## A Method for Comprehensive and Quantitative Proteome Analysis Using 1–D and 2–D DIGE Combined with MicroSol IEF Prefractionation

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Current methods for quantitatively comparing complex protein profiles such as twodimensional gel electrophoresis (2–DE), 2–D differential in–gel electrophoresis (DIGE), and liquid chromatography (LC)–mass spectrometry (MS) have still limited resolution and dynamic ranges. In this study, we introduce a new method of 1–D/2–D DIGE combined with microscale solution isoelectric focusing (MicroSol–IEF) fractionation. The method has advantages over sample prefractionation and conventional 2–D DIGE technique including high reproducibility, high resolution, and a much wide linear dynamic ranges for detection. To illustrate its utility, this method was applied to analysis of human melanoma cell lines and mouse lung tissue extracts. Thus it is a powerful method for more comprehensive and quantitative comparison of protein profiles of very complex proteomes. [This work was supported by Basic Science Research Program through the National Research Foundation of Korea(NRF) funded by the Ministry of Education, Science and Technology(2010–0008826)]