

A formulation and evaluation of medicated chewing gum for cough

Pragati chute, Pooja Ghutke

Maharashtra Institute of Pharmacy (B.Pharm) Betala, Bramhapuri.

Date of Submission: 20-01-2025

Date of Acceptance: 30-01-2025

ABSTRACT-

Chewing gum is a mobile drug delivery system. It is potentially use for administrating drugs either locally or systematically via, the oral cavity. Herbal chewing gum has through the year gained increased acceptance as a drug delivery system. Several ingredients are now incorporated in herbal chewing gum. The aim of this study was to formulate herbal chewing gum using cinnamon oil and to prove its antibacterial property effectively at low dosage. Because cinnamon oil is a greater antibacterial agent against the Streptococcus mutans that causes tooth decay. The herbal chewing gum of cinnamon oil is prepared using different concentrations of cinnamon oil and yastimadhu. The prepared chewing is evaluated for different parameters like color, stickiness, weight variation, hardness, and in vitro drug release. In vitro drug release was performed by modification dissolution apparatus. It was seen that in formulation f2 drug release was increased compared to formulation f1 and f3. The formulation shows maximum drug release of 93% in 14 min. compared to all other formulations. So, chewing gum is used in the treatment of local disease in the oral cavity.

Keywords: Chewing gum, Cinnamon oil, Antibacterial

I. COUGH

A cough is the rapid expulsion of air from the lungs when the air passages are irritated. Cells within the air passages become irritated by fluids, mucus, or material, and a cough reflex is initiated, forcing air out of the lungs under air mass to clear the irritating substances and protect the lungs. A cough reflex is often voluntary or involuntary. We reassess the several sorts of coughs and their common causes and discuss symptoms you mustn't ignore after you have a cough.^[1]

TYPES OF COUGHS

A cough is taken into account as 'acute' if it lasts but three weeks and 'chronic' if it lasts longer than eight weeks in adults or four weeks in children. There are various kinds of coughs and

therefore the conditions related to those coughs can overlap.

Chesty cough

This is referred to as a wet cough or phlegmy cough; this makes a person's chest feel heavy and also the cough brings up mucus or phlegm. Each cough can produce a clump of mucus and thus these styles of coughs are called 'productive coughs'. Chesty coughs are caused by viruses from colds and flu and they can happen after an inflammatory disease. Chesty coughs are worse in the mornings as you tend to cough less while asleep, during which period the mucus produced by the cells within the air passages builds up overnight, resulting in excessive coughing and mucus expelling once you get up. Other more serious causes of chesty coughs include asthma, heart condition or bronchitis.^[2]

Dry cough

A dry cough, another non-productive cough, occurs because of irritants within the air passages, and causes can include colds, flu, hay fever, asthma, acid reflux, bronchitis, and certain medications used for treating high pressure.

Whooping cough

This can be a heavy infection that causes violent coughing fits. A vaccine against respiratory disease is mostly administered as a part of the vaccination program but the respiratory disorder can spread from non-immunized people to those that haven't yet received the vaccination including young babies. The 'whooping' sound occurs when the kid gasps for breath after a coughing fit. Babies might not cough or make the whooping sound but may gag and gasp.

Choking

Anyone presenting with a sudden episode of coughing or choking should seek immediate medical attention if a foreign body is present. For the patient's convenience, we prepared medicated chewing gum for cough. Due to the acceptance of oral drug delivery systems among people, chewing

gums soon became friendly to people all around the world because of their convenient administration. Besides its enjoyable taste and good feeling, it provides proven health, nutrition, and cognitive benefits.^[3]

CHEWING GUM

Chewing gum could be a soft, cohesive substance designed to be chewed without being swallowed. Modern chewing gum consists of gum base, sweeteners, softeners/plasticizers, flavours, colours, and a tough or powdered polyol coating. Its texture is paying homage to rubber because of the physical- chemical properties of its polymer, plasticizer, and resin components, which contribute to its elastic- plastic, sticky, chewy characteristics.



Figure.1

Chewing gum is being used worldwide since ancient times after man experienced the pleasure of chewing a variety of substance. One thousand years ago the Mayan Indians chewed tree resin from the sapodilla tree in order to clean their teeth and freshen their breath. Shortage of natural gum bases during World War II enhanced development of the synthetic gum bases that are used. Today Chewing gum can be used as a convenient modified release drug delivery system. Medicated chewing gums are currently available for pain relief, smoking cessation, travel illness, and freshening of breath. In addition, a large number of chewing gum intended for prevention of caries, xerostomia alleviation and vitamin / mineral supplementation are currently available.

The first commercial chewing gum —State of Maine pure spruce gum— was marketed in 1948 in the U.S.A. The first patent was filed in 1869. The gum was intended as dentifrices but it

has never been marketed. The first Medicated chewing gum —Aspergum— was launched in 1928. This chewing gum is still available and contains acetylsalicylic acid. Another commercially available medicated chewing gum is dimenhydrinate – containing chewing gum for motion sickness. However, chewing gum did not gain acceptance as a reliable drug delivery system until 1978, when nicotine chewing gum became available. Extended knowledge has made it possible to develop and manufacture medicated-chewing gum with pre-defined properties. Consequently, today chewing gum is a convenient drug delivery system, which is appropriate for a wide range of active substances.

Medicated chewing gum offers advantages in comparison to conventional oral mucosal and oral dosage forms both for (a) local treatment (b) systemic effect after absorption through the buccal and sublingual mucosal and from the gastrointestinal tract. Chewing gum can be retained in the oral cavity for a long period and, if the drug is readily absorbed across oral mucosa, chewing gum can provide a fast onset time for a systemic effect and the potential for avoidance of gastrointestinal and hepatic first – pass metabolism of susceptible drugs. Generally, medicated chewing gum has a good stability, the medicine can be taken easily and directly without the prerequisite of water, and if required, prompt discontinuation of medication is possible. Physiochemical properties of the drug like aqueous stability.

This is the foremost common sort of cough. It's caused by irritation within the throat and they tend to be the foremost annoying variety of cough non-productive cough because it produces little or no phlegm. Sometimes a post-nasal drip is felt which is caused by inflamed tissue within the nose producing excess mucus, which then drips down into the throat. This post-nasal drip triggers the cough reflex.

Foremost common causes are colds, flu, hay fever, or coryza. If the symptom is the reason for the tickly cough, then over-the-counter decongestants and antihistamines can help alleviate symptoms.

History of Chewing gum

Chewing gum could be a drug delivery system that's visiting advanced more and more in nowadays research. It seems to urge more standardization within the future industry because it can deliver either pharmaceuticals or nutrients referred to as medicated chewing gum (MCG). Ancient Greeks want to get a chewable resin from a

tree called mastic but thanks to archaeological diggings chewing gum-like substances or masticatory resins back to 5000 years ago. Resin pieces have even been found with teeth traces in Finland and Sweden. The first marketing of chewing gums was in 1848 when chicle from the Manilkara zapota was sapped. John Curtis and his son boiled spruce tree sap and added sugar, flavours and fillers, then rolled it and first made masticatory sticks which they wrapped in papers and sold them.

