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3.3.1

**Number of research papers
published per teacher in the
Journals notified on UGC care list
during the last five years**



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3.3.1: Number of research papers published per teacher in the Journals notified on UGC care list during the last five years

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LIST OF PUBLISHED PAPER

Sr. No	Title of Paper	Name of Author	Name of Journal
ACADEMIC YEAR 2022-23			
1	Formulation and evaluation of Fluconazole topical gel by using <i>ocimum basilicum</i> seed mucilage as agelling agent	Nilesh Sancheti, Dr.Pratima Shinde	World Journal of Pharmacy and Pharmaceutical Sciences
2	Review on formulation of microemulsion as novel drug delivery system	Prathmesh Kulkarni, Dr.Pratima Shinde	World Journal of Pharmacy and Pharmaceutical Sciences
3	Review of comparative study of OTF with conventional drug delivery system	Shravani Havale, Dr.Pratima Shinde	World Journal of Pharmacy and Pharmaceutical Sciences
4	Hutchinson-Gilford Progeria Syndrome	Apeksha Pangare, Dr.Pratima Shinde	World Journal of Pharmacy and Pharmaceutical Sciences
5	Design, Docking, Insilco ADME Prediction Of Novel Indole Based Benzamide Scaffolds Targeting For Estrogen Receptor Alfa In Af-2 Domain For Effective Anticancer Treatment	Dr. Swati N. Deshmukh	Journal of Pharmaceutical Negative Results
6	Study Of The Properties And Behaviors Of Nanoparticles And Their Potential Applications In Medicine And Catalysis	Dr. Swati N. Deshmukh , Vanita Gade, Sunita Shewalkar, Payal Pansare , Trupti Kajale, Aarti Gaikwad	Journal of Pharmaceutical Negative Results
7	Novel Film Forming Spray from Tea Tree Leaves with Special Emphasis on Development, Formulation and Evaluation	Dr. Swati N. Deshmukh , Vanita Gade, Sunita Shewalkar, Dumbre R K, Girme S	Journal of Positive School Psychology
8	Editorial: Medicinical Chemistry Actual Teaching Aesthetics	Amit G.Nerkar	Current Trends in Pharmacy and Pharm. chem
9	<u>Ethnopharmacological review of <i>boswellia serrata</i> for anticancer activity</u>	Amit G.Nerkar and Pallavi Gade	Current Trends in Pharmacy and Pharm. chem
10	Formulation and evaluation of Fluconazole topical gel by using <i>ocimum basilicum</i> seed mucilage as agelling agent	Nilesh Sancheti, Dr.Pratima Shinde	World Journal of Pharmacy and Pharmaceutical Sciences
11	<u>Ethnopharmacological review of <i>vinca</i> plant for anticancer activity</u>	Amit Gajanan Nerkar, Nagarkar Rushikesh,	Current Trends in Pharmacy and Pharm. chem




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12	<u>Ethnopharmacological review of kalmegh for anticancer activity</u>	Amit Gajanan Nerkar, Shubhangi Badar	Current Trends in Pharmacy and Pharm. chem
13	<u>Ethnopharmacological review of ginger for anticancer activity</u>	Amit G.Nerkar and Srushti Ghadge	Current Trends in Pharmacy and Pharm. chem
14	<u>Recent advances in pharmaceutical, chemical and biological sciences</u>	Amit G.Nerkar, Dumbre R K	Current Trends in Pharmacy and Pharm. chem
15	Editorial: Pharmaceutical Organic Chemistry Actual Teaching Aesthetics	Amit G.Nerkar	Current Trends in Pharmacy and Pharm. chem
16	Formulation and Evaluation of Herbal Syrup of Bhilawa Seed Extract	Amit G.Nerkar and Ashutosh Pansare	Current Trends in Pharmacy and Pharm. chem
17	Formulation and Evaluation of Herbal Syrup of Turmeric Extract	Amit G.Nerkar , Rushikesh Nagarkar and Shubhangi Badar	Current Trends in Pharmacy and Pharm. chem
18	Formulation and Evaluation of Herbal Syrup of Indian Mulberry (Noni) Extract	Amit G.Nerkar and Pallavi Gade	Current Trends in Pharmacy and Pharm. chem
19	Formulation and Evaluation of Herbal Syrup of Kalmegh Extract	Amit G. Nerkar, Shubhangi Badar and Rushikesh Nagarkar	Current Trends in Pharmacy and Pharm. chem
20	Ethnopharmacological Review of Turmeric for anticancer activity	Amit Gajanan Nerkar, Rushikesh Nagarkar and Shubhangi Badar	Current Trends in Pharmacy and Pharm. chem
21	Ethnopharmacological Review of Arjuna	Amit G.Nerkar, Rahul K. Dumbre and Shubhangi Badar	Current Trends in Pharmacy and Pharm. chem
22	A review on emerging trends in targeted drug delivery systems: events, approaches and specific drug targeting	DR. Gita Chaurasia, swati Kale and Payal Pansare	Journal of emerging technologies and Innovative research
23	Microwave Induced Solid Dispersion as a Novel Technique for Enhancing Solubility of Rifampicin	Dr. Swati Jogdand, Pooja Jadhav	Neuroquantology
24	Formulation development of sustained release swellable matrix tablet of venlafaxine HCl	Dr. Swati Jogdand, Pooja Jadhav	NeuroQuantology: An Interdisciplinary Journal of Neuroscience and Quantum Physics
25	Analytical Method Development And Validation Of Niclosamide By RP-HPLC	Dr. Narendra Gowekar, Shivam Kale , Dr. Swati Jogdand , Sujata Shinde , Dr. Rahul Dumbre	IJFANS International Journal of Food and Nutritional Sciences
26	Determination Of Total Flavonoid Content, Total Steroids Content, Total Tannin Content And Antioxidant Activity Of Fruits Of Hydro Alcoholic Extract Of Piper Longum Linn.	Dr. Swati Jogdand , Dr. Narendra Gowekar , Mrs. Sujata Shinde , Dr Rahul Dumbre	IJFANS International Journal of Food and Nutritional Sciences
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29	An Overview Novel Drug Delivery System	Ms. Swati B. Kale, Shubhangi Thopate	IJFMR International Journal for Multidisciplinary Research

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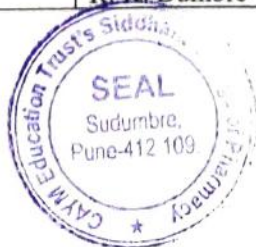

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33	Review: Medicinal Plants on Anti-Diabetic Activity	Roshan Phanse, Dr. Narendra Gowekar	Journal of Emerging Technologies and Innovative Research
34	Review: Topical Drug Delivery System	Dhanshree Kanawade, Dr. Narendra Gowekar	International Journal of Pharmaceutical Research and Applications
35	Formulation and evaluation of Fluconazole topical gel by using ocimum basilicum seed mucilage as agelling agent	Nilesh Sancheti, Dr. Pratima Shinde	International Journal of Advanced Science and Technology
36	Review on formulation of microemulsion as novel drug delivery system	Dr. Pratima Shinde, Prathmesh Kulkarni,	World Journal of Pharmacy and Pharmaceutical Sciences
37	Review of comparative study of OTF with conventional drug delivery system	Dr. Pratima Shinde, Shravani Havale	World Journal of Pharmacy and Pharmaceutical Sciences
38	Hutchinson-Gilford Progeria Syndrome	Dr. Pratima Shinde, Apeksha Pangare,	World Journal of Pharmacy and Pharmaceutical Sciences
39	A Review on: 3D Printing in Pharmaceutical Industry	Devendra Gorle, Dr. Pratima Shinde	International Journal of Pharmaceutical Research and Application
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49	A Short Review on: Definition, Principle and Applications of HPLC	S. D. Varhadi*, V. A. Gaikwad, R. R. Sali, Vikas Kandekar	International journal of Pharmacy & Pharmaceutical Research
50	Novel Film Forming Spray from Tea Tree Leaves with Special Emphasis on Development, Formulation and Evaluation	Swati N. Deshmukh, Vanita Gade, Rahul Dumbre, Swappali Girme, Sunita	Journal of Positive School Psychology (JPSP)




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55	A Review on: Natural Gums and Mucilages: Used as Excipients and Pharmaceutical Sciences	Dr. Pratima Shinde	International Journal of Pharmaceutical Research and Applications
56	A Review on: Effectiveness of Various Types of Face masks	Dr. Pratima Shinde	International Journal of Pharmaceutical Research and Applications
57	A Review on: Retinoids in cosmetics and dermatological treatment and its detrimental side effects	Dr. Pratima Shinde	International Journal of Pharmaceutical Research and Applications
58	A Review On:3D Printing In Pharmaceutical Industry	Dr. Pratima Shinde	International Journal of Pharmaceutical Research and Applications
59	Rp-hplc method development and validation for estimation of chlorpheniramine maleate in liquid dosage form	Dr. R. K. Dumbre, Mr. Vikas Kandekar	Journal of Emerging Technologies and Innovative Research
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Abstract

FORMULATION AND EVALUATION OF FLUCONAZOLE TOPICAL GEL BY USING OCIMUM BASILICUM SEED MUCILAGE AS A GELLING AGENT

Nitesh Nitin Sancheti* and Dr. Pratima Shinde

ABSTRACT
Fluconazole is an imidazole derivative used for the treatment for local and systemic fungal infection. The oral use of fluconazole is not recommended as it has many side effects. The present study was design to formulate and evaluate different formulae of topical gel containing fluconazole for treatment of fungal infection of skin. The gel was formulated by using natural seed mucilage obtained from ocimum basilicum with different concentration of acacia and glycerin. Three different formulae were prepared and characterized physically in term of color, spreadability, pH and drug content. In-vitro drug release in phosphate buffer pH 6.8 and permeation study through cellulose membrane using a modified franz diffusion cell were performed. Candida albicans was used as model fungus to evaluate the antifungal activity of prepared formulae. The results of in-vitro drug release and its permeation studies showed that the highest values was form F2 (89.18% of drug release after 4 hr) also F2 shows highest antifungal activity.

Keywords: Fluconazole, Ocimum Basilicum, Glycerin, Acacia.


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Abstract

REVIEW ON FORMULATION OF MICROEMULSION AS NOVEL DRUG DELIVERY SYSTEM

Prathamesh P. Kulkarni* and [Pratima S. Shinde](#)

ABSTRACT


Microemulsion is defined as the isotropic, thermodynamically stable transparent system of oil-water-surfactant and co-surfactant which having very small droplets size (20-200nm). Microemulsions are isotropic, thermodynamically stable transparent (or translucent) systems of oil, water and surfactant. Frequently in combination with a cosurfactant with a droplet size usually in the range of 20-200 nm. They can be classified as oil-in-water (o/w), water-in-oil (w/o) or bicontinuous systems depending on their structure and are characterized by ultra low interfacial tension between oil and water phases. To date microemulsions have been shown to be able to protect labile drug, control drug release, increase drug solubility, increase bioavailability and reduce patient variability. Furthermore, it has proven possible to formulate preparations suitable for most routes of administration. Since the discovery of microemulsions, they have attained increasing significance both in basic research and in industry. Due to their unique properties, namely, ultra-low interfacial tension, large interfacial area, thermodynamic stability and the ability to solubilize otherwise insoluble liquids, uses and applications of microemulsions have been numerous. Microemulsions are readily distinguished from normal emulsions by their transparency, low viscosity and more fundamentally their thermodynamic stability. Microemulsions are shown to be effective dermal delivery mechanism for several active ingredients for pharmaceutical and cosmetic applications. Topical microemulsions allow rapid penetration of active molecules due to the large surface area of the internal phase, and their components reduce the barrier property of stratum corneum. Microemulsions thereby enhance dermal absorption compared with conventional formulations and are therefore a promising vehicle due to their potential for transdermal drug delivery.

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Abstract

REVIEW OF COMPARATIVE STUDY OF OTF WITH CONVENTIONAL DRUG DELIVERY SYSTEM

*Shravani A. Havale and [Dr. Pratima S. Shinde](#)

ABSTRACT

Fast dissolving oral films (FDOFs) are the most advanced form of oral solid dosage form due to more flexibility and comfort. It improve the efficacy of APIs by dissolving within minute in mouth after the contact with less saliva as compared to fast dissolving tablets, without chewing and no need of water for administration. The FDOFs place as an alternate within the market thanks to the consumer's preference for a fast-dissolving product over conventional tablets / capsules. The oral thin-film technology remains within the beginning stages and has bright future ahead because it fulfills all the necessity of patients. Eventually, film formulations having drugs are going to be commercially launched using the oral film technology. However, for future growth point of view the oral thin film sector is well-positioned. In US market the OTC films of pain management and sinusitis are commercialized. More importantly, prescription OTFs have now been approved in US, EU and Japan which are the three major regions. These approved Rx films, have potential to dominate over other oral dosage forms of the same drugs. It seems that the worth of the general oral thin film market will grow significantly.

Keywords : Oral thin film (OTF), Conventional drug delivery, Oral disintegrated formulation.

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Abstract

HUTCHINSON – GILFORD PROGERIA SYNDROME
 Apekha S. Pangare* and Dr. Pratima S. Shinde

ABSTRACT
 Hutchinson – Gilford Progeria Syndrome (HGPS) is an Autosomal dominant, rare, fatal pediatric segmental pro-mature aging disease. Hutchinson – Gilford Progeria Syndrome, a rare genetics disorder associated with a characteristic aged appearance very early in life. The review on Hutchinson- Gilford Progeria Syndrome summarizes the clinical characteristics of this disease and therefore the underlying mutation within in the lamin A (LMNA) gene that leads to produce abnormal lamin A. Le Progerin; this disrupts the nuclear membrane and alters transcription. Accumulation of this progerin proteins in the normal cell which leads to nuclear morphology defects, decreased lifespan and premature cell death occurs. Based on the positive outcomes from the studies based on statins, aminobisphosphonates and FTIs, two clinical trials have been performed in children with HGPS.

Keywords: Hutchinson – Gilford Progeria Syndrome, Progerin Protein, Lamin A/C gene

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Design, Docking, Insilco ADME Prediction Of Novel Indole Based Benzamide Scaffolds Targeting For Estrogen Receptor Alfa In Af-2 Domain For Effective Anticancer Treatment

B. J. Werude , Dr. V. A. Chatpallwar, S. N. Wagh, Dr. V. S. Neharkar, Dr. S. R. Deshmukh, Dr. R. Mhetre and Dr. A.A.A.Garud

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ABSTRACT

Aim: To discover some novel indole based benzamide scaffold and their screening through in silico approach.

