Oro-dispersible tablets of ayurvedic powder for improving taste, compliance, ease and accuracy of administration

Deepak Khobragde*, Arun Kotha, K. Ravalika, Richa Gupta and P. Vasu Kumar

Vijaya College of Pharmacy, Munganoor, Hyderabad-501 511, Telangana, India

*Correspondence Info:
Dr. Deepak S. Khobragade
Professor; Pharmaceutics
Vijaya College of Pharmacy
Munuganoor (v), via Sanghi Nagar post,
Hyderabad-501511, R.R. Dist, (Telangana)
E-mail: ksdeepak31@yahoo.co.in

Abstract
Most of the ayurvedic medicines are in the form of powder. Being in powder form the administration of accurate dose with ease is a problem. They may have some kind of unacceptable bitter taste. Furthermore it needs water or honey for administration and chances of spoilage and waste are more. Oro-dispersible tablet which rapidly disintegrating in mouth will be the best remedy for efficient use of ayurvedic powders. The aim of this study was to formulate oro-dispersible tablets of ayurvedic polyherbal powder Talisadi. Talisadi is a traditional Ayurvedic powder preparation well known and effective in various disorders of respiratory and digestive system. Using various excipients like super-disintegrants and sweeteners, different formulation of polyherbal drug Talisadi were formulated by direct compression method. The acceptable formulation, among all the developed formulations of oro-dispersible tablet was having a disintegrating time of 1 min and 10 sec and acceptable taste. Thus the study concludes that formulation of oro-dispersible tablets of ayurvedic powder preparation can be a good option to enhance acceptability, efficiency and easy and accurate administration of powder ayurvedic preparations like Talisadi. Keywords: Talisadi, oro-dispersible, ayurvedic, poly-herbal powder, cough.

1. Introduction
Most of the dosage forms formulated, prepared and prescribed to the patients are solid formulations like powders, tablets and capsules etc and are mostly preferred to be administered by oral route. Oral route of administration of drug is the most popular method because of patient compliance and convenience in administration of drug. But because of unpalatable taste of drugs and difficulty in administration and swallowing in case of geriatric, pediatric, bed ridden patients, the drug cannot be given as such in oral conventional formulations may it be tablet or powders. There are chances of spilling and spoiling and spreading all over with powder formulations especially bulk powders. Powders have also problem of administration of accurate dosage and with ease due to their fine particulate nature. This leads to decrease in patient compliance [1,2]. To overcome this, a new approach i.e. development of mouth dissolving/dispersing solid formulations have generated a lot of research interest [3,4]. In these formulations, some of the drug gets absorbed through pharynx and esophagus while it is travelling to the stomach thus increasing the bioavailability and making it more beneficial than other conventional tablet dosage forms. Oro-dispersible [5] tablets are one of the kinds of formulations. These tablets get disintegrated in the saliva and dissolve [6-8]. These tablets are formulated with special type of excipient called super disintegrants to aid rapid disintegration, sweetening agent to mask the bitter taste, making it palatable and flavoring agent to leave a pleasant taste and feel in the mouth.

Talisadi is a ayurvedic polyherbal powder used to treat cough, cold and other disorders of respiratory tract [9-11] and many gastrointestinal disorders. Most of the ingredients present in the powder like long pepper and cinnamon are reported to have immune-modulators activity also act as bio enhancer. These Ayurvedic powders are taken 1-1/2 teaspoon along with honey because of the problem in administrating powder as such an unpalatable taste. So to avoid the drug adherence issue, carrying drug and honey, and various other disadvantages which arise due to Ayurvedic formulation these powder formulations are converted into MDT’s which have sufficient hardness, fast disintegration rate and pleasant taste [12-14].
2. Materials and Methods

Talisadi powder was purchased from Wonder Herbals Ltd Hyderabad India. All the other excipients were procured from S. D. Fine Chemicals Mumbai India. All reagents were of AR grade and used as received.

2.1. Preparation of oro-dispersible tablets:
Oro-dispersible tablets of talisadi powder were prepared as per the formula given in Table 1. Accurately weighed quantities of drug and excipients mannitol, crospovidone and diluents were passed through sieve number 100 and mixed in a glass mortar. The drug and directly compressible excipient were mixed by adding small portion of each at a time and blending it to get a uniform mixture. The other ingredients were mixed in geometrical order but magnesium stearate and talc were added at the last and mixed for further two minutes the blend was air-dried and compressed using CADMACH 8 punch rotary tablet machine at a fixed compression force. The mean weight and diameter of the tablets were 650 mg and 9 mm, respectively.

