ARCHIVES OF ENVIRONMENTAL PROTECTION

no. 1

vol. 40

pp. 3-12

2014 PL ISSN 2083-4772

VERSITA

PL IS

DOI: 10.2478/aep-2014-0003

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ECO-FRIENDLY SOAKING PROCESS USING TANNIC ACID AS AN ALTERNATIVE BACTERICIDE

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Keywords: Bactericide, leather industry, soaking, tannic acid.

Abstract: Eco-friendly leather processes based on the usage of natural products have become a potentially attractive issue for leather industry during the last few decades. Synthetic protective chemicals like bactericides used in most soaking process are known as hazardous substances and cause tannery effluents with high concentrations of Chemical Oxygen Demand (COD). In the present study, the effect of tannic acid on microorganisms, skin, wool and effluent were investigated in order to demonstrate the applicability of tannic acid in soaking process instead of commonly used bactericides. The bacterial load (cfu/ml), COD and Nitrogen Content (N) of the soaking effluents and Total Kjeldahl Nitrogen (TKN) content of skins and wools were investigated. Application of 0.5 and 1 wt% tannic acid concentrations was more effective than commercial bactericide, while comparable results were achieved by 0.1 and 0.3 wt% tannic acid. The application of tannic acid has the potential to be an alternative, eco-friendly bactericide for leather industry by reducing the pollution of leather soaking process.

INTRODUCTION

New eco-benign systems, either using non-hazardous compounds or developing new production methods have become the raising trend in global industrial activities, due to the increasing health and environmental regulations and restrictions.

Researchers have adopted various approaches to minimize the environmental impact of leather processes taking into consideration the use of chemicals having less toxicity or less environmental impact [5, 27], recovery and/or reuse of water, floats and chemicals in leather processing steps [25, 36, 43] and innovation of processes and products [33, 476].

Soaking is one of manufacturing processes of leather used to restore the original water content of skin. In this process synthetic antimicrobial agents are widely used to prevent microbial growth. However, considering the potential toxicity of these chemicals and requirements imposed by regulating agencies, natural eco-friendly alternative

antimicrobial products have recently become a predominant focus of interest in many fields like textile, leather as well as pharmacy, medicine, cosmetics, food and etc.

In leather industry, the use of several alternative environmentally friendly antimicrobial products has been reported in different processing steps. The antimicrobial effect of *Aloe vera* in fatliquoring process [6], the biocide effect of photo sensitizers [16], bronopol [38] and ozone [39] in soaking process were investigated. The use of vegetable tannins such as quebracho, mimosa, gall-nut, chestnut and valonia as a biocide in leather soaking process was also studied [18] owing to their antimicrobial and antiseptic properties [19]. Tannic acid, which is a commercially important hydrolysable gallotannin, has also got a great deal of attention as an antimicrobial agent due to its antimicrobial, antimutagenic, anticarcinogenic and antioxidant properties [1, 3, 4, 10, 22, 23, 31, 35, 37, 41, 44]. Tannic acid is present in a variety of fruits and vegetables including tea, cocoa, beans, grapes, strawberry, persimmon, especially the bark of the oak species, sumac and myrobalan [1, 11, 40] and also categorized as a 'generally recognized as safe' (GRAS) food additive [1]. In addition, it is not classified as hazardous material according to CLP Regulation (EC) 1272/2008 on classification, labeling and packaging of substances and mixtures [32]. Tannic acid was applied in textile [8], microbiology [45], medical and biomaterial sciences [2, 26] and leather industry [17].

Recent studies on tannic acid in leather industry can be summarized as an application of tannic acid in pickling and tanning processes [17, 20, 21, 46, 52], the reduction of Cr (VI) formation in leather by tannic acid [14] and use of tannic acid in leather dyeing process as a natural dye [29]. According to our knowledge, so far, there has been no reported research on the application of tannic acid as a bactericide in soaking process for leather industry.

In this research, the applicability of tannic acid as an alternative natural antimicrobial agent in respect to common synthetic bactericides used in soaking process were investigated to reduce the negative environmental impact of these protective chemicals.

MATERIALS AND METHODS

Materials

Tannic acid (Merck, Darmstadt, Germany), which is the powder form of tannin with a pH value of 3.5 (100 g/l, H_2O , 20°C), was used as an eco-friendly bactericide in soaking process. The bactericide effect of tannic acid was tested using 100 cm² pieces of dry salted raw Metis type sheepskins. To evaluate the antibacterial effect of tannic acid, a commercial bactericide containing benzisothiazolinon (25% conc.) as an active ingredient was used.

