

Recent advances in the synthesis of quinolines: a review

Cite this: *RSC Adv.*, 2014, 4, 24463

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Quinolines have become important compounds because of their variety of applications in medicinal, synthetic organic chemistry as well as in the field of industrial chemistry. In recent years there are greater societal expectations that chemists should produce greener and more sustainable chemical processes. This review article gives information about the green and clean syntheses using alternative reaction methods for the synthesis of quinoline derivatives. The article includes synthesis by microwave, using clay or some other catalyst which could be recycled and reused, one-pot reaction, solvent-free reaction conditions, using ionic liquids, ultrasound promoted synthesis and photocatalytic synthesis (UV radiation).

Received 1st March 2014
Accepted 17th April 2014

DOI: 10.1039/c4ra01814a

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Introduction

Quinoline is one of the most important N-based heterocyclic aromatic compounds. Quinolines recently have been caught the attention of researchers because of their broad range of activities and of course for their wide applications too.

The main sources of quinoline include petroleum, coal processing, wood preservation and shale oil. The quinoline derivatives occur in various natural products, especially in alkaloids. In 1820, quinine (**1**) was isolated from the bark of the cinchona tree which replaced the use of crude bark for the treatment of malaria. Other quinoline derivatives (**2–9**) having

various activities were also isolated from different plant species.¹ (Fig. 1).

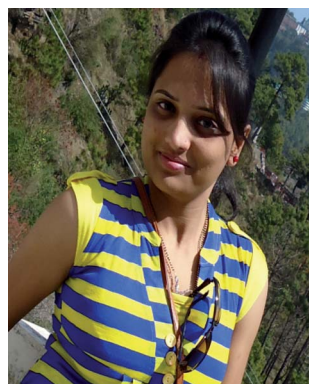
Quinoline was first extracted from coal tar in 1834 by Friedlieb Ferdinand Runge. Coal tar remains the principal source of commercial quinoline.²

Activities and applications

Quinoline derivatives in general are known to have a broad range of applications in medicinal, bioorganic, and industrial chemistry as well as in the field of synthetic organic chemistry.

Their derivatives have been found to possess various biological activities like anti-malarial, anti-bacterial, anti-fungal, anti-asthmatic, antihypertensive, anti-inflammatory, and anti-platelet activity.³ They also exhibit anti-tubercular⁴ and immune depressing⁴ activities. There are a few promising compounds

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with the quinoline ring system, like pamaquine (**10**), chloroquine (**11**), tafenoquine (**12**), bulaquine (**13**), quinine (**14**) and mefloquine (**16**) as antimalarial agents, and amodiaquine (**15**) as an antimalarial and anti-inflammatory agent (Fig. 2).^{5–7}

The 2-arylquinoline derivatives (**17a**) and (**17b**) show selectivity in binding to the estrogen receptor β (ER β), which plays an important role in the development, maintenance, and function of the mammalian reproductive system, as well as in non-sexual tissues.⁸

4-[2-(Diethylamino) ethylamino] quinolin-7-ol (**18**) containing nitrogen at position 4 exhibits antiplasmodial activity.⁹ Polysubstituted quinoline derivatives such as 8-hydroxyquinoline and quinoline-8-thiol have been used to produce metal complexes which emit light.¹⁰ 2-(4-Bromo-5-ethynylthiophen-2-yl)-6-ethynyl-4-phenylquinoline (**19**) has been applied in sensors and light emitting diodes.¹¹



Rajesh H. Vekariya was born in Ahmedabad (Gujarat) India in 1989. He obtained his BSc degree in Chemistry in 2010 and his MSc degree in organic chemistry in 2012 both from the Gujarat University, Ahmedabad (India). Currently, he is pursuing his PhD under the guidance of Dr Hitesh D. Patel at the Chemistry Department of the Gujarat University, Ahmedabad. His research focuses on synthesis of organic intermediates via green chemistry and their utilization in the synthesis of biologically active compounds. Mainly, he is working on highly important and basic industrial process such as nitration, bromination and oxidation reactions.

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Some quinoline derivatives such as quinaldine (**20**) and quinaldic acid (**21**) show activity as corrosion inhibitors for mild steel in hydrochloric acid.⁴ (Fig. 3).

In recent years some quinoline derivatives (**22**) shown below, which have explored the structural modification of Sorafenib, where synthesized as novel Raf kinase inhibitors with more potent and selective antitumor activities.¹²



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Aziridine-isoquinoline hybrids and their ring-opening products shown in Fig. 4 exhibit potent antiplasmodial activity.¹³

A new series of hybrid conjugates of *N*-(7-chloroquinolin-4-yl) piperazine-1-carbothioamide and 1,3,5-triazine derivatives have considerable antimalarial activity against both wild and mutant parasites with marked variations on changing the pattern of substitution. Such derivatives also show excellent antibacterial activity against several Gram-positive and Gram-negative microorganisms.^{14,15}



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4-Arylquinoline-2-carboxylate derivatives show antiprotozoal activity against the pathogenic parasite *Toxoplasma gondii*.¹⁶



Dr Hitesh D Patel was born in 1973 and received his PhD in Organic Chemistry from the South Gujarat University, Surat, Gujarat, India in 2005 under the supervision of Dr B. D. Mistry of B K. M. Science College, Valsad, Gujarat, India. He is working on synthesis of biologically active heterocyclic compounds, their characterization by various analytical techniques and their testing for activities. He joined as

a lecturer at St. Xavier's College, Ahmedabad Gujarat, India and had been associated with Xavier Research Foundation, Ahmedabad, India, during the period of 1998–2005. Since then he joined as an Associate Professor at Department of Chemistry, Gujarat University, Ahmedabad, Gujarat, India.



