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TITLE: BIOTRANSFORMATION OF INDUSTRIAL TANNINS BY FILAMENTOUS FUNGI

AUTHORS: Valeria Prigione^a, Federica Spina^a, Valeria Tigrini^a, Samuele Giovando^b and Giovanna Cristina Varese^{a*}

AFFILIATION: ^aDepartment of Life Sciences and Systems Biology, University of Turin, Viale Mattioli 25, 10125 Turin, Italy; ^b Centro Ricerche per la Chimica Fine Srl for Silvateam Spa, Via Torre 7, 12080 San Michele Mondovì (CN), Italy

***CORRESPONDING AUTHOR:** Phone: +39-011-6705984; E-mail: cristina.varese@unito.it

Abstract

Tannins are secondary metabolites widely distributed in the plant kingdom. They act as growth inhibitors towards many microorganisms: upon microbial attack, they are released helping to fight the infection of plant tissues. Extraction of tannins from plants is an active industrial sector with several applications from the beginning of the industrial era. Actually, tannins have many industrial applications in oenology, animal feeding, mining and chemical industry and, in particular, in the tanning industry. But tannins are also considered very recalcitrant pollutants in wastewaters of different origins. The ability to grow on plant substrates rich in tannins and on industrial tannin preparations is traditionally considered peculiar of some species of fungi that have developed mechanisms to tolerate the toxicity of tannins producing a complex enzymatic pattern active in the transformation of these substrates, mainly by hydrolysis and oxidation. Filamentous fungi capable of degrading tannins could have a strong environmental impact as bioremediation agents mostly in the treatment of tanning wastewaters.

Keywords: industrial tannins, filamentous fungi, tannery wastewater, biotransformation, bioremediation, laccases

Introduction

Tannins are a complex family of water-soluble polyphenolic compounds, synthesized as secondary metabolites by many plants (Haslam 1981). They are gathered from many types of trees and plants and can be present in barks, leaves, wood, as well in fruits and roots. Extraction of tannins is one of the first industrial activities related with nature, being an active industrial sector with several applications from the beginning of the industrial era. During the last two centuries several plants have been classified and proposed to be used as raw material, but only few are still of interest because really available and containing a sufficient quantity of tannins to provide economically sustainable processes. Usually, the production process is very simple, starting with a mere hot-water extraction, followed by some purification steps.

Today the total amount of industrially produced tannins in the world could be estimated around 170.000 tons per year, mainly extracted from: black wattle or mimosa (*Acacia mearnsii*) barks, quebracho colorado (*Schinopsis lorentzii* and *Schinopsis balansae*) hardwood, chestnut (*Castanea vesca* and *Castanea sativa*) hardwood, tara (*Caesalpinia spinosa*) pods extract, maritime pine (*Pinus pinaster*) barks, oak (*Quercus* spp.) hardwood, Chinese gallnut (*Rhus typhina semialata*) and Turkish gallnut or Aleppo galls (*Quercus infectoria*) (Food Chemical Codex 2003; FAO-JECFA 2005; European Pharmacopoeia 2005).

The wide application of tannins is mainly due to their ability to bind proteins, to the complexing capacity towards metals, and to the antioxidant capacity of the polyphenolic rings in their molecules. Traditionally, tannins are used in the tanning of leathers. This operation makes the animal hide a very resistant material to microbiological attack. Nowadays, tannins have many industrial applications: i) in oenology as processing aid in the clarification process they allow reducing the amount of SO₂ to stabilize the color; ii) in animal feeding as alternative to antibiotics to limit microbial proliferation; iii) in mining industry, they are used in rare elements recovery, as germanium used in the electronic industry; iv) in chemical industry, hydrolysable tannins are used as raw materials to produce gallic acid for the production of displays for smartphone and TVs.

On the other hand, industrial tannins, particularly when used in the tanning industry, can represent a serious environmental problem on a global level: even though vegetable tanning agents are natural materials, they are poorly biodegradable and act as growth inhibitors towards many microorganisms ultimately affecting the receiving ecosystem.

Filamentous fungi are known to populate usual and unusual ecological niches with even a strong anthropization pressure (Chambergo and Valencia 2016) and particular attention should be given to the strains isolated from matrixes rich in tannins since they may have developed an adapted metabolism indispensable to exploit tannins as source of nourishment. Actually, fungal strains capable to perform the biotransformation of polyphenolic substances contained in tannins could have a certain environmental impact as bioremediation agents. Moreover, biotransformed tannins could have a number of applications in agriculture, in the feed and wine industries and in the tanning process, for example improving the yields of tanning or the leather quality.

This work presents an overview on the interactions between filamentous fungi and industrial tannins: we focused on how tannins exert their action against fungi and, on the other hand, on the mechanisms used by fungi to biotransform tannins. Particular attention was given to the use of filamentous fungi for bioremediation of

wastewaters containing tannins, with the integration of the most recent data published on this application at industrial level.

Chemical composition of tannins

From a chemical point of view, it is difficult to define tannins due to their heterogeneity in terms of chemical composition and molecular weight (MW). Traditionally, tannins have been divided into two large groups: hydrolysable and condensed tannins (Haslam 1996; Scalbert 1991). Hydrolysable tannins are composed of a carbohydrate core whose hydroxyl groups are esterified with phenolic acids with a MW ranging from 300 to 5,000 Da (Mueller-Harvey and McAllan 1992). Depending on the substances produced following hydrolysis (by acids, basis or certain enzymes), hydrolysable tannins can be classified in gallotannins (yielding gallic acid) or ellagitannins (yielding ellagic acid). Tara and chestnut are representative of gallotannins and ellagitannins, respectively. Condensed tannins are oligomers where flavonoid units are condensed together with MW ranging from 1,000 to 20,000 Da. Quebracho extract is among the most industrially produced tannins predominantly composed by oligomers of profisetinidins.

