


 Cite this: *RSC Adv.*, 2024, **14**, 20365

Advances in organocatalyzed synthesis of organic compounds

 Ayesha Zafar,^a Muhammad Adnan Iqbal,^b  ^{ab} Ghazala Iram,^a Umar Sohail Shoukat,^a Faisal Jamil,^a Muhammad Saleem,^c Muhammad Yousif,^a Zain ul Abidin^a and Mohammad Asad^d

The recent advancements in utilizing organocatalysts for the synthesis of organic compounds have been described in this review by focusing on their simplicity, effectiveness, reproducibility, and high selectivity which lead to excellent product yields. The organocatalytic methods for various derivatives, such as indoles, pyrazolones, anthrone-functionalized benzylic amines, maleimide, polyester, phthalimides, dihydropyrimidin, heteroaryls, *N*-aryl benzimidazoles, stilbenoids, quinazolines, quinolines, and oxazolidinones have been specifically focused. The review provides more understanding by delving into potential reaction mechanisms. We anticipate that this collection of data and findings on successful synthesis of diverse compound derivatives will serve as valuable resources and stimulating current and future research efforts in organocatalysis and industrial chemistry.

Received 25th April 2024

Accepted 17th June 2024

DOI: 10.1039/d4ra03046j

rsc.li/rsc-advances

1 Introduction

Numerous techniques and effective catalysts have been developed to create synthetic organic compounds, which are in higher demand due to their low cost, simplicity of use, increased color stability, and resistance to light and pH changes. In order for modern civilization to successfully advance economically, it is essential that synthetic chemicals be produced in large quantities at low cost.¹ With the use of catalysts, it is theoretically possible to quickly and selectively synthesize desired chemical molecules without using additional energy or the catalyst itself.^{1,2} Catalysis is the current foundation for the majority of modern chemical processes. Growing environmental consciousness and increased pressure on the chemical sector to reduce waste generation both support the gradual replacement of conventional stoichiometric methods with cleaner catalytic substitutes. In contrast to the bulk chemicals and oil refining industries, which typically produce about 0.1 and 1–5 kg of byproducts per kilogram of product, respectively, the quantity of waste produced per kilogram of product, or E factor, is particularly high in the production of fine chemicals (up to 100 kg). This is partly

because fine chemistry uses multi-step synthesis, but it's also because stoichiometric rather than catalytic techniques are used more frequently.³

Recently, it has become more and more important to use tiny organic compounds as catalysts. Compared to the organometallic catalysts typically predominate in asymmetric synthesis, it has several advantages, such as decreased toxicity, greater stability, reduced activation energy, and friendly to the environment. Because of their low contamination and lack of metal traces in products, they facilitate the best reaction conditions, prevent metallic waste, and are perfect for the pharmaceutical industry.^{4–7} This article aims to quickly display certain classic and contemporary publications explaining, how these catalysts play crucial role in synthesis of organic components. In literature, recent work reported on organocatalysis includes organocatalysis of pyrrole and amino acids,^{8,9} dehydroabietane-type bifunctional organocatalysis,¹⁰ degradation of polluting dyes,^{11–13} asymmetric photocatalysis,¹⁴ proline derivatives catalyzed reactions in aqueous media,^{15–17} ionic liquid as organocatalysts,^{18,19} guanidinium catalyzed hypiodite mediated reactions,²⁰ asymmetric organocatalysis in drug discovery,²¹ enantioselective organocatalysis in disguise,²² asymmetric Michael addition of unactivated ketones and fluorination reaction.^{23,24} In this study, we gave a general overview of the methods and reaction mechanism (Table 1) for obtaining organocatalyzed synthetic organic compounds which have vast industrial applications such as sensors,^{25,26} fertilizers, pesticides,^{27,28} synthetic polymers, cosmetics, and platform chemicals are all utilized to improve people's standards of life.^{29,30}

^aDepartment of Chemistry, University of Agriculture Faisalabad, Faisalabad-38000, Pakistan. E-mail: adnan.iqbal@uaf.edu.pk

^bOrganometallic and Coordination Chemistry Laboratory, Department of Chemistry, University of Agriculture Faisalabad, Faisalabad-38000, Pakistan

^cDepartment of Basic and Applied Chemistry, Faculty of Sciences and Technology, University of Central Punjab, Lahore, Pakistan

^dCenter of Excellence for Advanced Materials Research (CEAMR), King Abdulaziz University, P.O. Box 80203, Jeddah 21589, Saudi Arabia



Table 1 Summary of discussed organocatalyzed organic reactions

Catalyst	Compound	Solvent	Reaction mechanism	Reaction time	Yield (%)
4,4'-bipyridyl	1–2	C ₆ D ₆	Fischer indole	24 h	95–92
	4–5	C ₆ D ₆	Deoxygenative reduction	22 h	60–12
Et ₃ N	6	1,4-Dioxane	1,4 Addition	24 h	98
Et ₃ N hydroiodide	7	Without solvent	Bronsted acid	20 min	83
Cinchona alkaloids	8	Toluene	Amination reaction	48 h	96
	9	CH ₃ Cl	Asymmetric Michael addition	48 h	86
Squaric acid	10	Water	Condensation	30 min	96
	11	DCM	Ring-opening polymerization	10 h	98.7
Taurine	12	Water	Cyclo nucleophilic addition	2–3 h	75–90
	13	Water	—	2–3 h	89–90
	14–15	Water	Aldol condensation	30–40 min	98–99
	16	EtOH	Biginelli cyclo-condensation	3 h	99
L-Proline	17	EtOH	Knoevenagel condensation	12 min	94
	18	EIOH	—	10 min	95
	19	EtOH	—	19 min	90
	20	Toluene	Diels–Alder reaction	10 h	82
	21	CH ₃ Cl	Atroposelective construction	15 min	87
CPA	22	Toluene	Atroposelective construction	24 h	88
	23	DMSO	EDA	14 h	52
THC	24	DMSO	—	—	84
	25	DMSO	—	—	73
	26	CH ₃ CN	Radical sulfonylation	30 min	67
Salicylic acid	27	DMSO	Oxidative condensation	48 h	71
	28	DMSO	Oxidative olefination	24 h	92
	29	DMSO	Cyclo condensation	20 h	92
DBU	30	DMSO	—	20 h	Traces

2 Organocatalyzed synthesis and reaction mechanisms

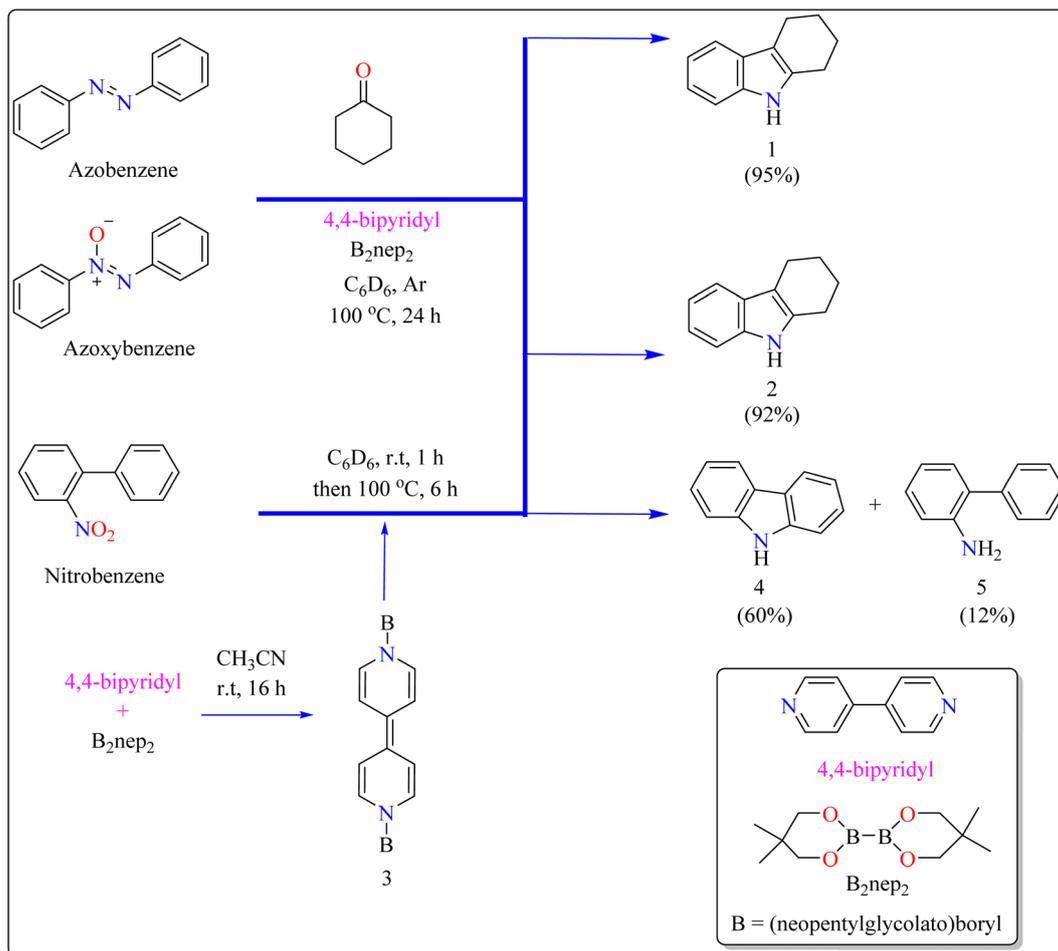
2.1. 4,4'-Bipyridyl catalyzed synthesis of indoles

Indole is a manufactured desirable structural component that is frequently used in a wide range of organic compounds, including drugs, agrochemicals, dyes, and³¹ the food sector has shown a great deal of interest in red colorant that is an indole-based dye.^{32,33} Indoles with different substituents were synthesized using a variety of simple processes that extensively studied over the past century.³⁴ Previously, 4,4'-bipyridyl was widely employed as a rigid linear bridging ligand in the design and fabrication of multidimensional coordination polymers. The literature has various homopolymetallic complexes containing 4,4'-bipyridyl, which is valued for its stiff, stable structure and two divergent binding sites, making it ideal for the formation of coordination networks. Despite the potential rotation of its pyridyl groups, the ligand's value is enhanced by its appropriate length and capacity to generate molecular cavities.^{35,36} However, in recent years, it has additionally been used as an organo-catalyst in the synthesis of indoles.³⁷ Bis(neopentylglycolato) diboron (B₂pin₂) served as a reductant to aid in the *trans*-1,2-diboration of carboxylates using 4,4'-bipyridyl.³⁸ Recently, it has been shown that, in the presence of catalytic amounts of 4,4'-bipyridyl, B₂nepe₂ also acts as a reductant for azobenzene, azoxybenzene, and nitrobenzene, resulting in increased yield and selectivity of indole derivative synthesis (Scheme 1).

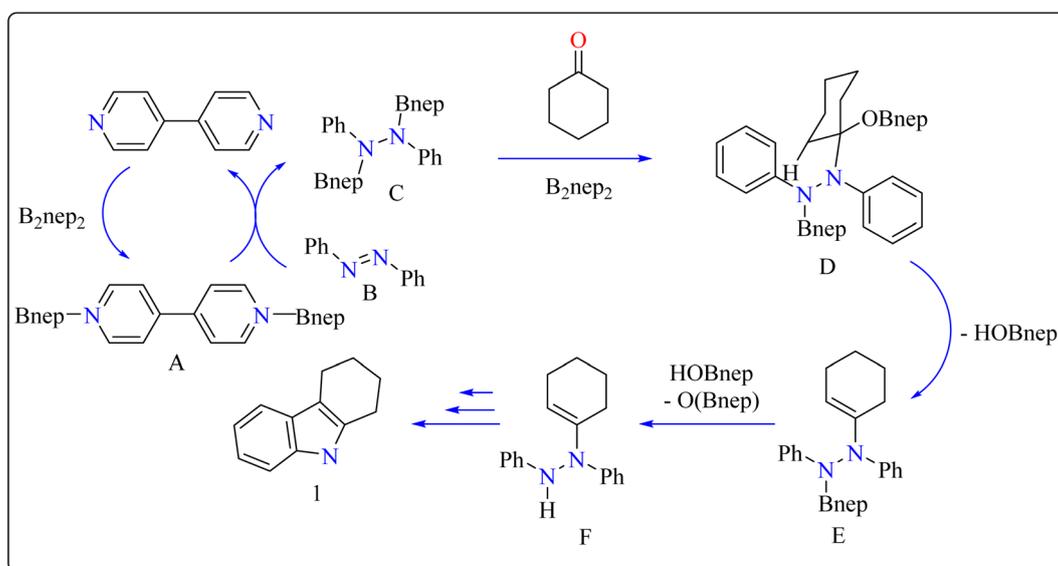
In the synthesis of indole, azobenzene interacts with cyclohexanone in the presence of B₂nepe₂ (2.2 equiv.) and a catalytic

quantity of 4,4'-bipyridyl. The reaction was performed in a variety of solvents, including C₆D₆, THF, and acetonitrile, in an argon environment at 100 °C for one day, with excellent results. During the first 10 minutes, the tube was gently flipped a few times to dissolve any particulates before being returned to the water at 100 °C. After the reaction was completed, the tube was allowed to cool to room temperature. The reaction mixture was then filtered through Celite and poured directly into a round-bottom flask. The tube was rinsed with ethyl acetate, and the solvent that was used is evaporated to obtain compound **1** with 95% yield.³⁹ The same technique was followed with azoxybenzene as a precursor and 3.3 equivalents of B₂nepe₂, giving compound **2** in 92% yield. For the synthesis of compound **4**, 4,4'-bipyridyl interacts with B₂nepe₂ in CH₃CN for 16 h at room temperature, yielding intermediate compound **3**. This intermediate (3.3 equiv.) subsequently interacts with nitrobenzene in C₆D₆, where the reaction mixture was heated to 100 °C for 6 h. This procedure yields two primary products: carbazole compound **4** (60% yield) and compound **5**, which yields 12%. Compound **4** was synthesized by inserting the aryl nitrene intermediate into the original compound's ortho C–H bond.⁴⁰ The suggested reaction process comprised the diborylation of azobenzene (**B**) by *in situ*-generated compound **A**, which yields diborylated compound **C** (Scheme 2). This interacted with cyclohexanone to produce enamine (**F**). The atom of nitrogen of compound **C** reacts nucleophilically with the carbonyl of cyclohexanone, for producing intermediate **D** and eliminating HOBnep to produce species **E**. This was quenched by HOBnep, which produced enamine **F** and O(Bnep)₂, followed by indole





Scheme 1 Synthesis of indole at optimized reaction conditions.



Scheme 2 Reaction mechanism for the formation of indole over 4,4'-bipyridial catalyst.



derivative **1** by aniline elimination, analogous to Fischer indole synthesis.³⁹

2.2. Triethylamine catalyzed reactions

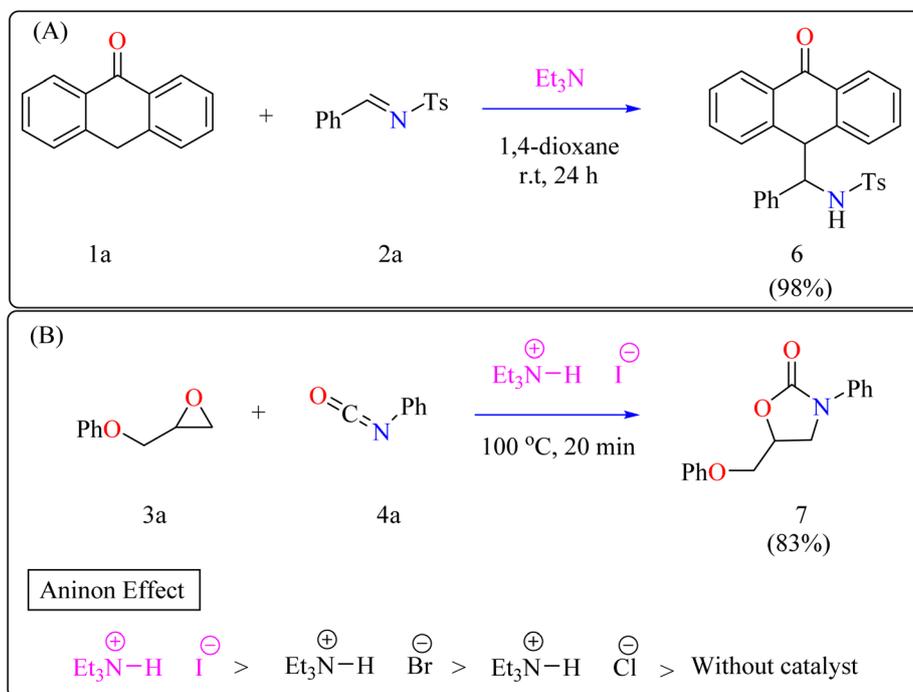
2.2.1. Synthesis of anthrone functionalized benzylic amines. Anthrone and its derivatives are common components in medications, dyes, and optoelectronic materials, as well as natural products and restorative drugs.^{41–44} They have high therapeutic potency, including antibacterial, diuretic, antipsoriatic, telomerase inhibitory, and selective anticancer activity.^{45–50} Functionalized anthrone derivatives can be synthesised by non-asymmetric and asymmetric 1,4-addition, as well as Diels–Alder reactions.^{51–54} While Kitamura *et al.* reported the coupling of 2-hydroxyindoline with anthrone using $\text{BF}_3 \cdot \text{Et}_2\text{O}$,⁵⁵ and White's group established a two-step technique employing $\text{Fe}(\text{PDP})$ -catalyzed oxidation followed by 1,2-addition of anthrone,⁵⁶ the 1,2-addition of imine remains undiscovered. Recently, researchers reported for the first time a Et_3N direct 1,2-addition of anthrone to *N*-sulfonylaldimines over moderate conditions, resulting in benzyl amine functionalized anthrone.

