Hexadecyldimethyl benzyl ammonium bromide: an efficient catalyst for a clean one-pot synthesis of tetrahydrobenzopyran derivatives in water

Tong-Shou Jin*, Ai-Qing Wang, Feng Shi, Li-Sha Han, Li-Bin Liu, and Tong-Shuang Li

College of Chemistry and Environmental Science, Hebei University, Baoding 071002, P. R. China

E-mail: jintongshou@yahoo.com.cn

Abstract

A clean and efficient method for the synthesis of 2-amino-4-aryl-3-cyano-5-oxo- 4*H*-5,6,7,8-tetrahydrobenzopyran derivatives using hexadecyldimethylbenzyl ammonium bromide (HDMBAB) as the catalyst is described. This method provides several advantages, such as simple work-up procedure, environmentally benign, neutral conditions and high yields. In addition, water was chosen as a green solvent.

Keywords: Tetrahydrobenzopyran, aromatic aldehyde, malononitrile, 1,3-cyclohexanedione, 5,5-dimethyl-1,3-cyclohexanedione; aqueous media

Introduction

In recent years, 4*H*-benzopyran and its derivatives have attracted strong interest due to their useful biological and pharmacological properties, such as anticoagulant, spasmolytic, diuretic, anticancer, antianaphylactin agents.¹ Some 2-amino-4*H*-pyrans can be employed as photoactive materials.² Furthermore, substituted 4*H*-pyrans also constitute a structural unit of a series of natural products.^{3,4} The synthesis of 4*H*-benzopyrans has been reported;⁵ the conventional methods are with base as catalyst (piperidine or triethylamine) in an organic solvent such as ethanol, acetic acid or DMF,⁶⁻⁸ and using ultrasonic irradiation.⁹ Each of the above methods has its own merit, but some are not entirely satisfactory, owing to such drawbacks as low yields, difficult work-up, problems of corrosiveness and effluent pollution. Consequently, there is a need to develop alternative methods for the synthesis of tetrahydrobenzopyran derivatives under mild and environmentally friendly conditions.

Water has been applied to organic reactions as a solvent, and it has several advantages such as its low cost, safety, non-polluting nature and operational simplicity.¹⁰ In 1983, Breslow discovered that the Diels-Alder reaction performed in water was subject to a huge rate

acceleration.¹¹ This observation led to increased interest from synthetic organic chemists in organic reactions conducted in water. Soon it was discovered that other organic reactions, like the Claisen rearrangement,¹² the aldol condensation,¹³ Diels-Alder reaction,¹⁴ the benzoin condensation,¹⁵ Mannich reaction¹⁶ and Michael reaction¹⁷ exhibit rate enhancements in water. To date, many more organic reactions have been carried out in water.¹⁸

In this manuscript, we wish to report a general and highly efficient route for the synthesis of tetrahydrobenzopyrans using an inexpensive, commercially available hexadecyldimethylbenzyl ammonium bromide (HDMBAB) catalyst. This is a one-pot reaction in aqueous media, which is not only operationally simple but also consistently gives the corresponding products in good to excellent yields (Scheme 1).

Scheme 1

In a typical general experimental procedure, an aqueous solution of 1,3-cyclohexanedione or 5,5-dimethyl-1,3-cyclohexanedione **1**, aromatic aldehyde **2** and malononitrile **3** was heated at 80-90 °C in the presence of a catalytic amount of HDMBAB. The corresponding 2-amino-4-aryl-3-cyano-5-oxo-4*H*-5,6,7,8-tetrahydrobenzopyran derivatives **4** were obtained in good to excellent yields and the results are summarized in Table 1.

As shown in Table 1, the three-component cyclocondensation reaction proceeded smoothly at 80-90 °C in water to give the corresponding products **4** in high yields. The electronic nature of the substituents on the aromatic ring did not show a strong effect in terms of yields under these reaction conditions. Both aromatic aldehydes containing electron-withdrawing groups (such as nitro group, halide) or electron-donating groups (such as alkyl group, alkoxy group) gave good to excellent yields of the corresponding 4*H*-tetrahydrobenzopyrans.