The chemical composition of tannins could be explained considering as representative three of the main tannins: chestnut tannin, quebracho tannin and tara tannin. Chestnut tannin was shown to be a ellagic-type hydrolysable tannin by MALDI-TOF mass spectrometry (Pasch and Pizzi 2002). It contains castalagin (Fig. 1), which represents, with the isomer vescalagin, around 30% of the product. These substances and their higher oligomers are quite stable in chestnut tannin, and derive from rearrangement of polygalloylglucose naturally occurring in the chestnut wood (Pasch and Pizzi 2002). The higher oligomers contain repeating units of polygalloylglucose chain where galloyl groups can be linked differently to each other (Pizzi et al. 2009). The chestnut tannin also contains digalloyl glucose, glucose and gallic acid (Radebe et al. 2013).

The composition of tara tannin is based on a polymeric structure where monomers are esters of gallic acid on a core of quinic acid. The predominant structure is a pentagalloyl quinic acid (Giovando et al. 2013). The tannin contained in Tara pods, after extraction and purification, is known as tannic acid. Tara tannin could be represented by the structure in Fig 2.

The molecular composition of condensed tannins has been studied by means of MALDI-TOF mass spectrometry. In particular, quebracho tannin is composed by polyflavonoids of the type of profisetinidins (Pasch et al. 2001). More details were reported by Venter and collaborators (2012), who identified quebracho tannin as a mixture of polyanthocyanidins oligomers consisting of linear structures as represented by the tetramer in Fig 3.

Antimicrobial activity of tannins and fungal adaptation

The antimicrobial properties of tannins are well known: they act as growth inhibitors towards many microorganisms including bacteria, yeasts, and molds. Tannins exert antibacterial effects on a wide number of foodborne pathogens and infectious microorganisms (Aguilera-Carbo et al. 2008; Akhtar et al. 2015). For example, pomegranate fruits are an important source of hydrolysable tannins (punicalin, punicalagin, pedunculagin) and flavonoids (catechin, epicatechin) largely employed in folk medicine for the treatment of

various diseases such as ulcers, fever, diarrhoea, and microbial infections. Their pharmacological properties have been extensively studied: methanol extract of pomegranate peel has strong broad-spectrum antimicrobial properties against both gram-positive and gram-negative bacteria (Kharchoufi et al. 2018; Prashanth et al. 2001; Voravuthikunchai et al. 2005). The antibacterial efficacy of tannins is probably due to the alteration of biological membrane functionality by oxidative phosphorylation uncoupling, because of their lipophilic properties, causing the loss of cytoplasmic content (Cabral et al. 2013; Pane et al. 2016; Gupta et al. 2017).

Regarding the fungicide action of tannins, Elsherbiny and collaborators (2016) demonstrated that the methanol extract of pomegranate peels has a significant antifungal activity on the spore germination and mycelial growth of *Fusarium sambucinum*, one of the causal agents of potato tubers dry rot. Scanning and transmission electron microscopy observations revealed morphological modifications of hyphae such as curling, twisting and collapse, and dramatic changes of the cytoplasm and organelles. These changes could be the consequence of the disruption of cell permeability, which resulted in imbalance in internal osmotic pressure (Plodpai et al. 2013; Elsherbiny et al. 2016). Analogously, Kharchoufi et al. (2018), in a recent study about the fungicide action of pomegranate peel extract on *Penicillium digitatum*, observed by optic microscopy noticeable changes in the shape of hyphae, which appeared wilted and coiled. Other filamentous fungi susceptible to the action of the same extract were *Penicillium italicum*, *Rhizopus stolonifer* and *Botrytis cinerea* (Tehranifar et al. 2011).

With reference to yeasts, Endo and collaborators (2010) reported an irregular budding pattern and pseudo-hyphae in *Candida albicans* when treated with punicalagin and remarkable morphological alterations, which were visible by electron microscopy such as thickened cell wall, changes in the space between cell wall and the plasma membrane, vacuoles, and a reduction in cytoplasmic content. Two other *Candida* species, *C. stellatoidea* and *C. guilliermondii* were found to be susceptible to the antifungal activity of tannins (Haslan 1996; Vasconcelos et al. 2003).

Despite tannins have such well-documented antimicrobial activities, some filamentous fungi, yeasts and bacteria are resistant to them by the development of adaptive mechanisms and pathways for their degradation in natural habitats, where they can exploit ecological niches precluded to other organisms. For example Panno and collaborators (2013) studied the microbial community associated to *Posidonia oceanica*, a seagrass known for the high amount of tannins ranging between 55 and 95 $\mu\text{g/g}$ of dry weight of plant material. A total of 88 taxa were isolated, mainly Ascomycota, and many strains were recognized as good producers of tannases active in presence of high salt concentrations. These fungi, hence, were able to colonize and exploit lignocellulose residues, playing important ecological roles in marine environments (thanks to the substrate detoxification operated by tanning degrading fungi, leaves and rhizomes of *P. oceanica* may become available to other organisms) but can also be very useful in different biotechnological areas.

Some phytopathogenic *Ascochyta* species produce tannases, which help the invasion into the host plant by hydrolyzing the complex polyphenolic materials present in the bark of plants that usually confer protection against the attacking pathogens (Lekha and Losane, 1997). On the other hand, in mycorrhizal symbiosis, the ability to degrade tannic acid by *Oidiodendron* sp. may be necessary for the successful colonization of root cortical cells of members of Ericaceae (Li et al. 2006).

The ability to grow on plant substrates rich in tannins and on industrial tannin preparations is traditionally considered peculiar of some species of the genera *Penicillium* and *Aspergillus*, and in particular, of species belonging to the “black *Aspergilli*” group. In addition to Ascomycota, this ability seems to be widespread in the fungal kingdom, involving also Basidiomycota and Mucoromycota.

