COMPARATIVE STEROLS COMPOSITION OF THE RED ALGA ASPARAGOPSIS ARMATA AND ITS TETRASPOROPHYTE FALKENBERGIA RUFOLANOSA

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ABSTRACT.—The sterol content ranged from 0.015% of the dry weight for Asparagopsis armata to 0.075% for Falkenbergia rufolanosa. Four samples at different reproductive states are analyzed by means of gas-liquid chromatography and gas-liquid chromatography—mass spectrometry. Cholesterol 2 is always the major sterol. In every case two cholesta-diene-diols 8a, 8b and two cholestene-diols 5, 6 have been identified.

The red alga Asparagopsis armata Harv. is a dioecious gametophytic plant that alternates in its life cycle with a heteromorphic tetrasporophyte known as Falkenbergia rufolanosa (Harv.) Schmitz. The comparative chemical composition of the gametophyte and the tetrasporophyte has been previously reported for halogenated compounds (1). We have now studied the sterol content of some samples of Asparagopsis armata and Falkenbergia rufolanosa at different reproductive states (table 1).

TABLE 1.	Samples	studied for	sterols.
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Samples	Date	Reproductive state
$\begin{array}{c} \mathbf{A_1}\\ \mathbf{A_2}\\ \mathbf{F_1}\\ \mathbf{F_2} \end{array}$	3rd Mar 1976 29th Jun 1976 14th Jan 1976 20th Dec 1976	A sparagopsis armata young without fructification A sparagopsis armata with fructifications Falkenbergia rufolanosa young without tetraspore Falkenbergia rufolanosa with tetraspores

MATERIALS AND METHODS

Male and female plants of Asparagopsis armata, which were not separated for this investigation, and the asexual Falkenbergia rufolanosa were obtained along the Catalan Mediterranean coast near Banyuls-sur-Mer in France. The freshly collected, wet plants were extracted with methanol, chloroform, and ethanol. Solvents were evaporated, and the residues were transferred into ethyl ether to give the lipid oil, which was saponified. Sterols were precipitated from the non-saponifiable fraction with digitonin (2). Further purification was obtained by tle with pentane-ethyl acetate (7:3 v/v) as eluant. In the same manner as previously described, sterols were identified as their trimethylsilyl ethers by means of gas chromatography-mass spectrometry (gc-ms) technique (3).

Sterol analyses were made on free sterols; a 5% OV-1 column at 280°C was used for quantitative determination (4).

RESULTS AND DISCUSSION

The tetrasporophyte contains significantly greater concentrations of total sterols than the gametophyte. Their amounts increase during fructification of the gametophyte and slowly decrease during appearance of tetraspores for the sporophytic plant (fig. 1).

Table 1 shows the estimation of each sterol.¹ Cholesterol 2 is the major sterol in all samples analyzed; desmosterol 3 is abundant only for the gametophyte and

¹For the identification of these sterols by gc-ms, see previous report (4).

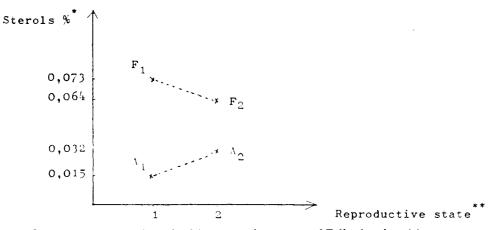


FIGURE 1. Amount of sterols of Asparagopsis armata and Falkenbergia rufolanosa. *Based on dry weight of algae. **1 sterile plant 2 frunctificated plant.

22-dehvdro-cholesterol 1 for the sterile *Asparagopsis armata*. The three diols 5, 6, 8 are present in both gametophytic and sporphytic plants.

Samples	Sterols** identified as $\%$ of total sterols analyzed								
	1	2	3	4	5	6	7	8	
$\begin{array}{c} \mathbf{A}_1 \\ \mathbf{A}_2 \\ \mathbf{F}_1 \\ \mathbf{F}_2 \end{array}$	$\frac{7}{2}$ $\frac{4}{3}$	$70 \\ 84 \\ 81 \\ 84$	$\begin{array}{c} \frac{4}{7} \\ 1 \\ 1 \\ 1 \end{array}$		$\begin{array}{c}4\\2\\5\\6\end{array}$	4 1 1 1	$\begin{array}{c}1\\1\\4\\1\end{array}$		

TABLE 2. Estimation of sterols.

**1, 22-dehydro-cholesterol; 2, cholesterol; 3, desmosterol; 4, brassicasterol; 5, 25-hydroxy-cholesterol; 6, 25-hydroxy-24-methylcholesterol; 7, fucosterol+ β situaterol; 8a, liagosterol and 8b, cholesta-5,25-dien-33, 24-diol (4).

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