Review

Covalent Organic Frameworks: Promising Materials as Heterogeneous Catalysts for C-C Bond Formations

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Abstract: Covalent organic frameworks (COFs) are defined as highly porous and crystalline polymers, constructed and connected via covalent bonds, extending in two- or three-dimension. Compared with other porous materials such as zeolite and active carbon, the versatile and alternative constituent elements, chemical bonding types and characteristics of ordered skeleton and pore, enable the rising large family of COFs more available to diverse applications including gas separation and storage, optoelectronics, proton conduction, energy storage and in particular, catalysis. As the representative candidate of next-generation catalysis materials, because of their large surface area, accessible and size-tunable open nano-pores, COFs materials are suitable for incorporating external useful active ingredients such as ligands, complexes, even metal nanoparticles deposition and substrate diffusion. These advantages make it capable to catalyze a variety of useful organic reactions such as important C-C bond formations. By appropriate pore-engineering in COFs materials, even enantioselective asymmetric C-C bond formations could be realized with excellent yield and ee value in much shorter reaction time compared with their monomer and oligomer analogues. This review will mainly introduce and discuss the paragon examples of COFs materials for application in C-C bond formation reactions for the organic synthetic purpose.

Keywords: covalent organic frameworks; heterogeneous catalysts; organic synthesis; C-C bond formation

1. Introduction

Nanoporous materials possess extremely large surface areas, ordered pore channel structure, tunable active sites and functional groups [1–20]. These properties rendered them as pivotal nano-platforms for various applications such as adsorption and separation [21–29], energy storage [30–49], proton conduction [47,48,50–52] and catalysis [6,53–61]. In particular, as the workhorse of current synthetic chemistry, catalytic reactions as well as catalysts are faced with the challenge of updating the use of more green, earth-rich and high-efficient materials and advanced assemblies to win the sustainable development. These new-type artificial nano-porous materials have been constantly developed and applied as pivotal catalysts for both fine-chemical and petrochemical industry for several decades [62–66]. After these 70 years exploration, the domain of nanoporous material has been greatly extended from traditional inorganic mesoporous materials zeolites [4,67–69] to organic-inorganic hybrid materials coordination polymer [70–76] and metal-organic-frameworks (MOFs) [2,77–85]. Although a number of highly porous organic
polymers with considerably large surface area have been designed and synthesized such as hyper-crosslinked polymers [86–91], polymers with intrinsic microporosity [92–100], conjugated microporous polymers [101–105], the linking chemistry totally relied on kinetically controlled irreversible coupling reactions. The irreversibility brought about the poor self-healing ability of these porous materials, which led to disordered pore and skeleton structure, and the formation of non-separable oligomers. The presence of structure disorders and oligomers severely limited their applications in catalysis. Organic porous materials which were conformed of long-range order structure and showed good crystallinity were severely desired.

Since the first seminal work by Omar Yaghi in 2005 [106], covalent organic frameworks (COFs) materials have been focused as a hot research field, explored and developed rapidly with great progress [22,23,107–110]. Compared with inorganic zeolites and silica porous materials, COFs materials possess higher porosity, tunable and larger pore size. These properties would facilitate the diffusion of the reactants and the desorption of the products. In this way, higher selectivity and yield can be obtained. Moreover, COFs materials usually own ultrahigh theoretical specific surface area and high chemical stability. These advantages render them very ideal heterogeneous catalyst for organic synthetic transformations. However, COFs materials are not perfect. They have some intrinsic disadvantages. Commonly, COFs materials show poorer crystallinity with more disordered and impure structures in comparison with zeolites and MOFs materials. And it is very difficult to obtain COF single crystal. The poor crystallinity and the difficulty in obtaining single crystalline COF all lead to the difficulty in confirming the accurate COF structure and the limitation in synthetic applications. Although COFs have ultrahigh theoretical specific surface area, it is very difficult to get high experimental specific surface area as MOFs due to the poorer crystallinity. And since the main skeleton structure of COFs materials is conformed of organic unit, the thermal stability could not be as high as inorganic crystalline zeolites materials. The major difference between COFs and other porous organic polymer materials is (i) that the key synthesizing reactions in the preparation of COFs materials are reversible. This endowed COFs forming process with high self-healing ability. And due to the self-healing and error-checking mechanism of the reversible and thermodynamically-controlled dynamic covalent chemistry, COFs material can form definitively long-range ordered crystalline structure [110]. The connection manner of organic building blocks in COFs is atomically precise 2D or 3D [107] extended polymeric structure with sufficient long-range order and retained crystallinity without compromising porosity; (ii) since the elements constitute COFs materials are all light elements (C, H, B, N, O, Si), the density of COFs are usually lower than other ordered micro- and mesoporous frameworks such as MOFs. Owing to the relative heavy metal ions as well their powerful hydrolysis nature, most MOFs materials lose their ordered structure and diffraction peaks in X-ray diffraction characterization experiments simply after contacting aqueous condition or even humid air. However, compared with its MOFs counterpart, many kinds of COFs display excellent stability in water, organic solvents and even extremely basic, acidic, reductive and oxidative condition [111–117]. Even long-time immersing in these harsh conditions does not deteriorate its long-ordered structure and crystallinity. This extraordinary stability is mainly attributed to that the skeleton structure was metal-free and constructed from strong covalent bonds, which possess much greater bond-dissociation-energy than coordinative bond in MOFs structure; (iii) moreover, the hydrogen bonding and π-π stacking interaction in COFs further strengthen the COFs skeleton and pore structure and prevent them from destruction by collective solvation, hydrolysis and redox presses in most catalytic cases; (iv) finally, and more important, intrinsic COFs materials themselves in fact have seldom active sites and/or catalytic activity as general regular catalysts to trigger C-H or C-X (X = Cl, Br, I, O, S, etc.) transformation for synthetic purpose. Therefore, appropriate active components such as Pd, Ir, Ru, etc. are often incorporated into the special pore or channel of COFs as catalytic active sites or groups [118]. In order to keep either the activity towards substrates and entire selectivity of reactions or COFs structure long-rang order, such an integrating of other active components with COFs pore structure through
binding or coordination has raised a lot of interesting impregnation model and topological structures, which greatly broadens the range of transition-metal catalysis application. Compared with common heterogeneous transition metal-catalyzed organic synthesis, COFs catalysis has displayed great flexibility and freedom to fit the various difficult organic synthesis tasks with high standard and high demand. Despite the fact that there are a few reviews on the application of COFs materials as heterogeneous catalysts in the catalyzed organic synthesis chemistry area. Some important organic reactions including oxidation, reduction, C-C and C-heteroatom coupling reactions have been successfully realized by COFs catalysis for a diversity of synthetic purposes.

