Nomenclature Correction of Certain Plant n-Alkanes

Sir:

Recently we initiated phytochemical studies on certain Euphorbia species because of the reported folkloric use of members of this genus (E. ingens, E. bupleurifolia, E. clavarioides, and E. gorgonis) as anticancer remedies (1). Indeed, certain of the euphorbias (E. amygdaloides, E. drummondii, E. marginata, E. pilulifera, and E. resinifera) have been shown to elicit antitumor activity in laboratory animals (2). During the course of one of our investigations, a considerable amount of waxy material previously described (8). Of the five samples analyzed, only the n-nonacosane from C. bulbosum (3) was reasonably pure (about 97%). The *n*nonacosane from E. watanabei (4) was actually 61% *n*-hentriacontane, and the *n*-nonacosane from A. patula (5, 8) was 60% n-hentriacontane. Samples of n-hentriacontane isolated from E. lathyris (6) and P. asiatica were complex mixtures, but each was more than 50% of the labeled hydrocarbon (Table I).

These studies re-emphasize the need for subjecting all *n*-alkanes isolated from plant sources to a gas chromatographic and/or mass spectrometric analysis before any identification of these compounds can be made, and their presence in plant material reported.

TABLE I-COMPOSITION OF n-ALKANES FROM SELECTED PLANTS

		n-Alkanes Found ^a										
n-Alkane Reported	Plant Source	Ref.	C25	C26	C27	C28	C29	C_{30}	C31	C32	C 33	C34
n-Nonacosane	Chaerophyllum bulbosum L.	(3)			1	2	97					
n-Nonacosane	Euphorbia watanabei Makino	(4)	< 1	1	3	4	20	5	61		5	
n-Nonacosane	Arctostaphylos patula Greene	(5)	<1	< 1	3	4	17	5	60	6	5	
<i>n</i> -Hentriacontane	Euphorbia lathyris L.	(6)			2	2	17	6	53	5	15	
<i>n</i> -Hentriacontane	Plântago asiatica L.	(7)	<1	1	2	1	16	6	61	6	3	5

^a As determined by gas chromatography (8), expressed to the nearest 1.0%. Estimations were made by planimetry.

was encountered, which readily deposited white plates from the usual crystallizing solvents. These plates gave a sharp melting point, indicative of either n-nonacosane or n-hentriacontane, and elemental analyses were in agreement with an identification as either of these *n*-alkanes. This prompted us to request samples of these two compounds from other investigators for comparative purposes. Samples of n-nonacosane isolated from Chaerophyllum bulbosum L. (3), Euphorbia watanabei Makino¹ (4), and Arctostaphylos patula Greene¹ (5), as well as n-hentriacontane from Euphorbia lathyris L. (6) and Plantago asiatica L. (7) were obtained.

Prior to using these samples for comparative purposes, they were each subjected to a gas chromatographic analysis using the conditions Watt, J. M., and Breyer-Brandwijk, M. G., "The Medicinal and Poisonous Plants of Southern and Eastern Africa," 2nd ed., E.&S. Livingstone Ltd., Edinburgh, Scotland, and London, England, 1962.
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¹ Tentative identifications, based solely on melting points, were reported in the original papers (4, 5).