The first MCG was launched in 1924 in the US of America which was called Asper gum but an admission of chewing gum as a drug delivery system failed to gain until nicotine chewing gum was released on the market. Thomas Adams first manufactured MCGs with a natural latex-based and issued the first patent for the chewing machine to render chicle kneaded, and smooth but modern chewing gums often consist of synthetic resins. There is a monograph in European Pharmacopeia (EP) that defines MCG but the term —chewing gum‖ was first listed in guidelines as a pharmaceutical dosage form in 1991 and approved by the commission of European communities[4,]

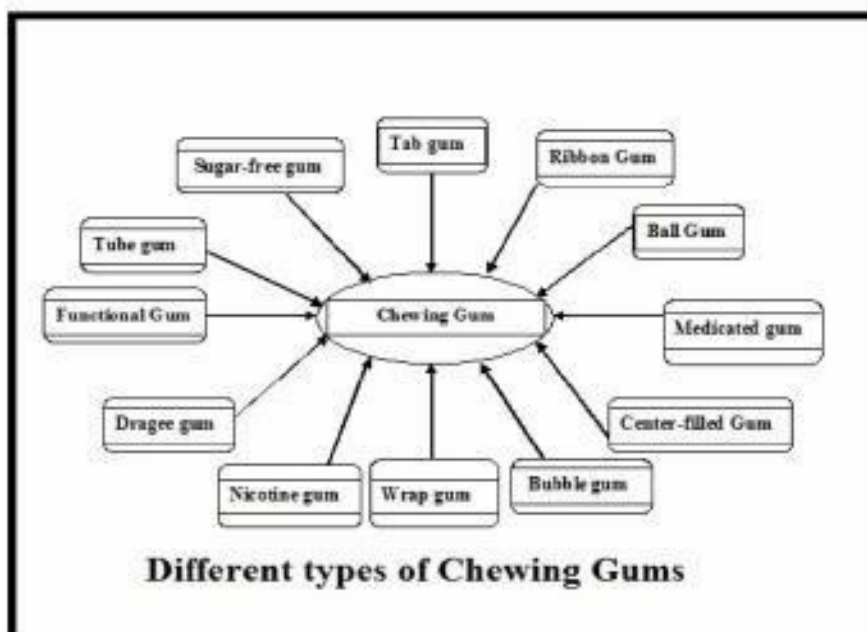


Diagram No.1 Different types of chewing Gums

II.

III. LITERATURE REVIEW

1. **Durga Devi. et.al.(2021),:-** The production of chewing gum in the global chewing gum market is rapidly increasing from USD 25

billion to an estimated USD 37 billion by the year 2023. The application of chewing gum in the drug delivery system has increased compared to pharmaceutical pre-dosage form.

However, gum bases are Split out after chewing which generates non-biodegradable waste causing harmful effects to the environment as well as living organisms. The review confers adequate information to increase the scale of production of biodegradable chewing gum for delivering bioactive compounds as well as zero-waste technology which sustainably reduces environmental pollution.

1. **Indumathi .et.al.(2020),:-** Chewing gums are mobile drug delivery systems. The extract of the herbal medicines may be incorporated into the chewing gum and may be utilized in the treatment of mouth ulcers. it was concluded that chewing gum is a wonderful drug delivery system for self-medication. Natural gum base is economical, safe, and environmentally utilized in the treatment of assorted mouth diseases.
1. **Azra Shake. et.al.(2017),:-** An attempt was made to formulate new chewing gum for Dolasetron. The new drug delivery system was obtained, at room temperature with conventional pharmaceutical equipment. The resulting chewing gum comprises a gum core combined with fillers, antioxidants, coloring agents, and plasticizers, which provide a smooth appearance and flexibility during storage and chewing. Drug release from a dosage form is the critical step in drug absorption and bioavailability, thus an experimental work has been designed to evaluate the efficiency of this kind of therapeutic system by verifying its capability to release the drug dose and by assessing the delivery of dolasetron for bypassing the hepatic first-pass effect. In the present study, an attempt has been made to formulate the chewing gum of Dolasetron. From this study, we can conclude that the medicated chewing formulation can be a better choice in the coming years which provides several benefits and also benefits commercially.
1. **Manasi paradhan. et. al(2016),:-** Medicated chewing gum (MCG) of Domperidone Maleate (DM) was developed by direct compression method to achieve quick onset of action and improve patient compliance. MCG containing DM was prepared by screening different concentrations of sweeteners, flavoring agents, softening agents, lubricants, and anti-adherents

by changing one variable at a time. Performance evaluation was carried out by evaluating size, shape, thickness, taste, scanning electron microscopy, texture analysis, and in vivo drug release study. The statistical analysis showed significant improvement in organoleptic properties such as chewable mass, product taste, product consistency, product softness, total flavor lasting time, and pharmaceutical properties. The developed formulation of medicated chewing gum can be a better alternative to mouth dissolving and conventional tablet formulation. It may be proved as a promising approach to improve bioavailability as well as improve patient compliance.

1. **Rahul Shetty. et.al (2016),:-**An attempt was made to formulate medicated chewing gum to prevent motion sickness using natural gum base for faster onset of action and easy administration, anywhere and anytime, without access to water. Chewing gum could be a soft, cohesive substance designed to be chewed without being swallowed. Modern chewing gum consists of gum base, sweeteners, softeners/plasticizers, flavours, colours, and a tough or powdered polyol coating. Its texture is paying homage to rubber because of the physical- chemical properties of its polymer, plasticizer, and resin components, which contribute to its elastic-plastic, sticky, chewy characteristics.
1. **Ritesh Kumar. et.al.(2014),:-** Explained that unlike chewable tablets medicated gums aren't alleged to be swallowed and should be off from the location of application without resorting to invasive means, and medicated chewing gum MCG could be a solid, single-dose preparation. Since it is taken anywhere, a chewing gum formulation is a wonderful choice for acute medication. the benefits for youngsters and for patients who find swallowing tablets difficult are obvious. The medicated chewing gums are solid, single-dose preparations with a base consisting mainly of gums that are intended to be chewed, but not swallowed. They contain one or more active substances, which are released by chewing and are intended to be used for local treatment of mouth diseases or systemic delivery after absorption through the buccal mucosa.

1. **Prashant P.Pagare.et.al.(2012),:-** Chewing gum is one of the very popular oral confectionery ducts. It's a potentially useful means of administering drugs either locally or systematically via, oral fissure. The medicated chewing gum has through the recent years gained increasing acceptance as a drug delivery system. It offers various advantages over conventional drug delivery systems. More over- medicated chewing gums require active and continuous masticatory activities for activation and continuation of drug release. An In-vitro apparatus was specially designed and constructed for release testing of medicated chewing gums. Medicated chewing gums are excellent mobile drug delivery systems for self- medication. It offers various advantages over conventional drug delivery systems.
1. **Vipul P. Patel. et.al. (2011),:-** Several ingredients are now incorporated in medicated chewing gum, ex. Fluoride for prophylaxis of dental caries, chlorhexidine as a local disinfectant, nicotine for smoking cessation, aspirin as an analgesic, and caffeine as a stay-alert preparation. In addition, a large number of chewing gum intended for the prevention of caries, xerostomia alleviation, and vitamin/mineral supplementation are currently .

