Acrylic hydrogels modified with bee pollen for biomedical applications

Bożena Tylišczak¹, Dorota Walczyk*, Sławomir Wilczyński²

¹Cracow University of Technology, Department of Chemistry and Technology of Polymers, 24 Warszawska St., 31-155 Krakow, Poland.
²School of Pharmacy with the Division of Laboratory Medicine in Sosnowiec, Medical University of Silesia in Katowice, Department of Basic Biomedical Science, Kasztanowa St. 3, Sosnowiec, Poland.

INTRODUCTION

Acrylic hydrogels are polymeric matrices with a three-dimensional structure, which is characterized by an extraordinarily high swelling degree in water and other fluids. Moreover, appropriate synthesis allows them to be bio-compatible, biodegradable and bioactive - all are traits of good biomaterials (Peppas and Hilt 2006; Serra et al., 2009). Although these materials have a vast number of possible applications, we focus solely on their use as wound dressings, which are especially used in treatment of slowly healing and chronic wounds, such as decubitus ulcers, venous ulcers, diabetic foot ulcers, as well as second-degree burns and post-operative wounds (Queen et al., 2004). Introduction of an active component into the polymer matrix would allow the hydrogel to exhibit additional properties, rendering it an even more suitable material for biomedical applications. Natural products are a promising source of new discoveries in the field of pharmacy and medicine, though a majority of them has been known for centuries. An excellent example is provided by apiproducts, such as honey, propolis and bee pollen, which have been known since the ancient times because of their unique properties. Bee pollen and propolis both exhibit a vast spectrum of biological and pharmacological properties, such as immunomodulatory, anti-inflammatory and antioxidant (Burdock, 1998; Catchpole et al., 2004; Kashkooli et al., 2011; Leja et al., 2007; Sforcin 2007; Sforcin and Bankova 2011). Bee pollen contains a variety of nutrients, such as amino acids, minerals, including phosphorus, potassium, magnesium, calcium, copper, manganese, iron and zinc, some trace minerals, all B vitamins, including B₁₂, folic acid, panthothenic acid, rutin and lecithin (Almeida-Muradian et al., 2005; Human and Nicolson 2006; Kashkooli et al., 2011; Kroyer and Hegedus, 2001; Sforcin, 2007). It appears highly beneficial for human organism to profit from these bee products, and one may see a great potential of their use especially in healing and wound dressing materials, as they can provide additional anti-inflammatory activity, thus leading to shorter healing time. We have followed this lead and we have focused on modifying hydrogels with bee pollen, hoping that these would constitute very interesting materials for biomedical applications.

ABSTRACT

The focus of this paper is to present the effect of bee pollen on the structure and properties of acrylic hydrogels, often used as materials for wound dressings. Honey, propolis and bee pollen have been very well known for their extraordinary healing and medical properties for thousands of years. Ancient Egyptians, Greeks and Romans have reportedly used bee pollen as a drug, especially for a number of skin diseases and lesions. Acrylic hydrogels are, on the other hand, rather recently discovered materials used for novel wound dressings. The idea of our research was to combine a synthetic polymer matrix with bee pollen, in order to obtain innovative materials which could be used as wound dressings exhibiting new properties. The obtained acrylic hydrogels containing bee pollen were investigated towards swelling ability in distilled water and different fluids, including pseudo-extracellular fluid, simulated body fluid and Ringer’s solution. Moreover, incubation in water and simulated body fluids (SBF) and Ringer’s solution was carried out. The morphologies of bee pollen and obtained acrylic hydrogels modified with bee pollen were characterized by means of scanning electron microscopy (SEM). The chemical structure of the polymer matrix was confirmed by ATR-FT-IR spectra.
MATERIALS AND METHODS

Acrylic acid (abbrv. AA; 99.5 %wt.), potassium hydroxide (KOH, aq. solution, volumetric standard 1M) and ammonium persulfate ((NH4)2S2O8; ≥98.0 %wt.) were all purchased from Avantor Performance Materials Poland. Poly(ethylene glycol) diacrylate (abbrv. PEGDA, Mn = 700) was purchased from Sigma-Aldrich and bee pollen (BP) was obtained from Appiol-Farma.

