Researches Concerning Chemical Modifications of Hair Keratin

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The work presents chemical procedures used to obtain a film or a hydrogel after hydrolyze treatment of hair at the temperature of 150°C and the pressure of 4.9 atm. In order to obtain the hydrogel, this treatment is followed by a stage of oxidation with hydrogen peroxide, and a stage of reduction with Na2S2O4 or, in another variant, a stage of oxidation with hydrogen peroxide and a treatment with NaOH in alcoholic medium, to form natrium cationic groups. The protein chains with sulphonic groups and natrium cations determine opposite electrostatic interactions, with the formation of a polymer loop that incorporate water. The evaluation of formed hydrogel was carried out by determining the swelling degree. We have used optical and SEM photomicrographs to determine morphological organization, and EDAX micropictures for chemical modifications. A cellulose film realized by growing Gluconoacetobacter xylinus bacteria was used as control sample.

Keywords: beta-keratose, gamma-keratose, hydrogel, film

Loosing the skin function due to a lesion or a disease determines a severe unbalance and a physiological incapacity that can be lethal. Skin substitutes manufactured in tissues have their limits related to clinical utilization. In this field of specialization, there is no therapeutic tissue substitute able to take over skin function through a sole procedure, and with an adequate thickness for a corresponding coating of thick wound [1].

In order to obtain material support, in medical applications one uses natural polymers, such as keratin which, by their biological and physico-chemical properties, transfer these properties to the designed materials [2-5]. Regardless its origin, keratin presents scientifically substantiated morphological details that needed over time special cognition efforts. The evolution of the understanding of structural details and the role played in fiber biology has as aiming points some reference names such as: Elod E., Speakman father and son, Zahn H., but there are also structural details for which the scientific research continues. The utilization of keratin hydrogel offers the biomaterial the possibility of a controlled drug release, as well as its utilization as a support for building the artificial skin [7, 8].

In the case of burnt wounds with a wide surface of the affected skin, there is a complex of factors that strongly affect the general patient condition. Namely, the severe burns determine destructions of skin and of other deep structures, with homeostasis (organism natural self-regulation) alteration, by maintaining the biochemical parameters within constant limits). Depending on the duration of heat aggression, it can result in enzymatic process alteration, protein necrosis, tissue caramelization and calcination.

The above mentioned phenomena are accompanied by diminution or loss of body capacity to maintain the heat and to preserve biologic fluids, with the severe alteration of circulatory, hepatic and renal functions, against a generalized sepsis. The wound evolution determines frequently an additional loss of derma tissue [9]. In literature, there are communications regarding cell proliferation in the burnt wounds. These communications refer to collagen or keratin biomaterials [10], factors of cell regeneration intensification [11,12], or cell engineering procedures [1].

The connection between keratin and skin regeneration is related to a tradition from the history of civilization, according to which, in order to cure the burnt wound, one needs to burn some of patient hair; the resulting solid mass in crushed and transformed in a powder, remedy for patient wound. In literature there are approaches related to the production of keratin-based biomaterials [13]. Besides the surgical and pharmaco-dynamic therapy, for the amelioration of patient general condition, it is imperiously necessary to solve the problem of skin deficit for wounds’ coating self-grafting. It has been mentioned that their limitation can be compensated by producing teguments through tissue engineering techniques.

The work presents the possibility of chemical keratin processing from patient human hair, to obtain a biomaterial under the form of hydrogel or film, destined to tissue growth with prospective to be used as skin substitute, in order to avoid immunity reactions.

Considerations on human keratin utilization as biomaterial

Human hair is a keratin fiber developed from skin follicle, having the role of protection agent, sensor and sexual attraction, being a characteristic common to both man and mammals. The human hair structure is chemically and morphologically similar to the sheep wool. The difference consists in the fact that for wool, the ortho-para structure is differentiated, with cell disposition under the form of two bands twisted along the fiber [14,15]. Keratin

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products are biocompatible mediums that facilitate the cell growth [5].

**Experimental part**

One has collected undyed hair, cut at lengths of 2-5 cm, from men regardless the age, delivered by a hair dresser; then the collected hair was cleaned for 24 h, in a Soxhlet by removing the lipids, using a mixture of CHCl₃ and CH₃OH in ratio of 2:1 (v/v). The hair is dried by night at 40°C. The cleaned hair and an amount of distilled water are introduced in a sealed stainless steel capsule, at a solid/liquid ratio of 2:35 (w/w). The capsule is introduced in an air-drier at 150°C for 2 h, as illustrated in figure 1.

**Hydrogel 1**

In order to obtain the hydrogel from beta keratose film according to procedure 1, after cooling the solution, the supernatant is extracted through centrifugation at 10,000 rpm for 10 min. The obtained solution is cooled and additionally oxidized with 4 mL H₂O₂ for 10 min, repeating the procedure 6 times. Then the solution remains for 72 h at room temperature. The solution gets slowly thicker, forming a cross-linked gel.

