

## MICROENCAPSULATION OF ELLAGIC ACID FROM POMEGRANATE HUSK AND KARAYA GUM BY SPRAY DRYING

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### ABSTRACT

**Objective:** The aim of this study was to obtain and characterize microcapsules with Ellagic Acid (EA) from pomegranate as core material and Karaya Gum (KG) as wall material.

**Methods:** EA was obtained from dry pomegranate peel powder via methanolysis and quantified by HPLC. Microcapsules were obtained preparing a dispersion containing KG and EA in phosphate buffer pH 8. The dispersion was processed in a spray dryer under specific conditions (inlet temperature at 150 °C, feed flow at 30% and aspirator at 100 %) for obtaining of microcapsules. Fourier transform infrared (FTIR) spectroscopy, differential scanning calorimetry (DSC) and scanning electron microscopy (SEM) were used for characterization.

**Results:** Obtained material contains 98.03±2.82 mg EA/g of pomegranate peel. FTIR showed that there were changes in the molecular structure of microcapsules referred to raw materials. SEM confirmed that particles obtained had micron-size (1-5 µm). DSC analysis showed that raw materials had glass transition temperatures of 79.58 and 83.41 °C and for microcapsules the value was 67.25 °C.

**Conclusion:** Methanolysis is a viable technique for the obtaining of EA from the peel of pomegranate. KG shows good potential for be used as wall material for EA microencapsulation.

**Keywords:** Pomegranate husk, Ellagic acid, Karaya gum, Spray drier, Microcapsules.

### INTRODUCTION

Ellagitannins belong to the class of hydrolyzable tannins, which are polyphenolic compounds that are chemically characterized by a group named hexahydroxydiphenic acid (HHDP) usually linked to

glucose as polyol unit [1-3]. When the ellagic tannins are exposed to acidic or alkaline conditions, the ester bonds are hydrolyzed realizing the HHDP group, then a spontaneous lactonization occurs to get a more stable form, giving origin to the molecule known as ellagic acid (EA) (fig. 1) [4-8].

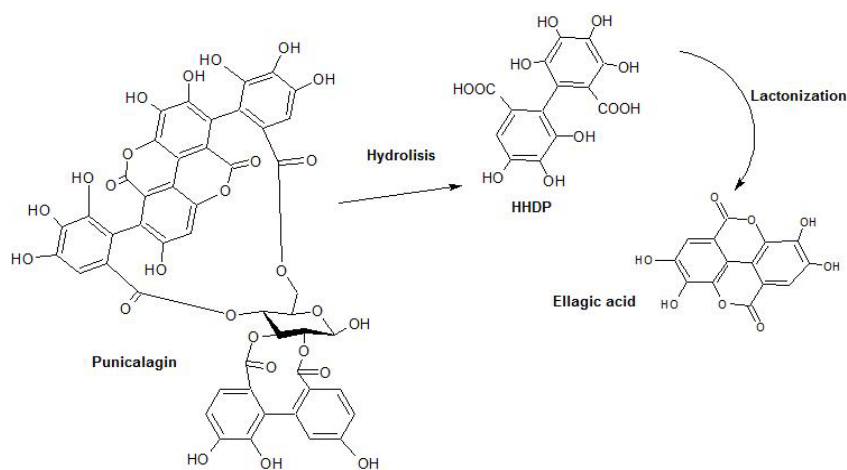


Fig. 1: Hydrolysis of an ellagitannin by Larrosa *et al.* [9]

The typical sources of these compounds are red fruits like strawberries, raspberries, blackberries and pomegranate, the content in these materials ranging from 1 to 1794 mg/100g [4, 10-

13]. Pomegranate peel is considered an agro industrial by product, however, several reports have demonstrated it contains great quantities of hydrolysable tannins, mainly punicalagin [14-18].

The EA or dilactone of HHDP acid, has generated many researches due to its important biological properties since they help to prevent cardiovascular and hypertension diseases, also has antitumor, antiviral, antimicrobial and antioxidant activity [19-22]. This knowledge has been used for development of functional foods and drugs. However, the effectiveness depends on the preservations of its stability and bioactive properties [23, 24]. An important method to prolong these properties is to use encapsulation technologies that help to protect the compounds of adverse conditions facilitating their storage and processing [25].

