

## Studies in the Flavin Series. Part XVII.<sup>1</sup>

By **S. Ghisla** and **P. Hemmerich**,\* *Fachbereich Biologie, University of Konstanz, Germany*  
**C. Jefcoate**, Department of Biochemistry, Edinburgh University Medical School, Teviot Place, Edinburgh EH8 9AG

The distribution of products from the reductive alkylation of 1,3,7,8-tetramethylalloxazine depends upon whether saturated (hard) or unsaturated (soft) alkylating agents are used. The former yield 5-alkylated 5,10-dihydroalloxazines and, under more drastic conditions, 5,5-dialkylated derivatives, while the latter (benzyl and allyl bromide) yield 4a-monoalkylated and 4a,5-dialkylated derivatives. The 5-alkyldihydroalloxazines are readily oxidised to the 5-alkylalloxazinium salts, which either undergo hydrolytic dealkylation to the initial alloxazine or, if the 5-substituent is allyl or benzyl, in part rearrange to a 6-alkylalloxazine. Oxygen, under acidic conditions, converts 4a-benzyl- (or allyl)-dihydroalloxazines into the 6-benzyl- (or allyl-)alloxazine *via* the 5-substituted alloxazinium salt. The mechanisms of these reactions are discussed and their relevance to group transfer reactions of flavocoenzymes is stressed.

PREVIOUS papers have dealt with the alkylations of three classes of (iso)alloxazines:† (i) 1,5-dihydroisoalloxazines (flavohydroquinones);<sup>1-4</sup> (ii) 5-acetyl-1,5-dihydroisoalloxazines;<sup>1,5,6</sup> and (iii) 5-acetyl-1,5-dihydroalloxazines.<sup>1,6</sup> In the present paper the alkylations of 5,10-dihydroalloxazines [N(5) unprotected] are described.

Previous results indicated that the alkylations of dihydro(iso)alloxazines † can be described well by competition between frontier orbital and charge control effects.<sup>1,7</sup> The atoms N(1), N(3), O(2), and O(4) operate as charge controlled sites, C(4a) and N(5) as frontier orbital controlled sites, while N(10) is probably intermediate in character. Thus the orientation of alkylation is determined by the hardness<sup>1</sup> of the alkylation agent and by the substituents at the dihydro(iso)alloxazine nucleus. The effect of acetylation at N(5) is particularly marked since the most reactive frontier orbital position is then eliminated. Recently, the remarkable mobility of alkyl groups at the dihydro(iso)alloxazine nucleus [at N(5), N(10), and C(4a)] has become apparent.<sup>1-3</sup> Thus 1,2-(5 → 4a) and 1,3-(10 → 4a) alkyl rearrangements have been observed together with photo-activated oxidation of active 4a-substituents by molecular oxygen. We now report further unusual rearrangements of alkylated alloxazines.

### RESULTS

*Alkylations.*—The 5,10-dihydroalloxazine (2) was prepared *in situ* by reduction of 1,3-dimethyl-lumichrome (1) with alkaline sodium dithionite and alkylations were carried out anaerobically with dimethyl sulphate, diethyl sulphate, allyl bromide, and benzyl bromide.

The reactions with saturated alkylating agents yielded mixtures of 5-monoalkylated dihydroalloxazines (3) and 5,5-dialkylated dihydroalloxazines (4) and could be performed so that either class of products predominated. The reactions with unsaturated ('activated') alkylating agents (R = benzyl or allyl) yielded mixtures of 4a-mono-

alkylated dihydroalloxazines (5) and 4a,5-dialkylated dihydroalloxazines (8) (see Scheme 1).

The structure of these compounds was ascertained as follows. The 5-monoalkylated compounds (3) are easily oxidised in acidic solution to yield alloxazinium salts (6) ( $\lambda_{\max}$  455 and 398 nm). These are dealkylated upon raising the pH to 6 to yield dimethyl-lumichrome (1) quantitatively (see Figure 1). This dealkylation proceeds without the intermediate formation of a 4a-hydroxy-adduct (pseudo-base), which has been observed in the isoalloxazine series.<sup>3</sup> The 5,5-dialkylated compounds (4) were isolated as the quaternary salts ( $\lambda_{\max}$  315 and 285 nm) which yielded mesoionic compounds (7) ( $\lambda_{\max}$  345 and 305 nm) upon neutralization [deprotonation at N(10)], with a *pK* of 6.6 (Figure 2). The salts as well as the mesoionic compounds were stable to oxygen. The 5,5-disubstituted compounds (4) were easily converted into the 5-mono-substituted alloxazinium salts (6) by oxidation with sodium nitrite in acid solution. Under more drastic conditions of oxidation (6N-HCl-NaNO<sub>2</sub>, 90°) the salts (6) were converted into dimethyl-lumichrome (1). The n.m.r. spectra of the alkylated compound (4; R = Me) show *one* sharp signal corresponding to two equivalent methyl groups, which indicates double substitution at a cationic centre. In the case of the diethyl compound (4; R = Et) the n.m.r. spectrum shows an ABX signal due to the non-equivalence of the ethylenic protons attached to a diastereoisotopic centre N(5). A more extensive discussion of this has been given for the analogous isoalloxazine compound.<sup>4</sup>

