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November 26, 2561 BC - "The most important thing is that the piece does not play in isolation, the piece does not play. For example, Vishy Anand drew everything his 3 games with Black in this.Â» - I. Rabinovich - â€œI would like to say the following about this book: I do not see any shortcomings in it, except for one thing - it is very long. I would suggest that it be a little shorter, at least two pages. And so everything is fine. I read it with great pleasure and am grateful to the author for this work. â€” R. Fischer â€” â€œIn my opinion, this book is an excellent example of what sports are made of. I read it with great interest and would very much like to see the same book in Russian.â€”

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1. Field of the Invention This invention relates to 1H-benz[de]isoquinoline compounds which have activity as cholecystokinin antagonists and their use as central nervous system agents.

2. Description of the Prior Art The octapeptide hormone cholecystokinin (CCK) was first isolated and identified as a major gastrointestinal regulatory peptide in 1975 by R. J. Giffin et al. See *Advances in Endocrinology and Molecular Biology*, Vol. 1, p. 165, published by Academic Press, Inc. (1976). It is also now known that the peptide is a major regulator of

pancreatic enzyme secretion and exocrine function of the gut. See Blumberg, B. A., *The Pharmacological Basis of Therapeutics*, 6th ed. p. 745, MacMillan Publishing Company (1980).

Molecules which are structurally homologous to CCK and are reported to exhibit CCK antagonist activity, include those disclosed in U.S. Pat. Nos. 4,182,763 and 4,182,764. The tetrapeptide derivatives described therein were derived from a related group of peptides, gastrin releasing peptides (GRP's) originally isolated from porcine and murine gastrointestinal tissue. See, Van Vugt, et al. *Science*, 1991, 254, 513-516. Other CCK antagonists of the above peptide type include those disclosed in U.S. Pat. Nos. 4,970,204 and 4,795,719, as well as EPO 0 245 191 A2. More recently, CCK

antagonists of the pyroglutamyl-glutamyl peptide type have been described in U.S. Pat. Nos. 5,093,329, 5,093,317 and 5,068,251. Two additional CCK antagonists of this type have also been described in U.S. Pat. No. 5,030,695. The present invention describes novel heterocyclic derivatives of CCK and their use as CCK antagonists. Contents In the video clip, the man states that he is representing Keith Fournier. Fourn c6a93da74d

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