Microencapsulation by spray drying is an encapsulation technology that has been widely used in the food industry since the late 1950's and early 1960's [26, 27], this method allows the active compound to get into a spherical and strong semi-permeable polymer matrix called microcapsule, in order to protect it from adverse conditions such as light, temperature and oxygen which can cause undesired reactions [28, 29].

The microcapsule has a diameter ranging from 0.2 to 5000  $\mu\text{m}$ . The material packaged is also called core material, coated material, active, internal phase or payload. On the other hand, the packaging material is called wall material, capsule, carrier, shell or coating material.

In microencapsulation by spray drying the most common wall materials are bimolecular of three types: polysaccharides, lipids, and proteins. The polysaccharide is the most commonly used, especially maltodextrins of different polymerization degree, measured as dextrose equivalent (D. E. 10, 11, 18, 20 and 21) and Arabic gum [30, 31].

Karaya gum (KG) has potential to be used as a cover material for the encapsulation of active compounds by spray drying, due it has a complex and branched structure [32] which provides it with qualities such as emulsifier, cohesive and adhesive. These properties are essential for application in the food and pharmaceutical industries [31, 33, 34].

The aim of this work was to use KG as cover material for microencapsulation of a hydrolyzate dreich in EA obtained from pomegranate peel, and the characterization of prepared EA microcapsules.

## MATERIALS AND METHODS

The active material of the capsules was extracted from pomegranate peel, which was provided by the Department of Food Research at the Chemistry Faculty of the Universidad Autónoma de Coahuila. The peel was heat dried for 48 h at 60 °C; once dehydrated, the raw material was pulverized in a mechanical mill to obtain small particles with an average size of 2 mm. HPLC grade EA was obtained from Sigma-Aldrich. The packing material used was commercial grade KG purchased from a local business.

### Extraction of ellagic acid by methanol sis

The extraction of EA was performed following the methodology described by Lei *et al.* [35] with modifications. Briefly, 11.9 g of dry pomegranate peel powder was placed in a sealed container with 595 ml of metabolic sulfuric acid reagent (sulfuric acid at 19% in methanol); this mixture was placed in an oven at 80 °C for 30 h for recovery of EA. Water was added to the extract and subjected to sonic vibration for 30 min and centrifuged at 6000 rpm for 20 min to remove water-soluble compounds discarding supernatant. The pellet was resuspended in 96% ethanol, subjected to sonic vibration for 30 min and centrifuged at 6000 rpm for 20 min and finally the supernatant was eliminated to obtain a powdered solvent-free pomegranate peel hydrolyzate dreich in EA.

### Quantification of ellagic acid content in pomegranate peel hydrolyzated by High-resolution liquid chromatography (HPLC)

EA quantification was performed by High-Performance Liquid Chromatography (HPLC). Equipment for HPLC Varian Pro Star 430 was used for this analysis; UV-Vis photodiode array detector was used for the analysis and a Grace Denali C18, 5 $\mu\text{m}$  (250 mm x 4.6 mm) column with a flow rate of 1.2 ml min<sup>-1</sup> was used for sample

separation. Mobile phase for compound quantification were: methanol (solvent A), acetonitrile (solvent B) and 3% acetic acid (solvent C). The injection method by Poupard *et al.* [36] was used with some modifications: briefly, the elution gradient was as follows: initial 97% C, 3% B; 0-5 min 91% C, 9% B; 5-15 min 84% C, 16%B; 15-30 min 67% C, 33% B; 30-33 min 10% C, 90% B; 33-35 min 10% C, 90% B; 35-40 min 3% C, 97% B; 40-42 min 3% C, 97% B. To determine the amount of EA, a calibration curve using HPLC degree EA was constructed measuring standard solutions at different concentrations (100, 200, 300, 400 and 500 mg L<sup>-1</sup>) using A as solvent.