The catalyst plays a crucial role in the success of the reaction in terms of the rate and the yields. Taking the reaction of 4-nitrobenzaldehyde with 1,3-cyclohexanedione **2** and **3** as an example, the reaction could be carried out in the absence of the catalyst when the mixture was heated in water for 6h, but a very poor yield resulted (18%). We have tested some catalysts such as tetrabutyl ammonium bromide, benzyltrimethyl ammonium chloride, sodium dodecyl sulfate and hexadecyldimethylbenzyl ammonium bromide, and the yields using these four catalysts were 78%, 86%, 65% and 96%, respectively. From the yield data, it was found that

hexadecyldimethylbenzyl ammonium bromide (HDMBAB) was the best catalyst for this reaction.

Table1. Synthesis of 4H-tetrahydrobenzopyrans catalyzed by HDMBAB in water

Entry	Ar	R	Time (h)	Product	Yield ^a (%)	Mp(°C)	
						Found	Reported ¹⁹
1	2-ClC ₆ H ₄ 1a	H	6.0	4a	92	210-212	213-215
2	4-ClC ₆ H ₄ 1b	H	6.0	4b	94	224-226	226-229
3	2,4-Cl ₂ C ₆ H ₃ 1c	H	6.0	4c	95	223-225	225-227
4	3-NO ₂ C ₆ H ₄ 1d	H	5.5	4d	96	202-204	198-200
5	4-NO ₂ C ₆ H ₄ 1e	H	5.5	4e	96	234-236	234-235
6	4-CH ₃ OC ₆ H ₄ 1f	H	7.0	4f	88	192-194	193-195
7	3,4-OCH ₂ OC ₆ H ₃ 1g	H	7.0	4g	86	210-212	211-214
8	2-ClC ₆ H ₄ 1h	CH ₃	7.5	4h	88	216-218	215-216
9	3-ClC ₆ H ₄ 1i	CH ₃	7.5	4i	89	222-224	224-225
10	4-ClC ₆ H ₄ 1j	CH ₃	7.5	4j	90	237-239	239-241
11	2-NO ₂ C ₆ H ₄ 1k	CH ₃	7.0	4k	90	182-184	180-182
12	3-NO ₂ C ₆ H ₄ 1l	CH ₃	7.0	4l	92	212-214	213-214
13	4-NO ₂ C ₆ H ₄ 1m	CH ₃	7.0	4m	93	178-180	176-178
14	4-CH ₃ OC ₆ H ₄ 1n	CH ₃	8.0	4n	86	194-196	196-198
15	4-CH ₃ C ₆ H ₄ 1o	CH ₃	8.0	4o	84	218-220	220-222
16	3,4-OCH ₂ OC ₆ H ₃ 1p	CH ₃	8.0	4p	85	220-222	221-223

^a Isolated yield.

We have studied also the effect of the amount of the catalyst on these reactions. With the mixture **1e**, 1,3-cyclohexanedione and **3** in the presence of 3mol% HDMBAB, the product **4e** was obtained in 77% yield at 80-90 °C in water for 5.5h. Increasing the amount of catalyst to 5mol%, 8mol% and 12mol%, yields of 84%, 88% and 96%, respectively, were obtained. Use of just 12mol% HDMBAB is sufficient to push the reaction forward and higher amounts of the catalyst did not improve the results to any greater extent. Thus, 12mol% HDMBAB was chosen as the amount of catalyst for these reactions.

We propose the possible following mechanism to account for the reaction. First, the aromatic aldehyde **2** is condensed with malononitrile (**3**) to afford the α -cyanocinnamitrile derivative **5**. The step (**2**+**3**→**5**) can be regarded as a rapid Knoevenagel reaction. Since, in a model reaction, the Knoevenagel reaction of malononitrile and aromatic aldehydes can be carried out in water without any catalyst, we conjecture that the second step requires the presence of HDMBAB. The active methylene of **1** reacts with the electrophilic C=C double bond in **5** giving the intermediate **6**, which tautomerizes into **7**. The latter is then cyclized by nucleophilic attack of the OH group on the cyano (CN) moiety, giving intermediate **8**. Finally, the expected product **4** is afforded by tautomerization (**8**→**4**). HDMBAB not only is a phase transfer catalyst, but it serves also as an emulsifier in this reaction process (Scheme 2).

