

Review Article

Quick Reference to the Matrix Forming Gums and Mucilage

U. Raghu¹, A. A. Hindustan², K. Haritha¹, J. Anji¹, G. Sudhakar¹, G. Vijayalakshmi¹

¹Department of Pharmaceutical Sciences, Jawaharlal Nehru Technological University-OTPRI, Ananthapuramu-515001, AP, India.

²Department of Industrial Pharmacy, Raghavendra Institute of Pharmaceutical Education and Research (RIPER), Ananthapuramu-515001, AP, India.

*Corresponding author's e-mail: abduhindustan@gmail.com

Abstract

The main aim of the present work is to explore the past work done on gums and mucilage as release retarding matrix forming material. Use of gums and mucilage as matrix forming material in drug delivery systems has been weighed down by the synthetic materials. Natural based excipients offered advantages such as non-toxic, less cost and abundant. The aqueous solubility of natural excipients plays an important role in their selection for designing sustained release formulations. This article provides an overview of natural gum and mucilage used as an excipient in dosage forms as well as in novel drug delivery systems as matrix forming materials. This article came with 65 different plant gums and mucilage successfully tried as release modifiers. From this article the researchers can save their valuable time by simply going in glance with the list of gums and mucilage successfully tried for making sustained release dosage forms.

Keywords: Gums; Mucilage; Plant; Sustained; Tablets.

Introduction

Gums are referred extracellular pathological products, formed by giving injury to the plant or due to unfavorable conditions (drought, the halt of cell walls). Mucilages are intracellular metabolic formations of plants. Exudates readily dissolve in water, whereas, mucilage forms slimy masses. Hydrolysis of gums and mucilages yields mixture of sugars and uremic acids [1,2].

Sustained release dosage form designed to achieve prolong therapeutic effect by continuously releasing medication over an extended period of time after administration of a single dose and also to control drug release profile at a specified rate to achieve desired drug concentration either in blood plasma or at the target site [3]. The sustained release dosage forms are of matrix (monolith) system or membrane (reservoir) system. In matrix system, the drug is homogeneously dispersed in a rate controlling medium [4]. Matrix systems are preferred as they show reduced drug fluctuations in plasma, decreased the total amount of drug in the dosage form, safety and improved patient

compliance [5]. The main pitfalls of these systems are cost of formulation, additional patient education and reduced potential for accurate dose adjustment [6]. Matrix systems are classified into different types as follows [7].

Hydrophilic matrix tablet

Hydrophilic matrix tablets are meant for oral administration. These can be compressed into tablets by direct compression. They retard the drug release, simple to prepare, economical and safe excipient. These systems deliver the drug over a defined period of time without disintegration.

Fat/wax Matrix systems

Fat/wax granulation involves spray congealing in the air, blend congealing in an aqueous media without the aid of a surfactant and spray drying technique. The medicament is blended with fat/wax matrix polymer by spray congealing technique (drug suspension melted with and melted with waxy polymers and allowed to solidify later sifted to get sustained release granules. These granules can be compressed into matrix tablets.

Plastic matrix systems

In this, the active medicament is dispersed in a tablet within a porous skeletal structure by direct compression. The medicament and the plastic polymer can be mixed/kneaded with organic solvent (for polymer). The drug release from these devices is delayed due to the diffusion of dissolved drug from the capillary pores of polymer. Use of hydrophobic matrices to sustain the drug release is usually delayed because the dissolved drug has to diffuse through a capillary network between the compacted polymer particles.

Biodegradable matrices

The monomers of these biodegradable polymers are biologically degraded /eroded by living cell enzymes or by metabolites of microbes.

Mineral matrices

These systems contain polymers which are obtained from various species of seaweeds.

Traditional excipients

Gums are pathological products of plants which readily dissolve in water, whereas, mucilage is physiological plant product [8]. The plant-based polymers are gaining importance nowadays in pharmaceuticals as a matrix system, film formers, mucoadhesive systems, micro/nano biphasic systems, implants, viscosity enhancers, stabilizers, disintegrants, solubilizers, and gel-forming agents are already proved effective. These traditional excipients are biodegradable, renewable, biocompatible, non-toxic, economical, eco-friendly, better patient tolerance, public acceptance and from edible sources [9]. The natural gums and mucilage used as release retention in dosage forms were shown in table 1 and table 2 respectively.