### Preparation of the feed liquid (Preparation of core material and wall Material dispersion)

The dispersion was prepared according to the ratio used by Medina-Torres *et al.*, in 2013 [37], which contained the rich-EA hydrolyzated as a core material and KG as wall material. KG was prepared at 1% per each L of phosphate buffer solution 0.2 M at pH 8, subsequently the dispersed phase contains 322.58 mg of hydrolyzated, added as an aqueous solution. The viscosity of the feed liquid was measured on a Viscometer Mark (Brookfield DV-II+Pro Extra) at 100 rpm and registered a viscosity of 37.87 centipoises (cP).

### Microencapsulation by spray drying

The feed liquid was processed in a Mini Spray Dryer Buchi 290 with an aspiration of 100% (35 m<sup>3</sup> h<sup>-1</sup>) using a nozzle with a diameter of 1.5 mm, in co-current flow, with an inlet temperature of 150 °C and pumping of 30 %.

### Fourier transforms infrared (FTIR) spectroscopy

FTIR spectra of Hydrolyzated, EA (HPLC grade), KG and microcapsules were obtained with a Perkin-Elmer 16 PC spectrometer (Perkin-Elmer, Boston USA), in Attenuated Total Reflectance mode (ATR) between 600 and 4500 cm<sup>-1</sup> with a spectral resolution of 4 cm<sup>-1</sup>. Each spectrum was base lined, and the absorbance was normalized between 0 and 1.

### Differential scanning calorimetry (DSC)

Differential Scanning Calorimetry (DSC) measurements were performed with a Shimadzu DSC-50 (Shimadzu Corporation Kyoto, Japan) calibrated using Indium as standard. Samples were weighed (approximately 5-10 mg of dry matter) in aluminum DSC pans. As a reference, an empty aluminum crucible was used. The heating rate was fixed at a temperature range of 40 to 200 °C for the hydrolyzated and karaka gum and from 30 to 300 °C for the microcapsules at a heating rate of 10 °C min<sup>-1</sup> in all assays.

### Scanning electron microscopy (SEM)

The morphology of the spray dried microcapsules was visualized by SEM with a Nova Nano SEM 200 scanning electron microscope (FEI, USA) with an acceleration voltage of 10 to 15 kV. The samples were mounted on a self-adhesive carbon tape and coated with gold before acquiring photomicrographs.

## RESULTS AND DISCUSSION

### Extraction, quantification and identification of ellagic acid in pomegranate peel

The pomegranate peel has shown a great potential for obtaining EA. Some authors have reported different concentrations of EA in this material: 1.3 [3] and 6.3 [13] mg of EA per gram of pomegranate peel respectively. This may be due that the first work reported only the free EA covered with a rapid, large-scale purification method, and the second one reported EA obtained from a fermentation process using pomegranate peel as substrate.

Our research group have an earlier publication [7], in this document 33.79 $\pm$ 7.43 mg of EA per gram of pomegranate peel obtained by methanol sis, whereas in the present study 98.03 $\pm$ 2.82 mg of EA per gram of pomegranate peel were obtained; this difference may be due that in the present study is a mixture of methanol and sulfuric acid was used as extraction solvent [35], instead of a mixture of methanol and HCl previously cited [7]. This difference detected may be due to the effectiveness of sulfuric acid to break of punicalagin

ester bounds is higher than hydrochloric, releasing larger amounts of HHD Pin the present in the raw material.

Fig. 2 shows the chromatogram of EA present in the hydrolyzated together with two points of the calibration curve