More than a century ago Knudson (1913) reported for the first time, that tannic acid could be degraded by an *Aspergillus niger* strain. In 1969, Lewis and Starkey reported that pure cultures of some terrestrial fungi grew on media containing tannins as sole carbon source, and some *Aspergillus* and *Penicillium* species grew on the surface of tannery pits and tannery wastes, and tannin-rich woods such as quebracho and chestnut. Moreover, different sources of tannins were compared, and both condensed and hydrolysable tannins were used as substrates. *Aspergillus*, *Penicillium*, *Fomes*, *Polysporus* and *Trametes* were shown to grow better on tannic acid (gallotannin) than on chestnut tannin (ellagitannin) or wattle tannin (condensed tannin) (Bhat et al. 1998).

The number of fungal species capable of growing in the presence of tannins, possibly using them as carbon source, is continuously increasing (Table 1). For example, Bhoite and Murthy (2015) cultured *Penicillium verrucosum* on coffee pulp as sole nutrient, and obtained up to 65% tannin degradation with up to 3.93-fold higher tannase concentration. More recently, Prigione and collaborators (2018), in a study on the mycobiota of industrial tannins, showed the ability of 125 isolates belonging to 10 fungal species to grow on culture media containing different types of industrial tannins as sole carbon source. In particular, five species (namely *Aspergillus acidus*, *Aspergillus costaricaensis*, *Aspergillus vadensis*, *Penicillium minioluteum*, *Talaromyces subinflatus*) had never been isolated from similar substrates, nor their ability to use tannins as the sole carbon source was reported before. Moreover, 10 fungal strains displayed a strong activity on chestnut and tara tannins in biotransformation tests by submerged fermentation.

The study of Prigione and collaborators (2018) highlighted that the choice of isolating autochthonous microorganisms is an appropriate practice to select strains with high applicative potential. Actually, tannases production is a strain-specific feature and fungi isolated from tannins are certainly adapted to these extreme environments, since the high tannin content exerts a selective pressure on the fungal community. Therefore, the isolates likely developed a unique adapted metabolism in order to exploit tannins as source of nourishment.

Biotransformation mechanisms

Some fungi have developed mechanisms to tolerate the toxicity of tannins, indeed producing a complex enzymatic pattern active in the transformation of these substrates. Hydrolysis and oxidation are the main reactions that lead to tannins biotransformation.

Tannases

Tannases are the main enzymatic class involved in the tannins hydrolysis. Tannases, namely tannin acyl hydrolase (EC 3.1.1.20), hydrolyze ester and depside bonds of hydrolysable tannins to produce glucose and gallic acid (Lekha and Lonsane 1997). Known to be produced also by bacteria (Reveron et al. 2017; Tomás Cortázar et al. 2018) and plants (Bains et al. 2009), fungal tannases are well known for their catalytic versatility.

Numerous fungi are known to be tannases producers, including strains belonging to *Penicillium*, *Aspergillus*, *Neurospora*, *Trichothecium*, *Fusarium*, *Trichoderma*, *Paecilomyces*, *Emericella*, *Lenzites* genera

(Schons et al. 2011; Ordonez et al. 2011; Bressan Gonçalves et al. 2013; Zakipour-Molkabadi et al. 2013; Herrera Bravo de Laguna et al., 2015). The choice of the fungus to be used is a critical point. For instance, Zakipour-Molkabadi and collaborators (2013) inoculated eight strains in the presence of tannic acid 1% as sole carbon source: tannase activity was always lower than 1 U/ml with the exception of *Penicillium* sp. that produced 4.33 U/ml after 96 h.

As regards their cellular localization, contrasting results have been reported. Known to be extracellular enzymes, de Böer and collaborators (2009) found specific secretion signal sequences on the yeast *Arxula adenivorans* tannase gene. On the contrary, *Aspergillus niger* and *Verticillium* sp. produced both extracellular and intracellular tannases (Aguilar et al. 2001; Kasieczka-Burnecka et al. 2007); the productivity of intracellular tannases was even higher than extracellular enzymes for *Aspergillus aculeatus* (Banerjee et al. 2001).

Purified tannases have been obtained from *A. niger* (Mata-Gómez et al. 2009, Ramos et al. 2011), *Aspergillus carbonarius* (Serrani Valera et al. 2015), *Aspergillus awamori* (Beena et al. 2011), *Aspergillus oryzae* (Abdel-Naby et al. 2016), *A. adenivorans* (Böer et al. 2009), *Lenzites elegans* (Ordonez et al. 2011), *Penicillium* sp. (Zakipour-Molkabadi et al. 2013) and the antarctic strain *Verticillium* sp. (Kasieczka-Burnecka et al. 2007) among others. Thanks to this information, the phenomenon at the base of tannase production has been clarified. Tannases production is an inducible process that can be obtained both in solid state and liquid fermentation. The process is mostly controlled by the source of C and the presence in the medium of structural constituents of tannic acid pyrogallol, methyl gallate and gallic acid (Aguilar et al. 2007; Zakipour-Molkabadi et al. 2013).

According to their ecological role, fungi respond to tannins-rich substrates by producing tannases: agricultural byproducts (i.e. tamarind seed powder, palm kernel cake, tea and coffee residues, olive mill wastewater, etc.) could be used for fungal fermentation to produce high-value products like tannases (Aissam et al. 2005; Jana et al. 2012; Baik et al. 2014; Sabu et al. 2015, Bhoite and Murthy 2015). These findings were confirmed by in-depth analysis of tannase genes. *ATANI* is the tannase gene of *A. adenivorans*, a common fungus model whose tannase was characterized and cloned. Its activation is controlled by a CAAT box-like sequence but no TATA box-like sequence were identified: expression profile revealed that tannase production is induced by tannic acid or gallic acid, whereas glucose did not trigger the same effect. When glucose feeding was shifted to tannic acid or gallic acid, the gene was rapidly activated and, after 2 h, the enzymes were already detectable (Böer et al. 2009). However, there are still few unclarified steps of the induction mechanism, including the role of tannin-related compounds as synthesis inducers. For instance, gallic acid enhanced tannase productivity of *A. niger* at higher extent in solid-state fermentation rather than ineffective submerged cultures (Aguilar et al. 2001).