The 4a-mono-substituted compounds (5; R = benzyl and allyl) were similar to the 4a-substituted dihydroisoalloxazine analogues<sup>2,3</sup> and were clearly identified from the n.m.r. and absorption spectra ( $\lambda_{\max}$  360 nm). Both 4a-alkylated dihydroalloxazines (5; R = benzyl and allyl) were easily photolysed by light in the presence of oxygen to yield 1,3-dimethylalloxazine (1) quantitatively, while the 4a,5-dialkylated products (8; R = allyl and benzyl) were stable. The 4a-substituted dihydroalloxazines (5) were not further alkylated under the standard alkylation

<sup>3</sup> W. H. Walker, P. Hemmerich, and V. Massey, *European J. Biochem.*, 1970, **13**, 258.

<sup>4</sup> P. Hemmerich, S. Ghisla, U. Hartmann, and F. Müller in 'Flavins and Flavoproteins,' ed. E. C. Slater, University Park Press, Baltimore, 1970.

<sup>5</sup> P. Hemmerich, B. Priejs, and H. Erlenmeyer, *Helv. Chim. Acta*, 1960, **43**, 372.

<sup>6</sup> K. H. Dudley and P. Hemmerich, *Helv. Chim. Acta*, 1960, **50**, 355.

<sup>7</sup> G. Klopman, *J. Amer. Chem. Soc.*, 1968, **90**, 223.

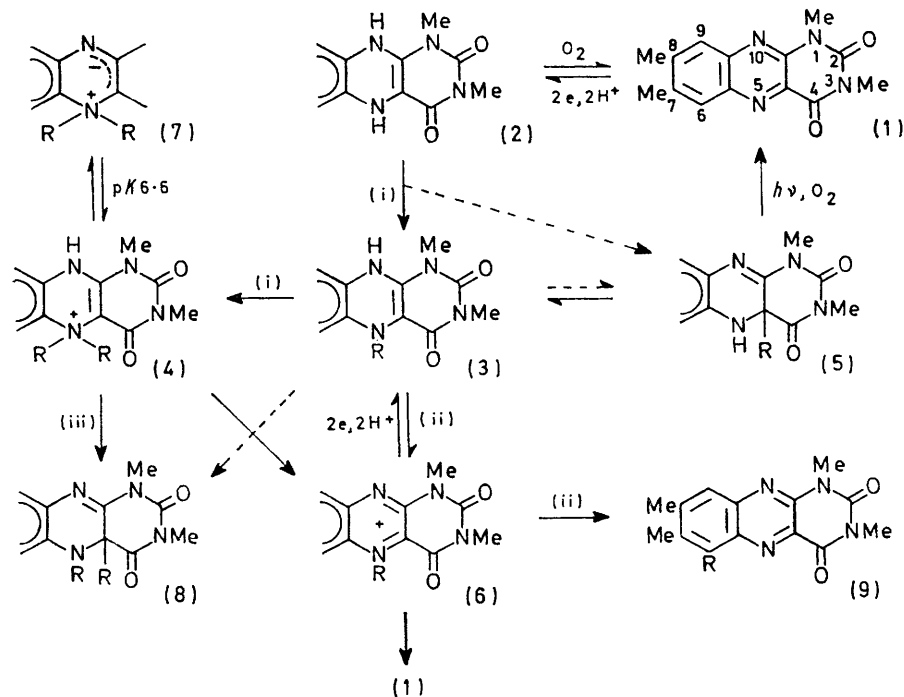
† Dihydro(iso)alloxazine has been used throughout this paper where statements apply to both dihydroisoalloxazines and to dihydroalloxazines.

<sup>1</sup> Part XVI, C. R. Jefcoate, S. Ghisla, and P. Hemmerich, *J. Chem. Soc. (C)*, 1971, 1689.

<sup>2</sup> W. H. Walker, P. Hemmerich, and V. Massey, *Helv. Chim. Acta*, 1967, **50**, 2269.

conditions and thus the 4a,5-dialkylated dihydroalloxazines (8), which were obtained from reductive benzylation and allylation of (1), were not formed by further alkylation of the 4a-alkylated product (5).

100 °C a yellow precipitate slowly formed, the spectral properties of which suggested extensive rearrangement. The i.r.  $[\nu(\text{CO})]$  and absorption spectra ( $\lambda_{\text{max}}$ , 385 and 351 nm) for (5; R = allyl) indicated a chromophore analogous



SCHEME 1 Reagents: (i), NaOH-RX; (ii),  $\text{O}_2$ -6N-HCl; (iii), heat

**Alkyl Migration.**—When the 4a-benzylidihydro- or 4a-allyldihydro-alloxazine (5; R = benzyl or allyl) was dissolved in an aqueous mineral acid-ethanol mixture at

to that of alloxazine (1). The following experiments have indicated that these rearrangement products were 6-substituted alloxazines (9; R = benzyl or allyl).

