

## Catalytic Action of Azolium Salts. II.<sup>1)</sup> Aroylation of 4-Chloroquinazolines with Aromatic Aldehydes Catalyzed by 1,3-Dimethylbenzimidazolium Iodide

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When a mixture of 4-chloroquinazoline (7), an aromatic aldehyde 6, sodium hydride, and a catalytic amount of 1,3-dimethylbenzimidazolium iodide (1) in tetrahydrofuran (THF) was refluxed with stirring for an appropriate time, the chlorine atom of 7 was replaced with the aroyl group, and the 4-aroylequinazolines 10 were obtained in excellent yields. Similar treatments of 4-chloro-2-methylquinazoline (8) and 4-chloro-2-phenylquinazoline (9) led to the 4-aroyle-2-methylquinazolines 11 and the 4-aroyle-2-phenylquinazolines 12, respectively.

Use of *N,N*-dimethylformamide (DMF) instead of THF as the reaction solvent in the above reaction reduced the reaction time and increased the yields of the ketones 10 and 12 as compared with those in THF.

**Keywords** chloroquinazoline; catalytic aroylation; aromatic aldehyde; aroylquinazoline; benzimidazolium iodide

In the previous paper,<sup>1)</sup> we reported that 1,3-dimethylbenzimidazolium iodide (1) is an effective catalyst for the preparation of the 4-aroyle-1 *H*-pyrazolo[3,4-*d*]pyrimidines 4 and 5 by treatment of the 4-chloro-1 *H*-pyrazolo[3,4-*d*]pyrimidines 2 and 3 with aromatic aldehydes 6 and sodium hydride (NaH) in refluxing tetrahydrofuran (THF) or in *N,N*-dimethylformamide (DMF), as shown in Chart 1. By analogy with the benzoin condensation process,<sup>2)</sup> the aroylation using aromatic aldehydes 6, which are the source of the aroyl group, is believed to proceed through the formation of the key intermediate B-2 via the ylide A generated by expulsion of the C<sup>2</sup> hydrogen of 1. It is considered that the important intermediate B-2 is equivalent to the aroyl anion (Ar-C<sup>-</sup>=O). Utilization of the intermediate B-2 for preparation of aroyl compounds by nucleophilic reaction should be a useful method in organic synthesis. We therefore attempted to extend the new aroylation method to 4-chloroquinazolines, and found that the aroylation took place, resulting in the formation of the 4-aroylequinazolines. In the present paper, we describe the aroylations of 4-chloroquinazoline (7),<sup>3,4)</sup> 4-chloro-2-methylquinazoline (8),<sup>5)</sup> and 4-chloro-2-phenylquinazoline (9)<sup>4)</sup> with aromatic aldehydes 6 catalyzed by 1.

In the presence of 1, 4-chloroquinazoline (7) with *p*-fluorobenzaldehyde (6b), *p*-chlorobenzaldehyde (6d), *p*-bromobenzaldehyde (6f), *p*-tolualdehyde (6h), *m*-anisaldehyde (6j), and 2-thiophenecarbaldehyde (6s) underwent nucleo-

philic aroylation in refluxing THF (method A) to give good yields of the expected 4-aroylequinazolines 10b, 10d,<sup>6)</sup> 10f, 10h,<sup>6)</sup> 10j,<sup>6)</sup> and 10s. The 4-chloroquinazoline 7 reacted with benzaldehyde (6g) under the same conditions to give 4-benzoylquinazoline (10g)<sup>6)</sup> as the main product in 96% yield together with 4-benzoyloxyquinazoline (13g) in only 1% yield. A similar result was obtained in the reaction of 7 with 1-naphthaldehyde (6q). Namely, both the ketone 10q and the 4-(1-naphthylmethoxy)quinazoline (13q) were obtained in 28% and 12% yields, when a mixture of 7, 6q, NaH, and a catalytic amount of 1 in THF was refluxed for 4 h. On the other hand, the above reaction for 30 min gave the ketone 10q in only 7% yield and the starting 7 was recovered in 53% yield. The reaction of 7 with *o*-chlorobenzaldehyde (6c) having a substituent at the *ortho* position of 6 in THF led to the ketone 10c,<sup>6)</sup> though the yield was low (21%) because of recovery of the starting 7 in 53% yield. Similarly, in the treatment with *o*-anisaldehyde (6i), starting 7 was recovered in 11% yield together with 35% yield of the ketone 10i.<sup>6)</sup> The low reactivity of *ortho*-substituted benzaldehydes might be caused by steric hindrance of the *ortho*-substituent to the aroylation. The 4-chloroquinazoline 7 was converted into the ketone 10r<sup>6)</sup> in the reaction with 2-furaldehyde (6r) in low yield (47%). On the other hand, an attempt at catalytic aroylation with *p*-nitrobenzaldehyde (6m) and *p*-cyanobenzaldehyde (6n), having a strong electron-withdrawing group at the *para*-position, resulted in recovery of the starting 7 in 67% and 46% yields, respectively. *p*-Nitrobenzoic acid (16m, 4%) and *p*-cyanobenzoic acid (16n, 9%) were formed together in the above reaction. But, the aroylation with *N,N*-dimethylaminobenzaldehyde (6o), having a strong electron-donating group, did proceed, leading to the ketone 10o in 31% yield. In the previous paper,<sup>1)</sup> we reported that use of DMF instead of THF as the reaction solvent for the catalytic aroylation was effective, and the reaction temperature and reaction time required to achieve the aroylation were reduced. Therefore, DMF was used as a solvent for the reaction of 7 with aromatic aldehydes 6, and it was found to reduce the reaction time and increase the yields of the ketones. That is to say, the conversion of the 4-chloroquinazoline 7 into the 4-aroylequinazolines 10h, 10i, and 10r could be accomplished by the reaction with 6h, 6i, and 6r at 80 °C for 5 to

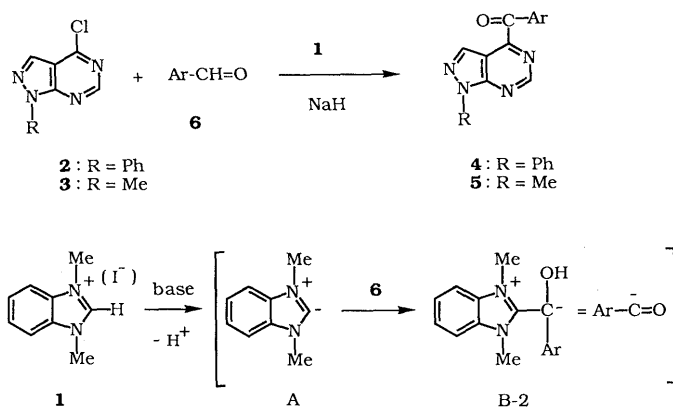


Chart 1

TABLE I. Aroylation of the 4-Chloroquinazolines **7**, **8**, and **9** with Aromatic Aldehydes **6** in the Presence of NaH Catalyzed by **1**

