ISSN: 2320-2882

IJCRT.ORG



INTERNATIONAL JOURNAL OF CREATIVE RESEARCH THOUGHTS (IJCRT)

An International Open Access, Peer-reviewed, Refereed Journal

A REVIEW ON SUPERDISINTEGRATING AGENT USED IN PHARMACEUTICAL FORMULATION

Disha P.Wankhede¹, Damini A. Mundhare², Laxmi S. Deshmukh³, Shravan S. Sawarkar⁴, Sayyed Tohid Sayyed Hakeem⁵, Mohammad Tausif Shaikh Qayyum⁶, Nandakishor Deshmukh⁷, and Dr. Swati P. Deshmukh⁸

> 1,2,3,4,5,6 Student Of Bachelor Of Pharmacy, Shraddha Institute of Pharmacy, Kondala Zambre, Washim 444-505. 7 Assistant Professor Department of Pharmaceutics, Shraddha Institute of Pharmacy, Washim 444-505. 8 Professor Department Of Pharmacology, Shraddha Institute of Pharmacy, Kondala Zambre, Washim 444505.

Abstract

Disintegrating agents are super helpful in tablets. They help break up the compacted mass so the active ingredients can dissolve or release when in contact with fluids. The most important function of disintegrants is resist the ability of the tablet binder and bodily forces that act below compression to structure the tablet. Superdisintegrants provide rapid dissolution disintegration because of blended impact of swelling and water absorption. Superdisintegrants promotes the wettability and dispersibility of the machine, as a consequence increasing the disintegration and dissolution. One gram of superdisintegrants absorb 10-40 gm of water or liquid solution. After the absorption it produce pressure and leads to the entire tablet shape to break a side. Their particles are typically tiny and porous, which permit for fast tablet disintegration inside the mouth without an objectionable mouth-sense from either large particles or gelling. The debris are also compressible which enhance the tablet hardness and its friability. Powerful superdisintegrants offer advanced compressibility, compatibility and don't have any negative impact on the mechanical strength of formulations containing excessive-dose drug. Great disintegrating tablets are novel drug delivery system that dissolves, break into small particle's or dispersed the active pharmaceutical elements in saliva inside few seconds with or with out consumption of water. The quicker the dissolution of drug into the answer, quicker is the absorption and onset of medical effect.

Keyword -disintegranting agents, superdisintegrants, tablets, disintegration, dissolution.

Introduction

Superdisintegrants are the amazing absorbing substance with the swelling property. Those substances aren't planned to absorb quantities of water or aqueous solutions, however deliberate to swell very rapid. Superdisintegrants are used as a structural weakener for the disintegrable stable dosage forms. They're bodily dispersed inside the matrix of the dosage form and could become large when the dosage shape's uncovered to the moist environment¹. These newer substances are extra effective at decrease concentrations with greater disintegrating performance and mechanical power². Superdisintegrants are usually used at a low level inside the solid dosage usually 1 - 10 % with the weight relative to the entire weight of the dosage unit³.

Superdisintegrants, are those substances, which facilitate the faster disintegration with smaller amount in comparison to disintegrants. The disintegration of dosage form are depend upon the physical factors of disintegranting agents /superdisintegrating agents which might be as observe:

1.Percentage of disintegrants present in the method.

- 2.Percentage of disintegrants used.
- **3.**Compatibility with different excipients.
- **4.**Presence of many surfactants.
- **5.**Tablet hardness.
- **6.**Behavior of drug materials^{4,5}.

Researchers nowadays are searching out for a brand new, safe and effective disintegrating agents that could fall apart. Tablets unexpectedly even at a tablet crushing energy of greater than 3.5 kg. On studying the behavior of disintegration time within the oral cavity in addition to wetting time by means of surface free energy. We came to recognise, that for a faster wetting a molecule must have high polar components of surface free energy and the retailers which meet those special requirements are called as superdisintegrants.⁶ the convenience of availability of those agents and the simplicity in the direct compression procedure advise that their use would be a more profitable opportunity in the training of odt than the state-of-the-art and patented strategies⁷.

