

UNDERSTANDING THE LIMITS OF IONIC LIQUID BASED SUPERCAPACITORS

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Ionic liquids are versatile solvents and electrolytes for many electrochemical applications, such as secondary batteries, capacitors, solar cells and electrochemical deposition. In order to rationally design and further develop such systems, however, a detailed understanding of the processes that govern the interfacial processes and stability of ionic liquid interfaces is required [1,2]. It is thus the focus of this work to explore the limits and provide new methods to increase both the power and energy characteristics of supercapacitors by applying different ionic liquids as the electrolytes for high specific surface area carbon-based supercapacitors [3].

A novel technique of *in situ* infrared absorption spectroscopy for the study of thin film graphite electrodes has been developed [1] and is applied for the study of the graphite | 1-ethyl-3-methylimidazolium tetrafluoroborate (EMImBF₄) ionic liquid interface as a model supercapacitor and graphite | polydicyanamide interface as a dielectric capacitor. The infrared absorption results show that, unlike previous theoretical considerations of the electrical double layer between semimetals and electrolytes, the strong interaction between the electronic and ionic part of the electrical double layer is of fundamental importance for the description of semimetal interfaces. Together with the data provided by *in situ* electroreflectance technique of the electronic and energetic structure of the graphite electrode and differential capacitance-potential dependence of the graphite | EMImBF₄ system [1,2], it is demonstrated that the screening of excess surface charge by the ionic liquid is strongly correlated with the potential dependent electronic properties of the electrode surface states, thus limiting the amount of charge stored.

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References

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