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longer contained the bands at 1719 and 2735 cm. $^{-1}$  attributable to the aldehyde function.

These results lead to the unequivocal conclusion that the olefinic double bond in gelsemine is present in a vinyl side chain, and not in an exocyclic methylene group as had heretofore been assumed.

THE DIVISION OF PURE CHEMISTRY

NATIONAL RESEARCH COUNCIL OF CANADA LÉO MARION OTTAWA, CANADA K. SARGEANT

RECEIVED AUGUST 22, 1956

## CODEINONE DIMETHYL KETAL AND ITS CONVER-SION TO THEBAINE

Sir:

Thebaine is prodigious among morphine alkaloids for the number and variety of its transformation products.<sup>1</sup> However, no morphine derivative as yet has been converted to thebaine, and it is this conversion we now wish to report.

Ketalization of codeinone (I) with trimethyl orthoformate, methanol, and sulfuric acid was expected to give codeinone dimethyl ketal (II) or neopinone dimethyl ketal (III), but the only iso-



lable product was 8-methoxy- $\Delta^6$ -dihydrothebaine (IV) [m.p. 190–191°;  $[\alpha]^{25}$ D –133° (c, 0.9, ethanol); anal. Calcd. for C<sub>20</sub>H<sub>25</sub>O<sub>4</sub>N: C, 70.0; H, 7.3; OCH<sub>3</sub>, 27.1. Found: C, 70.2; H, 7.1; OCH<sub>3</sub>, 27.5]. The structure of IV was established as follows. Degradation of the methiodide [m.p. 212– 213°;  $[\alpha]^{25}$ D –86° (c, 1, ethanol); anal. Calcd. for C<sub>21</sub>H<sub>28</sub>O<sub>4</sub>NI·H<sub>2</sub>O: C, 50.1; H, 6.0; I, 25.2. Found: C, 50.3; H, 5.8; I, 24.8] gave a methine [m.p. 124–125°,  $[\alpha]^{25}$ D –119° (c, 0.9, ethanol); anal. Calcd. for C<sub>21</sub>H<sub>27</sub>O<sub>4</sub>N: C, 70.6; H, 7.6; OCH<sub>3</sub>, 26.0. Found: C, 70.4; H, 7.5; OCH<sub>3</sub>, 25.6] which was stable to all attempts at alkaline isomerization, indicating the alicyclic double bond was not  $\Delta^7$ . With cyanogen bromide an N-cyanonor compound [m.p. 228–231°;  $[\alpha]^{25}$ D –185° (c, 1, pyridine); anal. Calcd. for C<sub>20</sub>H<sub>22</sub>O<sub>4</sub>N<sub>2</sub>: C, 67.8; H, 6.3; N, 7.9. Found: C, 68.0; H, 6.4; N, 7.9] was obtained, indicating the double bond was not  $\Delta^8$ . Acid hydrolysis gave a mixture of codeinone and 8-methoxydihydrocodeinone [m.p. 195–197°; anal. Calcd. for C<sub>19</sub>H<sub>23</sub>O<sub>4</sub>N: C, 69.3; H, 7.0; OCH<sub>3</sub>, 18.9. Found: C, 69.4; H, 6.8; OCH<sub>3</sub>, 19.4], and hydroxylation with osmium tetroxide gave 7-hydroxy-8-methoxydihydrocodeinone, characterized as the oxime [m.p. 251–253°;  $[\alpha]^{25}$ D –204 (c, 1, pyridine); anal. Calcd. for C<sub>19</sub>H<sub>24</sub>-O<sub>5</sub>N<sub>2</sub>: C, 63.3; H, 6.7; N, 7.8; OCH<sub>3</sub>, 17.2. Found: C, 62.9; H, 6.9; N, 7.9; OCH<sub>3</sub>, 16.9].

An alternative method for preparing codeinone dimethyl ketal was found in the dehydrobromination, using potassium *t*-amylate, of 7-bromodihydrocode inone dimethyl ketal (V), itself prepared by methyl hypobromite addition to  $\Delta^6$ -dihydrothebaine.<sup>2</sup> That the product was code inone dimethyl



