Preliminary investigations of a textile fabric used as support for a sarcophagus from Astra Sibiu Museum

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INTRODUCTION

In order to establish a correct and personalized protocol for the optimal maintenance conditions of the objects with historical and artistic value from museums, a thorough comprehension of the materials from which these objects are made of and their physical and chemical properties is imperious [1]. The aim of this scientific study is to present the results of a preliminary assessment of a textile fabric which has been used as a support for a wooden sarcophagus containing an Egyptian mummy which is estimated to be over 2000 years old.

MATERIALS AND METHODS

Materials

The sarcophagus containing the Egyptian mummy can be found at the Museum of Universal Ethnography Franz Binder, which is a department of the National
Museum Complex ASTRA Sibiu. The three samples that were investigated come from different areas of the textile support for the wooden sarcophagus (figure 1). From the collective memory, the textile support was most likely introduced under the sarcophagus in the 1990s.

The samples collected from the textile fabric are presented in table 1 below. In addition to the three textile samples, the dust residue that was present in sample C was also analyzed.

| Sample label | Sample appearance |
|--------------|------------------|
| A            |                  |
| B            |                  |
| C            |                  |

Table 1

Microbiological assessment

The three archeological samples were subjected to microbial isolation of main strains present on the fabrics. Approximately 5 grams of each fabric was chopped in smaller pieces (in order to maximize sample surface) and put in a volume of 300 mL sterile physiological saline solution with Tween 80 anionic detergent (Fisher Scientific, UK), a polysorbate surfactant which helps in the process of protein stabilization, allowing a proper dispersion of plated microbial cells (prior to incubation). Samples were agitated on a Promax 2020 shaker (Heidolph) for 5 h at 220 rpm, in order to dislocate the potentially existing microbial cells from the surface of the fabrics. Bioburden isolation was carried out on nine semi-synthetic nutritive media, as following: Czapek-Doxy semi-synthetic media (Scharlau, Spain), used for cultivation of fungi, containing nitrogen as a sole source of nitrogen, frequently used for isolation of fungal species such as *Aspergillus*, *Penicillium*, *Paeclomycetes*, *Saccharomyces* etc.; Potato-Dextrose-Agar (Scharlau, Spain) usually used for stimulating sporulation and growing of various fungi strains (species of *Aspergillus*, *Saccharomyces*, *Rhodotorula*, *Geotrichum*, *Penicillium*, *Trichophyton* etc.); Malt-Extract-Agar (Scharlau, Spain), a classic culture medium for moulds and yeast (*Aspergillus*, *Saccharomyces*, *Penicillium*, *Candida* etc.), with a high quantity of sugar (maltose, glucose, sucrose) that allows excellent growth and additional necessary growth factors provided by the gelatine peptone; Sabouraud 4% Dextrose Agar (Merck, Germany), a complex medium for cultivation and isolation of yeasts and moulds (*Trichophyton*, *Microsporum*, *Geotrichum*, *Penicillium*), with a high concentration in dextrose, which promotes the formation of Conidia and Sporangia spores (combined with low pH value), as well as pigments of yeasts and molds, along with the inhibition of bacterial growth; Bengalrot-Agar with chloramphenicol (Roth, Germany), a selective medium for the enumeration of moulds and yeasts, with additional selectivity against bacterial growth, by the incorporation of the heat-stable antibiotic Chloramphenicol and glucose as incorporated fermentable carbohydrate source, with enzymatic digest of animal and plant tissues providing the essential vitamins, minerals, amino acids, nitrogen and carbon; Nutrient-Agar (Scharlau, Spain), a solid culture medium for general purpose use with less fastidious organisms; Muller Hinton Agar (Oxoid, United Kingdom), a medium usually used for the isolation of pathogenic Neisseria species, inclusion of starch ensuring that toxic factors found during growth will be absorbed, thus allowing the development of microorganisms that are present in very small inocula; Simple Gelose (Sanimed, Romania), simple structure media for the isolation and growth of certain strains of bacteria. Except for the Bengalrot, Nutrient-Agar, Muller Hinton and Simple Gelose medias, all other

Methods

**Scanning Electron Microscopy (SEM)**

This method is widely used for the analysis of the morphology of fiber and fabric surfaces [2, 3]. For example, Cybulskom. et al. [4] applied this technique in the analysis of archaeological textile samples from the Roman period and the Middle Ages and they were able to identify the constituent fibers of the evaluated material.