The n.m.r. data shown in Table 1 (*cf.* refs. 8 and 9) suggest that the rearrangement product is a 6-substituted

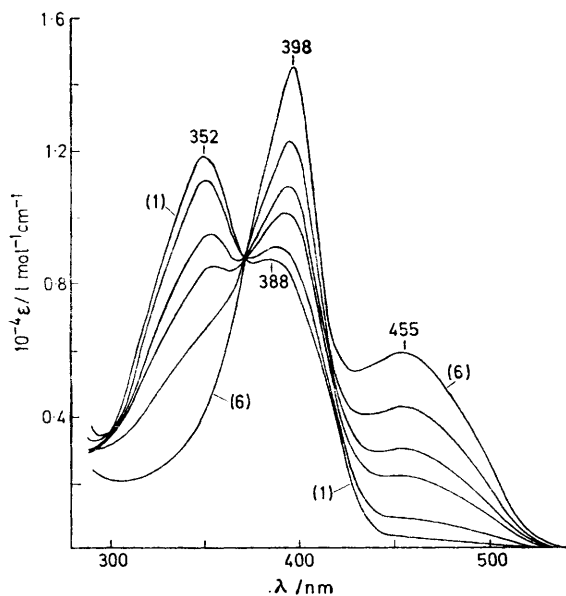


FIGURE 1 Direct hydrolysis of the alloxazinium salt (6; R = Me) to 1,3-dimethyl-lumichrome (1) in aqueous solution at 24° and pH 7, half-time 6 min.

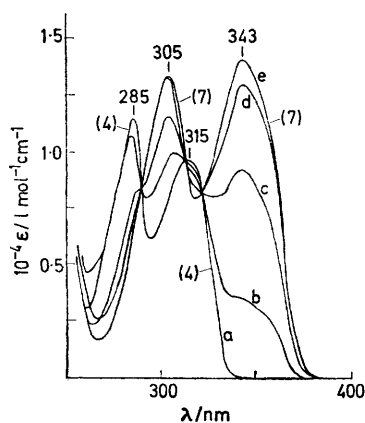


FIGURE 2 Absorption spectra of the conversion of the quaternary dihydroalloxazine salt (4) into the mesoionic form (7) in 0.1M-aqueous buffers: (a) sodium sulphate pH 2.0; (b) potassium phosphate pH 6.0; (c) pH 7.0; (d) pH 7.6; and (e) sodium borate pH 9.0

<sup>8</sup> J. Lauterwein and P. Hemmerich, unpublished results.

<sup>9</sup> F. J. Bullock and O. Jardetzky, *J. Org. Chem.*, 1965, **30**, 2056.

alloxazine. Previous work with deuterium-labelled isoalloxazines has indicated that the chemical shift of the 7-Me protons is upfield from that of the 9-Me protons. The signals due to 6-H, 9-H, 7-Me, and 8-Me were assigned by assuming that these signals retained the same relative shifts as in the isoalloxazine series. Comparison of the n.m.r. spectra of the allyl and benzyl rearrangement

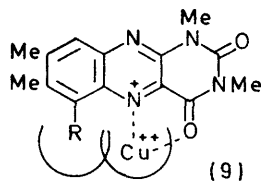
TABLE I  
Characteristics of n.m.r. spectra of (iso)alloxazine derivatives

Compound	Solvent	$\delta/p.p.m.$			
		7-Me	8-Me	9-H	6-H
1,3-Dibenzyl-7,8-dimethylalloxazine <sup>a</sup>	CDCl <sub>3</sub>	ca.	ca.	7.78	8.00
6-Allyl-1,3,7,8-tetramethylalloxazine (9)	CDCl <sub>3</sub>	2.48	2.58	7.75	
6-Benzyl-1,3,7,8-tetramethylalloxazine (9)	CDCl <sub>3</sub>	2.43	2.52	7.76	
9-Deuterio-7,8-dimethylisoalloxazine <sup>a</sup>	0.2M-NaOD	2.23	2.30	(6.89) <sup>b</sup>	7.08
8-Trideuterioflavin mononucleotide <sup>a</sup>	D <sub>2</sub> O <sup>c</sup>	2.30	(2.47) <sup>b</sup>	7.39	7.68

<sup>a</sup> Synthesised according to reference 5. <sup>b</sup>  $\delta$ -values of the corresponding <sup>1</sup>H compounds. <sup>c</sup> pH 7.

products with the spectrum of 1,3-dibenzyl-7,8-dimethylalloxazine suggests the probable loss of the 6-proton resonance for both rearrangement products, implying that the 6-proton has been replaced by allyl or benzyl substituents. Furthermore the 8-Me group appeared as a doublet ( $J$  2Hz) due to 1,3-coupling with 9-H,<sup>10</sup> while the 7-Me group showed a sharp singlet, which is consistent with the substitution at C(6). Clearly the small separations of the 6- and 9-proton resonances and the 7- and 8-Me resonances make these assignments uncertain.