| 4-Chloroquinazoline | <b>6</b>  | Reaction conditions   |            |                                | Products, yield (%) |        |                                            |
|---------------------|-----------|-----------------------|------------|--------------------------------|---------------------|--------|--------------------------------------------|
|                     |           | Solvent <sup>a)</sup> | Time (min) | Temperature <sup>b)</sup> (°C) | Ketone              | Others | Recovery                                   |
| <b>7</b>            | <b>6b</b> | THF                   | 60         | Refl.                          | <b>10b</b>          | 82     | 53                                         |
| <b>7</b>            | <b>6c</b> | THF                   | 60         | Refl.                          | <b>10c</b>          | 21     |                                            |
| <b>7</b>            | <b>6c</b> | DMF                   | 10         | r.t.                           | <b>10c</b>          | 93     |                                            |
| <b>7</b>            | <b>6d</b> | THF                   | 60         | Refl.                          | <b>10d</b>          | 85     |                                            |
| <b>7</b>            | <b>6d</b> | DMF                   | 15         | r.t.                           | <b>10d</b>          | 77     |                                            |
| <b>7</b>            | <b>6f</b> | THF                   | 60         | Refl.                          | <b>10f</b>          | 81     |                                            |
| <b>7</b>            | <b>6g</b> | THF                   | 60         | Refl.                          | <b>10g</b>          | 96     | <b>13g</b> 1                               |
| <b>7</b>            | <b>6h</b> | THF                   | 60         | Refl.                          | <b>10h</b>          | 79     |                                            |
| <b>7</b>            | <b>6h</b> | DMF                   | 10         | 80                             | <b>10h</b>          | 77     |                                            |
| <b>7</b>            | <b>6i</b> | THF                   | 60         | Refl.                          | <b>10i</b>          | 35     | 11                                         |
| <b>7</b>            | <b>6i</b> | DMF                   | 10         | 80                             | <b>10i</b>          | 65     |                                            |
| <b>7</b>            | <b>6j</b> | THF                   | 60         | Refl.                          | <b>10j</b>          | 69     |                                            |
| <b>7</b>            | <b>6k</b> | THF                   | 30         | Refl.                          | <b>10k</b>          | 35     | 38                                         |
| <b>7</b>            | <b>6k</b> | THF                   | 150        | Refl.                          | <b>10k</b>          | 79     |                                            |
| <b>7</b>            | <b>6k</b> | DMF                   | 15         | r.t.                           | <b>10k</b>          | —      |                                            |
| <b>7</b>            | <b>6k</b> | DMF                   | 8          | 80                             | <b>10k</b>          | 90     | 87                                         |
| <b>7</b>            | <b>6m</b> | THF                   | 60         | Refl.                          | <b>10m</b>          | —      |                                            |
| <b>7</b>            | <b>6m</b> | DMF                   | 30         | 80                             | <b>10m</b>          | —      |                                            |
| <b>7</b>            | <b>6n</b> | THF                   | 60         | Refl.                          | <b>10n</b>          | —      | 67 <sup>c)</sup><br>21<br>46 <sup>d)</sup> |
| <b>7</b>            | <b>6n</b> | DMF                   | 5          | r.t.                           | <b>10n</b>          | 59     |                                            |
| <b>7</b>            | <b>6n</b> | DMF                   | 5          | 80                             | <b>10n</b>          | 42     |                                            |
| <b>7</b>            | <b>6o</b> | THF                   | 60         | Refl.                          | <b>10o</b>          | 31     | 11<br>53                                   |
| <b>7</b>            | <b>6o</b> | DMF                   | 60         | 80                             | <b>10o</b>          | —      |                                            |
| <b>7</b>            | <b>6q</b> | THF                   | 30         | Refl.                          | <b>10q</b>          | 7      |                                            |
| <b>7</b>            | <b>6q</b> | THF                   | 240        | Refl.                          | <b>10q</b>          | 28     | <b>13q</b> 12<br><b>13q</b> 52             |
| <b>7</b>            | <b>6q</b> | DMF                   | 10         | r.t.                           | <b>10q</b>          | 23     |                                            |
| <b>7</b>            | <b>6q</b> | DMF                   | 7          | 80                             | <b>10q</b>          | 58     |                                            |
| <b>7</b>            | <b>6r</b> | THF                   | 60         | Refl.                          | <b>10r</b>          | 47     |                                            |
| <b>7</b>            | <b>6r</b> | DMF                   | 5          | 80                             | <b>10r</b>          | 93     |                                            |
| <b>7</b>            | <b>6s</b> | THF                   | 60         | Refl.                          | <b>10s</b>          | 86     |                                            |
| <b>8</b>            | <b>6d</b> | Dioxane               | 60         | Refl.                          | <b>11d</b>          | 83     |                                            |
| <b>8</b>            | <b>6g</b> | THF                   | 20         | Refl.                          | <b>11g</b>          | 85     |                                            |
| <b>8</b>            | <b>6k</b> | Dioxane               | 60         | Refl.                          | <b>11k</b>          | 51     |                                            |
| <b>9</b>            | <b>6a</b> | THF                   | 30         | Refl.                          | <b>12a</b>          | 88     | <b>14a</b> 2                               |
| <b>9</b>            | <b>6a</b> | DMF                   | 20         | r.t.                           | <b>12a</b>          | 79     |                                            |
| <b>9</b>            | <b>6b</b> | THF                   | 60         | Refl.                          | <b>12b</b>          | 51     |                                            |
| <b>9</b>            | <b>6b</b> | DMF                   | 7          | 80                             | <b>12b</b>          | 89     | <b>14b</b> 3<br><b>14d</b> 8               |
| <b>9</b>            | <b>6d</b> | THF                   | 60         | Refl.                          | <b>12d</b>          | 65     |                                            |
| <b>9</b>            | <b>6d</b> | DMF                   | 20         | 80                             | <b>12d</b>          | 76     |                                            |
| <b>9</b>            | <b>6e</b> | DMF                   | 20         | r.t.                           | <b>12e</b>          | 81     |                                            |
| <b>9</b>            | <b>6f</b> | DMF                   | 20         | r.t.                           | <b>12f</b>          | 85     |                                            |
| <b>9</b>            | <b>6f</b> | DMF                   | 10         | 80                             | <b>12f</b>          | 37     |                                            |
| <b>9</b>            | <b>6g</b> | THF                   | 60         | Refl.                          | <b>12g</b>          | 35     | 42                                         |
| <b>9</b>            | <b>6g</b> | DMF                   | 10         | 80                             | <b>12g</b>          | 95     |                                            |
| <b>9</b>            | <b>6h</b> | THF                   | 60         | Refl.                          | <b>12g</b>          | 22     |                                            |
| <b>9</b>            | <b>6h</b> | Dioxane               | 60         | Refl.                          | <b>12h</b>          | 64     | 90                                         |
| <b>9</b>            | <b>6h</b> | DMF                   | 30         | r.t.                           | <b>12h</b>          | —      |                                            |
| <b>9</b>            | <b>6h</b> | DMF                   | 20         | 80                             | <b>12h</b>          | 92     |                                            |
| <b>9</b>            | <b>6i</b> | THF                   | 60         | Refl.                          | <b>12i</b>          | —      | 56<br>86                                   |
| <b>9</b>            | <b>6i</b> | DMF                   | 40         | r.t.                           | <b>12i</b>          | —      |                                            |
| <b>9</b>            | <b>6i</b> | DMF                   | 20         | 80                             | <b>12i</b>          | 81     |                                            |
| <b>9</b>            | <b>6p</b> | THF                   | 60         | Refl.                          | <b>12p</b>          | —      | 62<br>85                                   |
| <b>9</b>            | <b>6p</b> | DMF                   | 20         | r.t.                           | <b>12p</b>          | —      |                                            |
| <b>9</b>            | <b>6p</b> | DMF                   | 20         | 80                             | <b>12p</b>          | 72     |                                            |
| <b>9</b>            | <b>6r</b> | THF                   | 30         | Refl.                          | <b>12r</b>          | 47     | <b>14r</b> 28                              |
| <b>9</b>            | <b>6r</b> | DMF                   | 10         | 80                             | <b>12r</b>          | 67     |                                            |

