUBLICATION

(21) Application No.201921009581 A

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(19) INDIA

(22) Date of filing of Application :12/03/2019

(43) Publication Date: 19/04/2019

(54) Title of the invention: MICROPARTICLES CONTAINING MONTELUKAST FOR INHALATION THERAPY.

(51) International classification (31) Priority Document No	:A61K 9/00 :NA	(71)Name of Applicant: 1)PANCHAL CHANDRAWADAN VISHWAMBHAR Address of Applicant: MAHARASHTRA COLLEGE OF PHARMACY, NILANGA, TALUKA-NILANGA, DIST.: LATUR, MAHARASHTRA, INDIA, PIN CODE: 413521. Maharashtra India 2)BANSODE HEMANT BALU
(32) Priority Date	:NA	3)DR. JOSHI SUMIT ASHOK
(33) Name of priority country	:NA	4)DR. DAMA GANESH YOGIRAJ
(86) International Application No	:NA	5)DR. AREHALLI S. MANJAPPA
Filing Date	:NA	6)GURAV Prashant B.
(87) International Publication No	: NA	7)JADHAV Sachin Manik
(61) Patent of Addition to Application Number	:NA	(72)Name of Inventor :
Filing Date	:NA	1)PANCHAL CHANDRAWADAN VISHWAMBHAR
(62) Divisional to Application Number	:NA	2)BANSODE HEMANT BALU
Filing Date	:NA	3)DR. JOSHI SUMIT ASHOK
		4)DR. DAMA GANESH YOGIRAJ
		5)DR. AREHALLI S. MANJAPPA
		6)GURAV Prashant B
		7)JADHAV Sachin Manik

(57) Abstract:

ABSTRACT The present invention relates to microparticles containing Montelukast for inhalation therapy, specifically microparticles containing Montelukast sodium loaded chitosan and sodium alginate and a process for preparation thereof.

No. of Pages: 17 No. of Claims: 10

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Ар	plication Details		
APPLICATION NUMBER	1943/MUM/2015		
APPLICATION TYPE	ORDINARY APPLICATION		
DATE OF FILING	18/05/2015		
APPLICANT NAME	1 . DISOUZA; JOHN INTRU 2 . PATIL; GANPATRAO DNYANDEV 3 . PATIL; AJIT BABURAO 4 . PATIL; ABHINANDAN RAVSAHEB		
TITLE OF INVENTION	A NOVEL HERBAL EXTRACT WITH ANTICANCER ACTIVITY		
FIELD OF INVENTION	PHARMACEUTICALS		
E-MAIL (As Per Record)			
ADDITIONAL-EMAIL (As Per Record)	poonamdhake@gmail.com		
E-MAIL (UPDATED Online)			
PRIORITY DATE			
REQUEST FOR EXAMINATION DATE	30/06/2017		
PUBLICATION DATE (U/S 11A)	28/04/2017		

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(12) PATENT APPLICATION PUBLICATION

(21) Application No.2021/MUM/2008 A

Page | 114/113

(19) INDIA

(22) Date of filing of Application :22/09/2008

(43) Publication Date: 02/04/2010

(54) Title of the invention: SYNTHESIS OF PHENYL NITRATE DERIVATIVES OF FREE CARBOXYLIC ACID GROUP CONTAINING NSAIDS AS CYCLOOXYGENZISE INHIBITOR FOR ANTI-FLAMMATORY, ANALGESIC AND SMOOTH MUSCLE RELAXANT ACTIVITY

(51) International classification (31) Priority Document No (32) Priority Date (33) Name of priority country (86) International Application No Filing Date (87) International Publication No (61) Patent of Addition to Application Number Filing Date (62) Divisional to Application Number Filing Date	(71)Name of Applicant: 1)BHATIA MANISH SUDESH Address of Applicant: 65, CHAVREKAR COLONY, CU7C225/22 1NA
---	--

(57) Abstract:

The present invention describes the synthesis of phenyl nitrate derivatives of free carboxylic acid group containing NSAIDs as cyclooxygenase inhibitor and nitric oxide donors (CINOD) for anti-inflammatory, analgesic and smooth muscle relaxant activity, novel cyclooxygenase 2 (COX-2) selective inhibitors and novel compositions comprising at least one cyclooxygenase 2 (COX-2) inhibitor, and, at least one compound that donates, transfers, releases nitric oxide and/or stimulates endogenous synthesis of nitric oxide and/or elevates endogenous levels of endothelium-derived relaxing factor or is a substrate for nitric oxide synthase, and/or at least one therapeutic agent. The invention also provides methods for treating inflammation, pain and fever; for treating and/or improving the gastrointestinal properties of COX-2 selective inhibitors; for facilitating wound healing, for treating and/or preventing other disorders resulting from elevated levels of cyclooxygenase-2; and for improving the cardiovascular profile of COX-2 selective inhibitors.

No. of Pages: 56 No. of Claims: 8

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