However, confirmatory evidence that the benzyl or allyl group had migrated to the 6- not to the 9-position, was provided by the observation that alloxazine (9) did



SCHEME 2

not complex with Cu<sup>2+</sup> ions (Figure 3), even in the presence of a 10-fold excess of Cu<sup>2+</sup> ions. In the control experiment, the alloxazine (1) (6-position unsubstituted) and its 9-bromo-derivative both showed a new absorption band ( $\lambda_{max}$  410 nm) due to unhindered complexing of Cu<sup>2+</sup> at the 4- and 5-position (Scheme 2).<sup>8,11</sup> The analogous 6-allylalloxazine (9; R = allyl), which was isolated by oxidation of the dihydroalloxazine (5; R = allyl) with oxygen in acid solution, also failed to complex with Cu<sup>2+</sup> ions.

The same alloxazine (9; R = benzyl) has been synthesised by an alternative procedure which gives more insight into the mechanism of the rearrangement. When

the 4a-benzylidihydroalloxazine (5; R = benzyl) was treated in acetic acid with a platinum-silica catalyst under hydrogen, the absorption spectrum of the product

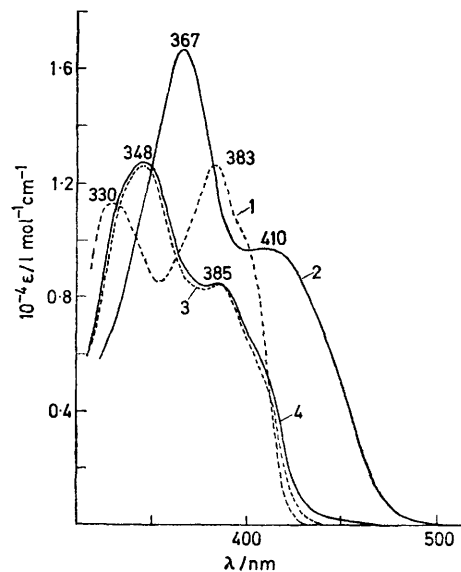


FIGURE 3 Effect of substitution at position 6 on complex formation by alloxazines with Cu<sup>2+</sup> in acetone. Spectrum 1 and spectrum 3: the absorption spectra of 1,3-dimethyl-lumichrome (1) and 6-benzyl-1,3,7,8-tetramethylalloxazine (9) in absence of Cu<sup>2+</sup>. Spectrum 2: The spectrum of 1,3-dimethyl-lumichrome (1) after addition of 2.0 mol. equiv. Cu(ClO<sub>4</sub>)<sub>2</sub>. Spectrum 4: The spectrum of the alloxazine (9) after addition of 2.0 mol. equiv. Cu(ClO<sub>4</sub>)<sub>2</sub>.

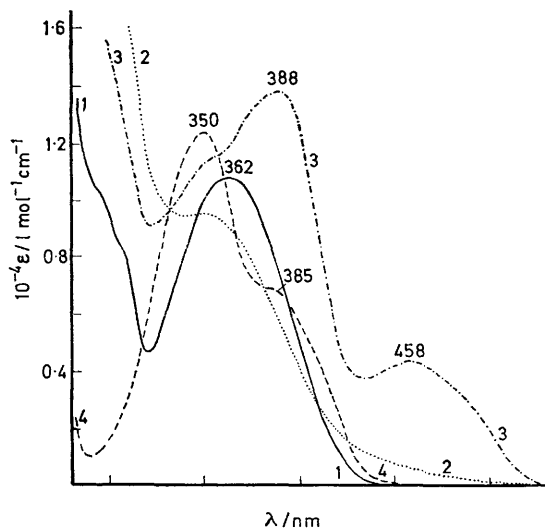


FIGURE 4 Isomerisation of 4a-benzylidihydroalloxazine (5) to 6-benzylalloxazine (9) followed by means of absorption spectra. Absorption spectrum of (5) in acetic acid (spectrum 1) and after reduction by H<sub>2</sub>-Pt (spectrum 2). The spectrum is typical of 5-substituted dihydroalloxazines;<sup>12</sup> probably a mixture of 5-benzyl-5,10-dihydroalloxazine (3) and 5,10-dihydroalloxazine (2). Spectrum 3: immediately after admission of oxygen showing instantaneous formation of a 5-substituted alloxazinium salt (6) (cf. Figure 1). Spectrum 4: 6-benzylalloxazine (9) after separation from lumichrome (1).

The total reaction sequence is (5)  $\xrightarrow{O_2}$  {(2) + (3)}  $\xrightarrow{O_2}$  {(1) + (6)}  $\xrightarrow{O_2}$  {(1) + (9)}

<sup>10</sup> M. Barfield and B. Chakrabarti, *Chem. Rev.*, 1969, **69**, 757.