a) THF = tetrahydrofuran, DMF = *N,N*-dimethylformamide. b) Refl. = reflux, r.t. = room temperature. c) *p*-Nitrobenzoic acid (**16m**) was obtained in 4% yield. d) *p*-Cyanobenzoic acid (**16n**) was obtained in 9% yield.

10 min in DMF (method B). Moreover, the aroylation with **6c** and **6d** at room temperature resulted in the formation of the corresponding ketones **10c** and **10d** in excellent yields. The treatment of **7** with **6k** at room temperature resulted in recovery of the starting **7**, while at 80 °C the aroylation proceeded, giving the ketone **10k**<sup>6)</sup> in 90%

yield. The reaction of **7** with **6q** in DMF at room temperature gave the ketone **10q** together with the 4-(1-naphthylmethoxy)quinazoline (**13q**) in 52% yield, and this is the same result as obtained in the reaction in THF. But, only the ketone **10q** was formed in the reaction at 80 °C for 7 min. The reaction with **6n** having a strong

electron-withdrawing group produced the ketone, 4-(4-cyanobenzoyl)quinazoline (**10n**), both at 80 °C and at room temperature, and there was no appreciable difference between the yields of the ketone obtained under the two conditions described above. With **6m**, the reaction at 80 °C for 30 min resulted in recovery of the starting **7**, and the result was similar to that obtained in the reaction using THF as the reaction solvent.

Next, the above arylation was extended to **8** and **9**. When the 4-chloroquinazoline **8** was treated with **6d** and NaH in the presence of **1** in refluxing dioxane for 30 min, the arylation took place, resulting in the formation of the ketone **11d** in 83% yield. Similar treatment with **6g** in refluxing THF for 20 min gave the expected ketone **11g** in 85% yield. A 51% yield of the ketone **11k** was obtained by the reaction of **8** with **6k** in dioxane.

The conversion of **9** into the corresponding ketones **12a**, **12b**, and **12d** was achieved by treatment with **6a**, **6b**, and **6d** in refluxing THF catalyzed by **1**, together with a small amount of the corresponding 4-(arylmethoxy)quinazolines **14a**, **14b**, and **14d**. A similar result was obtained in the reaction of **9** with **6r** in refluxing THF for 30 min: the ketone was obtained in 47% yield together with 4-fur-

furyloxy-2-phenylquinazoline (**14r**) in 28% yield. In the reaction with **6g** for 60 min in refluxing THF, the ketone **12g** was obtained in low yield (35%) with recovery of a large amount of starting **9** (42%). Furthermore, when the arylation was done with **6l** and **6p**, which have an electron-donating group at the *para*-position, in refluxing THF, the starting **9** was recovered in quantitative yield. These results prompted us to examine the new arylation of **9**, whose 4-position may have weaker nucleophilic reactivity than that of **7**. The arylation of **9** in DMF by the same method as described for the reaction of **7** with aromatic aldehydes **6** provided the ketone in excellent yield under mild conditions. Thus, the treatment of **9** with **6a**, **6d**, **6e**, and **6f**, bearing halogen substituent, in DMF at room temperature for 20 min gave the corresponding ketones **12a**, **12d**, **12e**, and **12f** in good yields. But, when the 4-chloroquinazoline **9** was treated with aromatic aldehydes **6h**, **6l**, and **6p**, which have an electron-donating group at the *para*-position, under similar conditions, none of the corresponding ketones was obtained and the starting **7** was recovered in quantitative yield. In contrast, when the same reactions were examined at 80 °C for 10 to 20 min, the expected ketones **12h**, **12l**, and **12p** were formed in

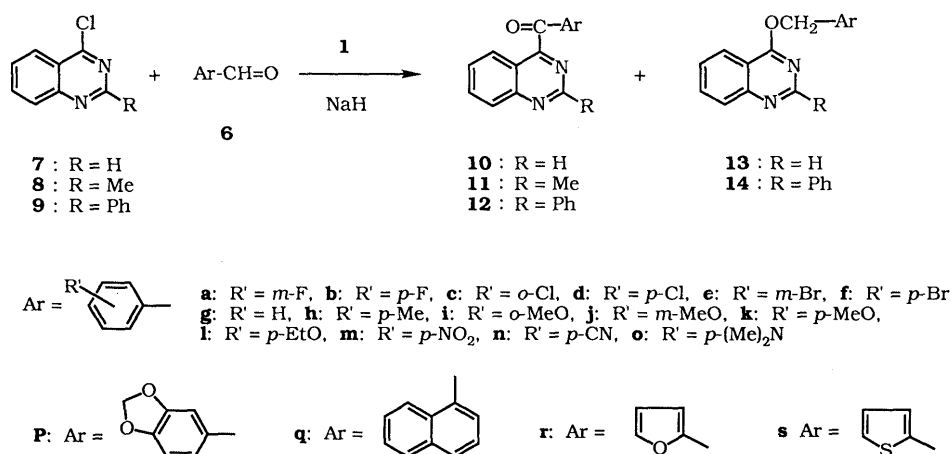


Chart 2

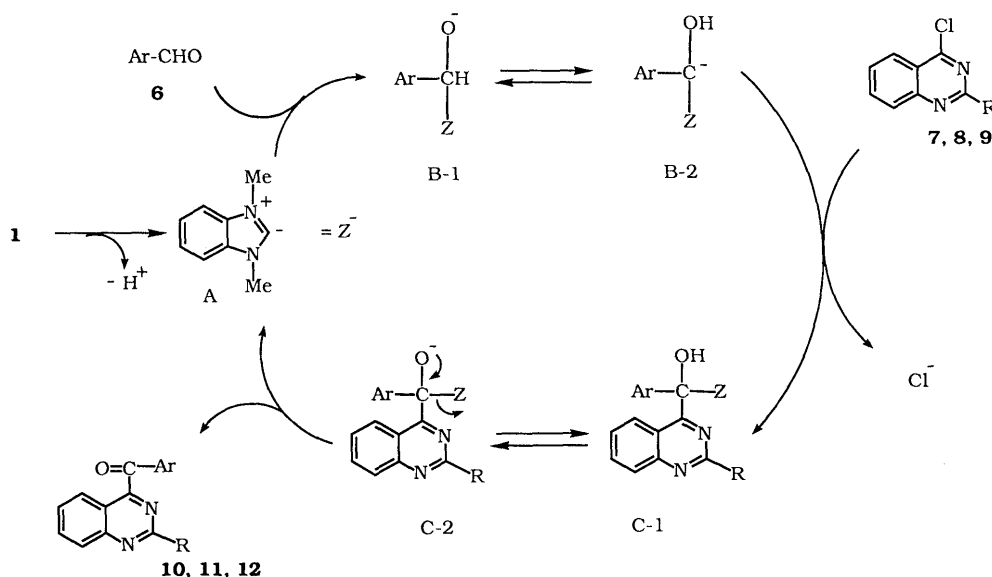


Chart 3

TABLE II. Melting Points, Mass Spectral Data, and Elemental Analyses for **10**, **11**, and **12**