Selection of superdisintegrants

- 1. Low solubility.
- 2. Low gel formation.
- 3. Good moisturization capacity.
- 4. Good moulding and flow properties.
- 5. No tendency to form complexes with the drugs.
- 6. Good mouth feel.
- 7. It should also be suitable with the other excipients and have desirable tableting properties^{8,9}.

Method of addition of superdisintegrant

Internal addition

In the moist granulation technique, the disintegrant is introduced to different excipients before wetting the powder with the granulating fluid. Thereby, the disintegrant is included within the granules. In a dry granulation approach, the disintegrant is introduced to other excipients before compressing the powder between the rollers. In a computer-optimized experiment, the examine suggests the impact of incorporating a disintegrant, croscarmellose sodium, intragranular, greater granularly or allotted equally between the two stages of a tablets in which a low soluble drug constituted as a minimum 92.5% of the method.

External addition

In each moist and dry granulation method, the super disintegrants are introduced to the granules throughout dry blending before compression. The impact of mode of incorporation of excellent disintegrants (croscarmellose sodium, sodium starch glycolate, and crospovidone) at the dissolution of three version tablets with various aqueous solubility (carbamazepine, acetaminophen, and cetirizine hcl) from the irrespective pill formulations by way of moist granulation turned into studied. It is proved that crospovidone is effective in enhancing the dissolution of the drugs in an extra granular mode of addition appears to be the excellent mode of incorporation, no matter the solubility of the principle tablets factor.

Internal and external addition

On this method, the disintegrant is divided into portions. One portion is brought before granule formation (intra) and the final portion is delivered to granules (greater) with mixing earlier than compression¹⁰.

Advantages

- High-quality tendency on wetting causing fast disintegration.
- No lump formation on disintegration.
- Well suited with normally used healing retailers and excipients.
- Does not persist with the punches and dyes.
- Effective in decrease concentrations.
- Much less impact on compressibility and flow ability.
- Extra powerful intra granularly.
- Some are anionic and may purpose a few slight in vitro binding with cationic capsules.
- Biodegradable^{11,12,13}.

Disadvantages

- Costly.
- Time consuming and easily break.
- Extra sensitive and water absorbing in nature¹⁴.

Ideal properties

- 1. They not require water or any other fluid at the time of administration.
- 2. They should easily break into small particles and easily mix.
- 3. It should masks or get better the unacceptable taste of medicine or drug.
- 4. They should have excessive drug loading.
- 5. They must have a pleasant experience in their mouths.
- 6. They should have minor or no residue oral cavity after administration.
- **7.** They should have minimum sensitivity in opposition to environmental situations like moisture temperature and so on.
- 8. They should easy of administration for sufferers who're mentally unwell damage and uncooperative.
- 9. They should be portable without fragility worries.
- **10.** They must be manufactured the use of conventional tablets processing and packing instrumental minimum cost¹⁵.
- **11.** It should to produce speedy disintegration.
- **12.** It must produce suitable molding and flowing properties.
- 13. It should have to a very good particle size excellent hydration ability and a compressibility index.
- 14. It should have to low water solubility.
- **15.** It should have to produce compactable less friable tablets.
- **16.** It should be effective at very low concentrations and should have more disintegrating performance.
- 17. It must be nontoxic and must have a good mouth experience.
- **18.** It should have no longer tendency to form the complexes with the drugs.
- **19.** It should be well matched with the opposite excipients and have to applicable tableting properties^{16,17,18,19}.

Types of superdisintegrants

- **1.** Natural super disintegrants
- 2. Synthetic super disintegrants
- **3.** Co-processed super disintegrants

1.Natural superdisintegrants

Ispaghula husk mucilage (plantago ovata)

Ispaghula husk includes dried seeds of the plant that's referred to as plantago ovata and it includes mucilage that is presentinside theepidermis of the seeds.the mucilage of plantagoovata has distinct functions like binding, disintegrating and maintaining properties. Mucilage is a super disintegrating agent that is used to formulate fast dissolving tablets because the percentage of swelling index is very excessive(around $89\pm2.2\%$ v/v) as compared to the other superdisintegrants^{20,21,22}.

