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RNA Interference Using c-Myc–Conjugated Nanoparticles Suppresses Breast and Colorectal Cancer Models

The newly generated Newcastle disease virus rNDV-B1/Fas encodes the human TNF receptor Fas. Human and murine cancer cells infected by rNDV-B1/Fas displayed a modified cell death response compared with the wild-type rNDV-B1, characterized by an earlier and enhanced apoptosis response due to the overexpression of Fas following the coactivation of both extrinsic and intrinsic apoptosis pathways. The enhanced cytotoxicity shown in vitro correlated with an improved oncolytic activity and therapeutic effect of rNDV-B1/Fas virus in intratumoral-treated melanoma-bearing mice. The cover image shows murine B16-F10 melanoma cells infected by rNDV-B1/Fas virus. The human Fas receptor (red) is expressed on the surface of infected cells (green) but also internalized following induction of apoptosis. DNA was counterstained with Hoechst. For details, see the article by Cuadrado-Castano and colleagues on page 1247.
Molecular Cancer Therapeutics

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