gy (J < 0). This behaviour opposes entirely to the one of Dioxin possessing a combination of four π electron Spins with a singlet ground state (S=0) and a positive exchange energy (J > 0). Special binding complexes of Dioxin to XD* structures have been also observed with a negative exchange energy (J < 0). These [XD*. Dioxin] complexes appear with a very high probability, about 10^6 times of the one of the [Hemoglobin - Dioxin] complex in human blood and of the [Mn-Porphyrinoprotein - Dioxin] complex in human liver. The properties of these interaction complexes corresponding with the results achieved in some respective experimental clinical studies over a long period of time suggest that the specific natural XD* structures are able to conquer and arrest potently Dioxin molecules, i.e. to interact concurrently contrary to the toxicity of Dioxin, namely to its carcinogenic activity. This activity depends on many factors such as the concentration and the lifetime of the XD* state, the kinetic conditions of the surrounding and the interaction.

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Hydrochlorides of N-(1-Naphthyl)-N' -(3-ethylphenyl) guanidine (NEGH) and an N'-methyl analog aptiganel (CERESTAT®), which are N-methyl-D-aspartate ion channel blockers, have anomalous physical properties. The melting point of crystals of the salts differ by 70°, their densities differ by 0.12g/cm^3, and they exhibit significant differences in solubility. In general, high melting points suggest greater stability. Higher melting compounds of analogous composition might be expected to have higher density and lower solubility than the lower melting compounds. In this case the opposite relationships are observed. The higher melting salt is more soluble and less dense than the lower melting salt. Due to the addition of a methyl group in aptiganel there is a larger volume per molecule. However, the increased volume per molecule (66Å^3) is greater than might be expected. In think state NEGH has an extended conformation with the planar groups well away from each other. The methyl substitution in aptiganel appears to induce or stabilize a conformation in which the rings are oriented toward one another. Due to its extended conformation planar rings of NEGH are sufficiently exposed to permit stacking that contributes to closer packing of the molecules and higher density but not to a higher melting point. Although there were three NH...Cl hydrogen bonds in each structure, those in NEGH are engaged in forming dimers whereas in aptiganel one of the hydrogen bonds links the molecules into infinite chains. This difference in hydrogen bonding provides the explanation for the anomalous relationship between melting points and density and solubility. Upon initial melting only van der Waals forces must be overcome to melt NEGH crystals, while in aptiganel the crystals do not begin to melt until hydrogen bonds between adjacent molecules are broken. The broader melting point observed for NEGH crystals is due to the fact that initial melting involves breaking van der Waals contacts followed by the disruption of hydrogen bonds as dimers dissociate to form monomers.

1Registered trademark of Cambridge Neuroscience.

Chiral Compounds of Industrial Interest

MS11.08.01 CONFORMATIONAL STUDIES OF VITAMINS D DERIVATIVES. Kinga Suwinska and Andrzej Kutner, Institute of Physical Chemistry, Polish Academy of Sciences, Warsaw, Poland Pharmaceutical Research Institute, Warsaw, Poland

The fat-soluble vitamins D play essential role in stimulation of intestinal calcium transport, bone calcification, bone calcium mobilization, and phosphate transport as well as stimulation of cell differentiation and inhibition of cell proliferation. Vitamins D2 and D3 were found to exist in two almost equinucler equilibrium conformations in both liquid and solid states. The biologically active form of vitamin D3 1.25-hydroxylated derivative was found to exist in solid state as the beta conformer only [1]. To explain the influence of 1-hydroxy substituent on the conformation of the vitamins D the 1-hydroxy vitamin D2 and 1-hydroxy vitamin D3 was crystallized and the structures of those two compounds were solved. For both compounds only one conformer was found in the solid state. The results of crystal structure determinations were compared with the semi-empirical calculations and will be discussed.


MS11.08.02 STRUCTURAL ASPECTS OF DIASTEREOEMERIC DISCRIMINATION. Edward J. Valente, Christopher Miller, Gerard Ruggiero, Department of Chemistry, Mississippi College, Clinton, MS 39058, and Drake S. Eggleston, Department of Physical and Structural Chemistry, SmithKline Beecham, Co., King of Prussia, PA, 19401

Systems showing two kinds of diastereoelscutive solubility disparities are described. In the system of warfarin and its derivatives and (+)-quinidine, a pattern in resolution success (= formation of a separable salt) as a function of warfarin substituent position has been found. Warfarin with a single phenyl substituent in the 3- or 4'-positions and 2,3,4' -disubstituted warfarins usually form less-soluble salts between their (-)-(S)-isomers and (+)-quinidine; their diasteroemers are oils. The solid state structures of the less-soluble salts show that a single, strong N-H...O interaction linking warfarin enolate anions with protonated quinuclidine of quinidine. Packing motifs are remarkably similar over the range of substituents demonstrating that how a dominant interion interaction may serve as the basis for successful resolutions within a series of related substances. In the system of mandelic acid and its phenyl substituted derivatives and (+)-ephedrine, both less-soluble and more soluble solids can be isolated from 95% ethanol. The uniformly unsolvated salts vary from little to strong solubility disparities in aqueous ethanol, properties which correlate with their temperatures and heats of fusion. Six kinds of packing motifs have been found. These demonstrate a considerable range in packing efficiency while retaining correlations with configuration of the mandelate anion. Interion H-bonding type is one of the important organizing principles. Links between some of the packing groups have been found in the form of polymorphs and solid-solid phase transitions in some of the salt phases.