Xanthan gum

Xanthan gum that's derived from xanthomonas campestris is official in usp with excessive hydrophilicity and poor gelling tendency. It has poor water solubility and large swelling properties for faster disintegration²³.

Gellan gum

The disintegration of tablet may be due to the on the spot swelling characteristics of gellan gum when it comes in contact with water and due to its high hydrophilic nature. The complete disintegration of tablet is located inside 4 minutes with gellan gum at a concentration of 4 % $w/w^{24,25}$.

Chitin/chitosan-silicon di oxide

Naturally chitin is extracted from the shell wastes of shrimp, crab, lobster, krill and squid used for the manufacturing of chitosan through a deacetylation response in alkaline medium.chitosan is the excellent recognized herbal polysaccharide used for its versatile applications in pharmaceutical industry^{26,27}.

2. Synthetic super disintegrants

Modified starch (sodium starchglycolate, primojel)

Sodium starch glycolate is the sodium salt of carboxymethyl ether of starch. These are changed starches made by way of cross linking of potato starch as itoffers the product with the better disintegrating properties^{28,29}

Cross-linked polyvinyl pyrrolidone (crospovidone)

Crospovidone quick wicks saliva into the tablet to generate the volume expansion and hydrostatic pressure important to offerfast disintegration within the mouth³⁰.

Modified celluloses (croscarmellose sodium)

It's far insoluble in water, even though it fastly swells to 4-8times its unique volume on contact with water. It specific surface area is zero.81-zero.83 m2/g and swelling index is $65\pm1.7\%$ v/v.in tablets formulations, croscarmellose sodium can be utilized in both direct compression and wet-granulation processes³¹.

3. Co-processed super disintegrants

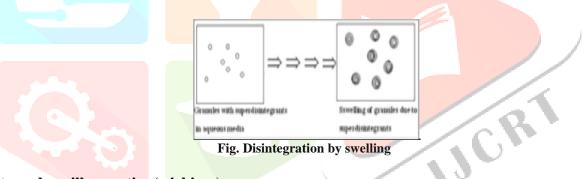
That is primarily based on the radical idea that 2-3 excipients engage at particle level, the objective is used to offer a synergy of functionality development and change the taste of the undesired properties of individuals.coprocessing excipients result in the formation of excipient granules with advanced properties. As compared with physical combinations of additives like improved flow property and compressibility. Better dilution capacity full uniformity and decreased lubricant sensitivity³².

Mechanism of action of superdisintegrants

- 1. Swelling
- 2. Porosity and capillary action(wicking)
- 3. Deformation
- 4. Enzymatic reaction
- 5. Due to disintegrating particle/particle repulsive forces

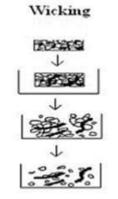
1.swelling

Swelling is thought to be a mechanism in which positive disintegrating agents (such as starch) gives the disintegrating effect. By swelling come in contact with water, the adhesiveness of different components in a tablet is overcome causes the tablet to disintegrate. E.g. sodium starch glycolate, platagoovata^{34,35,36}.



2.porosity and capillary action(wicking)

tablet porosity provides the pathways for the penetration of liquid into tablets. The disintegrant debris (with poor cohesiveness & compressibility) they act to enhance the porosity and provide these pathways into the tablet. E.g. crospovidone, croscarmellose sodium³⁷.



Disintegrant pulls water into the pores and reduces the physical bonding forces between particles

Fig. Disintegration by wicking

3. Deformation

starch grains are usuallythought to be "elastic" in nature meaning that grains that are deformed beneath strain will goback to their uniqueshapewhile that stress is eliminated. The capability for starch to swell is higher in energy rich starch grains than it's far for starch grains which have not been deformed underneath stress³⁸.

