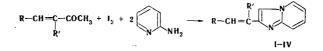
## $\beta$ -SUBSTITUTED 2-VINYLIMIDAZO[1,2-a]PYRIDINES FROM UNSATURATED KETONES

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2-Substituted and 2,3-disubstituted imidazo[1,2-a]pyridines can be obtained by heating alkyl aryl ketones, iodine, and 2-aminopyridine in an organic solvent with subsequent treatment of the resulting  $\beta$ -ketoalkylpyridinium iodides with sodium bicarbonate [1].

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In the present communication, we demonstrate that the use of  $\alpha$ ,  $\beta$ -unsaturated alkyl ketones in this reaction gives the previously undescribed 2-[ $\beta$ -aryl(hetaryl)vinyl[imidazo]1,2-a]pyridines.



The compounds (I-III) obtained by this method are identical to samples prepared by the Chichibabin method [2] by reaction of the appropriate unsaturated halo ketones [3, 4] with 2-aminopyridine. The presence of absorption bands at 700 cm<sup>-1</sup> in the IR spectra of III and IV indicates that they have the cis conformation. The PMR spectrum of III also confirms the assumed structure.

## EXPERIMENTAL

A 10-mmole sample of an  $\alpha$ ,  $\beta$ -unsaturated methyl ketone, 10 mmole of iodine, and 20 mmole of 2aminopyridine were stirred in 100 ml of benzene for 3 h. The benzene solution was then decanted, and the residual mass was heated and treated with 30 g of NaHCO<sub>3</sub> and 500 ml of water. The reaction products were removed by filtration and recrystallized from dimethylformamide (I, II, and IV) or alcohol (III) (see Table 1).

Com- pound	R	R'	Obtained by the new method mp, °C  yield, %		Obtained by the Chichibabin meth- od mp, °C   yield, %		Empirical formula
I	C4H2NO3b	Cl	212—213	80	214—215	33	C <sub>13</sub> H <sub>8</sub> ClN <sub>3</sub> O <sub>3</sub>
II	C4H2NO3b	CH₃	186—188	45	189—191	49	C <sub>14</sub> H <sub>11</sub> N <sub>3</sub> O <sub>3</sub>
III	C6H5	H	152—154	23 c	152—154	46	C <sub>15</sub> H <sub>12</sub> N <sub>2</sub>
IV	4-O2NC6H4	H	250—252	49	—	—	C <sub>15</sub> H <sub>11</sub> N <sub>3</sub> O <sub>2</sub>

TABLE 1. Characteristics of the Compounds Obtained<sup>a</sup>

<sup>a</sup>Satisfactory analytical data were obtained for all of the compounds. <sup>b</sup>5-Nitro-2-furyl. <sup>C</sup>Extracted from the reaction mixture with hot n-octane.

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