Scheme 2

In conclusion, we have described a procedure for the preparation of 4*H*-benzopyran derivatives catalyzed by HDMBAB, using a three-component condensation in heated water. In addition, it is possible to apply the tenets of green chemistry to the generation of biologically interesting products using aqueous media approaches that are less expensive and less toxic than

those with organic solvents. Moreover, the procedure offers several advantages including high yields, operational simplicity, clean reaction conditions and minimum pollution of the environment, which makes it a useful and attractive process for the synthesis of these compounds.

Experimental Section

General Procedures. IR spectra were recorded on a Bio-Rad FTS-40 spectrometer (KBr). ^1H NMR spectra were measured on a Bruker AVANCE 400 (400 MHz) spectrometer using TMS as internal reference and $\text{DMSO-}d_6$ as solvent. Elemental analyses were determined using Perkin-Elmer 2400 II elemental analyzer.

General procedure for synthesis of tetrahydrobenzopyran derivatives

A mixture of 5,5-dimethyl-1,3-cyclohexadione or 1,3-cyclohexadione (**1**, 5 mmol), aromatic aldehyde (**2**, 5 mmol), malononitrile (**3**, 5 mmol), and HDMBAB (12 mol%) in water (30 mL) was stirred at 80-90 °C for 5.5-8.0 h. Then the mixture was cooled to room temperature, solid was filtered off and washed with H_2O (40ml). The crude products were purified by recrystallization from ethanol (95%) to afford pure products **4**. Data for all compounds are shown below:

2-Amino-4-(2-chlorophenyl)-3-cyano-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran (4a). IR (KBr): $\nu_{\text{max}} = 3312, 3200, 2200, 1696, 1600, 1512, 1360, 1210, 1000, 750 \text{ cm}^{-1}$. ^1H NMR ($\text{DMSO-}d_6$): $\delta_{\text{H}} = 1.88\text{-}2.02$ (m, 2H, CH-7), $2.16\text{-}2.34$ (m, 2H, CH-8), $2.50\text{-}2.68$ (m, 2H, CH-6), 4.72 (s, 1H, CH-4), 7.00 (s, 2H, NH_2), $7.18\text{-}7.28$ (m, 3H, ArH), $7.34\text{-}7.36$ (m, 1H, ArH) ppm. Anal. calcd. for $\text{C}_{16}\text{H}_{13}\text{ClN}_2\text{O}_2$: C 63.90, H 4.36, N 9.31; found C 64.02, H 4.32, N 9.37 %.

2-Amino-4-(4-chlorophenyl)-3-cyano-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran (4b). IR (KBr): $\nu_{\text{max}} = 3416, 3340, 2200, 1684, 1606, 1600, 1495, 1360, 1260, 1206, 1000, 912, 780, 750 \text{ cm}^{-1}$. ^1H NMR ($\text{DMSO-}d_6$): $\delta_{\text{H}} = 1.88\text{-}2.00$ (m, 2H, CH-7), $2.22\text{-}2.34$ (m, 2H, CH-8), $2.60\text{-}2.72$ (m, 2H, CH-6), 4.22 (s, 1H, CH-4), 7.02 (s, 2H, NH_2), $7.18\text{-}7.20$ (m 2H, ArH), $7.32\text{-}7.36$ (m, 2H, ArH) ppm. Anal. calcd. for $\text{C}_{16}\text{H}_{13}\text{ClN}_2\text{O}_2$: C 63.90, H 4.36, N 9.31; found C 64.00, H 4.38, N 9.27 %.

