5-SULFOSALICYLIC ACID (5-SSA): AN EFFICIENT ORGANOCATALYST FOR THE SYNTHESIS OF 4-METHYLCOUMARINS VIA Pechmann Condensation

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ABSTRACT

An efficient synthetic route for the Pechmann condensation reaction of substituted phenols and β-ketoesters under solvent-free conditions using a Bronsted acid 5-sulfosalicylic acid (5-SSA) as an organocatalyst leading to the formation of coumarin derivatives is described. Using microwave heating, reaction times were reduced from 45 min to 6 min. The attractive features of this protocol are mild reaction conditions, short reaction time, metal-free environment, easy workup procedure and improved synthesis which avoid chromatographic purification.

Keywords: Coumarin, Metal-free Environment, Microwave Irradiation, Organocatalyst, Green Chemistry etc.

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INTRODUCTION

In the plant kingdom, an important group of naturally occurring compounds, coumarin moieties are widely distributed. Coumarins (benzo-α-pyrones) are members of benzopyrone systems, which are found in many vegetables, seeds, fruits like bilberry, coffee, tea, nuts, cinnamon bark oil and wine.1,2 Synthetic coumarins are widely used as aroma chemicals because of their stability to alkali, odor strength and relatively cheap price.3 A coumarin as heterocyclic nucleus present in many natural products; therefore synthesis of coumarins has attracted considerable attention from the medicinal and organic chemist for many years. They occupy an important position, in synthetic organic chemistry, because of their potential applications in fragrance, pharmaceutical and agrochemical industries.

They are used as anticoagulants, cosmetics, additives in food, dispersed fluorescent laser dyes and optical brighteners and the preparation of insecticides.4-6 Many compounds containing the coumarin moiety possesses anti HIV and anticancer activity.7,8 In the synthesis of chromenes, furocoumarins, 2-acylresorcinol and coumarones, coumarin also acts as intermediates.9 Thus the synthesis of heterocyclic nucleus, i.e. coumarins, is of much current interest.10,11 Wedelolactone and Novobiocin (Fig.-1) are naturally occurring products that are used as a venomous snake-bite antidote and as an antibiotic respectively.12,13

![Wedelolactone](Image)

![Novobiocin](Image)

Fig.-1: Some Biologically Active Coumarin-based Compounds

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Pechmann, Reformatsky, Knovenegel, Perkin, and Wittig reactions are reported for the synthesis of coumarins.\textsuperscript{14-20} Pechmann reaction requires simple and inexpensive starting materials, i.e., phenols and β-ketoesters, and therefore the Pechmann reaction because of its simplicity, is a widely used method for synthesis of coumarin derivative. Conventionally, the Pechmann reaction is carried out in the presence of conc. H\textsubscript{2}SO\textsubscript{4}, phosphorus pentaoxide, aluminium chloride, trifluoroacetic acid, ionic liquids, and PVP-HWP, Zn[(L)-Proline]\textsubscript{2} as a catalyst.\textsuperscript{21-29} Long reaction time, tedious workup procedure, the requirement of special apparatus and use of organic solvents, all the above methods suffer from one or more disadvantages even these methods have their advantages. To reduce waste and costs, and to simplify the synthetic process and handling, the reactions are carried out under solvent-free condition.\textsuperscript{30} Development of organic synthesis in which the reactions are catalyzed by organocatalyst is an important area for the green procedure.\textsuperscript{31} However to the best of our knowledge, the organocatalyzed synthesis of coumarins has not been accomplished. Thus in the view of the importance of coumarin compounds, the development of an efficient, simple and versatile method for the preparation of coumarins by Pechmann condensation is an active research area and there is a scope for further improvement in the reaction condition and product yields. In continuation of our research work in the development of the green procedure, for the synthesis of biologically active moieties, we have developed an organocatalytic synthesis of coumarin derivatives.\textsuperscript{32-35}

