

## Novelty Seeking - Characterization and evaluation of novel environmental microbial strains sampled from extreme habitats.



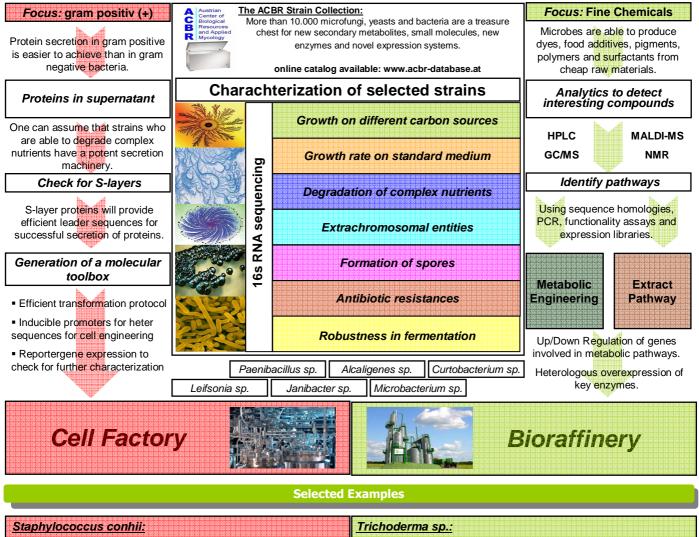
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## Background

Advances in molecular biology, genomics and bioinformatics, and cultivation technologies herald a new age of exploration of the microbial world. Prokaryotic life is able to adapt to environments that would prove fatal to most other life forms. The Austrian Center of Biological Resources and Applied Mycology (ACBR) harbors over 5000 uncharacterized bacterial and fungal isolates from extreme habitats. This projects aims at screening a sub-group of these, so far uncharacterized isolates, for their biotechnological potential. During this work we plan to establish a screening procedure that allows for the selection of strains that either a) secrete proteins efficiently in the extracellular space or b) that produce compounds with a biotechnological value. Additionally, we try to identify extrachromosomal entities like plasmids or phages that can be applied for the expression of recombinant proteins in the corresponding bacterial species and that allow for the generation of a molecular biology toolbox to modify these strains.

## Establish a screening procedure

To date most recombiannt proteins produced in bacteria are based on Escherichia coli expression systems. Due to limitations in secretion and folding capacity alternative prokaryotic systems are under investigation, e.g. Corynebacterium sp. (Srivastava and Deb 2005), Streptomyces sp. (Lussier et al. 2010), Bacillus sp. (Chan et al. 2002), Lactococcus sp. (Miyoshi et al. 2002), Lactobacillus sp. (Savijoki et al. 1997). However, the world of microorganisms comproses a vast diversity of bacteria.



Sequencing currently in progress, Secretes a number of proteins that have to be characterized production of exopolysaccharide and anti-microbials	posses a mid to high copy number plasmid,	production of coconut-aroma, purity analyzed by HPLC
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References Chan A, Chan M, Lo H, Leung Y, Lim B. 2002. A dual protein expression system in Bacillus sublits. Protein Expr Purif 26(3):337-42. Lussier F, Dens F, Shareck F. 2010. Adaptation of the highly productive 17 expression system to Streptomyces lividans. Appl Environ Microbiol 76(3):967-70. Miyoshi A. Poquet I, Azevedo V, Commissaire J, Bernudez-Humana L, Domakova E, Le Lioir Y, Oliveira S, Gruss A, Langela P. 2002. Controlled production of stable heterologous proteins in Lactococcus lactis. Appl Environ Microbiol 68(6):3141-6. Savijoti K, Kahala M, Haha A. 1937. High level heterologous protein production in Lactococcus and Lactobacillus using a new secretion system based on the Lactobacillus brevis S-layer signals. Gene 186(2):255-62. Srivastava P, Deb J. 2005. Gene expression systems in corynebacteria. Protein Expr Purif 40(2):221-9.