Background: Designed 7-substituted -1-4-(piperidine-1-yl) methoxybenzyl)-1H-indole-3-carboxamide derivatives targeting on ERe modulators. several interactions between the ligand and amino acid residues that would probably elicit fruitful modulation of the receptor using 4X33 pdb of ERe.





Objective: Studied in silico novel molecules of 7-substituted -1-4-(piperidine-1-yl) methoxybenzyl)-1H-indole-3-carboxamide derivatives



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Study Of The Properties And Behaviors Of Nanoparticles And Their Potential Applications In Medicine And Catalysis

June 2023
DOI: 10.47758/ncr.2023.14.03.477

Authors:
 **Nidhi Jain**
 Bharati Vidyapeeth's College of Engineering
 **N Swati**  **Deshmukh**
 **Vanita Gade**

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Novel Film Forming Spray from Tea Tree Leaves with Special Emphasis on Development, Formulation and Evaluation

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Swati N. Deshmukh, Vanita Gade, Aniket Garud, Rahul Dumbre, Bhagyashri Warude, Sunita Mahara, Swapnali Girmie, Sunita Shewalkar

Abstract

The dictum of this study was to develop topical film forming spray having tea tree oil which might increase wound healing. Film-forming sprays supply several benefits compared to standard topical preparations as a result of they will give uniform drug distribution and dose, increased bioavailability, lower incidence of irritation, continuous drug unleash, and accelerated wound healing through wet management. Film-forming sprays comprises polymers and excipients that improve the characteristics of preparations and enhance the soundness of active substances. every style of chemical compound and excipient can turn out films with completely



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November 2022 *Current Trends in Pharmacy and Pharmaceutical Chemistry* 4(4):135-138
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 **Amit Gajanan Nerkar**

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Abstract

Medicinal chemistry has been the backbone of the pharmacy profession. Medicinal chemistry may be defined as chemistry of medicines. This allied branch with pharmacology is being taken as a cumbersome and troublesome subject. This review deals with the mostly incorrect practice and assumptions of Medicinal chemistry as a theoretical subject being taught at the college level for diploma, undergraduate, postgraduate, and doctoral studies.

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Ethnopharmacological review of boswellia serrata for anticancer activity

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Author Details: Amit Gajanan Nerkar*, Pallavi Gade

Volume : 4, **Issue :** 4, **Year :** 2022

Article Page : 144-147

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Ethnopharmacological review of vinca plant for anticancer activity

Review Article

Author Details: Amit Gajanan Nerkar*, Rushikesh P Nagarkar, Shubhangi Badar

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Abstract

The strive of this assessment shows that Vinca rosea, many obviously grown vegetation round us which can be used for medicinal purposes. It has many known names like vinca Rosea, Madagascar periwinkle, vibrant eyes, Cape periwinkle, graveyard plant, old maid, crimson periwinkle, rose periwinkle myrtle. Ayurveda is the Indian conventional device of medication which focuses on the scientific capability of plant life. Catharanthus roseus is one plant recognized nicely in Ayurveda. It is known for its antitumour, anti-diabetic, anti-microbial, anti-oxidant and antimutagenic effects. It is an evergreen plant first originated from islands of Madagascar. The flowers can also range in color from red to pink and

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Ethnopharmacological review of Kalmegh for anticancer activity

Review Article

Author Details: Amit Gajanan Nerkar*, Shubhangi Badar

Volume : 4, Issue : 4, Year : 2022

Article Page : 152-157

<https://doi.org/10.18231/j.ctppc.2022.027>

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Abstract

The cultivation of Kalmegh (*A. paniculata*), a plant with numerous therapeutic benefits, has increased its significance as a medicinal plant. Given the plant's beneficial properties, it can be strongly recommended as a secure, crucial medicinal herb for people. There are many different chemical components in this herb, but the primary ones are lactones, diterpenoids, diterpene glycosides, flavonoids, and flavonoid glycosides. It possesses a wide range of pharmacological effects, including antibacterial, hepatoprotective, antitumor, anticancer, hypoglycemic, immunomodulatory, and hypotensive actions. This study describes the past and status of research on *Andrographis paniculata* plant's therapeutic utilisation. It also emphasizes at compiling vast pharmacological applications to make the potential image of *A. paniculata* as a multipurpose medicinal agent. This ethnopharmacological review deals with anticancer activity of Kalmegh or *A. paniculata*.



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Amit Gajanan Nerkar*,
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Ethnopharmacological review of ginger for anticancer activity

Review Article

Author Details : Amit Gajanan Nerkar*, Srushti Ghadge

Volume : 4, **Issue :** 4, **Year :** 2022

Article Page : 158-164

<https://doi.org/10.18231/j.ctppc.2022.028>

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Abstract

Zingiber officinale is a plant found locally in India that has been widely used as a flavouring agent in savoury dishes such as curries and sweets such as cakes and cookies, alcoholic beverages as well as in alcoholic beverages, like in tea. Ginger is a well-known herb, commonly used in traditional medicine all over the world. Ginger has been used for thousands of years to treat colds, nausea, arthritis, migraines, and high blood pressure. The many pharmacological activities of ginger are antiemetic, antidiabetic, analgesic, anti-inflammatory, anticancer, antioxidant, anticoagulant, antibacterial, anti-inflammatory, estrogenic and cardiovascular activities. Chemical irritants and an unsaturated phenolic ketone liquid C₁₇H₂₄O₃ are responsible for the spicy taste of ginger. The main components of ginger are aromatic essential oils, antioxidants and pungent resins. These aromatic or pungent compounds have been identified as C₆H₅C(O)CH₃, known as a chemical irritant, liquid unsaturated phenolic ketones C₁₇H₂₄O₃ and Vanillylacetone.

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Recent advances in pharmaceutical, chemical and biological sciences

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Abstract

Webinar was organized on Recent Advances in Pharmaceutical, Chemical and Biological Sciences on 15 Oct 2022. There were at total 8 speakers including the Chief Guest for the Webinar. Dr. Rahul K. Dumbre, Principal of CAYMET Siddhant College of Pharmacy, Sudumbare, Pune, M.S., India was the Convener for the Webinar. Dr. Amit G. Nerkar, Professor and Research Head, Siddhant College of Pharmacy and Founder and Director, Ateos Foundation of Science Education and Research, Pune, M.S., India was the Co-convener for the Webinar. At total 100 participants attended 8 sessions. The function started with Inauguration and concluded with Vote of Thanks by the Co-convener.

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Rahul M. Dumbre
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Formulation and evaluation of herbal syrup of bhilawa seed extract

A. Nehra, Ashishali Parashar · Published 15 April 2022 · Current Trends in Pharmacy and Pharmaceutical Chemistry

Semecarpus anacardium Linn (Family: Anacardiaceae), commonly known as 'Bhalataka' or 'Bhilwa', has been used in various traditional systems of medicine to treat various ailments since ancient times. Its nuts contain many bioactive compounds such as biflavonoids, phenolics, bhilawanols, minerals, vitamins and amino acids, which exhibit various healing properties. Fruit and seed extracts exhibit various activities such as anti-atherogenic, anti-inflammatory, antioxidant, antibacterial, anti...

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Title
A REVIEW ON EMERGING TRENDS IN TARGETED DRUG DELIVERY SYSTEMS: EVENTS, APPROACHES AND SPECIFIC DRUG TARGETING

Authors
Gita Charanada
Shweta Kalia
Payal Parmare

Abstract
Recently, Targeted Drug Delivery System (TDDS) is an advanced tool towards innovation in the field of Pharmaceutical science. It targets a particular cell or tissue or organ rather than the whole body to deliver the drug through a carrier to achieve its therapeutic effectiveness. It has developed various advanced methods to deliver the drug in a certain amount for a prolonged period of time to target sites to treat chronic or lethal diseases in the body. The drug is delivered in a specific location that is the combination of fields like molecular biology, polymer science, biochemistry, pharmacology, microbiology etc. The need of TDDS is to overcome the problems associated with conventional drug delivery systems. Some problems are related with administration, distribution, metabolism and elimination of drugs and carriers. This approach helps to increase the drug therapeutic index and decreases the side effects of drugs during multiple interactions and non-targeted actions. In this review concept of TDDS, Study of its generation, types, Advantages over Conventional drug delivery systems, disadvantages and application was summarized. Events and biological processes involved in drug Targeted delivery mechanisms through various biological processes are

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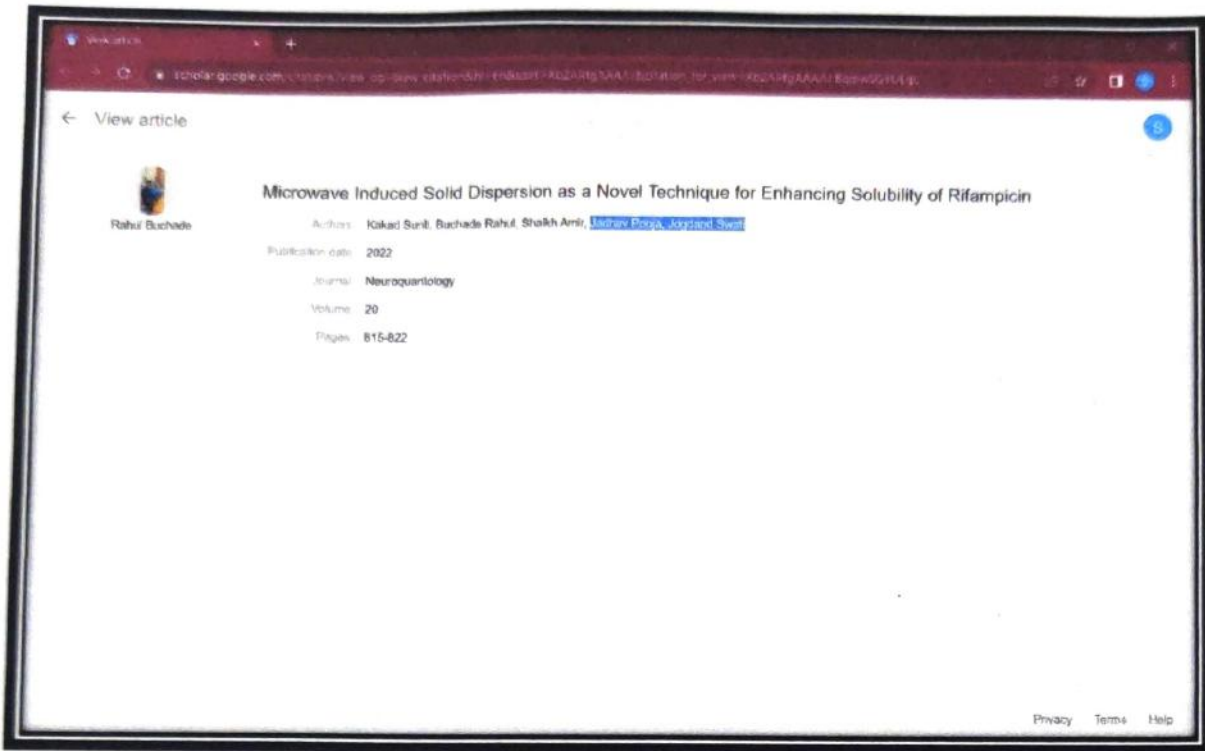
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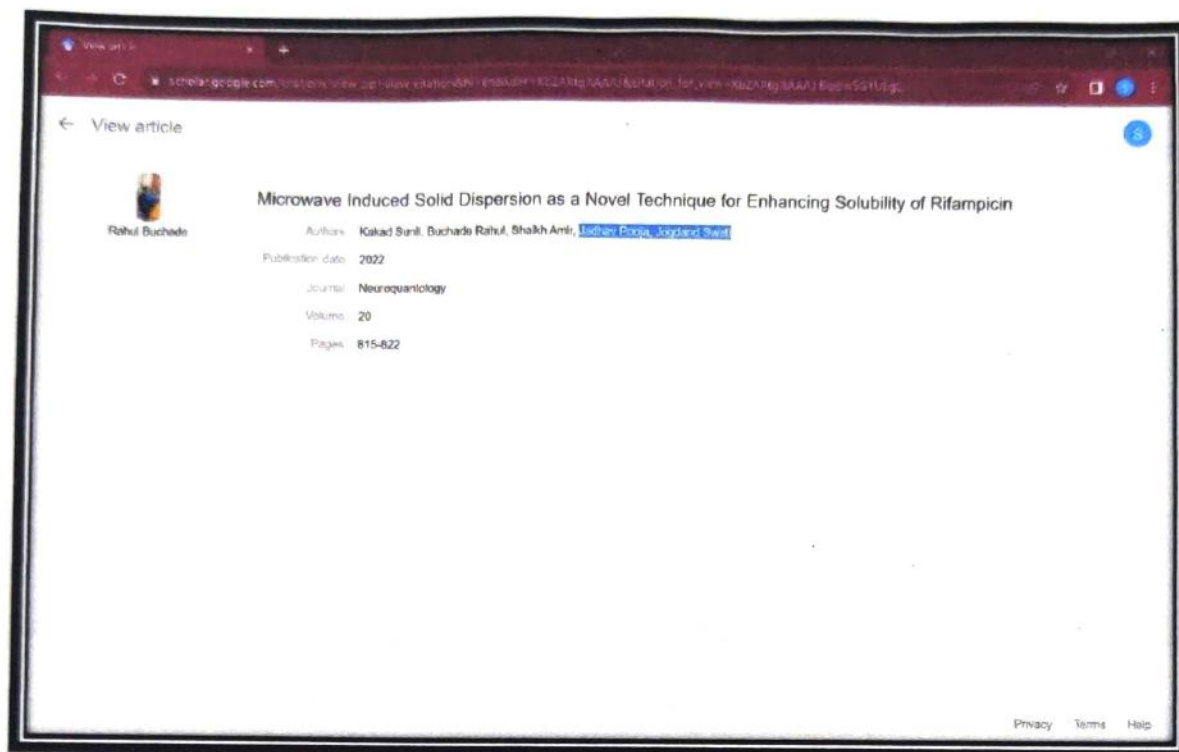
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Formulation development of sustained release swellable matrix tablet of venlafaxine HCl using a combination of hydrophilic and hydrophobic polymers

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Analytical Method Development And Validation Of Niclosamide By RP-HPLC.