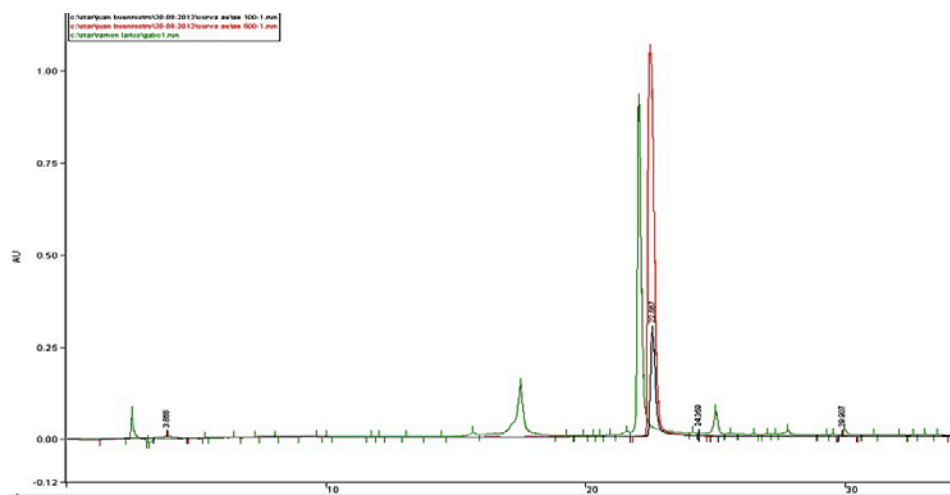


Fig. 2: Chromatograms of ellagic acid standard to different concentration (red 500 mg<sup>-1</sup>L and black 100 mg<sup>-1</sup>L), and chromatogram of hydrolyzed (green)

### Scanning differential calorimetry (DSC)

Glass transition temperature ( $T_g$ ) is a very important parameter in spray drying process since this property has been linked to the agglomeration and adhesion of the material in the drying chamber [38]. Different authors [38-40] mentioned that sticking occurs when working temperatures are 20 °C above  $T_g$  of the material.

In our process, the outlet temperature was 71 °C for microcapsules production. The  $T_g$  of the KG, hydrolyzated and microcapsules are shown in table 1.  $T_g$  values of the wall and core materials are larger than  $T_g$  of microcapsules; this may be due to possible interaction formed between components that may affect the glass transition temperature.

Table 1: Glass transition temperatures of the materials used

Materials	$T_g$ (°C)
Karaya gum	79.58
Hydrolyzate	83.41
Microcapsules	67.25

Li *et al.* [41] observed a similar behavior preparing amorphous solid dispersions (ASD) with EA as active material and carboxymethyl cellulose acetate butyrate as wall material, but when they used polyvinyl pyrrolidone (PVP) as wall material  $T_g$  was increased, concluding that PVP may be acting as a plasticizer in microcapsules formation process.

Medina-Torres *et al.* [37] used cactus mucilage as the wall material and gallic acid as an active material; they obtained  $T_g$  values of 48 and 60 °C respectively for raw materials. One of the most used carriers in microencapsulation process by spray drying is maltodextrin, which had reported  $T_g$  of 160-162 °C [42, 43]. Other coating materials used are chitin, chitosan and whey protein with  $T_g$  of 51, 59 and 162 °C respectively [43, 45].

### Analysis of FTIR

The spectrum of pomegranate peel extract obtained by methanol sis was compared with commercial EA. Spectra of standard compound showed important signals at 3500, 3200, 1750, 1600, 1550, 1300, 1200, 1050 and 800 cm<sup>-1</sup>; sharp signal at 3500 cm<sup>-1</sup> indicates the presence of hydroxyl groups while broadband at 3200 cm<sup>-1</sup> represent stretching of the hydrogen's in an aromatic ring; another

important signal is shown at 1750 cm<sup>-1</sup> which is due to the torsion of the C=O lactones carbonyl group; signals at 1600 and 1550 cm<sup>-1</sup> are caused by stretching of C=C and indicate the presence of aromatic rings; the mixed signals 1300 and 1050 cm<sup>-1</sup> are produced by the symmetric and asymmetric stretching of C-O bond, the peak at 1200 cm<sup>-1</sup> represents the stretch C-O phenolic bond and the signal at 800 cm<sup>-1</sup> shows the presence of an aromatic ring with substitution in 5 carbon atoms (Figure 3). These results agree with those reported previously by our research group [3, 44] and other authors [53]. The same characteristic signals found in the commercial compound present in the active material (signals at 3500, 3200, 1750, 1600, 1550, 1300, 1200, 1050 and 800 cm<sup>-1</sup>); signal at 3200 cm<sup>-1</sup> presents a deformation that may be caused to the accumulation of hydration water in the hydrolyzated.