Preparation

The acrylic hydrogels modified with bee pollen (BP) were prepared by free-radical polymerization in solution in presence of a small amount of initiator (1 % wt. (NH4)2S2O8 solution). Firstly, an appropriate amount of acrylic acid (20 ml) was mixed with 40 % wt. solution of potassium hydroxide (16.5 ml). After reaching the ambient temperature (25°C), bee pollen (2.5 % wt. or 5 % wt.), initiator (0.075 % wt.) and crosslinking agent - poly(ethylene glycol) diacrylate (PEGDA, Mn = 575, 4.5 % wt.), were added.

The reacting mixture was mixed intensively on a magnetic stirrer at 60°C for 30 min. Afterwards the mixture was poured into Petri dishes, which were left at 37°C for 24h. The compositions of formed acrylic hydrogels modified by appropriate amounts of bee pollen 0, 2.5 and 5 %wt. respectively are shown in Table 1 presented below.

Table 1: The compositions of acrylic hydrogels modified by bee pollen.

<table>
<thead>
<tr>
<th>Samples</th>
<th>Acrylic acid (ml)</th>
<th>Bee pollen (wt %)</th>
<th>PEGDA (wt %)</th>
<th>APS - initiator (wt %)</th>
</tr>
</thead>
<tbody>
<tr>
<td>PAA</td>
<td>20</td>
<td>0</td>
<td>4.5</td>
<td>0.075</td>
</tr>
<tr>
<td>PAA</td>
<td>20</td>
<td>2.5</td>
<td>4.5</td>
<td>0.075</td>
</tr>
<tr>
<td>PAA</td>
<td>20</td>
<td>5</td>
<td>4.5</td>
<td>0.05</td>
</tr>
</tbody>
</table>

Swelling behaviour

The dried samples of obtained acrylic hydrogels, each weighing 0.5 g, were immersed in the following fluids: water, 0.9 % wt. NaCl solution, 0.9 % wt. CaCl2 solution, Ringer’s solution (pH = 5.75; Na+ = 147.2 mmol-dm-3, Cl− = 155.7 mmol-dm-3, K+ = 4.0 mmol-dm-3, Ca2+ = 2.2 mmol-dm-3) and simulated body fluid (SBF, pH = 7.40) and in pseudo-extracellular fluid (PECF) (pH = 7.4±0.2), simulating the wound fluids. The comparison of concentrations of ions such as: Na+, K+, Ca2+, Mg2+, HCO3-, Cl−, HPO42- and SO42- that are present in accordingly: human plasma, SBF and PECF are summarized below in Table 2 (Kokubo and Takada 2006, Vallet-Regi and Arcos 2008).

The swelling capacity (Q) was calculated according to the following formula:

\[ Q = \frac{W_f - W_i}{W_i} \]

where: \( W_i \) and \( W_f \) refer to the weights [g] of the hydrogel in swollen and dried states, respectively.

Table 2: Human plasma and ions concentration [mM] of SBF and PECF (Kokubo and Takada, 2006; Vallet-Regi and Arcos, 2008).

<table>
<thead>
<tr>
<th>Ions →</th>
<th>Na⁺</th>
<th>K⁺</th>
<th>Ca²⁺</th>
<th>Mg²⁺</th>
<th>HCO₃⁻</th>
<th>Cl⁻</th>
<th>HPO₄²⁻</th>
<th>SO₄²⁻</th>
</tr>
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<tbody>
<tr>
<td>SBF</td>
<td>142</td>
<td>5</td>
<td>2.5</td>
<td>1.5</td>
<td>27</td>
<td>103</td>
<td>1</td>
<td>0.5</td>
</tr>
<tr>
<td>PECF</td>
<td>145</td>
<td>5</td>
<td>2.5</td>
<td>1.5</td>
<td>4.2</td>
<td>148</td>
<td>1</td>
<td>0.5</td>
</tr>
</tbody>
</table>

Incubation studies

In the incubation studies 0.5 g of prepared acrylic hydrogels were placed in 50 ml of an appropriate medium: water, SBF and Ringer’s solution for a period of 32 days. During this time pH values were measured periodically.