**Hydrogel 2**

Beta keratose is subjected to an oxidation treatment with boiling hydrogen peroxide for 3 h, using 4 g solid phase and H₂O₂ at a ratio of 1:4 (w/v). Then it is washed with 96% ethanol and squeezed by pressing with filter paper, treated with 18 mL of 96% ethanol with a content of NaOH reported to keratose quantity of 0.14: 2 (w/w) to obtain a pH= 8.5-9. It is treated at 98°C for 3 h and then stirred cooled for 5 h. It is then detached from filter paper and washed six times with 100 mL of distilled water. One obtains a hydrogel consisting of swollen hair filaments.

**Hydrogel characteristics**

Swelling degree

The swelling degree is evaluated in percentages, based on gravimetric tests:

\[
SD (\%) = \left( \frac{M_f - M_{in}}{M_{in}} \right) \times 100
\]

where: \( M_f \) - weight of material after swelling, \( M_{in} \) - weight of the initial (dry) material.

In the case of performed determinations, the initial mass of beta-keratose subjected to treatment was 2 grams.

Table 1 presents the swelling degree values obtained through gravimetric measurements in the two cases of hydrogel.

Elemental composition determinations

Chemical composition of the elements was determined in percentages through EDAX technique. The obtained data are summarized in table 2.

Morphological aspects

For morphological visualizations, one has prepared the following biomaterials:

- Hydrogel 1 is illustrated in figure 2, witness sample consisting of bacterial cellulose film is presented in figure 3, biomaterial consisting of cellulose film and beta-keratose is illustrated in figure 4, while figure 5 presents the biomaterial consisting of bacterial cellulose and gamma-keratose film, and figure 6 represents a micrograph of hydrogel 2 after drying.

<table>
<thead>
<tr>
<th>No.</th>
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<th>C</th>
<th>N</th>
<th>O</th>
<th>S</th>
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<td>-</td>
<td>51.12</td>
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<tr>
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<td>2.39</td>
<td>47.65</td>
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<tr>
<td>3</td>
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<td>2.98</td>
<td>41.75</td>
<td>4.52</td>
</tr>
<tr>
<td>4</td>
<td>Hydrogel</td>
<td>64.93</td>
<td>2.18</td>
<td>25.70</td>
<td>7.19</td>
</tr>
</tbody>
</table>
Fig. 2 - Optic microscopy image x100 of hydrogel 1 from beta-keratose in wet condition

Fig. 3. SEMx1000 micrograph of witness sample made of bacterial cellulose

Fig. 4. SEMx1000 micrograph of biomaterial consisting of cellulose film and beta-keratose

Fig. 5. SEMx1000 micrograph for the film of bacterial cellulose and gamma-keratose
Results and discussions

Taking into account that the skin represents a human organ with a big surface [17], its regeneration implies a series of peculiarities. According to the opinion of the authors of this study and to other bibliographic references [2, 5], keratins preparation with the formation of a biomaterial constitutes a basic sustaining element for artificial tegument realization.

The treatment of hair in the metallic capsule at 150°C and pressure of 4.9 atm. in the presence of water determines a hydrolytic process of protein material splitting in two parts: a solid part, considered to be beta-keratose, and a water-soluble phase, namely gamma-keratose [5,18,19].

In the case of the studied hydrogel, one has obtained different swelling degrees, as the result of procedures differences. Under the action of temperature and pressure as common treatment, both procedure participate in hair hydrolytic degradation in the two types of keratose, beta and gamma. This treatment is followed by an oxidative treatment with hydrogen peroxide, which facilitates the splitting of cystine disulphide bonds in sulphonic groups, as illustrated in figure 7.

According to the above presented images, one can obtain a hydrogel that follows the organization of the hair in quasi-fibrillary state (images from figs 2 and 6) or of more homogeneous formations (figs.3-5). Depending on keratin treatment and processing procedure, one can obtain more compact (beta-keratin and cellulose) or more porous formations.

According to EDAX data presented in Table 2, where the witness sample is a cellulose structure of bacterial origin, the presence of sulphur in the structure can be exclusively due to Acetobacter xylinum bacteria. For the other representatives illustrated in table 2, sulphur comes from hair cystine structure, while the nitrogen certifies the protein structure as the result of the presence of constitutive amino acids.

Different oxidation degrees indicate differences at the level of the implied chemical modifications and of keratin sources, from beta-keratin and gamma-keratin respectively.

Conclusions

The keratin structure of the human hair is susceptible of chemical transformations that permit to obtain a film or hydrogel precursors for the generation of a biomaterial as tegument substitute.

The obtained hydrogels, as well as the cellulose film are precursors for biomaterials meant for tegument regeneration.

References

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