Table 1. Natural gums release retention

Common name	Botanical name	Family
Acacia gum	<i>Acacia catechu</i>	Leguminosae [10]
Albizia gum	<i>Albizia zygia</i>	Leguminosae [11]
Almond gum	<i>Prunus communis</i>	Rosaceae [12]
Ayoyo gum	<i>Cochorus olitorius</i>	Tiliaceae [13]
Badam gum	<i>Prunus amygdalus</i>	Rosaceae [14]
Bael gum	<i>Aegle marmelos</i>	Rutaceae [15]
Bihul gum	<i>Grewia occidentalis</i>	Malvaceae [16]
Carrageenan gum	<i>Chondrus crypsus</i>	Gigartineae [17]
Cashew gum	<i>Anacardium occidentale</i>	Anacardiaceae [18]
Cederela gum	<i>Cedrela odorata foliage</i>	Meliaceae [19]
Drumstick gum	<i>Moringa olifera</i>	Moringaceae [20]
Galbanum gum	<i>Ferula gummosa</i>	Apiaceae [21]
Gellan gum	<i>Pseudomonas elodea</i>	Leguminosae [22]
Ghatti gum	<i>Anogeissus latifolia</i>	Combretaceae [23]
Grewia gum	<i>Grewia mollis</i>	Malvaceae [24]
Indian Cherry gum	<i>Cordia obliqua</i>	Baraginaceae [25]
Jack fruit gum	<i>Artocarpus heterophyllus</i>	Moraceae [26]
Karaya gum	<i>Sterculia urens</i>	Sterculiaceae [27]
Khaya gum	<i>Khaya grandifolia</i>	Meliaceae [28]
Kondagogu gum	<i>Cochlospermum gossypium</i>	Bixaceae [29]
Leucaena seed gum	<i>Leucaena leucocephata</i>	Fabaceae [30]
Malva nut gum	<i>Scaphium scaphigerum</i>	Sterculiaceae [31]
Mango gum	<i>Mangifera indica</i>	Anacardiaceae [32]
Moi gum	<i>Lannea coromandelica</i>	Anacardiaceae [33]
Neem gum	<i>Azadiracta indica</i>	Meliaceae [34]
Odina gum	<i>Odina wodier</i>	Anacardiaceae [35]
Okra gum	<i>Abelmoschus esculentus</i>	Malvaceae [36]
Olibanum gum	<i>Frankincense</i>	Burseraceae [37]
Plum gum	<i>Prunus domestica</i>	Rosaceae [38]
Tamarind gum	<i>Tamarindus indica</i>	Fabaceae [39]
Tawa gum	<i>Beilschmiedia tawa</i>	Lauraceae [40]
Tragacanth gum	<i>Astragalus gummifer</i>	Leguminosae [41]
Welan gum	<i>Alcaligenes species</i>	Alcaligenaceae [42]
Xanthan gum	<i>Xanthomonas campestris</i>	Xanthomonadaceae [43]