ketal (II) [m.p. 138–139°;  $[\alpha]^{21}D - 233°$  (c, 0.5, ethanol); anal. Calcd. for C<sub>20</sub>H<sub>25</sub>O<sub>4</sub>N: C, 70.0; H, 7.3; OCH<sub>3</sub>, 27.1. Found: C, 69.8; H, 7.3; OCH<sub>3</sub>, 27.2] derives from the following reactions. Acid hydrolysis gave codeinone, while hydrogenation led to dihydrocodeinone dimethyl ketal [m.p. 122–123°;  $[\alpha]^{25}D - 151°$  (c, 0.9, ethanol); anal. Calcd. for C<sub>20</sub>H<sub>27</sub>O<sub>4</sub>N: C, 69.6; H, 7.9; OCH<sub>3</sub>, 26.9. Found: C, 69.7; H, 8.0; OCH<sub>3</sub>, 26.5], identical with the product formed from dihydrocodeinone and trimethyl orthoformate, methanol, and acid. The methiodide (m.p. 193–195°) was degraded to a  $\Delta^{7,9}$ -methine [m.p. 71–72°;  $[\alpha]^{25}D - 328°$  (c, 0.9, ethanol);  $\lambda_{\text{ethanol}}^{\text{ethanol}}$ 274 m $\mu$ ,  $\epsilon$  8,500; anal. Calcd. for C<sub>21</sub>H<sub>27</sub>O<sub>4</sub>N: C, 70.6; H, 7.6; OCH<sub>3</sub>, 26.0. Found: C, 70.7; H, 7.5; OCH<sub>3</sub>, 26.2] which was isomerized with alcoholic alkali to a  $\Delta^{8(14),9}$ methine [oil;  $[\alpha]^{25}D + 301°$  (c, 1.2, ethanol);  $\lambda_{\text{max}}^{\text{ethanol}}$ 318 m $\mu$ ,  $\epsilon$  9,000; anal. Found: C, 70.6; H, 7.5; OCH<sub>3</sub>, 25.8].

When codeinone dimethyl ketal was treated with a dried solution of *p*-toluenesulfonic acid in chloroform, there was obtained a 40% yield of thebaine which after crystallization and sublimation was identical in m.p. and mixed m.p. (192–194°) and ultraviolet spectrum ( $\lambda_{max}^{thanol}$  283 m $\mu$ ,  $\epsilon$  7,500;  $\lambda_{min}$ 256 m $\mu$ ,  $\epsilon$  3,700) with an authentic sample.

In a formal sense, this may be considered to constitute a synthesis of thebaine, since the  $\Delta^6$ -dihydrothebaine used in the preparation above can be prepared from dihydrocodeinone,<sup>8</sup> and this in turn is easily made from codeine,<sup>1,4</sup> which has been synthesized.<sup>5</sup> Since thebaine recently has been converted to neopine,<sup>6</sup> the latter also may be considered as synthesized.

(2) We are greatly indebted to Dr. Lyndon F. Small for the details of this reaction. Our bromoketal melted at 116–117°, in agreement with Dr. Small's value.

(3) A. H. Homeyer, J. Org. Chem., 21, 370 (1956).

(4) H. Rapoport, R. Naumann, E. R. Bissell and R. M. Bonner, *ibid.*, **15**, 1103 (1950).

(5) M. Gates and G. Tschudi, THIS JOURNAL, 78, 1380 (1956).
(6) H. Conroy, *ibid.*, 77, 5960 (1955).

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## A NEW ALKYLATION OF CARBONYL COMPOUNDS. II<sup>1</sup> Sir:

We have submitted the new reaction we recently described for the alkylation and acylation of ketones

(1) Part I: G. Stork. R. Terrell and J. Szmuszkovicz, THIS JOURNAL, 76, 2029 (1954).

<sup>(1)</sup> L. F. Small and R. E. Lutz, "Chemistry of the Opium Alkaloids," Suppl. 103, Public Health Reports, U. S. Government Printing Office, Washington, D. C., 1932; K. W. Bentley. "The Chemistry of the Morphine Alkaloids," Clarendon Press, Oxford, England, 1954.

via their pyrrolidine enamines (I) to further scrutiny.

With respect to the introduction of alkyl groups  $\alpha$ to a carbonyl two reactions may be distinguished: The first utilizes alkyl halides, and the second employs reactive  $\alpha,\beta$ -unsaturated nitriles, esters, ketones, etc. (II, X = CN,  $CO_2R$ ,  $O = C - R \dots$ ). The important practical difference between these two reactions is that the undesirable N-alkylation of the enamine is *reversible* in the case of the latter which is therefore the more general reaction.



We have in fact found it of wide applicability to cyclic and acyclic ketones, as well as aldehydes, provided the carbonyl compound has at least one  $\alpha$ methylene group. The following examples will illustrate the usefulness of the new method:<sup>2</sup>

From cyclopentanone<sup>3</sup> were obtained the following alkylated ketones: III (n = 0, R = H):  $R_1 =$  $R_2 = H, X = CN$ ; b.p. 144–147° (13 mm.), in 65% yield with *acrylonitrile*<sup>4a</sup>;  $R_1 = R_2 = H, X =$ CO<sub>2</sub>CH<sub>3</sub>; b.p. 127-130° (11 mm.), in 55% yield with methyl acrylate.4a