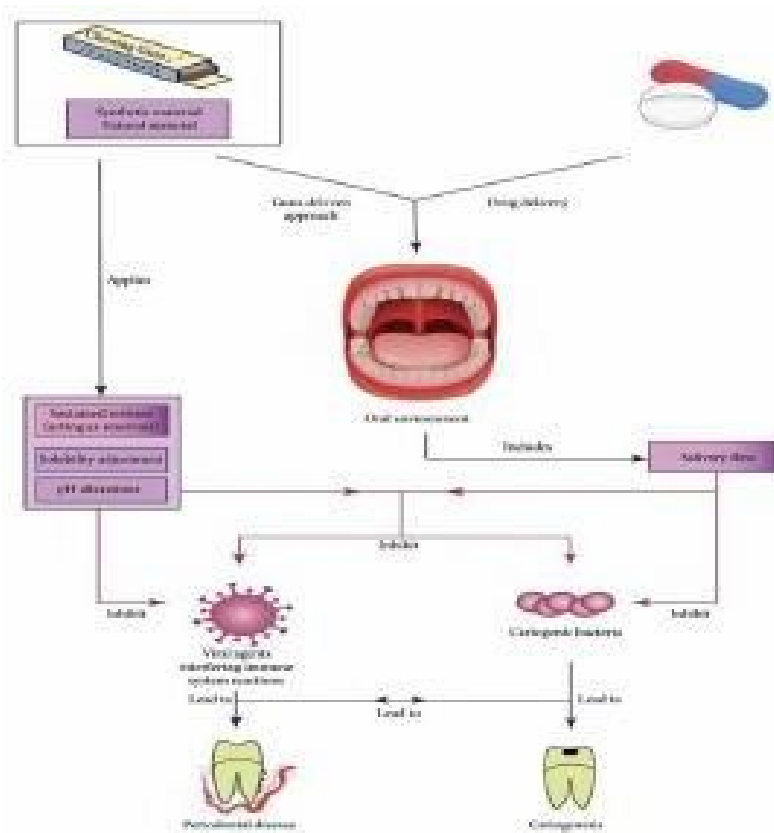


Figure No.3

Advantages of medicated chewing gums

1. Increased rate of effectiveness rather than another oral dosage form.
2. Use for both systemic and local action.
3. Does not require water to swallow, hence can be taken anywhere.
4. Avoid first-pass metabolism and thus increases the bioavailability of the drug.
5. Fast onset of action due to rapid release of the drug.
6. High acceptance by children and teenagers.
7. Good stability against light, oxygen, and moisture.
8. Convenient -promoting higher compliance .
9. Discreet less stigmatization.
10. Excellent for acute medication .^[5]

Disadvantages of medicated chewing gums

1. Getting choked by swallowing gum in under-aged children.
2. Risk of overdose.
3. Prolong chewing on gum may result in pain in facial muscles.
4. It may adhere to enamel dentures and fillers.

5. May cause stomach irritation and gastric ulcers.
6. Different release profiles because of chewing style differences.
7. Teeth decay through being coated with sugar.
8. Highly acceptable by children.
9. Pleasant taste .
10. Candidacies and caries.^[6]



Figure No.4

MATERIAL AND METHODS :-

DRUG PROFILE :

- Structural formula:

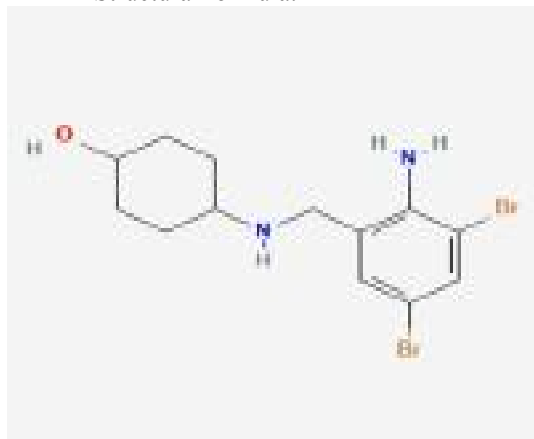


Figure No.2 Structural Formula of Ambroxol HCl

- **Synonym:** rac-cis-Ambroxol
- **Empirical formula:** C₁₃H₁₈Br₂N₂O
- **Chemical name:** 4-((2-amino-3,5-dibromobenzyl)amino)cyclohexane-1-ol
- **Boiling point:** 468.6°C at 760 mmHg
- **Melting point:** 235 - 240°C^[17]
- **Mechanism of action:** Ambroxol HCL is a clinically proven systemically active mucolytic agent. When administered orally onset of action occurs after about 30 minutes. The breakdown of acid mucopolysaccharide fiber makes the sputum thinner and less viscous and therefore more easily removed by coughing. Although sputum volume eventually decreases, its viscosity remains low for as long as treatment is maintained.^[18]

Gum For Cough .

- **Ambroxol HCL Indications:** All forms of tracheobronchitis, emphysema with bronchitis, pneumoconiosis, chronic inflammatory pulmonary conditions, bronchiectasis, bronchitis with bronchospasm asthma. During acute exacerbations of bronchitis, it should be given with the appropriate antibiotic.
- **Ambroxol HCL Contraindications:** There are no absolute contraindications but in patients with gastric ulceration relative caution should be observed.
- **Ambroxol HCL Side effects:** Occasional gastrointestinal side effects may occur but these are normally mild.
- **Precautions:** It is advisable to avoid use during the first trimester of pregnancy.^[11]

