

Immobilisation of laccase from Trametes versicolor on a hydrophilic PVDF microfiltration membrane: characterization of the grafted support and application in the removal of a phenylurea pesticide in waste water

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Claude Jolivalt, Eliane Caminade, Christian Mougin, Maxime Pontié. Immobilisation of laccase from Trametes versicolor on a hydrophilic PVDF microfiltration membrane: characterization of the grafted support and application in the removal of a phenylurea pesticide in waste water. Euromembrane 2000, Sep 2000, Jérusalem, Israel. 2000. hal-01600755

HAL Id: hal-01600755

https://hal.science/hal-01600755

Submitted on 12 Mar 2023

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Immobilisation of laccase from T. versicolor on a microfiltration membrane and application in the removal of a phenylurea pesticide

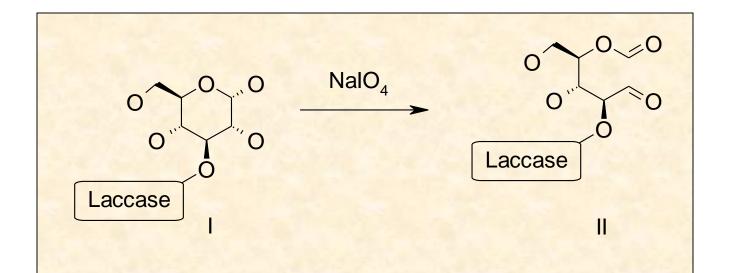
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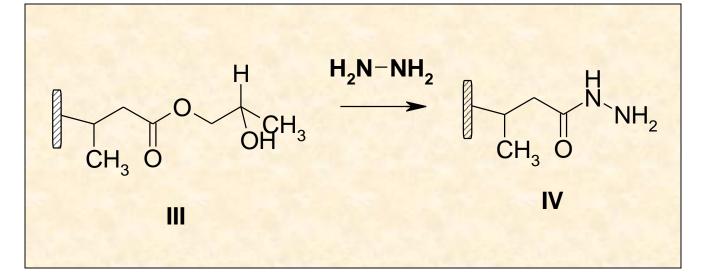
OBJECTIVES

In the present work, laccase from Trametes versicolor was covalently immobilised on a hydrophilic PVDF microfiltration membrane using a site-directed immobilisation technique. The protein-grafted membrane was used in a filtration module to transform N',N'-(dimethyl)-N-(2-hydroxyphenyl)urea (2-HF), an herbicide derivative known to be oxidised into insoluble compound in the presence of laccase. The characterisation of the microfiltration membrane was performed by means of physico-chemical methods (contact angle and streaming potential measurements, XPS) throughout its modification procedure.

