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Development and

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Medoxomil ...  
Research Article .  
Int. J. Pharm. Sci.  
Rev. Res., 51(2),  
July - August 2018;  
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Pages: 110-115  
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Doshi MET Institute  
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lipid nanoparticles  
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using one type of  
lipid (homolipid)

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Rev. Res., 63(1),  
July - August 2020;  
Article No. 21,  
Pages: 125-128  
ISSN 0976 - 044X

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Evaluation of Fast  
Disintegrating  
Tablets of  
Salbutamol  
Sulphate, Cetirizine  
Hydrochloride in  
Combined  
Pharmaceutical  
Dosage Form: A  
New Era in Novel  
Drug Delivery for  
Pediatrics and  
Geriatrics Deepak  
Sharma,<sup>1</sup> Gurmeet

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Singh,<sup>2</sup> Dinesh  
Kumar,<sup>3</sup> and  
Mankaran Singh<sup>4</sup> 1

Development

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Development and -  
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The main criteria of  
the present work is  
formulation

development of  
Clarithromycin

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Research

topical gel by using  
four types of  
gelling agents Na  
CMC, Hydroxy  
propyl cellulose,  
Guar gum,  
Poloxamer 407 and  
study the gelling  
agents affecting on  
the release of  
drug.<sup>3,4</sup>

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Volume 5, Issue 3,

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November -

December 2010;

Article-011 ISSN

0976 - 044X

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and Research Page

65 ...

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FORMULATION

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## DEVELOPMENT AND EVALUATION OF ...

The analytical chemistry or the analytical development lab often executes the majority of the formulation tasks, as they are already the 'experts' in the analytical assays, most often high

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performance liquid chromatography (HPLC) assays, which are an integral part of product development.

Types of Projects

Formulation

development

encompasses a

very wide range of activities.

Traditionally,

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formulation covers  
such functions as  
pre-formulation,  
including analytical  
assay de-  
velopment and ...

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be dexterous to  
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Available online at  
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December -  
January, 2016, Vol.  
5, No.1, pp

1963-1973 ISSN

(P): 2393-932X,

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2278-0238

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FORMULATION

DEVELOPMENT

AND EVALUATION

OF MATRIX TYPE

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TOPICAL PATCH

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116. 14.

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EVALUATION AND  
ANTI...

Our formulation  
development team  
has extensive  
experience in  
different dosage  
form development  
for both New  
Chemical Entity  
(NCE) as well as  
generic product  
development  
applying our

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technical skills,  
efficiency and  
quality  
consciousness to  
develop robust  
products. Our  
expertise lies in  
overcoming the  
challenges faced  
during formulation  
development and  
offer solutions to  
overcome  
flowability,

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solubility,  
wettability,  
dissolution,  
degradation,  
bioavailability  
issues encountered  
during  
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effective parenteral  
formulation  
containing  
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Trometamol in a  
hydro alcoholic  
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of Pharmacy, Al-  
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Zinc sulphate tablets are indicated for the management of diarrhoea in children regardless of the cause. In Tanzania, there is only one pharmaceutical industry manufacturing zinc sulphate tablets

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and only 44% of children in need of zincsulphate tablets get access to them. Fast-disintegrating tablets of zinc sulphate were prepared by direct-compression method after incorporating the ...

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JCPRC5 88

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Formulation

development and  
optimization of

controlled release

microspheres of

Aceclofenac using

response surface

methodology Ketan

J. Patel 1\* and

Abhay Dharamsi 2

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Research Article

ISSN : 0975-7384

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Research

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JCPRC5

The efficacy of formulation was evaluated in patients by subjective assessment, gamma scintigraphic approaches, and confocal microscopy.

Methods:

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Nifedipine-loaded different formulations such as sucrose bead, pellets, and microparticles (slugging method, ionotropic gelation, and chemical denaturation) were designed. The studies were performed on 50 subjects, of which

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30 subjects were treated with optimized nifedipine loaded microcapsules while 20 subjects were given capsule becosule-Z as a ...

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Formulation development and evaluation of nifedipine as ...



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In the development of the co-transfer formulation, the nano-formulation exhibits good bioavailability and compatibility. For example, polymeric micelles (PMs) are self-assemblies of block copolymers providing numerous opportunities for

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drug delivery.

Besides, the nano-encapsulated anticancer agent targeting specific tumor tissues can significantly optimize the therapeutic efficacy of the drug.

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