

AN RECENT ADVANCEMENT IN TOPICAL DOSAGE FORMS: A REVIEW

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Abstract

There are many semisolid dosage forms but creams, pastes, emulsions, ointments, gels, and rigid foams etc are main examples of this category of dosage form. They serve as carriers for drugs that are topically, administered by route of the skin (derma), retina, rectal tissue, nasal mucosa, vagina, buccal tissue, urethral membrane, and external ear lining. Extant definitions of lotions, gels, creams and ointments vary depending on literature source, market history or traditional use. This often leads to confusion when deciding which topical Dosages forms to prescribe and purchase. The purpose of this study is to obtain a scientifically based, regular classification of Dosages forms for topical drugs. A variety of prescriptions and over the counter topical products currently marketed as lotions, gels, creams, and ointments are evaluations using variant techniques including rheology, viscosity and shear rate versus shear stress, loss on drying, specific gravity, surface tension, thermogravimetric analysis (TGA), water absorption, dilution properties, microscopic evaluation, transmittance of visible light, features (appearance) or conformation (composition). Skin is the largest organ of the humanitarian corpus and plays the most important role in protecting against microorganisms and foreign body. The important instruments such as topical, regional and Transdermal are widely used for delivery of variant dosages forms. The main use of semi-solid dosage form additives are controlling the development of absorption, maintaining the viscosity, improving the stability as well as organoleptic (sense organs) Property and increasing the bulk of the (synthesis) formulation.

Keywords: rigid foams, ointments, absorption, viscosity, thermogravimetric

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INTRODUCTION

Drug semisolid arrangements might be characterized as skin items planned for application on the skin or available mucous films to give confined and some of the time foundational impacts at the site of utilization. As a rule, semisolid measurements structures are complicated definitions having complex underlying elements. The utilization of a medication

containing detailing to the skin or mucous film, to treat explicit cutaneous problems (for example skin break out) or cutaneous signs of a summed-up infection (for example psoriasis), with the aim of containing the pharmacological impact of the medication just to the surface or inside the layers of skin or mucous (derma)membrane. Semi solids are the

skin dose structure utilized for the restorative, defensive or corrective capacity. They might be applied to the skin, or utilized nasally, vaginally, or rectally...Semisolid dose structures: are results of semi-strong consistency and applied to skin or mucous films for a remedial or defensive activity or corrective capacity.[1-4]

ADVANTAGES OF SEMI-SOLID DOSAGES FORMS:

- It is utilized remotely.
- Probability of incidental effect can be diminishing.
- First pass gut and hepatic digestion is kept away from.

- Local activity and Site explicit activity of medication on influenced region.
- Convenient for oblivious patient or patient experiencing issues on oral organization.
- Suitable dose structure for harsh medications.
- More stable than fluid dose structure.
- Site explicit application

DIFFERENT TYPES OF SEMI SOLID:

It includes ointments, creams, pastes, gels and many more. {5-6}



OINTMENT:

Ointments (Balms) are homogenous, clear, goeey semi-strong arrangements, most ordinarily an oily, thick (oil 80% - water 20%) planned for outside application to the skin or mucous layer. Drug fixings can be broken down, emulsified or suspended in

the salve base. Balm primarily use as an emollient or defensive for skin. They are utilized as Emollients or for the utilization of dynamic Ingredients to the skin for defensive, restorative, or prophylactic reason and where a level of Occlusion is wanted.[7-9]



CREAM:

These are semi-strong emulsions for counter use. There are lipophilic (W/O) and hydrophilic (O/W) creams, contingent upon the constant stage...There are two kinds of aqueous and oily creams in which the emulsions are (oil in water and water in

oil) respectively. The oil in the water type is somewhat non-greasy. The creams are exceptionally mainstream type of outside medication. Consist of a lipophilic stage and a fluid phase. There are lipophilic (W/O) and hydrophilic (O/W) creams, contingent upon the constant stage.[10-12]