| Compd.     | mp (°C)                                                | Formula                                                       | Analysis (%)     |              |                | MS $m/z$<br>$M^+$ |
|------------|--------------------------------------------------------|---------------------------------------------------------------|------------------|--------------|----------------|-------------------|
|            |                                                        |                                                               | Calcd (Found)    | C            | H              | N                 |
| <b>10b</b> | 125.5—126 <sup>a,b</sup>                               | C <sub>15</sub> H <sub>9</sub> FN <sub>2</sub> O              | 71.42<br>(71.37) | 3.60<br>3.52 | 11.11<br>11.07 | 318               |
| <b>10c</b> | 74—76 <sup>c,b</sup><br>(lit. <sup>6</sup> ) 80—81     | C <sub>15</sub> H <sub>9</sub> ClN <sub>2</sub> O             |                  |              |                |                   |
| <b>10d</b> | 130—131 <sup>a,b</sup><br>(lit. <sup>6</sup> ) 128—129 | C <sub>15</sub> H <sub>9</sub> ClN <sub>2</sub> O             |                  |              |                |                   |
| <b>10f</b> | 126—127 <sup>a,b</sup>                                 | C <sub>15</sub> H <sub>9</sub> BrN <sub>2</sub> O             | 57.53<br>(57.53) | 2.90<br>2.83 | 8.95<br>8.94   | 378, 380          |
| <b>10g</b> | 98—99 <sup>d,e</sup><br>(lit. <sup>6</sup> ) 97—98     | C <sub>15</sub> H <sub>10</sub> N <sub>2</sub> O              |                  |              |                |                   |
| <b>10h</b> | 130—131 <sup>e,f</sup><br>(lit. <sup>6</sup> ) 129—130 | C <sub>16</sub> H <sub>12</sub> N <sub>2</sub> O              |                  |              |                |                   |
| <b>10i</b> | 98—100 <sup>b,g</sup><br>(lit. <sup>6</sup> ) 100—101  | C <sub>16</sub> H <sub>12</sub> N <sub>2</sub> O <sub>2</sub> |                  |              |                |                   |
| <b>10j</b> | 122—123 <sup>b,g</sup><br>(lit. <sup>6</sup> ) 122     | C <sub>16</sub> H <sub>12</sub> N <sub>2</sub> O <sub>2</sub> |                  |              |                |                   |
| <b>10k</b> | 120—121 <sup>a,b</sup><br>(lit. <sup>6</sup> ) 119—120 | C <sub>16</sub> H <sub>12</sub> N <sub>2</sub> O <sub>2</sub> |                  |              |                |                   |
| <b>10n</b> | 162—163 <sup>d,b</sup>                                 | C <sub>16</sub> H <sub>9</sub> N <sub>3</sub> O               | 74.12<br>(74.46) | 3.50<br>3.43 | 16.21<br>15.93 | 258               |
| <b>10o</b> | 159 <sup>b,h</sup>                                     | C <sub>17</sub> H <sub>15</sub> N <sub>3</sub> O              | 73.63<br>(73.72) | 5.45<br>5.47 | 15.15<br>15.21 | 344               |
| <b>10q</b> | 151.5—152 <sup>b,h</sup>                               | C <sub>19</sub> H <sub>12</sub> N <sub>2</sub> O              | 80.27<br>(79.95) | 4.25<br>4.37 | 9.85<br>9.78   | 368               |
| <b>10r</b> | 165.5—166 <sup>a,b</sup><br>(lit. <sup>6</sup> ) 162   | C <sub>13</sub> H <sub>8</sub> N <sub>2</sub> O <sub>2</sub>  |                  |              |                |                   |
| <b>10s</b> | 136—138 <sup>b,i</sup>                                 | C <sub>13</sub> H <sub>8</sub> FN <sub>2</sub> OS             | 64.98<br>(64.89) | 3.36<br>3.31 | 11.66<br>11.65 | 256               |
| <b>11d</b> | 141—142 <sup>b,i</sup>                                 | C <sub>16</sub> H <sub>11</sub> ClN <sub>2</sub> O            | 67.97<br>(67.83) | 3.92<br>3.87 | 9.91<br>9.84   | 256               |
| <b>11g</b> | 142—143 <sup>b,g</sup>                                 | C <sub>16</sub> H <sub>12</sub> N <sub>2</sub> O              | 77.40<br>(77.36) | 4.87<br>4.88 | 11.28<br>11.14 | 247               |
| <b>11k</b> | 142—144 <sup>b,g</sup>                                 | C <sub>17</sub> H <sub>14</sub> N <sub>2</sub> O <sub>2</sub> | 73.37<br>(73.14) | 5.07<br>5.08 | 10.07<br>9.96  | 266               |
| <b>12a</b> | 153—154 <sup>b,i</sup>                                 | C <sub>21</sub> H <sub>13</sub> FN <sub>2</sub> O             | 76.82<br>(76.79) | 3.99<br>3.98 | 8.58<br>8.53   | 329               |
| <b>12b</b> | 172—173 <sup>a,b</sup>                                 | C <sub>21</sub> H <sub>13</sub> FN <sub>2</sub> O             | 76.82<br>(77.04) | 3.99<br>3.91 | 8.53<br>8.53   | 328               |
| <b>12d</b> | 186—186.5 <sup>b,i</sup>                               | C <sub>21</sub> H <sub>13</sub> ClN <sub>2</sub> O            | 73.15<br>(73.05) | 3.80<br>3.71 | 8.12<br>8.07   | 345               |
| <b>12e</b> | 181—184 <sup>e,i</sup>                                 | C <sub>21</sub> H <sub>13</sub> BrN <sub>2</sub> O            | 64.80<br>(65.11) | 3.37<br>3.43 | 7.20<br>7.68   | 388, 390          |
| <b>12f</b> | 181—183 <sup>b,i</sup>                                 | C <sub>21</sub> H <sub>13</sub> BrN <sub>2</sub> O            | 64.80<br>(65.02) | 3.37<br>3.36 | 7.20<br>7.25   | 388, 390          |
| <b>12g</b> | 151.5—152 <sup>e,i</sup>                               | C <sub>21</sub> H <sub>14</sub> N <sub>2</sub> O              | 81.27<br>(81.40) | 4.55<br>4.56 | 9.03<br>9.00   | 256               |
| <b>12h</b> | 148—148.5 <sup>b,i</sup>                               | C <sub>22</sub> H <sub>16</sub> N <sub>2</sub> O              | 81.46<br>(81.29) | 4.97<br>4.95 | 8.64<br>8.46   | 324               |
| <b>12l</b> | 147—148 <sup>b,i</sup>                                 | C <sub>23</sub> H <sub>18</sub> N <sub>2</sub> O <sub>2</sub> | 77.95<br>(77.68) | 5.12<br>5.16 | 7.90<br>7.88   | 354               |
| <b>12p</b> | 162.5—163 <sup>b,j</sup>                               | C <sub>22</sub> H <sub>14</sub> N <sub>2</sub> O <sub>3</sub> | 74.56<br>(74.68) | 3.98<br>3.93 | 7.98<br>7.90   | 355               |
| <b>12r</b> | 151—152 <sup>b,j</sup>                                 | C <sub>19</sub> H <sub>12</sub> N <sub>2</sub> O <sub>2</sub> | 75.99<br>(75.84) | 4.03<br>4.02 | 9.03<br>9.35   | 301               |