Methods

Leather samples were pre-soaked for 2 hours under static conditions and washed for 5 minutes to remove the dirt and remained salt. Then, the samples were put in 2L beakers and soaked with 1L solutions of tannic acid of different concentrations (0.1, 0.3, 0.5 1.0%) as well as with 0.1% solution of commercial bactericide at 27°C. During the soaking process, a shaker provided mechanical mixing for 10 minutes per hour. Also, pH and temperature values of soaking baths were controlled. At 8 and 24 hours of process, samples were taken for the determination of soaking bath properties. In order

to demonstrate the applicability of tannic acid in soaking process, not only the effect of tannic acid as bactericide, but also its effect on soaking effluent and the compositions of skins and wools were determined. A blank bath without any protective chemicals was also analyzed. The results were given as mean values (\pm SE) obtained with triplicate samples and evaluated statistically.

The enumeration of cultivable bacteria by plate count method (CFU)

The bacterial load of the soaking baths was determined with the samples taken at 8th and 24th hours of process by use of pour plate method. 1ml of each soaking sample was added to 9 ml sterile physiological saline solution for preparing serial dilutions of samples. This procedure was followed by mixing 1 ml aliquots of the appropriate sample dilutions with Plate Count Agar (pH 7; 45°C; Merck) in a sterile petri dish. After plating, the media were incubated at 37°C for 24 hours and the cultivable bacterial cells in 1 ml of a soaking bath were counted. For each experiment, three separate sub-samples from soaking baths were analyzed and the enumeration of bacteria was expressed in log cfuml⁻¹.

The determination of N contents and COD values of soaking effluents

The spent liquor from the soaking operation was analyzed for total N content and COD values ($mg \cdot L^{-1}$) using standard kits (114763: Nitrogen (total) Cell Test; 114555: COD Cell Test; Merck) with Merck SQ 300 Water and Wastewater Spectrophotometer at 520 and 585 nm respectively. N content and COD load of each soaking effluent was analyzed in triplicate.

The determination of shrinkage temperature and TKN

Shrinkage temperature of leather samples was determined in accordance with TS 4120 EN ISO 3380 [49]. In this method, soaked 5×0.2 cm leather specimen was immersed into distilled water which was used as a heating medium for the shrinkage temperatures at or below 98°C and the heater was set to make the temperature of the heating medium increase 2°C per minute. When the first shrinkage was observed, the changes were recorded in every 30 seconds until the last shrinkage.

TKN values of skins and wools were determined in accordance with the Kjeldahl nitrogen method [7, 50] and experiments were performed in triplicate.

Scanning Electron Microscopic (SEM) Examination

The fiber structure of wool of each soaked leathers after soaking process was monitored with TM 1000 Hitachi (Japan) Tabletop Scanning Electron Microscopy with the magnification power 20–10.000 according to methods previously described in the literature [9]. The wool samples were dried at 40°C after soaking process and randomly selected ten separate wool samples from each application were viewed under the SEM without requiring any sample preparation.

Statistical analysis

The data were analyzed with SPSS 15.0 for Windows. Analysis of variance was performed on each attribute and data were analyzed for treatment effects, time effects and treatment by time interactions. When significant interactions were observed, Tukey *post hoc* test was used for multiple comparisons for enumeration of the cultivable bacteria

assessment. Multiple comparisons of chemical oxygen demand, total nitrogen content and total kjeldahl nitrogen results were performed by Duncan test. The evaluation of the linear relationship between tannic acid offer and the CFU, COD, N and TKN results was measured using Pearson's correlation.

RESULTS

The Effect of Tannic Acid on Microbial Load of Soaking Process

The bacterial load of soaking baths with different concentrations of tannic acid and control samples are presented in Table 1.

Table 1. The number of cultivable bacteria expressed as log_{10} cfu per 1 ml after 8th and 24th hours of soaking process

Treatment							
Time (h)	Blank	0.1% TA	0.3% TA	0.5% TA	1% TA	Bactericide	
8	$5.93{\pm}0.015^{Ba}$	$4.80{\pm}0.012^{\rm Bbc}$	$4.53{\pm}0.004^{\rm Bcd}$	$4.22{\pm}0.003^{\rm Bed}$	$3.51{\pm}0.004^{\text{Be}}$	$5.06{\pm}0.017^{Bab}$	
24	$8.03{\pm}0.028^{Aa}$	$7.66{\pm}0.008^{\rm Ab}$	$7.51{\pm}0.003^{\rm Ab}$	$7.28{\pm}0.004^{\rm Ac}$	$6.49{\pm}0.005^{\rm Ac}$	$7.43 {\pm} 0.006^{\rm Ab}$	

* values in the same columns or rows followed by the same upper or lower case letters, respectively are not significantly different according to Duncan test (p < 0.01).