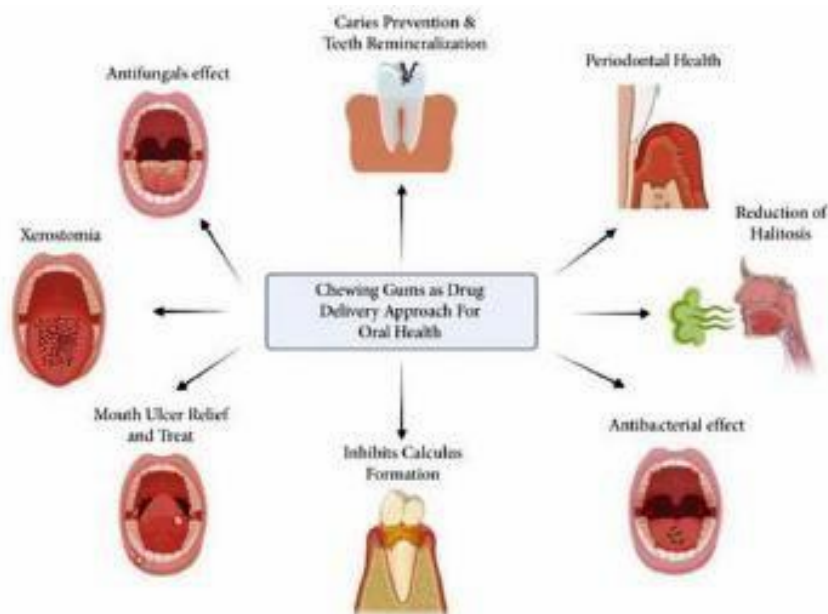


Figure No.5

V. METHOD FORMULATIN CHEWINGUM

Fusion method

The first step of a typical process for manufacturing chewing gum is to melt and soften the gum base at about 60°C and place it in a very kettle mixer, within which blades soften the bottom, then other ingredients like sugar, glycerine, sweeteners, taste-masking agent are added to the softened base, lately, the flavouring agent is added within the mixing procedure at 40°C, then cooling and rolling steps would be done, and also the rolled chewing gum would then delve pieces of desired shapes and sizes. A coating agent should be sprayed to make an identical surface to form a coated gum tablet.

The second form of this method is somehow different: the first step of preparation is to line up a mixer (the mixer may well be a sigma blade or other varieties of mixers), if a sugar-containing gum is required, the primary step is to feature syrup to the mixer, so finely granulated sugar is added gradually. Sugar, employed in this step, can be powdered sucrose, dextrose, fructose, syrup solids, or a mixture of them. Glycerine is the most preferred plasticizer used. Other components specified could be added to the matrix according to required characteristics, such as fillers, colorants, and flavouring agents. But it is recommended that

IV.

flavors be added to the matrix at the end of procedures when the gum base is totally and completely homogenized because most flavors are relatively volatile. The proportions of components in the matrix are variable between sources and depend on desired characteristics. But powdered sugar has approximately the most proportion.^[4,7]

The mechanical forces of the mixer, that is, compressive and shear, and warmth can ease the softening process. When no heat is applied, a better power is demanded. the blending process continues until a standardized mass is created. the blending process should last about 8 min. otherwise blending ingredients is to feature sugar gradually till the top of adding other components. After matrix preparation and completely mixing, the commercially prepared particles of gum base are added to the chamber all at once. But it is believed that these particles should have been heated and mixed before adding all other ingredients to the mass of the gum base. In this stage, mixing will continue for 10-20 min^[7]

Cooling, grinding, and tableting method
One other method to supply a chewing gum with the specified taste, color, and therefore the flavors is to combine the gum base with favorable and suitable sweeteners, corn syrups, starches, flavoring agents, and colorants, then refrigerate and funky it

with a freezer apparatus or by contracting with a coolant like CO₂ to a temperature below

-15°C which is therefore crushed and pulverized with a cutter or grinding apparatus to get minute particles then these finely ground particles are heated to a temperature which makes them adhere to every other and form a slick and uniform bulk with consistent texture and low relative density. If the fragments are such they are not self-adhesion and could be applied manually or mechanically before they're warmed to the conventional temperature to thereby promote self-adhesion. The cooling and grinding steps can be combined by cooling the grinding apparatus. After the grinding step, we can let the coolant (if used) evaporate and disappear from our desired composition. The minute particles may be coated with edible substances or premixed with powdery materials.^[8]

Direct compression

A new technology to create a chewing gum tablet is direct compression and tableting at the high-speed standard machine, but as explained during a patent, this manner of forming chewing gum tablets provides a quickly dissociable chewing gum, but after some seconds of chewing, particles adhere together to make an identical and homogenous mass. during this method; we'd like a granulating agent, preferably sorbitol, which might also act as a sweetener. A lubricant like magnesium stearate, talc, octadecanoic acid, hydrogenated vegetable oils, and sodium stearyl fumarate is added to the formulation before tableting. the primary step of this method is dry mixing of gum base, granulating agent, and a minimum of one processing material then adding an energetic ingredient, sweeteners, and other needed ingredients to the formulation within the free-flowing wine sort of materials then directly compressing the chewing gum tablets.^[9]

PREPARATION AND PROCEDURE :-

1. AIM AND OBJECTIVE

1. AIM

The experiment aims to prepare suitable or more convenient dosage forms for children or adults.

1. OBJECTIVE

1. To fasten the onset of action.
2. To get quick relief from cough.
3. To make a more suitable dosage form for pediatrics.
4. To increase patient compliance.



Figure No.7

VI. MATERIALS AND METHODS

4.1 Materials and Composition:

Sr. no.	Contents	Quantity	Functions
01.	Gum acacia	2.5 gm	Provides all the basic textural and masticatory properties.
02.	Xanthan gum powder	2 gm	As a thickening agent emulsifier and stabilizer .
03.	Sweetener(Mentha oil)	5 gm	To improve the taste of the gum base.
04.	Preservative (butylated hydroxytoluene)	0.1 gm	to extend the shelf life of the product.
05.	Antioxidant (ascorbic acid)	0.3 ml	to protect the gum base and flavors from oxidation.
06.	Colouring agent	0.2 ml	to give a required color.
07.	The active component Ambroxol HCL	10 Mg	to relieve cough and throat irritation.

Table.1

4.2

4.3 Method of preparation of MCG

Various methods are used to prepare chewing gum, that is mentioned in the introduction section but we prepare to chew gum with the help of the following steps :

1. Initially gum acacia and xylitol are ground with the help of a mortar and pestle to form a homogenous mixture.

2. later on we added a flavoring agent, antioxidant, Preservative, and coloring agent
3. In the third step we added glycerol to bind the above ingredients and made a dough.
4. Incorporation of the drug into chewing gum: we incorporate drugs in the liquid form in chewing gum by making cavity or molding chewing gum mass.



4.4 Dose of the drug: 10 mg per chewing gum

Initially, we take gm of chewing gum mass, then we added 8 mg of Ambroxol HCL in the chewing gum mass and the remaining 2 mg added in the liquid form by making a cavity or molding it.

FORMULATION AND EVALUTION :-

Evaluation of Gum Base Color: The color of gum base is RED GREEN color

[1]. Solubility Studies of Gum Base: The swelling power and solubility of gum base were determined according to leach et. (1959) with some modification as follows. An appropriate amount of dried sample was transferred into a per weighted centrifuge tube. Distilled water was added to give a total weight of mixture equivalent to (90g) and the mixture was stirred by a magnetic stirrer at (200rpm) to suspended the sample completely. The centrifuge tuber was then immersed in a water bath maintained at desired temperature for 30min. with continuous stirring. The stirring was removed and rinsed into the tube with sufficient distilled water to make the total weight of water equal. The tube was then stopped, inverted several times to mix the content, and centrifuged for one hour at 300rpm. The clear supernatant solution was taken decanted. An aliquot (25ml) of the supernatant was transferred quantitatively to a pre weighted porcelain evaporating dish. It was then evaporated to dryness on a steam bath and dried to constant weight at 70 (Table 2). As the gum base has showed very negligible solubility in phosphate buffer, it can be concluded that the procured gum base was the best for use as the base herbal chewing gum preparation.