a) Colorless needles. b) Recrystallized from MeOH. c) Slightly yellow granules. d) Pale yellow needles. e) Recrystallized from petroleum benzene. f) Slightly yellow scales. g) Slightly yellow prisms. h) Yellow prisms. i) Slightly yellow needles. j) Yellow needles.

92%, 81%, and 72% yields. Similarly, the treatment of **9** with **6b**, **6d**, and **6r** in DMF at 80 °C for 7 to 10 min resulted in the formation of the ketones **12b**, **12d**, and **12r** in good yields. The reaction with **6f** at room temperature for 20 min gave the ketone **12f** in 85% yield, while the yield of the ketone decreased in the same reaction at 80 °C. This may be a result of further reaction, such as aryl migration,<sup>7)</sup> proceeding through the ketone. The results obtained in this paper are summarized in Table I.

The structures of the newly obtained ketones were sup-

TABLE III. IR and <sup>1</sup>H-NMR Spectral Data for **10**, **11**, and **12**

| Compd.     | IR $\nu_{\text{KBr}}^{\text{max}}$ cm <sup>-1</sup> | <sup>1</sup> H-NMR (CDCl <sub>3</sub> ) $\delta$ (ppm)                                                                                                                                                                                                                                                                                      |
|------------|-----------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <b>10b</b> | 1665 (CO)                                           | 9.28 (1H, s, C <sup>2</sup> -H), 6.90—8.05 (8H, m, aromatic H)                                                                                                                                                                                                                                                                              |
| <b>10f</b> | 1660 (CO)                                           | 9.28 (1H, s, C <sup>2</sup> -H), 7.40—8.10 (8H, m, aromatic H)                                                                                                                                                                                                                                                                              |
| <b>10n</b> | 1670 (CO)                                           | 9.25 (1H, s, C <sup>2</sup> -H), 7.10—8.10 (8H, m, aromatic H)                                                                                                                                                                                                                                                                              |
| <b>10o</b> | 2220 (CN)<br>1640 (CO)                              | 9.25 (1H, s, C <sup>2</sup> -H), 7.50—8.10 (6H, m, aromatic H), 6.51 (2H, d, $J = 8$ Hz,  , 3.05 (6H, s, N(CH <sub>3</sub> ) <sub>2</sub> )                                                                                                              |
| <b>10q</b> | 1650 (CO)                                           | 9.25 (1H, s, C <sup>2</sup> -H), 8.80—9.00 (1H, m,  , 7.30—8.15 (10H, m, aromatic H)                                                                                                                                                                     |
| <b>10s</b> | 1640 (CO)                                           | 9.30 (1H, s, C <sup>2</sup> -H), 7.10 (1H, dd,  , $J = 6$ , 2 Hz), 7.50—8.30 (6H, m, aromatic H)                                                                                                                                                         |
| <b>11d</b> | 1670 (CO)                                           | 7.20—8.00 (8H, m, aromatic H), 2.92 (3H, s, C <sup>2</sup> -Me)                                                                                                                                                                                                                                                                             |
| <b>11g</b> | 1670 (CO)                                           | 7.38—8.04 (9H, m, aromatic H), 2.93 (3H, s, C <sup>2</sup> -Me)                                                                                                                                                                                                                                                                             |
| <b>11k</b> | 1655 (CO)                                           | 7.39—7.95 (6H, m, aromatic H), 6.89 (2H, d, $J = 9$ Hz,  , 3.85 (3H, s, OMe), 2.92 (3H, s, C <sup>2</sup> -Me)                                                                                                                                           |
| <b>12a</b> | 1675 (CO)                                           | 8.58—8.60 (2H, m), <sup>a)</sup> 7.35—8.20 (11H, m, aromatic H)                                                                                                                                                                                                                                                                             |
| <b>12b</b> | 1670 (CO)                                           | 8.58—8.60 (2H, m), <sup>a)</sup> 7.20—8.20 (11H, m, aromatic H)                                                                                                                                                                                                                                                                             |
| <b>12d</b> | 1670 (CO)                                           | 8.35—8.55 (2H, m), <sup>a)</sup> 7.20—8.05 (11H, m, aromatic H)                                                                                                                                                                                                                                                                             |
| <b>12e</b> | 1670 (CO)                                           | 8.45—8.62 (2H, m), <sup>a)</sup> 7.35—8.30 (11H, m, aromatic H)                                                                                                                                                                                                                                                                             |
| <b>12f</b> | 1675 (CO)                                           | 8.40—8.65 (2H, m), <sup>a)</sup> 7.30—8.10 (11H, m, aromatic H)                                                                                                                                                                                                                                                                             |
| <b>12g</b> | 1670 (CO)                                           | 8.35—8.55 (2H, m), <sup>a)</sup> 7.20—8.05 (12H, m, aromatic H)                                                                                                                                                                                                                                                                             |
| <b>12h</b> | 1665 (CO)                                           | 8.60—8.62 (2H, m), 7.15—8.10 (11H, m, aromatic H), 2.40 (3H, s, Me)                                                                                                                                                                                                                                                                         |
| <b>12l</b> | 1660 (CO)                                           | 8.45—8.70 (2H, m), <sup>a)</sup> 7.35—8.15 (9H, m, aromatic H), 6.90 (2H, d, $J = 8$ Hz,  , OC <sub>2</sub> H <sub>5</sub> ), 4.10 (2H, q, $J = 7$ Hz, OCH <sub>2</sub> CH <sub>3</sub> ), 1.46 (3H, t, $J = 7$ Hz, OCH <sub>2</sub> CH <sub>3</sub> ) |
| <b>12p</b> | 1650 (CO)                                           | 8.60—8.62 (2H, m), <sup>a)</sup> 7.50—8.15 (9H, m, aromatic H), 6.83 (1H, d, $J = 8$ Hz,  , 6.01 (2H, s, OCH <sub>2</sub> O)                                                                                                                           |
| <b>12r</b> | 1650 (CO)                                           | 7.40—8.70 (11H, m, aromatic H), 6.60 (1H, dd, $J = 6$ , 2 Hz,  )                                                                                                                                                                                       |



ported by the elemental analyses, infrared (IR) spectra, and proton (<sup>1</sup>H-) and carbon (<sup>13</sup>C-) nuclear magnetic resonance (NMR) spectra, as shown in Tables II, III, and IV. The structures of the 4-(arylmethoxy)quinazolines **13** and **14** were determined by mixed melting point determination with authentic samples.