PDF

Dr. Narendra Gowekar, Shivam Kale, Dr. Swati Jogdand, Sujata Shinde, Dr. Rahul Dumbre, Dr. Savita Yedav

Keywords: Niclosamide, RP-HPLC, Recovery, Analysis.

Abstract

This research covers the development of an RP-HPLC method for estimating niclosamide. The developed method was verified in terms of Specificity, Accuracy, Linearity, LOD, LOQ and Robustness according to ICH requirements. The precision findings for inter-day and intra-day were excellent enough to demonstrate that the proposed Method was exact and reproducible. The Assay experiment revealed that the niclosamide content measured in the tablet dose was free of excipient interference, indicating that the devised approach was specific. The recovery

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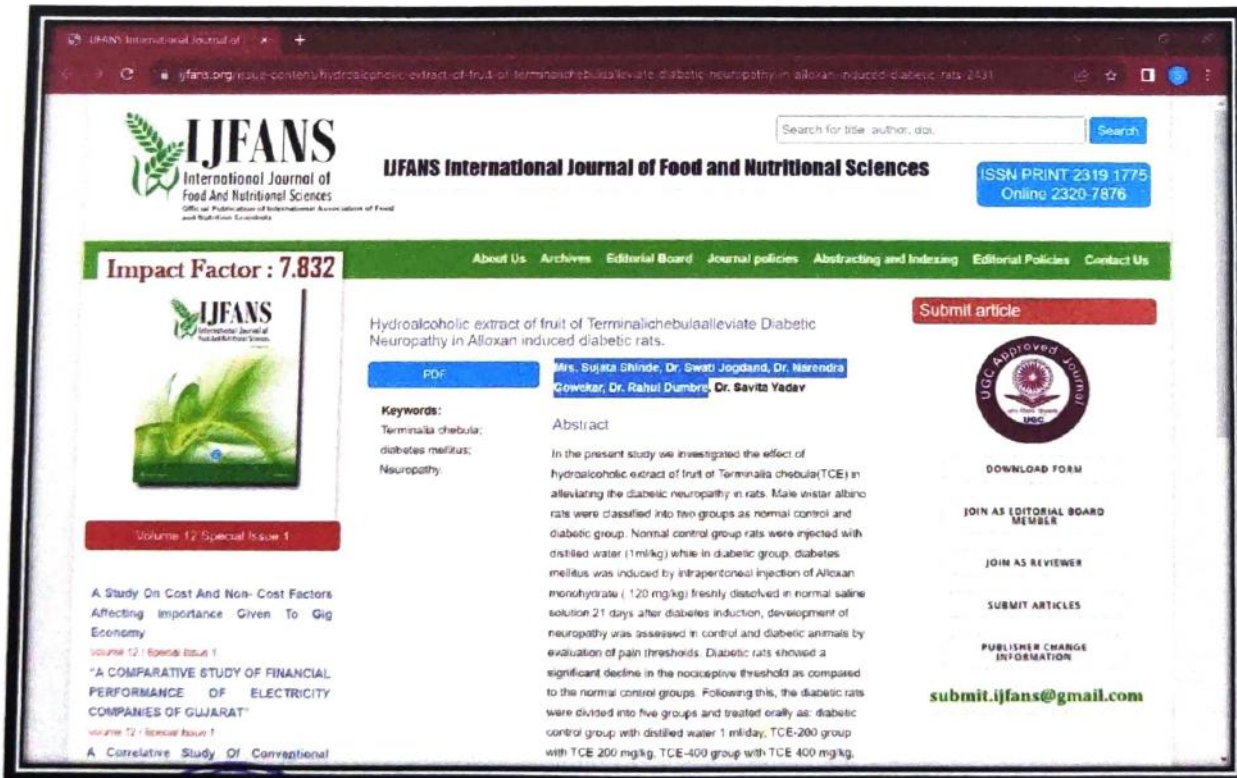
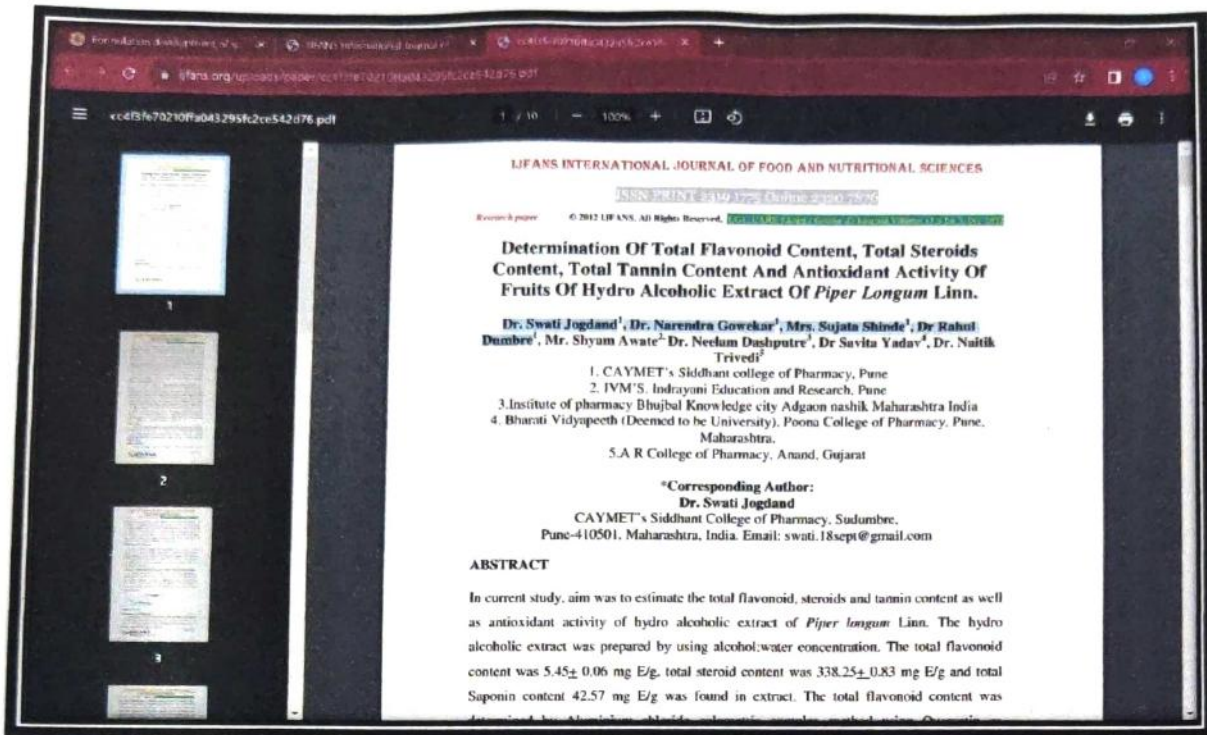
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Research Article | Published: 31 March 2023

Acute and sub-acute toxicity studies of hydro-alcoholic extract of dried fruits of *Piper longum* Linn in Wistar rats

[Swati Vinod Jagtap](#) [Ghanshyam B. Jadhav](#) & [Yogesh P. Talekar](#)

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Abstract

The current study evaluated the prospective toxicity of hydroalcoholic extract of *Piper longum* L. (HEPL) dried fruits with acute and sub-acute oral administration in Wistar rats. During acute toxicity study, female rats were orally administered with HEPL at a single dose of 300 mg/kg and repeated dose of 2000 mg/kg (OECD guidelines-423). Subacute toxicity of HEPL (250, 500, and 1000 mg/kg p.o.) was studied with the control group (1% CMC) by daily dosing in Wistar rats of both sexes for 28 days. To assess reversibility, other satellite acute toxicity study showed no

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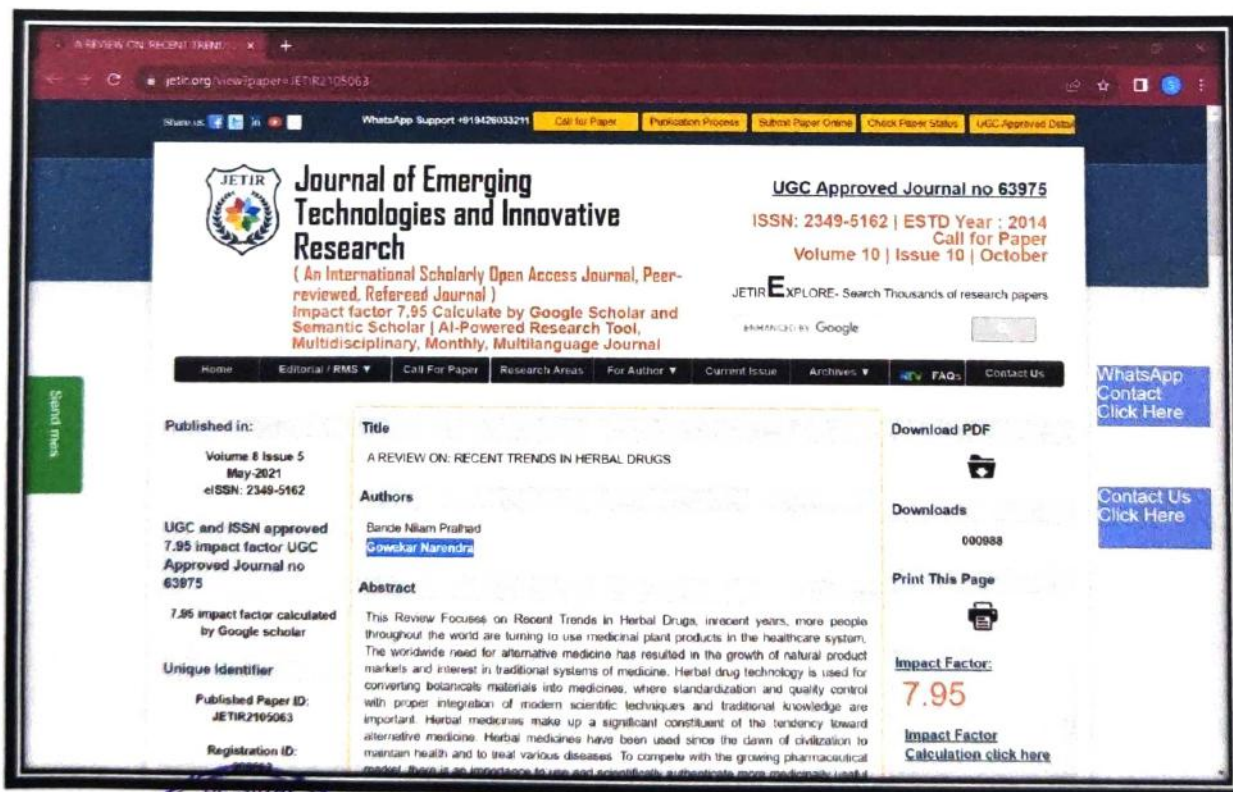
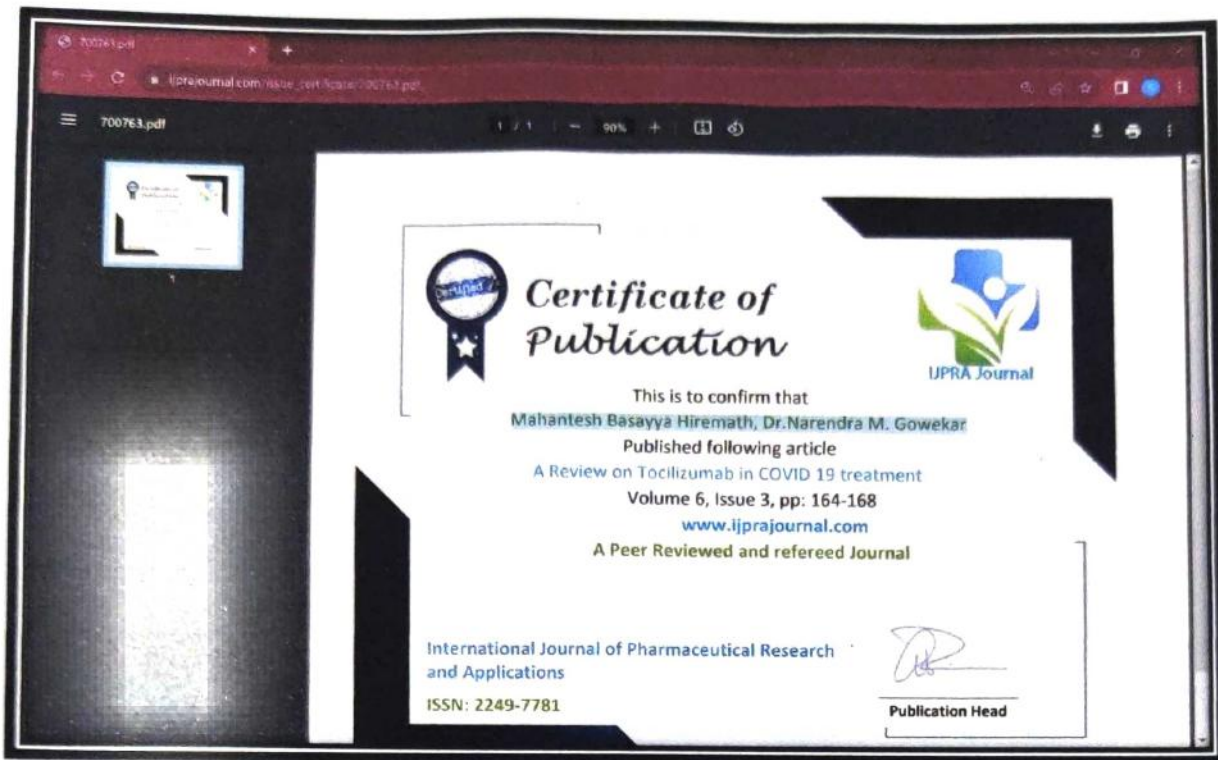
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International Journal of Pharmaceutical Research and Applications
Volume 6, Issue 3 May - June 2021, pp: 596-613 www.ijprajournal.com ISSN: 2249-7781

Review: Medicinal Plants on Anti-Diabetic Activity

Roshan Phanse*1, Dr. Narendra Gowekar*2

*Student, Department of Quality Assurance Siddhant College of Pharmacy, Sudumbare, Pune, Maharashtra, India
**Professor, Department / HOD Of Quality Assurance Siddhant College of Pharmacy, Sudumbare, Pune, Maharashtra, India

Date Of Submission: 15-05-2021 Date Of Acceptance: 26-05-2021

ABSTRACT: Rich herbs for antidiabetic drugs are suggested as a potential but undiscovered potential source. Diabetes Insulin deficiency has been used since ancient times to detect sugar in the blood and urine. Many synthetic substances were found either alive or indirectly from plant sources.

