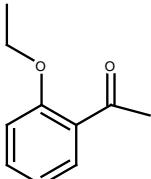
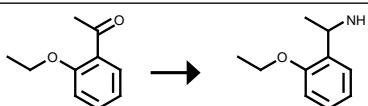


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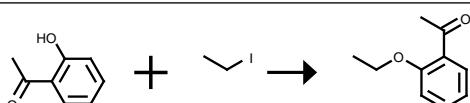
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1. Query		35 reactions	2011-09-12 10h:44m:30s (EST)

Search as: As drawn, No isotopes, No radicals



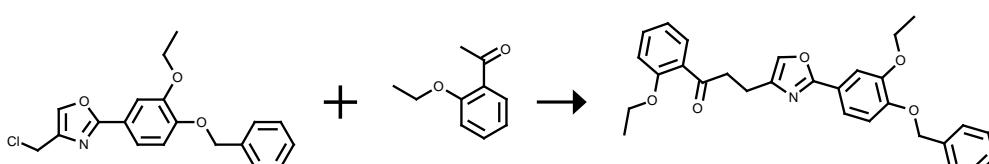
Rx-ID: 30426357 [View in Reaxys](#)

Yield	Conditions & References
	<p>Example Name 4.1          Synthesis of N-ethyl-N-[1-(2-ethoxyphenyl)ethyl]-ethane-1,2-diamine (1) 2'-Ethoxyacetophenone (2.0 g) was dissolved in methanol (20 ml), ammonium acetate (9.4 g) was added to the solution, and the resulting mixture was stirred at room temperature for 1 hour. Sodium cyanoborohydride (536 mg) was added to the mixture, and the resulting mixture was stirred at room temperature for 2.5 days. Saturated aqueous sodium hydrogencarbonate and 5 N aqueous sodium hydroxide were added to the reaction mixture, and the resulting mixture was extracted with chloroform. The organic layer was dried over anhydrous sodium sulfate, and filtered. The filtrate was concentrated under reduced pressure, and the resulting residue was purified by silica gel column chromatography (chloroform to chloroform:methanol:28 percent aqueous ammonia = 10:1:0.1) to obtain an amine compound (1.46 g)</p> <p><b>Stage 1:</b> With ammonium acetate in methanol, Time= 1h, T= 20 °C  <b>Stage 2:</b> With sodium cyanoborohydride in methanol, Time= 60h, T= 20 °C  <b>Stage 3:</b> With water, sodium hydrogencarbonate, sodium hydroxide in methanol</p> <p><b>Patent:</b> Taisho Pharmaceutical Co., Ltd.; Meiji Seika Kaisha, Ltd.; EP2287173; (2011); (A1) English  <a href="#">View in Reaxys</a></p>



Rx-ID: 2412788 [View in Reaxys](#)

Yield	Conditions & References
89 %	<p><b>With</b> potassium phosphate in dimethyl sulfoxide, T= 60 °C</p> <p><b>Horaguchi, Takaaki; Tsukada, Chikara; Hasegawa, Eietsu; Shimizu, Takahachi; Suzuki, Tsuneo; Tanemura, Kiyoshi;</b> Journal of Heterocyclic Chemistry; <b>vol.</b> 28; nb. 5; (1991); p. 1261 - 1272  <a href="#">View in Reaxys</a></p>
	<p>(i) aq. &lt;Et<sub>4</sub>N&gt;F, DMF, (ii) /BRN= 505934/, Multistep reaction</p> <p><b>Miller, J.M. et al.;</b> Canadian Journal of Chemistry; <b>vol.</b> 57; (1979); p. 1887 - 1889  <a href="#">View in Reaxys</a></p>

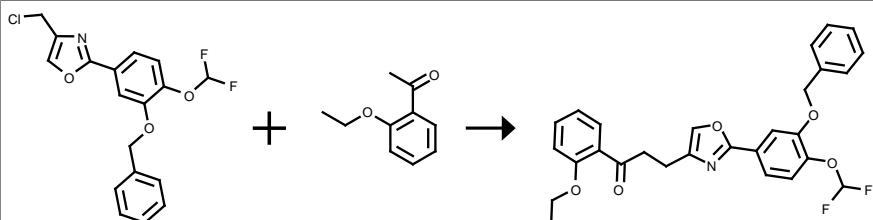


Rx-ID: 10882459 [View in Reaxys](#)