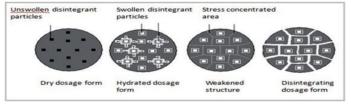


Fig. Disintegration by deformation

4.by enzymatic reaction

enzymes are present inside the body and also act as disintegranting agents. These enzymes gives the binding action of binder and helps in the disintegration.³⁹

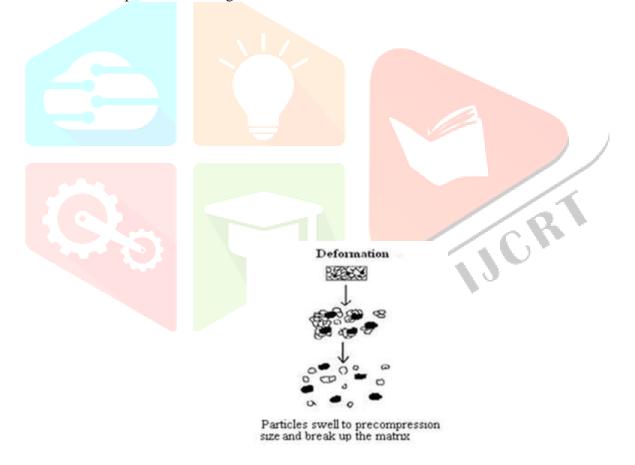


Fig. Disintegration by enzymatic reaction

5.due to disintegrating particle/particle repulsive forces

guyot-hermann has proposed a particle repulsion concept based on the commentary that nonswelling particle also cause disintegration of tablets. The electrical repulsive forces between debris are the mechanism of disintegration and water is needed for it. Researchers found that repulsion is secondary to wicking. It's miles believed that no single mechanism is responsible for the movement of most disintegrants⁴⁰.

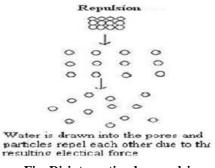


Fig. Disintegration by repulsion

Applications of super disintegrants

Mouth dissolving tablet- it is evaluated by the soy polysaccharide as a disintegrant in drugs made by the direct compression using lactose and di-calcium phosphate di-hydrate as fillers.

Fast disintegrating tablets- the preparation and evaluation of rapidly fast-dissolving tablets of metoclopramide by using the novel co-processed super disintegrant. Inside the present novel co-processed superdisintegrants had been advanced by a solvent. Evaporation approach the usage of crospovidone and sodium starch glycolate in different ratios (1:1 1:2 & 1:three) to be used in the fast-dissolving tablet formulations.

Rapidly disintegrating tablets- prepared olanzapine brief dispersing tablets by direct compression method. The impact of super disintegrant crospovidone on wetting time disintegration time drug content and in vitro released has been studied.

Disintegrating loadable tablets: a disintegrating loadable tablet product in compressed dosage form. A disintegrant or a aggregate of disintegrants has a) porosity of 45% v/v or extra b) a hardness of at the least 20 newtons and c) a loading ability of at least 30% of a liquid⁴¹.

Conclusion

The natural and synthetic superdisintegrants have a betterimpact on fast dissolving tablet. To increases the rate of drug release from tablet decrease dissolution and disintegration time and, natural polymers had been used as binder superdisintegrants and diluents. They may beeffectivelyavailable at a low price, are utilized in poor concentrations, and offerdietarysupplements becausethey may benaturally extracted. While disintegrants are moist, they expand and dissolve, causing the tablets to break apart and release the active ingredient for digestion. Natural polymers had beenalsoemployed as a fastener superdisintegrant and diluent because they increased the rate of drug discharge from the tableteven asdecreasing disintegration and dissolution. Water-insoluble superdisintegrants have a higher degradation property than the water dissolvable professionalsdue to the fact they do no longer swell. The tablets and capsules that arecurrently the maximumpopulartherapeutic dosage form for oral administration and have a few drawbacks for patientsundergoing chemotherapy and those with dysphagia. To conquer this trouble, mouth dissolving tablets containing guar gum and xanthan gum that should be considered the right detailing and shipping framework to improve affected person consistency, for the reason thatpercent drug transport and disintegration time are notablysuperiorto traditional dosage forms.

