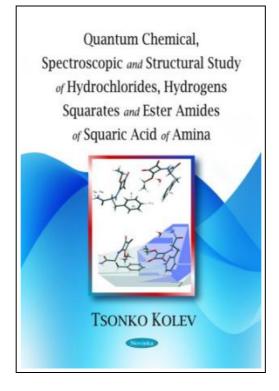
Quantum Chemical, Spectroscopic and Structural Study of Hydrochlorides, Hydrogens Squarates and Ester Amides of Squaric Acid of Amina



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Reviews

This book will never be easy to start on looking at but quite entertaining to read. It is actually packed with wisdom and knowledge It is extremely difficult to leave it before concluding, once you begin to read the book. (Ms. Missouri Satterfield DVM)

QUANTUM CHEMICAL, SPECTROSCOPIC AND STRUCTURAL STUDY OF HYDROCHLORIDES, HYDROGENS SQUARATES AND ESTER AMIDES OF SQUARIC ACID OF AMINA



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Nova Science Publishers Inc. Paperback. Book Condition: new. BRAND NEW, Quantum Chemical, Spectroscopic and Structural Study of Hydrochlorides, Hydrogens Squarates and Ester Amides of Squaric Acid of Amina, Tsonko Kolev, The interest to amino acid amides arises from their biological important role. Some C-nGBP\- amidated amino acids Ile, Val, Thr, Ser, Met, Trp, Gln and Arg have been studied by single crystal X-ray diffraction and their bioactivity have been compared with the corresponding amino acids due to most of mammalian peptide hormones as calcitonin, gastrin, neurokinins or neuropeptides possess a C-GBP\-terminal-amides. The most C-GBP\- amides are much more biologically active, comparing with the corresponding C-GBP\- terminal free acids. For example the "potency ratio" of peptide amide towards the corresponding peptide free acid in neurocinin is more than 40 000. Since the protonated forms of amino acid amides and C-GBP\-amidated peptides exists in the living cell their investigation could provide an understanding of their biological role. The choice of the acidity agent for the in vitro investigations are based manly of its own biological activity as for example squaric acid (H2Sq). Its application for synthesis of optically active amino acid derivatives with potential non-linear optical and electro-optical properties is well known, but its important biological role is intensively studied in last five years. A large number of medications based on H2Sq derivatives are effective inhibitors of protein tyrosine phosphatases or DNA polymerases from several viruses. H2Sq diamides replaced a phosphate diester linkage in oligodeoxynucleotide. Selective antagonist of ionotropic glutamate receptors is obtained by replacing of GBP^-carboxylic acid of a glutamate residue within a polyamine toxin with squaric acid derivatives. Some H2Sq-based peptides are inhibitors of matrix metalloprotease V1. These facts provoked the systematic investigations of hydrochlorides, hydrogensquarates and ester amides of squaric acid

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