Table 2. Mucilage release retention

Common name	Botanical name	Family
Aloe mucilage	<i>Aloe species</i>	Asphodelaceae [44]
Asario mucilage	<i>Lepidum sativum</i>	Brassicaceae [45]
Banana peel mucilage	<i>Musa paradisiaca</i>	Musaceae [46]
Basil seed mucilage	<i>Ocimum gratissimum</i>	Labiatae [47]
Bavchi mucilage	<i>Ocimum canum</i>	Lamiaceae [48]
Bidi leaf mucilage	<i>Bauhinia racemosa</i>	Fabaceae [49]
Broom creeper Mucilage	<i>Cocculus hirsute</i>	Menispermaceae [50]
Cactus mucilage	<i>Opuntia ficusindica</i>	Cactaceae [51]
Chinee apple mucilage	<i>Zizyphus mauritiana</i>	Rhamnaceae [52]
Date palm mucilage	<i>Phoenix dactylifera</i>	Palmaceae [53]
Fenurgreek mucilage	<i>Trigonella foenumgraenum</i>	Leguminosae [54]
Golden shower mucilage	<i>Cassia fistula</i>	Caesalpiaceae [55]
Hibiscus mucilage	<i>Hibiscus esculentus</i>	Malvaceae [56]
Humble plant mucilage	<i>Mimosa pudica</i>	Mimosaceae [57]
Ispagol mucilage	<i>Plantago psyllium</i>	Plantaginaceae [58]
Naga mucilage	<i>Brachystegia eurycoma</i>	Leguminosae [59]
Okra mucilage	<i>Abelmoschus esculentus</i>	Malvaceae [60]
Orange peel mucilage	<i>Citrus aurantium</i>	Rutaceae [61]
Red Cassia mucilage	<i>Cassia roxburghii</i>	Fabaceae [62]
Satavari mucilage	<i>Asparagus racemosus</i>	Asparagales [63]
Senna tora mucilage	<i>Cassia tora</i>	Caesalpiaceae [64]
Vanda mucilage	<i>Dendrophthoe falcate</i>	Loranthaceae [65]

Conclusions

There are great numbers of natural substances have been used in pharmaceutical preparations. Natural gifts viz., gums and mucilages can be used as matrix forming agents. They have been shown good potential as matrix former as well as other properties like fillers, disintegrating agent and sustain releasing agent. Natural gums and mucilages exposed good matrix property in wet granulation for the manufacturing of tablets/granules. Natural matrix formers are non-polluting renewable resources for sustainable supply of cheaper pharmaceutical excipients/product. Various applications of gums and mucilages have been established in the field of pharmaceuticals. However, there is a need to develop other natural sources as well as with modifying existing natural resources for the formulation of novel drug delivery systems, biotechnological applications and other delivery systems. From this article the researchers and young scientists can save their valuable time by simply going in glance with the list of gums and mucilage successfully tried in making sustained release dosage forms.

Conflicts of interest

No conflicts of interest are declared.

Acknowledgement

The authors wish to thank to Dr. Devanna, Director-JNTUA-OTPRI, Anantapur for his encouragement and support.

References

- [1] Kokate DCK, Purohit AP, Gokhale SB. Pharmacognosy, Nirali, 29th edition. 2008;29:51-57.
- [2] Liberman H, Lachman L. The theory and practice of industrial pharmacy. 3rd Ed. Verghese Publication House, Bombay, India. 1991.
- [3] Bhardwaj TR, Kanwar M, Lal R. Natural gums and modified natural gums as sustained-release carriers. Drug Development and Industrial Pharmacy 2000;26(10):1025-38.
- [4] Agarwal G, Kaushik A. Pharmaceutical Technology-II. 1st Ed. CBS Publishers, New Delhi. 2012.
- [5] Pundir S, Badola A, Sharma D. Sustained release matrix technology and recent advance in matrix drug delivery system: a review. Int J Drug Res Technol 2013;3 (1):12-20.