Yield	Conditions & References
	<p>Example Name 190          A 2 g quantity of 4-chloromethyl-2- (4-benzyloxy-3- ethoxyphenyl)oxazole obtained in Reference Example 63 and 0.96 g of 1- (2-ethoxyphenyl) ethanone were dissolved in 20 ml of tetrahydrofuran, and 0.47 g sodium hydride was added thereto. After foaming, the reaction mixture was heated and refluxed for 3 hours. After cooling, the reaction mixture was added to ice water, and extraction was performed with ethyl acetate. The organic layer was washed with water, dried over magnesium sulfate, and concentrated under reduced pressure. The residue was purified by silica gel column chromatography (n-hexane : ethyl acetate = 3 : 1) to give 0.4 g of colorless powdery 3- [2- (4-benzyloxy-3- ethoxyphenyl) oxazol-4-yl] -1- (2-ethoxyphenyl) propan-1-one. <sup>1</sup>H-NMR (CDCl<sub>3</sub>) δ: 7.70 (1H, dd, J = 7.5, 1.8 Hz), 7.55-7.30 (8H, &lt;n="139"/&gt;m), 6.97 (2H, t, J = 7.5 Hz), 6.93 (1H, d, J = 7.5 Hz), 5.19 (2H, s), 4.18 (2H, q, J = 6.9 Hz), 4.13 (2H, q, J = 6.9 Hz), 3.41 (2H, t, J = 6.9 Hz), 2.99 (2H, t, J = 6.9 Hz), 1.48 (3H, t, J = 6.9 Hz) , 1.47 (3H, t, J = 6.9 Hz)</p>

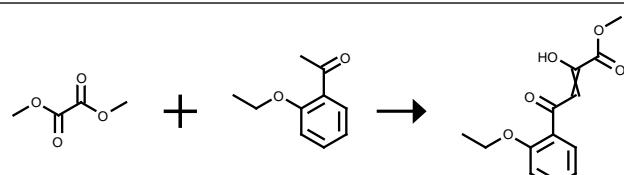
With sodium hydride in tetrahydrofuran, Time= 3h, Heating / reflux

Patent: OTSUKA PHARMACEUTICAL CO., LTD.; WO2007/58338; (2007); (A2) English  
[View in Reaxys](#)



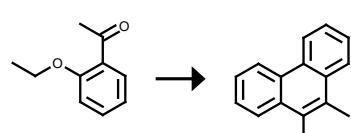
Rx-ID: 10882499 [View in Reaxys](#)

Yield	Conditions & References
	<p>Example Name 228</p> <p>A 65 mg quantity of sodium hydride was suspended in 5 ml of tetrahydrofuran. A 0.27 g quantity of I-(2- ethoxyphenyl) ethanone and 0.3 g of 2- (3-benzyloxy-4-difluoro methoxyphenyl)-4-chloromethyloxazole obtained in Reference Example 44 was successively added thereto with ice-cooling and stirring, and the mixture was stirred for 3 hours with heating and refluxing. An aqueous saturated ammonium chloride solution was added to the reaction mixture with ice-cooling and stirring. After stirring for 15 minutes, water was added thereto, and extraction was performed with ethyl acetate. The mixture was dried over anhydrous magnesium sulfate, and the solvent was removed. The obtained residue was purified by silica gel column chromatography (n-hexane : ethyl acetate = 4 : 1) to give 75 mg of colorless oily 3-[2- (3-benzyloxy-4-difluoromethoxyphenyl) oxazol-4-yl] -1- (2-ethoxyphenyl)propan-I-one. <sup>1</sup>H-NMR (CDCl<sub>3</sub>) δ: 7.72-7.69 (2H, m) , 7.59 (1H, dd, J = 8.1, 1.8 Hz), 7.47-7.32 (7H, m) , 7.00-6.92 (3H, m) , 6.61 (1H, t, J = 74.7 Hz), 5.20 (2H, s), 4.15 (2H, q, J = 7.2 Hz), 3.43 (2H, t, J = 7.2 Hz), 3.00 (2H, t, J = 7.2 Hz), 1.48 (3H, t, J = 7.2 Hz)</p> <p>With sodium hydride in tetrahydrofuran, Time= 3h, Heating / reflux</p> <p>Patent: OTSUKA PHARMACEUTICAL CO., LTD.; WO2007/58338; (2007); (A2) English  <a href="#">View in Reaxys</a></p>



Rx-ID: 30732532 [View in Reaxys](#)

