



## **Peptides in Oncology III: Somatostatin and LH-RH Analogues: Somatostatin and LH-RH Analogues v. 3 (Recent Results in Cancer Research)**

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Interference with protein-mediated intra- and intercellular pathways has become a major goal of preclinical and clinical research. A rapidly increasing number of peptides are known to be responsible for endo-, para- and autocrine signal transduction. These peptides and their receptors have been studied with regard to their cell growth stimulatory action and their impact on differentiation. In parallel, peptide antagonists are being investigated in terms of their potential role in preclinical and clinical application. Thus, biotherapy might improve the clinical outcome of patients with tumors that respond to the respective hormonal manipulation. Among the numerous peptides of interest somatostatin (somatotropin release inhibitory factor) and the luteinizing hormone releasing hormone (LH-RH) have been the topic of intensive research during recent years. In this third volume of peptides in oncology, experts in the field extensively review and update the mechanisms of action of somatostatin and LH-RH-analogues in oncology. Somatostatin and its various short- and long-acting analogues have the unique feature of suppressing and inhibiting a wide range of cellular processes including cell proliferation. Receptors for these peptides which belong to the family of neuropeptides or neurotransmitters, are widely distributed, a feature which is not in keeping with the general view of a growth hormone regulatory system. LH-RH analogues play an established role in curative (adjuvant) and palliative treatment of hormone sensitive tumors.

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