INTRODUCTION

Nyctanthes arbor-tristis is a traditional medicinal plant, which belongs to the family Oleaceae. The Nyctanthes arbor-tristis is a shrub or tree having fragrant flowers. The plant generally grows in the tropical and subtropical regions [1-3]. The Nyctanthes arbor-tristis having various names like Parijat, Night Jasmine, Gonal Jasmine, Harsinghar, etc. There are several Hindu religious stories related to the Nyctanthes arbor-tristis (Parijat). The holistic connection of the Parijat plant with Bhagwath Purana, The Mahabharata and Vishnu Purana. The Nyctanthes arbor-tristis is called as “Tree of Sorrows” because of the loss of the flowers their brightness during dark hours. The plant name arbor tristis means the sad tree [4, 5].

The present research is focused on the Arthritis, Nyctanthes arbor-tristis leaves were proven activity to cure Arthritis and gives relief from fever, pains and inflammation. The whole plant having various medicinal activities like anti-fungal, anti-diabetic, anti-oxidant [6, 7].

Arthritis means joint inflammation and it is chronic autoimmune disease occurs in any age group of peoples. The most common form of arthritis is Osteoarthritis and Rheumatic arthritis. Arthritis conditions tend to involves pain, aching, stiffness and swelling and in and around one of more joints. The symptoms can develop gradually or suddenly [8].

According to the Centers for Disease and Prevention (CDC) Arthritis is more common among adults aged 65 y or older, but it can affect peoples of all ages, including children. The cause of arthritis is the breakdown of cartilage. The function of Cartilage to protect a joint and allowing function smoothly. About 1% of the population is affected by arthritis women’s are three times more affected as compared to men’s. The prepared tablets were used to cure Arthritis and gives relief from fever, joint pain and inflammation [9].

MATERIALS AND METHODS

Materials

Nyctanthes arbor-tristis leaves were collected from the local area dried, powdered and used as an antipyretic, analgesic, anti-inflammatory to cure arthritis, joint pains etc. The excipients used in the formulation are Methylcellulose which is used as disintegrate, Magnesium stearate is used as a lubricant, Lactose is used as the diluent, Talc is used as a lubricant and gives the pleasant appearance to the tablet, and Acacia, HPMC-10, Sodium alginate these three excipients are used as the binder for the preparation of wet granulation.

Methods

Preparation of dry powder of nyctanthes arbor-tristis leaves

Collection of fresh leaves of Nyctanthes arbor-tristis from the local area. Clean the leaves by using distilled water. Leaves are dried at room temperature for a few days. The hot air oven is used for the complete drying of leaves. The dried leaves are collected and grind in a mixer to make a fine powder.

Preparation of 1% acacia solution

Take 100 ml distilled water in a beaker. Take 1 gm of acacia powder and mix in 100 ml distilled water. Stir continuously until all powder was mix properly.

Preparation of 1% HPMC-10 solution

Take 100 ml distilled water in a beaker. Take 1 gm of HPMC-10 powder and mix in 100 ml distilled water. Stir continuously until to form a jelly-like appearance.

Preparation of 1% sodium alginate solution

Take 100 ml alcohol in a beaker. Add 1 gm of Sodium alginate powder in 100 ml alcohol. Stir properly to mix well.

Formulation of herbal tablets

In this formulation, the dried leaves powder of Nyctanthes arbor-tristis was used to form a tablet dosage form. The formulation was done by following the wet granulation process and further compression by tablet punching machine.

Wet granulation method

Weigh all ingredients accurately, mix well and triturate by using mortar and pestle. The prepared 1% binding agent was added slowly to form a damp mass. Damp mass was transfer through sieves no. 22. Prepared granules are dried at room temperature. The well dried granules are ready for compression.
Table 1: Compression of formulation ingredients of tablet

| S. No. | Ingredients                | Quantity |
|-------|---------------------------|----------|
|       |                           | F1       | F2       | F3       |
| 1     | Nyctanthes arbor-tristis  | 250 mg   | 250 mg   | 250 mg   |
| 2     | Methyl cellulose          | 180 mg   | 180 mg   | 180 mg   |
| 3     | Magnesium stearate        | 20 mg    | 20 mg    | 20 mg    |
| 4     | Talc                      | 10 mg    | 10 mg    | 10 mg    |
| 5     | Lactose                   | 50 mg    | 50 mg    | 50 mg    |
| 6     | Acacia                    | 1%       | -        | -        |
| 7     | HPMC                      | -        | 1%       | -        |
| 8     | Sodium alginate           | -        | -        | 1%       |

Table 2: Scale of flowability

| Flow character | Hausner’s ratio | Carr’s index (%) | Angle of repose (°) |
|----------------|-----------------|------------------|---------------------|
| Excellent      | 1.00-1.11       | >10              | ≤25-30              |
| Good           | 1.12-1.18       | 11-15            | 31-35               |
| Fair           | 1.19-1.25       | 16-20            | 36-40               |
| Passable       | 1.26-1.34       | 21-25            | 41-45               |
| Poor           | 1.35-1.45       | 26-31            | 46-55               |
| Very poor      | 1.46-1.59       | 32-37            | 56-65               |
| Very, very poor| >1.60           | >38              | >66                 |