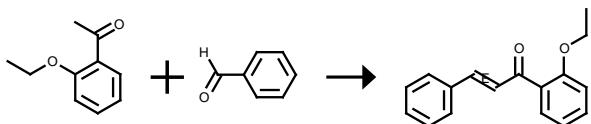
Yield	Conditions & References
46 %	<p><b>Stage 1:</b> With sodium methylate in methanol, Time= 0.25h, Claisen condensation</p> <p><b>Stage 2:</b> in methanol, Time= 0.083333h, T= 30 °C , Microwave irradiation, Claisen condensation</p> <p>Reddy, Tummala R. K.; Li, Chan; Guo, Xiaoxia; Myrvang, Helene K.; Fischer, Peter M.; Dekker, Lodewijk V.; Journal of Medicinal Chemistry; vol. 54; nb. 7; (2011); p. 2080 - 2094  <a href="#">View in Reaxys</a></p>



Rx-ID: 2116259 [View in Reaxys](#)

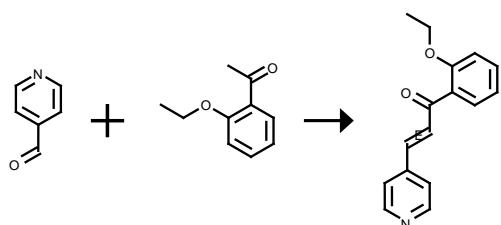
Yield	Conditions & References
80 %	<p>With pyridine, titanium(III) chloride, lithium in tetrahydrofuran, Time= 16h, Heating</p> <p>Kadam, Suresh M.; Nayak, Sandip K.; Banerji, Asoke; Synthetic Communications; vol. 25; nb. 2; (1995); p. 135 - 142</p>

	<a href="#">View in Reaxys</a>
38 %	<p><b>With</b> titanium(III) chloride, lithium <b>in</b> tetrahydrofuran</p> <p><b>Banerji, Asoke; Nayak, Sandip, K.</b>; Journal of the Chemical Society, Chemical Communications; nb. 20; (1991); p. 1432 - 1434</p> <a href="#">View in Reaxys</a>



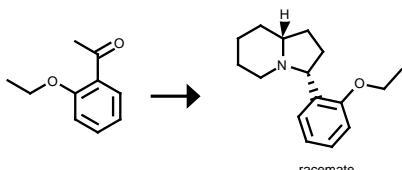
Rx-ID: 2680461 [View in Reaxys](#)

Yield	Conditions & References
69 %	<p><b>With</b> sodium hydroxide <b>in</b> ethanol, Time= 24h, Ambient temperature</p> <p><b>Batt, Douglas G.; Goodman, Robin; Jones, David G.; Kerr, Janet S.; Mantegna, Lisa R.; et al.</b>; Journal of Medicinal Chemistry; <b>vol.</b> 36; nb. 10; (1993); p. 1434 - 1442</p> <p><a href="#">View in Reaxys</a></p>



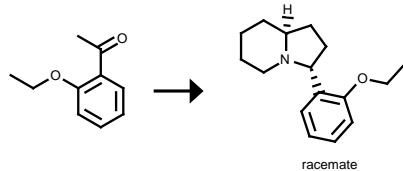
Rx-ID: 1524093 [View in Reaxys](#)

Yield	Conditions & References
43 %	<p><b>With sodium hydroxide in ethanol, Time= 24h, Ambient temperature</b></p> <p><b>Batt, Douglas G.; Goodman, Robin; Jones, David G.; Kerr, Janet S.; Mantegna, Lisa R.; et al.</b>; Journal of Medicinal Chemistry; <b>vol.</b> 36; nb. 10; (1993); p. 1434 - 1442</p> <p><a href="#">View in Reaxys</a></p>



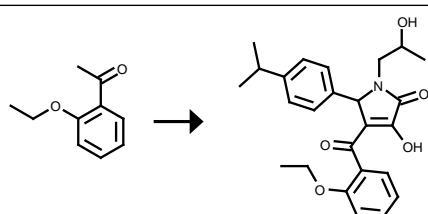
Bx-ID: 19593783 [View in Reaxys](#)

Yield	Conditions & References
	<p>Reaction Steps: 2 1: 39 percent / 0.42 h / Heating 2: 21 percent / H<sub>2</sub> / Pt <b>With</b> hydrogen, platinum</p> <p><b>Carson, John R.; Carmosin, Richard J.; Vaught, Jeffry L.; Gardocki, Joseph F.; Costanzo, Michael J.; et al.</b>; Journal of Medicinal Chemistry; <b>vol.</b> 35; nb. 15; (1992); p. 2855 - 2863</p> <p><a href="#">View in Reaxys</a></p>



