

# NEW SOURCE AND OPTIMIZED CONDITIONS FOR HYDROGEN PRODUCTION BY *RHODOBACTER SPHAEROIDES* DURING PHOTO-FERMENTATION

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Photosynthetic bacteria are often presented as perspective organisms for the biohydrogen (H<sub>2</sub>) production [1]. Purple non-sulfur bacteria *Rhodobacter sphaeroides*, isolated from Arzni (sodium-chloride type with pH 6.3–6.6) and Jermuk (sulphate-chloride type with pH 5.9–7.6) mineral springs in Armenian mountains [2], can produce H<sub>2</sub> in various conditions upon illumination. H<sub>2</sub> production by *R. sphaeroides* during “photo-fermentation” depends on different factors. of carbon (succinate, malate) and nitrogen (various amino acids, yeast extract) sources Yeast extract enhanced ~6 fold H<sub>2</sub> production (compared to glutamate) [3]. Various metal ions such as Fe<sup>2+</sup>, Mo<sup>6+</sup> and Mg<sup>2+</sup> had stimulating effect on H<sub>2</sub> production and F<sub>0</sub>F<sub>1</sub>-ATPase activity [4]. Regulations of H<sub>2</sub> production by external reducers and oxidizers and light-dark duration alternations have been also shown [5,6]. To develop biotechnology enhancing H<sub>2</sub> production cheap source and optimized conditions are of significance.

Using ethanol fermentation waste, distillers grains (DG), for H<sub>2</sub> production has the advantages of recycling wastes and protecting environment. H<sub>2</sub> production by pure and co-cultures of photo- (*R. sphaeroides*) and dark-fermentative (*Escherichia coli*) bacteria using DG has been investigated. During growth on diluted DG media at pH 7.0 H<sub>2</sub> production by pure and co-cultures was started at 24 h growth, whereas H<sub>2</sub> production by *R. sphaeroides* cells, grown on Ormerod medium, was detected after 48 h. Moreover, the co-culture has produced significantly more (~2-3-folds) H<sub>2</sub> from 2-5-folds diluted DG during 96 h growth compared with pure culture. This could be related with formation of reductive power and ATP synthesis during photo-fermentation [5,6]. The results obtained indicate that DG can be used as effective and valuable substrate in H<sub>2</sub> production by *R. sphaeroides* and other bacteria.

- References: 1. Trchounian A. (2015) *Crit Rev Biotechnol* 35:103–13.  
2. Paronyan A. (2002) *Appl Biochem Microbiol* 38:53-58.  
3. Hakobyan L., Gabrielyan L., Trchounian A. (2012) *Int J Hydrogen Energy* 37:6519–26.  
4. Hakobyan L., Gabrielyan L., Trchounian A. (2012) *Int J Hydrogen Energy* 37:17794–800.  
5. Gabrielyan L, Sargsyan H, Hakobyan L, Trchounian A. (2014) *Appl Energy* 131:20–5.  
6. Sargsyan H., Gabrielyan L., Hakobyan L., Trchounian A. (2015) *Int J Hydrogen Energy* 40:4084-91.