Three-component Mannich reactions, as well as synthesis of 1-amido alkyl 2-naphthol and 3,4-disubstituted isooxazol-5(4H)-ones using 5-sulfosalicylic acid catalyst, was reported.\textsuperscript{36,37} This work was attractive for us to test the feasibility of 5-SSA as a novel basic organocatalyst in the synthesis of Coumarin derivatives. The metal-free organocatalytic process can reduce the cost and avoid the pollution caused by metals therefore, metal-free catalysts are regarded as green catalysts. Microwave provides a powerful way to do synthetic chemistry in the green approach. Material may be heated, in a microwave oven with the use of high-frequency electromagnetic waves. Due to the interaction of the electric field component of the wave with the charged particle in the material, heating arises. The heat is generated through resistance to an electric current, through the conduction mechanism. An oscillation of electrons or ions in a conductor is generated due to the oscillating electromagnetic field, which results in an electric current. The electric current generated faces internal resistance, which then heats the conductor. Microwave irradiation was preferred for heating the reaction mixtures because utilization of microwaves generally reduces reaction time and improves yields.\textsuperscript{38} In this study, we have developed the synthesis of coumarins by Pechmann condensation of substituted phenols and ethyl acetoacetate using 5-SSA as a mild and efficient organocatalyst under solvent-free conditions using conventional heating as well as microwave irradiation.

**EXPERIMENTAL**

**Materials and Methods**

5-sulfosalicylic acid (Spectrochem), substituted phenols (Spectrochem and Thomas Baker), and ethyl acetoacetate (Thomas Baker) were used as received. An open atmosphere and dried glassware are used to carry out all reactions. A hot paraffin bath is used for recording the melting points of all synthesized compounds and is uncorrected. Synthesis of coumarin derivatives was carried out under solvent-free conditions using the Microwave Synthesizer System (Milestone Srl, Sorisol (BG), Italy). The IR spectra were recorded on Alpha100508, FT-IR spectrometer in the frequency range 500-4000 cm\textsuperscript{-1}. The NMR spectra were recorded on a Bruker Avance (400 MHz for \textsuperscript{1}H NMR and 100 MHz for \textsuperscript{13}C NMR) spectrometer using DMSO as solvent using Tetramethylsilane (TMS) as an internal standard. Chemical shifts (\(\delta\)) are expressed in parts per million (ppm).

**Thermal Heating**

A mixture of resorcinol (1 mmol), ethyl acetoacetate (1 mmol) and 20 mol% 5-SSA was mixed and magnetically stirred at 80\(^\circ\)C for time intervals (20-45 min.). The completion of the reaction was confirmed by TLC (n-hexane: ethyl acetate, 3:1) and then iced cold water was poured into the reaction mixture. A pure solid was obtained by scratching the beaker, which was filtered, dried and recrystallized from ethanol to obtain coumarins in good yield and high purity. The 5-SSA is water-soluble which is recovered by evaporating water from the filtrate under vacuum, dried, and then used for the next cycle.
Microwave Irradiation
A mixture of resorcinol (1 mmol), ethyl acetoacetate (1 mmol) and 20 mol% 5-SSA catalyst was irradiated in a microwave synthesizer system at 320 W for 6 min. Work-up was done as reported in the thermal method.

Spectral Data of Representative Compound
7-hydroxy-4-methylcoumarin (Table-4, 3a)
FT-IR (KBr) (cm⁻¹): υ = 3299, 3050, 2850, 1695.; ¹H NMR (400 MHz,CDCl₃): δ 10.25 (br.s, 1H, OH), 7.43-7.41 (J = 8.8 Hz), δ 6.64-6.65 (J = 8.8 Hz) (d, 3H), 6.71-6.74 (m, 3H), 5.97 (s, 1H), 2.32 (s, 3H); ¹³C NMR (100 MHz, CDCl₃) : δ 18.6, 102.7, 110.6, 112.4, 113.2, 126.0, 153.2, 155.2, 161.4; MS (EI):m/z 176 [M⁺].

7,8-Benzo-4-methylcoumarin. (Table-4,3b)
FT-IR (υ in cm⁻¹): 1590, 1566,1474, 1630.; ¹H NMR (400 MHz, CDCl₃): δ 8.5 (d, 1H, ), 7.8 (d, 1H, ), 7.5-7.6(m, 2H, ), 7.4 (d, 1H,), 6.9 (d, 1H,), 6.4 (s, 1H), 2.5 (s, 3H).