Attenuated total reflectance FT-IR spectroscopy analysis

All Attenuated Total Reflectance Fourier-Transform Infra-Red (ATR-FT-IR) spectra were recorded using Spectrum 65 (Perkin Elmer) spectrometer. The ATR probe was equipped with a diamond crystal. All spectra consisted of 32 scans at 4.0 cm⁻¹ resolution and were recorded at 25°C.

Scanning Electron Microscopy

In order to investigate the structure of the obtained hydrogels modified with bee pollen, as well as bee pollen alone, images using scanning electron microscopy were taken and analyzed (JEOL JSM 7500F with a Back Scattered Electrons detector).

RESULTS AND DISCUSSION

SEM analysis

In order to assess whether and how the hydrogel and bee pollen interact, SEM images of both bee pollen and the hydrogel modified with bee pollen were taken. Fig. 1 shows bee pollen in its raw form, as purchased. As it can be seen, bee pollen grains occur mostly in elliptic forms with two or three grooves. Moreover, these particles have a very uniform size distribution. This is due to the fact that the pollen grains were all collected from the same plant species.

Enlargement of a single particulate of bee pollen clearly shows a porous surface of the grain, with a very uniform distribution of the pores. Since the surface is not smooth but exhibits certain porosity, it is highly likely that adhesion to such a surface ought to be feasible and allow good bonding between the acrylic hydrogel matrix and the natural substance (see Fig. 1).

Images shown in Fig. 2, were collected using SEM imaging and show hydrogel modified with bee pollen. The image to the left shows fragments of the gel in reasonably small magnification, while the image in the centre shows clearly how the bee pollen particles are bound to the hydrogel. As expected, it appears that the gel at least partially coats the bee pollen particles.
Fig. 1: SEM images of bee pollen where A denotes raw bee pollen magnified 250x, B shows single particulate of raw bee pollen magnified 2500x and C shows enlargement of image A magnified 1000x.

Swelling behavior

Hydrogels are very well known for their excellent moisture sorption abilities which allow them to be used in a variety of biomedical and personal hygiene applications. Addition of any substance, such as bee pollen, may affect these properties therefore sorption capacity of modified hydrogel was tested using a variety of solutions, as it is shown in Fig.

The results show that the polymer matrix and modified pollen hydrogels exhibit the lowest sorption capacity in CaCl₂ solution. This is a consequence of presence of Ca²⁺, a multivalent cation which contributes to further cross-linking of the gel, leaving less void spaces in the macromolecule three-dimensional structure available for the solution to penetrate. The highest sorption values are achieved upon sorption of water, as expected. The water molecules can easily penetrate to the interior of the hydrogel and since they are not charged and are inert, they do not meet any hindrances.

It is impossible to define an overall relationship between the amount of added bee pollen and general sorption capacity. In case of swelling in SBF, addition of bee pollen caused a dramatic decrease, while other sorption capacities changed only insignificantly and could be both positive (PECF, Ringer’s solution, water) and negative (0.9%wt. NaCl). It is possible that the worsened sorption capacities are caused by the fact that there is less volume available to penetrate in the hydrogel matrix, as it is partially occupied by the bee pollen particles. Nevertheless, since sorption properties are not severely affected, there is a good possibility that hydrogel modified with bee pollen could be used in the same applications as the non-modified material.

Fig. 2: SEM micrographs of acrylic hydrogels modified with bee pollen where A denotes acrylic hydrogel modified with bee product magnified 150x, B and C show single particulate of modified hydrogel magnified 1000x.

Fig. 3: Sorption capacity of acrylic hydrogel modified with bee pollen.