Stickiness: The formulated herbal chewing gum was placed on plane surface. A mass of 250gm was hammered on it up to 10min, after 10min, sticking

of mass to hammered surface was observed [7].
Weight Variation Test: Chewing gum from each batch were individually weighted on analytical balance. The average weight and standard deviation were calculated which was found in acceptable unit [2].
Plasticity/Hardness: The hardness of chewing gum was determined by Monsanto hardness tester and the average hardness and standard deviation were reported



Filtration of Drug

[3]. Percentage Drug Content: One gram of formulation taken in mortar, to this about 20ml of 6.8 phosphate buffer was added and triturate. This was transferred to conical flask. About 30ml of 6.8 phosphate buffer was added to this and shaken well for about 3 hours using orbital shaking incubator at 100rpm. Then was filtered and the filtrate was made up to make with the same buffer solution. Suitable dilutions were made, and the drug concentration was determined by measuring the absorbance at 279nm

Drug soluble in water

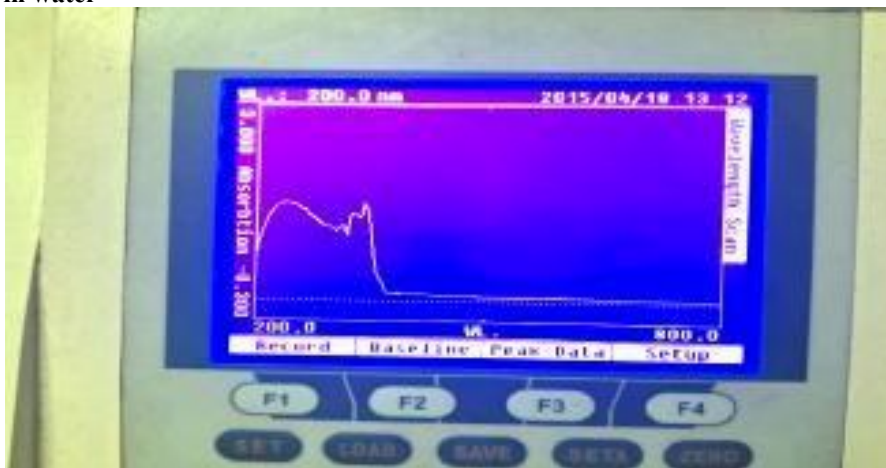
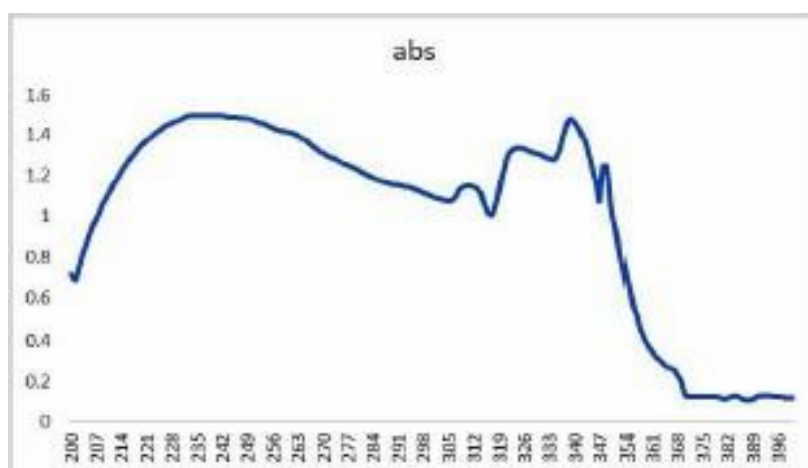


Figure No.1 Water



Graph : Water

Drug soluble in Ethanol

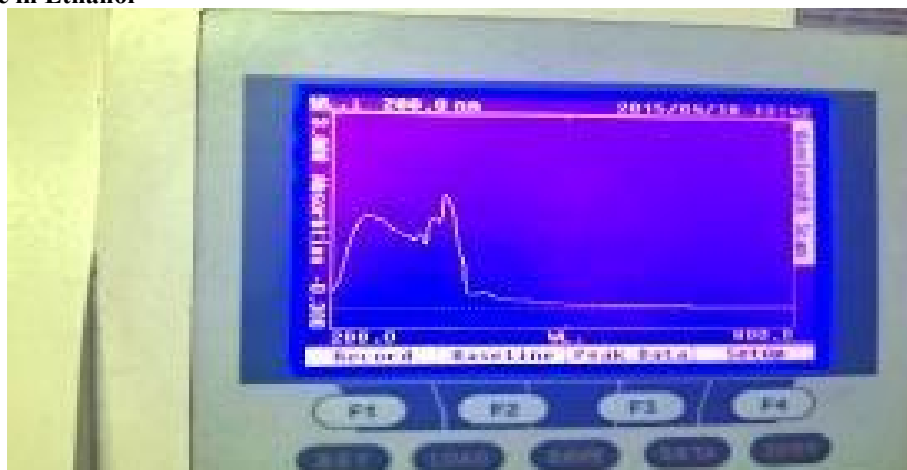
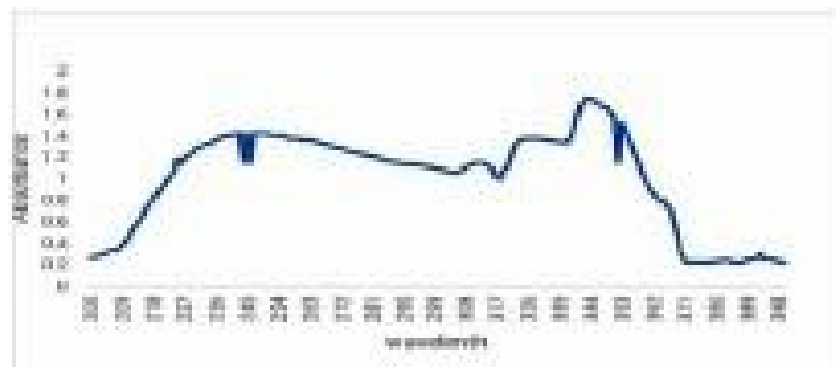