Table 3: Weight variation tolerance

| S. No. | Average weight of tablets(mg) | Max.% difference is allowed |
|--------|-------------------------------|-----------------------------|
| 1      | 80 or less                    | 10%                         |
| 2      | 80-250                        | 7.5%                        |
| 3      | More than 250                 | 5%                          |

Evaluation [10]

Pre-formulation study

Bulk density
Bulk density was carried out in 100 ml dried measuring cylinder. Pouring of dried granules in measuring cylinder and calculated by using the following formula:

$$\text{Bulk density} = \frac{\text{Mass of the granules}}{\text{Bulk volume of the granules}}$$

Tapped density
Tapped density was carried out by pouring of dried granules in 100 ml measuring cylinder. 100 tapping was done, note down the volume and calculate by using the following formula:

$$\text{Tapped density} = \frac{\text{Granules weight}}{\text{Volume of tapped granules}}$$

Hausner’s ratio
Hausner’s ratio is the ratio of the tapped density of granules to the bulk density of granules. Calculated by using the following formula.

$$\text{Hausner’s ratio} = \frac{\text{Tapped density}}{\text{Bulk density}}$$

Carr’s index
Carr’s index or compressibility index is determined by the following formula.

$$\text{Carr’s index} (%) = \frac{\text{Tapped density} - \text{Bulk density}}{\text{Tapped density}} \times 100$$

Table 2 shows the flow property of granules.

Table 3: Weight variation test

For each formulation, the hardness and thickness of 20 tablets were determined. Hardness test was determined by Monsanto hardness tester and the thickness of tablets was determined by Vernier Calipers. Results are shown in table 5.

Friability test
Friability of a tablets can determine in a laboratory by Roche friabilator. The friabilator consists of plastic chamber that rotates at 25rpm, dropping the tablets through a distance of six inches in the friabilator, which is then operated for 100 revolutions. The tablets are reweighed. Compress tablets loss less than 0.5% to 1.0% of the tablet weight are considered acceptable. Results are shown in table 5.

Disintegration time
This test was a time required for the tablet to separate into particles, the disintegration test measure only of the time required under a given set of aconditions for a group of tablets to disintegrate into particles. This test was performed to identify the disintegration of tablet in a specific time period.
RESULTS

The formulation was prepared by wet granulation method were tested for pre-formulation studies for the effective evaluation of tablets. All the evaluated pre-formulation parameters are shown in table 4. Based on the pre-formulation study the flow property of granules was good. The physical parameters of compressed tablets were shown in table 5. The compressed tablets color was Greenish white for F1 and F3, Dark greenish-white color for F2. The weight variation test, hardness, thickness, friability and disintegration time were shown in table 5.

| Table 4: Pre-formulation parameters for herbal tablets |
|--------------------------------------------------------|
| S. No. | Pre-formulation parameters | F1 | F2 | F3 |
| 1 | Bulk density | 0.267 g/cm³ | 0.29 g/cm³ | 0.18 g/cm³ |
| 2 | Tapped density | 0.325 g/cm³ | 0.33 g/cm³ | 0.23 g/cm³ |
| 3 | Carr’s index | 17.84% | 12.12% | 21.73% |
| 4 | Hausner’s ratio | 1.21 | 1.14 | 1.27 |
| 5 | Angle of repose | 27.29 | 25.74 | 26.22 |

| Table 5: Physical parameters for herbal tablets |
|------------------------------------------------|
| S. No. | Parameters | F1 | F2 | F3 |
| 1 | Weight variation test | 497±5% | 506±5% | 50.2±5% |
| 2 | Hardness (kg/cm²) | 3.3±0.17 | 3.13±0.01 | 3.13±0.01 |
| 3 | Thickness (mm) | 4.00±0.005 | 3.66±0.023 | 3.96±0.001 |
| 4 | Friability test (%) | 0.81% | 1.43% | 1.8% |
| 5 | Disintegration test (min) | 28 | 25 | 32 |

DISCUSSION AND CONCLUSION

Nyctanthes arbor-tristis was a traditional medicinal plant which having various medicinal activities but present research was focused on Arthritis and antipyretic, analgesic and anti-inflammatory activity. The leaves powder was used to formulate tablets. Wet granulation was done by using different binders and making three batches like F1,F2,F3. Pre-formulation study was carried out and gives good flow properties of prepared granules. The compression of prepared tablets, were evaluated and gives satisfactory results. The batch F3 was more disintegration time as compared to F1 and F2 batch. Based on the results it is concluded that the formulation and evaluation are good. The pharmacological evaluation is required for the treatment of Arthritis.

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AUTHORS CONTRIBUTIONS

All the authors have contributed equally.

CONFLICT OF INTERESTS

Declare none

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