Fungal tannases have a pH optimum at acidic values and are mildly tolerant to high pH and temperature. These data are strictly dependent of the microbial enzymatic source, according probably to isolation origin and adaptation skills evolved by each fungus. Tannase activity is usually maximal at pH 5.5 and 35°C and are stable at pH 5-6 and 25-35°C (El-Tanash et al. 2011; Zakipour-Molkabadi et al. 2013). Particular attention is given to those enzymes with better catalytic features. Tannase from *P. variotii* showed 95% of activity recovery in the range of 3.5-7.5 pH and 20-50°C (Schons et al. 2011). The Basidiomycota *L. elegans* secreted a tannase

effective from 40 to 60°C, whose activity was only halved at 80°C (Ordóñez et al. 2011). Promoting structural modifications on the active site, tannase activity is generally inhibited by heavy metals and metal ions (Serrani Valera et al. 2015; Yao et al. 2014), even though some exceptions can be found. Although being inhibited by metal ions, a tannase from *A. carbonarius* was not perturbed by Ag^+ : the activity even increased up to 32% according to the Ag_2SO_4 concentration (Serrani Valera et al. 2015). Interestingly tannases can be considered solvent- and salt-tolerant enzymes, making them good candidates for biotechnological application in wastewater treatment (Serrani Valera et al. 2015). Indeed tannase from *Penicillium* sp. maintained 50% of the activity in the presence of 1 M NaCl after 24 hours (Zakipour-Molkabadi et al. 2013). Activity recovery of tannase from *A. carbonarius* was always higher than 80% after exposition to detergents (SDS and Tween-20) and organic solvents (methanol, ethanol, acetone, isopropanol, n-butanol and glycerol) (Serrani Valera et al. 2015).

Enzymes may be used for industrial application, becoming an intriguing alternative to whole cell systems. Although a time-saving technology requiring very low operational control, enzymes are still considered as a high-cost system with room for improvement at their catalytic stability over time and stressors (Rao et al. 2014). Due to their biochemical properties, the demand for industrial enzymes is increasing, asking for the development of solutions feasible for industrial application. Stable and re-usable enzymes may overcome technical and economic drawbacks, driving to increasing interest in immobilized enzymes (Arca-Ramos et al. 2018). Tannases have been immobilized using several supports as Eupergit® (Crestini and Lange 2015), carbon nanotubes (Ong and Annuar 2018), gelatin (El-Tanash et al. 2011), sodium alginate, gellan gum and pectin (Schons et al. 2011), agarose (Bressan Goncalves et al. 2013), magnetic diatomaceous (Silva de Lima et al. 2018). Advanced techniques have been successfully applied as cross-linking (Ong and Annuar 2018), microencapsulation in calcium alginate membranes (Yu et al. 2004), etc. The choice of the immobilization method has a central role in the success of the treatment, because inadequate systems can influence the diffusion of molecules and products (El-Tanash et al. 2011). Respect to free tannases, immobilization allows repetitive cycles (Ong and Annuar et al. 2018) and enhances the enzymatic stability, enlarging the pH and temperature range of action (Yu et al. 2004; Schons et al. 2011; Kumar et al. 2015). Immobilization can also protect the enzymes from inhibition effects played by metal ions. Tannases from *A. aculeatus* were more stable by metal ions disturbance when immobilized on gelatin: in the presence of several ions, the activity recovery was always higher than free enzymes, with peaks of 20% activity increase for K^+ (El-Tanash et al. 2011). Positive effects can be also observed in the catalytic efficiency of the immobilized enzymes. For instance, tannase from *A. niger* on sepharose beads better converted methyl gallate into gallic acid: process yields increased from 30% up to 40% (Sharma et al. 2002). According to the authors, immobilized enzymes are less perturbed by the competitive inhibition by gallic acid, which is instead electrostatically pushed away from the positively charge support.

The competitiveness of tannase-based methodology lays the foundation on the rapidity of the catalyzed reactions, positively affecting the feasibility and the sustainability of the proposed methodology. *A. oryzae* tannase can hydrolyse tannic acid forming at first penta-, tetra- and tri- galloylglucose during the first 10 min, which are further transformed in gallic acid and mono galloylglucose after 15 min. The reaction continues having gallic acid as the sole end product after 35 min incubation (Abdel-Naby et al. 2016). Immobilized and free tannases can be indeed applied in several biotechnological fields as environmental bioremediation,

pharmaceutical and food industry (e.g. clarification of beverage, manufacture of instant tea, etc.). Tannase from marine *A. awamori* catalyzed transesterification reaction of tannic acid producing propyl gallate, an antioxidant used in the food industry with nutraceutical importance (Beena et al. 2011). Tannase from marine *A. niger* transformed green tea extracts, rich of polyphenols that are responsible for the formation of sediment and haze (Baik et al. 2014). *P. verrucosum* tannase treatment of pomegrade and aonla juice enhanced the clarity and removed astringency (Bhoite and Murthy 2015). It has been proved that tannase-treated straw have a reduced content of tannins, along with an improved nutritional content useful for cattle feed (Raghuwanshi et al. 2014).