Figure No.2



Graph :-Ethanol

Drug soluble in phosphate buffer 0.4

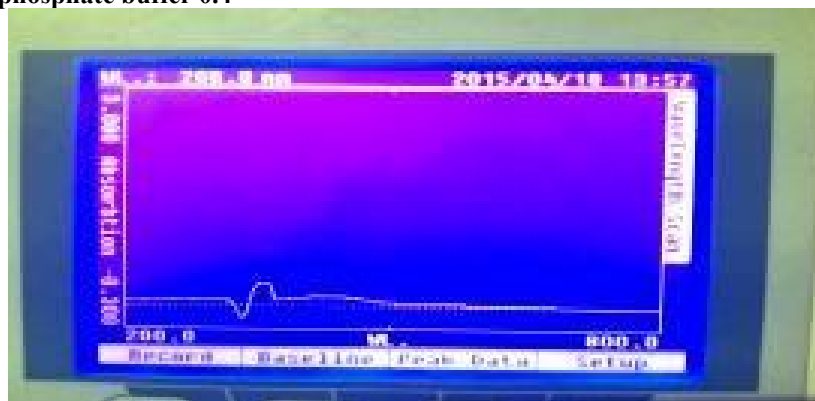


Figure No.3 phosphate buffer 0.4



Graph of phosphate buffer 0.4

Table of Ultra violet light

Sample	Absorption	Wavelength		
		F1	F2	F3
Water	1.449	242	241	240
Ethanol	1.740	344	-	-
Phosphate buffer 0.4	0.323	357	358	-

Table No.2



UV machine

[2] PH TEST:- The pH values of these gums are representative of six popular sugar gums (pH range 2.70–3.56) and seven sugar-free gums (pH range 6.58–7.57) investigated for this study. The enamel and dentine specimens were derived from surgically removed human third molars from 18-year-olds or older individuals of either gender



Figure No.7 Digital pH meter

4.5

4.6 Chewing gum packaging

The advantages of chewing gum packaging are clear to the world since it extends the shelf-life and gum stability. There are too many packaging methods with a wide range of options. In almost all the packaging types, we need a wrapping machine that receives and wraps the sticks of gums; in some cases, the wrapping machine seals the end of the package. In the following, a formed blister pack may be used then a foil will be heat-sealed at the back or a traditional packaging may be applied by lining the pellets up in a row and wrapping then sealing both ends.²⁵

The manufacturing and packing steps should be performed at about 20-25°C and relative humidity of 57%. Packaging has a substantial portion of the whole process both in terms of cost and time.²⁶

Undoubtedly, packaging increases the appearance of products among consumers, thus stylish design can attract more consumers to buy the specific product. Therefore, besides protecting the content, avoiding impurity, expediting transport, and improving storage, packaging can influence consumers.



Figure No.1 formulation chewing gum



Formulation 1



Figure.2 Formulation Chewing Gum



Formulation 2



Figure.3 formulation of chewing gum



Formulation 3

VII. RESULTS AND DISCUSSION

5.1 Result :-

We formulated medicated chewing gum by using gum acacia as gum bases and addition of excipients like coloring agent like coloring agent and flavoring agent thus taste was masked and ambroxol HCL was used as an active agent. We had a preception study we formulated it on a practical basics by the rough method. No subjecting of product on human volunteers was done as no clinical trial was brought into work. It is for providing an idea for future preparation that chewing gum can be used as a type of drug delivery system from as it is highly accepted and show an increased on set of action. So as per our view this types of dosage form can be for the formulation of any drug and polished products can be made further on an industrial level.

5.2 Stability

Chewing gum is a very stable product due to its low moisture content and less reactive nature than that other oral ingredients.²⁸

A major challenge in the production of chewing gum is its shelf life, storage conditions, and effect of some ingredients that impress stability. Water content can lead to the growth of microorganisms and chemical degradation, but water can be bound to other compounds, so that is not noticeably available to active agents, even if little water exists in the chewing gum is determined to annihilate and dangerous for other components, no water can be employed in the manufacturing method. To avoid oxidation of the drug, antioxidants are needed but due to the low content of water, the presence of preservatives is not essential.²⁹

Holding water content with no significant change at low or high concentrations of moisture in the atmosphere needs a major amount of gum base and xylitol as a bulking agent. Such made chewing gums are stable in extreme conditions and are capable of adding more desired ingredients and active agents. No stiffening, compacting, or softening is observed in terms of moisture stability in these kinds of gums.

Xylitol would also enhance storage stability in the gum; it means that in low or high humid conditions the gum's water content remains at its level and the gum flexibility, elasticity, splitting, and softness do not encounter major changes. The effect of xylitol on the crystalline structure and its water-binding properties are responsible for these changes.³⁰

In general, the stability of the active substance is good because chewing gum holding drug protects it from oxygen, light, and humidity. The high temperature for some heat-sensitive components to facilitate mixing can be avoided by increasing other powers, instead. Undesired interactions between different components can be prevented by encapsulation or coating some ingredients with suitable substances so that less contact between compounds occurs.³¹

One other important parameter involved in this topic is appropriate packaging and storage of gum to prevent water and moisture penetration and light exposure. The costly packaging and wrapping can be eliminated by spotting the above notes. Finally, the freshness and stability of gum would remain for a long period and the problem of staling, brittleness, and/or growth of microorganisms can be greatly reduced.³²

5.3 Factors affecting release rate and amount

In vivo and in vitro release of drugs from MCG is dependent not only on formulation factors but also on active ingredients' portion and individual chewing characteristics.

1. Water solubility: When the active ingredient is water-soluble, the release of the drug gets to the end rather than other active ingredients with slight water-soluble properties, and lipid-soluble drugs face further release problems than others because they are bound to lipophilic substances and gum bases and slowly released into the oral cavity.
2. Formulation factor: Mixing active ingredients with hydrophilic compounds or hydrophobic compounds affects the release of the drug. Sometimes faster release does not mean the more complete release of the drug; rather formulations with slower release profiles often show a more complete release of the drug.
3. Physicochemical properties: Ingredients more soluble in saliva will be immediately released within a few minutes of chewing, but highly lipid-soluble ingredients are first released into the gum base and then into saliva. The stability of gum base and its components to salivary enzymes, molecular mass, and ionization play an important role in the release and absorption of the drugs through the mucosa.
4. Individual characteristics: Speed, intensity, frequency, and type of chewing characteristics of different individuals affect the release of active ingredients; EP recommends 60 chews/min for the appropriate release of active ingredients. But these numbers of chews

depend also on the retention time of MCG in the mouth, which in clinical trials, the ordinary and suitable time is about 30 min of chewing. These differences lead to variable results of drug release.

5. The problem of adhering many lipophilic ingredients to gum base and other lipophilic compounds and, therefore, slow release of the drug into saliva may be solved by encapsulation of active ingredients or coating them with appropriate substances.^{33,34}



5.4 Problems associated with the manufacturing of chewing gum

1. Capping, lamination, picking, and sticking are the most common processing problems.
2. In the first method, one of the problems is that the inordinate content of moisture in the matrix may cause a low viscosity which reduces the shear and compressive forces, indeed more gum base particles are more likely to dissociate and float.
3. Heating and melting can make controlling the accuracy and uniformity of the drug difficult.
4. It is hard to provide sanitary conditions to make MCGs.
5. In the second method, the moisture content of chewing gum may cause the gum to jam to the blades and punches of apparatus, screens, surfaces, and chamber's wall.
6. In the second method caking and balling of the gum prevent the formation of gum fragments.
7. Forming a low-calorie chewing gum has resulted in gum with hard chew, poor texture, and bad taste or off-taste.
8. Sugar spots or lumps may appear in the final texture and cause an undesired feeling.
9. Some ingredients and active agents can irritate the mucosa.

