## Basic study of gene therapy for hepatocellular carcinoma

メタデータ	言語: jpn
	出版者:
	公開日: 2022-06-09
	キーワード (Ja):
	キーワード (En):
	作成者: Kobayashi, Kenichi
	メールアドレス:
	所属:
URL	https://doi.org/10.24517/00066223

This work is licensed under a Creative Commons Attribution-NonCommercial-ShareAlike 3.0 International License.



## 1996 Fiscal Year Final Research Report Summary

## Basic study of gene therapy for hepatocellular carcinoma

Research Project

Project/Area Number
07457133
Research Category
Grant-in-Aid for Scientific Research (B)
Allocation Type
Single-year Grants
Section
一般
Research Field
Gastroenterology
Research Institution
Kanazawa University
Principal Investigator
KOBAYASHI Kenichi Medical department, Kanazawa University, Professor, 医学部, 教授 (70019933)
Co-Investigator(Kenkyū-buntansha)
MATSUSHITA Eiki Department of Neurosurgery, Kanazawa University Assistant professor, 医学部・附属病院, 講師 (00242545) KANEKO Shuichi School of Medicine, Kanazawa University Associate Professor, 医学部, 助教授 (60185923)
Project Period (FY)
1995 – 1996
Keywords
gene therapy / hepatocellular carcinoma / retroviral vector / herpes simplex thymidine kinase / ganciclovir

**Research Abstract** 

Currently, clinical trials of suicide gene transfer with retroviral vector producer cells are being carried out for the treatment of brain tumors. The feasibility of applying this approach for the treatment of hepatocellular carcinoma is studied.

Methods: Five hepatoma cell lines were tested for retroviral mediated transduction efficiency with the herpes simplex thymidine kinase (HSV-TK) gene and for the induced drug sensitivity to ganciclovir. Different populations of HSV-TK transduced HuH7 cells were subcutaneously injected into athymic nude mice with non transduced HuH7 cells to analyze the "bystander" effect. The retroviral vector producer cells were intratumorally injected into pre-established HuH7 tumors to study the antitumor effect.

Results: Five hepatoma cell lines were transduced with the viral supernatant, and all resultant 50 transduced clones, except one, were sensitive to ganciclovir. Complete regression occurred in 21 out of 30 tumors when the inoculum consisted of as few as 10% of HSV-TK containing transduced cells. The producer cell injection into tumors and GCV treatment group was significantly smaller in size of the tumors compared to GCV treatment group (p<0.01).

Conclusions: This gene therapy approach could potentially be effective for the treatment of human hepatocellular carcinoma.

## Research Products (13 results)

All Other All Publications (13 results) [Publications] Shuichi Kaneko: "Adenovirus-mediated gene therapy of hepatocellular carcinoma using cancer-specific gene expression." Cancer Research. 55. 5283-5287 (1995) [Publications] 松下栄紀: "肝疾患の遺伝子治療はどこまで期待できるか。" medicina. 33. 544-546 (1996) [Publications] 松下栄紀: "肝細胞癌の遺伝子治療" 医学のあゆみ. 176. 963-966 (1996) [Publications] Shuichi Kaneko, et al: "Adenovirus-mediated gene therapy of hepatocellular carcinoma using cancer-specific gene expression." Cancer Research. 55(15). 5283-5287 (1995) [Publications] Eiki Matsushita, Shuichi Kaneko, Kenichi Kobayashi: "Possibility of gene therapy for liver diseases." Medicina. 33(3). 544-547 (1996) [Publications] Eiki Matsushita, Shuichi Kaneko: "Gene therapy for hepatocellular carcinoma." Journal of Clinical and Experimental Medicine (IGAKU NO AYUMI). 176 (13). 963-966 (1996) [Publications] Kenichi Kobayashi, Shuichi Kaneko, et al: "Clinical pathoepidemiology of hepatocellular carcinoma in japan." Hepatitis C virus and its involvement in the developement of hepatocellular carcinoma. pp67-74, edited by Kenichi Kobayashi, RH Purcell, K Shimotohno, and E Taber Princeton Scientific Publishing Co., Inc.(1995) [Publications] Yasunari Nakamoto, Shuichi Kaneko, Kenichi Kobayashi, et al: "B-Cell Epitopes in Hypervariable Region 1 of Hepatitis C Virus Obtained From Patients With Chronic Persistent Hepatitis." Journal of Medical Virology. 50. 35-41 (1996) [Publications] Yutaka Inagaki, Kenichi Kobayashi, et al: "Regulation of the a2 (I) Collagen Gene Transcription in Fat-Storing Cells Derived From a Cirrhotic Liver." Hepatology. 22. 573-579 (1995) [Publications] Gabriele Pozzato, Suichi Kaneko, Kenichi Kobayashi, et al: "Interferon Therapy in Chronic Hepatitis C Virus: Evidence of Different Outcome With Respect to Different Viral Strains." Journal of Medical Virology. 45. 445-450 (1995) [Publications] Naoki Ikeda, Shuichi Kaneko, Kenichi Kobayashi, et al: "Prevention of Endotoxin-Induced Acute Lethality in Propionibacterium acnes-Primed Rabbits by an Antibody to Leukocyte Integrin b2 with Concomitant Reduction of Cytokine Production." Infection and Immunity. 63. 4812-4817 (1995)[Publications] Kyosuke Kaji, Ken-ichi Kobayashi, et al: "Hemosiderin Deposition in Portal Endothelial Cells: A Novel Hepatic Hemosiderosis Frequent in Chronic Viral Hepatitis B and C." Human Pathology. 26. 1080-1085 (1995) [Publications] Rieko Saeki, Shuichi Kaneko, Kenichi Kobayashi, et al: "Effects of chenodeoxycholic and ursodeoxycholic acids on interferon-g

production by peripheral blood mononuclear cells from patients with primary biliary cirrhosis." Journal of Gastroenterology. 30. 739-744 (1995)

**URL:** https://kaken.nii.ac.jp/report/KAKENHI-PROJECT-07457133/074571331996kenkyu\_seika\_hokoku\_

Published: 1999-03-08