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G. Inzelt

Editor F. Scholz

Conducting Polymers

A New Era in Electrochemistry



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Conducting Polymers

Elements consisted at first of certain small and primary Coalitions of minute Particles of matter into Corpuscles very numerous and very like each other. It will not be absurd to conceive, that such primary Clusters may be of far more sorts than three or five; and consequently, that we need not suppose, that each of the compound Bodies we are treating of, there should be found just three sorts of such primitive Coalitions.

—Robert Boyle: The Sceptical Chymist, Oxford, 1680.

Preface

Conducting polymers have permeated many fields of electrochemical research. Like metals and alloys, inorganic semiconductors, molecular and electrolyte solutions, and inorganic electroactive solids, they comprise a group of compounds and materials with very specific properties; indeed, there is now a research field focusing on the electrochemistry of conducting polymers. Conducting polymers possess similarities from an electrochemical point of view to all of the other compounds and materials mentioned above, making them a highly fascinating research topic. Furthermore, such research has led to numerous new applications, ranging from corrosion protection to analysis. There are a huge number of electrochemical papers on conducting polymers, and a good number of books on this topic too. However, the editor of the present series of *Monographs in Electrochemistry* noted that there was no modern monograph on the market in which the electrochemistry of conducting polymers is treated with the appropriate balance of completeness and selectivity. Such a monograph should be written by an active electrochemist who is experienced in the field of conducting polymers, and who possesses a solid knowledge of the theoretical foundations of electrochemistry. Therefore, I am very happy that György Inzelt from the Eötvös Loránd University in Budapest, Hungary, agreed to write this monograph. I hope that graduate students in electrochemistry, the chemistry and physics of materials, industrial chemists, and researchers at universities and industry alike will find this monograph enjoyable and stimulating, as well as helpful for their work.

March 2008

Fritz Scholz
Editor of the series *Monographs in Electrochemistry*

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Chapter 1

Introduction

Polymers have long been thought of and applied as insulators. Indeed, not so long ago, any electrical conduction in polymers—mostly due to loosely bound ions—was generally regarded as an undesirable phenomenon. Although the ionic conductivity of polymer electrolytes (macromolecular solvents containing low-molar-mass ions) and polyelectrolytes (macromolecules containing ionizable groups) have been widely utilized in electrochemical systems over the last few decades (e.g., in power sources, sensors, and the development of all-solid-state electrochemical devices), the emergence of electronically conducting polymers has resulted in a paradigmatic change in our thinking and has opened up new vistas in chemistry and physics [1].

This story began in the 1970s, when, somewhat surprisingly, a new class of polymers possessing high electronic conductivity (*electronically conducting polymers*) in the partially oxidized (or, less frequently, in the reduced) state was discovered. Three collaborating scientists, Alan J. Heeger, Alan G. MacDiarmid and Hideki Shirakawa, played a major role in this breakthrough, and they received the Nobel Prize in Chemistry in 2000 “for the discovery and development of electronically conductive polymers” [2, 4–8].

As in many other cases in the history of science, there were several precursors to this discovery, including theoretical predictions made by physicists and quantum chemists, and different conducting polymers that had already been prepared. For instance, as early as 1862, Henry Letheby prepared polyaniline by the anodic oxidation of aniline, which was conductive and showed electrochromic behavior [9].

Nevertheless, the preparation of this polyacetylene by Shirakawa and coworkers and the discovery of the large increase in its conductivity after “doping” by the group led by MacDiarmid and Heeger actually launched this new field of research.

Electrochemistry has played a significant role in the preparation and characterization of these novel materials. Electrochemical techniques are especially well-suited to the controlled synthesis of these compounds and for the tuning of a well-defined oxidation state.

The preparation, characterization and application of electrochemically active, electronically conducting polymeric systems are still at the foreground of research activity in electrochemistry. There are at least two major reasons for this intense

interest. First is the intellectual curiosity of scientists, which focuses on understanding the behavior of these systems, in particular on the mechanism of charge transfer and on charge transport processes that occur during redox reactions of conducting polymeric materials. Second is the wide range of promising applications of these compounds in the fields of energy storage, electrocatalysis, organic electrochemistry, bioelectrochemistry, photoelectrochemistry, electroanalysis, sensors, electrochromic displays, microsystem technologies, electronic devices, microwave screening and corrosion protection, etc.