Fig. 1 Quinoline derivatives isolated as natural products.



Many quinoline derivatives are found to have applications as agrochemicals¹⁷ as well as use in the study of bio-organic and bio-organometallic³ processes. They are also used in manufacturing dyes, food colorants, pH indicators and other organic compounds. In addition to this they have also been used as ligands for the preparation of OLED phosphorescent complexes¹⁸ and with conjugated polymers used as a selective chemo-sensors of the fluoride and metal ions.^{19,20}

Due to such a wide range of applicability, there has been increasing interest in the development of efficient methodologies for the synthesis of quinoline derivatives.

Conventional methods of synthesis

A number of preparations have been known since the late 1800s for the synthesis of quinoline and its derivatives (Fig. 5).

The structural core of quinoline has been generally synthesized by various conventional named reactions such as Skraup, Doebner-von Miller, Friedlander, Pfitzinger, Conrad-Limpach, Combes synthesis.¹

Though many of these methods are very effective, they often involve the use of various acids or reagents that are not environmentally compatible, produce a large amount of waste and require longer reaction times.¹⁷ Moreover, many of these methods give relatively large amounts of undesirable by-products whose removal is tedious and often wasteful and are also not satisfactory with respect to operational simplicity to isolate the yield. Thus, it has become very important to follow methods which could be considered as a better and eco-friendly viable 'green synthetic methods'.

This green chemistry approach provides a way to design products in a simplified manner and has less feedstock,



Fig. 2 A few promising compounds with the quinoline ring system.



Fig. 3 Some important quinoline derivatives.

minimum waste, low energy consumption, less hazardous, renewable materials, high atom economy, reduce reaction steps and green catalysts that improve the efficiency of the reaction.

Clean chemical synthesis using alternative reaction methods includes:

(1) Alternative reaction media for the synthesis using supercritical fluids, ionic liquids, water, polyethylene glycol and solvent-free or grinding methods.

(2) Alternative energy sources like microwaves (fast and homogeneous heating by microwave irradiation), ultrasound and sunlight/UV.



Fig. 4 Aziridine-isoquinoline hybrids and their ring-opening products.



Fig. 5 Various conventional routes for the synthesis of quinoline derivatives.

(3) Catalysts which may be clay or other green catalysts which could be recycled and reused.

Various reaction schemes

Safari *et al.* described a procedure for preparation of quinaldine derivatives (35) from aniline derivatives (33) and acetaldehyde (34) under microwave irradiation without any solvent

(Scheme 1).²² In this method they tried different Bronsted acids but found hydrochloric acid appeared to be the best catalyst for this reaction, showing the highest yield. Moreover the yield of the product was not affected by the nature of substituent in this reaction.

The method presents a simple and useful synthetic process for quinaldines because of high yields, short reaction time, a straight forward, easy work-up procedure, the use of microwave



Scheme 1

irradiation as the novel efficient source of energy and the use of molecular oxygen as a green oxidant.

A method for the synthesis of quinoline derivatives and bis-quinoline derivatives (**39**) including a microwave-assisted, one-pot-three-component reaction between aromatic amines, (**36**) aromatic aldehydes (**37**) and phenylacetylene (**38**) in the presence of catalytic amounts of potassium dodecatungstocobaltate trihydrate ($\text{K}_5\text{CoW}_{12}\text{O}_{40} \cdot 3\text{H}_2\text{O}$) has been demonstrated by Anvar and his co-workers (Scheme 2).²³ They were first to report the use of polyoxometalates (POMs) as catalysts for the one-pot three-component synthesis of quinolines and bis-quinolines under microwave irradiation.

The catalyst could be easily recovered by simple filtration and could be reused for several cycles without any significant loss of its catalytic activity. Moreover no metal was detected in the final product which confirms the green nature of the present method. This makes the method useful and attractive for the synthesis of quinoline derivatives.

A synthesis of steroidal and nonsteroidal quinoline derivatives (**41**) has been established by Gogoi *et al.* (Scheme 3).¹⁷ In this method steroidal quinoline derivatives were synthesized from a one-pot reaction of steroidal β -bromovinyl aldehydes (**40**) and arylamines in high yield using microwave irradiation without the use of a catalyst and in a solvent-free condition. This methodology offers an environment friendly 'green' alternative organic synthesis.

Facile microwave-assisted processes suitable for the preparation of a series of quinoline-4-carboxylic acids (**44**) have been introduced by Zhu and co-workers (Scheme 4).²⁴ In this Pfitzinger type of reaction a condensation reaction between isatins (**42**) and sodium pyruvate to give quinoline-2,4-dicarboxylic acid (QDC) (**43**) is carried out under microwave conditions which optimise reaction solvent, time and temperature. The subsequent decarboxylation reaction of QDCs in water instead of toxic nitrobenzene under MW was also promoted successfully.



