



Analysis *in silico* of polysaccharides degrading enzymes of *Aspergillus flavipes* FP-500 found in complex carbon sources.

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Introduction. Aspergillus is capable to produce a lot of carbohydrate degrading enzymes (CDE) which applied in a great variety of industrial processes such as feed and food production. Aspergillus flavipes posses interesting capabilities to produce those enzymes. However, its entire genome is not sequenced yet. However, the sequence of Aspergillus terreus genome is available: According to the literature the family flavipes and terreus are so similar that can be joined¹. Some of (CDE) have been reported to possess carbohydrate binding domains (CBD's); which are sequences for specific union to some polysaccharides².

The aim of this work is the identification to analyze the enzymes secreted by *A*. *flavipes* growing on complex carbon sources and analyze the probable CBD's in these enzymes by comparison with sequences of *Aspergillus terreus*, polysaccharide degrading enzymes.

Methods. Secreted proteins from *Aspergillus flavipes* FP-500 cultures with corn cobs as complex carbon source were analyzed by LC/MS/MS coupled system. Proteins identification and CBD's search were made using Conserved Domain Database tool (CDD) from NCBI or Uniprot databases.

Results. Proteins were identified as very similar to Aspergillus terreus proteins (table 1). Glucoamylase acts on starch hydrolysis, while α -galactosidase is involved in degradation of galacto(gluco)mannan³. On the other hand, cellobiohydrolase acts on the backbone of cellulose while xylanase and acetylxylan esterase belong to xylanolytic system. Many references indicate synergistic effect of different enzymatic systems; such as the combined action of Cellobiohydrolases with xylanase and acetylxylan esterase gives an important

increase of depolymerization of corn stover due to synergistic action of these enzymes⁴. We looked for CBD's and some of them were found using the CDD tool and Blastp protein sequences. Unlike the first three enzymes, xylanase A precursor and Acetylxylan esterase didn't show CBD's when were analyzed by CDD tool but shows identities near to 57 and 67% respectively to fungal cellulose binding domain (A1CCD3 _ASPCL Uniprot).The presence of CBD's on these enzymes suggests that may be necessary at least one of each with CBD's as a strategy used by the fungus to improve the polysaccharide hydrolysis.

 Table 1.Identified enzymes on Aspergillus flavipes FP-500 cultures on corn cobs with probable Carbohydrate Binding Domain sequences

Enzyme	NCBI and Uniprot Accesion	Score	e-value
Glucoamylase precursor	XP_001213553.1	49.4	4x10 ⁻⁶
	Q0CPK9	109	9x10 ⁻⁵
α-galactosidase	Q0CVX4.2	208	2X10 ⁻¹⁷
Cellobiohydrolase	AAW68437.2	69.8	10 ⁻²
	Q0CMT2	157	6X10 ⁻¹¹
Endo-1,4-beta-xylanase A	XP_001216082.1	53.7	2X10 ⁻⁷
precursor	Q0CFS3	119	6X10 ⁻⁸
Acetylxylan esterase	Q0CNM5	103	9X10 ⁻⁵

Conclusions. It's possible to identify CDE on proteins secreted by *A. flavipes* on corn cobs. Some of these enzymes likely to own CBM's which were possible to identify by comparison with *A. terreus* proteins.

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