Anthrone **1a**, *N*-benzylidene-*p*-toluenesulfonamide **2a**, and catalytic amount of Et_3N (20 mol%) were mixed in an oven-dried round-bottom flask with anhydrous 1,4-dioxane, flushed with N_2 , and agitated for 24 h at room temperature (approx. 32 °C). The crude reaction mixture was then concentrated over reduced pressure and refined directly *via* silica gel column-chromatography, using petroleum ether/DCM/EtOAc as eluent, providing compound **6** with a 98% yield (Scheme 3A). Thus, Et_3N was shown to be the best catalyst, yielding the desired product in 98% of the reactions. However, an observational study showed that the organocatalyst plays a significant

part throughout the 1,2-addition process because the reaction only produced a little amount of product in its absence.⁵⁷

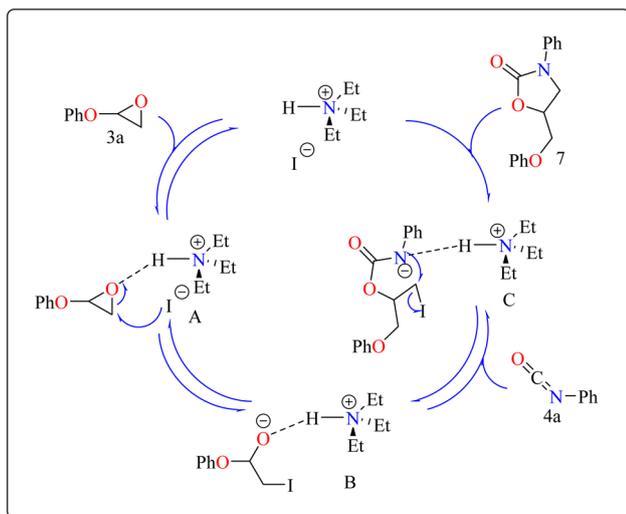
2.2.2. Synthesis of 2-oxazolidinones. Penta heterocycles, notably 2-oxazolidinones (**7**), are important in drug design, with several medications, like the anticoagulant rivaroxaban, built around this core.^{58–61} Previous literature demonstrated viable methods for synthesizing five-membered cyclic carbonates utilizing epoxides and carbon dioxide *via* the use of Et_3N hydroiodide as an excellent bifunctional catalyst.^{62–66} The catalytic activity of Et_3N hydroiodide, renowned for its acidic hydrogen on the atom of nitrogen, was investigated for coupling epoxides and isocyanates to produce 2-oxazolidinones. This simple and affordable catalyst acts as a bifunctional agent, effectively facilitating the necessary reactions.^{67,68}

Developing effective methods for the synthesis of compound **7** is critical.^{69,70} Recent research shown that Et_3N hydroiodide can efficiently catalyze the coupling of epoxides and aryl isocyanates and produced compound **7** without any byproduct. The reaction takes place under basic conditions: glycidyl phenyl ether **3a**, phenyl isocyanate **4a**, and Et_3N hydroiodide (10 mol%) are exposed to heat to 100 °C (Scheme 3B). After 20 minutes of stirring, the goal product **7** is achieved in 83%. The catalytic efficiency of ammonium salts varies with halide anions; iodide is the most efficient, generating **7** in 83%, while bromide and chloride yield 44% and 12%, respectively. Previous experiments involving cyclic carbonate synthesis have shown that no product formed without a catalyst. According to Scheme 4, the catalytic procedure using Et_3N hydroiodide includes epoxide **3a** activation *via* hydrogen bonding with the catalyst (intermediate **A**).⁶² The activated epoxide is nucleophilically attacked by an iodide anion, resulting in intermediate **B**. This intermediate's alkoxide



Scheme 3 Et_3N catalyzed synthesis of (A) Anthrone functionalized benzylic amines and (B) 2-oxazolidinones.





Scheme 4 Schematic route for the Et_3N catalyzed synthesis of compound 7.

anion subsequently attacks **4a**, resulting in intermediate **C**. The target product **7** is produced through intramolecular ring-closing of **C**, which also regenerates the catalyst. The iodide anion's efficacy stems from its low coordination with hydrogen that is acidic and high leaving group ability.⁷¹

2.3. Cinchona alkaloid catalyzed reactions

Cinchona alkaloids (quinine) have emerged as key asymmetry inducers in enantioselective catalysis over the last decade. These compounds are effective agents because of their structural characteristics, particularly a strong and easily functionalize chiral framework.⁷² Cinchona alkaloids are a truly green commodity because they are obtained from biomass and are renewable resources. They have been used as chiral Lewis bifunctional catalysts, chiral bases, and quaternized ammonium salts are utilized in phase-transfer catalysis for the past 20 years.^{73,74}

2.3.1. Synthesis of pyrazolones. The pyrazolones, classified as nitrogen-containing five-membered heterocyclic compounds, are frequently utilized as synthetic substrates in chemical and pharmaceutical chemistry as well as chelating, coloring, and pharmacological agents.^{75–79} A common pyrazolones interaction is the synthesis of heterocyclic dyes having long conjugated systems.⁸⁰ Fascinatingly, prior to 2008, there were no organocatalytic methods for the enantioselective production of functionalized pyrazolones.⁸¹ In 2009, Zhao published the initial instance of enantioselective reactions organocatalyzed by utilizing pyrazolone analogues. After this different organocatalysts (Thiourea, Gadolinium) used to catalyze enantioselective reactions.⁸² Here discussed quinine as organocatalyst for amination of pyrazolones. Quinine has been noticed as an appropriate catalyst and showed enantioselectivity up to 97%.

General procedure for the synthesis of functionalized pyrazolones proceeded with substituted pyrazol-5-one **5a** and quinine mixed in 2 mL of toluene and agitated for ten minutes

at ambient temperature (Scheme 5). The reaction mixture was cooled to $-40\text{ }^\circ\text{C}$. The next step involved adding azodicarboxylate **6a**, stirring was done in the reaction mixture. The reaction mixture was agitated at $-40\text{ }^\circ\text{C}$ until complete conversion. After that, a silica gel column directly loaded with the reaction mixture. Column chromatography (hexanes/EtOAc) produced the desired product **8**. When quinine is used to catalyze the reaction, pyrazolones might undergo the catalytic mechanism as shown in Scheme 5. In initial step pyrazolone deprotonated and proton transferred to catalyst, in second step added to azodicarboxylate, carbon–nitrogen (C–N) bond formation step which predicted to serve as the reaction's step that determined stereoselectivity, and at the last proton transferred from the catalyst to adduct. As seen in (Fig. 1), the optimized transition states of pyrazolone. The 'R' product was produced when pyrazolone (Re-face) attacked by azodicarboxylate through **TS-(R)**, while 'S' product was produced when pyrazolone (Si-face) added to azodicarboxylate *via* **TS-(S)**. There are three key interactions based on H-bonding with (catalyst)NH \cdots N(azodicarboxylate), (catalyst)NH \cdots O(pyrazolone), and (catalyst)OH \cdots O(azodicarboxylate) were found to be effective for charge stabilization on transition states. In addition, the frameworks made of pyrazolone and azodicarboxylate also take advantage of extra stabilization, bonded *via* hydrogen interactions in the **TS-(R)**.⁸³

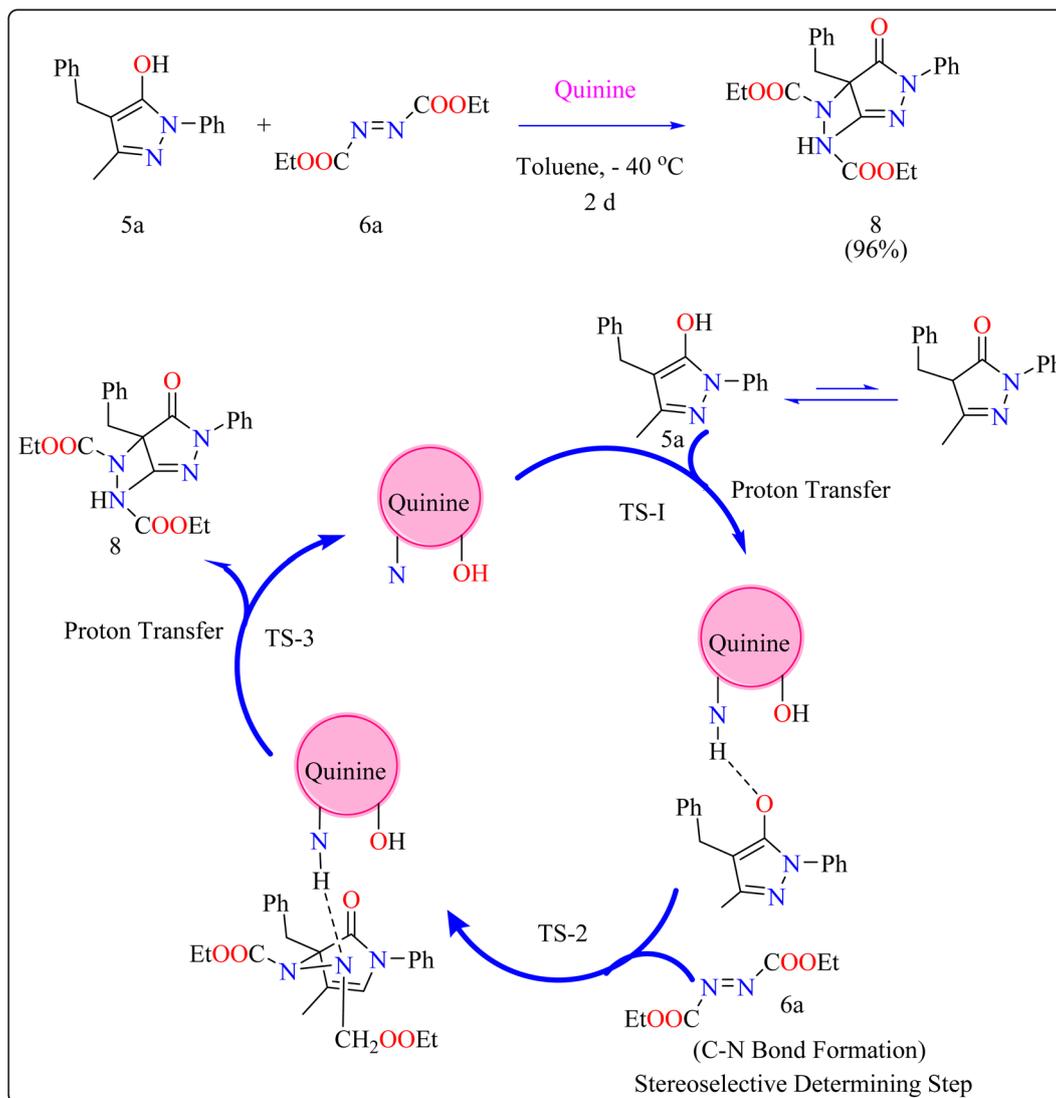
2.3.2. Synthesis of chiral maleimide. Maleimide derivatives have excellent physical and biological features, making them suitable frameworks.^{84–87} Although extensively investigated as Michael acceptors in asymmetrical Michael addition, which produces succinimides, their application as nucleophiles is uncommon.^{88,89} Some operations have used maleimides to produce maleimide-containing compounds by directly introducing the maleimide moiety. However, reactions using α -aminomaleimides as asymmetric Michael donors have not yet been documented. This study demonstrated the first asymmetric Michael addition of α -aminomaleimides (**8a**) as Michael donors to β -nitrostyrenes (**7a**) utilizing a bifunctional quinine organocatalyst. This expands the nucleophilic uses of maleimides.

Researchers used **7a** as a Michael acceptor to optimize the organocatalyst. Using natural quinine and cinchonidine, they were able to produce the Michael adduct **9** in moderate to good yield. However, the use of benzoyl-protected quinine decreased the yield. The incorporation of urea and thiourea molecules increased yield even further. Notably, **C** catalyst, generated from quinine and containing a 3,5-bis(trifluoromethyl)phenyl moiety, obtained 86% yield and 88% enantioselectivity, making it the best catalyst. As shown in Scheme 6, the reactions were conducted under the following circumstances: The substrate **7a** and **8a** were mixed in equimolar concentrations (0.1 mmol each), containing 20 mol% catalyst **C** in 1 mL of CHCl_3 . The combinations were swirled at ambient temperature for two days in an open-air setting.⁹⁰

2.4. Squaric acid catalyzed reactions

Squaric acid (SA), a dibasic organic acid which has a decomposition temperature of $245\text{ }^\circ\text{C}$, functions as a green





Scheme 5 Synthesis and proposed reaction mechanism for quinine catalyzed compound **8**.

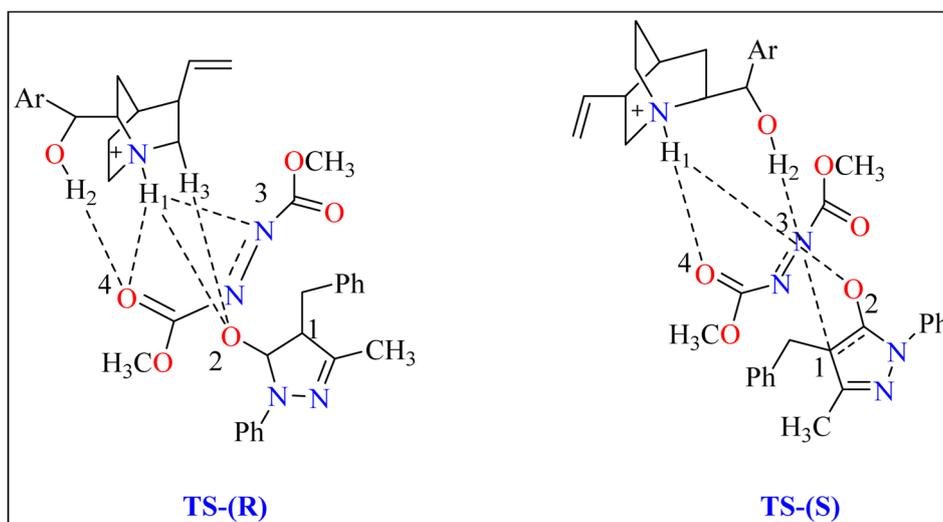
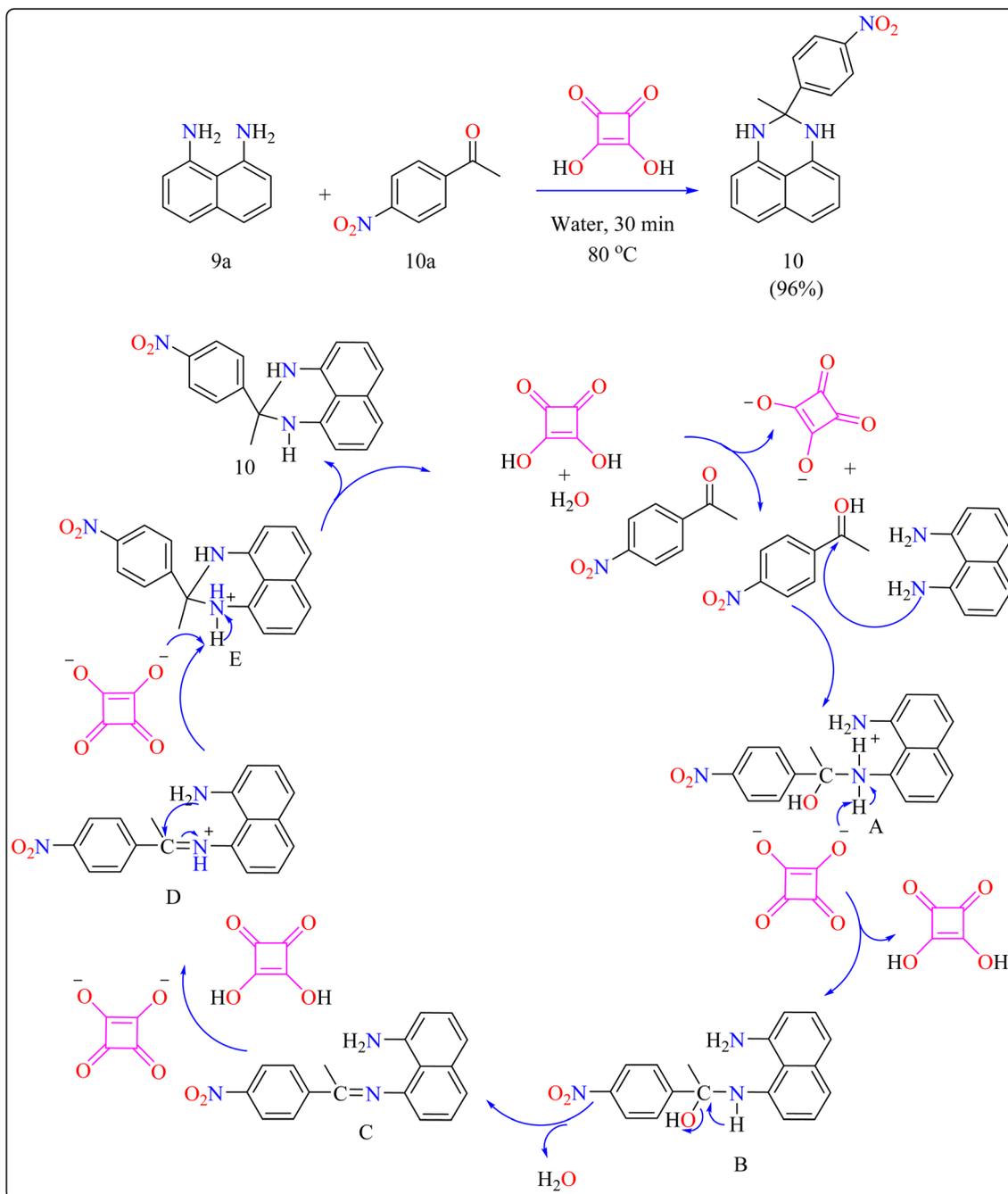


Fig. 1 Representation of two significant transition states of compound **8**, **TS-(R)** and **TS-(S)**.