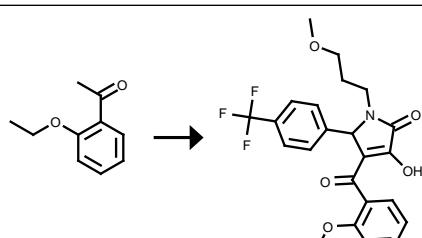
Rx-ID: 19593784 [View in Reaxys](#)

Yield	Conditions & References
	<p>Reaction Steps: 2</p> <p>1: 39 percent / 0.42 h / Heating</p> <p>2: H<sub>2</sub> / Pt</p> <p><b>With</b> hydrogen, platinum</p> <p><b>Carson, John R.; Carmosin, Richard J.; Vaught, Jeffry L.; Gardocki, Joseph F.; Costanzo, Michael J.; et al.</b>; Journal of Medicinal Chemistry; <b>vol.</b> 35; nb. 15; (1992); p. 2855 - 2863  <a href="#">View in Reaxys</a></p>



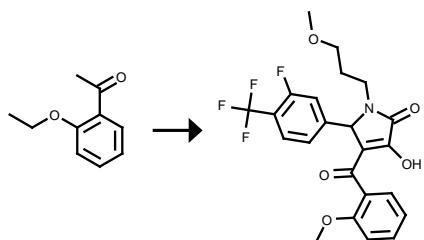
Rx-ID: 30732609 [View in Reaxys](#)

Yield	Conditions & References
	<p>Reaction Steps: 2</p> <p>1.1: sodium methylate / methanol / 0.25 h</p> <p>1.2: 0.08 h / 30 °C / Microwave irradiation</p> <p>2.1: 1,4-dioxane / 20 °C</p> <p><b>With</b> sodium methylate <b>in</b> 1,4-dioxane, methanol, 1.1: Claisen condensation / 1.2: Claisen condensation</p> <p><b>Reddy, Tummala R. K.; Li, Chan; Guo, Xiaoxia; Myrvang, Helene K.; Fischer, Peter M.; Dekker, Lodewijk V.</b>; Journal of Medicinal Chemistry; <b>vol.</b> 54; nb. 7; (2011); p. 2080 - 2094  <a href="#">View in Reaxys</a></p>



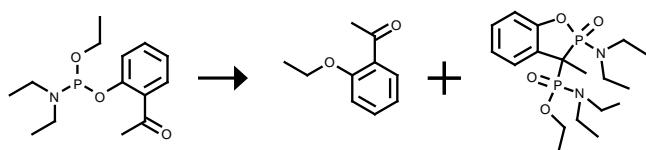
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Yield	Conditions & References
	<p>Reaction Steps: 2</p> <p>1.1: sodium methylate / methanol / 0.25 h</p> <p>1.2: 0.08 h / 30 °C / Microwave irradiation</p> <p>2.1: 1,4-dioxane / 20 °C</p> <p><b>With</b> sodium methylate <b>in</b> 1,4-dioxane, methanol, 1.1: Claisen condensation / 1.2: Claisen condensation</p> <p><b>Reddy, Tummala R. K.; Li, Chan; Guo, Xiaoxia; Myrvang, Helene K.; Fischer, Peter M.; Dekker, Lodewijk V.</b>; Journal of Medicinal Chemistry; <b>vol.</b> 54; nb. 7; (2011); p. 2080 - 2094  <a href="#">View in Reaxys</a></p>



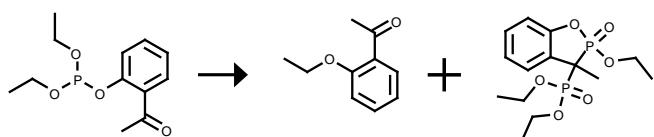
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Yield	Conditions & References
	<p>Reaction Steps: 2</p> <p>1.1: sodium methylate / methanol / 0.25 h</p> <p>1.2: 0.08 h / 30 °C / Microwave irradiation</p> <p>2.1: 1,4-dioxane / 20 °C</p> <p><b>With</b> sodium methylate <b>in</b> 1,4-dioxane, methanol, 1.1: Claisen condensation / 1.2: Claisen condensation</p> <p><b>Reddy, Tummala R. K.; Li, Chan; Guo, Xiaoxia; Myrvang, Helene K.; Fischer, Peter M.; Dekker, Lodewijk V.</b>; Journal of Medicinal Chemistry; <b>vol.</b> 54; nb. 7; (2011); p. 2080 - 2094</p> <p><a href="#">View in Reaxys</a></p>