I. INTRODUCTION
Diabetes mellitus According to WHO, the term diabetes mellitus is defined as a metabolic disorder of multiple etiology characterized by chronic hypoglycemia with disturbances of carbohydrate, fat and protein metabolism resulting from defects in insulin secretion, insulin actions or both. The effects of diabetes mellitus include long-term damage, dysfunction and failure of various organs. Diabetes mellitus may have the characteristic symptoms such as thirst, polyuria, blurred vision and loss of weight [1].

Current review of plants with antidiabetic properties. Although many trees it is recommended that further pharmacological and chemical research be carried out to clarify the exact mechanism Hypoglycaemic action.
Keywords: diabetes mellitus, herbs, antidiabetic destruction of beta cells in type-1 diabetes is caused by islet cell autoantibodies, insulin directed antibodies, autoantibodies towards Glutamic Acid Decarboxylase or GAD or antibodies directed towards the tyrosine phosphatases. When diagnosed with Type-1 diabetes, multiple antibodies towards pancreatic cells are detected in the patient's blood. Association with HLA gene is also suspected to act as a predisposing factor for the cause of this type of diabetes. Destruction of Beta cells in type-1 diabetes may vary from patient to patient. Destruction may be rapid in some individuals as in case of infants and children or

Review Topical Drug Delivery System.pdf

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Review: Topical Drug Delivery System

Dhanshree Kanawade*1, Dr. Narendra Gowekar*2

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Date Of Submission: 15-05-2021 Date Of Acceptance: 26-05-2021

ABSTRACT: Clinical evidence suggests that advanced gel is the safest and most effective treatment option for use in skin-related disease management and is used for topical measures to reduce side effects associated with other common dose forms. Advanced drug delivery systems include a large variety of drug dose forms such as semisolid, liability preparations, emulsion and solid powder. Many of the widely used semisolid preparations for distribution of advanced drugs include gels, creams and ointments. Gel is a polymer network connected to the inflammatory polymer in the middle of the fluid its assets rely greatly on the interaction between a solid government polymer and the liquid part. Gel does not show the continuous flow of the government. The conversation between the breakdown of polymer sand liquid forms a three-dimensional network interacting with scattered phase particles. The forms, attempts has being made to utilize drug carriers that ensure adequate localization or penetration of the drug within or through the skin in order to enhance the local and minimize the systemic effects, or to ensure adequate Percutaneous absorption. Topical preparation prevents the GI-irritation, prevent the metabolism of drug in the liver so as increase the bioavailability of the drug. Topical preparations give its action directly at the site of action. A gel is a two-component, cross linked three-dimensional network consisting of structural materials. The structural materials that form the gel network can be composed of inorganic particles or organic macromolecules, primarily polyoxines.

U.S.P. defines gels as a semisolid system consisting of dispersion made up of either small inorganic particle or large organic molecule enclosing and interpenetrated by liquid. Gels



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Abstract

FORMULATION AND EVALUATION OF FLUCONAZOLE TOPICAL GEL BY USING OCIMUM BASILICUM SEED MUCILAGE AS A GELLING AGENT
 Nitesh Nitin Sancheti¹ and [Dr. Pratima Shinde](#)

ABSTRACT
 Fluconazole is an imidazole derivative used for the treatment for local and systemic fungal infection. The oral use of fluconazole is not recommended as it has many side effects. The present study was design to formulate and evaluate different formulae of topical gel containing fluconazole for treatment of fungal infection of skin. The gel was formulated by using natural seed mucilage obtained from ocimum basilicum with different concentration of acacia and glycerin. Three different formulae were prepared and characterized physically in term of color, spreadability, pH, and drug content. In-vitro drug release in phosphate buffer pH 6.8 and permeation study through cellulose membrane using a modified Franz diffusion cell were performed. Candida albicans was used as model fungus to evaluate the antifungal activity of prepared formulae. The results of in-vitro drug release and its permeation studies showed that the highest values was form F2 (80.18% of drug release after 4 hr) also F2 shows highest antifungal activity.

Keywords: Fluconazole, Ocimum Basilicum, Glycerin, Acacia.

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Abstract

REVIEW ON FORMULATION OF MICROEMULSION AS NOVEL DRUG DELIVERY SYSTEM
 Prathamesh P. Kulkarni¹ and [Pratima S. Shinde](#)

ABSTRACT
 Micro-emulsion is defined as the isotropic thermodynamically stable transparent system of oil water surfactant and co surfactant which having very small droplets size (20-200nm). Microemulsions are isotropic, thermodynamically stable transparent (or translucent) systems of oil, water and surfactant, frequently in combination with a cosurfactant with a droplet size usually in the range of 20-200 nm. They can be classified as oil-in-water (o/w), water-in-oil (w/o) or bicontinuous systems depending on their structure and are characterized by ultra low interfacial tension between oil and water phases. To date microemulsions have been shown to be able to protect labile drug, control drug release, increase drug solubility, increase bioavailability and reduce patient variability. Furthermore, it has proven possible to formulate preparations suitable for most routes of administration. Since the discovery of microemulsions, they have attained increasing significance both in basic research and in industry. Due to their unique properties, namely, ultralow interfacial tension, large interfacial area, thermodynamic stability and the ability to solubilise otherwise immiscible liquids, uses and applications of microemulsions have been numerous. Microemulsions are readily distinguished from normal emulsions by their transparency, low viscosity and more fundamentally their thermodynamic stability. Microemulsions are shown to be effective dermal delivery mechanism for several active ingredients for pharmaceutical and cosmetic applications. Topical microemulsions allow rapid penetration of active molecules due to the large surface area of the internal phase, and their components reduce the barrier property of stratum corneum. Microemulsions thereby enhance dermal absorption compared with conventional formulations and are therefore a promising vehicle due to their potential for transdermal drug delivery.

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REVIEW OF COMPARATIVE STUDY OF OTF WITH CONVENTIONAL DRUG DELIVERY SYSTEM

*Shravani A. Havale and [Dr. Prabha S. Shinde](#)

ABSTRACT


Fast dissolving oral films (FDOFs) are the most advanced form of oral solid dosage form due to more flexibility and comfort. It improves the efficacy of APIs by dissolving within minutes in the mouth after contact with less saliva as compared to fast-dissolving tablets, without chewing and no need of water for administration. The FDOFs place as an alternate within the market thanks to the consumer's preference for a fast-dissolving product over conventional tablets / capsules. The oral thin film technology remains within the beginning stages and has bright future ahead because it fulfills all the necessity of patients. Eventually, film formulations having drugs are going to be commercially launched using the oral film technology. However, for future growth point of view the oral thin film sector is well-positioned. In US market the OTC films of pain management and kinestosis are commercialized. More importantly, prescription OTFs have now been approved in US, EU and Japan which are the three major regions. These approved Rx films, have potential to dominate over other oral dosage forms of the same drugs. It seems that the worth of the general oral thin film market will grow significantly.

Keywords: Oral thin film (OTF), Conventional drug delivery, Oral disintegrated formulation.

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HUTCHINSON – GILFORD PROGERIA SYNDROME

[Apeksha S. Pangare](#) and [Dr. Prabha S. Shinde](#)


ABSTRACT

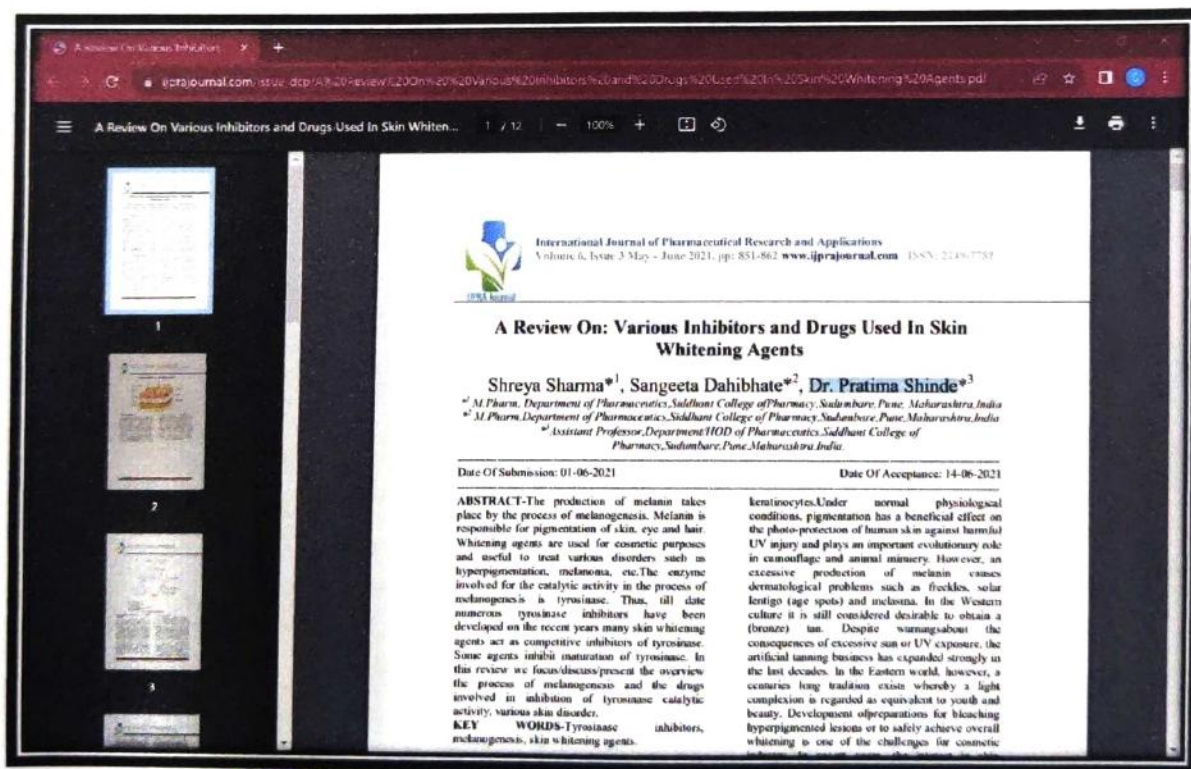
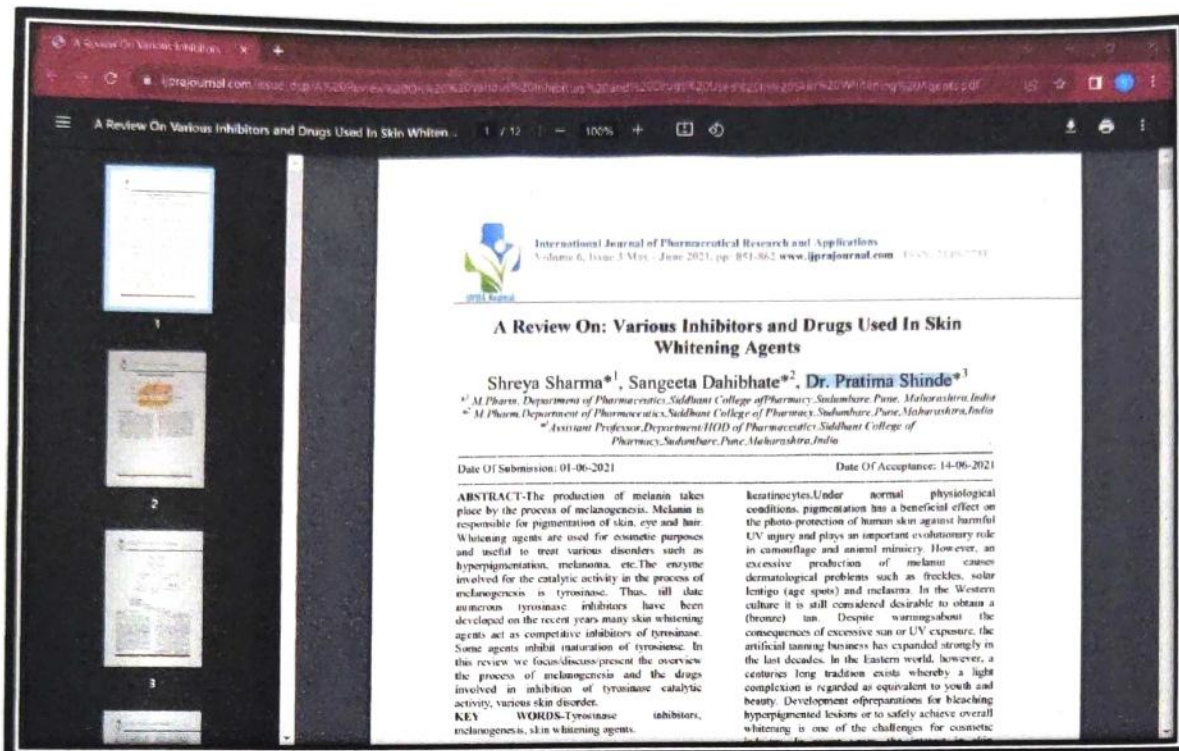
Hutchinson – Gilford Progeria Syndrome (HGPS) is an Autosomal dominant, rare, fatal pediatric segmental pre-mature aging disease. Hutchinson – Gilford Progeria Syndrome, a rare genetic disorder associated with a characteristic aged appearance very early in life. The review on Hutchinson – Gilford Progeria Syndrome summarizes the clinical characteristics of this disease and therefore the underlying mutation within the lamin A (LMNA) gene that leads to produce abnormal lamin A. The Progerin; this disrupts the nuclear membrane and alters transcription. Accumulation of this progerin proteins in the normal cell which leads to nuclear morphology defects, decreased lifespan and premature cell death occurs. Based on the positive outcomes from the studies based on statins, aminobisphosphonates and FTIs, two clinical trials have been performed in children with HGPS.