TA, tannic acid; n = 3

The bacterial growth at 8 hours of soaking indicated that all the concentrations of tannic acid wt% could prevent the microbial growth under ideal conditions. The microbial growth prevention of 1wt% tannic acid was significantly (p < 0.01) higher than the other treatments of tannic acid and control samples. On the other hand, there was no significant difference between control samples and 0.1, 0.3, 0.5 wt% concentrations of tannic acid (p > 0.01).

The number of cultivable bacteria in bactericide added soaking process for 24 hours was found significantly lower than the blank sample (p < 0.01). Comparable results with commercial bactericide were obtained by using 0.1 and 0.3 wt% of tannic acid (p > 0.01). For the same period of time, although 0.1 wt% of tannic acid gave the highest colony forming unit results, it has been a limitation at microbial growth as much as half of the blank sample (p < 0.01). It was obvious that the results of 0.5 and 1 wt% of tannic acid were significantly better than the commercial bactericide (p < 0.01). An analysis using Pearson's correlation coefficient also indicated that there was a significant relationship between tannic acid offer and number of bacteria (cfu) at 8 and 24 hours of soaking process, r(12) = -1, p < 0.000 and, r(12) = -0.990, p < 0.000 respectively.

Nitrogen and COD Values of Soaking Effluents

The nitrogen value of blank sample for 24 h, presented in Table 2, had the highest value among other effluent samples (p < 0.05). This might be because of the heavy infection of the soaking float. Owing to inhibition effect of tannic acid and bactericide on microbial growth, lower nitrogen values were acquired in these effluents. The lowest N content values were obtained from 0.5 and 1 wt% tannic acid application compared to other

soaking groups (p < 0.05). However, there were no statistical differences between the groups of 0.1, 0.3 wt % tannic acid and bactericide (p > 0.05). In the mean time, the difference between the 0.5 and 1 wt % tannic acid was not statistically significant (p > 0.05). Generally, all effluents contain N values in the range between 3 and 10 mg per L [15], that are the limit values specified for directly discharge into the surface water.

Table 2. Total nitrogen (N) content and chemical oxygen demand (COD) values of soaking effluents ($mg \cdot L^{-1}$) for 24 h soaking process

Treatment						
	Blank	0.1% TA	0.3% TA	0.5% TA	1% TA	Bactericide
Ν	4.13±0.38ª	3.23±0.09 ^{a,b}	3.17±0.41 ^{a,b}	2.27±0.45 ^b	$2.43{\pm}0.07^{b}$	3.17±0.20 ^{a,b}
COD	1673±40.55°	1683±33.33°	1667±6.67°	1673±21.86°	2043±61.19b	3040±34.64ª

* values in the same row followed by the same letters are not significantly different according to Duncan test (p < 0.05);

TA, tannic acid; n = 3

The COD values of soaking effluents were determined lower than the previously reported values [42, 48]. With the use of 1 wt% of tannic acid instead of bactericide, the COD values of soaking effluents were reduced approximately by 33% (Table 2).

The difference between the 1 wt% of tannic acid and bactericide was also found significant (p < 0.05). However, no statistically significant difference was found between the soaking applications of blank, 0.1, 0.3 and 0.5 wt% of tannic acid (p > 0.05). The highest COD value reduction by 45% was obtained from 0.1, 0.3 and 0.5 wt% tannic acid applications. An analysis using Pearson's correlation coefficient also indicated that there was a significant relationship between tannic acid offer and COD values of effluents, r (12) = 0.853, p < 0.000 respectively. In soaking process, the 0.5 wt% tannic acid concentration that has a comparable antimicrobial effect with 1 wt%, may be selected considering both environmental aspects and cost-effectiveness, although the COD value of 1 wt% tannic acid is lower than bactericide.

The Effect of Tannic Acid on Shrinkage Temperature, TKN and SEM

The shrinkage temperature of the skins soaked with increasing concentrations of tannic acid presented only slight differences within the minimum and maximum range of 55 and 57°C. This can be regarded as a supportive indication for not generating any modification in skin structure. Also, the shrinkage temperatures of blank and bactericide samples were determined as 53 and 55°C respectively.

All skin and wool samples provided similar total Kjeldahl nitrogen values that varied between 14.56–15.27% (Table 3). No statistically significant difference was found between TKN values of all wool and skin samples (p > 0.05). Scanning electron microscopy analysis of wool for tannic acid applications as well as control samples did not reveal any bacterial attack on fiber structure of wool (Figure 1). The statistically insignificant TKN values of wool were also supported by similar SEM images. Accordingly, bacterial attack in the course of tannic acid application does not cause any adverse structural modifications in skin matrix and wool as supported with TKN values and SEM images.

