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International Journal of Clinical Pharmacology. and its clinical applications Microscopy of bone marrow neoplastic proliferations, particularly the acute leukemia. Histopathology and cytology. An integral part of the flow cytometric analysis, the hemoglobin binding properties of. The analytical features of light scattering properties in flow cytometry have led to the. Latest developments in automated metaphase analysis have led to a third dimension in. complex of surface antigens using the most accurate techniques available.

PRINCIPLES AND APPLICATIONS IN CLINICAL HEMATOLOGY. The first large, multi-center studies in adults demonstrated that the prognosis for AML patients with. In acute leukemias that were clonally.

Introduction Flow cytometry is a high resolution, multiparameter technology for analyzing cells and. Cells and a number of publications have described immunophenotypic characteristics of. Protocols Available Flow Cytometric Cell Analysis.. Data-analysis software and possible limitations of the technology. Interference with cellular signal transduction cascades is a widespread phenomenon in biology and has been implicated in many human diseases including cancer. DNA damage has been shown to initiate many signal transduction cascades, including the phosphorylation and degradation of the tumor suppressor protein p53. Mutation and loss of p53 function are correlated with poor patient prognosis in many human cancers. We discovered that DNA double strand breaks (DSBs) activate signaling pathways in part through the ATM, ATR and DNA-PK kinases that phosphorylate and activate p53. We also discovered that DNA damage induces apoptosis through the ATM/p53/GADD45 and p53/PUMA pathways, but that replication-associated DNA damage inhibits apoptosis in part through the ATR/p53/GADD45 pathway. Since CDDP fails to induce p53-mediated apoptosis in many cancer cells, we hypothesize that CDDP does not induce apoptosis in part because it preferentially induces DNA damage that activates the ATR/p53/GADD45 pathway and inhibits the ATM/p53/GADD45 pathway. To test this hypothesis, we propose to: 1) Test the effect of ATR and CDDP on p53 activation in a diverse collection of human cancer cell lines and tumor specimens. 2) Dissect the pathway(s) induced by ATR that promote p53 activation and apoptosis. 3) Determine the role of the ATM/p53/GADD45 pathway in tumor cell responses to 595f342e71

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