2-Amino-4-(2,4-dichlorophenyl)-3-cyano-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran (4c). IR (KBr): $\nu_{\text{max}} = 3350, 3200, 2212, 1706, 1650, 1610, 1508, 1468, 1416, 1280, 1004, 810, 760 \text{ cm}^{-1}$. ^1H NMR ($\text{DMSO-}d_6$): $\delta_{\text{H}} = 1.94\text{-}2.00$ (m, 2H, CH-7), $2.24\text{-}2.34$ (m, 2H, CH-8), $2.56\text{-}2.66$ (m, 2H, CH-6), 4.70 (s, 1H, CH-4), 7.06 (s, 2H, NH_2), $7.22\text{-}7.24$ (m, 1H, ArH), $7.34\text{-}7.36$ (m, 1H, ArH), $7.50\text{-}7.52$ (m, 1H, ArH). Anal. calcd. for $\text{C}_{16}\text{H}_{12}\text{Cl}_2\text{N}_2\text{O}_2$: C 57.33, H 3.61, N 8.36; found C 57.42, H 3.66, N 8.43 %.

2-Amino-3-cyano-4-(3-nitrophenyl)-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran (4d). IR (KBr): $\nu_{\text{max}} = 3360, 3308, 2200, 2190, 1686, 1640, 1510, 1450, 1402, 1370, 1260, 1140, 850, 780, 750 \text{ cm}^{-1}$. ^1H NMR ($\text{DMSO-}d_6$): $\delta_{\text{H}} = 1.88\text{-}2.01$ (m, 2H, CH-7), $2.20\text{-}2.35$ (m, 2H, CH-8), $2.60\text{-}2.70$

(m, 2H, CH-6), 4.52 (s, 1H, CH-4), 7.15 (s, 2H, NH₂), 7.60-7.68 (m, 2H, ArH), 7.98 (s, 1H, ArH), 8.06-8.08 (m, 1H, ArH) ppm. Anal. calcd. for C₁₆H₁₃N₃O₄: C 61.73, H 4.21, N 13.50; found C 61.78, H 4.16, N 13.57 %.

2-Amino-3-cyano-4-(4-nitrophenyl)-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran (4e). IR (KBr): $\nu_{\max}$ = 3400, 3330, 2200, 1702, 1690, 1590, 1510, 1436, 1350, 1210, 1000, 820, 790, 740 cm⁻¹. ¹H NMR (DMSO-*d*₆): δ_{H} = 1.94-1.98 (m, 2H, CH-7), 2.28-2.32 (m, 2H, CH-8), 2.62-2.66 (m, 2H, CH-6), 4.38 (s, 1H, CH-4), 7.12 (s, 2H, NH₂), 7.44 (d, 2H, *J*=8.0Hz, ArH), 8.14 (d, 2H, *J*=8.0Hz, ArH) ppm. Anal. calcd. for C₁₆H₁₃N₃O₄: C 61.73, H 4.21, N 13.50; found C 61.81, H 4.26, N 13.57 %.

2-Amino-3-cyano-4-(4-methoxyphenyl)-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran (4f). IR (KBr): $\nu_{\max}$ = 3460, 3320, 2220, 1700, 1600, 1510, 1450, 1368, 1245, 1026, 840, 770, 750 cm⁻¹. ¹H NMR (DMSO-*d*₆): δ_{H} = 1.93-1.96 (m, 2H, CH-7), 2.26-2.30 (m, 2H, CH-8), 2.58-2.62 (m, 2H, CH-6), 3.74 (s, 3H, OCH₃), 4.20 (s, 1H, CH-4), 6.82 (d, 2H, *J*=8.0Hz, ArH), 6.98 (s, 2H, NH₂), 7.06 (d, 2H, *J*=8.0Hz, ArH) ppm. Anal. calcd. for C₁₇H₁₆N₂O₃: C 68.91, H 5.44, N 9.45; found C 69.00, H 5.47, N 9.49 %.