Keywords: Hutchinson – Gilford Progeria Syndrome, Progerin Protein, Lamin A/C gene

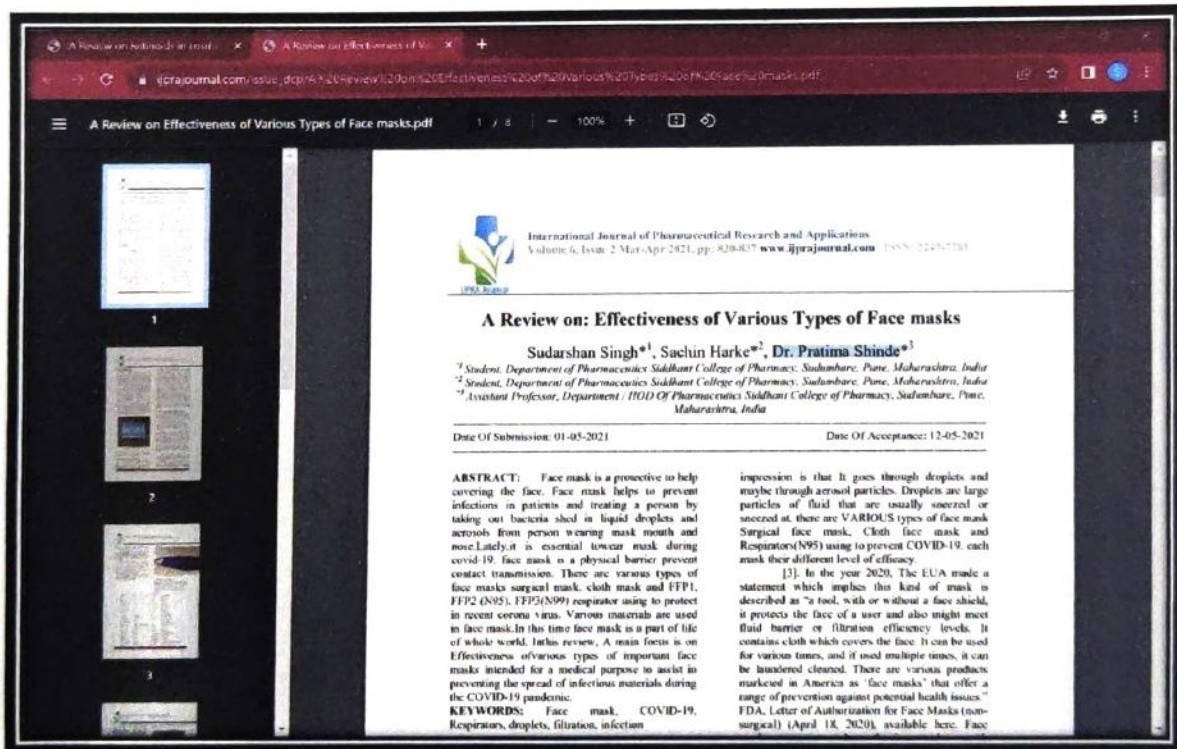
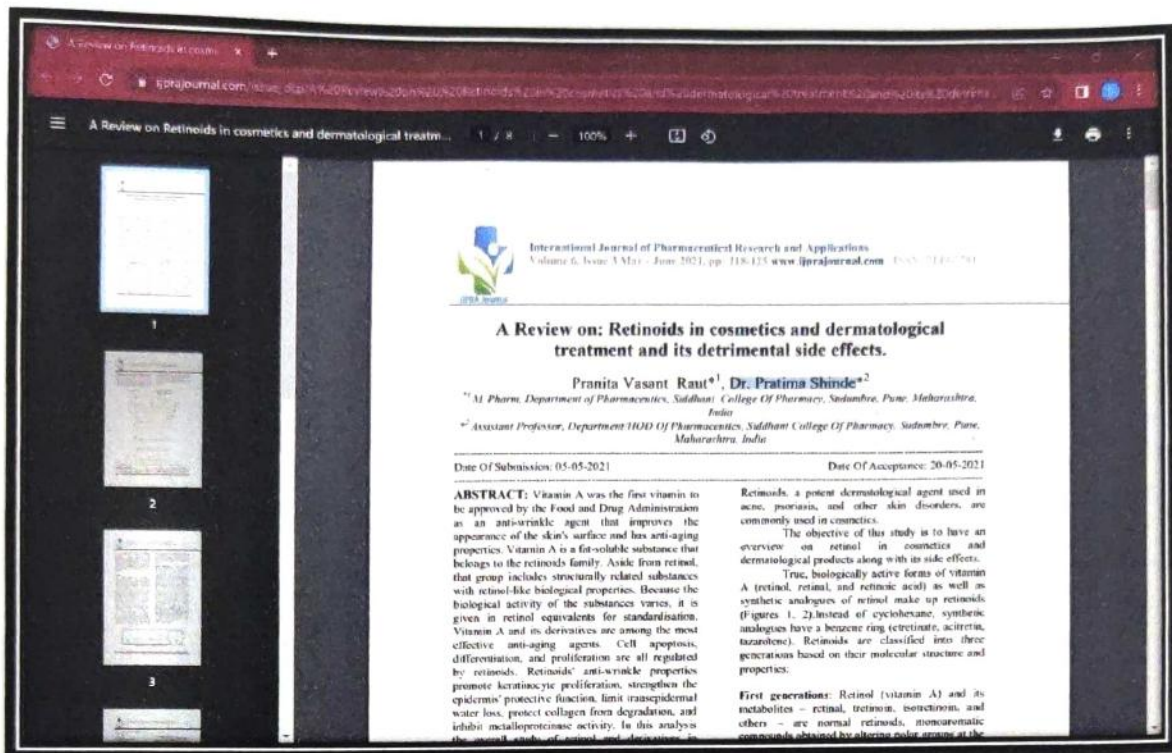
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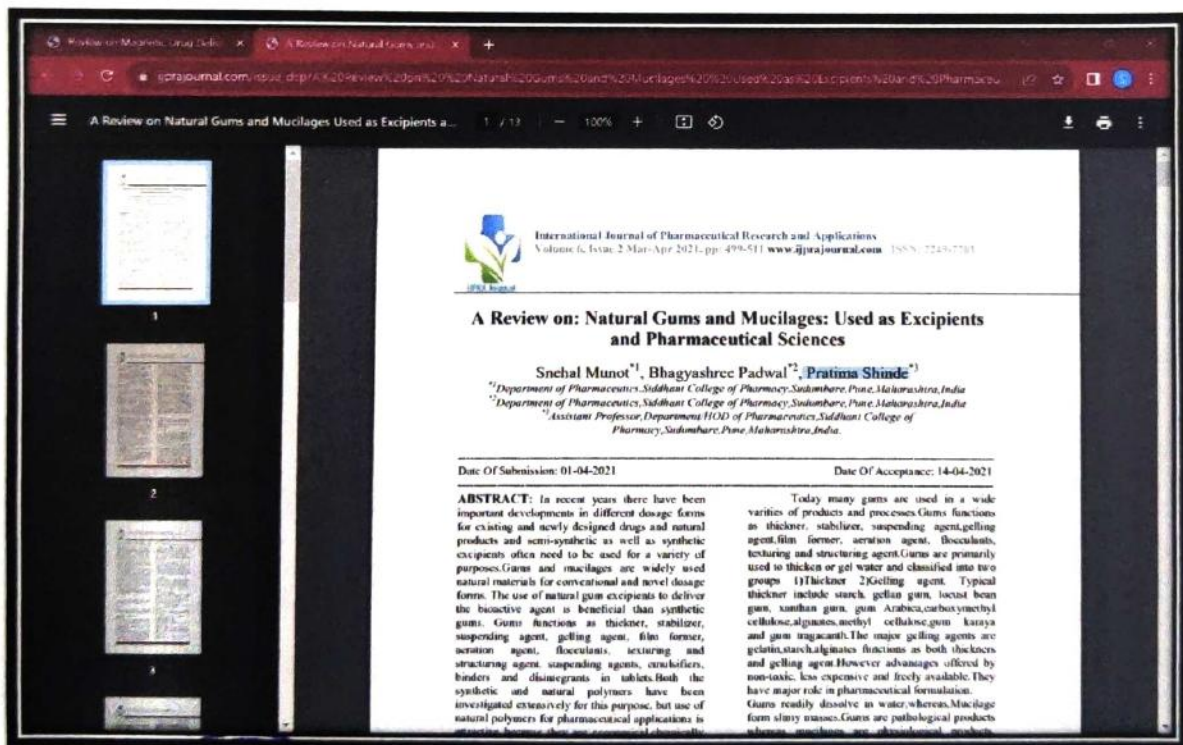
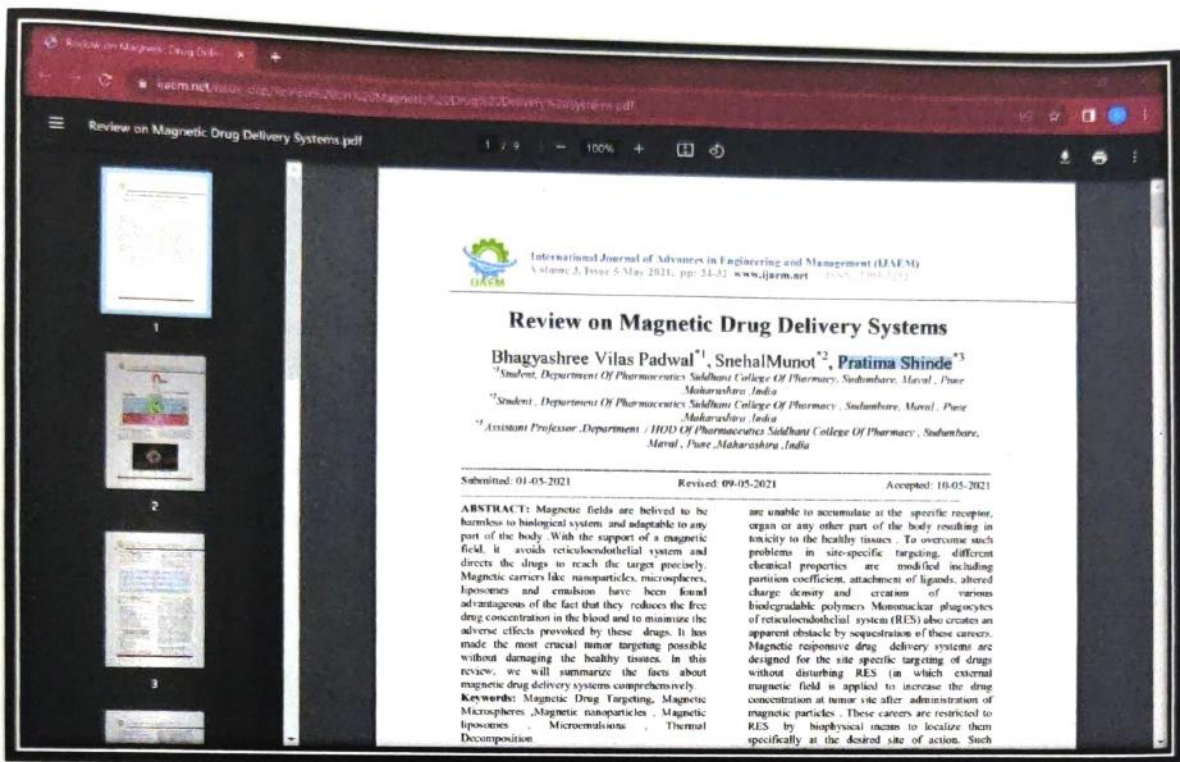

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Abstract

COMPARATIVE STUDY OF HPLC AND UPLC AN OVERVIEW
Pooja S. Kahade* and [Vikas Kanekar](#)

ABSTRACT
The Liquid Chromatographic procedures HPLC and UPLC are both used to separate the components of a substance. To understand the differences between HPLC and UPLC, we must first understand HPLC. This is due to the fact that UPLC is a variant of HPLC. Recent advances in pharmaceutical analysis have made chromatographic media with a 1.7 μm particle size, as well as a liquid handling system capable of operating such columns at significantly higher pressures, available. Ultra performance liquid chromatography (UPLC), which uses sub 2 micron particles and extremely high pressure (up to 100 MPa in a UPLC system), has shown to increase method sensitivity, resolution, and speed when compared to traditional HPLC. When compared to chromatographic systems using 5 μm size particle loaded analytical columns, the UPLC system can reduce analysis time by up to nine times. The analysis time was reduced by nearly three times when compared to 3μm particle packed analytical columns. The current review paper distinguishes between HPLC and UPLC, as well as analytical method validation, applications, and HPLC and UPLC advantages and disadvantages.

Keywords: Ultra performance liquid chromatography, High performance liquid chromatography, sensitivity, BEH technology, HSS Technology.

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Abstract

FORMULATION AND EVALUATION OF INDOMETHACIN PARACETAMOL FAST DISSOLVING TABLET
Sourabh Dilip Khalekar*, Bhagyshri Satish Rupenwar, Gopal K. Munde and [Prof. Vikas B. Kanekar](#)

ABSTRACT
Fast dissolving tablet Indomethacin with Paracetamol has been shown anti-inflammatory analgesic activity COX-2 inhibitor. Convenient mode of administration, there is need to develop fast disintegrating dosage form, particularly one that disintegrates and dissolve when contact with saliva and can be administered without water. Indomethacin paracetamol were prepared by direct compression method by using superdisintegrant croscopolvidone, sodium starch glycolate, mannitol, gum karaya, banana powder to enhance patient compliance prepared batches of tablet were evaluated for hardness, friability, drug content uniformity, wetting time, water absorption ratio and in vitro dispersion time, drug release, stability. Analytical technique used like FTIR, DSC and dissolution study. In vitro dissolution study carried out disintegration approach phosphate buffer PH 6.8 dissolution medium of F4 batch comparison gum karaya and banana powder shows better drug release. Post compression parameter like hardness 3.2kg/cm², stability 0.41 IPF4 showed improved drug dissolution 96.49% in 30 min and in-vitro dispersion 67 seconds. The F4 was the best among all formulation of fast dissolving tablet. The FTIR and DSC showed no interaction between the drug excipients. Optimized batch IPF4 showed good drug release.

