



Abstracts

XVI<sup>th</sup> National Congress of Biochemistry



Ponta Delgada, São Miguel, Açores  
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- S11.P9** *Petrotoga mobilis* possesses two pathways for the synthesis of the rare compatible solute mannosyl- $\alpha$ -1,2-glucoacylglycerate  
Chantal Fernandes, Nuno Empadinhas, Vitor Mendes, Joana Costa, Helena Santos, Milton S. da Costa
- S11.P10** Identification of genes putatively involved in the synthesis of cyanobacterial exopolysaccharides  
Sara Pereira, Ângela Brito, Andrea Zille, Pedro Moradas-Ferreira, Paula Tamagnini
- S11.P11** A unique combination of genetic systems for the synthesis of trehalose in *Rubrobacter xylanophilus*: Characterization of the first bacterial TreT  
Ana Nobre, Susana Alarico, Nuno Empadinhas, Milton S. da Costa
- S11.P12** Genetic diversity among azorean isolates of *Steinernema carpocapsae*  
L. Esteves, R. Montiel, N. Simões
- S11.P13** Metalloproteases produced by the entomopathogenic bacteria *Photorhabdus* sp. Az29: expression studies and characterization of the prtA and prtS genes  
C. M. Cabral, R. Montiel, G. Nascimento, N. Simões
- S11.P14** The sugar phosphotransferase system (PTS) is involved in the uptake of the quorum sensing molecule AI-2 in *Escherichia coli*  
Antota Santos, Catarina S. Pereira, João C. Marques, Karina B. Xavier

## Symposium 12: BIOTECHNOLOGY

### Invited Speaker

- S12.IS** Biotechnological approaches to fruit related allergens  
Margit Laimer

### Selected Communications

- S12.C1** A new analgesic peptide: From biophysical screening to in vivo tests  
Marta M. B. Ribeiro, M. Heras, E. Bardaj-, M. Pinto, I. Tavares, Miguel A.R.B. Castanho
- S12.C2** Molecular phytopathology: From plant pathogen diagnosis to the research on resistance mechanisms  
D. Mendonça, I. Foroni, M. S. Lopes, S. Leitão, A. da Câmara Machado
- S12.C3** Improvement of naringinase activity and stability via immobilization techniques  
M. I. Amaro, M. H. Ribeiro
- S12.C4** HE-TAIL PCR, a powerful tool for identification of *Trichoderma harzianum lip1* gene  
L. Jorge, F. González, E. Monte, A. Choupina

### Poster Presentations

- S12.P1** Maintenance of catalytic activity in complex steroid bioconversion  
Marco P.C. Marques, Filipe Carvalho, Gabriel A. Monteiro, Joaquim M.S. Cabral, Pedro Fernandes
- S12.P2** Aminopeptidase N from *A. thaliana*: expression and characterization  
Rui Cruz, Rosário Faro, Euclides Pires, Carlos Faro, Paula Ver-ssimo
- S12.P3** Identification of cytotoxic proteins of *Bacillus thuringiensis*  
Ana Duarte, Vera Gouveia, Rafael Montiel, Carla Cabral, Luisa Oliveira, Nelson Simões
- S12.P4** scFv antibodies selected by phage display technology that target gp41 and gp120 are strong candidates to inhibit HIV-1 fusion process  
Andreia Couto, Pedro Lucas, João Gonçalves

**S12 C4 HE-TAIL PCR, a powerful tool for identification of *Trichoderma harzianum lip1* gene**

<sup>1</sup>Jorge, L., <sup>2</sup>González, F., <sup>3</sup>Monte, E. & <sup>1</sup>Choupina, A.

<sup>1</sup>Instituto Politécnico de Bragança – Escola Superior Agrária, Campus de Santa Apolónia, Apartado 1172, Bragança, 5301-855, Portugal.

<sup>2</sup>Newbiotechnic, S. A. (NBT). Parque Industrial de Bollullos A-49 (PIBO). 41110, Bollullos de la Mitación. Sevilla, Spain.

<sup>3</sup>Centro Hispano-Luso de Investigaciones Agrarias (CIALE), Departamento de Microbiología y Genética, Universidad de Salamanca. Edificio Departamental, lab 208, Plaza Doctores de la Reina s/n, 37007 Salamanca, Spain.

*Trichoderma harzianum* is a widespread soil fungus, known as a biocontrol agent against soilborne plant pathogens. Species of *Trichoderma* are commercially applied as biological control agents against plant fungal pathogens based on different mechanisms, such as antibiosis, competition for nutrients and mycoparasitism.

One of the mechanisms involved is the production of several lytic enzymes. In *T. harzianum* has been identified several glucanases, celulasas, chitinases and proteases, but nothing are already known about its lipolytic system.

In this work we described the identification of *lip1* from *T. harzianum*, a gene encoding the first extracellular triacylglycerol lipase known in this species. A cDNA library of *T. harzianum* CECT 2413 cloned in pBluescript SK+ was screened, and the expressed sequence tags (ESTs) with more similarity against lipase gene sequences in international databases selected and sequenced. Elucidation of complete gene nucleotide sequence of *lip1*, including 52 bp of the open reading frame at the N-terminal region and 693 bp of the promoter region were achieved by high-efficiency thermal asymmetric interlaced (HE-TAIL) PCR (Michiels *et al.*, 2003), which confirmed to be a powerful tool to identify flanking regions from short known sequences.

*Lip1* codifies a 558 amino acids protein, with 59322.6Da and a calculated global iso-electric point value of 4.56.

The GenBank/EMBL/DBJ accession number for the sequence reported in this paper is AM18087. This work was supported by COMBATINTA/SP2.P11/02 Interreg IIIA (PC007) - 220201