2-Amino-3-cyano-4-(3,4-methylenedioxyphenyl)-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran (4g). IR (KBr): $\nu_{\max}$ = 3360, 3196, 2962, 2195, 1650, 1605, 1490, 1378, 1360, 1250, 1210, 1124, 1036, 850, 780 cm⁻¹. ¹H NMR (DMSO-*d*₆): δ_{H} = 0.98 (s, 3H, CH₃), 1.02 (s, 3H, CH₃), 2.12 (d, 1H, *J*=16.0Hz, CH-8), 2.22 (d, 1H, *J*=16.0Hz, CH-8), 2.46-2.56 (m, 2H, CH-6), 4.15 (s, 1H, CH-4), 5.98 (s, 2H, OCH₂O), 6.63 (d, 1H, *J*=8.0Hz, ArH), 6.66 (s, 1H, ArH), 6.82 (d, 1H, *J*=8.0Hz, ArH), 6.98 (s, 2H, NH₂) ppm. Anal. calcd. for C₁₇H₁₄N₂O₄: C 65.80, H 4.55, N 9.03; found C 65.87, H 4.46, N 9.11 %.

2-Amino-4-(2-chlorophenyl)-3-cyano-7,7-dimethyl-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran (4h). IR (KBr): $\nu_{\max}$ = 3330, 3186, 2962, 2200, 1660, 1560 cm⁻¹. ¹H NMR (DMSO-*d*₆): δ_{H} = 0.98 (s, 3H, CH₃), 1.02 (s, 3H, CH₃), 2.18 (d, 1H, *J*=16.0Hz, CH-8), 2.24 (d, 1H, *J*=16.0Hz, CH-8), 2.46-2.56 (m, 2H, CH-6), 4.50 (s, 1H, CH-4), 7.02 (s, 2H, NH₂), 7.16-7.36 (m, 4H, ArH) ppm. Anal. calcd. for C₁₈H₁₇ClN₂O₂: C 65.75, H 5.21, N 8.52; found C 65.71, H 5.32, N 8.58 %.

2-Amino-4-(3-chlorophenyl)-3-cyano-7,7-dimethyl-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran (4i). IR (KBr): $\nu_{\max}$ = 3390, 3290, 3180, 2190, 1660, 1584 cm⁻¹. ¹H NMR (DMSO-*d*₆): δ_{H} = 1.08 (s, 3H, CH₃), 1.12 (s, 3H, CH₃), 2.26 (s, 2H, CH-8), 2.46-2.48 (m, 2H, CH-6), 4.43 (s, 1H, CH-4), 6.58 (s, 2H, NH₂), 7.18-7.28 (m, 4H, ArH) ppm. Anal. calcd. for C₁₈H₁₇ClN₂O₂: C 65.75, H 5.21, N 8.52; found C 65.78, H 5.31, N 8.49 %.

2-Amino-4-(4-chlorophenyl)-3-cyano-7,7-dimethyl-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran (4j). IR (KBr): $\nu_{\max}$ = 3408, 3310, 3006, 2200, 1694, 1606 cm⁻¹. ¹H NMR (DMSO-*d*₆): δ_{H} = 0.97 (s, 3H, CH₃), 1.05 (s, 3H, CH₃), 2.12 (d, 1H, *J*=16.0Hz, CH-8), 2.22 (d, 1H, *J*=16.0Hz, CH-8), 2.48-2.54 (m, 2H, CH-6), 4.28 (s, 1H, CH-4), 7.04 (s, 2H, NH₂), 7.16 (d, 2H, *J*=8.0Hz, ArH), 7.34 (d, 2H, *J*=8.0Hz, ArH) ppm. Anal. calcd. for C₁₈H₁₇ClN₂O₂: C 65.75, H 5.21, N 8.52; found C 65.79, H 5.28, N 8.47 %.

