

Isolation and Characterization of Dinochrome A and B, Anti-carcinogenic Active Carotenoids from the Fresh Water Red Tide *Peridinium bipes*

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Two epimeric carotenoids, named dinochromes A (2) and B (3), were isolated from the fresh water red tide *Peridinium bipes*, as anti-carcinogenic compounds. The stereostructure of dinochrome A and B were characterized to be (3*S*,5*R*,6*R*,3'*S*,5'*R*,8'*R*)- and (3*S*,5*R*,6*R*,3'*S*,5'*R*,8'*S*)-5',8'-epoxy-6,7-didehydro-5,6,5',8'-tetrahydro- β , β -carotene-3,5,3'-triol 3-*O*-acetate, respectively by ¹H- and ¹³C-NMR, and circular dichroism (CD) data. Dinochromes A (2) and B (3) inhibit 12-*O*-tetradecanoyl phorbol 13-acetate (TPA)-stimulated ³²P-incorporation into the phospholipids of HeLa cells. Furthermore, dinochrome A was found to inhibit the proliferation of human malignant tumor cells, such as GOTO, OST and HeLa cells.

Key words *Peridinium bipes*; carotenoid; anti-carcinogenic activity; anti-tumor activity; dinochrome A; dinochrome B

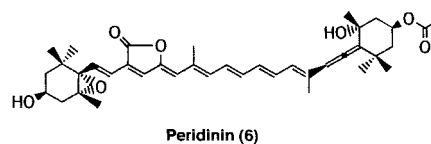
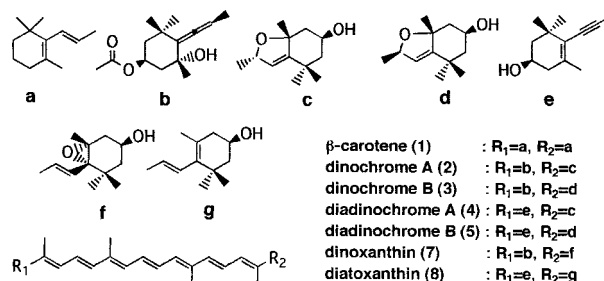
Peridinin is the main carotenoid in red tide,²⁾ and exhibits anti-tumor and anti-carcinogenic activities.^{3,4)} This prompted us to search for anti-carcinogenic active compounds from microalgae, and to isolate and characterize carotenoids that possess anti-carcinogenic activities from the fresh water red tide, *Peridinium bipes*. In this paper, we describe the isolation and the characterization of the stereostructure of two epimeric carotenoids, named dinochromes A and B. We also demonstrate the inhibitory effect of these carotenoids on 12-*O*-tetradecanoyl phorbol 13-acetate (TPA)-stimulated ³²P-incorporation into the phospholipids of HeLa cells; a mechanism that is used in the primary screening test for anti-carcinogenic compounds.⁵⁾ Furthermore, the inhibitory effects of dinochrome A on human malignant tumor cells were also studied.

Isolation of Carotenoids from *Peridinium bipes* The fresh water red tide *Peridinium bipes* (500 g), was extracted with acetone–methanol (7 : 3). After evaporation of the solvent, the residue was subjected to column chromatography on silica gel and preparative HPLC on octadecyl silica (ODS). Eight carotenoids were isolated; β -carotene (1) (40.5 mg), dinochrome A (2) (71.5 mg), dinochrome B (3) (13.5 mg), diadinochrome A (4) (13.5 mg), diadinochrome B (5) (1 mg), peridinin (6) (132.5 mg), dinoxanthin (7) (8.5 mg), and diatoxanthin (8) (34 mg). Compounds 1, 4, 5, 6, 7, and 8 were identified by comparing their spectral data with the values reported in the literature.^{6,7)}

Characterization of the Stereostructure of Dinochrome A and B Compound 2 (dinochrome A) exhibited visible light absorption maxima at 400, 423, and 450 nm. High resolution electron impact (HR-EI)-MS gave a molecular ion at *m/z* 642.4280, corresponding to the formula C₄₂H₅₈O₅. Acetylation of 2 produced a mono acetate with a molecular ion of *m/z* 684.4395. The presence of one secondary hydroxy group, one tertiary hydroxy group and one acetoxy group in 2 was confirmed by HR-EI-MS, ¹H-, ¹³C-NMR data. The remaining one oxygen is ascribed to a furanoxide group by ¹³C-NMR (δ 86.8, 87.7)⁷⁾ and a characteristic EI-MS fragment ion of M-80.⁸⁾ The presence of an allenic group was revealed by the characteristic ¹³C signal at δ 204.4 and the ¹H signal at δ 6.05.⁷⁾ The ¹H- and ¹³C-NMR data for 2, which