Scheme 7 Synthetic route for the squaric acid catalyzed synthesis of perimidines.

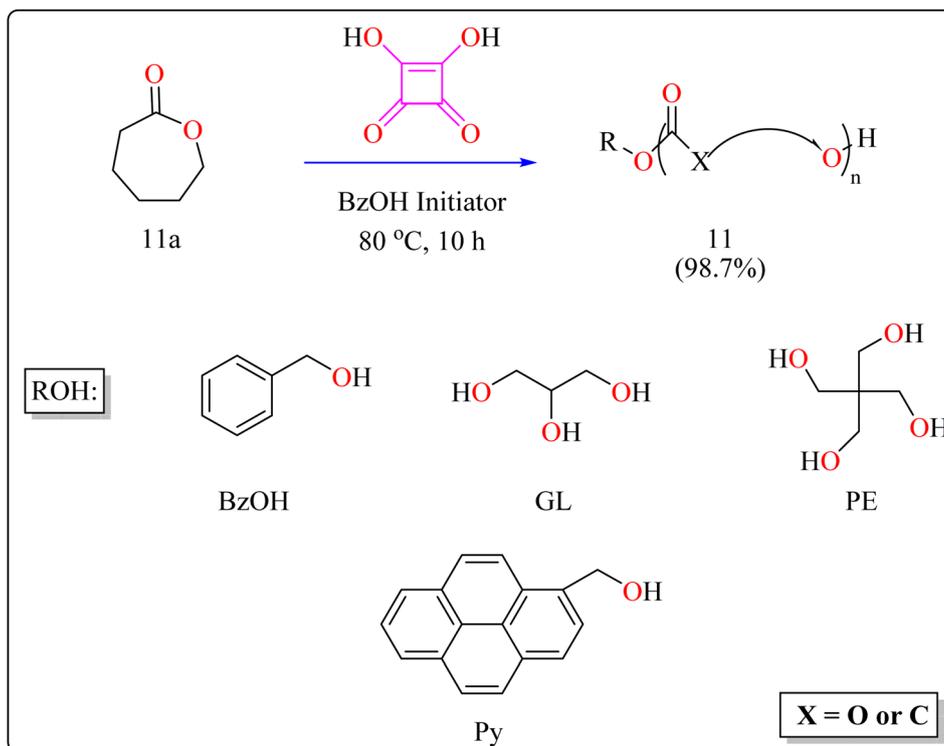
(Scheme 8). The polymerization then finally quenched by adding Amberlyst A21 for the specific time while the reactor was set at 80 °C. A specified amount of the polymerization mixture withdrawn. Dichloromethane (DCM) added to the reaction mixture after the allotted time to dissolve the polymerized PCL, and the SA catalyst has been removed by filtering. The filtrate was ultimately vacuum-evaporated in order to generate white solid of PCL **11**. Simply filtration, washing, and drying were used to isolate SA for each catalytic cycle. Interesting is that SA can be easily retrieved and utilized again for more than ten successive cycles without noticeably demolish its catalytic

activity.^{109,110} Scheme 9 showed the catalytic pathway for the synthesis of aliphatic polyester **11**.

2.5. Taurine catalyzed reactions

Taurine (2-aminoethanesulfonic acid), a commonly available amino acid that is also cheaply priced, has recently been employed to accelerate several chemical reactions.^{111,112} In nature, taurine is a common S-containing amino acid that can be obtained from a variety of living things.¹¹³ Due to the presence of the $-\text{SO}_3\text{H}$ group, taurine, which is widely accessible and moderately priced, has the ability to function as proton-acid



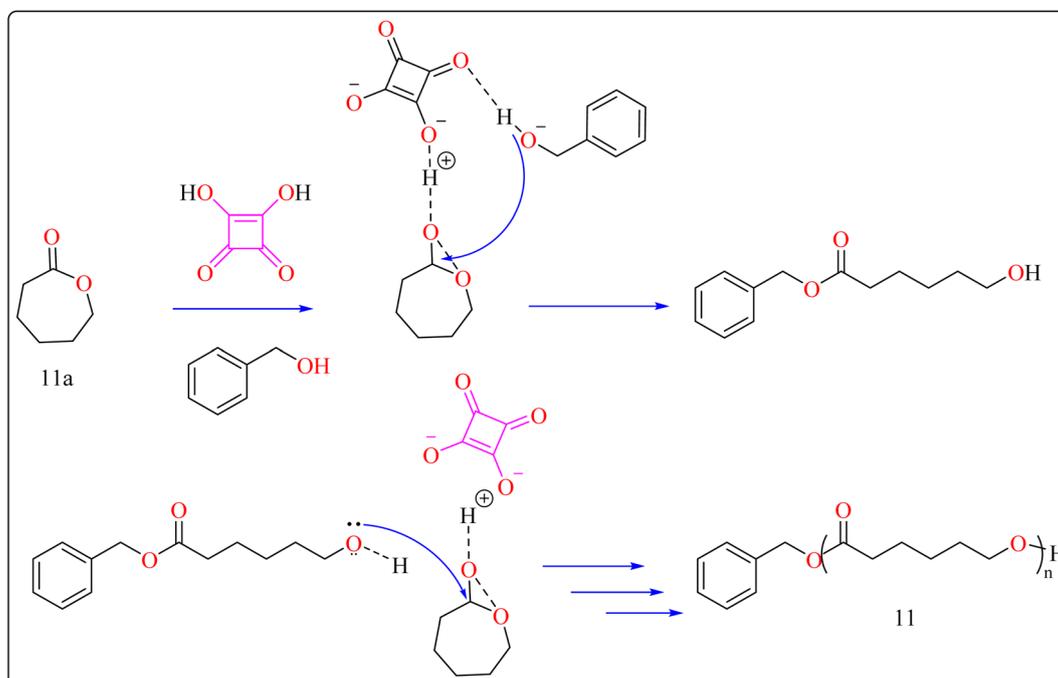


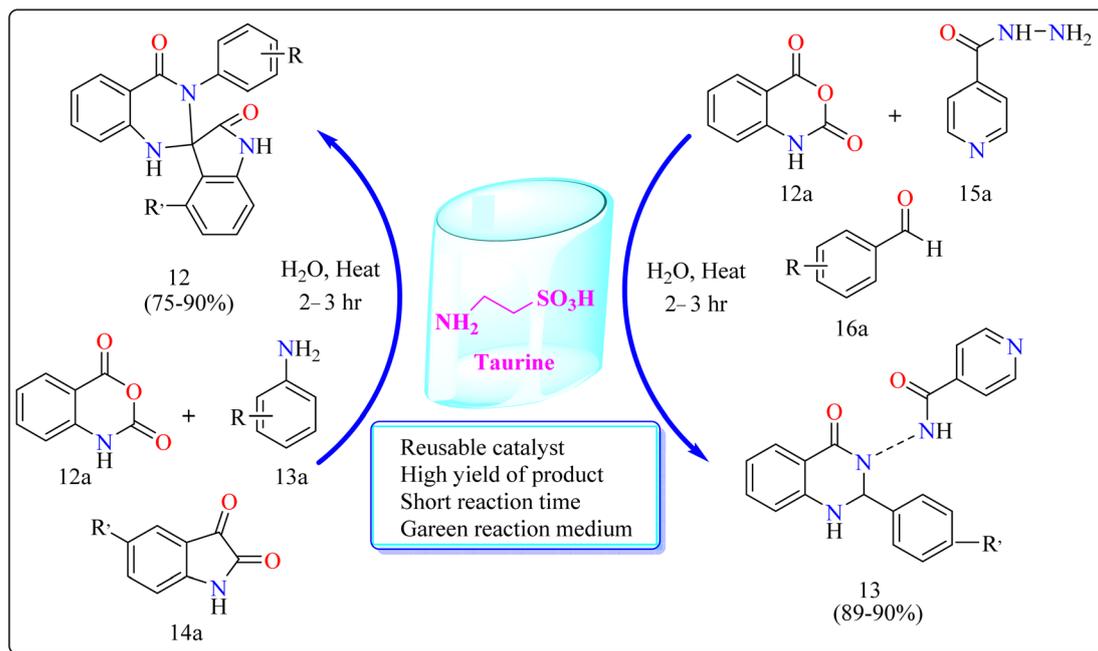
Scheme 8 Synthetic of polyester.

catalysts in the process of oxidation of sulphides towards sulphoxides utilizing aquatic peroxide of hydrogen to act as oxidant. Taurine has important biological roles in the body and is a key ingredient in energy drinks and supplements. Furthermore, it promotes the Knoevenagel reaction in aqueous solution and

helps for synthesizing derivatives of barbituric and thio-barbituric acid.^{114–116}

2.5.1. Synthesis of spirooxindole dihydroquinazolinones (12) and novel 1,2-dihydroquinazolinones-3(4H)isonicotinamides (13). The researchers evaluated the reusability of

Scheme 9 Mechanism for squaric acid catalyzed synthesis of poly(ϵ -caprolactones).



Scheme 10 Methodology for the synthesis of compound 12 and 13.

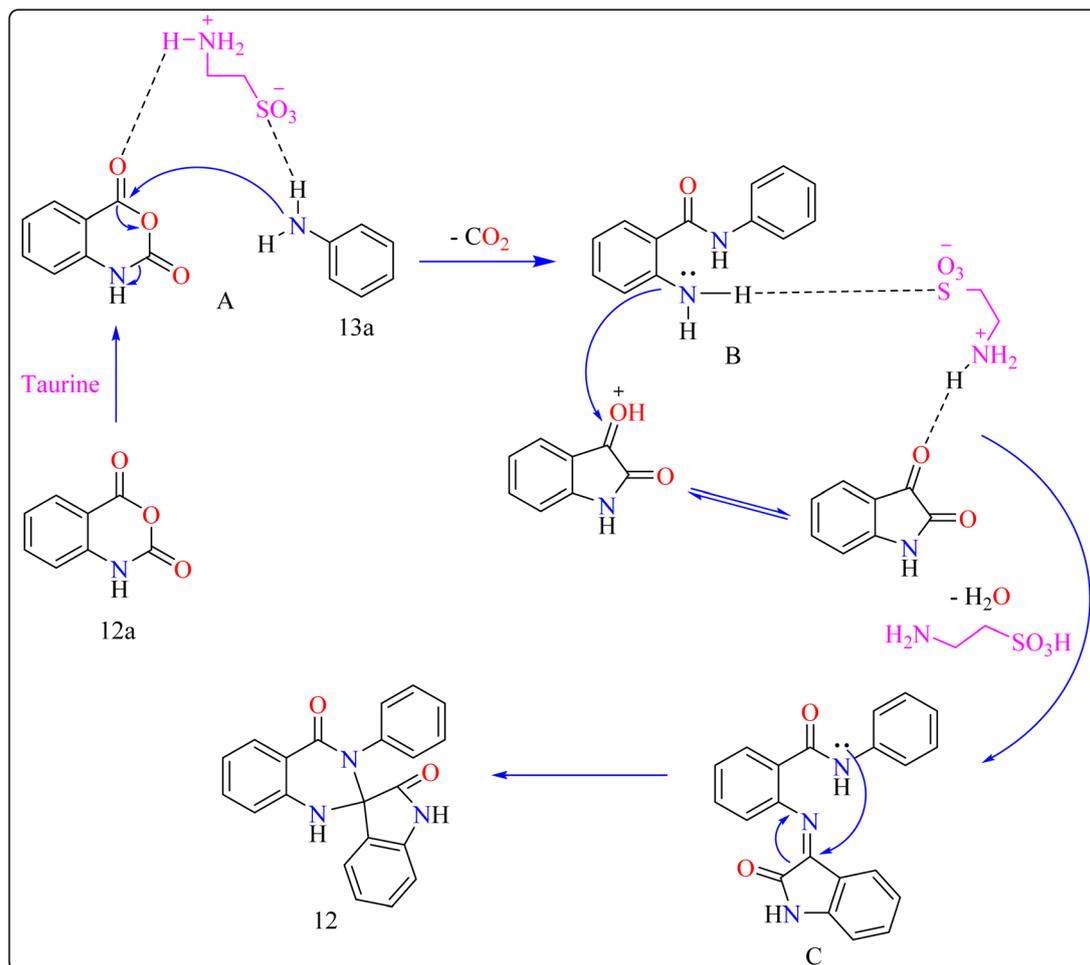
taurine for each of the reactions comprising compound 12 and 13. Without significantly affecting the product yield, it was simple to recover and reuse the 2-aminoethanesulfonic acid for at least three runs. Taurine catalyzed synthesis of 12 and 13 showed in Scheme 10. Taurine catalyst (15 mol%) was introduced to an equimolar mixture of isotonic anhydride (12a), aniline (13a), and isatins (14a) for the reaction, took 10 mL of H₂O inside a 100 mL round-bottom flask. For a period of six hours, the reaction mixture had been heated. After the reaction, that was monitored by TLC, finished, 10 mL of water was added to it and the mixture was stirred for 3 minutes. Then, it was allowed to cool. At this point, the material precipitated, and it was later separated by filtration. The isolated product had many water rinses. No further processing using column chromatography as well as the utilization of any kind of organic solvent was necessary after drying to produce the pure product. Here excellent yield of product was obtained in ranging from 75–90%. Similarly for the synthesis of compound 13. Taurine catalyst (15 mol%) was introduced to an equimolar mixture of 12a, isoniazid (15a) and aldehyde (16a) in 10 mL of water in a round bottom flask for the reaction. The reaction mixture was heated for 2.5 h. Next procedure for the synthesis of VIIC same as used for the synthesis of IVC. The product yield was obtained in the range of 89–90% (Scheme 10). Quinazolinone derivatives have multiple applications in the dyestuff enterprises as coloring ingredients in addition to their use in pharmaceuticals.^{117,118}

Reaction mechanism for the synthesis of compound 12 showed in Scheme 11. Here taurine functions as a donor–acceptor reagent with dual functions by activating 12a carbonyl position to produce intermediate A, which may allow 13a nucleophilic attack on the carbonyl unit. 13a was added nucleophilically to 12a, which was then decarboxylated to form

2-aminobenzamide (B). Imine intermediate C was then produced by condensation of B with protonated isatin, which was made using taurine, and then underwent intramolecular cyclization to produce product 12.¹¹⁹

2.5.2. Synthesis of 3,4-dihydropyrimidin. The use of taurine as a green bio-organic catalyst enhances the Biginelli reaction, a one-pot multicomponent synthesis of bio-active 3,4-dihydropyrimidin-2(1H)-ones/thiones (14–15). This method employs ethyl acetoacetate (17a), an aromatic aldehyde (18a), and urea or thiourea (19a) in an aqueous solution of EtOH (1 : 3) as shown in Scheme 12. The reaction produced high yields, is simple, and environment friendly, with short reaction times and minimal costs. Without a catalyst, the process produces little product. Several catalysts (acetic acid, L-proline, *p*-toluenesulfonic acid, FeCl₃·6H₂O) were tried, and taurine produced the greatest yields and quickest reaction durations, especially at 20 mol%. Raising the total amount of taurine failed to considerably improve yield. Solvent investigations showed that a water–ethanol mixture outperformed other solvents such as DCM, EtOH, DMF, MeOH, 1,4-dioxane, MeCN, toluene, and pure water, all of which produced lower yields. The technique advantages from fast completion and easy product purification using ethanol recrystallization, thereby providing an effective and environmentally benign approach for synthesizing compound 14 and 15.¹²⁰ In Scheme 12, two approaches are presented on the basis of previous studies¹¹⁵ for synthesizing 14–15 products with taurine as a catalyst. Pathway A involves taurine promoting aldol condensation of 17a and 18a, followed by nucleophilic addition of 19a and hydrogen shift to produce compounds 14–15. Pathway B begins with the nucleophilic addition of 19a to 18a, followed by taurine-catalyzed imine production, and then addition and cyclization with 17a.¹²⁰





Scheme 11 Reaction mechanism for the synthesis of taurine catalyzed compound 12.

