

Biosynthesis and Interconversion of Phytoecdysones in *Sesuvium Portulacastrum* L.†

By A. T. SIPAHIMALANI, A. BANERJI,* and M. S. CHADHA

(*Bio-Organic Division, Bhabha Atomic Research Centre, Bombay-85, India*)

Summary Incorporation of mevalonic acid and cholesterol into ecdysone and ecdysterone and conversion of ecdysone into ecdysterone in the plant, *Sesuvium portulacastrum* L., have been demonstrated.

THE biosynthetic and metabolic transformations of ecdysones have been studied in several insects, but investigations on the biosynthesis in plants are few and restricted to lower plants.¹ The co-occurrence of ecdysone and ecdysterone in

† Presented in part at the 8th International Symposium on the Chemistry of Natural Products, New Delhi, 1972.

a higher plant, *viz.* *Sesuvium portulacastrum* L. (N.O. Aizoaceae),² presented an attractive possibility for a study of their biosynthesis and interconversion. We report here the incorporation of mevalonic acid and cholesterol into ecdysone and ecdysterone and also conversion of ecdysone into ecdysterone in *S. portulacastrum* L.

The labelled substrates (in aqueous or 10% aqueous ethanolic solution) were administered to the plant by the wick method. The plants were harvested after two weeks and worked up for the isolation of ecdysones.³ Ecdysone and ecdysterone could be separated by preparative t.l.c. The two compounds were identified by u.v. and t.l.c. com-

ecdysterone. Incorporation was rather low. Since Δ^4 -3-ketones are known to be intermediates in the transformation of Δ^6 - to 5β H steroids in animals⁴ as well as in plants,⁵ the possible incorporation of Δ^4 -[³H]cholesten-3-one into ecdysone and ecdysterone was explored. The absence of incorporation in the two phytoecdysones suggests that a different biosynthetic mechanism is operating in *S. portulacastrum* L.

In the insects, at least two modes of metabolic transformation of ecdysones have been reported. The experiments of Carlisle and Ellis on locusts suggest that ecdysone is deactivated by dehydroxylation.⁶ On the other hand,

Substrate	Activity given			Products				
	Total activity d.p.m.	Sp. activity d.p.m./mm	d.p.m.	Ecdysterone % Incorp.	Sp. activity d.p.m./mm	d.p.m.	Ecdysone % Incorp.	Sp. activity d.p.m./mm
Sodium [2- ¹⁴ C]acetate	5.5×10^8	5.1×10^{10}	2.6×10^4	0.005	3.8×10^8	3.1×10^3	0.0006	4.6×10^6
Mevalonic acid [2- ¹⁴ C]lactone	2.2×10^8	1.1×10^{10}	2.4×10^6	1.1	9.6×10^7	2.1×10^5	0.1	1.4×10^8
[4- ¹⁴ C]Cholesterol	1.1×10^8	1.2×10^{11}	1.0×10^4	0.009	1.4×10^5	5.2×10^3	0.005	1.3×10^6
[U- ³ H]- Δ^4 -Cholesten-3-one	5.7×10^8	3.1×10^{11}	No activity	—	—	No activity	—	—
[U- ³ H]Ecdysone	1.3×10^9	1.7×10^{12}	1.4×10^7	1.1	1.3×10^9	4.0×10^7	3.1	1.2×10^{10}
[U- ³ H]Ecdysterone	1.3×10^9	2.0×10^{13}	2.5×10^8	19.2	6.8×10^9	No activity	—	—

parisons with authentic samples.² Further, their acetates (Ac₂O-pyridine; 16 h at 30°) were identical with authentic samples. Radiochemical purity was checked by isotopic dilution.

In the first instance two general precursors, *viz.* [2-¹⁴C]-acetate and -mevalonic acid were fed to *S. portulacastrum* L.; ecdysterone and ecdysone were isolated and found to possess radioactivity. Plants administered with [4-¹⁴C]-cholesterol were also found to give labelled ecdysone and conversion of ecdysone into ecdysterone in arthropods pro-

vides evidence for hydroxylation at C-20 position.⁷ In our experiment clear evidence for the hydroxylation of C-20 was obtained as [³H]ecdysone was efficiently converted into [³H]ecdysterone. The possibility of equilibrium between ecdysone and ecdysterone is ruled out since ecdysone obtained from [³H]ecdysterone administered *S. portulacastrum* was inactive.

(Received, 20th March 1972; Com. 460.)

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