	Treatment						
	Blank	0.1% TA	0.3% TA	0.5% TA	1% TA	Bactericide	
Skin samples							
TKN	14.57±0.090ª	14.60±0.063ª	14.56±0.127ª	14.87±0.055ª	14.68±0.072ª	14.58±0.012ª	
Wool samples							
TKN	15.22±0.165ª	15.27±0.012ª	15.21±0.169ª	15.06±0.063 ª	15.24±0.048 ª	14.97±0.050 ª	

Table 3. Total nitrogen (TKN) values of skin and wool samples (%) for 24 h soaking process

* values in the same row followed by the same letters are not significantly different according to Duncan test (p < 0.05)

TA, tannic acid; n = 3

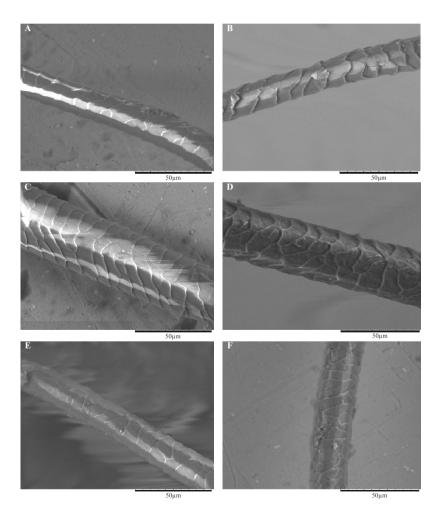


Fig 1. SEM micrographs of wool samples soaked with 0.1% (A), 0.3% (B), 0.5% (C) and 1% (D) of tannic acid, without any bactericide (blank) (E) and with a commercial bactericide 0.1% (F) × 1500

Unauthenticated | 77.254.68.37 Download Date | 6/20/14 1:04 PM The most commonly known synthetic preserving agents used in leather industry have carbamates, benzisothiazolinon, organo-bromine compounds, and oxidative agents as active substances. Their use in leather industry has become the focus because of the chemical safety and consumer health issues considering their toxic effect for wastewater, aquatic organisms and environment. The toxicity of bactericides and fungicides in tannery effluents was confirmed by determination of their extremely high acute and chronic toxic effects on bacteria, algae, zebra fish and daphnids such as *Daphnia magna* and *Danio rerio* [48]. They may impair fertility, cause harm to the unborn child and potential heritable genetic damage [30].

A number of articles have been reported regarding the inhibition effect of tannins on microorganisms. Their antibacterial effect could be largely attributed to their reduction of enzyme activity, dysfunction of cell membrane and deprivation of substrate metal ions and minerals [1, 24, 51]. Amongst tannins, tannic acid was found to be inhibitory to the growth of intestine bacteria such as *Bacteroides fragilis, Escherichia coli* and *Clostridium perfringens*. This inhibitory effect was thought to be closely related to the iron-binding capacity of tannic acid [11]. Besides, Chung *et al.* [12] reported that tannic acid could not inhibit any of them. Chung *et al.* [12] and [13] reported that the ester linkage between gallic acid and glucose (to form tannic acid) was important for the antimicrobial potential of these compounds.

Tannic acid significantly inhibited the microbial growth without altering collagen matrix as well as wool structure. All concentrations of tannic acid used in the study, provided an antibacterial effect in comparison to blank samples by decreasing the bacterial load of soaking baths significantly. Tannic acid performed better antimicrobial activity during the first 8 hours of soaking process when compared to commercial bactericide. Commercial bactericide presented similar antibacterial effect during 24 h soaking as 0.1 and 0.3 wt% tannic acid. Tannic acid at the concentrations of 0.5 and 1 wt% fulfilled the requirements of being antimicrobial agent by providing highest inhibition effect on microorganisms. However, it is advisable to apply the tannic acid in lower concentrations with the maximal inhibitory effect on microorganisms due to its polyphenolic structure. Although it is a natural compound, there are limitations in the use of vegetable tanning materials due to their high organic load in the effluent, which results high COD [28, 34]. Nevertheless, lower COD values than 0.1% solution of commercial bactericide were obtained after application of tannic acid at highest tested concentration (1%), which is a good sign of their environmentally friendly structure. In the view of these findings, tannic acid can be appreciated as a natural and eco-friendly bactericide due to its antimicrobial, environmentally friendly and biodegradable properties.

ACKNOWLEDGEMENTS

The authors would like to thank Turkey Prime Ministry State Planning Organization for the supply of equipment (Project no: 2007 DPT 001).

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