COMETABOLIC BIODEGRADATION OF TCE

BY BENZENE UTILIZING CONSORTIUM

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Introduction. Chlorinated pollutants including TCE are used in many industrial clean systems. TCE cause environmental pollution and a potent carcinogen. Aerobic degradation of TCE occurs through a process termed co-metabolism. Consortium B1 can degrade TCE by utilizing benzene as a substrate. Benzene was degraded, using benzene oxygenase (Bed), which catalyzes the oxidation of benzene to catechol. Bed gene consisted with BedA, BedB, BedC1 and BedC2. In this study, we want know interrelation of TCE degradation rate and gene expression. Induced overexpression of Bed gene, expected more high efficiency for benzene and TCE degradation rate.

Methods. Benzene and TCE degradation rate was analyzed using headspace method on GC-FID (Gas Chromatography-Flame Ionization Detector). Injected TCE and benzene, initial concentration of TCE was 1, 10 and 20 mg/L, benzene was 30 mg/L as substrate. And analyzed for gene expression, used Real-Time PCR method. For checking location of *Bed* gene which secreted enzyme and endo-enzyme. According to TCE and benzene degradation rate, checked expression of gene.

Results. In this study, consortium B1 degraded 1, 10 and 20 mg/L of TCE within 6, 74 and 111 hours, respectively.



Fig. 1. 1 mg/L of TCE degraded within 6 hours by B1.



Fig. 2. 10 mg/L of TCE degraded within 74 hours by B1.



Fig.3. 20 mg/L of TCE degraded within 111 hours by B1.

Injected benzene after 6 hours, *Bed* gene was expressed maximum. At D.O value lower than 0.5 mg/L inside the media, gene expression decreased.

Conclusions. *Bed* gene expression was increased on aerobic condition actively, and secreted to media. *Bed* gene was degraded benzene to catechol, catechol form detached from chloride of TCE, easily. Thus, according to overexpression of *Bed* gene, expected more efficiency for TCE degradation rate.

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