were assigned by double quantum filtered correlation spectroscopy (DQF-COSY), nuclear Overhauser effect spectroscopy (NOESY), heteronuclear single quantum coherence (HSQC), and heteronuclear multiple bond connectivity (HMBC), are presented in Table 1. These results revealed that the structure of 2 is 5',8'-epoxy-6,7-didehydro-5,6,5',8'-tetrahydro- β , β -carotene-3,5,3'-triol 3-*O*-acetate (dinochrome). The 5',8'-*trans* configuration in the furan ring was determined by the coupling constant of $J_{H-7'-H-8'}=0.7$ Hz^{7,9–11)} and the NOESY data. NOE between CH₃-18' (δ 1.62) and H-8' (δ 5.17) indicates that CH₃-18' and H-8' are located on the same side of the furan ring (Fig. 1).

Compound 3 (dinochrome B) showed the same UV–vis and EI-MS as compound 2. Acetylation of 3 also produced a mono acetate. The ¹H- and ¹³C-NMR data (Table 1) of 3 were consistent with the 8'-epimer structure of dinochrome A.^{7,9–11)} The coupling constant of $J_{H-7'-H-8'}=1.5$ Hz indicated a 5',8'-*cis* configuration in the furan ring. NOESY correlation between CH₃-18' and H-8' was not observed in compound 3. The absolute configurations of compounds 2 (3*S*,5*R*,6*R*,3'*S*,5'*R*,8'*R*) and 3 (3*S*,5*R*,6*R*,3'*S*,5'*R*,8'*S*) were determined by comparing the circular dichroism (CD) spectral data of the compounds with those of the stereoisomers of the neochromes that were reported by Marki-Fischer *et al.*¹¹⁾



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Table 1. ¹H- (500 MHz) and ¹³C-NMR (125 MHz) Data of Dinochrome A (**2**) and Dinochrome B (**3**) in CDCl₃

Position	Compound			
	2		3	
	$\delta^{13}\text{C}$	$\delta^1\text{H}$ mult. (J, Hz)	$\delta^{13}\text{C}$	$\delta^1\text{H}$ mult. (J, Hz)
1	35.7	—	35.7	—
2	45.4	β 1.41 dd (12.0, 12.0) α 1.91 dd (12.0, 4.0)	45.4	β 1.41 dd (12.0, 12.0) α 1.91 dd (12.0, 4.0)
3	67.8	5.38 m	67.8	5.38 m
4	45.2	β 1.51 dd (12.0, 12.0) α 2.28 ddd (13.5, 4.0, 2.0)	45.2	β 1.51 dd (12.0, 12.0) α 2.28 ddd (13.5, 4.0, 2.0)
5	72.7	—	72.7	—
6	117.4	—	117.4	—
7	204.4	—	204.4	—
8	103.4	6.05 s	103.4	6.05 s
9	131.7	—	131.7	—
10	128.6	6.12 d (11.0)	128.6	6.12 d (11.0)
11	124.7	6.54 dd (15.0, 11.0)	124.7	6.54 dd (15.0, 11.0)
12	137.4	6.34 d (15.0)	137.4	6.34 d (15.0)
13	138.0	—	138.0	—
14	132.5	6.24 m	132.5	6.24 m
15	130.0 ^{a)}	6.62 m	130.0 ^{a)}	6.62 m
16	32.1	1.07 s	32.1	1.07 s
17	28.9	1.35 s	28.9	1.35 s
18	29.2	1.38 s	29.2	1.38 s
19	13.9	1.80 s	13.9	1.80 s
20	12.8	1.96 s	12.8	1.96 s
CH ₃ -CO	21.4	2.04 s	21.4	2.04 s
CH ₃ -CO	170.4	—	170.4	—
1'	33.7	—	34.2	—
2'	46.7	α 1.52 dd (14.5, 3.5) β 1.77 ddd (14.0, 5.0, 1.5)	47.5	α 1.48 dd (14.5, 3.5) β 1.80 ddd (14.0, 5.0, 1.5)
3'	68.0	4.25 m	68.0	4.25 m
4'	47.3	α 1.99 dd (14.0, 4.0) β 2.13 ddd (14.0, 4.0, 1.5)	47.5	α 1.90 dd (14.0, 4.0) β 2.12 ddd (14.0, 4.0, 1.5)
5'	86.8	—	87.2	—
6'	154.1	—	153.3	—
7'	119.9	5.26 d (0.7)	118.8	5.31 d (1.5)
8'	87.7	5.17 br s	88.4	5.07 br s
9'	137.9	—	138.7	—
10'	127.2	6.19 d (11.0)	126.2	6.19 d (11.0)
11'	124.4	6.49 dd (15.0, 11.0)	124.4	6.51 dd (15.0, 11.0)
12'	137.6	6.32 d (15.0)	137.6	6.32 d (15.0)
13'	136.2	—	136.3	—
14'	132.6	6.24 m	132.6	6.23 m
15'	129.9 ^{a)}	6.62 m	129.9 ^{a)}	6.62 m
16'	31.4	1.17 s	31.2	1.20 s
17'	28.9	1.33 s	28.1	1.35 s
18'	29.0	1.62 s	30.6	1.69 s
19'	12.6	1.71 s	13.4	1.80 s
20'	12.8	1.95 s	12.8	1.95 s

a) Assignments may be interchangeable in each vertical column.