Table 1: Formula for various formulation of tablets (data in gm)

| Ingredients                  | F1  | F2  | F3  | F4  | F5  | F6  | F7  |
|------------------------------|-----|-----|-----|-----|-----|-----|-----|
| Talisadi                     | 500 | 500 | 500 | 500 | 500 | 500 | 500 |
| PVP                          | 15  | 15  | 15  | 15  | 15  | 15  | 15  |
| Mannitol                     | 20  | 20  | 20  | 20  | 20  | 18  | 18  |
| Lactose                      | 12  | 12  | 12  | 12  | 6   | 6   | 6   |
| Corn Starch                  | 66  | -   | -   | -   | -   | -   | -   |
| Sodium Starch Glycolate      | -   | 66  | -   | -   | 80  | -   | -   |
| Croscarmellose Sodium        | -   | -   | 66  | -   | 80  | -   | -   |
| Crospovidone                 | -   | -   | -   | 66  | -   | 80  | -   |
| Magnesium Stearate           | 6   | 6   | 6   | 6   | 6   | 6   | 6   |
| Talc                         | 6   | 6   | 6   | 6   | 6   | 6   | 6   |
| Trisodium Citrate            | 15  | 15  | 15  | 15  | 11  | 11  | 11  |
| Sodium Saccharine            | 10  | 10  | 10  | 10  | 8   | 8   | 8   |

2.2. Evaluation of Tablets:
All the formulation batches of oro-dispersible tablets were evaluated for the following parameters:

 **Weight variation:** Weight variation test can be a measure of uniformity of content. For this test, twenty tablets were randomly selected from each batch and individually weighed. The average weight of these selected tablets was calculated.

 **Hardness:** Tablet crushing strength, which is the diametric force required to break the tablet, was measured with a Pfizer tablet hardness tester. The hardness (crushing strength) of three tablets per batch was determined and mean taken.

 **Friability:** Friability is measure of strength of tablet. Tablet friability was measured using a ROCHE friabulator (USP) at 25 rpm for 4 min. The weight of ten tablets before and after completion of the test was recorded and friability was calculated by the following formula:

\[
\text{Percentage friability} = \left( \frac{\text{initial weight-final weight/initial weight}}{100} \right) \times 100 \quad \ldots \quad (1)
\]

 **Disintegration time:**
Three tablets per batch were evaluated for disintegration time by employing a digital disintegration apparatus. Water (1000 ml), maintained at 37±0.5 0C was used as disintegrating. Disintegration time was recorded when all the fragments of the disintegrated tablet passed through the screen of the basket.

 **Palatability Test:** The palatability of the formulated MDT’s was tested by Panel Method [15]. In this method 10 subjects were randomly selected and their mouth was cleansed with purified water. Tablets formulated were placed on each of the subject’s tongue and taste was evaluated after 10 seconds. The taste of a pure drug is used as a standard and the degree of bitterness of formulated tablets is judged by the volunteers. The results are classified as below, + = excellent taste masking; ++ = slightly bitter; +++ = very bitter.

4. Results and Discussions
All the results of various evaluator tests are shown in Table no. 2. From the results of the evaluation test of the tablets it can be observed that mannitol is a good diluents as it forms smooth and relatively strong tablets which disintegrate easily and rapidly when come in contact with disintegrating fluid. This is probably due to its water soluble nature as suggested by Debord et al[16].
It was also observed that among the various disintegrants used such as Corn starch, Crospovidone, Croscarmellose Sodium and Sodium Starch Glycolate, Crospovidone shows relatively faster disintegration time at same concentration as compared to others. It might be due to high water uptake capacity and low gelling capacity of Crospovidone when compared to others. Difference in swelling may also play a role in disintegrating agent efficiency It causes the tablet to disintegrate quickly.

**Table 3: Palatability results for Talisadi Tablets**

| Powder  | V1 | V2 | V3 | V4 | V5 | V6 | V7 | V8 | V9 | V10 |
|---------|----|----|----|----|----|----|----|----|----|-----|
| Tablet  |+++|+++|+++|++ |++ |++ |+++|+++|++ |++  |

+++ = very bitter; ++ = bitter; + = slightly bitter; 0 = no bitter taste

4. Conclusion
The study was successful in formulation of mouth dissolving tablets of Talisadi by direct compression method and using different super disintegrants. The aim of the study was to produce tablets that have palatable taste which is the major disadvantage of Ayurvedic formulations and this was successfully achieved. Thus it can be concluded that Ayurvedic formulations can be converted to ODT’s using different suitable excipients.

