



Contribution ID : 51

Type : **Poster**

Synchrotron radiation as a tool to determine chemical structure modification of wheat straw (*Triticum aestivum*) treated with an enzymatic extract obtained from Methylophilic Bacteria.

Wednesday, 12 August 2015 17:30 (1:00)

Abstract content

Vegetal biomass, also called lignocellulose, is distinguished by being refractory to physicochemical and enzymatic treatments [1]. Several factors are the responsible of this recalcitrance behavior as the crystallinity and composition of biomass (Cellulose, hemicellulose and lignin). The efficient enzymatic saccharification of cellulose at low cellulase (protein) loadings continues to be a challenge for commercialization of a process for bioconversion of lignocellulose to ethanol [2]. Cellulases are enzymes that hydrolyze the β -(1 \rightarrow 4)-linkages in cellulose. Though cellulase research has been concentrated mostly in fungi, actinomycetes, and protozoa, there is an increasing interest in cellulose production by bacteria [3-5]. In this context, pink pigmented facultative methylotrophs (PPFMs) are a physiologically interesting group of bacteria that are capable of growing on single carbon such as methanol and methylamine, as well as on C2, C3, and C4 compounds [6]. These bacteria can be found in plant phyllosphere and are not considered to be passive passengers on plant leaves, and is known that can stimulate seed germination and plant development. Though earlier studies indicate cellulase production by PPFMs [7] that can be used for transform cellulose in glucose in a bioethanol process. This work proposes the use μ -FTIR in order to understand the structural changes during enzymatic hydrolysis of lignocellulosic biomass that could lead to improved processes and cost reductions for bioethanol production.

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Summary

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Session Classification : Posters I