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STRUCTURE AND POLYMORPHISM OF THE HYDROCARBON CHAINS OF LIPIDS : A STUDY OF LECITHIN-WATER PHASES (*)

A. TARDIEU, V. LUZZATI

Centre de Génétique Moléculaire, CNRS
91190 Gif-sur-Yvette, France

and

F. C. REMAN †

Biochemisch Laboratorium der Rijks Universiteit
Vondelaan 26, Utrecht, Holland

Abstract. — This work describes the structure of a variety of lecithin-water phases observed below the « melting » temperature of the hydrocarbon chains, with special emphasis on the conformation of the chains. The lecithins studied in this work are the homologous series dioctanoyl to distearoyl, 2-decanoyl-1-stearoyl, and a preparation from hen eggs. The hydrocarbon chains are found to adopt a variety of conformations in additions to type α , the liquid-like organization observed above the melting temperature. Type β : the chains are stiff and parallel, oriented at right angles to the plane of the lamellae and packed with rotational disorder in a two-dimensional hexagonal lattice ($a \sim 4.85$ Å). Type β' : similar to β , but with the chains tilted with respect to the normal to the lamellae. Type δ : the chains are probably coiled into helices, whose axes are perpendicular to the plane of the polar groups and are packed with rotational disorder in a two-dimensional square lattice ($a \sim 4.80$ Å). α is the predominant conformation, common to most lipids in the presence of water and at sufficiently high temperature, and the one more relevant to membranes ; β is observed at lower temperatures in lipids whose chains are heterogeneous and in the presence of very small amounts of water ; β' is found in synthetic lecithins with identical chains, in the presence of variable amounts of water ; δ is observed in dry lecithins. A highly ordered *crystalline phase*, yet displaying rotational disorder of the chains, is observed in almost dry lecithins. Most of the phases are lamellar, and contain one lipid bilayer per repeat unit. Two phases display two-dimensional lattices : $P\delta$, formed by ribbon-like elements with the chains in the δ conformation ; $P\beta'$, formed by lamellae of type β' distorted by periodic ripples. The results emphasize the clear-cut difference between the liquid-like and the other types of partly ordered conformations, as well as the correlations which exist between the chemical composition and the structure of the lipids below the melting temperature of the chains.

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