Compound **16** was synthesized using important precursors such as substituted benzaldehyde (**20a**), thiourea (**21a**), and acetoacetanilides (**22a**). The reaction, which was carried out in ethanol at 100 °C, took 3 hours to complete (Scheme 13). The use of taurine as a catalyst considerably increased the efficiency of the reaction, generating compound **16** with a high productivity of 99%. This one-pot synthetic approach not only produces excellent yields but also streamlines the process, making it appropriate for large-scale industrial manufacturing.¹²¹ The usage of taurine corresponds with environmentally friendly techniques, increasing the method's overall sustainability. This synthesis's efficiency and speed, together with its high yield, make it a valuable technique for synthesizing 3,4-dihydropyrimidin derivatives, which have been shown to be bioactive. The fact that this technology can be applied to large-scale production highlights its potential for wider industrial adoption.¹²²

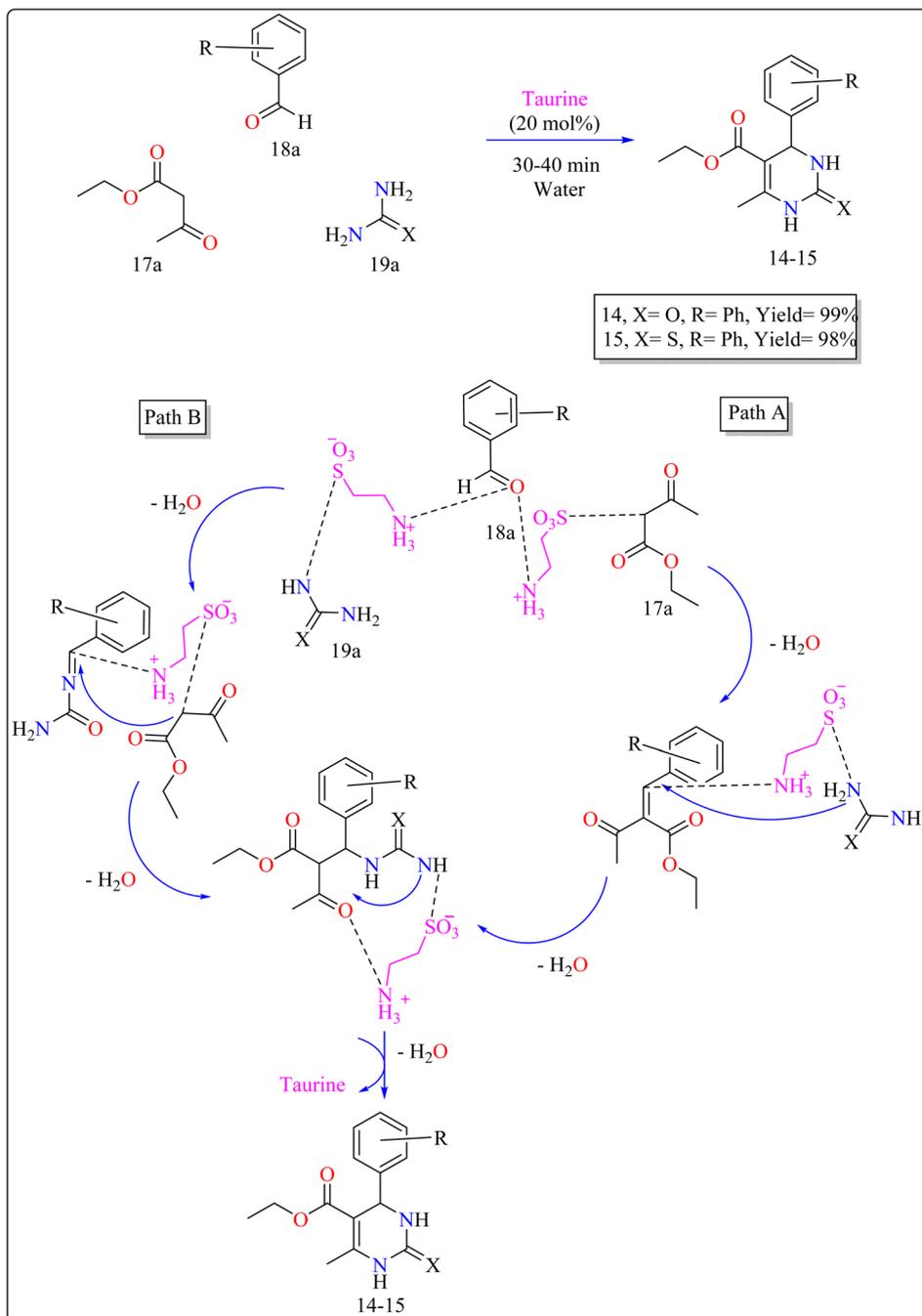
2.6. L-Proline catalyzed reactions

L-Proline, a naturally occurring amino acid having secondary amine functionality, operates as a bifunctional organocatalyst, with the amino group acting as a Lewis base and the group of

carboxylic acids serving as a Brønsted acid.^{123–125} It is a highly efficient, adaptable, and environmentally friendly catalyst for the synthesis of heterocyclic systems. L-Proline efficiently catalyzes a variety of asymmetric syntheses, such as Ullmann, Hantzsch, and Knoevenagel reactions, in addition to, aldol, Mannich, Robinson annulations, Michael, α -selenenylation, and Diels–Alder. Its dual role as a ligand and catalyst, combined with its excellent enantiomeric purity, increases its value in green chemistry.^{126–128}

2.6.1. Synthesis of benzil bis-hydrazones. Novel bioactive benzil bis-hydrazone derivatives (**17–19**) were synthesized by solvent-free mechanical grinding of benzil bis(cyanoacetylhydrazone) (**25a**) with aldehydes (**26a–c**) employing L-proline as an effective organocatalyst (Scheme 14). The precursor **25a** was synthesized by heating benzil (**23a**) and 2-cyanoacetylhydrazide (**24a**) in EtOH with HCl. The grinding method at room temperature with moist L-proline (5 mol%) produced high yields *via* Knoevenagel condensation. After assessing the completeness of the reactions after 10 minutes of grinding, researchers discovered that yields for compounds **17** and **19** declined from 94% to 83% and 90% to 89%, respectively, while yield for compound **18** remained constant. This shows that





Scheme 12 Synthesis and proposed mechanism for taurine catalyzed compound 14–15.

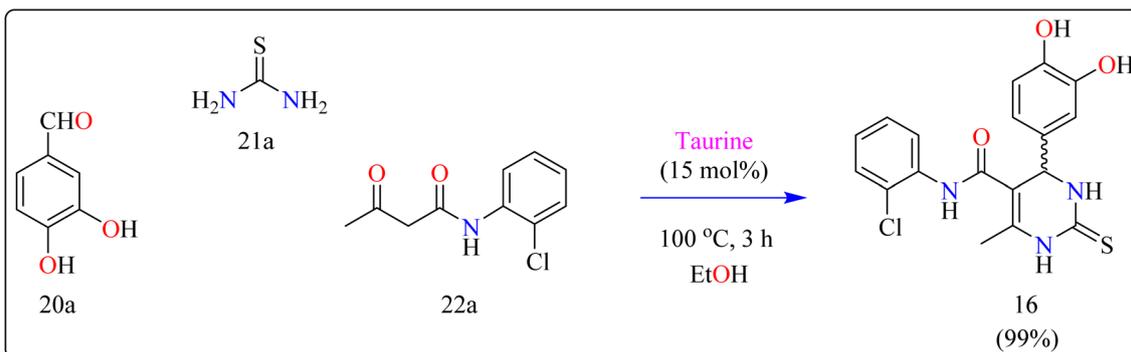
electron-donating and electron-withdrawing groups speed up the process, preventing major yield losses. In contrast, the yield of the unsubstituted derivative 17 dropped dramatically, showing a reduced reaction efficiency in the absence of substituents. Thus, substituent groups are critical for maintaining good yields during shorter reaction durations.¹²⁹

2.6.2. Synthesis of functionalized phthalimides. In the manufacture of beneficial substances including catalysts, dyes, polymers, along with fluorescence probes, derivatives of phthalimides are widely employed. The primary use of

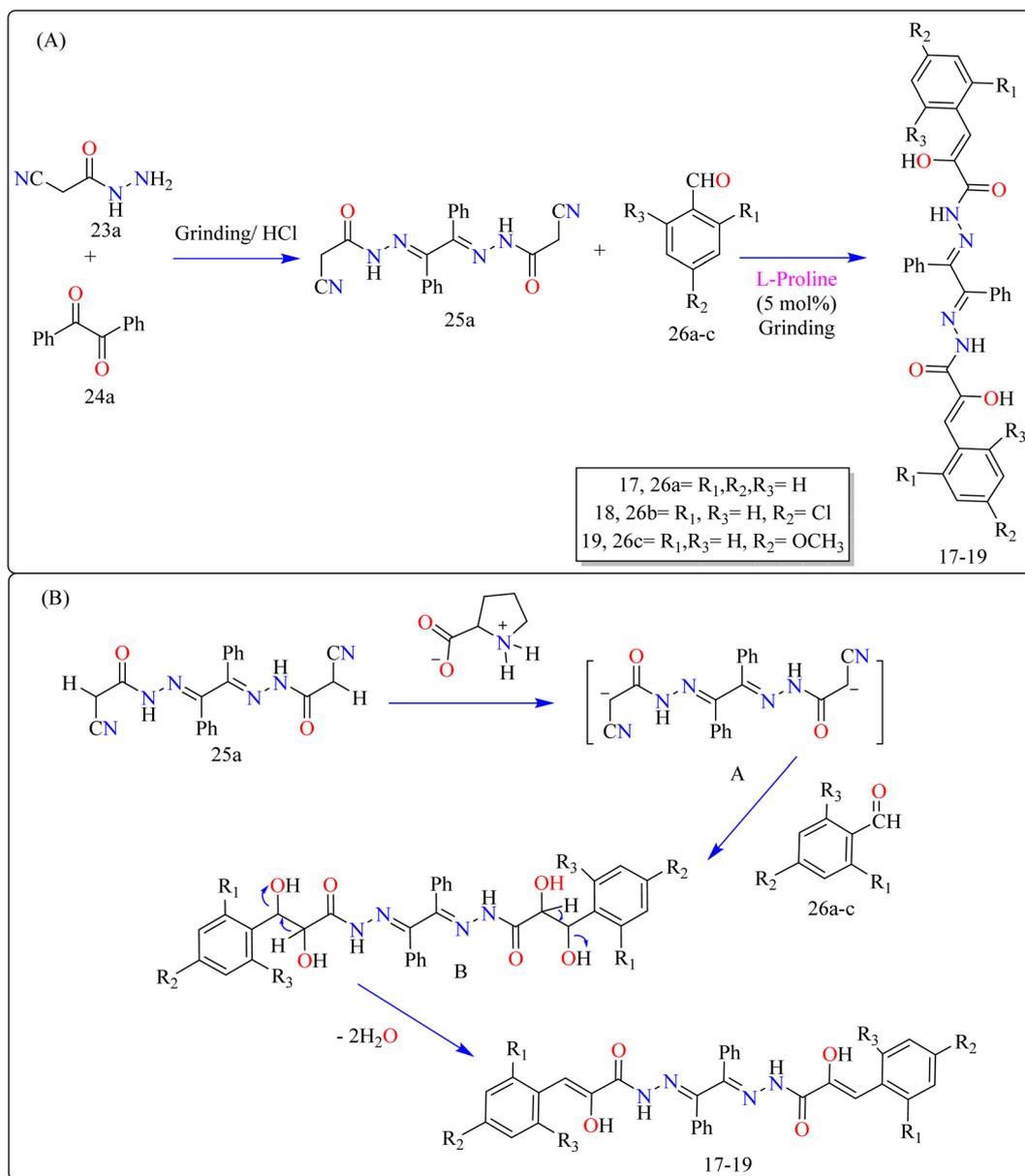
phthalimide structure in dispersed dyes is as a diazo or coupling component.^{130,131} More Effective coplanar aromatic rings are another feature of phthalimide disperse dyes. Increasing the plane shape of dye molecules is one of the best ways to improve their washing fastness because they result in higher intermolecular interaction energy, decreased temperature migration, and exceptional washing fastness.^{132,133}

Researcher presented the Diels–Alder reaction as a method for the *l*-proline-catalyzed benzannulation of different and multifunctionalized phthalimides using two dienophiles





Scheme 13 Synthesis of taurine catalyzed compound 16.



Scheme 14 (A) Methodology and (B) proposed route for the synthesis of L-proline catalyzed benzil bis-hydrazones.



instead of a diene and a dienophile (Scheme 15). This reaction allowed the transition-metal-free preparation of multiple phthalimides by an organocatalytic method involving malimides and α,β -unsaturated aldehydes. Here, with the addition of an *L*-proline catalyst, researcher used a number of additives (acetonitrile, acetic acid, pivalic acid, and benzoic acid) as well as a variety of solvents (THF, dioxane, and toluene). Fortunately, compound **20** was produced in a fast reaction time (10 h) with better yield of 82% using benzoic acid (10 mol%) as additive, with toluene serving as the solvent. The same way, *L*-proline and benzoic acid were added to a solution of 3-methyl-2-butenal (**27a**) and *N*-phenylmaleimide (**28a**) in toluene. The mixture was agitated for ten hours at 80 °C in an oil bath. When the reaction was determined to be finished by TLC, the product of the reaction had been evaporated using an evaporator with a rotary blade. Applying ethyl acetate/hexane (1 : 20) as an eluent, the residue was subsequently filtered with a silica gel column chromatography technique, producing the desired product **20** in 82% yielding as a yellow solid.¹³⁴

2.7. Axillary phosphoric acid catalyzed reactions

2.7.1. Synthesis of axially chiral 4-azaborinephenols.

Asymmetric catalysis and materials science rely heavily on axially chiral components found in chiral ligands and organocatalysts.¹³⁵ Research on heterobiaryl atropisomers particularly those with non-C–C or C–N axes is scarce compared to the vast study of biaryl atropisomers.^{136–140} To increase their use in the development of drugs and materials design, innovations in their preparation and characterization are necessary. A desymmetrization approach was taken,¹⁴¹ building on the understanding of atroposelective synthesis and effective electrophilic aromatic substitution.^{142–144} A 3,5-disubstituted phenol (**29a**) motif model substrate was developed. While methyl groups maintained the stereogenic axis for improved enantioinduction, hydroxyl groups served as directing groups and catalyst binding sites.¹⁴⁵

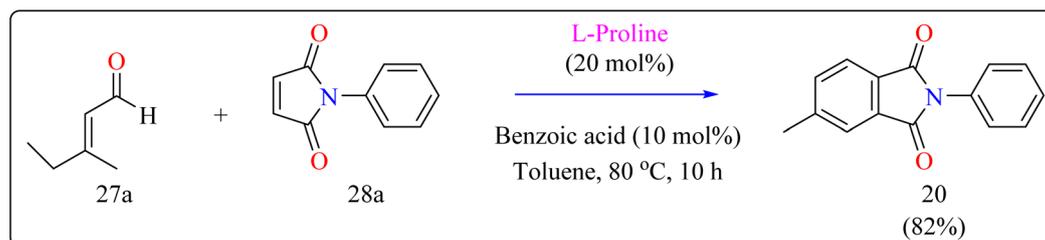
The difficulties were in finding a chiral organocatalyst that was appropriate for distant axial enantiocontrol, choosing an electrophilic reagent that was effective with phenol, and creating gentle reaction conditions to maintain stability. It was suggested to use CPAs for bifunctional activation through H-bonding interactions. Limited reactivity was seen in the initial substrate reactions with certain electrophiles. A close-by binding site is crucial for stereoselectivity, as demonstrated by the modest enantioselectivities of bromination and a particular reagent (E). Due to its advantageous qualities,

diazodicarboxamide **30a** which is reactive and compatible with CPA in terms of H bonds was selected. As shown in Scheme 16, in a 15 minutes reaction with **29a**, **30a**, and CPA (0.01 mmol) in 2 mL of CHCl₃ at room temperature, 87% of **21** was produced. Even after being heated to 120 °C for 24 hours, the product showed good configurational stability and no discernible racemization. This method effectively transfers stereochemical data obtained from the catalyst to the B–C linkage of azaborine derivatives by remotely setting the stereogenic axis. Asymmetric catalysis and materials science applications are being investigated.¹⁴⁶

