University of Nebraska - Lincoln

DigitalCommons@University of Nebraska - Lincoln

Papers in Reaction Kinetics

Chemical and Biomolecular Engineering Research and Publications

11-1-2003

A Macrokinetic Study of the High-Temperature Solid-Phase **Titanium-Carbon Reaction**

Julia Y. Kostagorva Department of Chemical Engineering, University of Nenraska-Lincoln

Hendrik J. Vilioen University of Nebraska-Lincoln, hviljoen1@unl.edu

Alexander S. Shteinberg ALOFT corporation, Berkely, California, 94708.

Follow this and additional works at: https://digitalcommons.unl.edu/chemengreaction



Part of the Chemical Engineering Commons

Kostagorva, Julia Y.; Viljoen, Hendrik J.; and Shteinberg, Alexander S., "A Macrokinetic Study of the High-Temperature Solid-Phase Titanium-Carbon Reaction" (2003). Papers in Reaction Kinetics. 2. https://digitalcommons.unl.edu/chemengreaction/2

This Article is brought to you for free and open access by the Chemical and Biomolecular Engineering Research and Publications at DigitalCommons@University of Nebraska - Lincoln. It has been accepted for inclusion in Papers in Reaction Kinetics by an authorized administrator of DigitalCommons@University of Nebraska - Lincoln.

A Macrokinetic Study of the High-Temperature Solid-

Phase Titanium-Carbon Reaction

Julia Y. Kostogorova and Hendrik J. Viljoen

Department of Chemical Engineering, University of Nebraska, Lincoln, Nebraska 68588

Alexander S. Shteinberg

ALOFT Corporation, Berkeley, California 94708

An experimental method, electro thermal explosion (ETE), is used to measure the macro kinetic parameters of the high-temperature titanium/carbon reaction. Different stages of the reaction have been identified, but the focus of this study is on the reaction between solid titanium and solid carbon, i.e., prior to the melting of titanium. The reaction has high activation energy, and an electric current is used to heat the cylindrically shaped sample to a specified temperature. The current is shut off at a temperature below the melting point of titanium; any further temperature rise is only due to reaction. The output of the ETE equipment is temperature time data that can be processed to recover the kinetic parameters. The activation energy and preexponential factor of the reaction rate constant are calculated and comprise 214 kJ mol-1 and (6.2 (1.5) 107 s-1, respectively. An important aspect of solid-phase reactions is the contact area between reactants. The contact area between titanium and carbon particles is calculated, and the reaction constant is

Published in Industrial and Engineering Chemistry Research 2003, Vol. 42 No 26, 6714-6719 © 2003 American Chemical Society:

Web Release Date: November 20, 2003

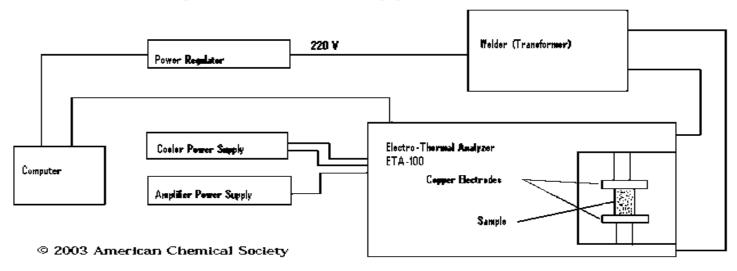
corrected for this effect.

The American Chemical Society allows the posting of only the title, abstract, tables, and figures from articles appearing in the Journal of American Chemical Society.

This article can be viewd at the publishers site: http://pubs.acs.org/cgi-bin/abstract.cgi/iecred/2003/42/i26/abs/ie030337k.html

DOI: 10.1021/ie030337k Copyright © 2003 American Chemical Society.

Figure 1. Schematic of ETE equipment.



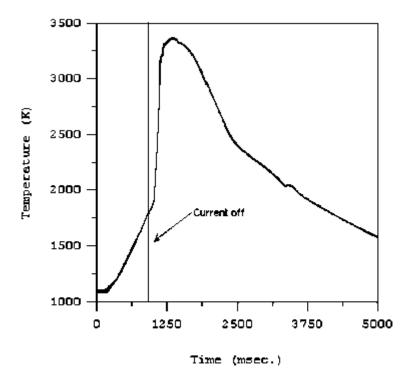
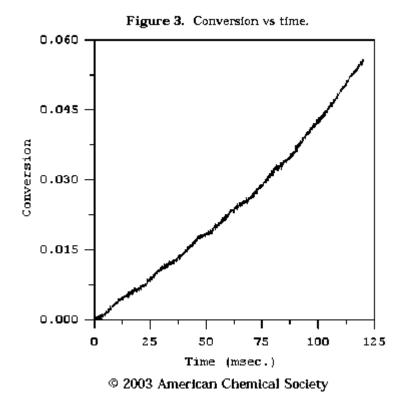
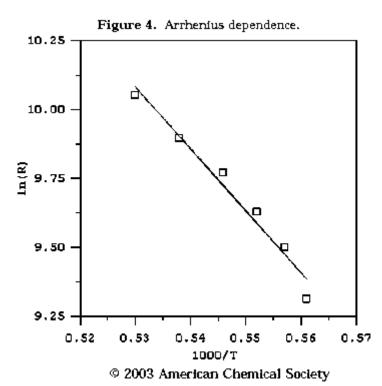


Figure 2. Experimental data of temperature (averaged over channels 9 and 10) vs time.

© 2003 American Chemical Society

A Macrokinetic Study of the High-Temperature Solid-Phase Titanium-Carbon Reaction Julia Y. Kostogorova, Hendrik J. Viljoen, Alexander S. Shteinberg





A Macrokinetic Study of the High-Temperature Solid-Phase Titanium-Carbon Reaction Julia Y. Kostogorova, Hendrik J. Viljoen, Alexander S. Shteinberg

Figure 5. Adsorption isotherm for titanium powder, 44 μ m.

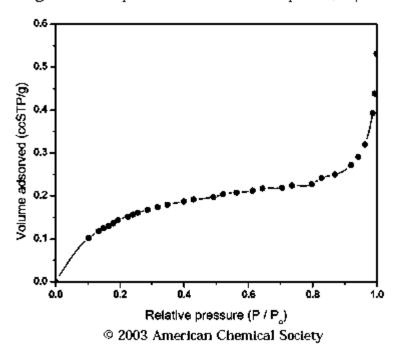


Figure 6. Adsorption isotherm for carbon powder, 44 μ m.

