Accepted Manuscript

Photoelectrochemical and photocatalytic systems based on titanates for hydrogen peroxide formation

Tomasz Baran, Szymon Wojtyła, Alberto Vertova, Alessandro Minguzzi, Sandra Rondinini

PII: S1572-6657(17)30472-1

DOI: doi: 10.1016/j.jelechem.2017.06.044

Reference: JEAC 3375

To appear in: Journal of Electroanalytical Chemistry

Received date: 30 January 2017 Revised date: 20 June 2017 Accepted date: 21 June 2017

Please cite this article as: Tomasz Baran, Szymon Wojtyła, Alberto Vertova, Alessandro Minguzzi, Sandra Rondinini, Photoelectrochemical and photocatalytic systems based on titanates for hydrogen peroxide formation, *Journal of Electroanalytical Chemistry* (2017), doi: 10.1016/j.jelechem.2017.06.044

This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting proof before it is published in its final form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.



ACCEPTED MANUSCRIPT

Photoelectrochemical and photocatalytic systems based on titanates for hydrogen peroxide formation

Tomasz Baran, a Szymon Wojtyła, b Alberto Vertova, Alessandro Minguzzi, a Sandra Rondininia, *

Abstract

The development of efficient, sustainable, cheap and safe methods for the synthesis of hydrogen peroxide is a very important challenge from the environmental viewpoint, since H_2O_2 can be used as a solar fuel or for degradation of organic, inorganic and biologic pollutants from wastewater. Efficient photocatalytic or photoelectrochemical formation of hydrogen peroxide from water and/or oxygen in an acidic medium has been obtained using nickel or cobalt titanates under visible light illumination. The concentration of H_2O_2 achieved in the photoreaction reached as high as 2.5 mM when nickel titanate was irradiated for 1 h in oxygen saturated suspension. In such conditions, NiTiO₃ provide active sides for both, oxygen reduction and water oxidation leading to H_2O_2 formation.

Introduction

Conversion of solar energy into easily storable chemicals has recently attracted considerable attention as strategic technology to provide a clean energy. Photoelectrochemical hydrogen production from water or photocatalytic carbon dioxide reduction to methane are considered as some of the most promising approaches for solar energy utilization [1,2]. The use of hydrogen peroxide (H_2O_2) as an alternative solar fuel is also very promising since the reduction of oxygen to H_2O_2 is thermodynamically more favorable than reduction of H^+ towards formation of hydrogen [3]. Moreover, H_2O_2 presents several great advantages in comparison with H_2 as a fuel [4]. It is liquid at room temperature and atmospheric pressure thus it has higher energy density and can be stored, transported and used without a high-pressure tank. The theoretical open circuit potential of the H_2O_2 fuel cell is 1.09 V [5], by consuming hydrogen peroxide as both an oxidant at an anode and a reductant at a cathode [6] according to:

$$H_2O_2 \rightarrow O_2 + 2H^+ + 2e^-$$
 (0.68 V vs NHE) (1)

$$H_2O_2 + 2H^+ + 2e^- \rightarrow H_2O (1.77 \text{ V vs NHE})$$
 (2)

The open circuit potential is thus just slightly lower than that of a hydrogen/oxygen fuel cell (1.23 V [6]).

Moreover, H_2O_2 is a clean oxidant, with water as the only byproduct. In industry, it is commonly used for pulp bleaching as well as for disinfection due to its antimicrobial activity [7].

^a Department of Chemistry, University of Milan, via Golgi 19, 20133 Milan, Italy

^b SajTom Light Future LTD, Wezerow, Poland

Download English Version:

https://daneshyari.com/en/article/6662308

Download Persian Version:

https://daneshyari.com/article/6662308

<u>Daneshyari.com</u>