Scheme 3

Ranu *et al.* developed a simple and efficient procedure for the synthesis of 4-alkylquinoline derivatives (**46**) by a one-pot reaction of anilines (**36**) with alkyl vinyl ketones (**45**) on the surface of a silica gel inseminated with indium(III) chloride under microwave irradiation without any solvent (Scheme 5).²⁵ The main advantages of this procedure are: operational simplicity, fast reaction, high yield and general applicability. These advantages accommodate a variety of substitution patterns.

Microwave-assisted solid acid-catalyzed syntheses of substituted quinoline derivatives (**48**) have been discussed by Kulkarni and his colleagues (Scheme 6).²⁶ The quinoline derivatives were synthesized by a multicomponent reaction of anilines (**36**), aldehydes (**37**) and terminal aryl alkynes (**49**). The reaction was catalyzed by montmorillonite K-10, a strong and environmentally benign solid acid. The multicomponent approach yields products with nearly 90% atom economy in excellent yields in a matter of minutes. The use of microwave activation reduces the reaction time significantly.

Quinoline derivatives (**51**) were synthesized by employing amino acetophenone (**41**) and phenylacetylene (**50**) in the presence of $\text{Zn}(\text{OTf})_2$ as an effective catalyst under microwave irradiation, as presented by Praveen and co-workers (Scheme 7).²⁷

The advantage of this method is that the reaction is amenable to alkynes containing alkyl, aromatic and hetero-aromatic groups. Good yields, a short reaction time, operational simplicity, low catalyst loading and wide substrate scope are the significant advantages of this reaction from synthetic viewpoint.

Naik *et al.* have reported a rapid and efficient method for the synthesis of various carbonitrile quinoline/benzo[*h*]quinoline derivatives (**55**) by utilizing benzaldehyde (**52**), methyl cyanoacetate (**53**) and aromatic amine (**54**) with nanostructured TiO_2 photocatalysts under solvent-free conditions under microwave irradiation (Scheme 8).²⁸

3-Unsubstituted 4-hydroxyquinolin-2(1*H*)-one (**57**) was synthesized using substituted aromatic amine (**36**) and malonic



Scheme 2



Scheme 4



Scheme 5



Scheme 6



Scheme 7

acid (**56**) under microwave irradiation in dimethylformamide,²⁹ without employing any solvent³⁰ and using polyphosphoric acid (PPA)³¹ (Scheme 9). Operational simplicity and high yield in significantly very short reaction time make this procedure a useful and attractive alternative to the currently available methods.

Reddy *et al.* introduced a three-component one-pot reaction between 3,4-dimethoxyaniline (**58**), aldehydes and ethyl-3,3-

diethoxypropionate (**59**) to a quinoline derivative (**60**) by using montmorillonite K-10 (Mont K-10) as a green catalyst by utilizing the oxygen of air and water (Scheme 10).²¹ Montmorillonite K-10 (Mont K-10) was found to be more effective compared to other Lewis acids as the expected product was isolated in good yield. Moreover montmorillonite K-10 was recovered by simple filtration and reused easily, while the use of water as a solvent makes this reaction eco-friendly.

Montmorillonite K10 clay-catalyzed synthesis of quinoline derivative (**63**) has been disclosed by Nagendrappa *et al.* by employing aniline derivatives (**61**) and cinnamaldehyde (**62**) (Scheme 11).³² The mechanism follows a domino process involving cyclization followed by dehydration and then after oxidation delivers quinolines. The reaction was carried out under solvent-free conditions and with the assistance of microwave irradiation.

A new, convenient, efficient and eco-friendly protocol for the synthesis of a polysubstituted quinoline derivative (**66**) under mild condition has been described by Jida *et al.* (Scheme 12).³³ A wide variety of new products were readily prepared in the presence of propylphosphonic anhydride (T₃P) in short reaction times and in excellent yields. Here T₃P is used as a mild water scavenger catalyst in this coupling reaction. In addition, this non-toxic T₃P offers several advantages over traditional reagents, such as low toxicity, commercial scale availability, low price, low epimerization tendency, high selectivity yields, excellent purity, a broad functional group, and easy work-up



Scheme 9



Scheme 8



Scheme 10



Scheme 11



Scheme 12

procedures. The simplicity and cost-effectiveness of this methodology are attractive for large scale synthesis.