2-Amino-3-cyano-7,7-dimethyl-4-(2-nitrophenyl)-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran

ran (4k). IR (KBr): $\nu_{\max}$ = 3440, 3310, 3006, 2200, 1698, 1604 cm^{-1} . ^1H NMR (DMSO- d_6): δ_{H} = 0.96 (s, 3H, CH₃), 1.06 (s, 3H, CH₃), 2.14 (m, 2H, CH-8), 2.47 (s, 2H, CH-6), 5.02 (s, 1H, CH-4), 6.96 (s, 2H, NH₂), 7.46-7.94 (m, 4H, ArH) ppm. Anal. calcd. for C₁₈H₁₇N₃O₄: C 63.71, H 5.05, N 12.38; found C 63.80, H 5.11, N 12.43 %.

2-Amino-3-cyano-7,7-dimethyl-4-(3-nitrophenyl)-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran (4l). IR (KBr): $\nu_{\max}$ = 3400, 3300, 3196, 2200, 1696, 1600 cm^{-1} . ^1H NMR (DMSO- d_6): δ_{H} = 1.00 (s, 3H, CH₃), 1.06 (s, 3H, CH₃), 2.26 (s, 2H, CH-8), 2.48-2.52 (m, 2H, CH-6), 4.54 (s, 1H, CH-4), 5.98 (s, 2H, NH₂), 7.42-7.96 (m, 4H, ArH) ppm. Anal. calcd. for C₁₈H₁₇N₃O₄: C 63.71, H 5.05, N 12.38; found C 63.80, H 4.98, N 12.43 %.

2-Amino-3-cyano-7,7-dimethyl-4-(4-nitrophenyl)-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran (4m). IR (KBr): $\nu_{\max}$ = 3400, 3300, 3190, 2200, 1690, 1600 cm^{-1} . ^1H NMR (DMSO- d_6): δ_{H} = 1.02 (s, 3H, CH₃), 1.06 (s, 3H, CH₃), 2.26 (s, 2H, CH-8), 2.48-2.52 (m, 2H, CH-6), 4.55 (s, 1H, CH-4), 6.04 (s, 2H, NH₂), 7.46-8.06 (m, 4H, ArH) ppm. Anal. calcd. for C₁₈H₁₇N₃O₄: C 63.71, H 5.05, N 12.38; found C 63.76, H 5.11, N 12.33 %.

2-Amino-3-cyano-7,7-dimethyl-4-(4-methoxyphenyl)-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran (4n). IR (KBr): $\nu_{\max}$ = 3380, 3188, 2962, 2200, 1680, 1656, 1598 cm^{-1} . ^1H NMR (DMSO- d_6): δ_{H} = 0.96 (s, 3H, CH₃), 1.02 (s, 3H, CH₃), 2.12 (d, 1H, J =16.0Hz, CH-8) 2.22 (d, 1H, J =16.0Hz, CH-8), 2.46-2.56 (m, 2H, CH-6), 3.72 (s, 3H, OCH₃), 4.46 (s, 1H, CH-4), 6.84 (d, 2H, J =8.0Hz, ArH), 6.90 (s, 2H, NH₂), 7.04 (d, 2H, J =8.0Hz, ArH) ppm. Anal. calcd. for C₁₉H₂₀N₂O₃: C 70.35, H 6.21, N 8.64; found C 70.39, H 6.27, N 8.68 %.

2-Amino-3-cyano-7,7-dimethyl-4-(4-methylphenyl)-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran (4o). IR (KBr): $\nu_{\max}$ = 3420, 3330, 2950, 2190, 1678, 1640, 1600 cm^{-1} . ^1H NMR (DMSO- d_6): δ_{H} = 0.97 (s, 3H, CH₃), 1.04 (s, 3H, CH₃), 2.10 (d, 1H, J =16.0Hz, CH-8) 2.22 (d, 1H, J =16.0Hz, CH-8), 2.26 (s, 3H, CH₃), 2.46-2.56 (m, 2H, CH-6), 4.32 (s, 1H, CH-4), 6.94 (s, 2H, NH₂), 7.02 (d, 2H, J =8.0Hz, ArH), 7.08 (d, 2H, J =8.0Hz, ArH) ppm. Anal. calcd. for C₁₉H₂₀N₂O₂: C 74.00, H 6.54, N 9.08; found C 74.10, H 6.57, N 9.13 %.

