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# Identification of nitrifying bacteria contained in a commercial inoculant using molecular biology techniques

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# Identification of Nitrifying Bacteria Contained in a Commercial Inoculant Using Molecular Biology Techniques



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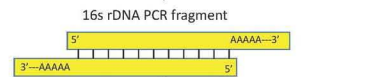
## Introduction

Nitrifying bacteria play an important role in the aquatic and terrestrial nitrogen cycle. Nitrification, one of the processes of the nitrogen cycle, refers to the oxidation of ammonia to nitrate. This process requires two types of chemoautotrophic bacteria: ammonia-oxidizing bacteria (AOB) and nitrite-oxidizing bacteria (NOB). These bacteria are essential as they supply nitrate for the growth of plants and aquatic organisms.

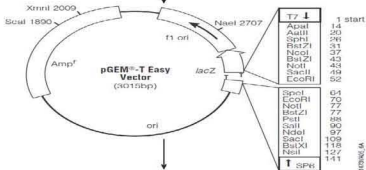
Current applications of nitrifiers include: inoculants for aquaria, biofertilizers, and nitrogen removal in wastewater treatment plants. Previous studies have shown that Fritz-zyme Turbostart 700, a commercial freshwater inoculant, has been successfully used in a semi-hydroponic system, i.e., zeoionics. In our laboratory, preliminary data have shown that Fritz-zyme contains more than the specific nitrifying bacteria. In order to determine an optimal consortium for zeoionics, it is necessary that we know exactly what bacteria are present. Using 16s rDNA universal primers and pGEM®-T Easy Vector Cloning Kit (Promega), we amplified the 16s rDNA genes from Fritz-zyme and cloned them into the pGEM®-T Easy *E. coli* vector plasmid. The cloned plasmids were transformed into competent *E. coli* cells and sequenced to identify the bacteria present in each sample. In this study, we determined whether the current enrichment techniques being used are sufficient to eliminate the heterotrophic and spore-forming bacteria present in Fritz-zyme.

## Materials and Methods

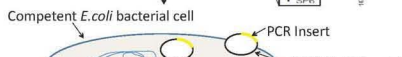
### Whole-cell PCR using universal and specific 16s rDNA primers



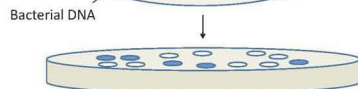
Whole-Cell PCR followed by PCR Purification using QIAquick PCR Purification Kit (Qiagen)



TA Cloning via Ligation using pGEM-T Easy Vector Kit (Promega)



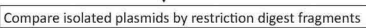
Transformation using Monserate MON1 competent cells



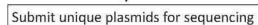
Blue/white colony screen on Luria Bertani agar containing Ampicillin, X-Gal and IPTG



Plasmid Prep using Qiagen Mini-Prep Kit



Restriction Digest using SAC I



Sequencing using primers T7 and SP6 at UNLV Genomics Center

## Materials and Methods (continued)

Primer	5'→3'	Specificity
27f	AGAGTTTGATCCTGGCTCAG	Bacterial 16s rDNA gene
1492r	ACGGTACCTGTACGACTT	Bacterial 16s rDNA gene
EUB338f	ACTCCTACGGGAGGACG	Bacterial 16s rDNA gene
Nso1225r	CGCCAATTGATTACGTGTGA	AOB 16s rDNA gene
NIT3r	CCTGTGCTCATGCTCCG	Nitrobacter 16s rDNA gene
Ntspa685r	CGGGAATCCGCGCTC	Nitrospira 16s rDNA gene

## Results

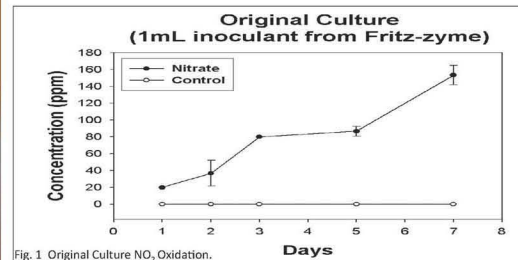


Fig. 1 Original Culture NO<sub>2</sub> Oxidation.