- [6] Kumar A, Raj V, Riyaz Md. Review on sustained release matrix formulations. *Int J Pharm Integrated Life Sci* 2013;3:1-14.
- [7] Indian Pharmacopoeia Commission. The Indian pharmacopoeia, 6th Ed. Ghaziabad, India. 2010.
- [8] Pal SK, Shukla Y. Herbal medicine. *Asian Pacific J Cancer Prevention* 2003;4:281-88.
- [9] Singh P, Mahmood T, Shameem A, Bagga P, Ahmad N. A review on Herbal Excipients and their pharmaceutical applications. *Sch Acad J Pharm* 2016;5(3): 53-7.
- [10] Khare B, Dubey N, Sharma A. Antiulcer activity of aqueous extract of *Acacia catechu wild* on rodent models by controlled release formulation. *Int J Curr Pharm Res* 2018;10(5);25-31.
- [11] Oluwatoyin A. Odeku. Assessment of *Albizia zygia* gum as a binding agent in tablet formulations. *Acta Pharm* 2005;55:263-76.
- [12] Sarojini S, Manavalan R Effect of natural almond gum as a binder in the formulation of diclofenac sodium tablets. *Int J Pharm Sci Res* 2010;1(3):55-60.
- [13] Muazu J, Alpha A, Mahammed GT. Isolation and release retardant properties of a plant gum obtained from ayoyo. *Carib J Sci Technol* 2014;2:301-13.
- [14] Ramesh KV, Shah F, Kiranmayi BH, Kumar KV. Design of sustained release matrix tablet of levofloxacin employing almond gum. *Int J Chem Sci* 2014;12(3):762-72.
- [15] Jindal M, Kumar V, Rana V, Tiwary AK. Physico-chemical, mechanical and electrical performance of bael gum-chitosan IPN films. *Food Hydrocolloids* 2013;30(1):192-9.
- [16] Vijay JK, Sati OP, Ranjit S. A potential natural tablet binder from *Grewia optiva*. 2011;3(3):120-27.
- [17] Baratam SR, Ratnam BV. Formulation and Optimization of Sustained Release Tablets of Rosuvastatin. Using HPMC K4M, HPMC K100M and Carrageenan gum. *Int J Chem Technol Res* 2018;11(5):376-86.
- [18] Maciel JS, Paula HCB, Miranda MAR, Sasaki JM, de Paula RCM. Reacetylated cashew gum: preliminary study for potential utilization as drug release matrix. *J Appl Polym Sci* 2006;99(1):326-34.
- [19] Michael AO, Adepeju OB, John OA. Evaluation of Cederela gum as a binder and bioadhesive component in ibuprofen tablet formulations. *Braz J Pharm Sci* 2013;49 (1):95-105.
- [20] Patel MT, Jitendra KP, Umesh MU. Assessment of various pharmaceutical excipients properties of natural Moringaoleiferagum. *Int J Pharm Life Sci* 2012;3(7):1833-47.
- [21] Shahbazi A, Lotfi M, Mostafavi KH, Asadian G, Heidarian AR. Effect of Persian galbanum (*Ferula gummosa* L.) extract on seed germination and growth of some weeds. *Afr J Agric Res* 2011;6(22):5106-11.
- [22] Kale VV, Kasliwal R, Parida SK. Formulation and release characteristics of guar gum matrix tablet containing metformin HCl. *Int J Pharm Expt* 2004;45:75-80.
- [23] Valluru R, Shivakumar HG. Investigation of kondagogu gum and ghatti gum as binders in formulating metoprolol tartrate tablets. *Res J Pharm Bio Chem Sci* 2013;4(2):1110-21.
- [24] Vijay JK, Sati OP, Ranjit S. A potential natural tablet binder from *Grewia optiva*. *Der Pharmacia Lettre* 2011;3(3):120-27.
- [25] Gupta R, Gupta GD. A review on plant *Cordia obluqua wild*. *Pharmacogn Rev* 2015;9(18):127-31.
- [26] Narkhede SB, Atul R. Bendale, Anil GJ. Isolation and Evaluation of Starch of *Artocarpus heterophyllus* as a Tablet Binder *Pharm Technol* 2011;3:836-40.
- [27] Reddy MM, Reddy JD, Moin A, Shivakumar HG. Formulation of Sustained-Release Matrix Tablets Using Cross linked Karaya gum. *Topical J Pharma Res* 2012;11(1):28-35
- [28] Odeku O, Fell JT. Evaluation of Khaya gum as a directly compressible matrix system for controlled release. 2004;56(11):1365-70.
- [29] Valluru R, Shivakumar HG. Investigation of kondagogu gum and ghatti gum as binders in formulating metoprolol tartrate tablets. *Res J Pharm Bio Chem Sci* 2013;4(2):1110-21.
- [30] Jeevanandham S, Sekar M, Dhachinamoorthi D, Muthukumaran M,