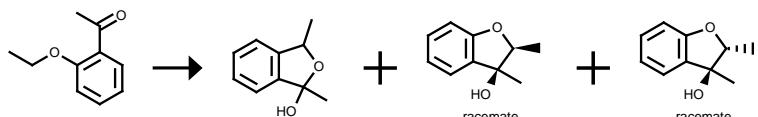
Rx-ID: 2225806 [View in Reaxys](#)

Yield	Conditions & References
72 %, 53 %	<p>Time= 6.5h, T= 110 °C</p> <p><b>Mukhametov, F. S.; Korshin, E. E.; Nekhoroshkov, V. M.; Efremov, Yu. Ya.</b>; J. Gen. Chem. USSR (Engl. Transl.); <b>vol.</b> 59; nb. 6.1; (1989); p. 1309 - 1320,1159 - 1169</p> <p><a href="#">View in Reaxys</a></p>
53 %, 72 %	<p>Time= 6.5h, T= 110 °C</p> <p><b>Mukhametov, F. S.; Korshin, E. E.; Nekhoroshkov, V. M.; Efremov, Yu. Ya.</b>; J. Gen. Chem. USSR (Engl. Transl.); <b>vol.</b> 59; nb. 6.1; (1989); p. 1309 - 1320,1159 - 1169</p> <p><a href="#">View in Reaxys</a></p>



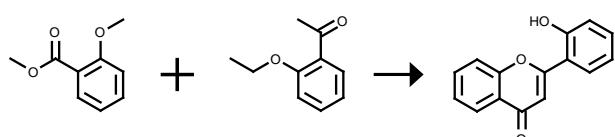
Rx-ID: 3663545 [View in Reaxys](#)

Yield	Conditions & References
60 %, 61.5 %	<p>Time= 1.33333h, T= 100 - 110 °C</p> <p><b>Mukhametov, F. S.; Korshin, E. E.; Korshunov, R. L.; Efremov, Yu. Ya.; Zyablikova, T. A.</b>; J. Gen. Chem. USSR (Engl. Transl.); <b>vol.</b> 56; nb. 8; (1986); p. 1781 - 1789,1574 - 1580</p> <p><a href="#">View in Reaxys</a></p>



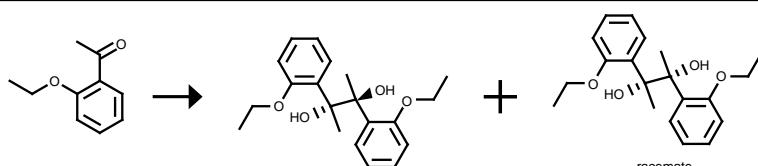
Rx-ID: 2116260 [View in Reaxys](#)

Yield	Conditions & References
46 %	<b>in acetonitrile, Time= 1h, Irradiation, Yields of byproduct given</b> <b>Horaguchi, Takaaki; Tsukada, Chikara; Hasegawa, Eietsu; Shimizu, Takahachi; Suzuki, Tsuneo; Tanemura, Kiyoshi;</b> Journal of Heterocyclic Chemistry; <b>vol. 28; nb. 5; (1991); p. 1261 - 1272</b> <a href="#">View in Reaxys</a>
46 %	<b>in acetonitrile, Time= 1h, Irradiation, Yield given. Yields of byproduct given</b> <b>Horaguchi, Takaaki; Tsukada, Chikara; Hasegawa, Eietsu; Shimizu, Takahachi; Suzuki, Tsuneo; Tanemura, Kiyoshi;</b> Journal of Heterocyclic Chemistry; <b>vol. 28; nb. 5; (1991); p. 1261 - 1272</b> <a href="#">View in Reaxys</a>



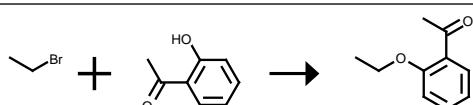
Rx-ID: 332971 [View in Reaxys](#)