2.7.2. Synthesis of axillary chiral *N*-aryl benzimidazoles.

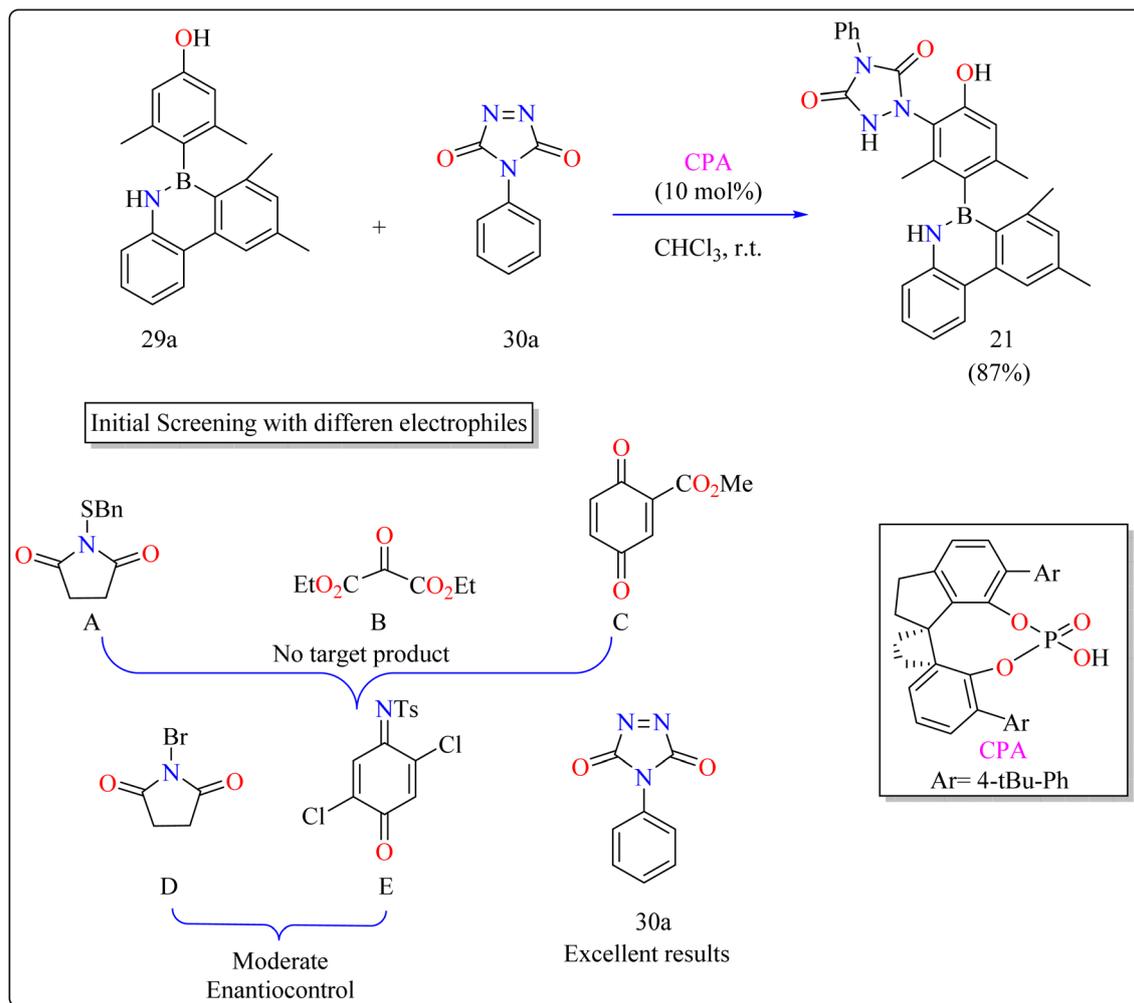
There are many biological active compounds that include *N*-aryl benzimidazoles for instance, they are used as inhibitors of certain enzymes like nonpeptide thrombin.^{147,148} They are also important components of high-temperature polymers, veterinary medications, fungicides, herbicides, and dyes. Previously reported, dyes containing benzimidazole as structural component are utilized to protein fibers like wool and silk fabric, having properties including color features, color appearance in daytime and UV light, antibacterial activity, insect repellency, and to enhance wet adhesion properties.^{149–151} Axially chiral compounds play crucial structural roles in biologically active chemicals and natural products. Additionally, chiral ligand and catalytic components are preferred. Significant progress has recently been made in the asymmetric method for synthesizing axially chiral skeletons. To the greatest extent of our understanding, the atroposelective production of *N*-aryl benzimidazoles has been a very challenging process up to this point. A crucial organocatalyst in the asymmetric synthesis is the axially chiral phosphoric acids (CPAs).^{152–154} Researcher developed an organocatalytic, atroposelective process for making axially chiral *N*-aryl benzimidazoles that involved the breakage of carbon (C–C) bonds in the presence of CPAs. The CPAs showed a number of distinct advantages, including environmental friendliness, ease in cleaving the carbon–carbon bonds of multicarbonyl substances, excellent efficiency along with enantioselectivity, a wide range of substrate tolerance, as well as gram-scale chemical reactions without departing reactivity and enantioselectivity (ee).

For the synthesis of compound **22**, combined *N*¹-(naphthalen-1-yl)benzene-1,2-diamine (**31a**) along with acetylacetone (**32a**) with addition of 10 mol% of CPA ((*R*)-C1), magnesium sulphate (MgSO₄) in toluene, two additives sodium sulphate (Na₂SO₄) and 3 Å molecular sieve (MS), have been used for higher enantioselectivity at 60 °C for 20 h and



Scheme 15 Optimized reaction conditions for *L*-proline catalyzed compound **20**.





Scheme 16 Methodology for the synthesis of CPA catalyzed compound 21.

three angstrom MS produced 88% yield and 92% enantioselectivity (Scheme 17). It was showed that **31a** reacted with three-carbonyl molecules: **32a**, ethyl acetoacetate (**32b**), and 1-phenylbutane-1,3-dione (**32c**). **32a** and **32c** exhibited a greater degree of reactivity (78–88% yields) as well as excellent ee (91 or 92%). According to Scheme 18, intermediate **A** was generated when two substrates, **31a** and **32a**, were treated with chiral phosphoric acid (R)-C₁. Then, **B** is produced by the nucleophilic attack of the amino in **31a** on the carbonyl in **32a**, regenerating (R)-C₁, and **B** was dehydrated to produce intermediate **C**. An intramolecular nucleophilic strike of the imino group on the atom of carbon found in the imine in **D** causes the catalyst to be released after the catalyst and **C** have reacted to form intermediate **D**. The target product, **22**, was produced when **F** is separated from **E** by cleaving the C–C bond.¹⁵⁵

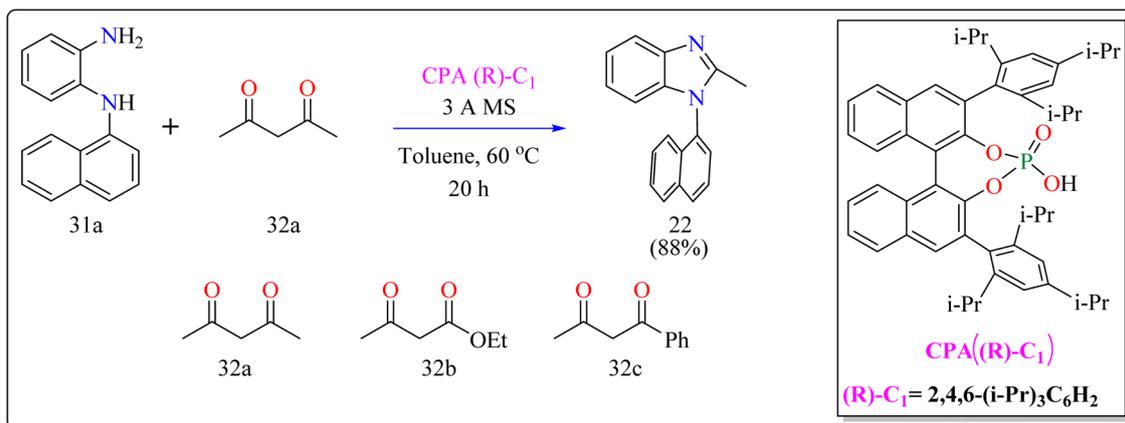
2.8. Tetrahydrocarbazole catalyzed synthesis of biaryl and aryl-heteroaryls

Agrochemicals, fragrance, pharmaceuticals, and dyes are just a few of the many industries that use biaryl and aryl-heteroaryl complexes as important structural components.^{156,157} Biaryl or

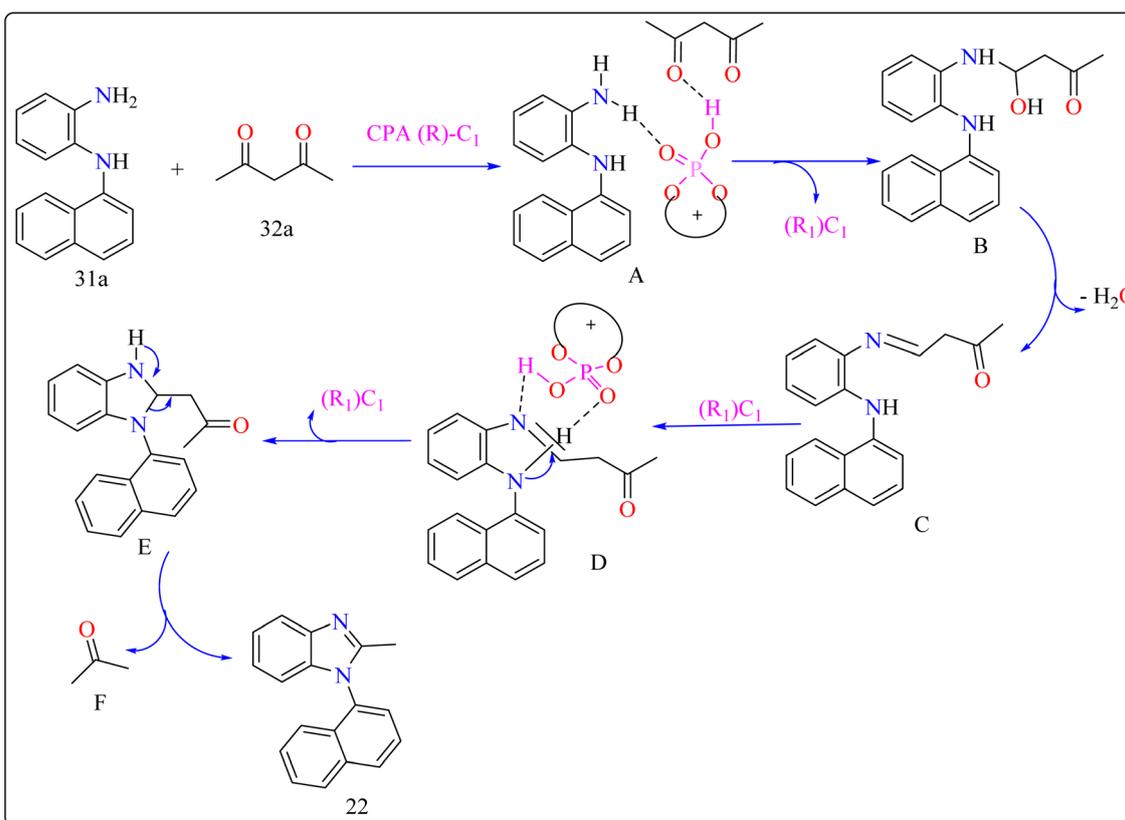
heteroaryl azo dyes are extensively researched for their potential uses in the textile industry, various clinical and biological studies, photochromic materials, indicators, sensors, and photo-sensitizers, as well as for their anti-microbial, antiseptic, and antioxidant properties.¹⁵⁸ Furthermore, researcher discussed the preparation of biaryl and aryl-heteroaryl compounds utilizing an electron donor acceptor (EDA) complex¹⁵⁹ produced by the reaction between the oxidant diazonium salt (**33a**) and the reductant tetrahydrocarbazole (THC), which has a high electron density. In particular, it was found that at milder conditions, EDA complex formed in catalytic levels was functional and facilitated a single electron transfer procedure without any photoactivation.

For synthesis, Biaryl used one equivalent of **33a**, Benzene (**34a**), and 10 mol% THC (**37a**), all of which were combined in a dark environment with N₂ at the ambient temperature for 14 h. Similarly for synthesis of aryl-heteroaryl repeated same procedure with 20 equiv. of heteroarene (thiophene (**35a**), furan (**36a**)). It was shown that THC (**37a**) surpassed other catalysts with comparable structural characteristics like simple **37b**, **37c**, **37d**, **37e**, **37f**, and **37g** during catalytic screening (Scheme 19).





Scheme 17 Optimized reaction conditions for CPAs catalyzed synthesis of auxiliary *N*-Aryl benzimidazoles.

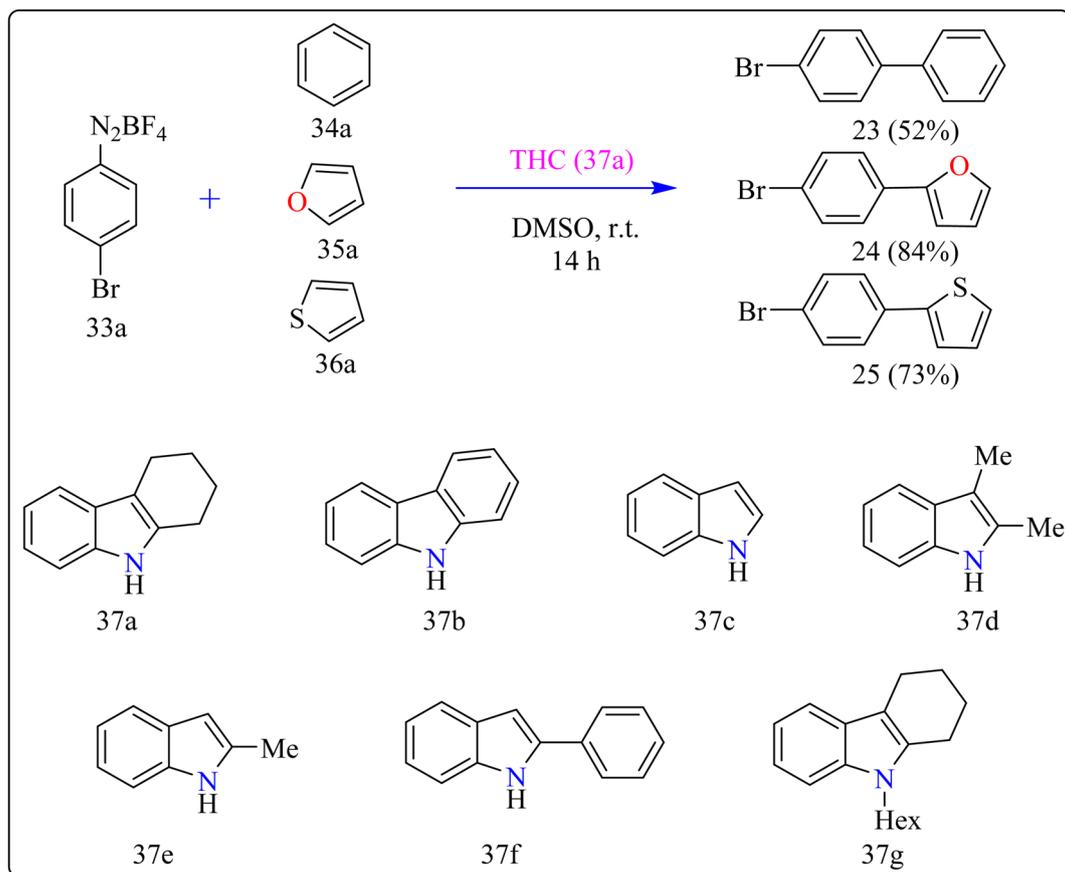


Scheme 18 Proposed reaction mechanism for CPA catalyzed compound **22**.

DMSO proved to be the most effective of the various solvents tested. The 10 mol% of **37a** catalyst provided maximum yield, conversely the yield decreased with 5 mol% of **37a** but the 20 mol% had no apparent effect on the final yield. As shown in Scheme 20, the creation of the EDA complex (**A**) between **33a** and **37a** initiates the process. Without any photoexcitation, the created complex **A** was passed through an easy Single Electron Transfer (SET). The **37a**, an organic reductant that contains nitrogen and is one of the elements that make up EDA, probably giving an electron to the component's **33a**. Then generated

radical cation **B**, radical anion species, and radical adduct likely breakdown into radical cation **C**, arylradical **D**, and nitrogen gas. The radical cationic entity **C** could be easily stabilized due to the existence of resonance patterns that include the stable tertiary free radical. Following a Hemolytic Aromatic Substitution (HAS) reaction with **35a**, the arylradical would produce the radical species **E**. Currently, the intermediate **C** has the ability to remove one electron from the radical arylated species **E** and revert to its original catalytic form, the **37a** molecule. After the borontetrafluoride anion deprotonated the cationic species **F**,





Scheme 19 Optimize reaction conditions for the synthesis of biaryl and aryl-heteroaryls.

the arylated compound 24 and HBF₄ were produced as byproducts.¹⁶⁰

2.9. Salicylic acid catalyzed reactions

Salicylic acid is a phenolic analogs, which are found in many plants, are essential for food preservation, cosmetics, medicines, and plant immunity.^{161,162} They play an important role as ligands in the catalysis of metal complexes in chemistry.¹⁶³ Additionally, their application breadth in the fields of medicine and material science is broadened by the use of organocatalysis in a variety of synthetic transformations, including arylation, hydrodeamination, oxidative acylation, along with the synthesis of different biologically active derivatives.^{164–166}

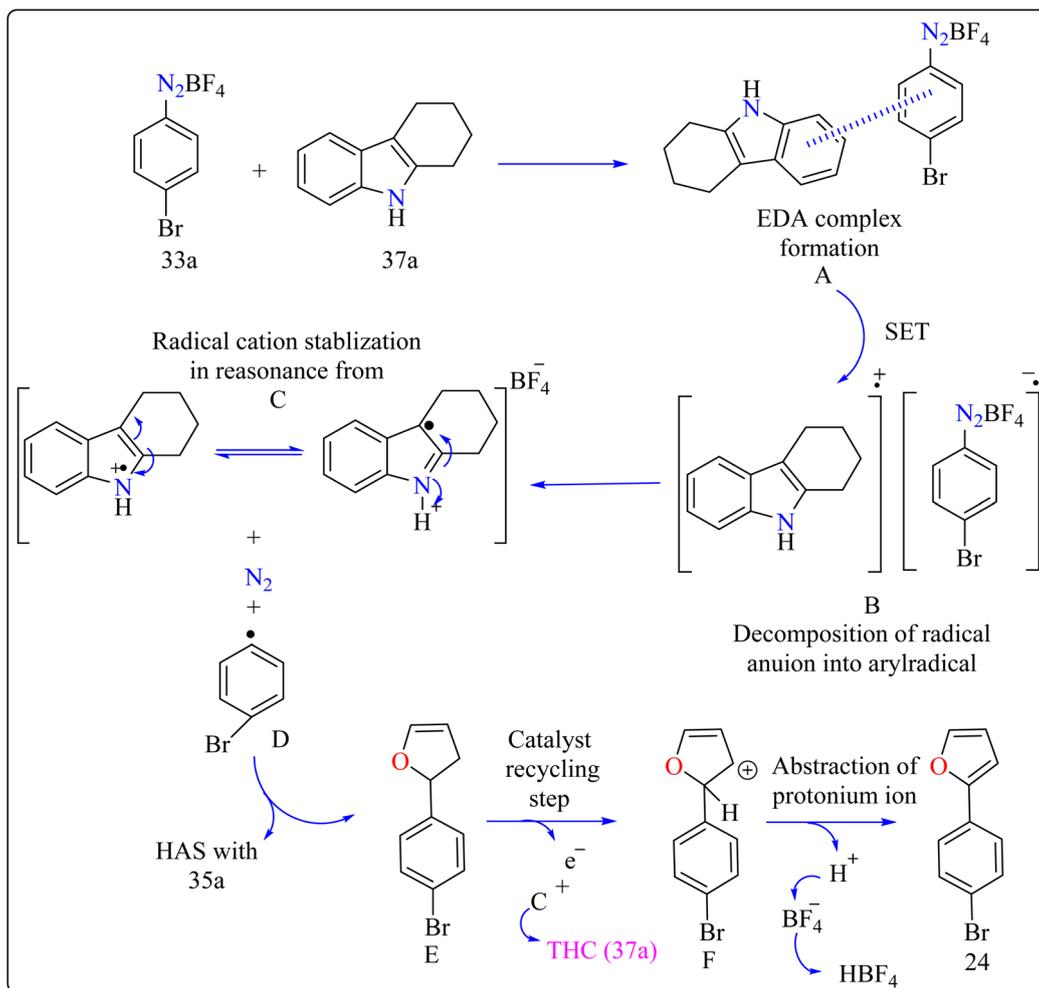
2.9.1. Synthesis of α -sulfonyl ketoximes. α -Sulfonyl ketoximes are useful in chemical synthesis and biologically active due to their variety of functional groups. They function as adaptable building blocks with a wide range of synthetic possibilities for different organic chemistry applications.^{167,168} A number of synthesis techniques have been described for α -sulfonyl ketoximes, mainly based on radical-mediated oximation and arylsulfonylation.^{169,170} Few organocatalytic techniques are known, and metal-catalyzed procedures predominate. Salicylic acid has been shown to have the ability to act as an organocatalyst in the synthesis of aryl radicals from arylamines and *t*-butyl nitrite, providing a new route for this reaction. Furthermore, DABSO improves ease and safety in these reactions by

acting as a secure replacement for sulfur dioxide.¹⁷¹ These biologically active molecules require more investigation into organocatalytic production from easily accessible starting materials.