Keywords: Croscopolvidone, Sodium starch glycolate, Mannitol, Banana powder, Gum karaya.

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Maribavir Antiviral Agent Overview for Treatment of Cytomeg...

International Journal of Pharmaceutical Research and Applications
Volume 7, Issue 3 May-June 2022, pp: 576-582 www.ijprajournal.com

Maribavir: Antiviral Agent Overview for Treatment of Cytomegalovirus

Jidnyesh P. Gharat*, Mr. Vikas Kandekar*

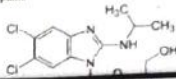
*1 Mpharm, Department of Quality Assurance Techniques, Siddhant College Of Pharmacy, Sudumbare, Pune, Maharashtra, India.
*2 Professor, Department of Quality Assurance Techniques, Siddhant College Of Pharmacy, Sudumbare, Pune, Maharashtra, India.

Submitted: 15-05-2022 Revised: 20-05-2022 Accepted: 25-05-2022

ABSTRACT:
Infections with the cytomegalovirus (CMV) are a frequent complication in solid organ transplant (SOT) and hematopoietic stem cell transplant (HSCT) recipients, increasing morbidity and death. Although currently available treatment approaches have lowered infection burdens, their use is constrained by side effects such as nephrotoxicity and/or myelosuppression, as well as the emergence of resistance. It is critical to expand our present arsenal against CMV infection. Here, we look at maribavir, an emerging medicine, and its safety and efficacy in the prevention and treatment of CMV infections, including resistant/refractory illness.

Keywords: Maribavir, Cytomegalovirus, efficacy, safety.

satisfy study goals. However, the Phase III trial's dosage may have been too low to be effective. Maribavir prophylaxis showed good antiviral effect in a Phase II research, as indicated by a statistically significant reduction in the rate of CMV reactivation in patients of hematopoietic stem cell bone marrow transplantation. The number of participants who required pre-emptive anti-CMV medication was statistically significantly lower with maribavir compared to placebo in an intent-to-treat analysis of the first 100 days following the transplant.



INTRODUCTION

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A COMPREHENSIVE REVIEW ON LIPOSOMES: A NOVEL DRUG DELIVERY SYSTEM

Tejasvini Vasant Anandrao¹ and Vikas Kandekar²

¹Student, Department of Quality Assurance, Siddhant College of Pharmacy, Sudumbare Maharashtra India.
²Professor of Quality Assurance, Siddhant College of Pharmacy, Sudumbare Maharashtra India.

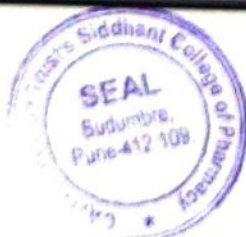
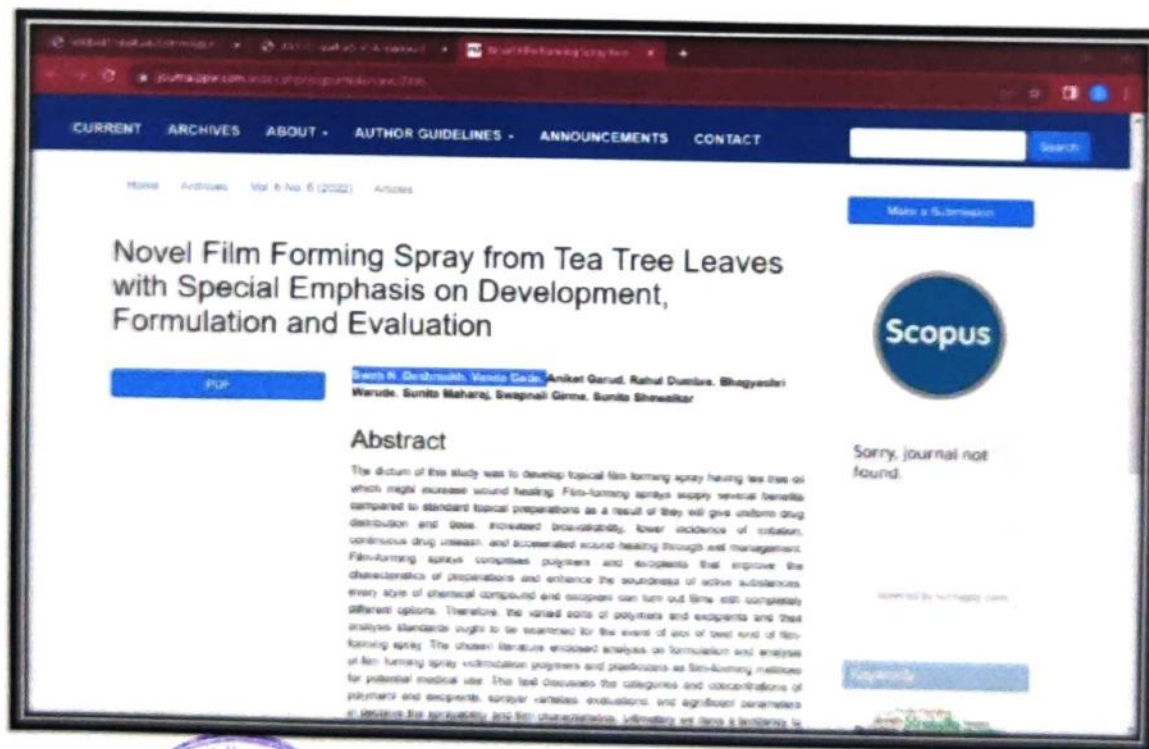
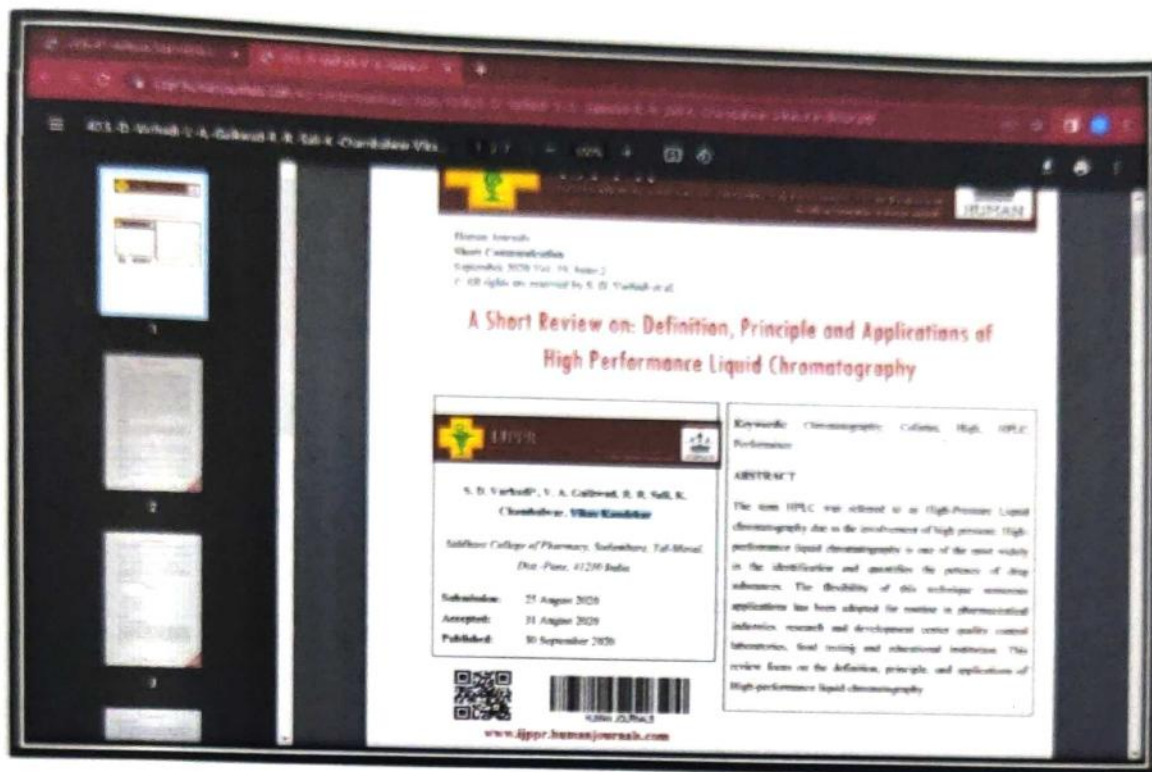
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DOI: 10.20959/wjpr.2022.26345

***Corresponding Author**
Tejasvini Vasant Anandrao
Student, Department of Quality Assurance, Siddhant College of Pharmacy, Sudumbare Maharashtra India.

ABSTRACT
The function of belayed vesicles as productive transporters for drugs, immunizations, indicative specialists, and other bioactive operators has prompted a fast headway in the liposomal drug conveyance system. The pharmacy- elements and pharmacokinetics properties are altered for the liposomal delivery system, which on the whole leads to an increased therapeutic index with decreased toxicity. The liposome can be named multilamellar vesicles or unilamellar vesicles, which can be additionally named large unilamellar vesicles (LUV) or small unilamellar vesicles (SUV). The part of liposome as a medication



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Abstract

PRINCIPLE, FORMULATION, CHARACTERIZATION AND APPLICATION OF NANOFIBERS IN PHARMACEUTICALS-A REVIEW
 Rabhya Patel* and Swati Deshmukh

Abstract
 Conventional oral dosage forms exhibit poor/bioavailability due to incomplete release of drug and short residence time at the absorption site. The benefits of the fibrous carriers are site specific delivery of drugs to the body, nanofibers are an exciting new class of material produced using an innovative manufacturing process technology. At present, there are three techniques available for the synthesis of Nanofibers: Electro spinning, self-assembly, and phase separation, out of these Electro spinning is the most widely used technique. Electrospinning has been recognized as an efficient technique for the fabrication of polymer nanofibers. Various polymers have been successfully electrospun into ultrafine fibers in recent years mostly in solvent solution and some in melt form. In this article, we discussed principle, process of manufacturing, characterization, application of nanofibers.

Keywords: nanofiber, Electrospinning, Tissue engineering etc.

[Full Text Article]

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Nanogels: An overview of properties, biomedical applications, future research trends and developments

Rabhya Patel, Swati Deshmukh,
 CAYMETS Siddhant College of Pharmacy, Pune.

Abstract- A nanoparticle which is composed of a hydrogel with a cross linked hydrophilic polymer network is known as "Nanogel". The term "nanogels" defined as the nanosized particles formed by physically or chemically cross-linked polymer networks that is swell in a good solvent. Nanogel is a nanoscale drug delivery systems with enhanced surface area not only are capable of transmitting hydrophilic drugs in vitro and in vivo with improved drug bioavailability but also help to reduce side effects of drug. Here we summarize emerging research of nanogels for biomedical applications and provide an overview of the state-of-the-art, recent developments as well as emerging trends in the field of nanogel. Nanogels have enabled enlargement of functionalized nanoparticles, which act as a drug carriers that can be loaded with drugs and other active material to be released in a controlled manner at specific site.

Keywords- Nanogel, controlled drug release

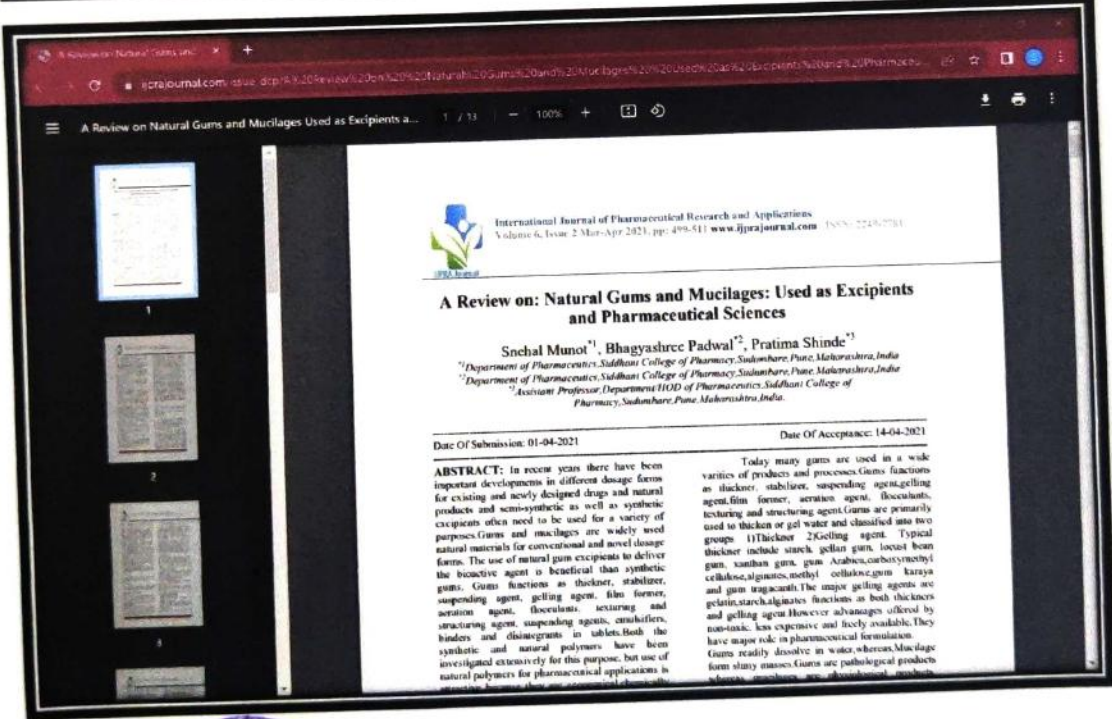
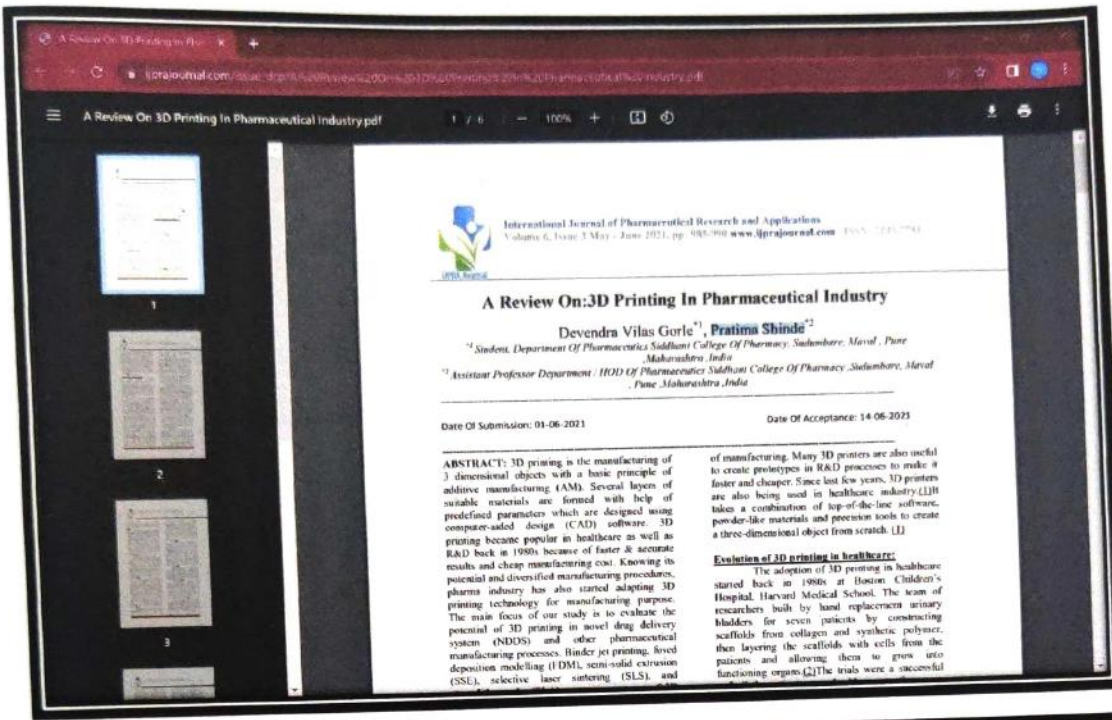
Introduction-
 Nanogels are currently considered as promising nanosized drug delivery carriers. Nanogels are made of a crosslinked polymeric network which could encapsulate both hydrophilic and hydrophobic drugs due to their tunable nature. The ability of nanogels to control drug release is vastly described in the literature and researchers are constantly improving the control of drug release from nanogel by designing new polymers having specific sensitivity to a chemical or physical stimulus. In this review, we briefly discuss the definition of nanogels, their release profiles, their specific gel-based characteristics and the pathways of drug release from nanogels, we have focused on the stimuli responsive nanogels and their release profile. This compilation opens the window for understanding the influence of chemical composition and design of various nanogel on their release in the presence and absence of corresponding stimuli such as temperature, pH, enzymes and others. The uniqueness of this review is that it highlights the data of release profiles in terms of the different nanogel composition and triggers. It also points the high potential of nanogels in the list of candidates for drug delivery systems, thanks to their properties regarding drug encapsulation and release, combined advantages of nano-size and swelling characteristics of hydrogel.