Yield	Conditions & References
	<b>With</b> sodium, $T = 130\text{ }^{\circ}\text{C}$ , Behandeln des entstandenen 2-Methoxy-2'-aethoxy-dibenzoylmethans mit Jodwasserstoffsaeure <b>Pistermann; Tambor;</b> Chemische Berichte; <b>vol. 45; (1912); p. 1241</b> <a href="#">View in Reaxys</a>



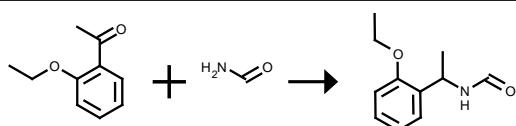
Rx-ID: 10402990 [View in Reaxys](#)

Yield	Conditions & References
115 mg, 100 mg	<b>With</b> triethylamine <b>in</b> ethanol, Irradiation <b>Singh, Rajinder; Garg, P. K.; Hundal, M. S.; Ishar, M. P. S.;</b> Indian Journal of Chemistry, Section B: Organic Chemistry Including Medicinal Chemistry; <b>vol. 45; nb. 2; (2006); p. 506 - 509</b> <a href="#">View in Reaxys</a>



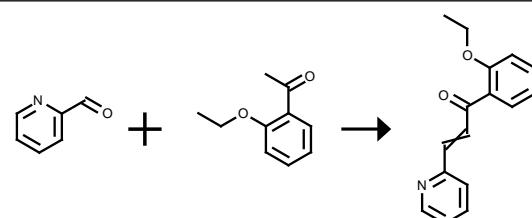
Rx-ID: 10414040 [View in Reaxys](#)

Yield	Conditions & References
2.5 g	<b>With</b> potassium carbonate <b>in</b> acetone, Time= 10h, Heating <b>Singh, Rajinder; Garg, P. K.; Hundal, M. S.; Ishar, M. P. S.;</b> Indian Journal of Chemistry, Section B: Organic Chemistry Including Medicinal Chemistry; <b>vol. 45; nb. 2; (2006); p. 506 - 509</b> <a href="#">View in Reaxys</a>



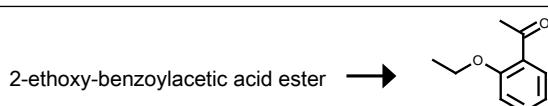
Rx-ID: 29783693 [View in Reaxys](#)

Yield	Conditions & References
	<p>in formic acid, Time= 30h, T= 130 °C , Leuckart-Wallach reaction</p> <p><b>Musatov, D. M.; Starodubtseva, E. V.; Turova, O. V.; Kurilov, D. V.; Vinogradov, M. G.; Rakishev, A. K.; Struchkova, M. I.</b>; Russian Journal of Organic Chemistry; <b>vol. 46</b>; nb. 7; (2010); p. 1021 - 1028; Zhurnal Organicheskoi Khimii; <b>vol. 46</b>; nb. 7; (2010); p. 1023 - 1029</p> <a href="#">View in Reaxys</a>



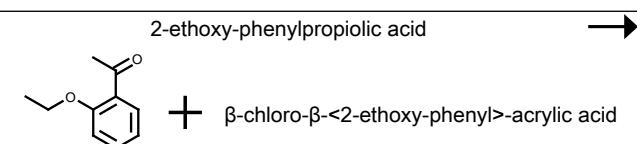
Rx-ID: 1523580 [View in Reaxys](#)

Yield	Conditions & References
39 %	<p>Time= 0.416667h, Heating</p> <p><b>Carson, John R.; Carmosin, Richard J.; Vaught, Jeffry L.; Gardocki, Joseph F.; Costanzo, Michael J.; et al.</b>; Journal of Medicinal Chemistry; <b>vol. 35</b>; nb. 15; (1992); p. 2855 - 2863</p> <a href="#">View in Reaxys</a>



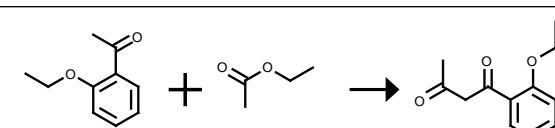
Rx-ID: 7388666 [View in Reaxys](#)

Yield	Conditions & References
	<p><b>With</b> sulfuric acid</p> <p><b>Besthorn; Banzhaf; Jaegle</b>; Chemische Berichte; <b>vol. 27</b>; (1894); p. 3036</p> <a href="#">View in Reaxys</a>



Rx-ID: 7388667 [View in Reaxys](#)

