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Controlled drug delivery concepts and advances. 1st ed. Jan 2002; 218-219; S P Vyas; R K Khar; Vyas SP, Khar RK. Controlled drug delivery concepts and advances. 1st ed. 2002. p. 218-9.

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### **Oral Controlled Release Drug Delivery System A Review**

A desirable characteristic of controlled release delivery system is that the duration of drug action should be dictated by the design property of drug molecules. There are different mechanistic aspects for design of oral controlled release drug delivery systems such as matrix, reservoir, osmotic pressure, ion exchange resins, altered density etc.

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drug delivery system or by modifying the molecular structure and /or physiological parameters [2]. Advantages of Controlled Drug Delivery [3-6] Maintenance of drug levels within a desired range.

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Supramolecular Hydrogel of a d-Amino Acid Dipeptide for Controlled Drug Release in Vivo. Langmuir 2009, 25 (15) , 8419-8422. DOI: 10.1021/la804271d. Emily Gullotti and Yoon Yeo . Extracellularly Activated Nanocarriers: A New Paradigm of Tumor Targeted Drug Delivery.

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### **Preparation and evaluation of magnetic microspheres of**

Conflict of interest statement We declare that we have no conflict of interest. References [1] Alagusundaram M, Madhusudana CC, Umasharkari K, Badrinath AV, Lavanya C, Ramkanth S. Microspheres as a novel drug delivery system. Int J Chem Tech Res 2009; 1(3): 526-534. [2] Vyas SP, Khar RK. Controlled drug delivery concepts and advances.

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Drugs that have narrow absorption window in the gastrointestinal tract (GIT) will have poor absorption. For these drugs, gastro retentive drug delivery systems offer the advantage in prolonging the gastric emptying time. Atenolol is an antihypertensive drug, which has low elimination half-life: 3 4 hrs. The floating tablets of Atenolol were prepared to increase the gastric retention and to

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### **9 controlled drug delivery systems issuu com**

29. Vyas SP, Khar RK. Controlled Drug Delivery: Concepts and Advances. 1st ed. vallabh prakashan, 2002:156-189. A model for the drug release from a polymeric matrix tabletseffect of swelling

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### **Formulation and Evaluation of Captopril Transdermal**

The absorption of drugs via transdermal route improves the bioavailability of drugs that could have might otherwise been metabolized by first-pass way (pre-systemic drug elimination) by gastrointestinal tract. Drug absorption is mainly through passive diffusion through the lipoidal membrane. Thus TDDS have attracted the attention worldwide for optimizing the drug delivery.

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INTRODUCTION IDEAL SYSTEM Agent independent Flexibility Monitoring & decision- making Targeting Vyas SP, Khar RK. Controlled drug delivery: Concepts and advances. 2nd edition. Vallabh Prakashan. 2012, 397-398. 4.

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#### **Oral Sustained Release Tablets An Overview with a special**

Vyas SP, Khar RK. Controlled drug delivery concepts and advances. 1st Ed.VallabhPrakashan, New Delhi. 2010;1-12. Kamboj S, Gupta GD. Matrix tablets: An important tool for oral controlled-release dosage forms. Pharmaceutical Rev. 2009;7(6):1-9. Shargel L, Yu ABC. Modified release drug products.

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#### **PROTEINS AND PEPTIDES authorSTREAM**

structure of proteins and barriers affecting protein and peptide drug delivery system- authorSTREAM Presentation. 36 REFERENCES Vyas S.P. And Khar Roop K., Controlled Drug Delivery Concepts and Advances ; Vallabh Prakashan , Delhi. Page no. 503-570. Satyanarayana U., Chakrapani U., Biochemistry , 3 rd edition, 2011, Books and

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#### **PAPER IV ADVANCES IN DRUG DELIVERY SYSTEMS GOAL OBJECTIVE**

ADVANCES IN DRUG DELIVERY SYSTEMS. GOAL . To train the students in the area of novel drug delivery systems. Concepts, advantages and disadvantages, structure of oral mucosa, transmucosal Vyas SP., Khar RK., Controlled drug delivery-concepts and advances, Vallabh Prakashan, New Delhi, first edition 2002.

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The objective of this study was to design oral controlled release matrix tablets of lamivudine using hydroxypropyl methylcellulose (HPMC) as the retardant polymer and to study the effect of various

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Controlled release (CR) tablet formulations are much desirable and preferred for such therapy because they offer better patient compliance, maintain uniform drug levels, reduce dose and side effects and increase safety margin of high-potency drugs . Several polymers have been used in the formulation of matrix based CR drug delivery systems.

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Vyas SP, Roop K, Khar: Controlled Drug Delivery Concepts & Advances. 383-409. Rathore KS, Nema RK: Formulation and Evaluation of Ophthalmic Films for Timolol Maleate. *Planta Indica*, 2008; 4:49-50.

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The objective of this study was to design oral controlled release matrix tablets of lamivudine using hydroxypropyl methylcellulose (HPMC) as the retardant polymer and to study the effect of various formulation factors such as polymer proportion, polymer viscosity, and compression force on the in vitro release of drug.

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REFERENCE Controlled drug delivery, concepts & advances; Suresh.P.Vyas ,Roop.K.Khar, Pg no: 257-300 Controlled & novel drug delivery, by N.K Jain , Pg no: 353-380 Singh.J, Deep.P; A review article on mucoadhesive buccal drug delivery system, International journal of pharmaceutical sciences & research, 2013, vol 4(3), 916-927 28 29.

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### **Controlled Release Drug Delivery Systems**

Controlled Drug Delivery (CDD) occurs when a polymer, whether natural or synthetic, is judiciously combined with a drug or other active agent in such a way that the active agent is released from the material in a pre-designed manner. The release of the active agent may be constant over a long period, it may be cyclic over

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The drug release performance was greatly affected by the materials used in microparticle preparations, which allow absorption in the intestinal tract. Key words Ethyl cellulose, eudragit, kinetic drug release, microparticles, ranolazine. REFERENCES 1. S.P. Vyas, K.R. Khar, Controlled drug delivery concepts advances. 1st ed 2002 1 51. 2.

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### **ORAL CONTROLLED RELEASE DRUG DELIVERY SYSTEM A REVIEW**

Drugs existing largely in ionized form are poor candidate for oral controlled release drug delivery

system because absorption rate of ionized drug is 3-4 times less than that of unionized form. The pKa range for acidic drug whose ionization is pH sensitive is around 3.0-7.5 and for basic drug whose ionization is pH sensitive is around 7.0-11.0

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