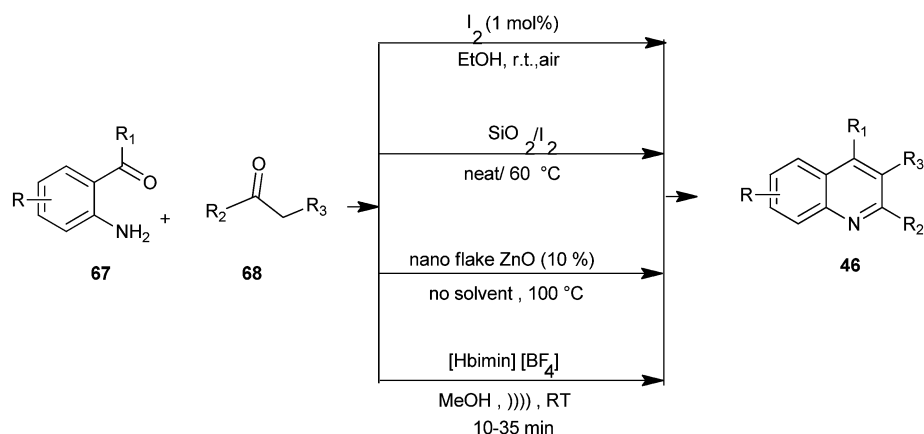
Mild and efficient routes for the synthesis of quinoline derivatives and polycyclic quinoline derivatives (**46**) by utilizing substituted *o*-amino acetophenone derivative (**67**) and enolisable ketone (**68**) with molecular iodine as a catalyst in ethanol,³⁴ combining iodine and silica gel under solvent-free conditions,³⁵

a Friedlander heteroannulation method by using nano ZnO as a mild, non-volatile, non-corrosive and efficient catalyst which provides regiospecific synthesis under solvent-free conditions,³⁶ and using ionic liquid [Hbim][BF₄] under ultrasound at room temperature.³⁷ These methods avoid the use of hazardous acids or bases and harsh reaction conditions. The advantages of these methods include good substrate generality, the use of inexpensive reagents and catalysts under mild conditions, and experimental operational ease (Scheme 13).

Gibson *et al.* used near critical water (NCW) as a medium for the organic synthesis of nitrogen heterocyclic compounds. The potential for solvent recycling was also demonstrated. In this method carbonyl compound with isatin (**69**) in NCW form the substituted quinoline derivative (**70–71**) *via in situ* decarboxylation (Scheme 14).³⁸ Hot pressurizing NCW to make it more ionized makes it a very good dehydrating media suited for isatin opening, condensation and cyclodehydration with carbonyls.

A highly efficient, cost-effective and environmentally benign protocol for the synthesis of 2,4,6-trisubstituted quinoline derivative (**73**) by using NaHSO₄·SiO₂ as a heterogeneous and reusable catalyst has been disclosed by Vu and co-workers (Scheme 15).⁸ This procedure is operationally simple and can be an alternative to the existing protocols for the synthesis of trisubstituted quinoline derivatives. From a practical point of view, the catalyst is easier to prepare from readily available reagents and like other heterogeneous catalysts, it can also be reused.

A microwave-assisted procedure for quinoline derivative (**76**) has been reported by Shekouhy and co-workers (Scheme 16).³⁹ The reaction between 2-aminoaryl ketones (**74**) and carbonyl



Scheme 13



Scheme 14



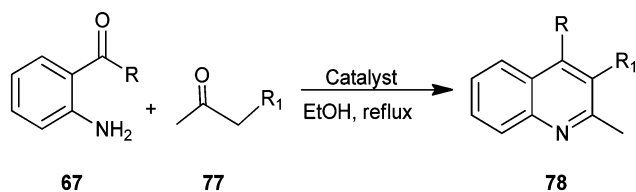
Scheme 15



Scheme 16

compounds (75) in the presence of silica nano-particles (NPs) as catalysts under microwave irradiation give high yields of quinoline derivatives. Silica nano-particles gave best results compare to CaO, MgO, Al₂O₃ and SiO₂. Silica NPs, a highly microporous solid, offer a wide range of active sites and often can be regenerated if deactivated during the reaction. On the other hand, reducing catalytic substances to nanometers in size greatly increases the surface area available per gram and ultimately the catalytic activity. The NPs catalyst can be reused without loss of activity even after recycling fourteen times.

A one-pot, mild, efficient, and environmentally benign protocol has been developed by Chermahini *et al.* for the synthesis of a quinoline derivative (78) catalyzed by Montmorillonite K-10, zeolite, nano-crystalline sulfated zirconia (nano-crystalline SZ) in high yields (Scheme 17).⁴⁰ The mild reaction conditions, easy work-up, clean reaction profiles, lower catalyst loading and cost efficiency make this approach an interesting alternative to the existing methods.

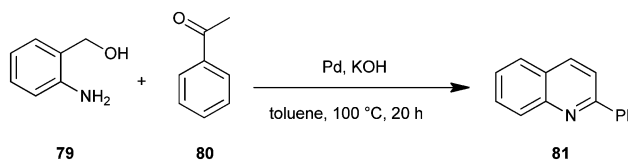


Scheme 17

Ren *et al.* have disclosed a method in which 2-aminobenzyl alcohol (79) reacts with ketones (80) in toluene or polyethylene glycol (PEG-2000) by employing a palladium catalyst along with KOH to isolate the corresponding quinoline derivative (81) in good yields (Scheme 18).⁴¹ The catalytic system could be recovered and reused five times without any loss of catalytic activity.

The use of some new metal dodecyl sulfates as a catalyst in the synthesis of quinoline derivatives (83) by a Friedlander protocol in aqueous media have been developed by Salehi *et al.* (Scheme 19).⁴² In this synthesis metal dodecyl sulfates were used in reaction between *o*-aminoaryl ketones (82) and ketones or β -diketones (72). The authors prepared some combined Lewis acid-surfactant catalysts (LASC) and tried to use them for the synthesis. They found zirconium tetrakis(dodecyl) sulfate Zr(DS)₄ superior to all other prepared metal dodecyl sulfates. Zr(DS)₄ was recovered easily through separation by a centrifuge and reused in aqueous media.