- Sriram N, Joysaruby J. Sustain-Release of Various Drugs from *Leucaena leucocephala* Polysaccharide. *J Young Pharma* 2010;2(1):15-20.
- [31] Ho YC, Norli I, Abbas F, Alkarkhi M, Morad M. Extraction, characterization and application of malva nut gum in water treatment. *Journal of Water and Health* 2015;13(2):489-99.
- [32] Anoop KS, Vipul KS, Panner RS, Sivakumar T. Evaluation of *Mangifera indica* gum as tablet binder. *International Journal of PharmTech Research* 2010;2(3):2098-2100.
- [33] Nayak BS, Nayak UK, Patro KB, Rout PK. Preparation and in Vitro evaluation of lamivudine entrapped MOI microspheres for oral administration. *Res J Pharm Technol* 2008;1(4):437-41.
- [34] Ogunjimi AT, Alebiowu G. Neem Gum as a Binder in a Formulated Paracetamol Tablet with Reference to Acacia Gum BP. *AAPS Pharm Sci Technol* 2014;15(2):500-10.
- [35] Dinda SC, Mukherjee B, Samata A. Odina gum: A novel matrix forming material for sustained drug delivery. *Orient Pharm Exp Med* 2011;11:131-6.
- [36] Ahad HA, Chitta SK, Ravindra BV, Sasidhar CGS, Harika B. Fabrication and in vitro evaluation of gliclazide *abelmoschus esculentus* fruit mucilage prolonged release matrix tablets. *Journal of Pharmacy Research* 2011;4(1):118-20.
- [37] Kebebe D, Belete A, Mariam TG. Evaluation of two olibanum resins as rate controlling matrix forming excipients in oral sustained release tablets. *Ethiop Pharm J* 2010;28:95-109.
- [38] Rahim H, Azam MK, Amin B, Kamran AC. Evaluation of *Prunus domestica* gum as a novel tablet binder. *Braz J Pharma Sci* 2014;50(1):195-202.
- [39] Phani KGK, Gangarao B, Kotha NSLR. Isolation and evaluation of tamarind seed polysaccharide being used as a polymer in pharmaceutical dosage forms. *Res J Pharm Bio Chem Sci* 2011;2(2):274-90.
- [40] Smale MC, Kimberley MO. Growth of naturally regenerated *Beilschmiedia tawa*. *New Zealand Journal of Forestry Science*. 1986;7:131-41.
- [41] Owen SC. Gum Tragacanth. In: Raymond CR, Paul JS, Paul JW. ed. *Handbook of Pharmaceutical Excipients*. The Pharmaceutical Press and the American Pharmaceutical Association, Washington, DC, USA. 2008.
- [42] Abhijeet P, Swati J, Jui J, Bhanudas K, Aniruddha C. In situ-gelling gellan formulation as vehicle for oral sustained delivery of famotidine. *Inventi Rapid/Impact: NDDS* 2012;3:1-4.
- [43] Dhopeshwarkar V, Zatz JL. Evaluation of xanthan gum in the preparation of sustained release matrix tablets. *Drug Dev Ind Pharm* 1993;19:999-1017.
- [44] Hindustan AA, Sreeramulu J, Hima BV, Padmanabha RY. Formulation and Evaluation of *Aloe barbadensis* Miller Mucilage Based Controlled Release Matrix Tablets of Glimepiride, *Asian Journal of Chemistry* 2009;21(8):6271-6.
- [45] Ramling P, Sachinkumar P, Suchita K. Evaluation of *Lepidium sativum* Mucilage as Suspending Agent in Paracetamol Suspension. *Indo American Journal of Pharm Research* 2011;1(3):181-6.
- [46] Bansal J, Malviya R, Malaviya, Bhardwaj V, Sharma PK. Evaluation of Banana Peel Pectin as Excipient in Solid Oral Dosage Form. *Global Journal of Pharmacology* 2014;8(2):275-8.
- [47] Saeed M, Semnani KM, Semnani J, Bazargani MH, Amin G. Evaluation of *Ocimum basilicum* L. seed mucilage as rate controlling matrix for sustained release of propranolol HCl. *Pharm Biomed Res* 2015;1(1):18-25.
- [48] Patel MM, Chauhan GM, Patel LD. Mucilage of *Lepidium sativum*, Linn (Asario) and *ocimumcanum*, sims. (Bavchi) as emulgents. *Indian J Hosp Pharm* 1987;24:200-2.
- [49] Gangurd AB, Boraste SS. Preliminary evaluation of *Bauhinia racemosa* Lam caesalpinaceae seed mucilage as tablet binder. *Int J Pharm* 2012;2(1):80-3.
- [50] Rao KM, Gnanaprakash K, Badarinath AV. Preparation and evaluation of flurbiprofen gel mucilage of *Cocculus hirsutus* leaf powder as gel base. *Int J Pharm Technol Res* 2010;2(2):1578-83.
- [51] Gebresamuel N, Mariam TG. Comparative Physico-Chemical Characterization of the Mucilages of Two Cactus Pears (*Opuntia* spp.) Obtained from Mekelle, Northern

