

Webster K. Cavenee, Webster Cavenee, Nicholas Hastie, Eric J. Stanbridge

Recessive Oncogenes and Tumor Suppression Webster K. Cavenee, Webster Cavenee, Nicholas Hastie, Eric J. Stanbridge, 1989 Contributors to the meeting held at the Lab in March 1989 review mapping and cloning of cancer genes retino blastoma new putative tumor suppressor genes and the ways in which interactions between viral transforming proteins and cell proteins may lead to loss of growth control No index Annotatio Cancer Cells ,1990 National Library of Medicine Current Catalog National Library of Medicine (U.S.),1990 Cancer Research .1990 Current Cataloa **Directory of Published** National Library of Medicine (U.S.), First multi year cumulation covers six years 1965 70 Oncogenes and Tumour Suppressors Dr. Gordon Peters, Karen H. Vousden, 1997 Oncogenes and **Proceedings** .1991 tumour suppressor genes have been the focus of much research because of their role not only in cancer but also in normal cell growth and differentiation Oncogenes and Tumour Suppressors is a unique volume that brings together a team of leading researchers to present and critically assess our current knowledge The book covers two major areas of interest oncogenes and signal transduction and tumour supressors and cell cycle control Topics covered include the mechanisms of oncogene perturbation growth factors and their receptors in cell transformation oncogenic cytoplasmic protein tyrosine kinases the RAS RAP ERK signal transduction pathway oncogenic transcription factors mammalian cell cycle control the retinoblastoma gene product and its relatives the tumour suppressor gene p53 tumour suppressors and the inheritance of cancer the clinical relevance of oncogenes Oncogenes and Tumour Suppressors is a major review work that provides an unparalleled summary of our current understanding of this field as viewed by some of its leading researchers It is of interest to all those involved in research on the cell cycle signal transduction and cancer biology The book is also an excellent reference source with over 1800 selected citations to the primary literature **Tumor Suppressor Genes** Wafik S. El-Deiry, 2008-02-03 It has become clear that tumors result from excessive cell proliferation and a corresponding reduction in cell death caused by the successive accumulation of mutations in key regulatory target genes over time During the 1980s a number of oncogenes were characterized whereas from the 1990s to the present the emp sis has shifted to tumor suppressor genes TSGs It has become clear that oncogenes and TSGs function in the same pathways providing positive and negative growth regulatory activities The signaling pathways controlled by these genes involve virtually every process in cell biology including nuclear events cell cycle cell death cytoskeletal cell membrane angiogenesis and cell adhesion effects Mu tions in tumor suppressor genes have been identified in familial cancer syndromes and the same genes in many cases have been found to be mutationally inactivated in sporadically occurring cancers In their normal state TSGs control cancer development and progression as well as contribute to the sensitivity of cancers to a variety of therapeutics Understanding the classes of TSGs the biochemical pa ways they function in and how they are regulated provides an essential lesson in cancer biology We

cannot hope to advance our current knowledge and to develop new and more effective therapies without understanding the relevant pathways and how they influence the present approaches to therapy Moreover it is important to be able to access not only the powerful tools now available to discover these genes but also their links to cell biology and growth control

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