Gallic acid decarboxylases

Among tannin-transforming enzymes, tannases are not the only enzymes involved. Following gallic acid production by tannases, gallic acid decarboxylases (gallate decarboxylase, EC 4.1.1.59) indeed catalyze its decarboxylation to pyrogallol. Known to be produced mainly by bacteria (Zeida et al. 1998; Jiménez et al. 2013), very few of these enzymes have been characterized because highly unstable and known to be sensitive to oxygen (Zeida et al. 1998). Recently also fungi have been identified as gallate decarboxylase producers. The first report of fungal nonoxidative decarboxylation of tannins is the study of Sietmann and collaborators (2010) on the yeast *A. adenivorans*: pyrogallol was the main reaction product of gallic acid decarboxylation. Studying the transformation pathway, the authors suggested pyrogallol as the substrate for further ring fission, forming as end-product the 2-hydroxymuconic acid. This enzyme was further purified and characterized (Meier et al. 2017). The optimal pH and temperature is between 5.6 and 7.1, and between 25 and 45°C, respectively. The enzyme was almost completely inhibited by buffer and some metal ions.

Gallic acid decarboxylases expression is specifically controlled by gallic acid, whereas other hydroxybenzoic acids do not induce its production (Meier et al. 2017). Noteworthy, gallic acid induce also catechol-1,2-dioxygenase, even though its role in the tannins degradation is currently unclear (Meier et al. 2017). This observation may suggest that these enzymes are specifically involved in the transformation pathway of gallic acid. Kinetic analysis confirmed this assumption: both bacterial and fungal gallic acid decarboxylases have a high affinity for gallic acid, catalyzing reactions against gallic acid but not on structural analogues as benzoic acid derivatives (Zeida et al. 1998; Meier et al. 2017).

Laccases

Fungi also catalyze oxidation reactions leading to the transformation of phenolic compounds as tannins by extracellular oxidative reductases as laccases (EC 1.10.3.2) and peroxidases (EC 1.11.1.x) that are among the most known and studied fungal enzymes (Janusz et al. 2017). Although not constitutively produced by fungi, gene transcription is usually regulated by metal ions and catabolic byproducts of lignin and tannin (Martinez 2002; Piscitelli et al. 2011). Natural lignin-based materials rich of tannins as corn cob, wheat straw, saw dust and evergreen oak litter can be used as growth medium to trigger extracellular secretion of oxidoreductases (Tagger et al. 1998; Sahay et al. 2009).

Laccases are involved in the gallotannin, ellagitannin and proanthocyanidins biosynthesis by plants (Niemetz and Gross 2005; Hu et al. 2013) whereas in fungi may trigger oxidation cascades against tannins and

their derivatives. Indeed, laccases have a high affinity for phenol monomers including catechol, pyrogallol and gallic acid (Sahay et al. 2009; Itoh et al. 2016; Zerva et al. 2016). The reaction often leads to the production of dimers and phenol polymers (Zerva et al. 2016).

Although hydrolyzable tannins are usually a better substrate for laccase than condensed tannins (Widsten et al. 2009), laccase from *M. thermophila* was used to perform the grafting of polyphenols (condensed and hydroxypropylated tannins) to improve wood preservation reducing the water absorption and enhancing the hydrophobic properties (Filgueira et al. 2017). Laccases from *Myceliophthora thermophila* were also applied in the fiberboard manufacturing, providing an ecological sustainable alternative to common process that use formaldehyde-based adhesives: formaldehyde emissions pose indeed important health concern (Widsten et al. 2009). Laccases activated tannin and wood fibers, by forming radicals that later create covalent bonds among fibers by radical coupling: when the interfiber bonds are widespread on the surface, the adhesive effect can be considered similar to synthetic resins (Widsten et al. 2009; González-García et al. 2011). This reaction is at the base of wood decay prevention strategies, where tannins are under study as natural preservatives against wood decaying fungi. Laccase from *Trametes hirsuta* catalyzed polymerization reactions to fix phenolic preservatives as tannin acid and vanillin acid to the wood fibers: the treatment enhanced the resistance to microbial attack by the brown-rot fungus *Coniophora puteana* and the white-rot fungus *Coriolus versicolor* (Ratto et al. 2004).

Peroxidases

The role of peroxidases in tannins transformation has been mostly unexplored, even though they are recognized among the most powerful fungal enzymes. Despite the redox potential of peroxidases falls in the range of 1100–1500 mV, significantly higher than laccases, stability issues mostly limited their application (Ravichandran and Sridhar 2017). Post-translational modifications are under study to enhance this feature (Sáez-Jiménez et al. 2015). Even though gallic acid is involved in peroxidase metabolism as production inducer or oxidation cascade mediator (Mishra and Jana 2017; Mishra et al. 2017), very few reports studied their actual involvement in tannin degradation. Heme peroxidases from *Phanerochaete chrysosporium* were capable to degrade tannic acid and condensed tannins, and removed tannin components from spent tan liquor. Studying the enzymatic kinetic, authors suggested that complexation reactions were indeed followed by the cleavage of ethereal linkage, hydroxylation reaction, aromatic ring opening and ultimately oxidation to the end products (Gnanamani et al. 2001).

Fungal bioremediation of wastewaters containing tannins

Tannins are considered very recalcitrant pollutants in wastewaters of different origins, e.g. winery and beer wastewater (Moreira et al. 2015; Yague et al. 2000), pulp and paper mill effluents (Vepsäläinen et al. 2011), olive mill wastewaters (Aissam et al. 2005), landfill leachates (Bardi et al. 2017b) and mainly tannery wastewaters (He et al. 2007). The core of wastewater treatment plants (WWTPs) consists of engineered biological ecosystems (activated sludge) based on bacteria that allow an economical and sustainable treatment of wastewaters (Leyva-Díaz et al. 2017). However, bacteria are ineffective in the removal of compounds characterized by high toxicity and recalcitrance, such as tannins (Lofrano et al. 2013; He et al. 2007). A

substantial percentage of the Total Organic Carbon (TOC) in the effluent of the biological treatment phase in tannery WWTPs is attributable to natural and synthetic tannins (Munz et al. 2009). Condensed tannins, in particular, are more resistant to microbial attack than hydrolysable ones and have stronger toxic effect on microbial populations (Mutabaruka et al. 2007). Despite a direct inhibiting effect cannot be associated with the tannin concentration in the effluent, a lower specific growth rates of ammonium and nitrite oxidising bacteria indicates a strong environmental pressure due to the tannery wastewater (Munz et al. 2009). In this regard, the WWTP implementation with a fungal treatment could be useful to achieve better depuration results (Zhang et al. 2015).