An air-mediated metal-free one-pot and three-component green synthetic method for obtaining a quinoline derivative (86)



Scheme 18



Scheme 19



Scheme 20



Scheme 21



Scheme 22

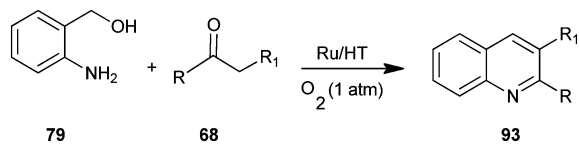
by utilizing 4-substituted aniline (**54**), aldehyde derivative (**84**), ethynylbenzene (**85**) with molecular iodine as a catalyst has been reported by Li *et al.* (Scheme 20).⁴³ Here molecular iodine acts as a mild Lewis acid.

The synthesis of quinoline derivatives and polycyclic quinoline derivatives (**88–89**) *via* a Friedlander condensation between 2-aminoarylketones (**87**) with enolisable ketones by using the recyclable heterogeneous catalyst sulfamic acid have been described by Rao and colleagues (Scheme 21).⁴⁴ The ease of recovery and reuse of this novel catalyst makes this method quite simple, convenient and environmentally benign

for the synthesis of highly functionalized quinoline derivatives.

The one-pot synthesis of 2- or 3-mono-substituted, 2,3-disubstituted quinoline derivatives and also other bi- or tricyclic quinoline derivatives has been presented by Li *et al.* (Scheme 22).⁴⁵ The reaction was applicable to a wide variety of ketones and aldehydes. The biggest advantage of this method is use of inexpensive, readily available reagents and solvents that do not require oxygen or moisture-free conditions.

Motokura *et al.* have presented a one-pot process for the synthesis of quinoline derivative (**93**) by using (2-aminophenyl)



Scheme 23



Scheme 24



Scheme 25

methanol (**79**) and enolisable ketone (**68**) with Ru-grafted hydrotalcite (Ru/HT), a multifunctional heterogeneous catalyst (Scheme 23).⁴⁶ The presented catalytic system improves on previously reported catalytic systems through advantages like high catalytic activity, wide applicability to various carbonyl compounds, no need for homogeneous bases and the use of molecular oxygen as a green oxidant. This is the first reported one-pot quinoline synthesis using heterogeneous catalysts.

A one-pot, solvent free, microwave-assisted, multi-component reaction for the synthesis of a quinoline derivative (**95**) without any solvent by utilizing substituted aniline (**94**), acetone and benzaldehyde (**52**) on the surface of alumina impregnated with hydrochloric acid has been developed by Mirza *et al.* (Scheme 24).⁴⁷ Hydrochloric acid appeared to be the best catalyst for this reaction, showing the highest yield and was preferred compared to sulfuric acid because of its higher safety, environmental friendliness and lower price. The advantages like high yield, short reaction time, straight forward and easy

work-up procedure makes the method completely eco-friendly, fast, simple, and highly efficient.

An efficient microwave assisted, one-pot, solvent-free synthesis of substituted quinoline derivative (**96**) from *o*-nitrobenzaldehyde (**90**) and enolizable ketones (**65**) using $SnCl_2 \cdot 2H_2O$ as the reductant has been introduced by Chaudhuri and co-workers (Scheme 25).⁴⁸ This method is relatively faster and it affords the desired products in respectable yields.

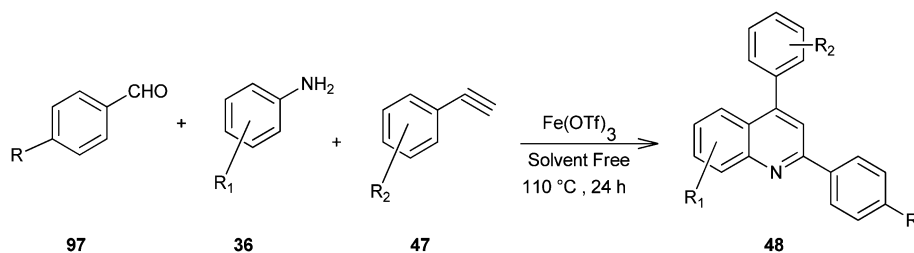
Yao *et al.* have described a novel application of highly stable $Fe(OTf)_3$ as an efficient catalyst for carbon-carbon bond formation *via* the activation of a terminal alkyne C-H bond under solvent-free conditions for the synthesis of a quinoline derivative (**48**) (Scheme 26).⁴⁹ Furthermore, the catalyst was easily recovered from the reaction mixture and reused many times with only a little loss of activity. Other catalysts such as $FeCl_3$, $Fe(OTf)_2$, $FeSO_4$, $Fe(NO_3)_3$ and $Fe_2(SO_4)_3$ were examined in this reaction, but they did not behave as efficiently as $Fe(OTf)_3$ in terms of yield of the products. It was also noted that the reaction in solvent-free conditions was more favorable compared to reactions with various organic solvents.