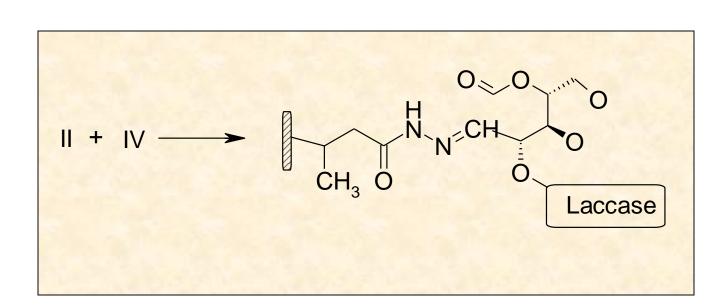
Immobilisation procedure



Step 1: Oxidation of laccase 80 % of the initial laccase activity retained

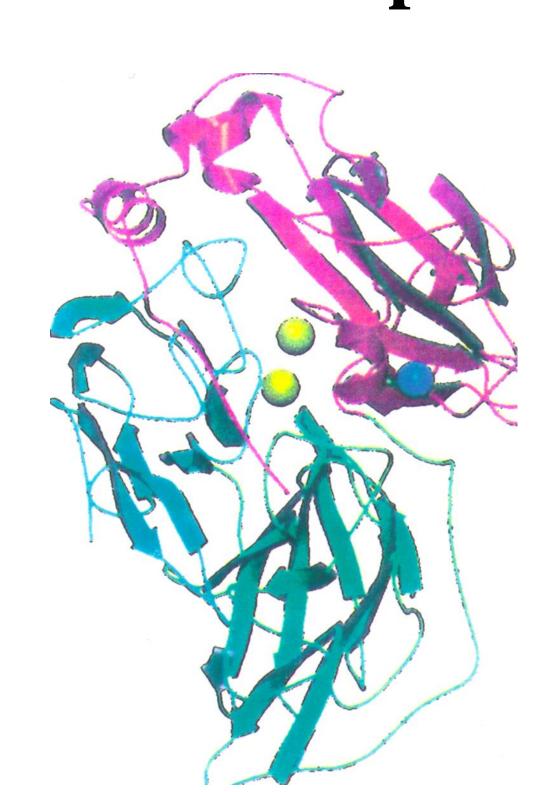


Step 1bis: Activation of the hydrophilic PVDF membrane (Millipore, Durapore 0.45µm) by hydrazine Grafting rate: 60% from XPS measurements



Step 2: Covalent coupling of laccase to the PVDF membrane Activity measurements in batch: 25 % recovered Evidence for diffusion limitations in batch experiments

Properties of Laccases

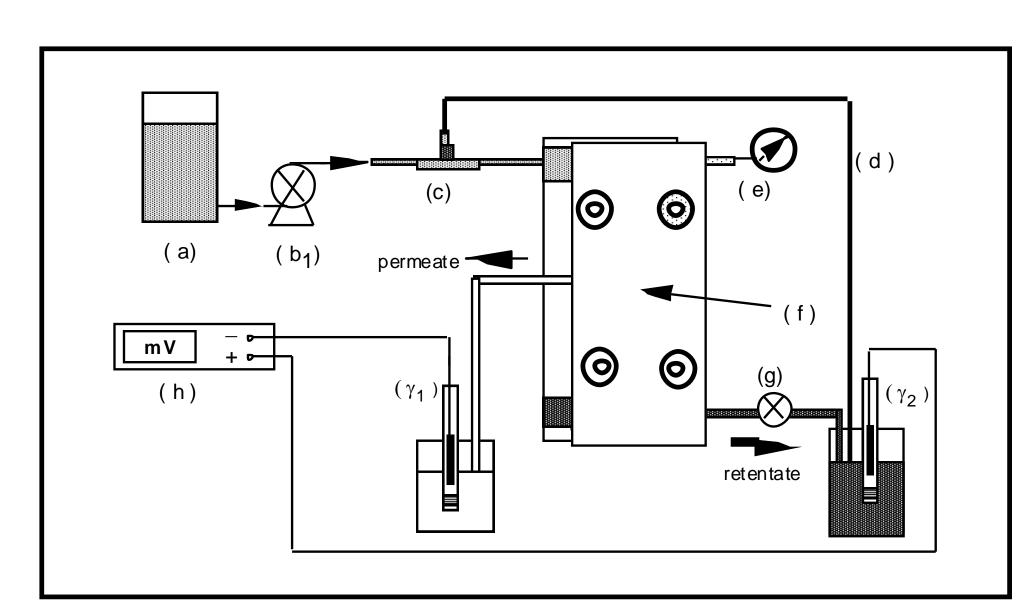


Laccases (EC. 1.10.3.2) catalyses the oxidation of various compounds, which concomitantly reducing molecular oxygen to water

Production of laccases:

- * 7-days fermentation of *Trametes versicolor*
- (culture collection of the unité de phytopathologie, INRA, Versailles)
- **★** Induction by 2,5-xylidine