Fungal treatment aimed to the tannin removal can be based on three mechanisms: adsorption (Natarajant and Manivasang, 2018), degradation (Zhang et al. 2015), and precipitation induced by extracellular enzymatic polymerisation (Yamaguchi et al. 1998). These mechanisms could run at the same time or be sequential (Song and Burns 2005).

In general, biosorption occurs within the first few hours but a change in the process conditions or long term operations could cause backward transformation or desorption (Tigini and Varese, 2018). Besides, this option requires an adequate disposal or further treatment of the loaded biosorbent, since it consists in the transfer and concentration of pollutants from one medium to another. For these reasons, fungal biosorption appears more adequate for the treatment of chromium tannery wastewaters, aimed to the recovery of chromium ions (Hintermeyer and Tavani, 2013).

Biodegradation is the most investigated strategy in fungal bioremediation of vegetable tanning wastewaters. Despite enzymes can be applied, economic drawbacks have discouraged their investigation in this field (Aguilar et al. 2007, Govindarajan et al. 2016). A whole-cell system can degrade tannins, despite the lack of information about the acting metabolic pathways (Chávez-González et al. 2012). Several fungal species, mainly belonging to *Aspergillus*, *Penicillium*, *Paecilomyces* and *Fusarium* genera have been reported to utilize both hydrolysable and condensed tannins as sole carbon source. However, no report of their application at real scale on vegetable tannery wastewaters is reported (Chaudhary et al. 2017). An application to a real scale encounters several issues to be addressed, since typical conditions of conventional WWTPs are unfavourable to fungal growth. The scale up of an effective fungal treatment is still an actual challenge.

The first goal is to obtain a stable system in real operative conditions where allochthonous fungi are often not able to compete with autochthonous microorganisms (Svobodová and Novotný, 2018). In particular, during the start-up period the faster growth rate of bacteria can cause the fungal growth suppression and mycelium disgregation, that turn in outclass of fungi (Rene et al., 2010). Moreover, bacterial colonisation can induce a drop in enzymatic activity, deeply affecting the treatment effectiveness (Gao et al. 2008). Autochthonous fungi can also affect the efficiency of the inoculum development in bioreactors as happens in the presence of *Trichoderma* spp., mycoparasites able to degrade tannins, and consequently often present in tannery wastewaters (Badia-Fabregat et al. 2017; Tigini et al. 2018). Therefore, many studies were indeed carried out in sterile conditions, but sterilization is impracticable in a full-scale reactor system, due to high operational costs (Gao et al. 2008).

Different strategies have been suggested to enhance the resistance of fungi in non-sterile conditions. Mature and well-developed fungal inoculum, possibly immobilised in an adequate support, is recommended (Spina et al. 2012; Bardi et al. 2017). The control of the environmental conditions (e.g. pH adjustment, nitrogen sources) may support the success of fungal bioreactors (Bardi et al. 2017b; Wu et al. 2018). To avoid negative effects on fungal metabolism and biomass growth, these operations must be carefully optimised (Badia-Fabregat et al. 2017; Wu et al. 2018).

The addition of co-substrates as selective carbon source, not exploitable by bacteria or with antibacterial effect, may help fungi to compete with them (Badia-Fabregat et al. 2017; Bardi et al. 2017b). Moreover, alternative carbon sources could be required by white rot fungi for their growth, as they often degrade pollutants with a co-metabolism mechanism, as well as enhancing the tannase production (Ordóñez et al. 2011). This is true also for fungi that produce oxidative enzymes with the aim of detoxifying their environment causing tannin precipitation by polymerisation. For instance, malt extract strongly enhanced quebracho removal by *Aspergillus tubingensis* (Spennati et al. 2017). This species produces high amount of laccases in addition to tannases (Spennati et al. 2016; Iqbal et al. 2018). Alternatively, lignocellulose material would be a strategic choice, operating a selective pressure against bacteria and supporting fungal co-metabolism at the same time (Spennati et al. 2016). However, the co-substrate addition should be evaluated in order to be a sustainable option when applied in a WWTP. Besides, strong catabolite repression of tannase synthesis in *A. niger* can occur when alternative carbon source is added, probably due to changes in carbon/nitrogen ratio (Aguilar et al. 2001). This must be kept in mind in the case of treatments specifically aimed to tannin degradation by means of tannases.

Lastly, bioreactor configurations and elevated retention time (1-3 days) are also useful to acquire a better understanding of the process, which facilitates the up-scaling and highlights possible unexpected effects like the ecotoxicity increase or excessive biomass growth (Spina et al. 2012; Chanda et al. 2016; Spennati et al. 2017).

Another future challenge is the investigation of microbial relationships, with the aim to enhancing the exploitation of fungal natural degradative ability towards tannins. In nature as well in wastewater oxidative biological treatment, microorganisms do not exist in isolated form, and they coexist with each other, establishing biological competences among the interacting species. Biotransformation in nature is commonly a combination of metabolic pathways from different microorganisms (Ijoma and Tekere 2017). Moreover, microbial consortia are more vigorous to environmental variations and better survive in nutrient limitation thanks to metabolites exchanging or by trading molecular signals. Each biological entity identifies the presence of other organisms in the consortium and acts in response to that (Chaudhary et al. 2017). Microbial consortia can serve unknown functions potentially inducing alternative degradation pathways and metabolites in bioreactors. In an established consortium, several enzymatic isoforms from different species may be complementary in the degradation of complex matrices, and mutualistic or antagonistic relationships may ultimately enhance enzymatic production (Ijoma and Tekere 2017). Furthermore, different species showed specific activity dependent on the tannin to be treated (Prigione et al. 2018). Likewise some strains are more effective against the Chemical Oxygen Demand (COD) and other on the total phenol content (Perovano et al. 2011). The development of a consortium is sometimes recommended, in order to achieve better yields in the bioremediation of tannins wastewaters.