<sup>11</sup> F. Müller, P. Hemmerich, and A. Ehrenberg, *European J. Biochem.*, 1968, **5**, 158; P. Bamberg and P. Hemmerich, *Helv. Chim. Acta*, 1961, **44**, 1001.

changed towards that of the 5,10-dihydroalloxazine structure (4; R = benzyl) (Figure 4).<sup>12</sup> Admission of oxygen immediately produced a new absorption at 458 nm, which then decreased over the next 30 min. These spectral changes suggest that the 5-benzylalloxazinium salt (6; R = benzyl) is an intermediate [for comparison Figure 1 shows the spectrum of (6; R = Me)]. T.l.c. showed two spots correspondingly to the 1,3-dimethylalloxazine (1) and the 6-benzyl-1,3-dimethylalloxazine (9; R = benzyl), both of which were isolated and identified.

#### DISCUSSION

The alkylations in the dihydroalloxazine series (see Table 2) are largely analogous to the corresponding reactions in the dihydroisoalloxazine series.<sup>4-6</sup> It was previously suggested<sup>1</sup> that this distinction between

exclude the possibility that the 4a,5-dialkyldihydroalloxazines (8; R = benzyl or allyl) were formed by a direct alkylation at C(4a) of the 5-monoalkylated intermediates (3; R = benzyl or allyl).

The proportion of the 4a-alkylated dihydroalloxazine (5; R = benzyl or allyl) which is formed respectively by direct alkylation at C(4a) and by a rearrangement of an initial N(5) isomer, has not been determined. The 5-alkylated dihydroalloxazines (3; R = benzyl or allyl) have not been detected, even though the dihydroisoalloxazine analogues have been isolated. However even the latter readily undergo 5 → 4a migration at 50 °C.<sup>2</sup> On the other hand saturated alkyl substituents do not show this 5 → 4a migration in the dihydroalloxazine series, while migration occurs under more drastic conditions in the iso-series.<sup>1</sup>

TABLE 2

Reductive monoalkylation of (iso)alloxazines

Prealkylated positions	Protective group at N(5)	Alkylating agent	Alkylated product yield (%)				Ref.
			N(5)	N(10)	C(4a)	O(2)	
Isoalloxazine (Flavin)	None	MeI—Me <sub>2</sub> SO <sub>4</sub>	80—100 *		0—20 *		4
	None	PhCH <sub>2</sub> Br—CH <sub>2</sub> :CH·CH <sub>2</sub> Br			>95		4
	Acetyl	MeI				>95	1,5,6
	Acetyl	PhCH <sub>2</sub> Br—CH <sub>2</sub> :CH·CH <sub>2</sub> Br				>95	1,5,6
Alloxazine (Lumichrome)	None	Me <sub>2</sub> SO <sub>4</sub>	>95				
	None	PhCH <sub>2</sub> Br—CH <sub>2</sub> :CH <sub>2</sub> Br		95	5		1,6
	Acetyl	MeI—Me <sub>2</sub> SO <sub>4</sub>		30	70		1
	Acetyl	PhCH <sub>2</sub> Br—CH <sub>2</sub> :CH·CH <sub>2</sub> Br		70	30		

\* Product rate depending on polarity of the solvent used.

hard (alkyl) and soft (benzyl, allyl) alkylating agents was due to an increase in hardness of N(5) compared to C(4a), which was caused by the bending of the molecule about the N(5)—N(10) axis. However, the benzylation and allylation of the dihydroalloxazine (2) were complicated by the ease of alkyl migration between N(5) and C(4a). The 5-benzylalloxazine (3; R = benzyl) was established as an intermediate in benzylation of (2) by the isolation of the 4a,5-dibenzyl-dihydroalloxazine (8; R = benzyl) from the reaction. Since the 4a-benzylalloxazine (5; R = benzyl) could not be further benzylated to this product, the 4a,5-dibenzyl compound (8; R = benzyl) must have been formed by alkylation of the 5-benzyl intermediate (3; R = benzyl), either directly or more probably by rearrangement of an intermediate dibenzyl quarternary salt (4; R = benzyl). The dialkyl quarternary salt (4; R = Et or Me), which was isolated from reductive alkylation of (1), rearranged at 150° to the 4a,5-dialkyldihydroalloxazine (8; R = Et or Me). Since these alkyl substituents have a far lower migratory aptitude than benzyl or allyl substituents, we may conclude that the dibenzyl or diallyl quarternary salts (4; R = benzyl allyl) will rearrange rapidly, even at the lower temperatures of the alkylation (20 °C). However, we cannot

The 5,5-dialkyldihydroalloxazine salts (4) were readily characterised by the pH sensitivity of the absorption spectra (Figure 2). The spectral changes indicated a pK of 6.6 for loss of 10-H and formation of a zwitter-ion. The unusually high acidity of the 10-proton can in part be attributed to the electron-withdrawing effect of the quarternary nitrogen group.

The alloxazinium salts (6; R = allyl or benzyl) appear to be formed by acid oxidation of 4α-alkyldihydroalloxazines (5; R = allyl or benzyl), but are only intermediates, since they either further rearrange to the 6-alkylalloxazine (9; R = allyl or benzyl) or undergo hydrolysis to alloxazine (1). The 4a → 5 rearrangement requires an initial oxidation step, since acid, in the absence of oxygen, did not isomerize (5) into (4). The most likely sequence for the rearrangement is that (5) is first oxidised to an unstable 4a-substituted alloxazine radical, which then rearranges to a more stable 5-substituted alloxazine radical.<sup>13</sup> A further oxidation of the 5-substituted radical then produces the alloxazinium salt (6).