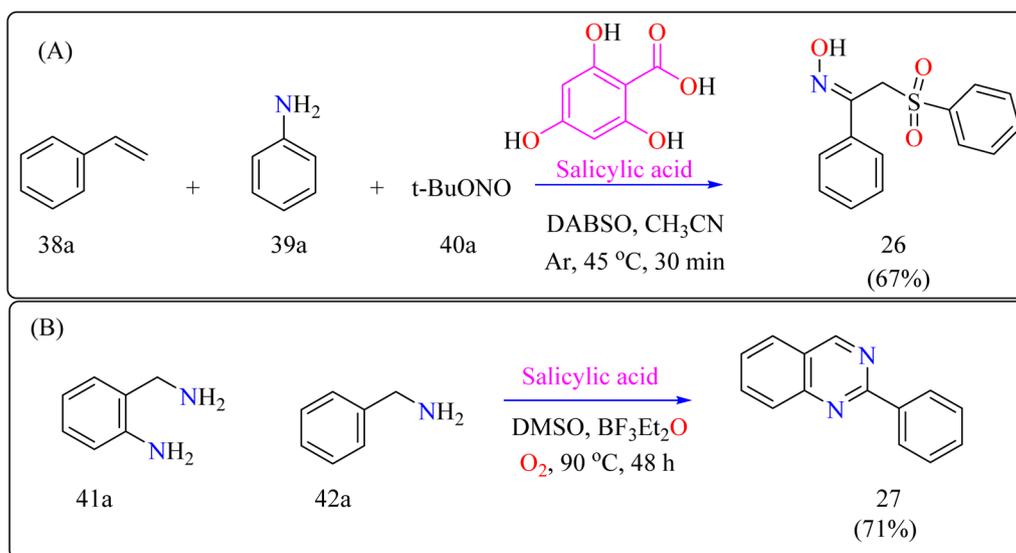
The first organocatalytic synthesis of α -sulfonyl ketoximes was reported by the researchers (Scheme 21A). This was achieved using a four-component reaction including styrene derivative (38a), anilines (39a), *t*-butyl nitrite (40a), and DABSO, all of which were catalyzed by salicylic acid.¹⁷² Salicylic acid (10 mol%) aided in the production of product 10a in 51% yield in acetonitrile under argon at the ambient temperature in 30 minutes, supporting its catalytic role. Maintaining 1.5 equiv. of DABSO and 3.0 equiv. 40a, optimization yielded a 67% yield at 45 °C. Optimal conditions were demonstrated by additional modifications to the reaction temperature and reactant loadings. Other examined solvents fared worse than acetonitrile. These results define effective parameters for this novel organocatalytic synthesis.¹⁷³

2.9.2. Synthesis of quinazolines. Modern developments in materials science and medicine require more environmentally friendly synthetic processes and fundamental chemicals with greater purity.^{174–176} Using oxygen and salicylic acid, a metal-free oxidation catalyst method allows the synthesis of a variety of functional compounds from unstable imines in a single pot.¹⁷⁷ This approach achieves great selectivity and environmental sustainability, making it suitable for multistep reactions such





Scheme 20 THC complex catalyzed reaction mechanism for compound 24.

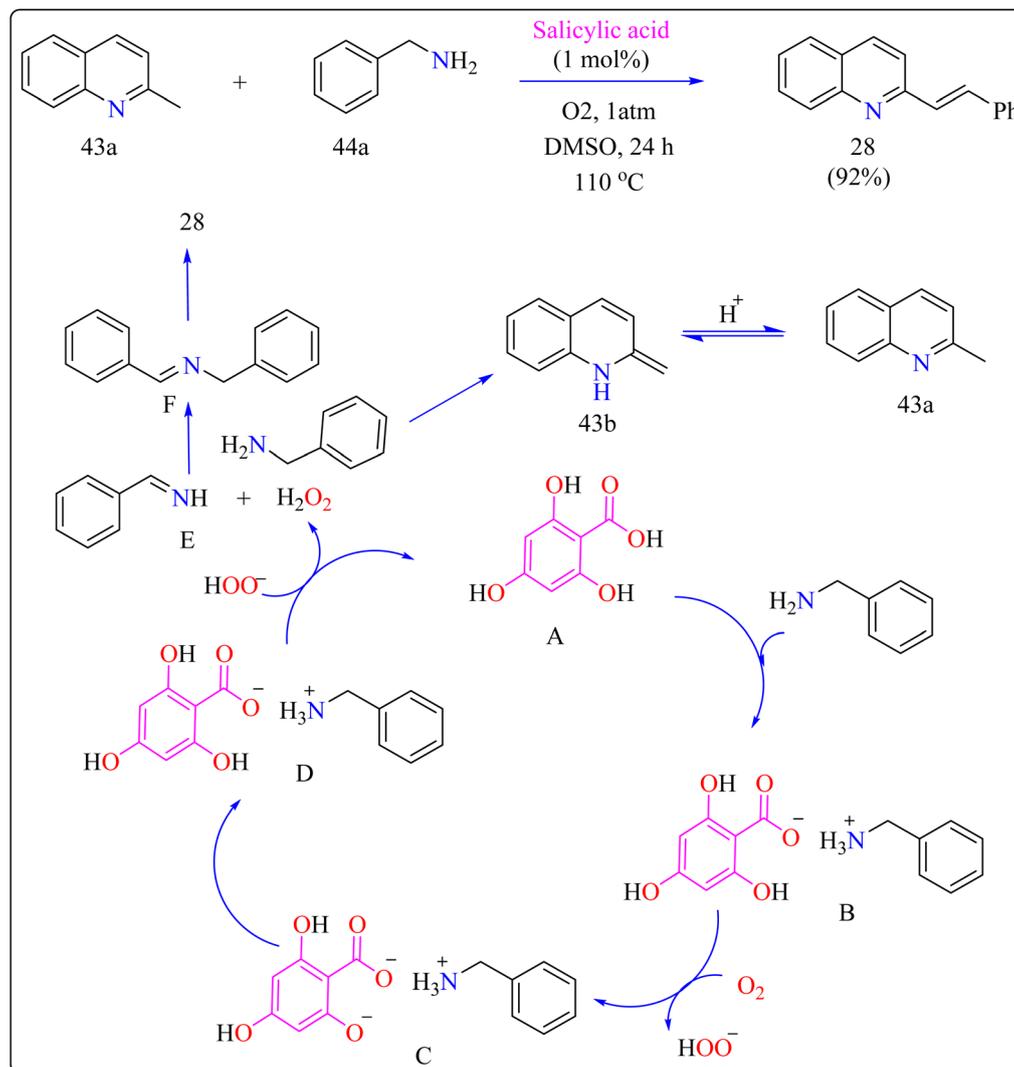
Scheme 21 At optimized reaction conditions methodology for the synthesis of (A) α -sulfonyl ketoximes (26) and, (B) quinazolines (27).

as the synthesis of quinazolines.¹⁷⁸ For industrial and medicinal chemistry, the development of scalable, metal-free synthesis techniques for quinazolines is essential, highlighting the need for better environmental conditions and wider use.¹⁷⁹ For synthesizing compound **27** as shown in Scheme 21B, salicylic acid (5 mol%), $\text{BF}_3\text{-Et}_2\text{O}$ (10 mol%) as an additive, 2-amino-benzylamine (**41a**), benzylamine (**42a**), and DMSO as a solvent (1.0 mL) were all mixed in a 10 mL two-necked flask with an O_2 atmosphere at 25 °C as part of a reaction setup. In an O_2 environment, the mixture was agitated for 48 hours at 90 °C. Using the activated alumina as the filler and iso-hexane for elution, column chromatography was used for purification following the reaction to produce product **27**.¹⁸⁰

2.9.3. Synthesis of *N*-heteroaryl stilbenoids. Stilbenoids are the structural component of squaraine dyes that are widely used in solar cell pigments, materials for non-linear optics, two-photon absorption compounds, NIR emitting fluorescent dyes, vivid violet, blue, or blue-green dyes came to be of enormous significance, components for biosensors and fluorescence imaging materials.¹⁸¹ For the production of medications,

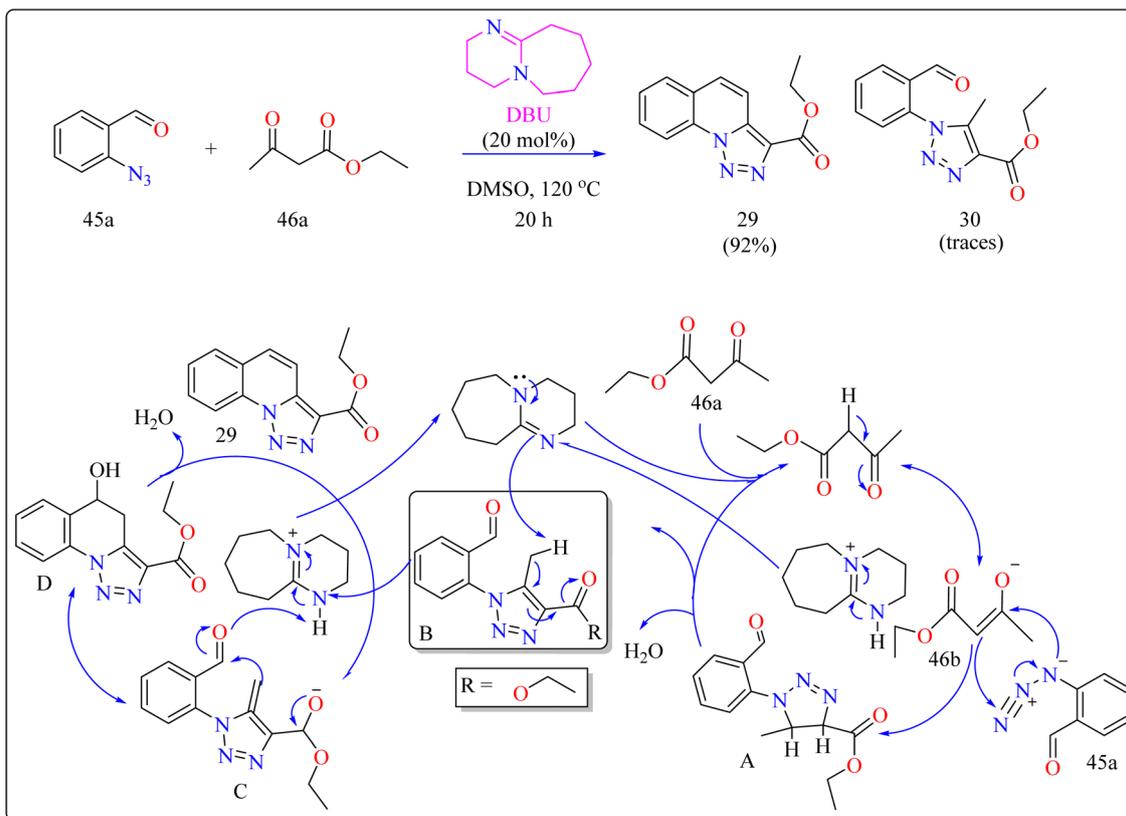
imaging agents, and materials such as organic light-emitting diodes, methyl *N*-heteroarenes must be olefinated.¹⁸²⁻¹⁸⁴ Aldehyde condensation or multi-step reactions are used in traditional procedures.¹⁸⁵⁻¹⁸⁷ Recent methods include using secondary or benzyl amines for metal-free deaminative olefination; however, more widespread use of environmentally friendly and effective catalytic systems free of metal residues is still desired.¹⁸⁸⁻¹⁹⁰

Scheme 22 showed, 2-methylquinoline (**43a**), benzyl amine (**44b**), 4,6-dihydroxysalicylic acid (**A**), and DMSO (4 mL) were added to a 10 mL Schlenk tube and agitated at room temperature in the presence of an O_2 balloon. Next, a warmed oil bath at 110 °C was used to hold the reaction mixture for 24 h. Further, the mixture was cooled with 5 mL of H_2O and extracted with ethyl acetate. Drying over sodium sulphate and subsequent filtering and concentration of the mixed organic layer. Compound **28** was obtained by purifying the residue using column chromatography on silica gel in which petroleum ether/ethyl acetate served as an eluent. Optimized reaction experiments concluded that, 92% of the **28** yield was obtained by



Scheme 22 Schematic diagram for Gram-scale preparation of (*E*)-2-styrylquinoline.





Scheme 23 Methodology and reaction route for the synthesis of DBU catalyzed compound 29.

reducing the salicylic acid catalyst to 1 mol%. The reaction was hampered by lower temperature and shorter times. Yield was somewhat impacted by extending the reaction time to 30 h. Utilizing solvents such as DMF, toluene, and CH₃CN, as well as reducing **44a** equiv., decreased yield. An excess of DMSO and a lack of O₂ reduced output.^{191,192}

2.10. DBU catalyzed synthesis of quinolines

In organic synthesis, 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU) an inexpensive, easily accessible organic base with a pK_a of 12 and due to low nucleophilicity is frequently utilized for multi-component reactions, cyclizations, and chemical derivatizations.^{193,194} It is useful in various reactions due to its steric hindrance, stability, and ease of handling. Additionally recoverable, DBU finds application as a complexing ligand, catalyst, and base in a variety of practical applications.¹⁹⁵

Quinolines are a type of heterocycles that exist both naturally and artificially and have fascinating biological properties.¹⁹⁶ Because of their many pharmacological properties, including anti-malarial, anti-oxidants, anxiolytic, and antitumoral effects, compounds with quinolinic cores in their structures have garnered a lot of attention in the field of material science and has the potential to act as an antileishmanial agent.^{197–200} In previous years, researcher reported triazolo quinoline-based synthesis of complexes catalyzed by copper iodide, copper(II), palladium catalyst and oxidative catalyst-free N–N coupling reaction was used to synthesize triazole derivatives.^{201–204} The

creation of more effective catalytic methods is still a major issue in synthetic organic chemistry, despite the notable progress made towards the preparation of quinoline-triazole hybrids and catalyzed by metal free DBU organocatalyst. This is a straightforward and effective method involving 2-azidobenzaldehyde (**45a**) and carbonyl compounds (**46a**) and DBU (20 mol%) as a catalyst was used for the organocatalytic synthesis of several fused quinolines **29** and traces of compound **30**, shown in Scheme 23. For synthesizing compound **29**, the enolate **46b** was formed by DBU eliminating one methylene acidic proton from molecule **46a**. Then, a [3 + 2] cycloaddition event occurs between enolate **46b** and **45a**, producing the intermediate triazoline **A**, which dehydrates to generate the intermediate **B**. The intermediate **B** then reacts with a catalyst, which extracts a proton from the methyl group at the 5-position of triazole **B**, with electron displacement by the triazole nucleus and the carbonyl, generating the enolate intermediate **C**. The negative charge in the oxygen then migrates, causing the carbon linked at position 5 of the triazole to undertake a nucleophilic attack on the pendent *ortho*-carbonyl group, resulting in the intermediate **D**, which after dehydration formed the compound **29**.²⁰⁵

3 Conclusion

As evident by the examples included in this paper, organocatalysis has already attained an advanced state of sophistication. The use of multidisciplinary research will undoubtedly



result in the growth of this field of catalysis, in which organo-catalyzed synthesis of various compounds. Because of the adaptability of organic synthesis and generally accepted theories on the effect of substituents on the electronic characteristics of organic compounds, organic components allow a significant degree of design flexibility with tunable qualities that are typically not feasible in inorganic materials. The idea can be supported theoretically by the growing number of synthetic compounds, and in the coming years, we might observe wider improvements in this field. This review's goal is to provide materials scientists, organic and industrial chemists, and other researchers with an overview of these developing advanced organocatalysts for simple eco-friendly organic transformations or building blocks of organic complexes. The basic idea of organo-catalysis originated from the use of simple and common organic molecules, readily available in the laboratory, to be used as catalysts for organic and inorganic transformations. It is highly recommended to the chemists to adopt this advanced idea and technology for chemical synthesis.