**PASTES:**

Pastes (Glue) are essentially salves into which a high level of insoluble strong has been added - The unprecedented measure of particulate matter hardens the framework. - Pastes are less entering and less macerating and less warming than salve. Pastes are essentially salves into which a high level of insoluble strong has been added. The phenomenal measure of particulate matter solidifies the framework

through direct Interactions of the scattered particulates and by adsorbing the fluid hydrocarbon division (part). The vehicle on the molecule surface pastes makes especially great defensive hindrance when put on the skin for, as well as Forming a whole film, the strong they contain can retain and subsequently kill certain poisonous synthetic compounds before they at any point arrive at the skin.[13]

**GELS (JELLIES):**

Gels are semisolid framework in which a fluid stage is compelled inside a 3-D polymeric lattice (comprising of regular or

gum) having a serious level of physical or compound cross-connecting. Gels are framed by utilizing manufactured

polymers, for example, carbomer 934 and cellulose like Hydroxypropyl cellulose and hydroxy propylmethyl cellulose. Some are just about as straightforward as water itself, a stylishly satisfying state, other are turbid, as the polymer is available in colloidal totals that scatter light. They are

utilized for drug, oil and some various applications like transporter for spermicidal specialists to be utilized intra vaginally with stomachs as an adjunctive method for contraception(contraceptive). [14]



POULTICES:

It is delicate, thick, pale groundwork for outer use. They are applied to skin while they are hot. Poultice should hold heat for a significant time frame since they are expected to supply warmth to aroused pieces of body.[15]

RIGID FOAMS:

Froths are framework in which air, or some different gas is emulsified in fluid stage to the reason behind hardening. For example, shaving creams, whipped creams and sprayed shaving creams.

RECENT ADVANCEMENT IN SEMI-SOLID DOSAGE FORMS:

The plan of a reasonable semisolid measurements structure includes the determination of a fitting medication transporter framework, with an exceptional accentuation on the medication's physicochemical properties and required Therapeutic application. Medication conveyance through semisolid measurement structures has seen new difficulties in the beyond couple of years as far as adjusted medication discharge

profiles just as the upgraded steadiness of dynamic drug fixings (APIs).[16]

A Cream Containing Liquid Nanoparticles:

For upgraded infiltration of skin drugs, impediment of skin is the superb model. This Requirement can be achieved easily by the incorporation of large quantities of fats and oils, especially liquid and such as Carbomers which confer a stylishly satisfying, clear shining appearance to the item and are handily washed off the skin with water. Gels are two-part Semisolid's frameworks wealthy in fluids. In an ordinary polar gel, a characteristic or engineered polymer constructs a Three-dimensional framework all through a hydrophilic fluid. Typical polymers used include the Natural gums tragacanth, carrageenan, pectin, agar, and alginic acid; semisynthetic materials such as methylcellulose, hydroxyethyl cellulose, hydroxyl propyl methylcellulose, and carboxymethyl Cellulose; and the synthetic polymer, carbopol may be used. Certain muds like bentonite, Veegum, and laponite given that the medication doesn't tie to the

polymer or mud. Such gels discharge Medicaments well; the pores permit generally free dissemination of particles, which are not very huge.[17]

Gels with permeation enhancers

Skin can go about as a boundary to the more profound entrance of medication atoms. With the presentation of different infiltration enhancers, nonetheless, fundamental medication conveyance through the transdermal course has acquired significant equilibrium (balance). These synthetics, joined in an appropriate medication conveying semisolid vehicle, enhance the measure of medication saturation through skin either by reversibly cluttering the lamellar pressing of layer corneum or by expanding the thermodynamic action of the medication. One more class of entrance enhancers acts by expanding the measure of medication in solubilized structure at the skin surface, bringing about the upgraded penetrability of lipophilic medication particles. Countless synthetics have been read for infiltration improvement movement. The quest proceeds for new synthetic compounds with helpful action at low focuses and with negligible cutaneous disturbance potential. Notwithstanding the utilization of entrance enhancers alone, their mix with cosolvents that convey a medication in solubilized structure has prompted the accomplishment of higher medication porousness (porosity).[12-17]

New Treatment for Faecal Incontinence using the Zinc-Aluminium Ointment:

A randomized, twofold visually impaired preliminary was performed. Patients who met the consideration rules were Randomized to get the treatment or a fake treatment. All were assessed previously and 3 weeks after Ointment application, utilizing the Wexner incontinence score and the Fecal Incontinence Quality of Life (FIQL) score. The investigation shows that the zinc-aluminum based balm diminishes waste Incontinence altogether contrasted and fake treatment.[18]

Controlled Release Gel:

Conveyance to nasal or Ocular mucosa for one or the other neighborhood or foundational activity faces numerous impediments. These courses are ensured by successful instruments. Gel details with appropriate rheological and mucoadhesive properties increment the contact time at the site of assimilation. Notwithstanding, drug Release from the gel should be maintained in case benefits are to be acquired from the drawn out contact time. Gelrite gels were framed in reenacted tear liquid at groupings of the polymer as low as 0.1%. And it was shown that sodium was the main gel advancing particle in-vivo. Rheology, Although it very well might be a sketchy method for assessing mucoadhesive properties of polymers, Showed that communications among mucin and polymers were probably going to be seen with powerless gels (sticky substance). It was feasible to control the arrival of uncharged medication substances by including surfactants that Form micelles in the gel. This delivery relied upon lipophilic communications between the medication and the Polymer and the micelles. Controlled-discharge plans of charged medications could be planned by Mixing the medications with oppositely charged surfactants in specific proportions. Thusly, vesicles in Which the medication and surfactant comprised the bilayer framed unexpectedly. The vesicle development Was influenced by the presence of polymer, and tiny vesicles that gave a sluggish delivery rate were Formed when a lipophilically altered polymer was (spend)utilized. The gels were additionally assessed in the Using chamber utilizing porcine nasal mucosa. The pace of transport of medications through the mucosa Could be constrained by the pace of delivery from the detailing. Besides, the Using chamber Could be utilized to assess the likely poisonousness of definitions.[12-14]

Oleo-hydrogel systems:-

Oleo-hydrogel frameworks for limited skin activity have been investigated successfully. Transdermal Permeation utilizing different vehicle frameworks to keep away from fundamental incidental effects and gastrointestinal Irritation from ketoprofen upon oral organization. When contrasted and traditional gel or Plaster plans, the oleo-hydrogel frameworks were observed to be the ideal definition since It diminished foundational dissemination of the drug and extended limited exercises(activity). The specialists Examined an oleo-hydrogel framework that comprised of ketoprofen consolidated into an emulsion of Oil and carbomers hydrogel combination, with N-methylpyrrolidone as a pervasion enhancer(improve). The Greater bioavailability of ketoprofen in the oleo-hydrogel framework was credited to acceptable medication discharge Properties, higher emulsion drops steadiness of the Carbomers gel, and the entrance upgrading Effect of Methylpyrrolidone. A serious level of relationship was seen between in vitro Permeation and in vivo percutaneous assimilation boundaries. The plan of ketoprofen oleo Hydrogel that showed greatest percutaneous assimilation was one that contained 3% ketoprofen, 1% carbomers, 10% N-Methylpyrrolidone, 10% oils, 8% surfactant, and water changed in accordance with pH 4. 6 Utilizing triethanolamine.[12-14]

Deoxycholate hydrogels:

Sodium deoxycholate (a low sub-atomic weight drug transporter) was observed to be a superior option in contrast to High sub-atomic weight polymers as a gelling specialist. It additionally goes about as an entrance enhancer for Topically controlled medication particles. Sodium deoxycholate offers benefits, for example, low dissolve Viscosity, likely biocompatibility and biodegradability, and nonappearance of harmful pollutions from Synthesis deposits

like natural solvents, impetuses, and initiators. At the point when it interacts with Excess cradle frameworks, it shapes a gooey thixotropic gel with improved film porousness. Sodium deoxycholate gels leave no buildup after application, and in view of their thixotropic Behavior, they are not difficult to apply on huge skin regions. The surfactant activity of sodium deoxycholate Facilitates the solubilization of a few medications by framing blended micelles. This framework has been Studied for its improved ingestion of progesterone and prednisolone through smooth mouse skin by delivering underlying changes in the layer (stratum)corneum.[12]

Lamellar Faced Creams:

They are fluid paraffin in water emulsion arranged from cetrimide/greasy liquor like blended Emulsifiers and ternary framework shaped by scattering the blended emulsifier in required an amount of water. The cationic emulsifying wax showed sensational enlarging in water, and this expanding Was because of electrostatic shock, which can be stifled by the expansion of salt and can be Reduced by changing surfactant counter particle.[19-20]

Nanospheres Gel:

Tyrosine-inferred nanospheres have shown potential as viable transporters for the effective Delivery of lipophilic atoms. Gel definition containing nanospheres was created for Effective skin application and upgraded pervasion. Carbopol and HPMC hydrophilic gels were Evaluated for scattering of these nanospheres. Sparingly water-solvent diclofenac sodium (DS) And lipophilic Nile Red were utilized as model mixtures. DS was utilized to decide the ideal Polymer type, consistency, and delivery properties of the gel while fluorescent Nile Red was utilized in-vitro and in-vivo skin dissemination contemplate (consider). Likewise, the impact of an entrance enhancer, azone, on the Skin conveyance was examinations.[21]

Hydrophilic Gels:

Hydrophilic gels are bicoherent frameworks made out of the inward stage made of a polymer delivering a sound three-dimensional net-like design, which fixes the fluid vehicle with respect to the outer stage. Intermolecular powers tie the atoms of the dissolvable to a polymeric net, in this manner diminishing the portability of these particles and delivering an organized framework with expanded thickness. The physical and compound bonds restricting the particles of the inner stage give a moderately steady construction, which can begin by enlarging of strong polymers, or by diminishing the dissolvability of the polymer in an answer(response). A significant gathering of gels utilized in drug store is hydrophilic gels, or hydrogels, normally made of hydrophilic polymers, which under specific conditions and polymer focus, Jellify.

Consideration of drug research currently focuses principally on hydrophilic gels, as this measurement's structure is by all accounts forthcoming for the advancement of present-day drugs dependent on frameworks with drawn out and controlled arrival of dynamic Ingredients(agent).[22]

Microsphere-Based Improved Sunscreen Formulation of Ethylhexyl Methoxycinnamate:**REFERENCE:**

1. Phaechamud T, Mahadlek J, Charoenteeraboon J and Choopun S: Characterization and antimicrobial activity of N-Methyl-2-pyrrolidone-loaded ethylene oxide-propylene Oxide block copolymer thermosensitive gel. Indian Journal of Pharmaceutical Sciences 2012; 74(6): 498-04.
2. Allen L, Popovich N and Ansel H: Pharmaceutical DosageForms and Drug Delivery Systems. Lippincott Williams &Wilkins, 8th edition, 2005: 276-97.
3. Shinde U, Pokharkar S and Modani S: Design andEvaluation of

Polymethylmethacrylate (PMMA) microspheres of Ethylhexyl methoxycinnamate (EHM) were Prepared by emulsion dissolvable dissipation technique to work on its photostability and adequacy as a sun screening specialist. The PMMA microspheres of EHM were fused in a water removable cream base. The fuse of EHM stacked PMMA microsphere into cream base had significantly expanded the viability of sunscreen definition roughly multiple times.[23]

CONCLUSIONS:

The primary benefits of novel semisolid measurement structures are non-oily since they are comprised of water launder able bases, simple application, fast definition, and capacity to topically convey a wide assortment of medication atoms. Presently a day, more prominent endeavors has been made for accomplishing controlled delivery definitions by utilizing distinctive transporter frameworks, dispensing with the cosmetically horrible characteristics of the traditional semisolid measurement's structures. Fitting excipient choice and security assessment are significant in the advancement of semisolid measurement structure.

- microemulsion gel system of Nadifloxacin. Indian Journal of Pharmaceutical Sciences 2012; 74(3):237-47.
4. Gupta P and Garg S: Recent advances in semisolid dosageForm for dermatological application. PharmaceuticalTechnology 2002; 144-62.
5. Jenning V, Schafer-korting M and Gohla S: Vitamin A loaded solid lipid nanoparticles for topical use: drugRelease properties. Journal of Controlled Release 2000;66(2-3): 115-26.