The study of autochthonous fungal consortia could enhance their exploitation in *in-situ* bioaugmentation strategy, too. The presence of fungi in activated sludge is established (Evans and Seviour 2012; Fathi et al. 2017) as the activity specificity according to pollutants of interest (Tigini et al. 2018). Nevertheless, during normal operational condition, hyphal colonisation of flocks in activated sludge is not predominant, so their role has been underestimated up to now (Liu et al. 2017). Besides high tolerance and adaptation to pollutant toxicity and microorganisms' competition (Herrero and Stukeley 2015), fungi could be involved both in the direct degradation of pollutants, and in the formation/stabilisation of adequate ecosystems for the development of degrading organisms (Liu et al. 2017).

Some possible configurations for tannin fungal treatment were hypothesized for future applications based on different tannin concentrations (Spennati et al. 2018). Firstly, vegetable tannin bath would be treated with an optimized process based on fungal bioreactors, before the conventional activated sludge process, exploiting high selective pressure given by high tannin concentration as an important factor for fungal biofilm stability. However, the maximum load of tannins treatable by fungi should be established for evaluating a dilution of tannery wastewater in case. Another possible strategy could be the treatment in a side-stream reactor with a recirculation system in active sludge, with the aim to remove the residual recalcitrant COD. The adjustment of nutrient load in the effluent could allow obtaining a stable fungal biofilm. Fungi predominate in the microbiome of WWTPs treating nitrogen deficient wastes, indeed (Evans and Seviour 2012). In any case, scientists are surely called to face the challenge of implementing of WWTPs with real scale fungal treatment in the next future.

Conclusions

Filamentous fungi are undoubtedly endowed with high application potential in the field of industrial tannins biotransformation. The more tannins are finding new applications the more fungi will play an increasingly important role both in biotransformed tannin production and in tannin biodegradation. Nevertheless, even though there are numerous references to the isolation of fungi from soils, plant material or extreme environments capable of degrading tannins, the report of successful applications at real scale are still missing. The actual challenge in the next future is to harness the potential of fungi in order to open the way for tannin exploitation, taking advantage of the infinite applicative resources of these heterogeneous compounds.

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Conflict of Interest: The authors declare that they have no conflict of interest.

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Table 1 Filamentous fungi capable of producing tannases and/or of using tannins as sole carbon source.

Fungus	Reference
<i>Ascochyta boltshauseri</i>	Lekha and Losane 1997
<i>Ascochyta pisi</i>	Lekha and Losane 1997
<i>Ascochyta viciae</i>	Lekha and Losane 1997
<i>Aspergillus acidus</i>	Prigione et al. 2018
<i>Aspergillus aculeatus</i>	Banerjee et al. 2001; El-Tanash et al. 2011
<i>Aspergillus aureus</i>	Bajpai and Patil 1997
<i>Aspergillus awamori</i>	Bradoo et al. 1996; Mahapatra et al. 2005; Beena et al. 2011; Prigione et al. 2018
<i>Aspergillus caespitosus</i>	Batra and Saxena 2005
<i>Aspergillus carbonarius</i>	Serrani Valera et al. 2015
<i>Aspergillus carneus</i>	Ganga et al. 1977
<i>Aspergillus costaricaensis</i>	Prigione et al. 2018
<i>Aspergillus ficuum</i>	Ma et al. 2015
<i>Aspergillus fischeri</i>	Bajpai and Patil 1997
<i>Aspergillus flavipes</i>	Ganga et al. 1977
<i>Aspergillus flavus</i>	Yamada et al. 1968; Batra and Saxena 2005
<i>Aspergillus foetidus</i>	Banerjee et al. 2001
<i>Aspergillus fumigatus</i>	Batra and Saxena 2005
<i>Aspergillus japonicus</i>	Ganga et al. 1977; Bradoo et al. 1996
<i>Aspergillus nidulans</i>	Ganga et al. 1977
<i>Aspergillus niger</i>	Knudson 1913; Haslam and Stangroom 1966; Lekha and Lonsane 1994; Bradoo et al. 1996; Rana and Bhat 2005; Cruz-Hernandez et al. 2005; Murugan et al. 2007; Prigione et al. 2018
<i>Aspergillus oryzae</i>	Iibuchi et al. 1967; Doi et al. 1973; Bajpai and Patil 1996; Bradoo et al. 1996; Huang et al. 2007; Abdel-Naby et al. 2016
<i>Aspergillus parasiticus</i>	Ganga et al. 1977; Bajpai and Patil 1996
<i>Aspergillus tamarii</i>	Lekha and Losane 1997
<i>Aspergillus terreus</i>	Ganga et al. 1977; Bajpai and Patil 1997
<i>Aspergillus tubingensis</i>	Prigione et al. 2018
<i>Aspergillus ustus</i>	Ganga et al. 1977
<i>Aspergillus vadensis</i>	Prigione et al. 2018
<i>Aspergillus versicolor</i>	Batra and Saxena 2005
<i>Chaetomium globosum</i>	Lekha and Losane 1997
<i>Cryphonectria parasitica</i>	Farias et al. 1994
<i>Cunninghamella</i> sp.	Bradoo et al. 1996
<i>Emericella nidulans</i>	Bressan Gonçalves et al. 2013
<i>Fusarium oxysporum</i>	Bradoo et al. 1996
<i>Fusarium solani</i>	Bajpai and Patil 1996; Bradoo et al. 1996
<i>Helicostylum</i> sp.	Bradoo et al. 1996
<i>Lentinus edodes</i>	Zheng and Shetty 2000; Vattem and Shetty 2003
<i>Lenzites elegans</i>	Ordóñez et al. 2011
<i>Mucor</i> sp.	Belmares et al. 2004
<i>Myrothecium verrucaria</i>	Lekha and Losane 1997
<i>Neurospora crassa</i>	Bradoo et al. 1996
<i>Paecilomyces variotii</i>	Mahendran et al. 2005; Battestin and Alves-Macedo 2007; Schons et al. 2011; Herrera Bravo de Laguna et al. 2015; Prigione et al. 2018
<i>Penicillium crustosum</i>	Batra and Saxena 2005
<i>Penicillium charlesii</i>	Bradoo et al. 1996
<i>Penicillium chrysogenum</i>	Rajakumar and Nandy 1983; Bajpai and Patil 1996; Bradoo et al. 1996; Karpe et al. 2015
<i>Penicillium citrinum</i>	Bradoo et al. 1996