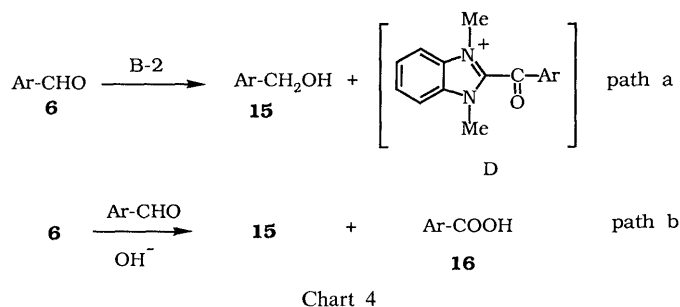
As illustrated in Chart 3, the formation of the ketones involving the catalytic action of **1** is considered to proceed through a similar process to that reported for the formation of 4-aryl-1 *H*-pyrazolo[3,4-*d*]pyrimidines.<sup>1)</sup> The for-

TABLE IV.  $^{13}\text{C}$ -NMR Spectral Data for **10**, **11**, and **12**

| Compd.                 | $^{13}\text{C}$ -NMR ( $\text{CDCl}_3$ ) $\delta$ ppm                                                                                                                                                                                                                                     |
|------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <b>10b<sup>a</sup></b> | 115.5 (d), 116.5 (d), 122.0 (s), 125.8 (d), 128.8 (d), 129.1 (d), 130.6 (s), 133.2 (d), 133.7 (d), 134.6 (d), 151.3 (s), 153.7 (d), 160.8 (s), 163.5 (s), 172.2 (s), 191.2 (s, CO)                                                                                                        |
| <b>10f</b>             | 122.0 (s), 125.6 (d), 128.8 (d), 129.1 (d), 129.9 (s), 132.0 (d), 134.0 (s), 134.6 (d), 151.4 (s), 153.7 (d), 163.8 (s), 191.7 (s, CO)                                                                                                                                                    |
| <b>10n</b>             | 117.2 (s, CN), 121.9 (s), 125.5 (d), 129.2 (d), 131.0 (d), 132.3 (d), 134.9 (d), 138.4 (s), 151.6 (s), 153.5 (d), 161.7 (s), 191.4 (s, CO)                                                                                                                                                |
| <b>10o</b>             | 40.0 (q, -NMe), 110.8 (d), 122.3 (s), 123.0 (s), 126.3 (d), 128.3 (d), 128.8 (d), 133.0 (d), 134.4 (d), 150.9 (s), 154.1 (d), 154.3 (s), 166.1 (s), 190.5 (s, CO)                                                                                                                         |
| <b>10q</b>             | 122.2 (s), 124.1 (d), 125.9 (d), 126.8 (d), 128.7 (d), 128.9 (d), 129.0 (d), 131.3 (s), 132.3 (s), 133.9 (d), 134.4 (d), 134.9 (d), 151.3 (s), 154.8 (d), 165.2 (s), 195.3 (s, CO)                                                                                                        |
| <b>10s</b>             | 121.7 (s), 126.1 (d), 128.4 (d), 129.0 (d), 134.4 (d), 136.8 (d), 137.1 (d), 141.5 (s), 151.7 (s), 153.6 (d), 161.7 (s), 184.4 (s, CO)                                                                                                                                                    |
| <b>11d</b>             | 26.4 (q, Me), 119.7 (s), 125.5 (d), 127.8 (d), 128.4 (d), 129.1 (d), 132.0 (d), 133.6 (s), 134.6 (d), 141.1 (s), 151.6 (s), 163.3 (s), 163.7 (s), 191.8 (s, CO)                                                                                                                           |
| <b>11g</b>             | 26.4 (q, Me), 119.8 (s), 125.6 (d), 127.7 (d), 128.3 (d), 128.8 (d), 130.6 (d), 134.49 (d), 134.54 (d), 135.2 (s), 151.4 (s), 163.4 (s), 164.6 (s), 193.2 (s, CO)                                                                                                                         |
| <b>11k</b>             | 26.4 (q, Me), 55.6 (q, OMe), 114.1 (d), 119.8 (s), 125.8 (d), 127.5 (d), 128.2 (d), 133.1 (d), 134.5 (d), 151.3 (s), 163.4 (s), 164.7 (s), 165.2 (s), 191.6 (s, CO)                                                                                                                       |
| <b>12a<sup>a</sup></b> | 116.9 (d), 117.3 (d), 120.5 (s), 121.1 (d), 121.5 (d), 125.6 (d), 126.8 (d), 126.9 (d), 128.2 (d), 128.7 (d), 129.4 (d), 130.3 (d), 130.4 (d), 131.0 (d), 134.6 (d), 137.3 (s), 152.2 (s), 159.5 (s), 160.9 (s), 163.1 (s), 164.5 (s), 191.7 (s, CO)                                      |
| <b>12b<sup>a</sup></b> | 115.8 (d), 116.1 (d), 120.5 (s), 125.7 (d), 128.1 (d), 128.7 (d), 129.2 (s), 129.3 (d), 131.0 (d), 131.9 (s), 133.5 (d), 133.7 (d), 134.6 (d), 137.3 (s), 152.2 (s), 159.5 (s), 163.5 (s), 164.6 (s), 168.4 (s), 191.4 (s, CO)                                                            |
| <b>12d</b>             | 120.5 (s), 125.7 (d), 128.1 (d), 128.6 (d), 128.7 (d), 129.0 (d), 129.3 (d), 131.0 (d), 132.2 (d), 133.8 (s), 134.6 (d), 137.3 (s), 140.9 (s), 152.2 (s), 159.5 (s), 163.1 (s), 191.8 (s, CO)                                                                                             |
| <b>12e</b>             | 120.5 (s), 123.0 (s), 125.6 (d), 125.8 (s), 128.2 (d), 128.65 (d), 128.70 (d), 128.9 (s), 129.4 (d), 129.6 (d), 130.2 (d), 131.0 (d), 131.1 (s), 133.4 (d), 134.6 (d), 137.0 (d), 137.2 (s), 152.3 (s), 193.6 (s, CO)                                                                     |
| <b>12f</b>             | 120.6 (s), 125.7 (d), 128.0 (d), 128.6 (d), 128.7 (d), 129.3 (d), 130.8 (d), 130.9 (d), 131.8 (s), 134.3 (d), 134.5 (d), 135.4 (s), 137.5 (s), 152.1 (s), 159.6 (s), 164.1 (s), 193.1 (s, CO)                                                                                             |
| <b>12g</b>             | 120.5 (s), 125.7 (d), 127.9 (d), 128.6 (d), 128.7 (d), 129.3 (d), 130.8 (d), 130.9 (d), 134.3 (d), 134.4 (d), 135.4 (s), 137.4 (s), 152.0 (s), 159.6 (s), 164.1 (s), 193.1 (s, CO)                                                                                                        |
| <b>12h</b>             | 21.9 (t, Me), 120.6 (s), 125.8 (d), 127.9 (d), 128.4 (s), 128.6 (d), 128.7 (d), 129.2 (d), 129.4 (d), 130.85 (d), 130.90 (d), 134.4 (d), 137.5 (s), 145.5 (s), 152.0 (s), 159.6 (s), 164.5 (s), 192.7 (s, CO)                                                                             |
| <b>12l</b>             | 14.6 (q, -OCH <sub>2</sub> CH <sub>3</sub> ), 63.9 (t, -OCH <sub>2</sub> CH <sub>3</sub> ), 114.4 (d), 120.6 (s), 125.9 (d), 127.8 (d), 128.2 (s), 128.6 (d), 128.7 (d), 129.2 (d), 130.8 (d), 133.2 (d), 134.4 (d), 137.5 (s), 151.9 (s), 159.6 (s), 164.0 (s), 164.8 (s), 191.6 (s, CO) |
| <b>12p</b>             | 102.1 (t, -OCH <sub>2</sub> O-), 108.1 (d), 109.4 (d), 120.6 (s), 125.8 (d), 127.9 (d), 128.6 (d), 128.7 (d), 129.2 (d), 130.2 (s), 130.9 (d), 134.4 (d), 137.5 (s), 148.4 (s), 151.9 (s), 153.1 (s), 159.6 (s), 164.7 (s), 191.2 (s, CO)                                                 |
| <b>12r</b>             | 112.9 (d), 120.4 (s), 124.4 (d), 125.9 (d), 128.3 (d), 128.6 (d), 128.8 (d), 129.2 (d), 131.0 (d), 134.5 (d), 137.4 (s), 148.8 (d), 151.5 (s), 152.5 (s), 159.5 (s), 161.3 (s), 179.9 (s, CO)                                                                                             |

