## Angiogenic switch in the development of colon cancer

メタデータ	言語: jpn
	出版者:
	公開日: 2021-09-10
	キーワード (Ja):
	キーワード (En):
	作成者: Takahashi, Yutaka
	メールアドレス:
	所属:
URL	https://doi.org/10.24517/00063970
	This work is licensed under a Creative Commons

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



## 2000 Fiscal Year Final Research Report Summary

## Angiogenic switch in the development of colon cancer

**Research Project** 

Project/Area Number
11671217
Research Category
Grant-in-Aid for Scientific Research (C)
Allocation Type
Single-year Grants
Section
一般
Research Field
Digestive surgery
Research Institution
Kanazawa University
Principal Investigator
TAKAHASHI Yutaka Cancer Research In stitute, Kanazawa University Surgical Oncology, Associate Professor, ガス研究所, 助教授 (10179541)
Project Period (FY)
1999 – 2000
Keywords
angiogenesis / colon cancer / VEGF / PD-ECGF / MMP-7 / angiogenic switch

**Research Abstract** 

We have already reported that vessel count, vascular endothelial growth factor (VEGF) produced by cancer cell and platelet derived endothelial cell growth factor (PD-ECGF) produced by infiltrating cells correlate with metastasis in human colon cancer. We studied whether there is anigiogenic switch in the development of colon cancer. We studied vessel count, VEGF, another angiogenic factor, basic figroblast growth factor (bFGF) and matrix metalloproteinase (MMP)-7 which is well known to be important for colon cancer, expressions in cancer cells and PD-ECGF expression in infiltrating cells in 25 adenomas, 35 mucosal cancers (in situ), 29 submucosal invasive cancers (sm) and 33 muscle propria invasive cancers (mp) by immunostaining.

The vessel density was  $12.7 \text{\AA} 6.7$  (SD) in adenoma,  $11.8 \text{\AA} 8.3$  in in situ,  $35.0 \text{\AA} 17.5$  in sm, and  $35.2 \text{\AA} 18.8$  in mp. There was significant difference between in situ and sm (p<0.001). The intensity of VEGF expression was  $0.6 \text{\AA} 0.4$ ,  $0.9 \text{\AA} 0.7$ ,  $1.7 \text{\AA} 0.9$ , and  $1.8 \text{\AA} 0.8$ , respectively. There was also significant difference between in situ and sm (p<0.001). There were also significant differences in the intensity of the expression of MMP-7 and PD-ECGF between in situ and sm shown as table. These results suggest that angiogenic switch "on" may occur between in situ and sm, in other word, start of invasion, in the development of colon cancer.

		Α	ll Othe	r
	All	Pub	lications	s
[Publications] Takahashi Y, et al: "DFMO induces apoptosis as well as anti-angiogenesis in the inhibition of tumor growth and Metastasis"Int.J.Cancer. 85. 243	3-247	(200	0) 🗸	,
[Publications] 高橋豊: "癌治療の新たな戦略Tumor Dormany Therapy"医学書院,東京. 1-172 (2000)			~	
[Publications] Takahashi Y, Mai M and Nishioka K: "a-Difluoromethylornithine Induces Apoptosis as well as Anti-angiogenesis in the Inhibition of Tumor Growt Metastasis in a Human Gastric Cancer Model."International Journal of Cancer. 85. 243-247 (2000)	:h anc	ł	~	

URL: https://kaken.nii.ac.jp/report/KAKENHI-PROJECT-11671217/116712172000kenkyu\_seika\_hokoku\_

Published: 2002-03-25