6. Bhowmik D, Gopinath H, Kumar P, Duraivel S and Kumar S: Recent advances in novel topical drug delivery system. *The Pharma Innovation* 2012; 1(9): 22-27.
7. Segers J, Zatz J, Shah V: In-vitro release of phenol from Ointment formulations. *Pharmaceutical Technology* 1997; 21(1): 70-81.
8. Halligan B, Ruotti V, Jin W, Laffoon S, Twigger S and Dratz E: ProMoST (Protein Modification Screening Tool): A web-based tool for mapping protein modifications on Two-dimensional gels. *Oxford journals, Nucleic Acids Research* 2004; 32(2): W638-W644.
9. Rabilloud T, Chevallet M, Luche S and Lelong C: Two-dimensional gel electrophoresis in proteomics: Past, Present and future. *Journal of Proteomics* 2010; 73(11): 2064-77.
10. Gohel M and Parikh R: Novel semisolid dosage form. *Int. J. Pharm.* Vol. 203. No. 1-2. August 10. 2000. 127-39.
11. Basu S, Chakraborty S, Bandyopadhyay AK, Development and Evaluation of a Mucoadhesive Nasal Gel of Midazolam Prepared with *Linum usitatissimum* L. Seed Mucilage, *Scientia Pharmaceuticals*, 2009; 77: 899-910.
12. Sharma AK, Naruka P S, Soni SL, Sarangdewot YS, Khandelwal M, Aman S, Formulation, Development and Evaluation of Luliconazole Organogel; *International Journal of Current Pharmaceutical Review and Research*. 12(1).
13. Alsarra A et al., Mucoadhesive Polymeric Hydrogels for Nasal Delivery of Acyclovir, *Drug Development and Industrial Pharmacy*, 2009; 35: 352-62.
14. Agrawal D, Goyal R, Bansal M, Sharma AK, Khandelwal M, Aman S, Development and Evaluation of Econazole Organogel; *International Journal of Current Pharmaceutical Review and Research*, 13(2), Pages: 15-23.
15. Varshosaz J, Sadrai H, Heidari A, Nasal Delivery of Insulin Using Bioadhesive Chitosan Gels, *Drug Delivery*, 2006; 13: 31-6.
16. Pisal SS, Reddy P, Paradkar AR, Mahadik KR, Kadam SS, Nasal Melatonin gels using Pluronic PF-127 for chronobiological treatment of sleep disorder, *Indian Journal of Biotechnology*, July 2004; 3: 369-77.
17. Sharma AK et al. pharmaceutical gel: A review, *International Journal of Pharmacy & Technology*, Dec. 2020. 12(4), 7223-7233.
18. Majithiya RJ, Ghosh PK, Umrethia ML, Murthy RSR., Thermoreversible-mucoadhesive Gel for Nasal Delivery of Sumatriptan. *AAPS PharmSciTech*. 2006; 7(3): Article 67.
19. Nandgude T, Thube R, Jaiswal N, Deshmukh P, Chatap V, Hire N, Formulation and Evaluation of pH Induced In-situ Nasal Gel of Salbutamol Sulphate, *International Journal of Pharmaceutical Sciences and Nanotechnology*, July –September 2008; 1(2): 177-83.
20. Mahajan HS, Gattani S, In situ gels of Metoclopramide Hydrochloride for intranasal delivery: In vitro evaluation and in vivo pharmacokinetic study in rabbits, *Drug Delivery*, 2010; 17(1): 19-27.
21. Rathnam G, Narayanan N, Ilavarasan R, Preparation and Evaluation of Carbopol Based Nasal Gels For Systemic Delivery Of Progesterone, *International Journal Of Periodontics and Restorative Dentistry*, March 2005; 2(1).
22. Tang Y, Zhou C, Luan J, Hu J. (2012). Improvement of length of survival of expanded flap by application of topical papaverine cream. *J Plast Surg Hand Surg* 46: 389-92.
23. Tanriverdi ST, Ozer O. (2013). Novel topical formulations of Terbinafine-HCl for treatment of onychomycosis. *Eur J Pharm Sci* 48: 628-36.