- Ethiopia. Journal of Biomaterials and Nano biotechnology 2012;3:79-86.
- [52] Dalapath G, Kumar CS. Exploring Ziziphus Sps fruit mucilage as a pharmaceutical excipients in Novel Drug Delivery Systems. Journal of Innovation in Pharmaceutical Sciences 2018;2(2):17-21.
- [53] El-Far AH, Shaheen HM, Mohamed M, Mousa. Date Palm (Phoenix dactylifera): Protection and Remedy Food. Journal of Nutraceuticals and Food Science 2016;1:2-9.
- [54] Kuppusamy G, Kulkarni GT, Muthukumar A, Suresh B. Evaluation of fenugreek mucilage as gelling agent. Int J Pharma Excip 2002;2:16-9.
- [55] Choudhary PD, Pawar HA. Recently Investigated natural gums and mucilages as pharmaceutical excipients. Journal of Pharmaceutics 2014;2014: Article ID 204849.
- [56] Ahad HA, Rajesh V, Raghavendra GMV, Lasya DN, Harish N, Khamartaz M. Fabrication and in vitro Evaluation of Glimepiride Hibiscus esculentus Fruit Mucilage Sustained Release Matrix Tablets. Int J Pharm Technol Res 2010;2(1):78-83.
- [57] Singh K, Kumar A, Langyan N, Ahuja M. Evaluation of Mimosa pudica seed mucilage as sustained-release excipient. AAPS Pharm Sci Technol 2009;10(4):1121-7.
- [58] Deokar G, Kshirsagar S, Deore P, Deore H. Pharmaceutical benefits of Plantago ovate (Isabgol seed). Pharm Boi Eval 2016;427-33.
- [59] Olayemi O, Oremeyi J. Preliminary evaluation of Brachystegia eurycoma seed mucilage as tablet binder. Int J Pharm Res Inn 2011;3(1):1-6.
- [60] Narahari N, Palei, Santhosh K. Mamidi, Rajangam J. Formulation and evaluation of lamivudine sustained release tablet using okra mucilage. Journal of Applied Pharmaceutical Science 2016;6(9):69-75.
- [61] Reddy MR, Manjunath K. Evaluation of pectin derived from orange peel as a pharmaceutical excipient. Int J Drug Dev Res 2013;5(2):283-94.
- [62] Girpunje K, Kumaran A, Pal R, Maski N, Thirumoorthy N. A novel binding agent for pharmaceutical formulation from cassia roxburghii seeds. Int. J. P. Pharm. Sci. 2009; 1: 1-5.
- [63] Alok S, Jain SK, Verma A, Kumar M, Mahor A, Sabharwal M. Plant profile phytochemistry and pharmacology of Asparagus (shatavari). Asian Pac J Trop Dis 2013;3(3):242-51.
- [64] Singh, Pharmaceutical characterization of Cassia tora seed mucilage in tablet formulations. Der Pharmacia Lettre 2010;2(5):54-61.
- [65] Karthikeyan A, Rameshkumar R, Sivakumar, Amri S, Pandian SK, Ramesh S. Antibiofilm Activity of Dendrophthoe falcata against Different Bacterial Pathogens. Planata Medica 2012;78:1918-26.