The synthesis of quinoline-2-carboxylate derivative (**98**) by $Cu(OTf)_2$ catalyst *via* the intermolecular addition of alkyne (**50**) onto imines (**99**) and subsequent intermolecular ring closure by arylation have been described by Huang *et al.* (Scheme 27).⁵⁰ In addition, various catalyst such as $Cu(OAc)_2$, $Cu(acac)_2$, $Cu(tmhd)_2$, CuI , $Cu(OTf)_2$ were examined by authors in this reaction with different organic solvents like DCM, 1,4-dioxane, toluene, furan, DMSO and DMF. However the best result was obtained when the reaction was carried out using $Cu(OTf)_2$ in DCM. The efficiency of this system allowed the reactions to be carried out at room temperature.

A method for producing a quinoline derivative (**100**) using ultrasound and an ionic liquid has been established by Kowsari *et al.* a two-component, one-pot, condensation reaction of isatin (**69**) with enolisable ketone (**68**) yields the quinoline derivative



Scheme 27



Scheme 26



Scheme 28



Scheme 30



Scheme 29



Scheme 31

(**100**) (Scheme 28).⁵¹ Authors examined the various solvents' effect on the reaction. The results showed that the reaction is favored in water. Also, the ionic liquid could be recovered and reused many times without loss of its activity. In addition, a simple process, high selectivity, short reaction time, use of cheap and environmentally benign solvent and the reusability of the aqueous media are the superior advantages of this protocol.

Austin *et al.* have described the photosynthesis of a substituted annulated quinoline derivative (**102–104**) from (**101**) and (**103**) (Scheme 29).⁵² This photocyclization–elimination process provides a convenient route for the synthesis of a variety of substituted 2,3-dihydro-1*H*-cyclopenta[*b*] quinoline derivatives from readily accessible precursors. Reactions of *meta*-substituted precursors were highly regioselective, with alkyl substituents, which afforded 5-substituted 2,3-dihydro-1*H*-cyclopenta[*b*]quinolines. In addition, substrates containing powerful electron-donating substituents generally afford 7-substituted products in high yields.

The photochemical cyclization method for the synthesis of 2,4-diiodoquinoline derivative (**105**) by the cyclization of *o*-alkynylaryl isocyanide (**106**) with iodine has been developed by Mitamura and co-workers (Scheme 30).⁵³ Which are difficult to synthesize by using other existing methods. For a comparative study authors employed various organic solvents such as chloroform, methanol, acetone, acetonitrile, ethyl acetate,

tetrahydrofuran, toluene, *n*-hexane and triethyl amine. However the best results were obtained in terms of the yield of the products when chloroform was used as a solvent. This photochemical reaction was also carried out with various cyclizing agents such as Br₂, *N*-chlorosuccinimide (NCS), *N*-bromosuccinimide (NBS) and *N*-iodosuccinimide (NIS) but they were not found to be efficient catalysts for this protocol. This eco-friendly green reaction proceeds under mild conditions and readily affords the appropriate products in high to excellent yields.

A one-pot synthesis of 4-ethoxy-1,2,3,4-tetrahydroquinoline (**109**) from a heterogeneous solution of nitroarene (**107**), ethanol and TiO₂ upon irradiation of UV light has been reported by Joo and colleagues (Scheme 31).⁵⁴ In addition, substrates have either oxygen or amino substituents like *m*-nitroanisole *etc.*, in such cases the reaction proceeds rather slowly compared to those having alkyl substituents under the same reaction conditions. An added advantage of this protocol is that quinoline derivatives were directly synthesized from the nitro arenes instead of aminoarenes using environmentally friendly conditions.

Selvam *et al.* have demonstrated the photo catalytic conversion of nitrobenzene (**110**) to 2-methylquinoline derivative (**113**) in absolute ethanol using TiO₂ as a photo catalyst. In this reaction excited electrons are trapped by the metal which is used to dope TiO₂; this enhances the charge separation between hole and electron. Holes cause the oxidation of alcohol (**112**) to aldehyde (**111**) and the trapped electrons are consumed for the reduction of nitrobenzene to aniline by H⁺ formed during the oxidation of alcohol. The further reaction of aniline and aldehyde gives quinaldine. This photo reaction is only possible in the presence of TiO₂. In 2009 Selvam *et al.* prepared substituted quinoline derivatives and other heterocycles by TiO₂ nanoparticles *via* this photocatalytic process.⁵⁵ In 2010 they synthesized quinaldines Au-loaded TiO₂.⁵⁶ In 2011 they synthesized nanosized silver particles loaded TiO₂ to get materials with enhanced adsorption and photocatalytic performance.⁵⁷ In 2012 N-doped TiO₂ using a new nitrogen precursor hydrazine hydrate has been synthesized by a simple wet method by Selvam *et al.* (Scheme 32). They concluded that N-TiO₂ is more efficient than

Scheme 32 Photosynthesis using photocatalyst TiO_2

other metal-doped catalysts in quinaldine synthesis under visible light. Therefore, this process has the potential to enable a more sustainable quinaldine synthesis from nitrobenzene in UV and visible light.⁵⁵

Conclusion

Since quinoline and its derivatives possess a wide spectrum of pharmacological activities and are also utilized as ligands in various biologically-modelled transition metal complexes, a number of methods have been developed from time to time for their synthesis by microwave-assisted, ultrasound-promoted, or heterogeneous acid-catalyzed methods, in UV light or solvent-free conditions and many more. We have made here efforts to compile most of these methods that have been reported in the literature. This review will be very useful to the researcher working in this field, and it would help them to develop a new eco-friendly, efficient and economical method. This is necessary from today's point of view as we need an environmentally clean protocol for the large scale production of such an important biological moiety, which may be used further in many reactions to develop a potent pharmacophore for the future.