6,7-Benzo-4-methylcoumarin. (Table-4,3c)
IR (υ in cm⁻¹): 1601,1512,1464 , 1630.; ¹H NMR(400 MHz, CDCl₃) : δ 7.7 (s, 2H), 7.6 (d, 1H), 7.2-7.4(m, 2H), 7.1 (d, 1H), 5.2 (s, 1H,), 1.7 (s, 3H).

8-Nitro-4-methylcoumarin. (Table-4,3d)
IR (υ in cm⁻¹): 1510, 1450(aromatic ring), 1600 (C=O), 1591,1335 (NO2), 1295, 1195(C-O).; ¹H NMR(400 MHz, CDCl₃) : δ 7.8 (d, 1H, ) , 7.1 (d of d, 1H), 6.9 (d, 1H), 3.3 (s, 1H ), 2.5(s,3H).

RESULTS AND DISCUSSION
An efficient protocol for the synthesis of coumarin using 5-SSA as catalyst under solvent-free conditions is reported. The structure of 5-Sulfoisalicylic acid is given below:

Optimization of Reaction Conditions
For the optimization of reaction condition, we investigated our study by choosing resorcinol and ethylacetoacetate (EAA) as a model substrate by using various mmols of catalyst and MW power levels for 4 to 6 min. When there is no catalyst, we got the low yield of the desired product which indicates the vital role of the catalyst.
20 mol% of 5-SSA catalyzes the reaction smoothly and optimized trials were summarized in Table-1. The reaction continuity was checked by taking TLC (Ethyl acetate/ n-hexane, 2:1 as a solvent system) at various time intervals. We observed that higher catalyst amount,25 mol% and 320 W, 40 %, MWI power yielded desired product 7-hydroxy-4-methylcoumarin in a short time 4 min with 90 % yield but more suitable reaction condition to obtain the desired product in acceptable yield, 90 % was using 20 mol% of 5-SSA as a catalyst and 320 W, 40 % MWI for 6 min (Table-1).

The efficiency of 5-SSA was also checked by the conventional heating method and same model reaction conditions with variable temperature and time and the results were given in Table-2. It was observed that in conventional heating satisfactory results for the synthesis of 7-hydroxy-4-methyl coumarin were
obtained at 80°C for 20 min with 85% yield (Table-2, entry 5). The yield obtained for model reaction in MWI (yield 90 %) is higher than conventional heating and reaction time reduced from 20 min to 6 min.

Table-1: Optimization of Reaction Conditions Using the Amount of 5- SSA at Different MWI

| mol% of 5-SSA and Microwave Power | Time (s) | TLC | mol% of 5-SSA and Microwave Power | Time (s) | TLC |
|----------------------------------|----------|-----|----------------------------------|----------|-----|
| 0 240 W (30%)                    | 120      | NP  | 0 320 W (40%)                   | 120      | NP  |
|                                  | 240      | NP  |                                  | 240      | NP  |
|                                  | 360      | NP  |                                  | 360      | NP  |
|                                  | 480      | NP  |                                  | 480      | TP  |
| 10 240 W (30%)                   | 120      | NP  | 10 320 W (40%)                  | 120      | TP  |
|                                  | 240      | NP  |                                  | 240      | TP  |
|                                  | 360      | TP  |                                  | 360      | CRM |
|                                  | 480      | TP  |                                  | 480      | CRM |
| 20 240 W (30%)                   | 120      | NP  | 20 320 W (40%)                  | 120      | CRM |
|                                  | 240      | TP  |                                  | 240      | P+CRM |
|                                  | 360      | TP  |                                  | 360      | P+MRI |
|                                  | 480      | CRM |                                  | 480      | P+MRI |
| 25 240 W (30%)                   | 120      | TP  | 25 320 W (40%)                  | 120      | CRM |
|                                  | 240      | TP  |                                  | 240      | P+MRI |
|                                  | 360      | CRM |                                  | 360      | P+MRI |
|                                  | 480      | CRM |                                  | 480      | P+MRI |

TP: Trace Product NP: No Product, CRM: Complex Reaction Mixture, P: Product, MRI: Minor Reaction Impurities

Table-2: Conventional Heating Methods to Optimize the Reaction Conditions

| Entry | Temperature ( 0°C ) | Time ( min ) | Yield ( % ) |
|-------|---------------------|--------------|-------------|
| 1     | RT                  | 90           | 48          |
| 2     | 45                  | 60           | 52          |
| 3     | 60                  | 45           | 65          |
| 4     | 70                  | 45           | 70          |
| 5     | 80                  | 20           | 85          |