Many excellent monographs on and reviews of the knowledge accumulated regarding the development of conducting polymers, polymer film electrodes and their applications have been published, e.g., [1, 10–53]. Beside these comprehensive works, surveys of specific groups of polymers [40, 49], methods of characterization [50–54] or areas of application [18, 21, 34, 36–38, 47, 48] have also appeared. These novel materials with interesting and unanticipated properties have attracted workers across the scientific community, including polymer and synthetic chemists [13, 14, 22, 23], material scientists [14, 20, 31, 32], organic chemists [17], analytical chemists [16, 21, 36, 45], as well as theoretical and experimental physicists [8, 31, 32].

After 30 years of research in the field, the fundamental nature of charge propagation is now in general understood; i.e., the transport of electrons can be assumed to occur via an electron exchange reaction (electron hopping) between neighboring redox sites in redox polymers, and by the movement of delocalized electrons through conjugated systems in the case of so-called intrinsically conducting polymers (e.g., polyaniline, polypyrrole). (In fact, several conduction mechanisms, such as variable-range electron hopping and fluctuation-induced tunneling, have been considered.) In almost every case, the charge is also carried by the movement of electroinactive ions during electrolysis; in other words, these materials constitute mixed conductors. Owing to the diversity and complexity of these systems—just consider the chemical changes (dimerization, cross-linking, ion-pair formation, etc.) and polymeric properties (chain and segmental motions, changes in the morphology, slow relaxation) associated with them, the discovery of each new system brings new problems to solve, and much more research is still needed to achieve a detailed understanding of all of the processes related to the dynamic and static properties of various interacting molecules confined in a polymer network.

Although the conductivity of these polymers is an interesting and an utilizable property in itself, their most important feature is the variability of their conductivity, i.e., the ease with which the materials can be reversibly switched between their insulating and conducting forms.

In this work, the topics that are presently of greatest interest in this field, along with those that may be of much interest in the future, are discussed. Some of the most important experiences, existing models and theories are outlined, and the monograph also draws attention to unsolved problems. Some chapters are also devoted to the most typical representatives of this group of materials and the most important techniques used for the characterization of these systems. Last but not least, abundant instances of the applications of conducting polymers are described.

The examples presented and the references recommended herein have been selected from more than ten thousand research papers, with emphasis placed on both classical and recent works. It is hoped that this monograph will be helpful to colleagues—electrochemists and non-electrochemists alike—who are interested in this swiftly developing field of science.

Considering the rapidly increasing number of applications of polymers in electrochemical cells, it can be declared that electrochemistry is currently moving out of the Bronze Age (i.e., typically using metals) and into the era of polymers.

Lecturi salutem!

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Chapter 2

Classification of Electrochemically Active Polymers

Electrochemically active polymers can be classified into several categories based on the mode of charge propagation (note that insulating polymers are not considered here except for those with variable conductivity). The mode of charge propagation is linked to the chemical structure of the polymer. The two main categories are electron-conducting polymers and proton (ion)-conducting polymers. We will focus on electron-conducting polymers here.

We can also distinguish between two main classes of electron-conducting polymers based on the mode of electron transport: redox polymers and electronically conducting polymers.

In this chapter we provide examples of each type of electron-conducting polymers, listing some of the most typical and widely studied of these polymers, as well as several new and interesting representatives of this class of materials. Some sections are also devoted to combinations, such as electronically conducting polymers containing redox functionalities and copolymers. Composites are briefly discussed too.

2.1 Redox Polymers

Redox polymers contain electrostatically and spatially localized redox sites which can be oxidized or reduced, and the electrons are transported by an electron exchange reaction (electron hopping) between neighboring redox sites if the segmental motions enable this. Redox polymers can be divided into several subclasses:

- Polymers that contain covalently attached redox sites, either built into the chain, or as pendant groups; the redox centers are mostly organic or organometallic molecules
- Ion-exchange polymeric systems (polyelectrolytes) where the redox active ions (mostly complex compounds) are held by electrostatic binding.