Reference

1.omidian h and park k: swelling agents and devices in oral drug delivery. Journal of drug delivery science and technology 2008; 18 (2): 83-93.

2. Bhardwaj s, jain v, sharma s, jatrc and jain s: orally disintegrating tablets: a review. Drug invention today 2010; 2(1): 81-88.

3. Beletmh and derle dv: analysis of patents pertaining to superdisintegrants used in tablet manufacturing. Journal of intellectual property rights 2008; 13: 601-604.

4. schimidt p c, and brogramann b., pharmaceutical technology. 1988 (34), 22.

5. Cohen y, and lach j l, journal of pharmaceutical sciences. 1963(52), 122

6.goel h, vora n, rana v: a novel approach to optimize and formulate fast disintegrating tablets for nausea and vomiting. Aapspharmscitech 2008; 9(3): 774-781.

7. Mohanachandranps, sindhumolpg and kirants: superdisintegrants: an overview. Journal of pharmaceutical sciences review and research 2011; 6(1): 105-109.

8.omidian h and park k: swelling agents and devices in oral drug delivery. Journal of drug delivery science and technology 2008; 18 (2): 83-93.

9. Iyad r, mayyasar, eftaiha aa and badwan a: chitin-silicon dioxide coprecipitate as a novel superdisintegrant. Journal of pharmaceutical sciences 2008; 97(11): 4955-69.

10.bhusnure o.g, gholves.b., giram p.s., thontes.s., mane j.m., kazip.a ., bhangem.a.; role of super disintegrating's in fast dissolving tablet; international journal of pharmacy & pharmaceutical research;2015,vol.4,issue:2.

11.vaibhay s, mahaveerpk, gupta k, agarwal d, sharma n, orally disintegrating tablet, friendly dosage form, international journal of research in ayurveda and pharmacy 1(2), 2010, 399-407.

12. Shaoorahmad, veshveshwaripanditsanjaydoshi, drshaikhsiraj r, afrozapatel, reviwe on the superdisintigrant and there phenomenon, world journal of pharmaceutical research, 7(17), 2013, 511-522.

13. Digpatiroy, debjitbhowmik, sampathkumarkp, a comprehensive review on superdisintegrants used in orodispersible tablet, indian journal of research in pharmacy and biotechnology, 2(4), 2014, 1297-1303.

14. Mohammed ismail, shaikkareemulla, mohamedabdulraheem, mohammedahmed, shaikgousebasha, zohebanjum, shafiqurrahman, formulation and evaluation of mouth dissolving tablets of amiodarone hcl by using natural superdisintegrants, international journal of current research, 9(2), 2017, 46761-46778.

15.kuchekar bs, bhisesb, arumugam v, design of fast disintegrating tablets. Indian j pharm educ. 2001;35(4):150-2.

16.digpati roy, debjitbhowmik, sampathkumarkp, a comprehensive review on superdisintegrants used in orodispersible tablet, indian journal of research in pharmacy and biotechnology, 2(4), 2014, 1297-1303.

17. Dhiraj a, khairnar, sanjay p, anantwar, chetan s, chaudhari, pravin, shelke, superdisintegrants, an emerging paradigm in orodispersible tablets, international journal of biopharmaceutics, 5(2), 2014, 119-128.

18. Omidian h, park k, swelling agents and devices in oral drug delivery, journal of drug delivery science and technology, 18(2), 2008, 83-93.

19. Iyad r, mayyasar, eftaiha aa, badwan, chitin-silicon dioxide co precipitate as a novel superdisintegrants, journal of pharmaceutical sciences, 97(11), 2008, 4955-69.

20.deveswaran s, bharath s, furtadobv, basavaraj s, abraham, madhavan, studies on the disintegrant properties of mucilage and seed powder of plantagoovata, international journal of chemical technology and research, 1, 2009, 621-626.