In contrast to the 'iso'-series,<sup>3</sup> a stable 'pseudo-base' is not formed upon neutralization of the alloxazinium salt (6), and instead immediate dealkylation occurs yielding 1,3-dimethyl-lumichrome (1) (cf. Figure 1).

<sup>12</sup> K. H. Dudley, A. Ehrenberg, P. Hemmerich, and F. Müller, *Helv. Chim. Acta*, 1964, **17**, 1354.

<sup>13</sup> F. Müller, P. Hemmerich, A. Ehrenberg, G. Palmer, and V. Massey, *European J. Biochem.*, 1970, **14**, 185.

This is similar to a known reaction in the phenazine series.<sup>14</sup> Thus, alloxazinium salts (6) can function as strong alkyl donors to appropriate nucleophiles. This behaviour may provide a model for flavin-dependent biological dehydrogenations, as well as hydrofolate-dependent monocarbon-transfer.<sup>4, 15, 16</sup>

#### EXPERIMENTAL

M.p.s were determined with a Kofler hot-stage apparatus and are corrected. Thin layer chromatography (t.l.c.) was carried out on MN-Silicagel S (Macherey, Nagel Co., Düren, Germany) with chloroform-ethanol (7:3, v/v). Absorption spectra were measured with a Cary 14 spectrophotometer. I.r. spectra were taken on a Beckman IR 8 or on a Perkin-Elmer 621 and <sup>1</sup>H n.m.r. spectra on a Varian A-60 A instrument.

*General Procedures for Reductive Alkylation of 1,3-Dimethyl-lumichrome (1).*—A stirred suspension of 1,3-dimethyl-lumichrome (1) (1.0 g) in a mixture of ethanol (50 ml) and aqueous 2N-sodium hydroxide (40 ml) was reduced anaerobically with a small excess of solid sodium dithionite. The alkylating agent (10 equiv. in 10 ml ethanol) was added to the reddish yellow solution over 30 min. The reaction was followed by t.l.c. and at completion the products were isolated as described below.

(a) *Methylation with dimethyl sulphate.* During the reaction a yellow precipitate of the sodium salt of 5,10-dihydro-1,3,5,7,8-pentamethylalloxazine (3; R = Me) gradually formed, which dissolved completely on acidification with acetic acid. After evaporation of the ethanol *in vacuo*, 2N-perchloric acid (50 ml) and solid sodium perchlorate (2 g) were added and the mixture was oxidised with solid sodium nitrite (500 mg). An orange product was formed, filtered off, and subsequently washed with water, methanol, chloroform, and ether, and dried *in vacuo*, to give 1,3,5,7,8-pentamethylalloxazinium perchlorate (6; R = Me) (0.75 g), m.p. 238–242° (Found: C, 46.7; H, 4.4; N, 14.5. C<sub>15</sub>H<sub>17</sub>ClN<sub>4</sub>O<sub>6</sub> requires C, 46.8; H, 4.45; N, 14.55%), λ<sub>max</sub> (6N-HCl) 455 and 398 nm (log ε 3.81 and 4.19), λ<sub>max</sub> (KBr) 1725 (4-C=O) and 1095 cm<sup>-1</sup> (ClO<sub>4</sub><sup>-</sup>), δ(CF<sub>3</sub>CO<sub>2</sub>H) 8.19 and 8.3 (6- and 9-H), 5.27 (5-Me), and 2.71 and 2.77 p.p.m. (7- and 8-Me).

The combined washings were concentrated *in vacuo* to yield 5,10-dihydro-1,3,5,7,8-hexamethylalloxazinium perchlorate (4; R = Me) (0.27 g), m.p. 268–270° (from methanol-acetone) (Found: C, 47.95; H, 5.2; N, 13.9. C<sub>16</sub>H<sub>21</sub>ClN<sub>4</sub>O<sub>6</sub> requires C, 47.95; H, 5.3; N, 14.0%), λ<sub>max</sub> (6N-HCl) 315 and 287 nm (log ε 3.97 and 4.07), λ<sub>max</sub> (pH 9) 343 and 305 nm (log ε 4.13 and 4.12).

