

CXCR1/CXCR2受体拮抗剂—G31P抑制前列腺癌血管新生的体内实验

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Inhibition of G31P:Chemokine Receptor CXCR1/CXCR2 Antagonist,in Angiogenesis of Human Prostate Cancer Cells in vivo

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摘要

目的

探讨G31P (CXCR1/CXCR2受体拮抗剂) 对人前列腺癌PC-3细胞的体内血管新生的抑制作用。方法建立体内绿色荧光蛋白(GFP)标记的人雄激素非依赖性前列腺癌细胞PC-3的裸鼠原位移植瘤模型, 观察G31P对裸鼠前列腺原位移植瘤血管新生的影响。结果与对照组(1.26±0.46)相比, G31P处理组明显抑制前列腺肿瘤的血管新生(0.49±0.12, P<0.05), 与对照组相比, G31P处理组VEGF

(P<0.01) 和NF-κB (P<0.01) 的表达具有统计学意义(免疫组织化学法)。结论在裸鼠原位移植瘤模型中G31P对人雄激素非依赖性前列腺癌的血管新生有明显抑制作用。

关键词: G31P 前列腺癌 血管新生 CXCR1 CXCR2

Abstract:

Objective

To investigate the inhibition of G31P on the angiogenesis of the prostate cancer PC-3 cell in vivo.MethodsThe effect of G31P on angiogenesis of human prostate tumor of nude mice were observed in nude mice by building a human androgen-independent prostate cancer PC-3 (GFP-labeled) orthotopic transplantation tumor cells model.ResultsThe tumor angiogenesis of G31P treated group (1.26±0.46) was significantly reduced (0.49±0.12, P<0.05) compared with the control group.VEGF (P<0.01) and NF-κB (P<0.01) expression of G31P treated groupwas significantly reduced (immunohistochemistry) compared with the control group.ConclusionG31P could inhibit the angiogenesis of the prostate cancer PC-3 cell in vivo.

Key words: G31P Prostate cancer Angiogenesis CXCR1 CXCR2

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