Incubation studies

In order to assess stability of the obtained hydrogels modified with bee pollen, pH measurements were carried out in the course of 32 days of incubation in the fluids mentioned in the Experimental section above. The results of these measurements performed on hydrogel modified with 2.5 %wt. and 5% wt. of bee pollen are shown in Fig. 4 a and b, respectively.

The conducted pH measurements during the incubation period show significant changes in pH values only for the first four days, after which, regardless of the type of solution used, a plateau is reached. In case of Ringer's fluid and SBF, this initial change which took place over the first four days is far more significant than for samples immersed in de-ionized water. This is certainly associated with presence of metal cations in both SBF and Ringer’s solution and their absence in de-ionized water. As the results show, amount of introduced bee pollen has no significant effect on the pH changes during the incubation. The results for both polymer matrices containing 2.5% wt. and 5% wt. of bee pollen are very similar. Lack of changes over an extended period of time appears to confirm that the hydrogel matrix would not degrade in conditions typical for biomedical applications.
Attenuated total reflectance FT-IR spectroscopy analysis

The structure of the hydrogel matrix and presence of functional binding groups such as ester and ether groups, as well as carbonyl, hydroxyl and carboxyl groups have been confirmed by means of Attenuated Total Reflectance Fourier-Transform Infra-Red (ATR-FT-IR) spectroscopy. The resulting spectra are shown in Fig. 5 above. In all spectra a wide signal is observed in the range of 3700-3100 cm\(^{-1}\), and it is resulting from stretching vibrations of hydroxyl groups and hydrogen interactions (formation of hydrogen bonds with hydroxyl groups, most likely). Presence of -CH\(_2\)- groups is confirmed by the signal found at 2930-2900 cm\(^{-1}\), resulting from valence vibrations of these groups. The band from the range of 1553-1556 cm\(^{-1}\) corresponds to the vibrations of the carbonyl group of poly (acrylic acid).

A signal caused by presence of ester groups of poly(ethylene glycol) diacrylate is observed at around 1704 -1717 cm\(^{-1}\). Signals corresponding to symmetrical scissor mode vibrations of the -CH\(_2\)- groups are observed at a wavelength of ca. 1450 cm\(^{-1}\). The structure of PEGDA is confirmed by peaks corresponding to C-O-C stretching, which are visible at 1165 and 1014 cm\(^{-1}\). Furthermore, in case of acrylic hydrogels modified by 5 %wt. of bee pollen several subtle changes in the overall shape of the spectrum in the range of 1405-800 cm\(^{-1}\) are observed.

For example, C-O-C stretching vibrations from PEGDA were observed at different wavelengths, i.e. 1190 and 1045 cm\(^{-1}\). In addition, an extra peak at 985 cm\(^{-1}\) is observed and can be assigned to the C-OH out-of-plane bending and -CH\(_2\)- twisting modes (Silverstein et al., 2014).

Fig. 4: Results of incubation studies, expressed as pH variation, of hydrogel matrices modified with 2.5 %wt. bee pollen (a) and 5 %wt. bee pollen (b).

Fig. 5: The ATR-FT-IR spectra of acrylic hydrogels containing a) 0 %wt., b) 2.5 %wt. and c) 5 % wt. bee pollen.
CONCLUSIONS

The focus of this research was to obtain hydrogel acrylic matrices modified with bee pollen and to pre-evaluate their potential for biomedical applications. Synthesis proposed by the authors is straightforward and allows to obtain hydrogel matrices in which bee pollen particles are well distributed and bound to the acrylic material. The resulting materials exhibit good sorption capacity against different liquids, typically used in biomedical testing, such as Ringer’s solution, SBF, PECF. Series of incubation experiments proved that these materials are stable in body-simulated conditions, as well as in conditions mimicking wound environment, over an extended period of time. Bearing in mind that acrylic hydrogels are already in use as wound dressing materials, further tests should focus on the benefits of addition of bee pollen into hydrogel matrices. It is expected that such modified dressings would have anti-inflammatory and antiseptic properties, and would also enhance the healing time of the wound.

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