Advantages of nanogel

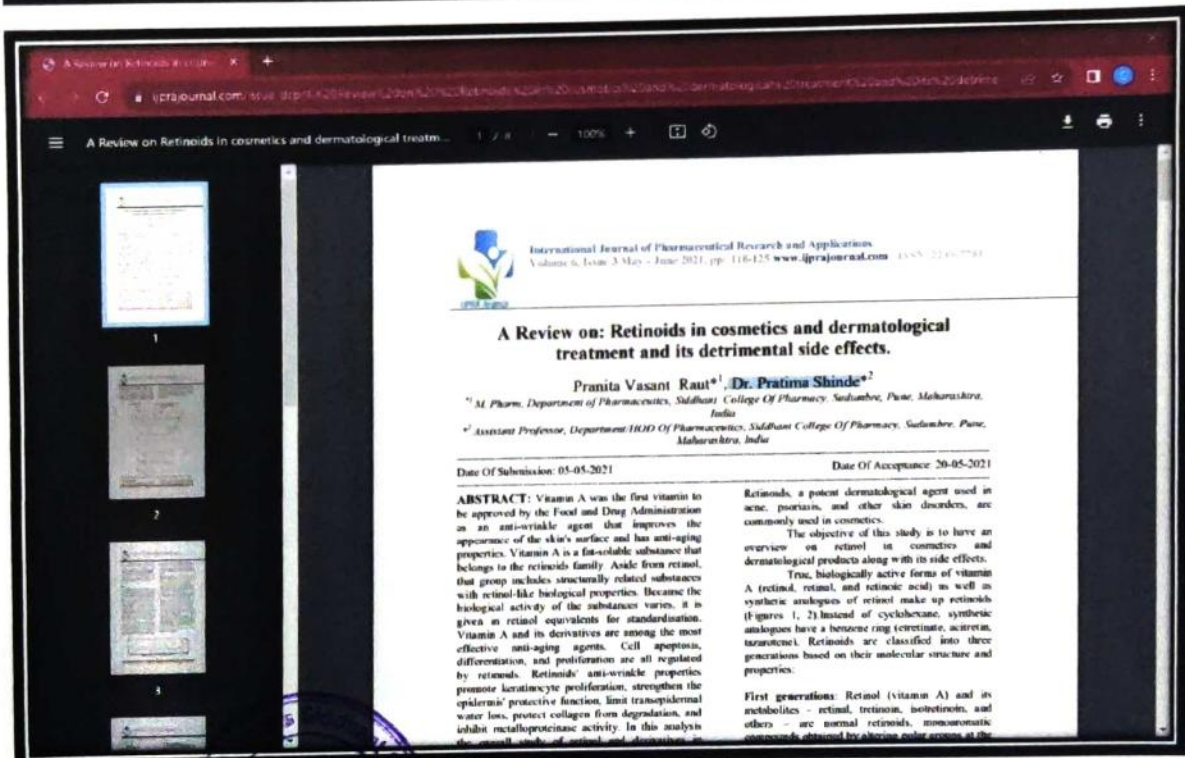
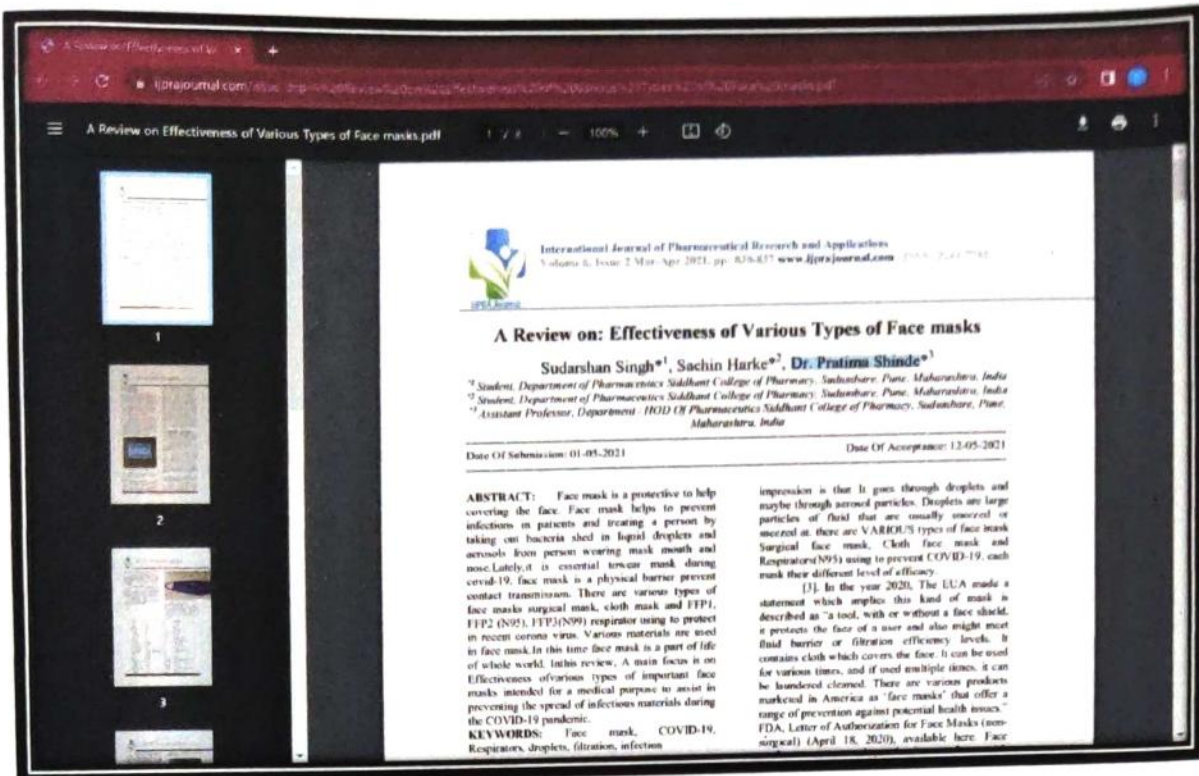
- Nanogels occur with high biocompatibility and biodegradable formulation.
- Nanogels can be controlled for sustained release of drug from the formulation by the addition of apolymeric network. Polymeric networks also control the particle size of the formulation⁽¹⁾.
- The free-flowing pearlescent solution of the nanogels is easily dispersed in aqueous media⁽²⁾.
- Nanogels can be easily administered in parenteral and mucosal administration⁽³⁾.
- The biggest advantage of nanogels is reduced preservatives, leakage of the drug from the solution⁽⁴⁾.



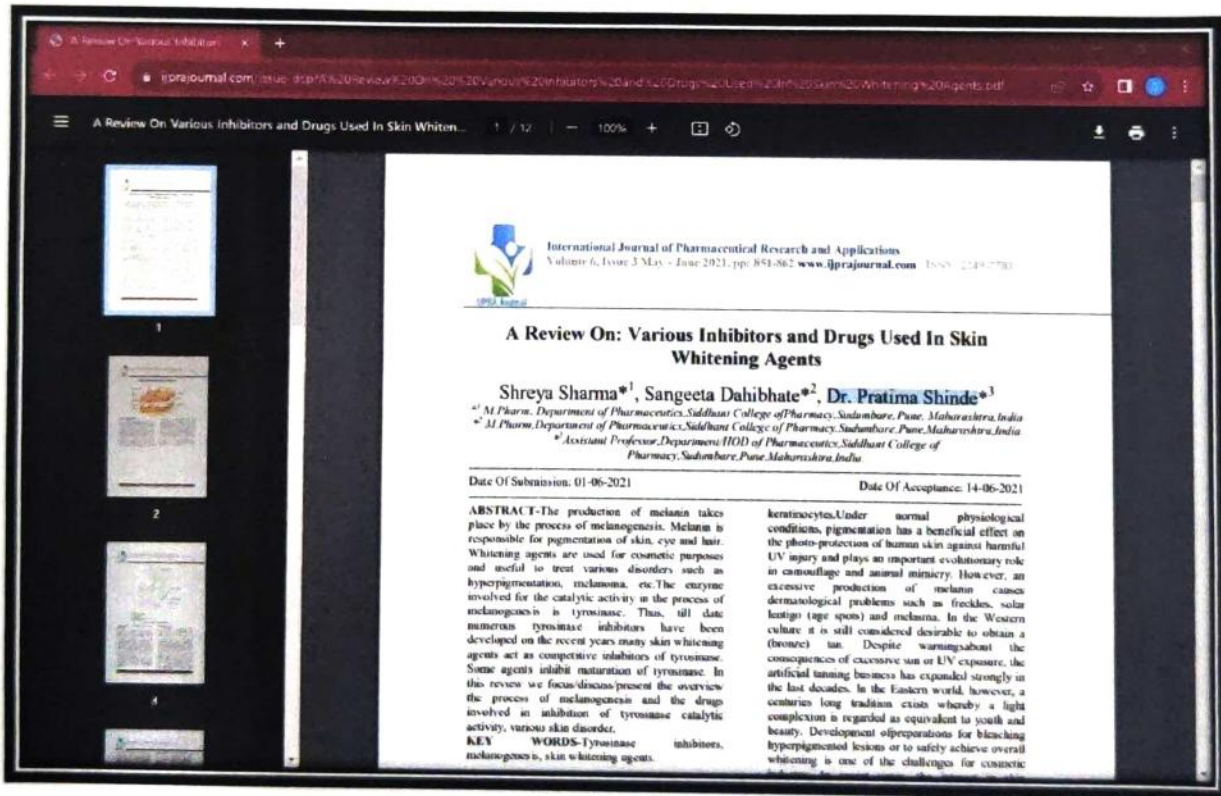
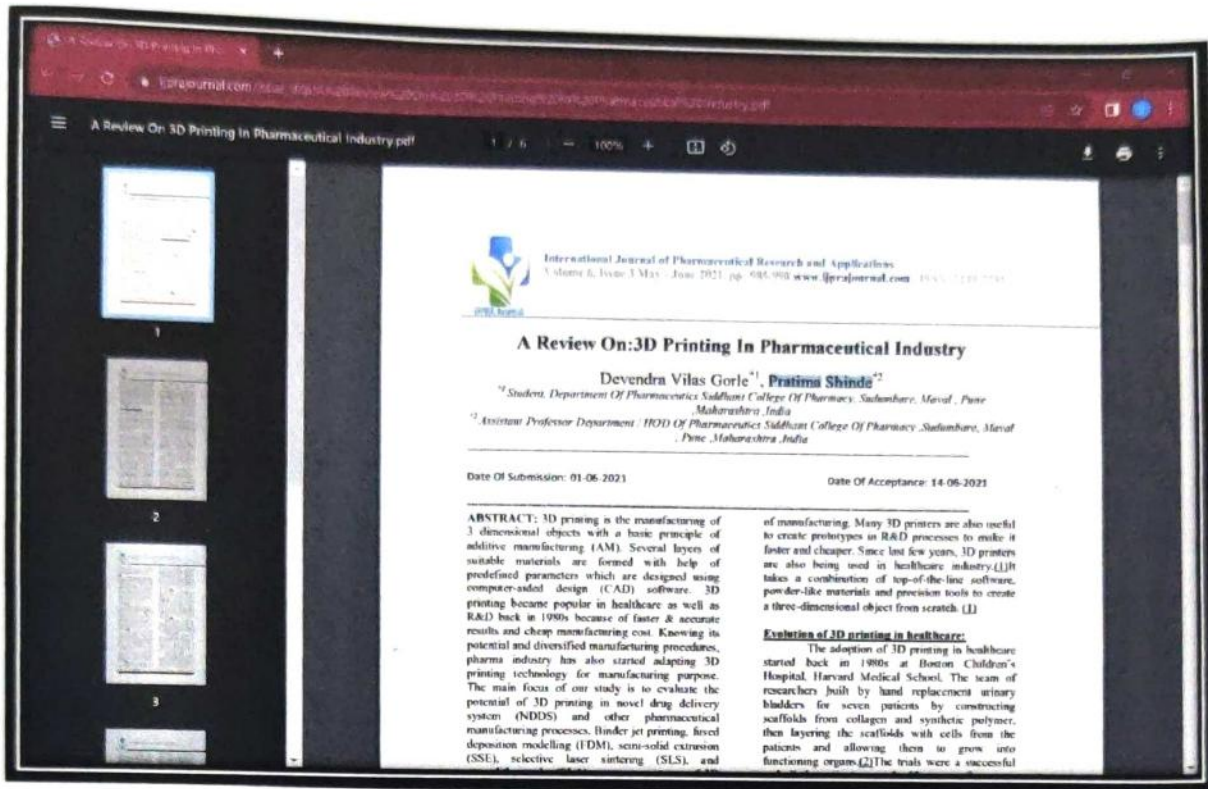
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Title
 RP-HPLC METHOD DEVELOPMENT AND VALIDATION FOR ESTIMATION OF
 CHLORPHENIRAMINE MALEATE IN LIQUID DOSAGE FORM