From cyclohexanone,<sup>3</sup> the following ketones III (n = 1, R = H) were formed:  $R_1 = R_2 = H$ , X = CN, in 85% yield with *acrylonitrile*<sup>1,4b</sup>;  $R_1 = R_2 = H$ , X = CO<sub>2</sub>CH<sub>3</sub>; b.p. 134–137° (11 mm.), in 65% yield with *methyl acrylate*<sup>4b</sup>;  $R_1 = CH_3$ ,  $R_2 = H$ , X = CO<sub>2</sub>C<sub>2</sub>H<sub>5</sub>; b.p. 165–170° (18 mm.), in 56% yield with *methyl acrylate*<sup>4b</sup>; R = CH<sub>3</sub>, R = CH<sub>3</sub>, R = H, X = CO<sub>2</sub>C<sub>2</sub>H<sub>5</sub>; b.p. 165–170° (18 mm.), in 56% yield with ethyl crotonate<sup>5</sup>;  $R_1 = H$ ,  $R_2 = CH_3$ ,  $X = CO_2CH_3$ ; b.p. 148-150° (18 mm.), in 81% yield with methyl methacrylate.5

From 2-methylcyclohexanone 42% III (n = 1,  $R = CH_3, R_1 = R_2 = H, X = CN$ , b.p. 132–133° (2 mm.), was obtained with acrylonitrile.5,6 Similarly, 2- $\beta$ -cyanoethylcyclohexanone gave, in only

- (3) Via the pyrrolidine enamine.
- (4) Refluxing in dioxane-(a) 16 hours, (b) 3 hours
- (5) Refluxing for 36 hours in dimethylformamide.

fair yield, III (n = 1,  $R = CH_2CH_2CN$ ,  $R_1 =$  $R_2 = H, X = CN$ ), b.p. 190–195° (1 mm.).<sup>5,6</sup> From cycloheptanone could similarly be ob-

tained III  $(n = 2, R = R_1 = R_2 = H, X = CN),$ b.p. 140–145° (10 mm.).

In the same manner, the N-methyl-N-cyclohexyl enamine of 2-heptanone and acrylonitrile4a yielded 45% of 3-β-cyanoethyl -2-heptanone, b.p. 145–147° (11 mm.) while from the enamine of heptaldehyde, acrylonitrile produced 50% of 2- $\beta$ -cyanoethylhept-aldehyde, b.p. 145–148° (12 mm.).<sup>4a</sup>

The reactions of pyrrolidine enamines with  $\alpha,\beta$ unsaturated ketones such as methyl vinyl ketone and its homologs proved especially interesting: The product in this case is initially of type III  $(X = O = C - CH_8)$ , but this intermediate undergoes cyclization and loss of pyrrolidine which then reacts with the resulting octalone (IV) to form its enamine. This relatively stable enamine can be decomposed to IV by *refluxing* with acetic acid-sodium acetate.7 The following examples will give an idea of the scope of this new reaction (equimolar amount in dioxane at room temperature, unless otherwise noted)



From cyclohexanone<sup>3</sup> were obtained: With methyl vinyl ketone (4 hr.), 75% IV (n = 1,  $R = R_1$  $= R_2 = R_3 = H$ ). With methyl isopropenyl ketone (14 hr. at reflux), 66% IV  $(n = 1, R = R_1 = R_3 = H, R_2 = CH_3)$ . With ethyl acetylacrylate (14 hr.), 75% IV  $(n = 1, R = R_1 = R_2 = H, R_3 = CO_2C_2-H_5)$ , m.p.  $50-52^\circ$ .

From 4-hydroxycyclohexanone benzoate<sup>3</sup> and MVK (10 hr. in dimethylformamide) 51% IV (n = 1,  $R = R_2 = R_3 = H$ ,  $R_1 = C_6H_5COO$ , m.p. 110-112°.

From 2-methylcyclohexanone<sup>3</sup> and MVK (14 hr.)

56% IV  $(n = 1, R_1 = R_2 = R_3 = H, R = CH_3)$ .<sup>6</sup> From cyclopentanone<sup>3</sup> and MVK (14 hr.) 42% IV  $(n = 0, R = R_1 = R_2 = R_3 = H)$ .<sup>8</sup>

(7) Cf., F. W. Heyl and M. E. Herr, THIS JOURNAL, 75, 1918 (1953).

(8) Compare the alternate method of synthesis, V. Prelog and M. Zimmermann, Helv. Chim. Acta. 32, 2360 (1945).

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## A NEW RING ENLARGEMENT SEQUENCE

Sir:

We have found a new method of ring enlargement which leads from a ketone to an unsaturated cyclic acid with two more carbon atoms. This is illustrated below with cyclohexanone.

<sup>(2)</sup> Boiling points or melting points are given for all new compounds. Analyses of these and/or their derivatives were satisfactory.

<sup>(6)</sup> Note that alkylation of an  $\alpha$ -substituted cyclohexanone gives further alkylation on the  $\alpha'$ -methylene by this method.