Acknowledgements

The authors are thankful to the Department of Chemistry, Gujarat University, Ahmedabad, for providing the necessary facilities and also thankful to UGC-Info net & INFLIBNET Gujarat University for providing e-source facilities.

References

- 1 V. V. Kouznetsov, L. Y. Mendez and C. M. Gomez, *Curr. Org. Chem.*, 2005, **9**, 141–161.
- 2 R. Heusch and B. Leverkusen, *Ullmann's Encyclopedia of Industrial Chemistry*, 2000, DOI: 10.14356007: a09_297.
- 3 U. Desai, S. Mitragotri, T. Thopate, D. Pore and P. Wadgaonkar, *ARKIVOC*, 2006, 198–204.
- 4 E. E. Ebenso, M. M. Kabanda, T. Arslan, M. Saracoglu, F. Kandemirli, L. C. Murulana, A. K. Singh, S. K. Shukla, B. Hammouti and K. Khaled, *Int. J. Electrochem. Sci.*, 2012, **7**, 5643–5676.
- 5 S. Bawa, S. Kumar, S. Drabu and R. Kumar, *J. Pharm. BioAllied Sci.*, 2010, **2**, 64–71.
- 6 M. ozyanik, S. Demirci, H. Bektas, N. Demirbas, A. Demirbas and S. A. Karaoglu, *Turk. J. Chem.*, 2012, **36**, 233–246.
- 7 P. R. Graves, J. J. Kwiek, P. Fadden, R. Ray, K. Hardeman, A. M. Coley, M. Foley and T. A. Haystead, *Mol. Pharmacol.*, 2002, **62**, 1364–1372.
- 8 A. T. Vu, S. T. Cohn, E. S. Manas, H. A. Harris and R. E. Mewshaw, *Bioorg. Med. Chem. Lett.*, 2005, **15**, 4520–4525.
- 9 C. H. Kaschula, T. J. Egan, R. Hunter, N. Basilico, S. Parapini, D. Taramelli, E. Pasini and D. Monti, *J. Med. Chem.*, 2002, **45**, 3531–3539.
- 10 Y. Tokoro, A. Nagai, K. Kokado and Y. Chujo, *Macromolecules*, 2009, **42**, 2988–2993.
- 11 G. Jégou and S. A. Jenekhe, *Macromolecules*, 2001, **34**, 7926–7928.
- 12 Y. Li, X. Shi, N. Xie, Y. Zhao and S. Li, *MedChemComm*, 2013, **4**, 367.
- 13 S. Vandekerckhove, S. De Moor, D. Segers, C. de Kock, P. J. Smith, K. Chibale, N. De Kimpe and M. D'Hooghe, *MedChemComm*, 2013, **4**, 724.
- 14 H. R. Bhat, U. P. Singh, P. Gahtori, S. K. Ghosh, K. Gogoi, A. Prakash and R. K. Singh, *RSC Adv.*, 2013, **3**, 2942.
- 15 H. R. Bhat, S. K. Gupta and U. P. Singh, *RSC Adv.*, 2012, **2**, 12690.
- 16 J. McNulty, R. Vemula, C. Bordon, R. Yolken and L. Jones-Brando, *Org. Biomol. Chem.*, 2014, **12**, 255–260.
- 17 S. Gogoi, K. Shekarrao, A. Duarah, T. C. Bora and R. C. Boruah, *Steroids*, 2012, **77**, 1438–1445.