Yield	Conditions & References
	<p><b>With</b> hydrogenchloride</p> <p><b>Fittig; Claus,R.</b>; Justus Liebigs Annalen der Chemie; <b>vol. 269</b>; (1892); p. 3</p> <a href="#">View in Reaxys</a>

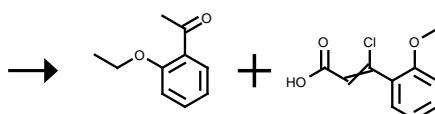


Rx-ID: 646579 [View in Reaxys](#)

Yield	Conditions & References
	<p><b>With</b> sodium ethanolate, Zersetzen das Natriumsalz in waessr.Loesung mit Essigsaeure</p> <p><b>Patent; Hoechster Farbw.</b>; DE79173; Fortschr. Teerfarbenfabr. Verw. Industriezweige; <b>vol. 4</b>; p. 1141</p>

[View in Reaxys](#)

Besthorn; Banzhaf; Jaegle; Chemische Berichte; vol. 27; (1894); p. 3036

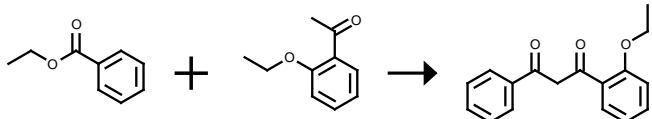
[View in Reaxys](#)

 Rx-ID: 8071939 [View in Reaxys](#)

Yield

Conditions &amp; References

**With** hydrogenchloride

**Fittig; Claus,R.;** Justus Liebigs Annalen der Chemie; vol. 269; (1892); p. 3

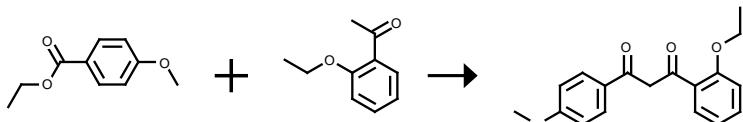
[View in Reaxys](#)

 Rx-ID: 281441 [View in Reaxys](#)

Yield

Conditions &amp; References

**With** sodium

**v.Kostanecki; Tambor;** Chemische Berichte; vol. 33; (1900); p. 333 Anm.2

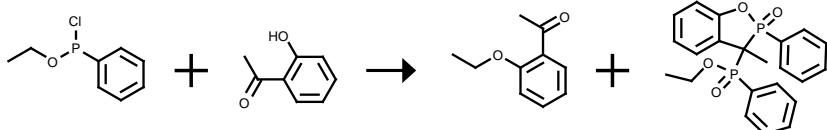
[View in Reaxys](#)

 Rx-ID: 334429 [View in Reaxys](#)

Yield

Conditions &amp; References

**With** sodium

**Tasaki;** Acta Phytochimica; vol. 3; (1927); p. 275; Chem. Zentralbl.; vol. 98; nb. II; (1927); p. 1949

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 Rx-ID: 2412700 [View in Reaxys](#)

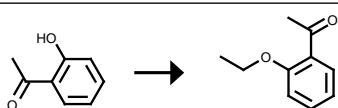
Yield

Conditions &amp; References

**With** triethylamine, 1.) ether, 1 d; 2.) 2 d, RT; 2 h, 80 deg C, Yield given. Multistep reaction. Yields of byproduct given

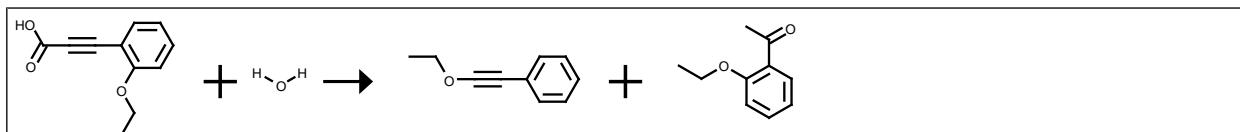
**Mukhametov, F. S.; Korshin, E. E.; Korshunov, R. L.; Efremov, Yu. Ya.; Zyablikova, T. A.**; J. Gen. Chem. USSR (Engl. Transl.); vol. 56; nb. 8; (1986); p. 1781 - 1789, 1574 - 1580