Data availability

No primary research results, software or code have been included and no new data were generated or analysed as part of this review.

Author contributions

A. Z.: writing – original draft, investigation. G. I.: review and editing. U. S. S.: data collection. F. J.: data curation and visualization. M. S.: formal analysis and validation. M. Y.: validation and review. Z. U.: data collection and formatting. M. A.: review and formatting. M. A. I.: conceptualization, resources, supervision, and overall guidance.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors are thankful to the higher education commission of Pakistan for awarding research grant NRPU # 8198 and Pakistan Science Foundation for awarding research grant PSF/CRP.Constr-676.

References

- 1 A. Zafar, T. H. Mawat, E. M. Atiyah, M. Adnan Iqbal, A. Majeed, G. Iram, R. Qureshi and S. Khalid, *ChemistrySelect*, 2024, **9**, e202304566.
- 2 T. Mou, H. S. Pillai, S. Wang, M. Wan, X. Han, N. M. Schweitzer, F. Che and H. Xin, *Nat. Catal.*, 2023, **6**, 122–136.
- 3 R. A. Sheldon, *Catal. Today*, 2011, **167**, 3–13.
- 4 V. d. G. Oliveira, M. F. d. C. Cardoso and L. d. S. M. Forezi, *Catalysts*, 2018, **8**, 605.
- 5 L. S. Hegedus, *J. Am. Chem. Soc.*, 2009, **131**, 17995–17997.
- 6 D. D. Gadge and P. S. Kulkarni, *J. Heterocycl. Chem.*, 2022, **59**, 1320–1332.
- 7 Y. Meng, J. Li, H. Liu, H. Wu and H. Li, *Chem. Commun.*, 2024, **60**, 1642–1645.
- 8 P. Pecchini, M. Fochi, F. Bartocchini, G. Piersanti and L. Bernardi, *Chem. Sci.*, 2024, **15**, 5832–5868.
- 9 B. Borah, K. D. Dwivedi and L. R. Chowhan, *RSC Adv.*, 2021, **11**, 13585–13601.
- 10 Z.-W. Zhang, S.-W. Liu, H.-P. Huang, Y.-H. Xie, R.-C. Huang, Y.-Q. Deng and N. Lin, *RSC Adv.*, 2023, **13**, 31047–31058.
- 11 A. Majeed, A. H. Ibrahim, S. S. Al-Rawi, M. A. Iqbal, M. Kashif, M. Yousif, Z. U. Abidin, S. Ali, M. Arbaz and S. A. Hussain, *ACS Omega*, 2024, **9**, 12069–12083.
- 12 M. Yousif, A. H. Ibrahim, S. S. Al-Rawi, A. Majeed, M. A. Iqbal, M. Kashif, Z. U. Abidin, M. Arbaz, S. Ali and S. A. Hussain, *RSC Adv.*, 2024, **14**, 16138–16149.
- 13 Z. U. Abidin, A. Majeed, M. A. Iqbal, M. Kashif, T. Fatima, M. Yousif, M. Arbaz, S. A. Hussain and M. Sajid, *Clean Technol. Environ. Policy*, 2024, 1–15.
- 14 W. Yao, E. A. Bazan-Bergamino and M. Y. Ngai, *ChemCatChem*, 2022, **14**, e202101292.
- 15 B. Vachan, M. Karuppasamy, P. Vinoth, S. Vivek Kumar, S. Perumal, V. Sridharan and J. C. Menéndez, *Adv. Synth. Catal.*, 2020, **362**, 87–110.
- 16 S. D. Pasuparthi and B. Maiti, *ChemistrySelect*, 2022, **7**, e202104261.
- 17 M. P. van Der Helm, B. Klemm and R. Eelkema, *Nat. Rev. Chem*, 2019, **3**, 491–508.
- 18 Y.-H. Oh, D. W. Kim and S. Lee, *Molecules*, 2022, **27**, 5702.
- 19 H. Li and S. Yang, *Int. J. Chem. Eng.*, 2014, 2014.
- 20 M. Odagi and K. Nagasawa, *Chem. Rec.*, 2023, e202300030.
- 21 A. J. Burke, *Expert Opin. Drug Discovery*, 2023, **18**, 37–46.
- 22 V. A. Larionov, B. L. Feringa and Y. N. Belokon, *Chem. Soc. Rev.*, 2021, **50**, 9715–9740.
- 23 F. Auria-Luna, S. Mohammadi, M. Divar, M. C. Gimeno and R. P. Herrera, *Adv. Synth. Catal.*, 2020, **362**, 5275–5300.
- 24 D. Rani, M. Bhargava and J. Agarwal, *ChemistrySelect*, 2020, **5**, 2435–2440.
- 25 A. Sofi, J. J. Regita, B. Rane and H. H. Lau, *Mech. Syst. Signal Process.*, 2022, **163**, 108113.
- 26 M. R. Miah, M. Yang, S. Khandaker, M. M. Bashar, A. K. D. Alsukaibi, H. M. Hassan, H. Znad and M. R. Awual, *Sens. Actuators, A*, 2022, **347**, 113933.
- 27 E. F. Durán-Lara, A. Valderrama and A. Marican, *Agriculture*, 2020, **10**, 41.
- 28 M. Chandrasekaran, M. Paramasivan and J. J. Sahayarayan, *Microorganisms*, 2022, **11**, 42.
- 29 R. Rana, S. Nanda, A. Maclennan, Y. Hu, J. A. Kozinski and A. K. Dalai, *J. Energy Chem.*, 2019, **31**, 107–118.
- 30 J. A. Okolie, S. Nanda, A. K. Dalai and J. A. Kozinski, *Waste Biomass Valorization*, 2021, **12**, 2145–2169.
- 31 K. Watanabe, N. Terao, T. Niwa and T. Hosoya, *J. Org. Chem.*, 2021, **86**, 11822–11834.
- 32 A. Santarangelo, F. Weber, S. Kehraus, J. S. Dickschat and A. Schieber, *Food Chem.*, 2023, 136473.



- 33 A. Zafar, C. Takeda, A. Manzoor, D. Tanaka, M. Kobayashi, Y. Wadayama, D. Nakane, A. Majeed, M. A. Iqbal and T. Akitsu, *Molecules*, 2024, **29**, 398.
- 34 E. Stempel and T. Gaich, *Acc. Chem. Res.*, 2016, **49**, 2390–2402.
- 35 K. Biradha, M. Sarkar and L. Rajput, *Chem. Commun.*, 2006, 4169–4179.
- 36 M. Indrani, R. Ramasubramanian, S. Kumaresan, S. K. Kang, M. Chen and M. Du, *Polyhedron*, 2008, **27**, 3593–3600.
- 37 J.-P. Zou, M.-H. Chen, L.-Z. Zhang, Q.-J. Xing and Z.-Q. Xiong, *J. Chem. Crystallogr.*, 2011, **41**, 1820–1833.
- 38 T. Ohmura, Y. Morimasa and M. Suginoe, *Chem. Lett.*, 2017, **46**, 1793–1796.
- 39 L. C. Misal Castro, I. Sultan, K. Nishi, H. Tsurugi and K. Mashima, *J. Org. Chem.*, 2021, **86**, 3287–3299.
- 40 H. Hosoya, L. C. Misal Castro, I. Sultan, Y. Nakajima, T. Ohmura, K. Sato, H. Tsurugi, M. Suginoe and K. Mashima, *Org. Lett.*, 2019, **21**, 9812–9817.
- 41 C. Fischer, F. Lipata and J. Rohr, *J. Am. Chem. Soc.*, 2003, **125**, 7818–7819.
- 42 J. DiGiovanni, P. Decina, W. Prichett, J. Cantor, K. Aalfs and M. Coombs, *Cancer Res.*, 1985, **45**, 2584–2589.
- 43 K. Srinivas, K. Yesudas, K. Bhanuprakash, V. J. Rao and L. Giribabu, *J. Phys. Chem. C*, 2009, **113**, 20117–20126.
- 44 R. Iwaura, M. Ohnishi-Kameyama and T. Iizawa, *Chem.–Eur. J.*, 2009, **15**, 3729–3735.
- 45 C. A. Kuttruff, H. Zipse and D. Trauner, *Angew. Chem., Int. Ed.*, 2011, **6**, 1402–1405.
- 46 Y. Koyama, R. Yamaguchi and K. Suzuki, *Angew. Chem., Int. Ed. Engl.*, 2008, **47**, 1084.
- 47 E. E. Podlesny and M. C. Kozlowski, *Org. Lett.*, 2012, **14**, 1408–1411.
- 48 F. Diaz, H.-B. Chai, Q. Mi, B.-N. Su, J. S. Vigo, J. G. Graham, F. Cabieses, N. R. Farnsworth, G. A. Cordell and J. M. Pezzuto, *J. Nat. Prod.*, 2004, **67**, 352–356.
- 49 H. Prinz, Y. Ishii, T. Hirano, T. Stoiber, J. A. Camacho Gomez, P. Schmidt, H. Düssmann, A. M. Burger, J. H. Prehn and E. G. Günther, *J. Med. Chem.*, 2003, **46**, 3382–3394.
- 50 H.-S. Huang, J.-M. Hwang, Y.-M. Jen, J.-J. Lin, K.-Y. Lee, C.-H. Shi and H.-C. Hsu, *Chem. Pharm. Bull.*, 2001, **49**, 969–973.
- 51 S. Das, A. Bhowmik, W. Sarkar, A. Mishra and I. Deb, *J. Org. Chem.*, 2021, **86**, 4131–4142.
- 52 A. D'Amato and G. Della Sala, *Catalysts*, 2021, **11**, 1545.
- 53 C. Zhou, Z. Liu, G. Liang, Y.-Q. Zhang, T. Lei, B. Chen, R.-Z. Liao, C.-H. Tung and L.-Z. Wu, *Org. Lett.*, 2024, **26**, 1116–1121.
- 54 R. Harrison and B. Rickborn, *Org. Lett.*, 2002, **4**, 1711–1713.
- 55 T. Kitamura, K. Harano and T. Hisano, *Chem. Pharm. Bull.*, 1992, **40**, 2255–2261.
- 56 T. J. Osberger, D. C. Rogness, J. T. Kohrt, A. F. Stepan and M. C. White, *Nature*, 2016, **537**, 214–219.
- 57 S. Das, A. Bhowmik, W. Sarkar, A. Mishra and I. Deb, *J. Org. Chem.*, 2021, **86**, 4131–4142.
- 58 N. A. McGrath, M. Brichacek and J. T. Njardarson, *J. Chem. Educ.*, 2010, **87**, 1348–1349.
- 59 M. Baumann, I. R. Baxendale, S. V. Ley and N. Nikbin, *Beilstein J. Org. Chem.*, 2011, **7**, 442–495.
- 60 M. R. Barbachyn and C. W. Ford, *Angew. Chem., Int. Ed.*, 2003, **42**, 2010–2023.
- 61 A. R. Renslo, G. W. Luehr and M. F. Gordeev, *Bioorg. Med. Chem.*, 2006, **14**, 4227–4240.
- 62 Y. Kumatabara, M. Okada and S. Shirakawa, *ACS Sustain. Chem. Eng.*, 2017, **5**, 7295–7301.
- 63 H. Büttner, J. Steinbauer and T. Werner, *ChemSusChem*, 2015, **8**, 2655–2669.
- 64 S. Liu, N. Suematsu, K. Maruoka and S. Shirakawa, *Green Chem.*, 2016, **18**, 4611–4615.
- 65 S. Shirakawa and K. Maruoka, *Tetrahedron Lett.*, 2014, **55**, 3833–3839.
- 66 S. Liu, Y. Kumatabara and S. Shirakawa, *Green Chem.*, 2016, **18**, 331–341.
- 67 K. Okuno, R. Nishiyori, M. Hiraki and S. Shirakawa, *Heterocycles*, 2021, **103**, 94–109.
- 68 K. Okuno, R. Nishiyori, K. Abe, T. Mori and S. Shirakawa, *Chirality*, 2022, **34**, 915–924.
- 69 I. Shibata, A. Baba, H. Iwasaki and H. Matsuda, *J. Org. Chem.*, 1986, **51**, 2177–2184.
- 70 G. P. Speranza and W. Peppel, *J. Org. Chem.*, 1958, **23**, 1922–1924.
- 71 R. Nishiyori, K. Okuno and S. Shirakawa, *Eur. J. Org. Chem.*, 2020, **2020**, 4937–4941.
- 72 M. Li, W. He and S.-Y. Zhang, *Mini-Rev. Org. Chem.*, 2022, **19**, 146–165.
- 73 E. P. Carreiro, A. J. Burke, G. J. Hermann and H.-J. Federsel, *Tetrahedron Green Chem.*, 2023, **2**, 100025.
- 74 H. Joshi and V. K. Singh, *Asian J. Org. Chem.*, 2022, **11**, e202100053.
- 75 G. Varvounis, *Adv. Heterocycl. Chem.*, 2009, **98**, 143–224.
- 76 J.-W. Byun, D.-H. Lee and Y.-S. Lee, *Tetrahedron Lett.*, 2003, **44**, 8063–8067.
- 77 J. S. Casas, M. S. Garcia-Tasende, A. Sanchez, J. Sordo and A. Touceda, *Coord. Chem. Rev.*, 2007, **251**, 1561–1589.
- 78 F. Lehmann, M. Holm and S. Laufer, *J. Comb. Chem.*, 2008, **10**, 364–367.
- 79 G. Mustafa, M. Zia-ur-Rehman, S. H. Sumrra, M. Ashfaq, W. Zafar and M. Ashfaq, *J. Mol. Struct.*, 2022, **1262**, 133044.
- 80 X.-L. Zhao, M. Wang, J. Geng and W. Huang, *Polyhedron*, 2021, **202**, 115208.
- 81 B. Mondal, R. Maity and S. C. Pan, *J. Org. Chem.*, 2018, **83**, 8645–8654.
- 82 S. Gogoi and C.-G. Zhao, *Tetrahedron Lett.*, 2009, **50**, 2252–2255.
- 83 B. Formánek, V. Šeferna, M. Meazza, R. Rios, M. Patil and J. Veselý, *Eur. J. Org. Chem.*, 2021, **2021**, 2362–2366.
- 84 D. L. Isac, A. Airinei, D. Maftai, I. Humelnicu, F. Mocci, A. Laaksonen and M. Pinteală, *J. Phys. Chem. A*, 2019, **123**, 5525–5536.
- 85 Y. Guo, L. Yao, L. Luo, H.-X. Wang, Z. Yang, Z. Wang, S.-L. Ai, Y. Zhang, Q.-C. Zou and H.-L. Zhang, *Org. Chem. Front.*, 2021, **8**, 239–248.