10. High temperature to facilitate the mixture of gum base, leads to spoiling other ingredients.
11. Water elimination from final formulation requires advanced techniques to avoid the hardness of gum.^{35,36}

5.5 Safety Aspects

Generally, today it is perfectly safe to chew chewing gum. Previously, hard chewing gum has caused broken teeth. Extensive chewing for a long period may cause painful jaws muscle, and extensive use of sugar alcohol containing chewing gum may cause diarrhea. Long-term frequent chewing of gum has been reported to cause increased release of mercury vapors from dental amalgam fillings. However, medicated chewing gum does not normally require extensive chewing or a consumption to great extent. Flavors, colors, etc. may cause allergic reactions. Overdosing by the use of chewing gum is unlikely because a large amount of gum has to be chewed in a short period to achieve this. Swallowing pieces of medicated chewing gum will only cause the minor release of the drug because the drug can only be released from the gum base by active chewing. As a general rule, medicated chewing gum (like other medicines) should be kept out of reach of children, if required; drug delivery may be promptly terminated by removal of the gum.³⁷

5.6 Future trends

The future of chewing gum will reveal all of the scientists' efforts for the development of chewing gum as a modern drug delivery system and the progress of chewing gum production technology. In the future, other attempts will be seen to formulate more drugs using chewing gum as a drug delivery system. Treatment of fungal diseases, prevention of caries and other dental health issues, smoking cessation, etc., are common health works of MCGs. But remineralization of teeth, cold relief, energy enhancement, anti-nausea and so many new advantages of this novel drug delivery system are going to play an important role in future studies. MCGs are admissible alternatives to chewable or standard tablets and oral disintegrating dosage forms.

It takes time for chewing gum to get acceptance by people as a drug delivery system, but we hope that MCG finds its real place in industry and market and among patients soon, through its numerous advantages.

Long-lasting flavored, filled gums, timed-release, and other new MCGs formulated for diseases that previous delivery systems have been used for, are

trendy products to be seen in the future as a new kind of chewing gum that is made biodegradable and can be dissolved in around 1 month. We predict a brighter future for MCG as a novel drug delivery system than previous oral systems.³⁸

5.7 Problems overcome during formulation

5.7.1 Why do we use chewing gum for drug delivery?

- We used chewing gum for drug delivery because of its acceptance in children, its bioavailability is also good, it has a pleasant taste, and it is easily available. 5.7.2. How does the drug maintained in chewing gum?
- Initially, we take 7 gm of chewing gum mass which contain 8 mg of Ambroxol HCL in the chewing gum mass and the remaining 2 mg added in the liquid form by making a cavity or molding it.

1. DISCUSSION

5.1 PROBLEMS ASSOCIATED WITH THE MANUFACTURING OF CHEWING GUM :

Capping, lamination, picking, and sticking are the most common processing problems. 2. Heating and melting can make controlling the accuracy and uniformity of the drug difficult. 3. It is hard to provide sanitary conditions to make MCGs. In the second method, the moisture content of chewing gum may cause the gum to jam to the blades and punches of apparatus, screens, surfaces, and chamber's wall. In the second method caking and balling of the gum prevent the formation of gum fragments. Forming a low-calorie chewing gum has resulted in gum with hard chew, poor texture, and bad taste or off-taste. Sugar spots or lumps may appear in the final texture and cause an undesired feeling. Some ingredients and active agents can irritate the mucosa. High temperature to facilitate the mixture of gum base, leads to spoiling other ingredient. Water elimination from final formulation require advanced technique to avoid the hardness of chewing gum.^[12]

5.2 PROBLEMS OVERCOME DURING FORMULATION :

- We used chewing gum for drug delivery because of its acceptance in children, its bioavailability is also good, it has a pleasant taste, and it is easily available
- Initially, we took 7 gm of chewing gum mass, then added 8 mg of Ambroxol HCL to the chewing gum mass, and the remaining 2 mg

was added in the liquid form by making a cavity and molding it.

- Chewing gum is generally categorized as food but the addition of drugs makes it medicated so it is categorized in medication.
- We overcome the leakage of the drug by increasing the thickness of chewing gum thus making it tough.
- We can't mask the taste of the drug but by increasing the amount of sucrose during formulation, we can minimize the bitterness of the drug.

a. SAFETY ASPECTS :

Generally, nowadays it is perfectly safe to chew chewing gum. Previously, hard chewing gum has caused broken teeth. Extensive chewing for a long period may cause painful jaws muscle, and extensive use of sugar alcohol containing chewing gum may cause diarrhea. Long-term frequently chewing of gum cause increased release of mercury vapors from dental amalgam fillings.

However, medicated chewing gum does not normally require extensive chewing or a consumption to great extent. Flavors, colors, etc. may cause allergic reactions. Overdosing by the use of chewing gum is unlikely because a large amount of gum has to be chewed in a short period to achieve this. Swallowing pieces of medicated chewing gum will only cause the minor release of the drug because the drug can only be released from the gum base by active chewing. As a general rule, medicated chewing gum (like other medicines) should be kept out of reach of children, if required; drug delivery may be promptly terminated by removal of the gum.^[14]

b. FUTURE TRENDS :

The future of chewing gum will reveal all of the scientists' efforts for the event of chewing gum as a contemporary drug delivery system and therefore the progress of chewing gum production technology. In the future, other attempts are seen to formulate more drugs using chewing gum as a drug delivery system. Treatment of fungal diseases, prevention of caries and other dental health issues, smoking cessation, etc., are common health works of MCGs. But remineralization of teeth, cold relief, energy enhancement, and anti-nausea so many new advantages of this novel drug delivery system are visiting play a very important role in future studies. It takes time for chewing gum to induce acceptance by people as a drug delivery system, but we hope

that MCG finds its real place in industry and market and among patients soon, through its numerous advantages. Long- lasting flavored, filled gums, timed-release, and other new MCGs formulated for diseases that previous delivery systems are used for, are trendy products to be seen within the future as a replacement reasonably chewing gum that's made biodegradable and may be dissolved in around 1 month. We predict a brighter future for MCG as a unique drug delivery system than previous oral systems.^[15,16]



Figure No.8



Figure No.9

VIII. SUMMARY

Thus, it can be concluded that chewing gum can be used, as a carrier for delivering the drugs in the mouth cavity to relieve cough. chewing gum produced local as well as systemic action on the throat and suppresses the cough by producing fast action. Chewing gum can be used without water, at any time. Medicated Chewing gums can also produce both local effects as well as systemic effects in the various oral cavity infections. They can be used for taste masking certain drugs too. thus, the use of medicated chewing gum is increasing day by day.