Morphological investigations of the textile samples and the dust in sample C were performed using a FEI Quanta 200 Scanning Electron Microscope. For each textile sample, a very small piece was cut and then it was placed on a specimen stub using double sided conductive carbon tape and analyzed using the following parameters: accelerated voltage: 20.00 kV; detector: GSED. In the case of the dust from sample C, the specimen stub covered with the double sided carbon tape was gently pressed onto the dust and analyzed using the same parameters as for the textile samples. All the SEM micrographs were taken using 1000X magnification.
nutritive medias had a concentration of 1g/L chloramphenicol (an antibiotic first isolated from cultures of *Streptomyces venezuelae*), for the inhibition of certain bacterial species. After samples agitation, 1000 µL were taken from each shaking solution and plated in duplicates on each nutritive media. Plates were then incubated at 28°C for 7 days and inspected for microbial morphological development.

**RESULTS AND DISCUSSION**

SEM analysis was used to investigate the microscopic appearance of the textile fiber surface, the type of fibers within the material, the diameter of these fibers, traces of insect and microorganisms attacks and the artifact residues (figure 2).

Following the SEM analysis, it can be said that the constituent fibers for all the samples are natural fibers of animal origin. However, the diameters of the constituent fibers vary widely, ranging between 15 and 90 microns. Due to the different appearance of the scales at the surface of the fibers and the very wide range of diameters, it can be preliminarily concluded that the fibers come from two different species of animals. Most likely, the fibers with more prominent and higher scales are wool fibers [5] and those with finer and less delimited scales are alpaca fibers [6].

For all of the textile samples, the extensive degradation caused by insects was highlighted. The most common insects that are known for attacking wool and other protein fibers are the larvae of the Tineidae (clothes moth) and Dermestidae (carpet beetle), although members of other species may attack wool incidentally [7].

Bioburden isolation from the archaeological samples highlighted various degrees of microbial development, based on both the originating archaeological sample, and the nutritive media used (figures 3–5), with certain samples having a higher bioburden degree, in terms of species, compared to others.

The highest bioburden load was registered by sample B (with 11 highlighted plates), followed by sample C (with 6 highlighted plates) and sample A (with 5 highlighted plates). The screening highlighted that all samples present both filamentous fungi specific structures and bacteria specific structures, with the nutritive media used for fungi screening (Czapek Dox, Sabouraud Agar, Bengalrot Agar and Malt Agar) highlighting more fungi-like morphological formations than the three media used for bacterial isolation (Nutrient Agar, Simple Gelose and Muller Hinton).

This separation was also strengthened by the presence of chloramphenicol in certain media, when compared to others that can also have a direct influence on the species of isolated strains.

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**Fig. 2.** SEM micrographs of the textile samples A, B and C, bar: 100 µm

**Fig. 3.** Bioburden isolation from archaeological sample A
CONCLUSIONS

The results obtained after SEM analysis show that the constituent fibers for all the samples are natural fibers of animal origin. Due to the particularity regarding the diameters, appearance and the surface of the fibers, it can be preliminarily concluded that the fibers come from two different species of animals.

Analyzing other existing data in the specialized literature, an estimation of the type of constituent fibers could be made, namely wool and alpaca fibers. In addition to the evaluation of the fiber type, with SEM it was possible to visualize the degree of degradation most likely caused by the attack of insects and different types of fungi and bacteria.
Bioburden isolation from the archaeological samples highlighted that the highest bioburden load was registered by sample B, followed by sample C and sample A.
The conclusion of the screening is that all samples present both filamentous fungi specific structures and bacteria specific structures [8].

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