

## ABSTRACT

### *MYCOBACTERIUM SMEGMATIS* MC<sup>2</sup> 155 *fbiC* AND MSMEG\_2392 ARE INVOLVED IN TRIPHENYLMETHANE DYE DEGRADATION AND COENZYME F<sub>420</sub> BIOSYNTHESIS

Triphenylmethane dyes are carcinogenic and widely used in the aquaculture and textile industries. Bioremediation, the use of microorganisms to degrade xenobiotic compounds, may be a more efficient alternative to conventional treatment methods for dye contaminated waste. To identify genes involved in triphenylmethane dye decolorization by mycobacteria, a transposon mutant library of *Mycobacterium smegmatis* mc<sup>2</sup> 155 was created and screened for mutants unable to decolorize the triphenylmethane dye Malachite Green. One gene identified was *fbiC*, which is essential for the biosynthesis of the electron carrier, coenzyme F<sub>420</sub>. Also identified was MSMEG\_2392, belonging to a superfamily without annotated function. High Pressure Liquid Chromatography revealed that F<sub>420</sub> was absent in both mutant strains, indicating that, like *fbiC*, MSMEG\_2392 is required for the biosynthesis of coenzyme F<sub>420</sub> and this cofactor is likely the electron donor for the reduction of Malachite Green. This is the first report of this coenzyme in dye decolorization.

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