- 18 R. Kwong, *J. Am. Chem. Soc.*, 2005, **127**, 1614–1615.
- 19 H. Tong, L. Wang, X. Jing and F. Wang, *Macromolecules*, 2003, **36**, 2584–2586.
- 20 G. E. Tumambac, C. M. Rosencrance and C. Wolf, *Tetrahedron*, 2004, **60**, 11293–11297.
- 21 T. R. Reddy, L. S. Reddy, G. R. Reddy, K. Yarbaga, Y. Lingappa, D. Rambabu, G. R. Krishna, C. M. Reddy, K. S. Kumar and M. Pal, *Green Chem.*, 2012, **14**, 1870–1872.
- 22 J. Safari, S. H. Banitaba and S. S. Samiei, *J. Chem. Sci.*, 2009, **121**, 481–484.
- 23 S. Anvar, I. Mohammadpoor-Baltork, S. Tangestaninejad, M. Moghadam, V. Mirkhani, A. R. Khosropour and R. Kia, *RSC Adv.*, 2012, **2**, 8713–8720.
- 24 H. Zhu, R. F. Yang, L. H. Yun and J. Li, *Chin. Chem. Lett.*, 2010, **21**, 35–38.
- 25 B. C. Ranu, A. Hajra and U. Jana, *Tetrahedron Lett.*, 2000, **41**, 531–533.
- 26 A. Kulkarni and B. Torok, *Green Chem.*, 2010, **12**, 875–878.
- 27 C. Praveen, P. DheenKumar, D. Muralidharan and P. T. Perumal, *Bioorg. Med. Chem. Lett.*, 2010, **20**, 7292–7296.
- 28 H. R. Prakash Naik, H. S. Bhojya Naik, T. R. Ravikumar Naik, T. Aravinda and D. S. Lamani, *Phosphorus, Sulfur Silicon Relat. Elem.*, 2009, **184**, 2109–2114.
- 29 K. Arya and M. Agarwal, *Bioorg. Med. Chem. Lett.*, 2007, **17**, 86–93.
- 30 J. H. Lange, P. C. Verveer, S. J. Osnabrug and G. M. Visser, *Tetrahedron Lett.*, 2001, **42**, 1367–1369.
- 31 J. Jampilek, R. Musiol, M. Pesko, K. Kralova, M. Vejsova, J. Carroll, A. Coffey, J. Finster, D. Tabak and H. Niedbala, *Molecules*, 2009, **14**, 1145–1159.
- 32 G. Nagendrappa, *Appl. Clay Sci.*, 2011, **53**, 106–138.
- 33 M. Jida and B. Deprez, *New J. Chem.*, 2012, **36**, 869.
- 34 J. Wu, H. G. Xia and K. Gao, *J. Org. Biomol. Chem.*, 2006, **4**, 126–129.
- 35 M. A. Zolfigol, P. Salehi, A. Ghaderi and M. Shiri, *J. Chin. Biochem. Soc.*, 2007, **54**, 267–271.
- 36 M. Hosseini-Sarvari, *J. Iran. Chem. Soc.*, 2011, **8**, 119–128.
- 37 M. R. Heravi, *Ultrason. Sonochem.*, 2009, **16**, 361–366.
- 38 T. A. Bryson, J. M. Gibson, J. J. Stewart, H. Voegtle, A. Tiwari, J. H. Dawson, W. Marley and B. Harmon, *Green Chem.*, 2003, **5**, 177–180.
- 39 A. Hasaninejad, M. Shekouhy and A. Zare, *Catal. Sci. Technol.*, 2012, **2**, 201–214.
- 40 A. Teimouri and A. Najafi Chermahini, *Arabian J. Chem.*, 2011, DOI: 10.1016/j.arabjc.2011.05.018.
- 41 C. S. Cho and W. X. Ren, *J. Organomet. Chem.*, 2007, **692**, 4182–4186.
- 42 M. A. Zolfigol, P. Salehi, A. Ghaderi, M. Shiri and Z. Tanbakouchian, *J. Mol. Catal. A: Chem.*, 2006, **259**, 253–258.
- 43 X. Li, Z. Mao, Y. Wang, W. Chen and X. Lin, *Tetrahedron*, 2011, **67**, 3858–3862.
- 44 J. S. Yadav, P. Purushothama Rao, D. Sreenu, R. S. Rao, V. Naveen Kumar, K. Nagaiah and A. R. Prasad, *Tetrahedron Lett.*, 2005, **46**, 7249–7253.
- 45 A. H. Li, E. Ahmed, X. Chen, M. Cox, A. P. Crew, H. Q. Dong, M. Jin, L. Ma, B. Panicker, K. W. Siu, A. G. Steinig, K. M. Stolz, P. A. Tavares, B. Volk, Q. Weng, D. Werner and M. J. Mulvihill, *Org. Biomol. Chem.*, 2007, **5**, 61–64.
- 46 K. Motokura, T. Mizugaki, K. Ebitani and K. Kaneda, *Tetrahedron Lett.*, 2004, **45**, 6029–6032.
- 47 B. Mirza and S. S. Samiei, *J. Chem. Chem. Eng.*, 2011, **5**, 644–647.
- 48 M. K. Chaudhuri and S. Hussain, *J. Chem. Sci.*, 2006, **118**, 199–202.
- 49 C. Yao, B. Qin, H. Zhang, J. Lu, D. Wang and S. Tu, *RSC Adv.*, 2012, **2**, 3759.
- 50 H. Huang, H. Jiang, K. Chen and H. Liu, *J. Org. Chem.*, 2009, **74**, 5476–5480.
- 51 E. Kowsari and M. Mallakmohammadi, *Ultrason. Sonochem.*, 2011, **18**, 447–454.
- 52 M. Austin, O. J. Egan, R. Tully and A. C. Pratt, *Org. Biomol. Chem.*, 2007, **5**, 3778–3786.
- 53 T. Mitamura and A. Ogawa, *J. Org. Chem.*, 2011, **76**, 1163–1166.
- 54 K. H. Park, H. S. Joo, K. I. Ahn and K. Jun, *Tetrahedron Lett.*, 1995, **36**, 5943–5946.
- 55 K. Selvam and M. Swaminathan, *RSC Adv.*, 2012, **2**, 2848–2855.
- 56 K. Selvam and M. Swaminathan, *Tetrahedron Lett.*, 2010, **51**, 4911–4914.
- 57 K. Selvam and M. Swaminathan, *J. Mol. Catal. A: Chem.*, 2011, **351**, 52–61.