(b) *Ethylation with diethyl sulphate.* No formation of a precipitate was observed. The mixture was acidified with acetic acid, the ethanol was distilled off under reduced pressure, and the aqueous residue was extracted five times with chloroform. The organic layer was washed twice with 2N-ammonia containing dithionite, with diluted mineral acid, and with a saturated sodium hydrogen carbonate solution, dried (MgSO<sub>4</sub>), and the solvent was distilled off under reduced pressure. The oily residue deposited a white precipitate on treating with chloroform-

isopropyl ether. This product was dissolved in methanol (10 ml), a solution of sodium perchlorate (0.5 g) in 2N-perchloric acid (5 ml) was added, and the methanol was slowly distilled off *in vacuo* until a solid was formed. The crystals were filtered and recrystallized from ethanol-2N-perchloric acid to give 5,5-diethyl-5,10-dihydro-1,3,7,8-tetramethylalloxazinium perchlorate (4; R = Et) (0.6 g), m.p. 235–238°, pK<sub>a</sub> 6.6 (Found: C, 50.3; H, 5.85; N, 13.0. C<sub>20</sub>H<sub>25</sub>ClN<sub>4</sub>O<sub>6</sub> requires C, 50.4; H, 5.9; N, 13.05%), λ<sub>max</sub> (6N-HCl) 315 and 287 nm (log ε 3.97 and 4.07); λ<sub>max</sub> (pH 9) 343 and 305 nm (log ε 4.14 and 4.12).

(c) *Benzylation with benzyl bromide.* The general procedure was repeated in tetrahydrofuran instead of ethanol (two-phase system). The pale yellow mixture was acidified with acetic acid, the tetrahydrofuran was distilled off under reduced pressure, and the residue was extracted thoroughly<sup>6</sup> with chloroform. The organic layer was washed twice with a solution of sodium dithionite in aqueous [N-potassium hydroxide, then with diluted acetic acid, hydrogen carbonate, and water, and was finally dried (MgSO<sub>4</sub>). The solvent was evaporated *in vacuo* and the oily residue crystallised from chloroform-isopropyl ether. A first fraction (690 mg) of pure 4a-benzyl-4a,5-dihydro-1,3,7,8-tetramethylalloxazine (5; R = benzyl) (m.p. 241–245°) was separated by filtration. This compound was identical with an authentic specimen.<sup>1</sup>

The mother liquor was evaporated *in vacuo* and the oily residue was crystallised twice from methanol to yield 4a,5-dibenzyl-4a,5-dihydro-1,3,7,8-tetramethylalloxazine (8; R = benzyl) (520 mg), m.p. 167–168° (from ethanol) (Found: C, 73.7; H, 6.15; N, 12.35. C<sub>28</sub>H<sub>28</sub>N<sub>4</sub>O<sub>2</sub> requires C, 73.4; H, 6.25; N, 12.4%), λ<sub>max</sub> (MeOH), 340 and 305 nm (log ε 3.68 and 3.85), λ<sub>max</sub> (6N-HCl) 355 and 312 nm (log ε 3.60 and 3.95), δ (CDCl<sub>3</sub>) 4.45 (5-PhCH<sub>2</sub>) and 3.00 p.p.m. (4a-PhCH<sub>2</sub>).

(d) *Allylation with allyl bromide.* The crude product was recrystallised from methanol to give pure 4a-allyl-4a,5-dihydro-1,3,7,8-tetramethylalloxazine (5; R = allyl) (300 mg), m.p. 143–146° and 210–230° (decomp.). This compound was identical to an authentic sample.<sup>1</sup>

6-Allyl-1,3,7,8-tetramethylalloxazine (7; R = allyl).—A mixture of 5-acetyl-4a-allyl-4a,5-dihydro-1,3,7,8-tetramethylalloxazine and 5-acetyl-10-allyl-5,10-dihydro-1,3,7,8-tetramethylalloxazine<sup>1</sup> (250 mg) were dissolved in ethanolic 6N-hydrochloric acid (1:1) under a very slow stream of nitrogen at 80°. After 48 h, the starting material had disappeared and formation of a new yellow fluorescent spot on t.l.c. was observed in addition to 1,3-dimethyl-lumichrome (1). The solution was diluted with the same volume of water and left at 0° until completion of crystallisation. Filtration yielded yellow needles of 6-allyl-1,3,7,8-tetramethylalloxazine (9; R = allyl) (20 mg), m.p. 223–226° (Found: C, 65.3; H, 5.85; N, 17.65. C<sub>17</sub>H<sub>18</sub>N<sub>4</sub>O<sub>2</sub> requires C, 65.8; H, 5.85; N, 18.05%), λ<sub>max</sub> (MeOH) 388s, 352, 255, and 222 nm (log ε 3.86, 4.10, 4.67, and 4.58), λ<sub>max</sub> (6N-HCl) 395 and 370 nm (log ε 4.08 and 4.07) ν<sub>max</sub> (CHCl<sub>3</sub>) 1718 (4-C=O) and 1675 (2-C=O), no ν(N-H), δ(CDCl<sub>3</sub>) 7.75 (9-H), 6.4–4.1 (5H, m, allyl residue), and 2.58 p.p.m. (d, J 1 Hz, 8-Me). The mother liquor contained only 1,3-dimethyl-lumichrome (1).

6-Benzyl-1,3,7,8-tetramethylalloxazine (9; R = benzyl).—

<sup>14</sup> 'Heterocyclic Compounds,' ed. R. C. Elderfield, vol. 6, Wiley, New York, 1956, p. 654.

<sup>15</sup> P. Hemmerich, G. Nagelschneider, and C. Veeger, *FEBS Letters*, 1970, **8**, 69.