**Reference**

[1] Seager H. Drug-delivery Products and the Zydis Fast-dissolving Dosage Form. *J Pharm Pharmacol* 1998; 50:375-382.
[2] Tejvir Kaur, Bhawandeep Gill, Sandeep Kumar, G.D. Gupta. Mouth dissolving tablets: A novel approach to drug delivery. *International J Current Pharm Res*; 2010; 3:1-11.
[3] Bradoo R, Shahani S, Poojary S, Deewan B, Sudarshan S. Fast Dissolving Drug Delivery Systems. *J American Medical Asso*, India 2001; 4(10): 27-31.
[4] Kuccherkar BS, Badhan AC, Mahajan HS. Mouth dissolving tablets: A novel drug delivery system. *Pharm Times* 2003; 35:3–10.
[5] Dey P, Maiti S. Orodispersible tablets: A new trend in drug delivery. *J Natural Sci, Bio Medicine*. 2010; 1(1):2-5.
[6] Nayak SM, Gopalkumar P. Design and optimization of fast dissolving tablets for promethazine theoclolate. *Indian Drugs* 2004; 41: 554–6.
[7] Toshifusa S, Hideshi S, Kenji H, Kunio I. Studies of rapidly disintegrating tablets in oral cavity using co ground mixtures of mannitol with crospovidone. *Chem Pharm Bull* 2002; 50(2): 193-198.
[8] Abdelhary G, Prinderre P, Eouani C, Joachim J, Reynier JP, Piccerelle PG. The preparation of orally disintegrating tablets using a hydrophilic waxy binder. *Int J Pharm*. 2004; 278:423–33.

**Palatability test:**
The prepared films were given to volunteers of different age groups between 20 to 40yrs and evaluated for effectiveness of taste masking. Results showed that excellent taste masking was achieved with all formulations. Taste evaluation study of mouth dissolving films by panel method revealed that about 90% of the volunteers sensed no bitter taste. The results of taste evaluation by panel method are shown in Table No.3.

[9] Sharma Mukesh, Dewangan Manish, Aftab M.D et al. Development of quality control parameters of an ayurvedic formulation: Talisadi Churn. *Int Res J Pharmacy* 2012; 3(11):137-138.
[10] Difference-between-talisadi-and-sitopaladi-churna available at [http://easyayurveda.com](http://easyayurveda.com) accessed on 30/04/2016.
[11] Talisadi-churna-ingredients-benefits-dosage-side-effects available at [https://www.ayurtimes.com](https://www.ayurtimes.com) accessed on 30/04/2016.
[12] Anoop Agnihotri, Vijender Singh Formulation development and evaluation of antidiabetic polyherbal tablet *The Pharma Innovation Journal* 2014; 3(6): 01-03.
[13] Vijaya SRR, Anithakumari, Ramesh RV, Selvakumar Duraipandi. Design and Development of Tablets Containing High Amount of Polyherbal Aqueous Extract with Improved Disintegration Time. *International J Pharm Bio Sci* 2011; 2(1):135-139.
[14] Anil Tatiya, Pathak GP, Sanjay Surana, Dhanraj Mahajan. Formulation Development And Evaluation Of Fast Dissolving Polyherbal Tablets On Bronchitis *World J Pharm Pharm Sci* 2015; 5(1): 1401-1410.
[15] Patil Arun, Chafle Sandip, Kho bragde Deepak, Umathe Sudhir, Avari Jasmine. Evaluation of Hot Melt Coating As Taste Masking Tool. *International Res Journal Pharm* 2011; 2230-8407.
[16] Debord B, Lefebvre C, Guyot Hermann AM, Bouche R, Guyot JC. Study of different crystalline forms of mannitol comparative behavior under compression. *Drug Dev Ind Pharm* 1987; 13:1533-1546.
[17] Najib NM, Suleiman M, Malakh A. Characteristics of the in vitro release of ibuprofen from polyvinylpyrrolidone solid dispersions. *Int J Pharm* 1986; 32:229-236.