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"NATURAL POLYMERS AND THEIR APPLICATIONS IN FAST DISSOLVING TABLETS: A COMPREHENSIVE REVIEW"

Present By: Meet Naliyadhara^{*a}, Riya chovatiya^b,

Shyam Vekariya^c, Deep Undhad^d, Dr.Sheetal Buddhadev^e

• Faculty of Pharmacy, Noble University

Introduction of FDT

Fast dissolving tablets (FDTs) offer rapid drug disintegration and absorption without water, addressing challenges for patients with swallowing difficulties, like pediatric and geriatric populations. FDTs improve patient compliance and provide faster therapeutic action. Traditional tablets often fail to meet these needs, making natural polymers a promising alternative. Natural polymers enhance safety, ensure faster disintegration, and improve drug release, offering significant advantages in pharmaceutical formulations.

Introduction Of Natural Polymer

Natural polymers are increasingly utilized in Fast Dissolving Tablets (FDTs) for their biocompatibility, safety, and ability to enhance disintegration. Polymers like Xanthan gum, Guar gum, and Plantago ovata promote rapid tablet breakdown in saliva, leading to faster drug release and absorption. Their incorporation not only boosts FDT performance but also meets the demand for safer, sustainable excipients, aiding the development of innovative drug delivery systems that improve patient compliance and therapeutic outcomes.

Mechanism Of Action

Disintegration: FDTs rapidly break apart in saliva without



Why Natural Polymers in FDTs

1.Biocompatibility: Safe, non-toxic for drug delivery.

- 2. Biodegradability: Eco-friendly, easily eliminated from the body.
- 3. Low Cost: Affordable, derived from abundant resources. 4. Non-allergenic: Lower risk of allergic reactions. 5. Patient Compliance: Suitable for all age groups due to safety. 6. Sustained Release: Allows for controlled and extended drug release. 7. Gel-forming: Quick disintegration in saliva.

- Dissolution: The drug is released and dissolves in saliva.
- Absorption: The drug is absorbed through the oral mucosa or GI tract.
- Therapeutic Effect: The drug enters the bloodstream quickly, by passing first-pass metabolism for faster action.

Introduction to Superdisintegrants

Superdisintegrants are key excipients in the formulation of Fast Dissolving Tablets (FDTs). They facilitate the rapid breakdown of tablets upon contact with saliva by promoting water uptake and swelling, which accelerates disintegration. Unlike conventional disintegrants, superdisintegrants work more efficiently at lower concentrations, ensuring quick disintegration without compromising tablet integrity. Their use is essential in improving drug release and absorption, making them vital for enhancing the performance and patient acceptability of FDTs.

Types of Superdisintegrants



Natural Superdisintegrants

• Example: Plantago ovata, Guar gum, Xanthan gum • Derived from natural sources, these are biodegradable, non-toxic,

Methods For The Preparation Of Fast Dissolving **Tablets**



Natural Polymer Utilized In Commercial Pharmaceuticals









Chitin and chitosan:

• Marketed drug: Cinnarizine

Agar and treated **Gum Karaya:**

- Marketed drug: Amlodipine
- **Fenugreek seed** mucilage: • Marketed
- drug: Metformin HCl

• Disintegration



Evaluation Parameters Used For Fast Dissolving Tablets

• Hardness Wetting Time • Taste Evaluation Water Absorption Ratio Mechanical Strength • Friability In-Vitro Dissolution Study • Weight Variation Porosity Stability Testing (under **Content Uniformity** • Thickness different temperature and Disintegration Time Moisture Uptake humidity conditions)

• Disintegration time: 60 sec

- Concentration used: 3% w/w
- Theophylline

• Marketed

drug:

agar:

- Disintegration time: 20 sec
- Concentration **used:** 1-2% W/W
- Disintegration time: 17.10 sec Concentration

used: 4% w/w

time: 15.6 sec • Concentration **used:** 4% w/w

Conclusion

Natural polymers in fast dissolving tablets (FDTs) offer a promising advancement in drug delivery, enhancing disintegration, drug release, and patient compliance. Their renewable nature provides a sustainable alternative to synthetic excipients while maintaining drug efficacy and bioavailability, making them a valuable option for future pharmaceutical formulations.

References

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