- 86 P. Sivaprakasam, A. Xie and R. J. Doerksen, *Bioorg. Med. Chem.*, 2006, **14**, 8210–8218.
- 87 Y. Fan, Y. Lu, X. Chen, B. Tekwani, X.-C. Li and Y. Shen, *Molecules*, 2018, **23**, 2878.
- 88 J. M. Ravasco, H. Faustino, A. Trindade and P. M. Gois, *Chem.–Eur. J.*, 2019, **25**, 43–59.
- 89 F. Yuan, W. Duan, Z. Li, X. Luo, M. Zhang, H. Deng and L. Song, *Synthesis*, 2019, **51**, 3345–3355.
- 90 N. Sakai, K. Kawashima, M. Kajitani, S. Mori and T. Oriyama, *Org. Lett.*, 2021, **23**, 5714–5718.
- 91 K.-S. Lee, J. J. Kweon, I.-H. Oh and C. E. Lee, *J. Phys. Chem. Solids*, 2012, **73**, 890–895.
- 92 R. West and D. L. Powell, *J. Am. Chem. Soc.*, 1963, **85**, 2577–2579.
- 93 N. Azizi, A. Davoudpour, F. Eskandari and E. Batebi, *Monatsh. Chem. Chem. Mon.*, 2013, **144**, 405–409.
- 94 N. Azizi, E. Gholibeghlo and Z. Manocheri, *Sci. Iran.*, 2012, **19**, 574–578.
- 95 N. Azizi and A. Davoudpour, *Catal. Lett.*, 2011, **141**, 1506–1510.
- 96 N. Azizi, E. Saki and M. Edrisi, *C. R. Chim.*, 2011, **14**, 973–977.
- 97 H.-J. Zhang, X.-Z. Wang, Q. Cao, G.-H. Gong and Z.-S. Quan, *Bioorg. Med. Chem. Lett.*, 2017, **27**, 4409–4414.
- 98 M. Brana, M. Garrido, M. L. Rodriguez, M. Morcillo, Y. Alvarez, Y. Valladares and G. Klebe, *Eur. J. Med. Chem.*, 1990, **25**, 209–215.
- 99 W. Wasulko, A. C. Noble and F. D. Popp, *J. Med. Chem.*, 1966, **9**, 599–601.
- 100 V. A. Kharlanov, *J. Photochem. Photobiol., A*, 1999, **122**, 191–197.
- 101 B. Alici, İ. Özdemir, K. Karaaslan, E. Çetinkaya and B. Çetinkaya, *J. Mol. Catal. A: Chem.*, 2005, **231**, 261–264.
- 102 R. Malherbe, G. Rist and D. Bellus, *J. Org. Chem.*, 1983, **48**, 860–869.
- 103 M. Azam, I. Warad, S. I. Al-Resayes, N. Alzaqri, M. R. Khan, R. Pallepogu, S. Dwivedi, J. Musarrat and M. Shakir, *J. Mol. Struct.*, 2013, **1047**, 48–54.
- 104 H. Alinezhad and M. Zare, *J. Chil. Chem. Soc.*, 2013, **58**, 1840–1841.
- 105 M. A. Bodaghifard, S. Asadbegi and Z. Bahrami, *J. Iran. Chem. Soc.*, 2017, **14**, 365–376.
- 106 S. Khopkar and G. Shankarling, *J. Chem. Sci.*, 2020, **132**, 1–10.
- 107 D. Y. Kim, M.-J. Kim, G. Sung and J.-Y. Sun, *Nano Convergence*, 2019, **6**, 1–24.
- 108 L. Quan, T. Sun, Y. Wei, Y. Lin, T. Gong, C. Pan, H. Ding, W. Liu and Z. Xie, *Colloids Surf., B*, 2019, **173**, 884–890.
- 109 B. Valeur and M. N. Berberan-Santos, *Molecular Fluorescence: Principles and Applications*, John Wiley & Sons, 2012.
- 110 J. Zhang, L. Xu, W. Xiao, Y. Chen, Z. Dong, J. Xu and C. Lei, *Eur. Polym. J.*, 2021, **157**, 110643.
- 111 S. Agarwal, P. Kalal, A. Sethiya and J. Soni, *Mini-Rev. Org. Chem.*, 2022, **19**, 617–628.
- 112 K. A. Chavan, M. Shukla, A. N. S. Chauhan, S. Maji, G. Mali, S. Bhattacharyya and R. D. Erande, *ACS Omega*, 2022, **7**, 10438–10446.
- 113 G. P. Salze and D. A. Davis, *Aquaculture*, 2015, **437**, 215–229.
- 114 J. Menzie, C. Pan, H. Prentice and J.-Y. Wu, *Amino Acids*, 2014, **46**, 31–46.
- 115 N. Daneshvar, F. Shirini, M. S. N. Langarudi and R. Karimi-Chayjani, *Bioorg. Chem.*, 2018, **77**, 68–73.
- 116 F. Shirini and N. Daneshvar, *RSC Adv.*, 2016, **6**, 110190–110205.
- 117 H. S. Bhatti and S. Seshadri, *Color. Technol.*, 2004, **120**, 151–155.
- 118 V. S. Patil, V. S. Padalkar and N. Sekar, *J. Fluoresc.*, 2014, **24**, 1077–1086.
- 119 A. V. Chate, P. P. Rudrawar, G. M. Bondle and J. N. Sangeshetti, *Synth. Commun.*, 2020, **50**, 226–242.
- 120 M. Abd El Aleem Ali El-Remaily and O. M. Elhady, *ChemistrySelect*, 2020, **5**, 12098–12102.
- 121 C. O. Kappe, *J. Org. Chem.*, 1997, **62**, 7201–7204.
- 122 M. P. Parmar, D. P. Vala, S. S. Bhalodiya, D. B. Upadhyay, C. D. Patel, S. G. Patel, S. R. Gandholi, A. H. Shaik, A. D. Miller and J. Nogales, *RSC Adv.*, 2024, **14**, 9300–9313.
- 123 M. R. R. Hormi Mecadon, I. Kharbangar, B. M. Laloo, I. Kharkongor, M. Rajbangshi and B. Myrboh, *Tetrahedron Lett.*, 2011, **52**, 3228–3231.
- 124 C. Mukhopadhyay, P. K. Tapaswi and R. J. Butcher, *Tetrahedron Lett.*, 2010, **51**, 1797–1802.
- 125 J. G. Hernández, V. García-López and E. Juaristi, *Tetrahedron*, 2012, **68**, 92–97.
- 126 G. Coulthard, W. Erb and V. K. Aggarwal, *Nature*, 2012, **489**, 278–281.
- 127 S. Rao, D. Mohan and S. Adimurthy, *J. Biomol. Res. Ther.*, 2016, **5**, 2.
- 128 A. Khalafi-Nezhad, S. Sarikhani, E. S. Shahidzadeh and F. Panahi, *Green Chem.*, 2012, **14**, 2876–2884.
- 129 A. R. Sayed, S. M. Gomha, H. M. Abd El-lateef and T. Z. Abolibda, *Green Chem. Lett. Rev.*, 2021, **14**, 180–189.
- 130 J. H. Choi, J. M. Jeon, M. H. Kim, A. D. Towns and C. Yoon, *Color. Technol.*, 2008, **124**, 92–99.
- 131 J. h. Choi, J. y. Choi, H. y. Lee, A. D. Towns and C. Yoon, *Color. Technol.*, 2008, **124**, 364–369.
- 132 T. B. Shah, R. S. Shiny, R. B. Dixit and B. C. Dixit, *J. Saudi Chem. Soc.*, 2014, **18**, 985–992.
- 133 Y. Zhan, X. Zhao and W. Wang, *Dyes Pigm.*, 2017, **146**, 240–250.
- 134 M. S. Akhtar and Y. R. Lee, *J. Org. Chem.*, 2020, **85**, 15129–15138.
- 135 Y.-M. Li, F.-Y. Kwong, W.-Y. Yu and A. S. Chan, *Coord. Chem. Rev.*, 2007, **251**, 2119–2144.
- 136 L. Pu, *Acc. Chem. Res.*, 2012, **45**, 150–163.
- 137 M. C. Kozłowski, B. J. Morgan and E. C. Linton, *Chem. Soc. Rev.*, 2009, **38**, 3193–3207.
- 138 M. Shibasaki and S. Matsunaga, *Chem. Soc. Rev.*, 2006, **35**, 269–279.
- 139 G. Ma and M. P. Sibi, *Chem.–Eur. J.*, 2015, **21**, 11644–11657.
- 140 E. Kumarasamy, R. Raghunathan, M. P. Sibi and J. Sivaguru, *Chem. Rev.*, 2015, **115**, 11239–11300.



- 141 N. Di Iorio, S. Crotti and G. Bencivenni, *Chem. Rec.*, 2019, **19**, 2095–2104.
- 142 J. K. Cheng, S.-H. Xiang, S. Li, L. Ye and B. Tan, *Chem. Rev.*, 2021, **121**, 4805–4902.
- 143 Y.-B. Wang and B. Tan, *Acc. Chem. Res.*, 2018, **51**, 534–547.
- 144 S. D. Vaidya, S. T. Toenjes, N. Yamamoto, S. M. Maddox and J. L. Gustafson, *J. Am. Chem. Soc.*, 2020, **142**, 2198–2203.
- 145 H. Lee, M. Fischer, B. K. Shoichet and S.-Y. Liu, *J. Am. Chem. Soc.*, 2016, **138**, 12021–12024.
- 146 J. Yang, J.-W. Zhang, W. Bao, S.-Q. Qiu, S. Li, S.-H. Xiang, J. Song, J. Zhang and B. Tan, *J. Am. Chem. Soc.*, 2021, **143**, 12924–12929.
- 147 M. Sabat, J. C. VanRens, M. J. Laufersweiler, T. A. Brugel, J. Maier, A. Golebiowski, B. De, V. Easwaran, L. C. Hsieh and R. L. Walter, *Bioorg. Med. Chem. Lett.*, 2006, **16**, 5973–5977.
- 148 N. H. Huel, H. Nar, H. Priepke, U. Ries, J.-M. Stassen and W. Wienen, *J. Med. Chem.*, 2002, **45**, 1757–1766.
- 149 S. Bait, S. Shinde, R. Adivarekar and N. Sekar, *J. Indian Chem. Soc.*, 2021, **98**, 100236.
- 150 G. Schwartz, K. Fehse, M. Pfeiffer, K. Walzer and K. Leo, *Appl. Phys. Lett.*, 2006, **89**, 083509.
- 151 J. A. Asensio and P. Gómez-Romero, *Fuel Cells*, 2005, **5**, 336–343.
- 152 M. Yamanaka, J. Itoh, K. Fuchibe and T. Akiyama, *J. Am. Chem. Soc.*, 2007, **129**, 6756–6764.
- 153 Y. X. Jia, J. Zhong, S. F. Zhu, C. M. Zhang and Q. L. Zhou, *Angew. Chem.*, 2007, **119**, 5661–5663.
- 154 J. Itoh, K. Fuchibe and T. Akiyama, *Angew. Chem.*, 2008, **120**, 4080–4082.
- 155 N. Man, Z. Lou, Y. Li, H. Yang, Y. Zhao and H. Fu, *Org. Lett.*, 2020, **22**, 6382–6387.
- 156 J.-P. Corbet and G. Mignani, *Chem. Rev.*, 2006, **106**, 2651–2710.
- 157 J. Hassan, M. Sévignon, C. Gozzi, E. Schulz and M. Lemaire, *Chem. Rev.*, 2002, **102**, 1359–1470.
- 158 A. K. Behera and A. Sen, *J. Photochem. Photobiol., A*, 2022, **433**, 114146.
- 159 A. d. A. Bartolomeu, R. C. Silva, T. J. Brocksom, T. Noël and K. T. de Oliveira, *J. Org. Chem.*, 2019, **84**, 10459–10471.
- 160 R. Saritha, S. B. Annes, S. Saravanan and S. Ramesh, *Org. Biomol. Chem.*, 2020, **18**, 2510–2515.
- 161 M. Shara and S. J. Stohs, *Phytother. Res.*, 2015, **29**, 1112–1116.
- 162 S. Hayat, B. Ali and A. Ahmad, *Salicylic Acid: A Plant Hormone*, 2007, pp. 1–14.
- 163 H. R. Kricheldorf and S. M. Weidner, *Eur. Polym. J.*, 2019, **119**, 37–44.
- 164 D. Felipe-Blanco and J. C. Gonzalez-Gomez, *Adv. Synth. Catal.*, 2018, **360**, 2773–2778.
- 165 D. Felipe-Blanco, F. Alonso and J. C. Gonzalez-Gomez, *Adv. Synth. Catal.*, 2017, **359**, 2857–2863.
- 166 M. Liu, X. Chen, T. Chen, Q. Xu and S.-F. Yin, *Org. Biomol. Chem.*, 2017, **15**, 9845–9854.
- 167 V. Padmavathi, B. C. O. Reddy, P. Thriveni and A. Nagendra Mohan, *Synth. Commun.*, 2007, **37**, 3127–3142.
- 168 F. Chen, N.-N. Zhou, J.-L. Zhan, B. Han and W. Yu, *Org. Chem. Front.*, 2017, **4**, 135–139.
- 169 N. Liu, P. Yin, Y. Chen, Y. Deng and L. He, *Eur. J. Org. Chem.*, 2012, **2012**, 2711–2714.
- 170 B. Wang, Z. Yan, L. Liu, J. Wang, Z. Zha and Z. Wang, *Green Chem.*, 2019, **21**, 205–212.
- 171 J. Yang, Y. Y. Liu, R. J. Song, Z. H. Peng and J. H. Li, *Adv. Synth. Catal.*, 2016, **358**, 2286–2292.
- 172 A. Dahiya, A. K. Sahoo, T. Alam and B. K. Patel, *Chem.-Asian J.*, 2019, **14**, 4454–4492.
- 173 S. Jakkampudi, N. Sakkani and J.-G. Zhao, *Tetrahedron Lett.*, 2021, **76**, 153229.
- 174 K. R. Campos, P. J. Coleman, J. C. Alvarez, S. D. Dreher, R. M. Garbaccio, N. K. Terrett, R. D. Tillyer, M. D. Truppo and E. R. Parmee, *Science*, 2019, **363**, eaat0805.
- 175 J. Garcia-Martinez, *Angew. Chem.*, 2021, **133**, 5008–5012.
- 176 I. T. Horváth, *Chem. Rev.*, 2018, **118**, 369–371.
- 177 C.-p. Dong, Y. Higashiura, K. Marui, S. Kumazawa, A. Nomoto, M. Ueshima and A. Ogawa, *ACS Omega*, 2016, **1**, 799–807.
- 178 Y. Yamamoto, S. Kodama, R. Nishimura, A. Nomoto, M. Ueshima and A. Ogawa, *J. Org. Chem.*, 2021, **86**, 11571–11582.
- 179 M. Faisal and A. Saeed, *Front. Chem.*, 2021, **8**, 594717.
- 180 Y. Yamamoto, C. Yamakawa, R. Nishimura, C.-P. Dong, S. Kodama, A. Nomoto, M. Ueshima and A. Ogawa, *Front. Chem.*, 2022, **9**, 822841.
- 181 H. Meier, *Z. Naturforsch. B*, 2019, **74**, 241–254.
- 182 M. Dhanawat, D. K. Mehta and R. Das, *Mini-Rev. Med. Chem.*, 2021, **21**, 1849–1864.
- 183 P. Das, M. D. Delost, M. H. Qureshi, D. T. Smith and J. T. Njardarson, *J. Med. Chem.*, 2018, **62**, 4265–4311.
- 184 W. Cieslik, J. Szczepaniak, A. Krasowska and R. Musiol, *Molecules*, 2020, **25**, 345.
- 185 S. Mahajan, N. Slathia, V. K. Nuthakki, S. B. Bharate and K. K. Kapoor, *RSC Adv.*, 2020, **10**, 15966–15975.
- 186 E. Liang, J. Wang, Y. Wu, L. Huang, X. Yao and X. Tang, *Adv. Synth. Catal.*, 2019, **361**, 3619–3623.
- 187 N. Biswas, K. Das, B. Sardar and D. Srimani, *Dalton Trans.*, 2019, **48**, 6501–6512.
- 188 B. Qian, S. Guo, J. Shao, Q. Zhu, L. Yang, C. Xia and H. Huang, *J. Am. Chem. Soc.*, 2010, **132**, 3650–3651.
- 189 B. Qian, S. Guo, C. Xia and H. Huang, *Adv. Synth. Catal.*, 2010, **352**, 3195–3200.
- 190 P. Zhang, D. Huang and T. R. Newhouse, *J. Am. Chem. Soc.*, 2019, **142**, 1757–1762.
- 191 C. Dong, Y. Wang, S. Chen, Y. Zhang, C. Wang, J. Li and Y. Zhou, *Eur. J. Org. Chem.*, 2023, e202300194.
- 192 C. Dong, Y. Wang, S. Chen, Y. Zhang, C. Wang, J. Li and Y. Zhou, *Eur. J. Org. Chem.*, 2023, **26**, e202300194.
- 193 N. Tewari, R. C. Mishra, V. K. Tiwari and R. P. Tripathi, *Synlett*, 2002, **2002**, 1779–1782.
- 194 N. S. Reddy, K. Ravinder, P. Krishnaiah and Y. Venkateswarlu, *Synlett*, 2001, **2001**, 0625–0626.
- 195 S. K. Boddu, N. Ur Rehman, T. K. Mohanta, A. Majhi, S. K. Avula and A. Al-Harrasi, *Green Chem. Lett. Rev.*, 2022, **15**, 765–795.



Review

- 196 A. R. Katritzky, C. A. Ramsden, J. A. Joule and V. V. Zhdankin, *Handbook of Heterocyclic Chemistry*, Elsevier, 2010.
- 197 L. Savegnago, A. I. Vieira, N. Seus, B. S. Goldani, M. R. Castro, E. J. Lenardão and D. Alves, *Tetrahedron Lett.*, 2013, **54**, 40–44.
- 198 S. M. Prajapati, K. D. Patel, R. H. Vekariya, S. N. Panchal and H. D. Patel, *RSC Adv.*, 2014, **4**, 24463–24476.
- 199 A. S. Reis, M. Pinz, L. F. B. Duarte, J. A. Roehrs, D. Alves, C. Luchese and E. A. Wilhelm, *J. Psychiatr. Res.*, 2017, **84**, 191–199.
- 200 A. Upadhyay, P. Kushwaha, S. Gupta, R. P. Dodda, K. Ramalingam, R. Kant, N. Goyal and K. V. Sashidhara, *Eur. J. Med. Chem.*, 2018, **154**, 172–181.
- 201 Y.-M. Cai, X. Zhang, C. An, Y.-F. Yang, W. Liu, W.-X. Gao, X.-B. Huang, Y.-B. Zhou, M.-C. Liu and H.-Y. Wu, *Org. Chem. Front.*, 2019, **6**, 1481–1484.
- 202 A. Kumar, S. S. Shinde, D. K. Tiwari, B. Sridhar and P. R. Likhari, *RSC Adv.*, 2016, **6**, 43638–43647.
- 203 Z.-Q. Liu, D. Zhu, B.-L. Luo, N. Zhang, Q. Liu, Y. Hu, R.-B. Pi, P. Huang and S. Wen, *Org. Lett.*, 2014, **16**, 5600–5603.
- 204 T. Hirayama, S. Ueda, T. Okada, N. Tsurue, K. Okuda and H. Nagasawa, *Chem.-Eur. J.*, 2014, **20**, 4156–4162.
- 205 G. P. da Costa, M. F. Bach, M. C. de Moraes, T. Barcellos, E. J. Lenardão, M. S. Silva and D. Alves, *Adv. Synth. Catal.*, 2020, **362**, 5044–5055.