21.shirsand s, suresh m, para p, swamy, kumardn, plantagoovata mucilage in the design of fast disintegrating tablets, indian journal of pharmaceutical science, 7(1), 2009, 41-45

22. Raghavakuchimanchikp, sureshkumar, a detailed study on disintegrating agents and an overview on oral disintegration tablet, international journal of research in pharmaceutical and nano sciences, 5(3), 2016, 117-126.

23. Mohanachandranps, sindhumolpg, kirants, superdisintegrants, on overview, international journal of pharmaceutical science reviews and research, 6(1), 2011, 105-109.

24. Minke r, blackell j, the structure of alpha-chitin, journal of molecular biology, 120, 1978, 167-181.

25. Lavanya b, shanmugam v, formulation and evaluation of bisoprolol fumarate optizorb dispersible tablet to improve tablet disintegration, world journal of pharmacy and pharmaceutical sciences, 4(1), 2014, 561-576.

26. Arshiyafirdous, mohammedabdulkhaleq, sanaahmed, a review on optizorb technology, international journal for pharmaceutical research scholars, 5(4), 2016, 113-126.

27. Bruscato fn, danti ag. 1978 us patent 4086365.

28.dhiraj a, khairnar, sanjay p, anantwar, chetan s, chaudhari, pravin, shelke, superdisintegrants, an emerging paradigm in orodispersible tablets, international journal of biopharmaceutics, 5(2), 2014, 119-128.

29.lavika gandhi, semimulakhtar, comparative study on effect of natural and synthetic superdisintegrants in the formulation of orodispersible tablets, journal of drug delivery & therapeutics, 9(2), 2019, 507-513.

30. Shashikant n, sharma, ravindra s, sonawane, role of superdisintegrants in immediate release tablets a review, journal of pharmaceutical and bio sciences, 2017.

31.camarco w, ray d, druffner a, selecting superdisintegrants for orally disintegrating tablet formulations, pharmaceutical technology, 2006, supplement, 5

32.uddhav s bagul, current status of tablet disintegrants: a review. Retrieved 2011 frompharmainfo.net.http://www.pharmainfo.net/reviews/current-status-tablet-disintegrantsa-review.

33. K. P. Raghavakuchimanchi*1 and e. Suresh kumar2; a detailed study on disintegrating agents and an overview on oral disintegration tablet; / international journal of research in pharmaceutical and nano sciences. 5(3), 2016, 117 - 126

34. D. Bikashapathi, k. Saikrishna, u. A. Kumar & g. Sabitha, fast dissolving table: an update. International research journal of pharmacy. 2(3) (2011) 45-53.

35. Formulationvinensia.com

36. N. G. R. Rao, t. Ketan, s. Bala. Formulation and evaluation of fast dissolving tablets of metoprolol tartrate using natural superdisintegrant, international journal of pharmaceutical and clinical research. 2 (2010) 40-45.

37.d.chougule, ghodkedhananjay, r.r. shah, rahulghaste. Fast dissolving tablets: an overview. (2010).

38. G. G. Gajare, s. R. Bakliwal, b. R. Rane, n. A. Gujrathi, s. P. Pawar, mouth dissolving tablet: a review. International journal of pharmaceutical research and development (ijprd). 6 (2011) 280-296.

39.g. P. Kumar, r. Nirmala, fundamental aspects of superdisintegrants: a concise review. Journal of global pharma technology. 4 (2012) 1-12.4516.r. Pahwa, n. Gupta. Superdisintegrants in the development of orally disintegrating tablets: a review. International journal of pharmaceutical science and research. Vol. 2 (2011) 2767-2780.

40.v. D. Kumar, i. Sharma, v. Sharma. A comprehensive review on fast dissolving tablet technology. Journal of applied pharmaceutical science. 01 (05); (2011) 50-58.

41.jyotiverma*, Dr. S. K Prajapati and Dr. R Irchhiaya., AN OVERVIEW ON SUPERDISINTEGRANTS: A REVIEW, EUROPEAN JOURNAL OF PHARMACEUTICAL AND MEDICAL RESEARCH, 2017,4(09),252-260.