2-Amino-3-cyano-7,7-dimethyl-4-(3,4-methylenedioxyphenyl)-5-oxo-4H-5,6,7,8-tetrahydrobenzopyran (4p). IR (KBr): $\nu_{\max}$ = 3370, 3206, 2938, 2204, 1650 1607 cm^{-1} . ^1H NMR (DMSO- d_6): δ_{H} = 0.96 (s, 3H, CH₃), 1.02 (s, 3H, CH₃), 2.12 (d, 1H, J =16.0Hz, CH-8) 2.24 (d, 1H, J =16.0Hz, CH-8), 2.46-2.56 (m, 2H, CH-6), 4.27 (s, 1H, CH-4), 5.96 (s, 2H, OCH₂O), 6.92 (s, 2H, NH₂), 7.02-7.16 (m, 3H, ArH) ppm. Anal. calcd. for C₂₀H₁₈N₂O₄: C 67.45, H 5.36, N 8.28; found C 67.52, H 5.43, N 8.34 %.

Acknowledgements

This project was supported by the National Natural Science Foundation of China, Educational Ministry of China, Educational Department of Hebei Province, Science and Technology Commission of Hebei Province.

References

1. (a) Bonsignore, L.; Loy, G.; Secci, D.; Calignano, A. *Eur. J. Med. Chem.* **1993**, 28, 517. (b) Bloxham, J.; Dell, C. P.; Smith, C. W. *Heterocycles* **1994**, 38, 399. (c) Nawwar, G. A. M.; Abdelrazek, F. M.; Swellam, R. H. *Arch. Pharm.* **1991**, 342, 875. (d) Zamocka, J.; Misikova, E.; Durinda, J. *Pharmazie* **1991**, 46, 610.
2. Armetso, D.; Horspool, W. M.; Martin, N.; Ramos, A.; Seaone, C. *J. Org. Chem.* **1989**, 54, 3069.
3. Hatakeyama, S.; Ochi, N.; Numata, H.; Takano, S. *J. Chem. Soc., Chem. Commun.* **1988**, 1202.
4. O'Callaghan, C. N.; Marry, T. B. H. *J. Chem. Res. Synop.* **1995**, 214.
5. (a) Kandeel, Z. E.; Farag, A. M.; Shaaban, M. R.; Elnagdi, M. H. *Heteroatom Chem.* **1996**, 7, 35. (b) Al-Ashmawi, M. I.; El-Sadek, M.; Sakr, S. M.; El-Sawah, M. *Egypt. J. Pharm. Sci.* **1992**, 33, 849. (c) Martin-Leon, N.; Quinteiro, M.; Seoane, C.; Soto, J. L. *Liebigs Annalen der Chemie* **1990**, 101. (d) Sharanina, L. G.; Nesterov, V. N.; Klokot, G. V.; Rodinovskaya, L. A.; Shklover, V. E.; Sharanin, Y. A.; Struchkov, Y. T.; Promonenkov, V. K. *Zhurnal Organicheskoi Khimii* **1986**, 22, 1315.
6. (a) Kuthan, J.; Sebek, P.; Bohm, S. *Adv. Heterocyclic Chem.* **1995**, 62, 19. (b) Elnagdi, M. H.; Aal, F. A. M. A.; Yassin, Y. M. *J. Prakt. Chem.* **1989**, 331, 971.
7. (a) Singh, K.; Singh, J.; Singh, H. *Tetrahedron* **1996**, 52, 14273. (b) Hassanien, A. A.; Zahran, M. A.; Gaby, M. S. A.; Ghorab, M. M. *J. Indian Chem. Soc.* **1999**, 76, 350.
8. Wang, X. S.; D. Shi, Q.; Tu, S. J.; Yao, C. S. *Synth. Commun.* **2003**, 33, 119.
9. Tu, S. J.; Jiang, H.; Zhuang, Q. Y.; Miao, C. B.; Shi, D. Q.; Wang, X. S.; Gao, Y. *Chin. J. Org. Chem.* **2003**, 23, 488.
10. (a) Li, C. J. *Chem. Rev.* **1993**, 93, 20. (b) Li, C. J. *Chem. Rev.* **2005**, 105, 3095. (c) Li, C. J.; Chen, L. *Chem. Soc. Rev.* **2006**, 35, 68. (d) Lubineau, A. *Chem. Ind.* **1996**, 123. (e) Meijer, A.; Otto, S.; Engberts, J. B. F. N. *J. Org. Chem.* **1998**, 63, 8989. (f) Ballin, R.; Bosica, G. *Tetrahedron Lett.* **1996**, 37, 8027.
11. Breslow, R.; Maitra, U.; Rideout, D. *Tetrahedron Lett.* **1983**, 24, 1901.
12. (a) Severance, D. L.; Jorgensen, W. L. *J. Am. Chem. Soc.* **1992**, 114, 10966. (b) Brandes, E. B.; Grieco, P. A.; Gajewski, J. J. *J. Org. Chem.* **1989**, 54, 515.
13. (a) Lubineau, A. *J. Org. Chem.* **1986**, 51, 2142. (b) Mori, Y.; Manabe, K.; Kobayashi, S. *Angew. Chem. Int. Ed.* **2001**, 40, 2815.
14. Li, C. J.; Chan, T. H. *Organic Reaction in Aqueous Media*, Wiley: New York, 1997.
15. Kool, E. T.; Breslow, R. *J. Am. Chem. Soc.* **1988**, 110, 1596.
16. (a) Armando, C.; Carlos, F. B. *Tetrahedron Lett.* **2003**, 44, 1923. (b) Yi, L.; Zou, J.; Lei, H.; Lin, X.; Zhang, M. *Org. Prep. Proc. Int.* **1991**, 23, 673. (c) Manabe, K.; Mori, Y.; Kobayashi, S. *Tetrahedron* **2001**, 57, 2537.
17. Silva, F. M.; Jones, J. J. *Braz. Chem. Soc.* **2001**, 12, 135.
18. (a) Wang, X. S.; Zhang, M. M.; Zeng, Z. S.; Shi, D. Q.; Tu, S. J.; Wei, X. Y.; Zong, Z. M.