a) The spectrum was observed with  $^{13}\text{C}$ - $^{19}\text{F}$  coupling.

mation of the ketones showed that the 4-chloroquinazoline react preferentially with the carbanion B-2 rather than the O-anion B-1, as an intermediate in the arylation. In this recycling process, 1,3-dimethylbenzimidazolium iodide (**1**) acts as a catalyst.



The formation of the 4-(arylmethoxy)quinazolines **13** and **14** may be considered to involve nucleophilic substitution between the arylmethoxide ion and the 4-chloroquinazolines **7** and **9**. The arylmethanol **15**, the source of the arylmethoxide ion, may be formed in the following two ways, involving oxidation-reduction reaction of the aldehydes: a) redox reaction between the intermediate B-2 and aromatic aldehyde, as reported in the previous paper,<sup>1)</sup> and b) Cannizzaro reaction<sup>8)</sup> between two molecules of the aromatic aldehyde with hydroxide ion derived from contaminating water in the reaction solvent and NaH. Our results support path b because the acids **16**, which might not be formed *via* path a, were obtained from the reaction of **7** with **6m** and **6n**, but it is not yet clear whether the arylmethanols **15** were formed through path a or path b. In fact, neither the arylmethanols **15** nor the 4-(arylmethoxy)quinazolines **13** were formed together with the acids **16** in the above reaction.

In conclusion, the nucleophilic arylation of the 4-chloroquinazolines **10**, **11**, and **12** with aromatic aldehydes **6** under the catalytic action of 1,3-dimethylbenzimidazolium iodide (**1**) provides a useful method for the preparation of the 4-aryloxyquinazolines **10**, **11**, and **12**, being superior to the other methods reported.<sup>6,7,9)</sup>

#### Experimental

All melting points are uncorrected. Infrared (IR) absorption spectra were recorded on a Jasco A-102 diffraction grating IR spectrometer.  $^1\text{H}$ -NMR spectra were measured at 60 MHz on a Hitachi R-24B high-resolution NMR spectrometer, and  $^{13}\text{C}$ -NMR spectra were obtained with a JEOL JNM-FX90Q FT-NMR spectrometer operating at 22.5 MHz. Chemical shifts are quoted in parts per million (ppm) with tetramethylsilane as an internal standard, and coupling constants ( $J$ ) are given in hertz (Hz). The following abbreviations are used: s=singlet, d=doublet, t=triplet, q=quartet, and m=multiplet. Mass spectra (MS) were recorded on a JEOL JMS D-100 mass spectrometer. Samples were vaporized in a direct inlet system. Column chromatography was carried out on  $\text{SiO}_2$ , Wakogel C-200. The 4-chloroquinazolines **7**,<sup>3,4)</sup> **8**,<sup>5)</sup> and **9**<sup>4)</sup> and 1,3-dimethylbenzimidazolium iodide (**1**)<sup>1)</sup> were prepared by the reported procedures.

**Reaction of the 4-Chloroquinazolines (**7**, **8**, and **9**) with Aromatic Aldehydes **6** in the Presence of NaH Catalyzed by 1,3-Dimethylbenzimidazolium Iodide (**1**)** General Procedure for Method A: A stirred solution of a 4-chloroquinazoline (**7**, **8**, or **9**, 4 mmol), an aromatic aldehyde (**6**, 4.8 mmol), and 1,3-dimethylbenzimidazolium iodide (**1**, 274 mg, 1.0 mmol) in THF (20 ml) or in dioxane (20 ml) was treated with NaH (50% in oil, 230 mg, 4.8 mmol) under an  $\text{N}_2$  atmosphere, and then the mixture was refluxed with stirring in an oil bath for an appropriate time (the progress of the reaction was checked by TLC, and the reaction conditions are shown in Table I). After cooling, the resultant mixture was poured into an ice-water mixture, and extracted with  $\text{CHCl}_3$  (100 ml  $\times$  2). The organic layer was dried over  $\text{Na}_2\text{SO}_4$  and concentrated to dryness. The residue was dissolved in a small portion of benzene, and chromatographed on a column of  $\text{SiO}_2$  with benzene, and then  $\text{CHCl}_3$ . The fraction eluted with benzene provided the starting 4-chloroquinazoline and the fraction eluted with  $\text{CHCl}_3$  gave the ketone (yield, recrystallized solvent, and

spectral data are shown in Tables I–IV).

In the reaction of **7** with **6g**, the fraction eluted with benzene gave 4-benzoyloxyquinazoline (**13g**, 10 mg, 1%) and the fraction eluted with  $\text{CHCl}_3$  gave 4-benzoylquinazoline (**10g**, 900 mg, 96%).

In the reaction of **7** with **6q**, the fraction eluted with benzene gave 4-(1-naphthylmethoxy)quinazoline (**13q**, 137 mg, 12%), and the fraction eluted with  $\text{CHCl}_3$  gave 4-(1-naphthyl)quinazoline (**10q**, 322 mg, 28%).

In the reaction of **7** with **6m**, after extraction with  $\text{CHCl}_3$  the  $\text{H}_2\text{O}$  layer was acidified with  $\text{AcOH}$ , and extracted with  $\text{CHCl}_3$ , then the extract was dried over  $\text{Na}_2\text{SO}_4$ . After removal of the  $\text{CHCl}_3$  by evaporation, the residue was recrystallized from MeOH to give 4-nitrobenzoic acid (**16m**, 32 mg, 4%).