<i>Penicillium corylophilum</i>	Bradoo et al. 1996
<i>Penicillium digitatum</i>	Bradoo et al. 1996
<i>Penicillium fellutanum</i>	Ganga et al. 1977
<i>Penicillium glabrum</i>	Van de Lagemaat and Pyle 2005
<i>Penicillium glaucum</i>	Lekha and Losane 1997
<i>Penicillium islandicum</i>	Ganga et al. 1977
<i>Penicillium minioluteum</i>	Prigione et al. 2018
<i>Penicillium montanense</i>	Silva de Lima et al. 2014
<i>Penicillium notatum</i>	Ganga et al. 1977
<i>Penicillium restrictum</i>	Batra and Saxena 2005
<i>Penicillium spinulosum</i>	Prigione et al. 2018
<i>Penicillium variabile</i>	Batra and Saxena 2005
<i>Penicillium verrucosum</i>	Bhoite and Murthy 2015
<i>Rhizopus oligosporus</i>	Vattem and Shetty 2002
<i>Rhizopus oryzae</i>	Hadi et al. 1994; Purohit et al. 2006
<i>Syncephalastrum racemosum</i>	Bradoo et al. 1996
<i>Talaromyces subinflatus</i>	Prigione et al. 2018
<i>Trichoderma hamatum</i>	Bradoo et al. 1996
<i>Trichoderma harzianum</i>	Bradoo et al. 1996
<i>Trichoderma viride</i>	Bajpai and Patil 1996; Bradoo et al. 1996
<i>Trichothecium roseum</i>	Lekha and Losane 1997
<i>Verticillium</i> sp.	Kasieczka-Burnecka et al. 2007

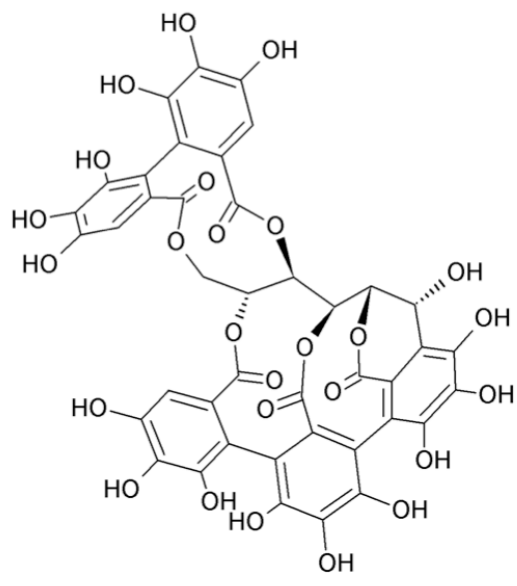


Fig. 1 Molecular structure of castalagin, the main component in the chestnut tannin

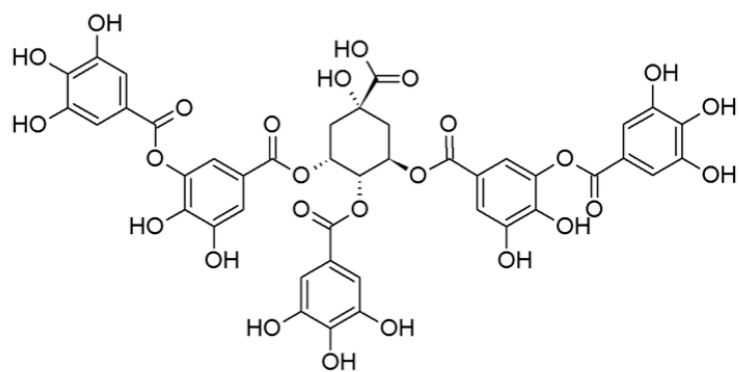


Fig. 2 Molecular structure of pentagalloyl quinic acid, the main component in the tara tannin

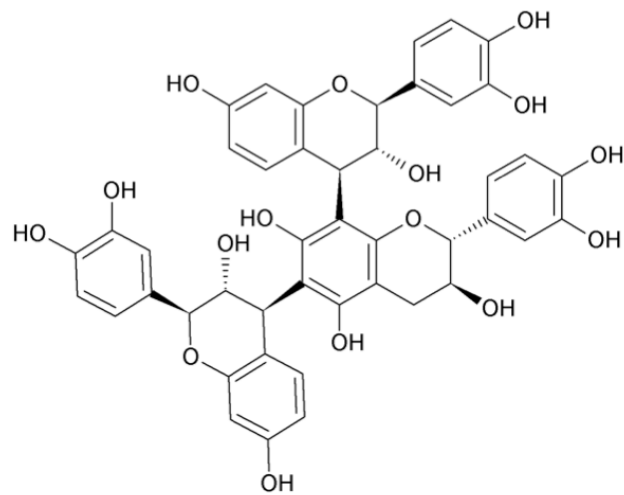


Fig. 3 Molecular structure of fisetinidin trimer, the main component in the quebracho tannin