FTIR analysis is effective for identifying drug-polymer interactions. Spectra of microcapsule and KG are also showed in fig. 3. FTIR spectrum from KG presented signals at 3400, 2950, 1731, 1616, 1423, 1350 and 1250 cm<sup>-1</sup>. Broad signal at 3400 cm<sup>-1</sup> corresponds to the stretching of O-H bond of the alcohol groups; the signal at 2950 corresponding to the stretching of the C-H bond of methyl groups in the structure of the gum; the band at 1731 cm<sup>-1</sup> represents the stretching of double bond between carbon and oxygen on the carbonyl group of the acetyl moiety and the presence of C=O stretching of the carboxylate group; the signal 1350 cm<sup>-1</sup> indicates the stretching of links C-O-C and bands at 1250 cm<sup>-1</sup> are produced by symmetrical and asymmetrical stretches in the C-H bonds of the methyl residue, while the bands at 1616 and 1423 cm<sup>-1</sup> are due to carboxylation of uronic acid residues in the structure of the gum [45].

Microcapsules spectral data shows a broadband at 2800-3700 cm<sup>-1</sup> indicating the presence of hydroxyl groups, besides there are bands at 1400 and 1600 cm<sup>-1</sup> similar to the KG which are due to the carboxylation of uronic acids (fig. 3). These changes indicate that the carbonyl bands of both material compounds are involved in hydrogen bonding interactions, these results were also observed by Li *et al.* [41]. When producing ASD by spray drying of EA and hydroxypropyl methylcellulose acetate succinate. The presence of characteristic peaks of EA can also be observed in microcapsules spectra such as the signals 1660 and 1550 (stretching C=C aromatic), 1200 (phenolic C-OH) and 800 cm<sup>-1</sup> (aromatic ring pentasubstituted). The modification of the location and the form of characteristic signals of both products demonstrates the presence of interactions between cover and active ingredients [46].

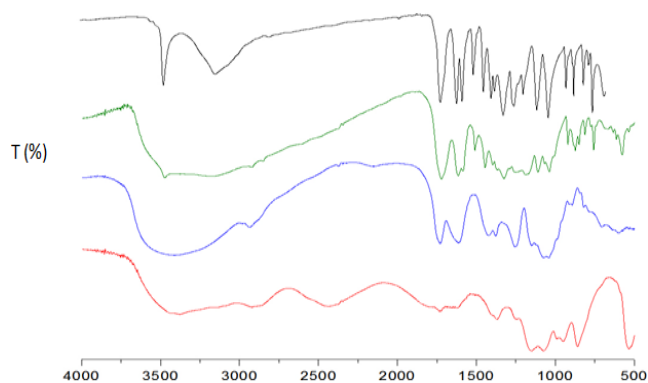


Fig. 3: Ellagic acid standard (black), hydrolyzed (green), karaya gum (blue) and microcapsule (red)

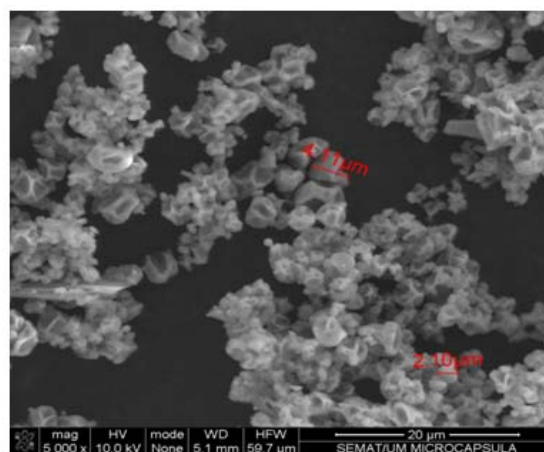
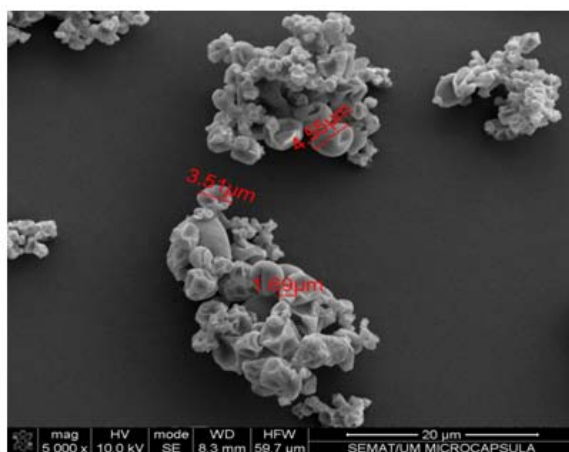