In the reaction of **7** with **6n**, work-up as described for the reaction with **6m** gave 4-cyanobenzoic acid (**16n**, 64 mg, 9%).

In the reaction of **7** with **6a**, the fraction eluted with benzene gave 4-(3-fluorobenzoyloxy)-2-phenylquinazoline (**14a**, 20 mg, 2%), and the fraction eluted with  $\text{CHCl}_3$  gave 4-(3-fluorobenzoyl)-2-phenylquinazoline (**12a**, 1160 mg, 88%).

In the reaction of **9** with **6b**, the first fraction eluted with benzene gave 4-(4-fluorobenzoyloxy)-2-phenylquinazoline (**14b**, 33 mg, 3%), and the second fraction eluted with benzene gave 4-(4-fluorobenzoyl)-2-phenylquinazoline (**12b**, 892 mg, 68%).

In the reaction of **9** with **6d**, the fraction eluted with benzene gave 4-(4-chlorobenzoyloxy)-2-phenylquinazoline (**14d**, 111 mg, 8%), and the fraction eluted with  $\text{CHCl}_3$  gave 4-(4-chlorobenzoyl)-2-phenylquinazoline (**12d**, 896 mg, 65%).

In the reaction of **9** with **6r**, the first fraction eluted with benzene gave 4-furfuryloxy-2-phenylquinazoline (**14r**, 338 mg, 28%), and the second fraction eluted with benzene gave 4-(2-furoyl)-2-phenylquinazoline (**12r**, 564 mg, 47%).

**General Procedure for Method B:** A stirred solution of a 4-chloroquinazoline (**7** or **9**, 4 mmol), an aromatic aldehyde (**6**, 4.8 mmol), and 1,3-dimethylbenzimidazolium iodide (**1**, 274 mg, 1.0 mmol) in DMF (20 ml) was treated with NaH (50% in oil, 230 mg, 4.8 mmol) under an  $\text{N}_2$  atmosphere and then the mixture was stirred in an oil bath under heating or at room temperature. The resultant mixture was poured into an ice-water mixture, extracted with  $\text{AcOEt}$  (100 ml  $\times$  2), dried over  $\text{Na}_2\text{SO}_4$ , and concentrated to dryness. The residue was chromatographed on a column of  $\text{SiO}_2$  with benzene and then  $\text{CHCl}_3$ . The fraction eluted with benzene gave 4-chloroquinazoline and the fraction eluted with  $\text{CHCl}_3$  gave the ketone.

In the reaction of **7** with **6q** at room temperature for 10 min, the fraction eluted with benzene gave **13q** (260 mg, 23%), and the fraction eluted with  $\text{CHCl}_3$  gave the ketone (**10q**, 593 mg, 52%).

**General Procedure for the Preparation of 4-(Arylmethoxy)quinazolines (13 and 14)** A stirred solution of a 4-chloroquinazoline (**7** or **9**, 3.0 mmol) and an arylmethanol (**15**, 3.6 mmol) was treated with NaH (50% in oil, 173 mg, 3.6 mmol), and the mixture was refluxed in an oil bath with stirring. After cooling, the resultant mixture was poured into an ice-water mixture, and extracted with  $\text{CHCl}_3$ . The extract was dried over  $\text{Na}_2\text{SO}_4$ , and concentrated to dryness. In order to purify it, the residue was passed through a short column of  $\text{SiO}_2$  with benzene to give the 4-(arylmethoxy)quinazoline (**13** or **14**).

**4-Benzoyloxyquinazoline (13g):** Yield 92% (650 mg). A colorless oil (picrate, yellow needles from MeOH, mp 164–165°C. *Anal.* Calcd for  $\text{C}_{21}\text{H}_{15}\text{N}_5\text{O}_8$ : C, 54.20; H, 3.25; N, 15.05. Found: C, 54.28; H, 3.24; N, 15.04).  $^1\text{H-NMR}$  ( $\text{CDCl}_3$ ): 8.72 (1H, s,  $\text{C}^2\text{-H}$ ), 7.15–8.05 (9H, m, aromatic H), 5.45 (2H, s,  $\text{CH}_2$ ).

**4-(1-Naphthylmethoxy)quinazoline (13q):** Yield 94% (810 mg). Colorless prisms (recrystallized from MeOH), mp 104–104.5°C. *Anal.* Calcd for  $\text{C}_{19}\text{H}_{14}\text{N}_2\text{O}$ : C, 79.70; H, 4.93; N, 9.78. Found: C, 79.59; H, 4.99; N, 9.71.  $^1\text{H-NMR}$  ( $\text{CDCl}_3$ ): 8.78 (1H, s,  $\text{C}^2\text{-H}$ ), 7.20–8.10 (11H, m, aromatic H), 6.01 (2H, s,  $\text{CH}_2$ ).

**4-(3-Fluorobenzoyloxy)-2-phenylquinazoline (14a):** Yield 96% (949 mg). Colorless needles (recrystallized from petroleum benzin), mp 123–123.5°C. *Anal.* Calcd for  $\text{C}_{21}\text{H}_{15}\text{FN}_2\text{O}$ : C, 76.35; H, 4.58; N, 8.48. Found: C, 76.12; H, 4.46; N, 8.51.  $^1\text{H-NMR}$  ( $\text{CDCl}_3$ ): 6.90–8.60 (13H, m, aromatic H), 5.65 (2H, s,  $\text{CH}_2$ ).

**4-(4-Fluorobenzoyloxy)-2-phenylquinazoline (14b):** Yield 93% (917 mg). Colorless needles (recrystallized from petroleum benzin), mp 92–93°C. *Anal.* Calcd for  $\text{C}_{21}\text{H}_{15}\text{FN}_2\text{O}$ : C, 76.35; H, 4.58; N, 8.48. Found: C, 76.12; H, 4.46; N, 8.51.

**4-(4-Chlorobenzoyloxy)-2-phenylquinazoline (14d):** Yield 88% (910 mg). Colorless prisms (recrystallized from MeOH), mp 132–133°C. *Anal.* Calcd for  $\text{C}_{21}\text{H}_{15}\text{ClN}_2\text{O}$ : C, 72.73; H, 4.36; N, 8.08. Found: C, 72.66; H, 4.37; N, 8.03.  $^1\text{H-NMR}$  ( $\text{CDCl}_3$ ): 8.40–8.58 (2H, m), 7.20–8.05 (11H, m, aromatic H), 5.50 (2H, s,  $\text{CH}_2$ ).

**4-Furfuryloxy-2-phenylquinazoline (14r):** Yield 86% (780 mg). Colorless prisms (recrystallized from MeOH), mp 90–91°C. *Anal.* Calcd for  $\text{C}_{19}\text{H}_{14}\text{N}_2\text{O}_2$ : C, 75.48; H, 4.67; N, 9.27. Found: C, 75.43; H, 4.63; N, 9.21.  $^1\text{H-NMR}$  ( $\text{CDCl}_3$ ): 7.27–8.70 (10H, m, aromatic H), 6.42 (1H, d,  $J=4\text{ Hz}$ ), 6.31 (1H, dd,  $J=4, 2\text{ Hz}$ ).

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