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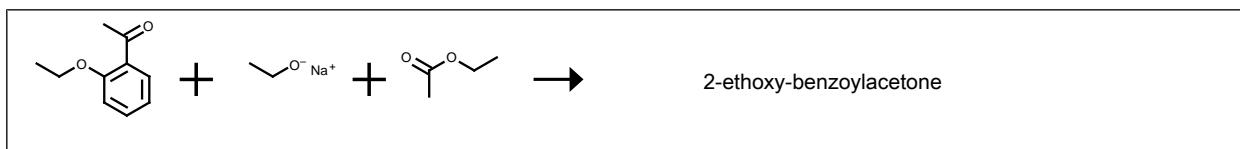
Rx-ID: 21973349 [View in Reaxys](#)

Yield	Conditions & References
	<p>Reaction Steps: 2          1: 77.5 percent / 1) 8h, 50 deg C, 50 mm, 2) 1h, 0.05 mm          2: 53 percent / 6.5 h / 110 °C</p> <p><b>Mukhametov, F. S.; Korshin, E. E.; Nekhoroshkov, V. M.; Efremov, Yu. Ya.</b>; J. Gen. Chem. USSR (Engl. Transl.); <b>vol. 59</b>; nb. 6.1; (1989); p. 1309 - 1320, 1159 - 1169  <a href="#">View in Reaxys</a></p>



Rx-ID: 7990352 [View in Reaxys](#)

Yield	Conditions & References
	<p>T= 140 - 150 °C</p> <p><b>Fittig; Claus,R.</b>; Justus Liebigs Annalen der Chemie; <b>vol. 269</b>; (1892); p. 3  <a href="#">View in Reaxys</a></p>



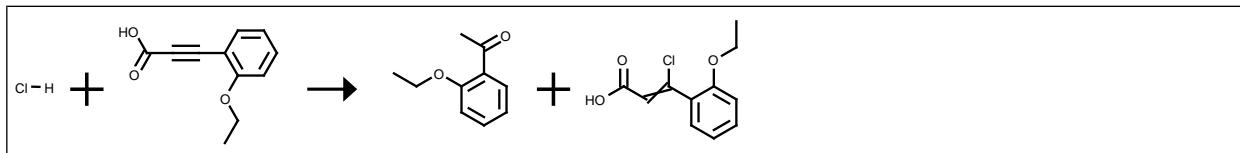
Rx-ID: 7434855 [View in Reaxys](#)

Yield	Conditions & References
	<p><b>Besthorn; Banzhaf; Jaegle</b>; Chemische Berichte; <b>vol. 27</b>; (1894); p. 3036  <a href="#">View in Reaxys</a></p>



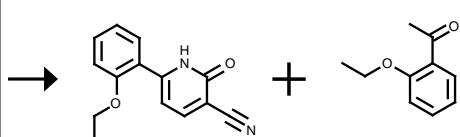
Rx-ID: 7990351 [View in Reaxys](#)

Yield	Conditions & References
	<p><b>Fittig; Claus,R.</b>; Justus Liebigs Annalen der Chemie; <b>vol. 269</b>; (1892); p. 3  <a href="#">View in Reaxys</a></p>



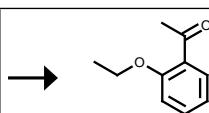
Rx-ID: 8071940 [View in Reaxys](#)

Yield	Conditions & References
	<p><b>Fittig; Claus,R.</b>; Justus Liebigs Annalen der Chemie; <b>vol. 269</b>; (1892); p. 3  <a href="#">View in Reaxys</a></p>



Rx-ID: 24295662 [View in Reaxys](#)

Yield	Conditions & References
	<p>Example Name 34  Example Title 3-Cyano-6-(2-ethoxyphenyl)-2(1H)-pyridinone  EXAMPLE 34  3-Cyano-6-(2-ethoxyphenyl)-2(1H)-pyridinone  In a similar manner to that of Example 26, 2-ethoxy-acetophenone (2 g) yielded the title compound, 0.32 g, m.p. 262.deg.-4.deg. C. (from dimethylformamide).</p> <p><b>Patent:</b> Cell Pathways, Inc.; US6046216; (2000); (A1) English  <a href="#">View in Reaxys</a></p>



Rx-ID: 7388665 [View in Reaxys](#)

Yield	Conditions & References
	<p>2-Acetyl-phenol, Et<sub>2</sub>SO<sub>4</sub>/NaOH</p> <p><b>Dhami; Stothers:</b> Canadian Journal of Chemistry; <b>vol.</b> 43; (1965); p. 479,482  <a href="#">View in Reaxys</a></p>