Experimental filtration device

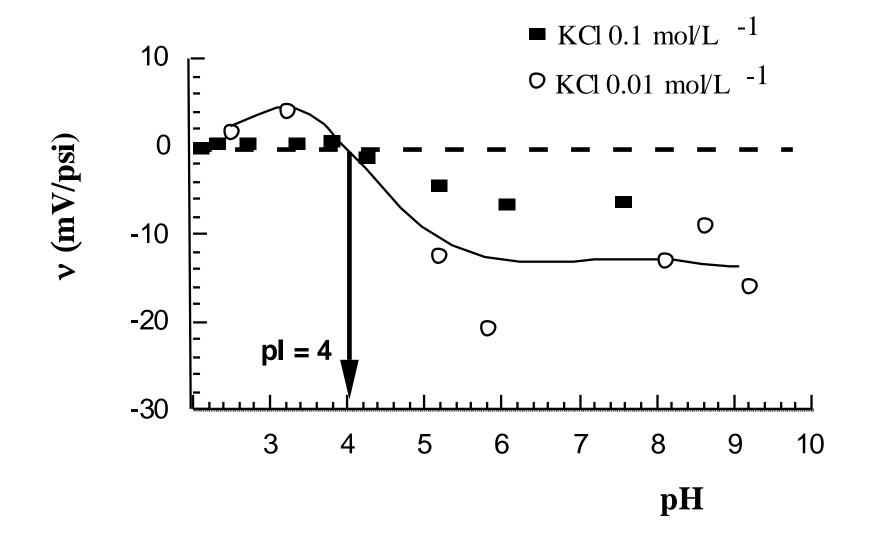


(a) reservoir (b) peristaltic pump (f)planar module $(\gamma 1, \gamma 2)$ saturated calomel electrodes (i) voltmeter (c) Vycor® fixing the junction potential

Used as shown for streaming potential measurements and without the potential measurements pieces for 2-HF filtration

RESULTS

Isoelectric point determination



Streaming potential measurements

Isoelectric point of the grafted membrane: $pI = 4 \oplus 0.5$ representative of the pI of the mixture of laccases

Wettability determination

Type of membrane	Contact angle, °
CH_3 O CH_3	0 °
Hydrophilic PVDF	
H_{1} H_{2} H_{2} H_{2}	123°
Hydrazine activated PVDF membrane	
CH ₃ O Laccase	0°
Laccase covalently grafted PVDF membrane	
Hydrazine activated PVDF membrane with adsorbed laccase	60 °

Sessile drop contact angle technique

M Covalent coupling of laccase to the membrane restores its hydrophilicity more efficiently than adsorption

ration, g/l 0,08 concen 0,06 0,02 250 100 150 200 **50** Filtration duration, minutes

→ 10 U - 50 U - 4,4 U

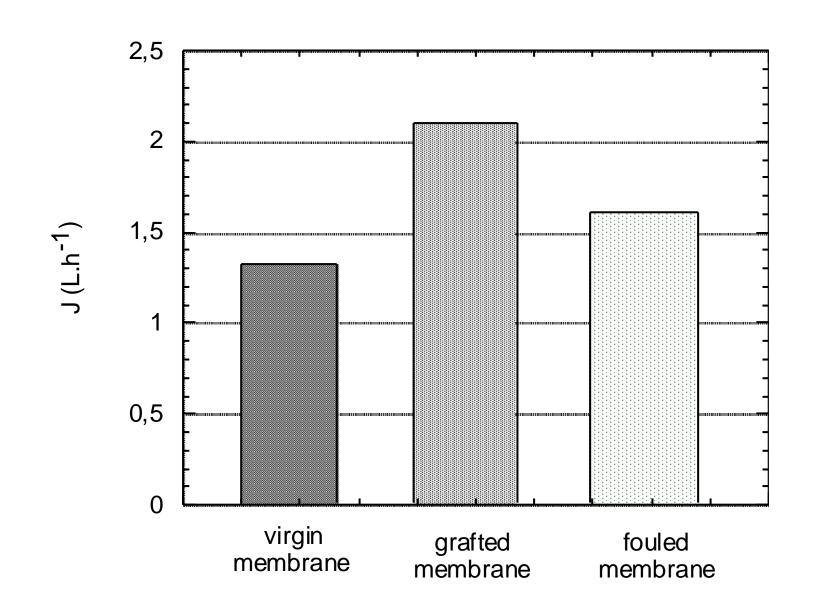
M Removal efficiency of 2-HF

in the filtrate increases with the amount of grafted laccases

Filtration module: Minitan S cm² planar membranes grafted with 4.4, 10, 50 and 220 laccase activity units. 2-HF initial concentration 0.1ss g/L in acetate buffer 0.05 M, pH 5 T=22°C.

Permeability measurements

Laccase degradation efficiency



M Laccase coating increases the flux through the membrane M The precipitation of the 2-HF degradation products does not induces significative fouling

Effective membrane surface: 60 cm² Grafted membrane with 50 laccase activity units Fouled membrane after treatment of 200 mL of 0.1 g/L 2-HF

Acknowledgement: we thank J.M. Siffre (ENSCP, Laboratoire de Physicochimie des Surfaces Métalliques) for performing the XPS measurements