Authors
 Suchitra Rane
 Dr. R.R. Dumbre
 Dr. Vikas Karkdekar

Abstract
 A high performance thin layer chromatographic method was developed and validated for determination of Chlorpheniramine maleate which comply with ICH Guidelines. The chromatographic development was performed using a Hypersil CN C18 (250x4.6mm, 5.0µm chromatographic particle size) and a mobile phase composed of sodium dihydrogen phosphate buffer-Acetone:nitric Methanol (35:34:31 v/v/v), pH 6.2 adjusted with triethylamine at 1.3 ml/min flow rate. The retention factor (Rf) was found to be 7.65 min. The linear calibration curves at concentration range 80-120% (r²=0.9999) for Chlorpheniramine maleate.

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Title
 Development and validation of analytical method for determination of bioactive drug in bulk and pharmaceutical dosage form

Authors
 Dipika marathe
 Vikas Karkdekar

Abstract
 To Develop and validate FTIR method for the simultaneous estimation of drug and Pharmaceutical Dosage form. Contain objective To develop FTIR method for the estimation of bulk Drug and pharmaceutical dosage form. And To validate above developed methods for estimation of bulk Drug and pharmaceutical dosage form. The Principle of Fourier-transform infrared spectroscopy is a technique used to obtain an infrared spectrum of absorption or emission of solid, liquid or gas. An FTIR Spectrometer simultaneously collects high-spectral-resolution data over a wide spectral range. In Method Validation study Linearity, Precision, Accuracy, Limit of Detection and Limit of Quantitation, Analytical Chemistry may be defined by the process and set of circumstances the composition of material in terms of the elements of



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Title: Development And Validation of stability indicating method for Raltegravir potassium.

Authors: Ruzesh R. Barde, Dr. Narentra Gorekar

Abstract: ABSTRACT Raltegravir potassium is an antiretroviral drug. A simple and rapid stability indicating HPLC method of Raltegravir potassium was successfully developed. This method is based on HPLC separation followed by UV detection at 315 nm. HPLC method was developed on a symmetry HICol C18 (150 x 4.6mm, id 5 µm) column with a mobile phase consisting of Methanol: Phosphate buffer: Acetonitrile 40:30:30 % v/v/v, pumped at 1.0 ml min⁻¹ flow rate. The pH of buffer was adjusted to 3.0 with ortho phosphoric acid. The column was maintained at ambient temperature and 20 µl of solutions were injected. The eluted compounds were detected by using PDA detector. Raltegravir potassium eluted at 3.3 ± 0.2 min. Stress degradation study shows that sample degraded with acid and base hydrolysis, oxidative, thermal and photolytic stress conditions.

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Title: A REVIEW ON RECENT TRENDS IN HERBAL DRUGS

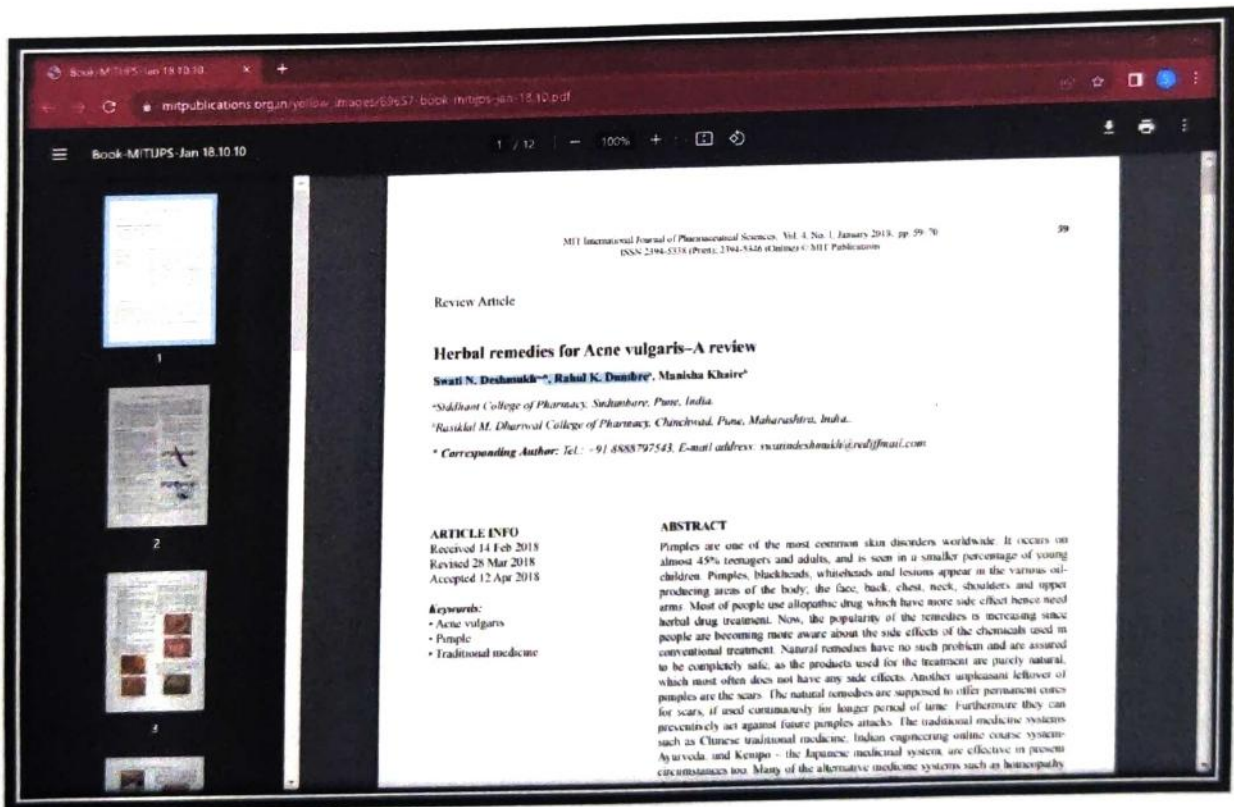
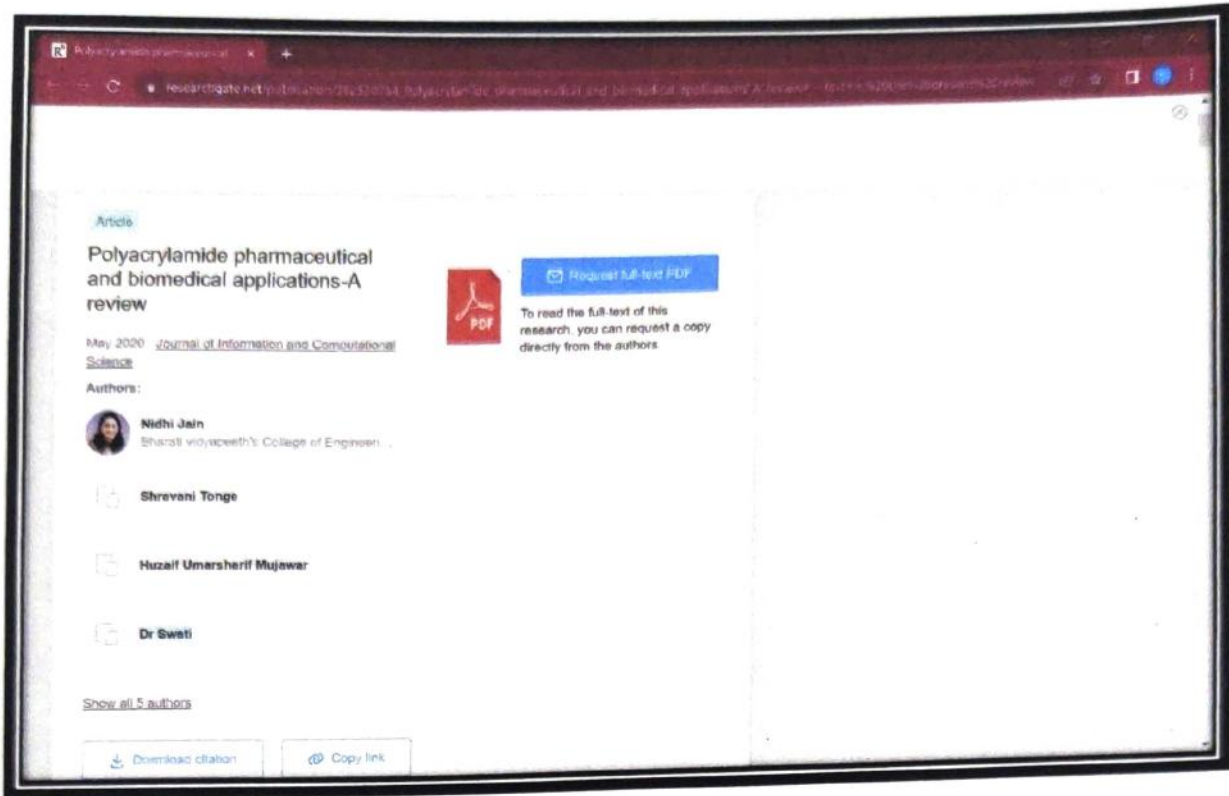
Authors: Bando Nilam Prahad, Gowkar Narentra

Abstract: This Review Focuses on Recent Trends in Herbal Drugs. In recent years, more people throughout the world are turning to use medicinal plant products in the healthcare system. The worldwide need for alternative medicine has resulted in the growth of natural product markets and interest in traditional systems of medicine. Herbal drug technology is used for converting botanicals materials into medicines, where standardization and quality control with proper integration of modern scientific techniques and traditional knowledge are important. Herbal medicines make up a significant constituent of the tendency toward alternative medicine. Herbal medicines have been used since the dawn of civilization to maintain health and to treat various diseases. To compete with the growing pharmaceutical market, there is an impetus to use and scientifically authenticate more medicinal plants.

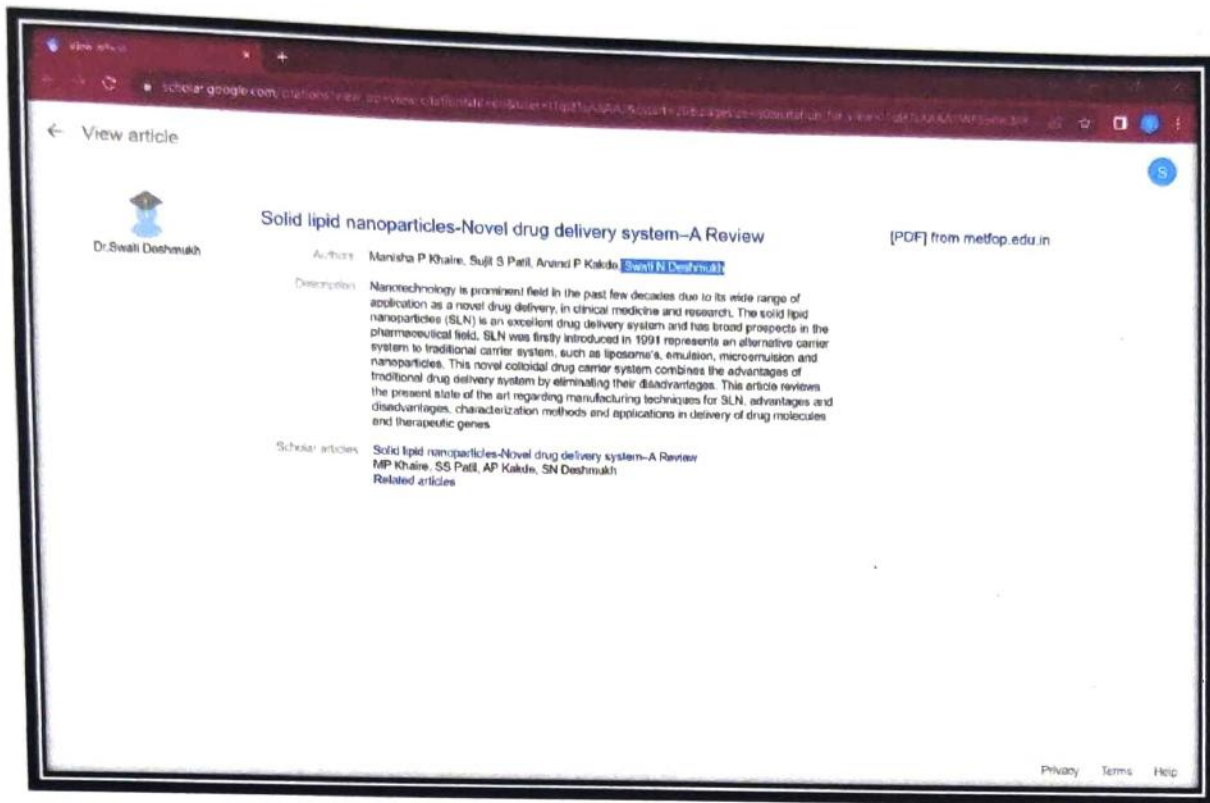
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