- Arkivoc* **2006**, 7(ii), 117. (b) Jin, T. S.; Zhao, R. Q.; Li, T. S. *Arkivoc* **2006**, 7(xi), 176. (c) Stefani, H. A.; Pereira, C. M. P.; Dorr, F. A.; Cell, R. *Arkivoc* **2005**, 6(ii), 19. (d) Jin, T. S.; Zhang, J. S.; Xiao, J. C.; Wang, A. Q.; Li, T. S. *Synlett* **2004**, 866. (e) Jin, T. S.; Wang, A. Q.; Cheng, Z. L.; Li, T. S. *Synth. Commun.* **2005**, 35, 2339. (f) Jin, T. S.; Zhang, J. S.; Guo, T. T.; Wang, A. Q.; Li, T. S. *Synthesis* **2004**, 2001. (g) Jin, T. S.; Wang, A. Q.; Cheng, Z. L.; Zhang, J. S.; Li, T. S. *Synth. Commun.* **2005**, 35, 137. (h) Naik, S.; Bharttcharjya, G.; Kavala, V. R.; Patel, B. K. *Arkivoc* **2004**, 5(i), 55.
19. (a) Shi, D. Q.; Mou, J.; Zhuang, Q. Y.; Wang, X. S. *J. Chem. Res.* **2004**, 821. (b) Sharanina, L. G.; Nesterov, V. N.; Klokot, G. V.; Rodinovskaya, L. A.; Shklover, V. E.; Sharanin, Y. T.; Struchkov, Y. T.; Promoonenkov, V. E. *Zh.Org. Khim.* **1986**, 22, 1315. (c) Jin, T. S.; Wang, A. Q.; Wang, X.; Zhang, J. S.; Li, T. S. *Synlett* **2004**, 871.