Fig. 4: Scanning electron micrographs of microcapsule

## CONCLUSION

Methanol sis is a viable technique for obtaining ellagic acid from pomegranate peel. Karaya gum is a polysaccharide with great potential for microencapsulation of active compounds, such as EA, using spray drying techniques, giving a cheaper option to reduce process cost. The obtained microcapsules by spray drying method had a range size of 1.69-4.55 mm. FTIR analysis made feasible to observe a possible interaction between the materials. This study shows the KG can be used as a wall material for microencapsulation of active compounds.

## CONFLICT OF INTERESTS

Declare None

## REFERENCES

1. Khanbabaee K, Ree Tv. Tannins: classification and definition. R Soc Chem 2001;18:641-9.
2. Ascacio-Valdés JA, Buenrostro-Figueroa JJ, Aguilera-Carbo A, Prado-Barragán A, Rodríguez-Herrera R, Aguilar CN. Ellagitannins: biosynthesis, biodegradation and biological properties. J Med Plants Res 2011;5:4696-703.
3. Robledo A, Aguilera-Carbó A, Rodríguez R, Martínez JL, Garza Y, Aguilar CN. Ellagic acid production by *Aspergillus niger* in solid state fermentation of pomegranate residues. J Ind Microbiol Biotechnol 2008;35:507-13.
4. Landete JM. Ellagitannins, ellagic acid and their derived metabolites: a review about source, metabolism, functions and health. Food Res Int 2011;44:1150-60.
5. Lei Z, Jervis J, Helm RF. Use of methanolysis for the determination of total ellagic and gallic acid contents of wood and food products. J Agric Food Chem 2001;49:1165-8.
6. Wilson TC, Hagerman AE. Quantitative determination of ellagic acid. J Agric Food Chem 1990;38:1678-83.
7. Aguilera-Carbo AF, Augur C, Aguilar CN, Prado-Barragan LA, Favela-Torres E. Extraction and analysis of ellagic acid from novel complex sources. Chem Papers 2008;62:440-4.
8. Ascacio-Valdés JA, Buenrostro JJ, Cruz RDI, Sepulveda L, Aguilera AF, Prado A, *et al.* Fungal biodegradation of pomegranate ellagitannins. J Basic Microbiol 2014;54:28-34.
9. Larrosa M, García-Conesa MT, Espín JC, Barberán FAT. Ellagitannins, Ellagic acid and vascular health. Phytochemicals and Cardiovascular Disease 2010;31:513-39.
10. Clifford MN, Scalbert A. Review. Ellagitannins-nature, occurrence and dietary burden. J Sci Food Agric 2000;80:1118-25.
11. García-Viguera C, Vicente AP. La granada. Alimento rico en polifenoles antioxidantes y bajo en calorías. Alimentación Nutrición y Salud 2004;11:113-20.
12. Häkkinen SH, Kärenlampi SO, Mykkänen HM, Heinonen I M, Törrönen AR. Ellagic acid content in berries: influence of domestic processing and storage. Eur Food Res Technol 2000;212:75-80.
13. Seeram N, Lee R, Hardy M, Heber D. Rapid large-scale purification of ellagitannins from pomegranate husk, a by-product of the commercial juice industry. Sep Purif Technol 2005;41:49-55.
14. Madrigal-Carballo S, Rodríguez G, Krueger CG, Dreher M, Reed JD. Pomegranate (*Punica granatum*) supplements: authenticity, antioxidant, and polyphenol composition. J Funct Foods 2009;1:324-9.
15. Çam M, Hisil Y. Pressurised water extraction of polyphenols from pomegranate peels. Food Chem 2010;123:878-85.
16. Ascacio-Valdés JA. Estudio de la hidrólisis microbiana de los elagitaninos. Saltillo: Universidad Autónoma de Coahuila; 2012.