<sup>16</sup> J. Straviniopoulos and L. Kaenicke, *European J. Biochem.*, 1967, **3**, 95; M. C. Archer and K. G. Scrimgeour, *Canad. J. Biochem.*, 1970, **48**, 278.

The same procedure as for (9; R = allyl) was performed with a mixture of 5-acetyl-4a-benzyl-4 $\alpha$ ,5-dihydro-1,3,7,8-tetramethylalloxazine and 5-acetyl-10-benzyl-5,10-dihydro-1,3,7,8-tetramethylalloxazine<sup>1</sup> (200 mg) to yield pure 6-benzyl-1,3,7,8-tetramethylalloxazine (9; R = benzyl) (80 mg), m.p. 271—275° (Found: C, 69.4; H, 5.7; N, 15.2. C<sub>21</sub>H<sub>20</sub>N<sub>4</sub>O<sub>2</sub> requires C, 70.0; H, 5.6; N, 15.55%),  $\lambda_{\max}$  (MeOH) 385 and 351 nm (log  $\epsilon$  3.86 and 4.11),  $\lambda_{\max}$  (6N-HCl) 395 and 370 nm (log  $\epsilon$  4.08 and 4.07),  $\delta$  (CF<sub>3</sub>CO<sub>2</sub>H) 8.19 (9-H), 4.92 (6-PhCH<sub>2</sub>), 2.78 (8-Me), and 2.52 p.p.m. (7-Me).

6-Benzyl-1,3,7,8-tetramethylalloxazine (9; R = benzyl) (*Alternative Procedure*).—4a-Benzyl-4a,5-dihydro-1,3,7,8-tetramethylalloxazine (5; R = benzyl) (400 mg) was dissolved in acetic acid (50 ml) and hydrogenated over platinum on silica gel. T.l.c. showed two spots corresponding to 1,3-dimethyl-lumichrome (1) (blue fluorescence) and 6-benzyl-1,3,7,8-tetramethylalloxazine (9; R = benzyl) (yellow fluorescence). The latter was separated by careful fractional crystallisation from aqueous acetic acid and from chloroform-isopropyl ether to yield the pure alloxazine (9; R = benzyl) (75 mg), m.p. 270—275°.

5,10-Dihydro-1,3,5,5,7,8-hexamethylalloxazine (7; R = Me).—The perchlorate (4; R = Me) (1.0 g) was dissolved in chloroform (50 ml) and this solution washed twice with a saturated solution of sodium hydrogen carbonate, then with water, was dried (MgSO<sub>4</sub>), and the solvent was distilled off under reduced pressure. The oily residue was treated with tetrahydrofuran-isopropyl ether and the crude product was collected by filtration (m.p. 186—190°).

This compound was recrystallised from chloroform-isopropyl ether to yield 5,10-dihydro-1,3,5,5,7,8-hexamethylalloxazine (7; R = Me) (0.68 g), m.p. 191—193° (Found: C, 63.8; H, 6.75; N, 18.4. C<sub>18</sub>H<sub>20</sub>N<sub>4</sub>O<sub>2</sub> requires C, 64.0; H, 6.7; N, 18.65%),  $\lambda_{\max}$  (MeOH) 343 and 305 nm (log  $\epsilon$  4.13 and 4.12). The corresponding 5,5-diethyl-5,10-dihydro-1,3,7,8-tetramethylalloxazine (7; R = Et) (m.p. 144—145°), which was prepared in the same way, has the same spectral characteristics as the dimethyl compound (7; R = Me).

4a,5-Dihydro-1,3,4a,5,7,8-hexamethylalloxazine (8; R = Me).—The alloxazine (4; R = Me) (0.5 g) in a small glass tube was heated at 190—195° for 30 s. The resulting yellow brown oil showed a single, yellow fluorescent spot on t.l.c. It was dissolved in methanol (2 ml), the solution was treated with charcoal, and water was added until the beginning of turbidity. Yellow-brown crystals separated after standing at 10° overnight and were recrystallised from methanol-water to yield 4a,5-dihydro-1,3,4a,5,7,8-hexamethylalloxazine (8; R = Me) (0.41 g), m.p. 131—133° (Found: C, 64.1; H, 6.7; N, 18.6. C<sub>18</sub>H<sub>20</sub>N<sub>4</sub>O<sub>2</sub> requires C, 64.0; H, 6.7; N, 18.65%),  $\lambda_{\max}$  (EtOH) 375s, 300, 260, and 215 nm (log  $\epsilon$  3.310, 3.890, 4.190, and 4.412),  $\lambda_{\max}$  (pH 7) 303 and 220 nm (log  $\epsilon$  4.000 and 4.392),  $\lambda_{\max}$  (6N-HCl) 311 and 215 nm (log  $\epsilon$  4.050 and 4.375),  $\delta$  (CDCl<sub>3</sub>) 7.08 and 6.88 (6- and 9-H), 3.53 and 3.37 (1- and 3-Me), 2.83 (5-Me), 2.25 (7- and 8-Me), and 1.49 p.p.m. (4a-Me).

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