The CD spectra of **2** and **3** were mirror-images each other and reflected chirality at C-8'.¹²⁾

The planer structures of dinochrome were postulated by Johansen *et al.*⁶⁾ This is the first report on the characterization of the stereostructure of dinochromes by NMR and CD spectral data and we named dinochrome A for **2** and dinochrome B for **3**, respectively. Dinochromes A and B were assumed to be acidic-catalyzed epoxy-furanoxide rearrangement products of dinoxanthin in *P. bipes*.

Anti-carcinogenic Activity of Carotenoids from *P. bipes*
The *in vitro* anti-carcinogenic activity of dinochrome A (**2**), dinochrome B (**3**), peridinin (**6**), and diatoxanthin (**8**) isolated from *P. bipes* was evaluated by assessing the inhibitory effect

on TPA-stimulated ³²P-incorporation into the phospholipids of HeLa cells.⁵⁾ All of the carotenoids investigated inhibited this incorporation potentially at a concentration of 25 $\mu\text{g}/\text{ml}$, as shown in Table 2. Dinochrome A and dinochrome B were found to be more effective than peridinin that has been studied in cancer prevention experiments using animal models.^{3,4)} Among the carotenoids examined,¹³⁾ dinochrome A showed the strongest inhibitory activity on TPA-stimulated ³²P-incorporation into the phospholipids of HeLa cells. Some carotenoids exhibit anti-carcinogenic and anti-tumor activities.^{3,4,13,14)} Thus, the inhibitory effects of dinochrome A on the proliferation of human malignant cells such as GOTO (neuroblastoma), OST (osteosarcoma) and HeLa cells were

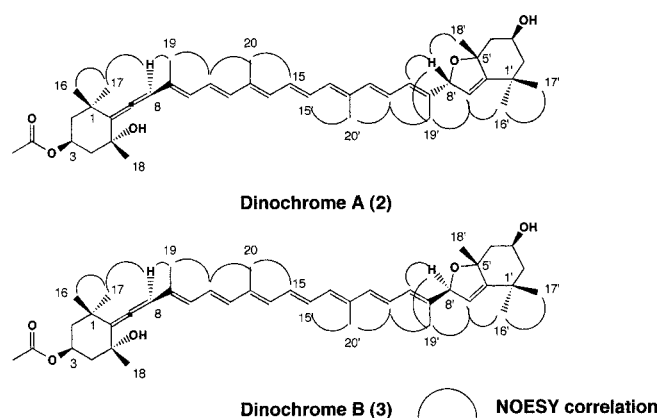


Fig. 1. Structures and Some Key NOE Correlations of Dinochrome A (2) and Dinochrome B (3)

Table 2. Effect of Red Tide Carotenoids on TPA-Stimulated ^{32}P -Incorporation into the Phospholipids of HeLa Cells

Carotenoid	Inhibition %
Dinochrome A (2)	72.1
Dinochrome B (3)	35.0
Peridinin (6)	28.2
Diatoxanthin (8)	48.2

TPA: 50 nM, Carotenoid: 25 $\mu\text{g}/\text{ml}$.

Table 3. Inhibitory Effects of Dinochrome A (2) on the Proliferation of Human Malignant Tumor Cells

Type of cells	Viable cell number ($\times 10^4/\text{dish}$)		
	Control	+Dinochrome A (5 $\mu\text{g}/\text{ml}$)	Inhibition %
GOTO	8.8	1.6	81.8
OST	18.3	4.5	75.4
HeLa	17.0	4.3	74.7

evaluated. Dinochrome A inhibited the growth of GOTO, OST and HeLa cells at the concentration of 5 $\mu\text{g}/\text{ml}$, as shown in Table 3. Therefore, dinochrome A exhibits a marked anti-carcinogenic activity and is an effective inhibitor of the proliferation of human malignant tumor cells *in vitro*.

Some marine carotenoids, such as fucoxanthin, peridinin and halocynthiaxanthin, which possess an allenic or an acetylenic group, exhibit anti-carcinogenic activity.^{3,4} In this study, we found that dinochromes A and B, which possess an allenic group, also exhibit marked anti-carcinogenic activity. These allenic carotenoids may be valuable as an anti-tumor promoter and chemopreventive agents in chemical carcinogenesis.