17. Aguilar CN, Aguilera-Carbó A, Robledo A, Ventura J, Belmares R, Martínez D, *et al.* Production of antioxidant nutraceuticals by solid-state cultures of pomegranate (*Punica granatum*) peel and creosote bush (*Larrea tridentata*) Leaves. *Food Technol Biotechnol* 2008;46:218-22.
18. Cruz-Quiroz Rdl, Herrera RR, Esquivel JCC, Aguilar CN, Aguilera-Carbó A, Barragán LAP. La granda: fuente de potentes agentes bioactivos *Ciencia Cierta*; 2011. p. 35-8.
19. Atta-Ur-Rahman, Ngounou FN, Choudhary MI, Malik S, Makhmoor T, Nur-E-Alam, *et al.* New antioxidant and antimicrobial ellagic acid derivatives from *Pteleopsis hydodendron*. *Planta Med* 2001;67:335-9.
20. Vattem DA, Shetty K. Ellagic acid production and phenolic antioxidant activity in cranberry pomace (*Vaccinium macrocarpon*) mediated by *Lentinus edodes* using a solid-state system. *Process Biochem Int* 2003;39:367-79.
21. Hamad A-WR, Al-Momani WM, Janakat S, Oran SA. Bioavailability of ellagic acid after single dose administration using HPLC. *Pak J Nutr* 2009;8:1661-4.
22. Sepúlveda L, Ascacio A, Rodríguez-Herrera R, Aguilera-Carbó A, Aguilar CN. Ellagic acid: biological properties and biotechnological development for production processes. *Afr J Biotechnol* 2011;10:4518-23.
23. Fang Z, Bhandari B. Encapsulation of polyphenols-review. *Food Sci Technol* 2010;21:510-23.
24. Murugesan R, Osrat V. Spray drying for the production of nutraceutical ingredients-a review. *Food Bioprocess Technol* 2011;5:3-14.
25. Champagne CP, Fustier P. Microencapsulation for the improved delivery of bioactive compounds into foods. *Sci Direct* 2007;18:184-90.
26. Chang TMS. Semipermeable microcapsules. *Sci Direct* 1964;146:524-5.
27. Desai KGH, Park HJ. Recent developments in microencapsulation of food ingredients. *Drying Technol* 2005;23:1361-94.
28. Rocha GA, Fávoro-Trindade CS, Grosso CRF. Microencapsulation of lycopene by spray drying: characterization, stability, and application of microcapsules. *Food Bioprod Process* 2012;90:37-42.
29. Bakowska-Barczac AM, Kolodziejczyk PP. Blackcurrant polyphenols: their storage stability and microencapsulation. *Ind Crops Prod* 2011;34:1301-9.
30. Haidong L, Fang Y, Zhihong T, Huanwei S, Tiehui Z. Use combinations of gum arabic, maltodextrin and soybean protein to microencapsulate ginkgo leaf extracts and its inhibitory effect on skeletal muscle injury. *Carbohydr Polym* 2012;88:435-40.
31. Shcherbina Y, Roth Z, Nussinovitch A. Physical properties of gum karaya-starch-essential oil patches. *AAPS PharmSciTech* 2010;11:1276-86.
32. Lujan-Medina GA, Ventura J, Lara-Ceniceros AC, Ascacio-Valdés JA, Boone-Villa D, Aguilar CN. Gum karaya: general topics and applications. *Macromol: Indian J* 2013;9:1-7.
33. Vinod VTP, Sashidhar RB, Sivaprasad N, Sarma VUM, Satyanarayana N, Kumaresan R, *et al.* Bioremediation of mercury (II) from aqueous solution by gum karaya (*sterculia urens*): a natural hydrocolloid. *Desalination* 2011;272:270-7.
34. Singh AV. A DSC study of some biomaterials relevant to pharmaceutical industry. *J Therm Anal Calorim* 2013;112:791-3.
35. Ascacio-Valdés JA, Aguilera-Carbó A, Rodríguez-Herrera R, Aguilar-González C. Análisis de ácido elágico en algunas plantas del semidesierto. *Mexicano Revista Mexicana*; 2013. p. 36-40.