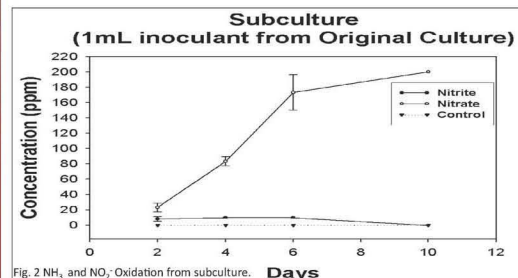


Fig. 2 NH<sub>3</sub> and NO<sub>2</sub> Oxidation from subculture.

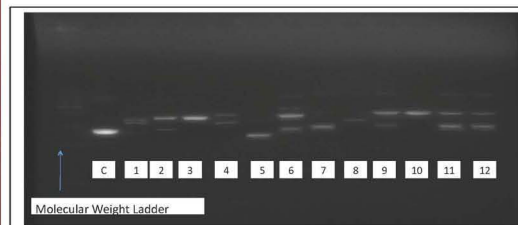


Fig. 3 Agarose Gel showing Cut Hybrid Plasmids using SAC I Restriction Endonuclease

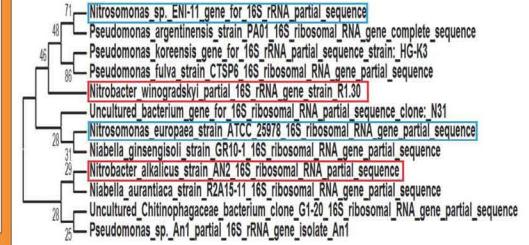


Fig. 4 Phylogenetic tree of cloned inserts created in MEGA 5.0 using results from BLASTn from NCBI Database. Contigs were created in DNA Baser Software. Values next to nodes represent percentage of replicate trees in which the associated taxa clustered together in the bootstrap test (500 replicates). Blue boxes represent Ammonia-oxidizing bacteria. Red boxes represent Nitrite-oxidizing bacteria.

## Conclusion

1. We confirmed the activity of nitrifying bacteria based on NH<sub>3</sub> and NO<sub>2</sub> oxidation using test strips.
2. Sequencing data showed the presence of Ammonia-oxidizing and Nitrite-Oxidizing bacteria in Fritz-zyme.
3. Sequencing data also showed the presence of non-nitrifying bacteria from the genera *Pseudomonas* and *Niabella*, which could indicate the presence of denitrifying bacteria as well as nitrifying bacteria.

## Future Research

- Measure oxidation of NH<sub>3</sub> and NO<sub>2</sub> using Ion-Selective Electrodes to gain a more accurate measurement of oxidation.
- Find Primers capable of amplifying 16s rDNA from sub-culture samples. Possible candidates include EUB338f and EUB338r along with specific 16s rDNA primers.
- Determine if sub-culturing techniques are suitable for isolating pure nitrifiers.

## References

Burrell, P.C., Phalen, C.M., and Hovanec, T.A. 2001. Identification of Bacteria Responsible for Ammonia Oxidation in Freshwater Aquaria. *Appl. Environ. Microbiol.* 67: 5791-5800.

Liesack, W., Weyland, H., and Stackebrandt, E. 1991. Potential Risks of Gene Amplification by PCR as Determined by 16s rDNA of a Mixed-Culture of Strict Barophilic Bacteria. *Microb Ecol.* 21: 191-198.

Regan, J.M., Harrington, G.W., and Noguera, D.R. 2002. Ammonia- and Nitrite-Oxidizing Bacterial Communities in a Pilot-Scale Chloraminated Drinking Water Distribution System. *Appl. Environ. Microbiol.* 68: 73-81.

## Acknowledgements

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