IX. CONCLUSION

Thus it can be concluded that the chewing gum can be used as a carrier for vast categories of drug were extended release and the local action is described . Chewing gum can be used without water at any time medicated chewing gum can produce both local effects as well as systemic effects in the oral cavity they can be used for the purpose of taste masking of certain drugs . Chewing gum not only of offerce clinical benefits but also is an attractive distinct and effect efficient drug delivery system . Clinical trais have conform the benefits that can be gainted by exploting the effects the chewing gum is of delivery and apporunitis for buccal cavity absorption and local effects . The current manuscript present many of the petented applicatin ued in the field of chewing gum.Managment of fungal infection prevention of carries, smoking cessation ,decaying gums .

REFERENCES

- [1]. Balbani A.P., Cough: Neurophysiology, Methods of Research, Pharmacological Therapy, and Phonoaudiology, International Archives of Otorhinolaryngology, 2012;16(2),259–268.
- [2]. Chung K. F., Bolser D., Davenport P., Fontana G., Morice A., & Widdicombe, J.Semantics and Types of Cough, Pulmonary Pharmacology & Therapeutics,2009; 22(2),139–142.
- [3]. Paradkar M, Gajra B, Patel B, Formulation Development and Evaluation of Medicated Chewing Gum of Anti-Emetic Drug, Saudi Pharmaceutical Journal, 2016; 24:153-164.
- [4]. Shah KR, Mehta TA, Medicated Chewing Gum- A Mobile Oral Drug Delivery System, International Journal of PharmTech Research, 2014; 6:(1)35-48
- [5]. Pathan JA, Nitalikar MM, Formulation and Evaluation of Medicated Chewing Gum Containing Antibacterial Agent. Journal of Current Pharma Research 2014; 4(4):1291-1296
- [6]. John F. Golding. Motion Sickness Susceptibility, Autonomic Neuroscience: Basic and Clinical. 2006; 129:67-76.
- [7]. Neil A. Minton. Volunteer Models for Predicting Antiemetic Activity of 5-HT3 Receptor Antagonists, British Journal of Clinical Pharmacology, 1994; 37:525-530.

- [8]. Ezhumlai K, Rajalakshmi A.N., Ilavarasan P, Sathiyaraj U., Murali Mugundhan R. Medicated Chewing Gum- A Novel Drug Delivery Technique For Systemic and Targeted Drug Delivery, International Journal of Pharmacy & Technology, 2011; 3:725-744.
- [9]. Patel VP. Medicated Chewing Gum: A Review, International Journal of Universal Pharmacy and Life Sciences, 2011; 1:111-128.
- [10]. Savaliya P, Karikar A, Ramana MV. Chewing Gum: A Modern Era of Drug Delivery, The International Research Journal of Pharmacy, 2011; 2(10):7-12.
- [11]. Kumar D, Rathi L, Tripathi A, Maddheshiya YP. A Review on Oral Mucosal Drug Delivery System, International Journal of Pharmaceutical Science and Research. 2010; 1:50-56.
- [12]. Surana AS. Chewing Gum: A Friendly Oral Mucosal Drug Delivery System, International Journal of Pharmaceutical Science Review and Research. 2010; 4:68-71.
- [13]. Jain H, Shah M, Shah B, Pasha TY, Medicated Chewing Gum: A Novel Oral Drug Delivery, International Journal of Drug Formulation & Research, 2010; 1:80-96.
- [14]. Venkat Matti U, Adla N, Rajakannan T, Valakkathala R. Insulin Chewing Gum: Need of the day for Diabetic patients, International Journal of Pharmaceutical Investigation. 2011; 1:131-134.
- [15]. Naik H, Gupta S. Medicated Chewing Gum- Updated Review, International Journal of Pharma Research & Development, 2010; 2:66-76.
- [16]. Pandey S, Goyani M, Devmurari V. Development, In-Vitro Evaluation and Physical Characterization of Medicated Chewing Gum: Chlorohexidine Gluconate, Scholars Research Library, 2009; 2:286-292.
- [17]. Jayachandar Gajendran, Johannes Kraemer, Stig Randers Knudsen. Product Performance Test for Medicated Chewing Gums, Pharmacopeial Forum, 2008; 34:843-847.
- [18]. Catharina Kvist L, Sven-Bo Rje Andersson, Johan Berglund, Bo Wennergren, Susan M. Fors. Equipment for Drug Release Testing of Medicated Chewing Gums, Journal of Pharmaceutical and Biomedical Analysis, 2000; 22:405-411.
- [19]. Akbal Ö, Cevher E, Araman AO. The development and in vitro evaluation of benzydamine hydrochloride medicated chewing gum formulations. iujp. 2017;47(2):45-51.
- [20]. Ravindra Semwal, Deepak Kumar Semwal, Ruchi Badoni: Chewing Gum: A Novel Approach for Drug Delivery. The Journal of Applied Research 2010; 10(3):124-131.
12. Navneet Sharma, Deepak Kaushik and Harish Dureja: Medicated chewing gums: A potential drug delivery system. Pharma buzz 2008;
- [21]. Michael Moore, Nathalie Hasler-Nguyen, and Geoffrey Saroea: In vitro tooth whitening effect of two medicated chewing gums compared to a whitening gum and saliva. BMC Oral Health 2008; 8(23):1-7.
- [22]. Athanikar Narayan K: Process for manufacturing a pharmaceutical chewing gum. US2000015282.2001.
- [23]. Aspirin Oral Gum-Aspergum. Available from: www.wikipedia.com/aspergum/
16. Nicorette-Stop Smoking Gum. Available from: www.nicorette.com
- [24]. Thomas Imfeld: Chlorhexidine-containing chewing gum. Schweiz monatschrz ahmed 2006; 116(5):476-83.
18. Stay Alert: Caffeine Chewing Gum. Available from: www.stayaalertgum.com
- [25]. Biradar S S, Bhagavti S T, Hukkeri V I, Rao K P, Gadad A P: Chewing Gum As Drug Delivery System. Scientific journal articles 2005. Available from: <http://www.vipapharma.com/>
- [26]. <http://www.fertinpharma.com>
21. T.Imfeld: Chewing Gum--Facts and Fiction: a Review of gumchewing and oral health. Critical reviews in oral biology & medicine 1999; 10(3):405- 419.
- [27]. Xerostomia. Published by National institute of dental and craniofacial research under National Institute of Health. Available from: www.nidcr.nih.gov/OralHealth/Topics/Dry Mouth C.
- Dawes: Absorption of urea through the oral mucosa and estimation of the percentage of secreted whole saliva inadvertently swallowed during saliva

- collection. Archives of Oral Biology 2006; 51:111—116. 24. Gary H. Kamimori a, Chetan S. Karyekar b, Ronald Otterstetter. [28]. Zoft Hoodia functional chewing gum for weight loss. Available from: www.zoft-hoodia-gum.com 29. Nazneen Z. Ratlam: Chewing gum as drug delivery system. Available from: <http://www.pharmainfo.net/> [29]. Hitesh Jain, Mansi Shah, Bhoomi Shah, and T. Y. Pasha: Medicated chewing gum: A Novel oral drug delivery.