36. Poupard P, Sanoner P, Baron A, Renard CMGC, Guyot S. Characterization of procyanidin B2 oxidation products in an apple juice model solution and confirmation of their presence in apple juice by high-performance liquid chromatography coupled to electrospray ion trap mass spectrometry. *J Mass Spectrom* 2011;46:1186-97.
37. Medina-Torres L, García-Cruz EE, Calderas F, Laredo RFG, Olivares GS, Gallegos-Infante JA, *et al.* Microencapsulation by spray drying of gallic acid white nopal mucilage (*Opuntia ficus indica*). *LWT-Food Sci Technol* 2013;50:642-50.
38. Bhandari BR, Howes T. Implication of glass transition for the drying and stability of dried food. *J Food Eng* 1999;40:71-9.
39. Bhandari BR, Datta N, Howes T. Problems associated with spray drying of sugar-rich foods. *Drying Technol* 1997;15:671-84.
40. Chiou D, Langrish TAG. Development and characterisation of novel nutraceuticals with spray drying technology. *J Food Eng* 2007;82:84-91.
41. Li B, Harich K, Wegiel L, Taylor LS, Edgar KJ. Stability and solubility enhancement of ellagic acid in cellulose ester solid dispersions. *Carbohydr Polym* 2013;92:1443-50.
42. Fang Z, Bhandari B. Comparing the efficiency of protein and maltodextrin on spray drying of bayberry juice. *Food Res Int* 2012;48:478-83.
43. Roos Y, Karel M. Water and molecular weight effects on glass transitions in amorphous carbohydrates and carbohydrate solutions. *J Food Sci* 1991;56:1676-81.
44. Ascacio-Valdés J, Burboa E, Aguilera-Carbo AF, Aparicio M, Pérez-Schmidt R, Rodríguez R, *et al.* An antifungal ellagitannin isolated from *Euphorbia antisyphilitica* Zucc. *Asian Pac J Trop Biomed* 2013;3:41-6.
45. Zhang J, Du Z, Xu S, Zhang S. Synthesis and characterization of Karaya Gum/Chitosan Composite Microspheres. *Iranian Polymer J* 2009;18:307-13.
46. Krishnaiah D, Sarbatly R, Nithyanandam R. Microencapsulation of morinda citrifolia L. extract by spray drying. *Chem Eng Res Des* 2012;90:622-32.
47. Ré MI. Microencapsulation by spray drying. *Drying Technol: Int J* 1998;16:1195-236.
48. Tonon RV, Grosso CRF, Hubinger MD. Influence of emulsion composition and inlet air temperature on the microencapsulation of flaxseed oil by spray drying. *Food Res Int* 2011;44:282-9.
49. Soottitantawat A, Bigeard F, Yoshii H, Furuta T, Ohkawara M, Linko P. Influence of emulsion and powder size on the stability of encapsulated D-limonene by spray drying. *Innovative Food Sci Emerging Technol* 2005;6:107-14.
50. Lokuwan J. Characteristics of microencapsulated  $\beta$ -caroteno formed by spray drying with modified tapioca starch, native tapioca starch, and maltodextrin. *Food Hydrocolloids* 2007;21:928-35.
51. Gharsallaoui A, Roudaut G, Chambin O, Voilley A, Saurel R. Applications of spray-drying in microencapsulation of food ingredients: an overview. *Food Res Int* 2007;40:1107-21.
52. Zheng L, Ding Z, Zhang M, Sun J. Microencapsulation of bayberry polyphenols by ethyl cellulose: preparation and characterization. *J Food Eng* 2011;104:89-95.
53. Hussein M, Al Ali S, Zainal Z, Hakim MN. Microencapsulation of bayberry polyphenols by ethyl cellulose: preparation and characterization. *Int J Nanomed* 2011;6:1373-83.