

MICROFLUIDIC CHIP FOR THE ANALYSIS OF VERSATILE BIOCHEMICAL REACTIONS

Chang-Soo Lee^{1*}, Sang-Ho Lee^{2,3}, Yong-Kweon Kim³, and Byung-Gee Kim¹

¹School of Chemical and Biological Engineering, ³School of Electrical Engineering
and Computer Science, Seoul National University, Seoul, Korea

*Present address: Department of Chemical Engineering, Chungnam National
University, Daejeon, Korea California NanoSystems Institute, University of
California, Santa Barbara, USA

TEL: +82-42-821-5896, FAX: +82-42-822-8995

The driving force for the successful miniaturization of microfluidic systems in biotechnology and bioanalysis is not only the reduction of sample volumes down to nano- and picoliter sizes but also leads to remarkably improved performance, such as higher separation efficiency, shorter analyzing times, and enhanced detection sensitivities. Integrated microfluidic devices containing pumps, valves, separation systems, and detectors follow the concept of the so-called total (bio-chemical) analysis systems, TAS. Ideally, each step of the analysis is realized on an integrated device. The steps include sample injection, transportation, mixing, (bio-) chemical reactions, separation, and their analytical identification. Especially, Interfacing microfluidic chips with external analytical instruments has been applied to most lab-on-a-chip application for parallel and multiple analyses since sensitive on-chip detection still remains challenge in analyzing extremely small volume and low concentration [1]. Mass spectrometry (MS) has become one of the most commonly used analysis methods for biomolecules due to its inherent simplicity, low sample consumption and high sensitivity [2].

In this study, we will present this concept as fully integrated biochemical microfluidic device, namely lab on a chip, for fluidic handling as well as reaction, separation, and detection.

References

1. M. Brivio, R. H. Fokkens, W. Verboom, D. N. Reinhoudt, N. R. Tas, M. Geobloded and A. Van den Berg, *Anal. Chem.*,**74**, pp. 39723976, (2002).
2. A. J. de Mello, *Lab on Chip* **1**, pp. 7N-12N, (2001).