Experimental

General Experimental Procedures The UV-vis and CD spectra were recorded in Et_2O at room temperature with a Shimadzu UV-240 spectrophotometer and a JASCO J-500C spectropolarimeter, respectively. The EM-MS spectra were recorded using a JEOL JMS-SX 102A and a JEOL JMS-GC-mate. The NMR spectra were measured with a Varian UNITY INOVA 500 (^1H : 500 MHz, ^{13}C : 125 MHz) instruments in CDCl_3 with tetramethylsilane (TMS) as internal standard. DQF-COSY, NOESY, HSQC, and HMBC were acquired using the standard Varian pulse programs. HPLC was performed on a Shimadzu LC-6AD instruments with a Shimadzu SPD-6AV spectrometer set at 450 nm. The column used was a Shim-pack Prep ODS with a mobile

phase of CH_2Cl_2 - CH_3CN (2:8). Chemical derivatizations such as acetylation were according to our routine method.¹⁵

Extraction and Isolation The red tide was collected by suction of surface water at Fuya dam in Nara Prefecture in August 1995. The red tide cells was harvested by centrifugation and dried over 40 $^\circ\text{C}$ for 2 d. The dried red tide (500 g) was extracted with acetone-methanol (7:3). After evaporation of the solvent, the brown green residue was subjected to successively column chromatography on silica gel using an increasing percentage of ether in hexane. The fraction eluted with hexane-ether (2:8) from silica gel column chromatography was further purified by preparative HPLC on ODS with CH_2Cl_2 - CH_3CN (2:8) to yield dinochrome A (2) (71.5 mg) and dinochrome B (3) (13.5 mg). The following six carotenoids were isolated; β -carotene (1) (40.5 mg), diadinoxanthin A (4) (13.5 mg), diadinoxanthin B (5) (1 mg), peridinin (6) (132.5 mg), dinoxanthin (7) (8.5 mg), and diatoxanthin (8) (34 mg).

Dinochrome A (2): A yellow crystal. Yield 71.5 mg. EI-MS m/z (rel. int., %): 642 $[\text{M}]^+$ (26), 624 $[\text{M}-\text{H}_2\text{O}]^+$ (25), 564 (15), 562 $[\text{M}-80]^+$ (13), 544 (45), 221 (69), 43 (100). HR-EI-MS: Calcd for $\text{C}_{42}\text{H}_{58}\text{O}_5$ (M^+): 642.4284. Found: 642.4280. UV-vis: λ_{max} (Et_2O) nm 400, 423, 450. CD: λ nm ($\Delta\epsilon$) 220 (-3), 250 (-6), 275 (0). ^1H - and ^{13}C -NMR: Table 1.

Dinochrome B (3): A yellow crystal. Yield 13.5 mg. EI-MS m/z (rel. int., %): 642 $[\text{M}]^+$ (25), 624 $[\text{M}-\text{H}_2\text{O}]^+$ (23), 564 (22), 562 $[\text{M}-80]^+$ (20), 544 (50), 221 (60), 43 (100). HR-EI-MS: Calcd for $\text{C}_{42}\text{H}_{58}\text{O}_5$ (M^+): 642.4284. Found: 642.4282. UV-vis: λ_{max} (Et_2O) nm 400, 423, 450. CD: λ nm ($\Delta\epsilon$) 225 (2), 250 (5), 275 (0). ^1H - and ^{13}C -NMR: Table 1.

TPA-Stimulated ^{32}P -Incorporation into the Phospholipids of HeLa Cells Radioactive inorganic phosphate (^{32}P) was obtained from the Japan Radioisotope Association, Tokyo and TPA was obtained from Sigma Chemical Co. (St. Louis, MO, U.S.A.). HeLa cells (human cervical cancer cells) were maintained as monolayers in Eagle's minimum essential medium, which was supplemented with 10% calf serum, in humidified air with 5% CO_2 . HeLa cells were incubated with the test samples (25 $\mu\text{g}/\text{ml}$) and after 1 h, ^{32}P (370 kBq per culture) was added, with or without TPA (50 nM). Following 4 h of incubation, the amount of radioactivity incorporated into the phospholipid fraction was measured.⁵

Assays of the Proliferation of Human Malignant Tumor Cells GOTO (neuroblastoma), OST (osteosarcoma) and HeLa cells were cultured in Dulbecco's modified Eagle's minimum essential medium, which was supplemented with 10% fetal bovine serum, in humidified air with 5% CO_2 at 37 $^\circ\text{C}$. The cells were subcultured at a density of 4×10^4 cells, in 2 ml of medium, in 35 mm diameter Petri dishes. Dinochrome A was dissolved in dimethyl sulfoxide (5 $\mu\text{g}/\text{ml}$) and was added to the cultured medium. Following incubation for 3 d, the number of viable cells was calculated. The control culture was treated with the vehicle alone.¹⁶

References and Notes

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