*Reaction conditions: Resorcinol (1 mmol), Ethylacetooacetate (1 mmol), and 5-SSA (20 mol %) at a given temperature, *Isolated yield

From an enviro-economic point of view, the reusability of the catalyst was highly significant so the study was carried out using resorcinol (1 mmol), ethyl acetooacetate (1 mmol) and 20 mol%, 5-SSA catalyst as model substrates. Subsequently, 5-SSA is water-soluble which is recovered by evaporating water from the filtrate under vacuum, dried, and then used for the next cycle. The reusability of 5-SSA was studied efficiently by using the same model substrates under optimized reaction conditions. Recyclability of the 5-SSA catalyst was checked to four consecutive cycles having yield 90-80% (Fig.-2). The reusability of 5-SSA was examined efficiently using Resorcinol as a model substrate. The recovered 5-SSA was reused directly for three consecutive cycles, and all the results are summarized in Table-3. After the third cycle 85% of the catalyst is recovered, and it gives 80% of the coumarin product.

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Table-3: Catalyst Recovery for the Synthesis of Coumarin Derivatives

| Entry | Number of Cycles | Catalyst Recovery | Yield (%) |
|-------|------------------|-------------------|-----------|
| 1     | Cycle 1          | 98                | 90        |
| 2     | Cycle 2          | 94                | 88        |
| 3     | Cycle 3          | 90                | 84        |
| 4     | Cycle 4          | 85                | 80        |

*Reaction Conditions: resorcinol (1 mmol), ethylacetoacetate (1 mmol), and 5-SSA (20 mol%) at a given temperature, 
iisolated yield

To further establish the scope of optimized reaction conditions and to generalize the synthetic procedure, a variety of substituted phenols were treated with ethylacetoacetate under conventional and microwave irradiation method. The presence of electron-donating and electron withdrawing groups on the aromatic rings does not affect the yield of the product. All the results are compiled in Table-4.

Table-4: Comparison between Classical and Microwave Heating for the Synthesis of Coumarin Derivatives

| Entries | Aryl Aldehyde | Product M.P. °C (References) | Classical Heating | Microwave Heating |
|---------|---------------|------------------------------|-------------------|-------------------|
|         |               | (a-i)                        | Time/min | Yield (%) | Time/min | Yield (%) |
| 1       | Resorcinol    | 183-185 [38]                | 20       | 85        | 6        | 90        |
| 2       | α-Naphthol    | 152-154 [38]                | 30       | 78        | 7        | 82        |
| 3       | β-Naphthol    | 187-189 [39]                | 30       | 72        | 8        | 75        |
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The comparison of 5-SSA with the other reported catalyst employed for the reaction of resorcinol with EAA is shown in Table-5. This result reveals that 5-SSA is a highly efficient catalyst under microwave conditions in terms of reaction time and yield for the synthesis of Coumarin Derivatives.
The plausible mechanism for the 5-SSA catalyzed synthesis of 7-hydroxy-4-methylcoumarin under solvent-free conditions is shown in Scheme-2. In this mechanism, 5-SSA activates EAA by protonation, making the carbonyl group susceptible to nucleophilic attack by resorcinol. 7-hydroxy-4-methylcoumarin was produced and the catalyst was regenerated simultaneously through the transesterification and dehydration reaction.

![Scheme-2: The Plausible Mechanism for the Synthesis of 7-hydroxy-4-methylcoumarin](image)

**CONCLUSION**

In summary, we have carried out the optimization of Pechmann reaction of resorcinol with ethyl acetoacetate by using 5-SSA as catalysts under solvent-free conditions using microwave irradiation for the synthesis of 7-hydroxy-4-methylcoumarin. All products obtained by the condensation reaction of substituted phenols with ethyl acetoacetate (EEA) under solvent-free conditions using microwave irradiation, gave good to excellent yields. The present method has several advantages such as short time, metal-free synthesis, high reaction rates, excellent yields, no side reaction, cost efficiency, ease of handling of the catalyst, and a simple experimental procedure.

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