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trophoblast cells is described. For this purpose, the human trophoblast cell line SGHPL-4 was transfected with a plasmid containing an SV40 enhancer/promoter linked to the green fluorescent protein (GFP) gene to permit the monitoring of cell fusion by analysis of GFP expression. Evidence of trophoblast cell fusion was obtained by the detection of GFP-positive cells. In addition, the expression of tumor necrosis factor-alpha was studied by reverse transcriptase-polymerase chain reaction. The expression of the tumor necrosis factor-alpha gene was found to be upregulated when the trophoblast cell line was exposed to fusion-inducing

agents such as tumor necrosis factoralpha and phorbol 12-myristate 13-acetate. To validate the model for studying gene expression during human trophoblast fusion, gene expressions that were previously shown to be involved in trophoblast cell fusion were studied by quantitative reverse transcriptasepolymerase chain reaction. Upregulated expression of the invasion-associated genes, urokinase-type plasminogen activator, leukocyte common antigen, and intercellular adhesion molecule-1 was observed. In addition, the fusionrelated genes, glucose transporter-1 and -3, the integrin alpha6beta4, and the adapter protein EAT-2 were also found

to be upregulated. The results presented in this study demonstrate that fusion-related genes are upregulated during trophoblast cell fusion. The role of pretreatment measurements in target delineation for external beam radiotherapy of prostate cancer. An MRI based image fusion method was applied for pre-treatment target delineation in prostate cancer patients. The aim of this study is the evaluation of

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