According to the organic building block’s extending dimension, COFs are divided into two categories, 2D and 3D COFs. In 2D COFs, the covalent bond connects the organic building block in sheets, while the interlayer connection interaction can be π-π stacking and Van-der-Waals force (see Figure 1a). 3D COFs stretch and extend the framework in space with sp3 tetrahedral carbon or silane building blocks with covalent bond connection in every dimension (see Figure 1b). Due to the interlayer π-π stacking and intralayer rigid aromatic motifs, 2D COFs commonly form columnar structure with linear 1D nano-pores channel penetrating the whole COF materials. The ordered column and direct penetrating open-pores permit the precise assembly of donor-acceptor structure in a highly ordered manner. This property renders 2D COFs much potential applications in functional optoelectronics such as sensing, luminescent, photoconductive, photovoltaic and photocatalytic materials. In comparison, 3D COFs possess highly developed porous structure in space and with their long-range order, e.g., crystallinity. Moreover, 3D COFs enjoy the present record highest surface area (larger than 4000 m2/g) and lowest density (0.17 g/cm3) among all COF materials, which is especially attractive as catalyst candidates.

There appeared several detailed excellent reviews covering COFs materials’ applications in CO2 capture, electrocatalysis, photocatalytic visible-light hydrogen evolution, etc., as one important branch of catalysis, 2D and 3D COFs materials are stepping into the catalyzed organic synthesis chemistry area. Some important organic reactions including oxidation, reduction, C-C and C-heteroatom coupling reactions have been successfully realized by COFs catalysis for a diversity of synthetic purpose. Even very challenging asymmetric organic synthesis have been accomplished by COFs materials modified with enantioselective organocatalysts. Compared with common heterogeneous transition metal-catalyzed organic synthesis, COFs catalysis has displayed great flexibility and freedom to fit the various difficult organic synthesis task with high standard and high demand. Despite the fact that there are a few reviews on the application of COFs materials as heterogeneous catalysts in the organic synthesis, there has remained not yet review focused on selective C-C formation reaction catalyzed by COF materials. In this article, we review recent advances in the utilization of COFs as catalysts to realize the most meaningful organic transformations, namely, C-C coupling reaction.

Figure 1. The structure of a typical 2D COF (a) and 3D COF (b). Hydrogen atoms are omitted for clarity. Carbon, boron, and oxygen atoms are represented as gray, orange, and red spheres, respectively, respectively, polyhedron in orange and triangle in blue, respectively. Reprinted with permission from [106], Copyright 2005, Science and [107] Copyright 2007, Science.
for C-C bond formation reactions. A series of important C-C coupling examples mediated by COFs catalysts are summarized for the synthesis purpose in analogues to Heck, Suzuki-Miyaura, Sonogashira cross-coupling, carbon-centered nucleophilic addition and cross-dehydrogenative couplings (CDC) with an emphasis on understanding the conversion mechanism. Especially we provide an outlook on the current challenges and promising opportunities in this burgeoning COFs application area. We believe that the advanced catalytic transformation strategies based on COFs catalysts will open up a new avenue to supplement the shortages of other transition-metal catalysis in C-C bond synthesis applications.

2. COFs Application as Heterogeneous Catalysis for C-C Bonds Coupling Reactions

Catalytic C-C coupling strategy is always the core of current organic synthesis chemistry field. ~85% synthesis projects in fine chemical production, pharmaceutical synthesis, agricultural chemicals and household chemicals etc. are involved in catalyzed C-C coupling movements. Now the catalytic C-C bond formation reactions often resort to homogeneous transition-metal complexes catalysis system such as the famous Pd-catalyzed Suzuki-Miyaura and Heck cross-coupling reactions [171]. The key of these success systems is that these d-block transition-metal complexes commonly have empty \( \pi^* \) anti-bonding orbital for \( \pi \) back-bonding and electron pair for coordination to sp\(^2\) or sp hybrid carbon-carbon multiple bond [172]. Moreover, these transition-metal centers are very adapted to the oxidative addition and reductive elimination. The low-valent transition-metals are stabilized by a variety of ligands bearing lone-pair electrons and \( \pi^* \) anti-bonding orbit. Thus, it is very facile to choose a number of different chiral ligand to render the transition-metal complex asymmetric activity [173]. These chiral organometallic compounds are often very powerful catalyst for asymmetric C-C coupling reactions. The realization of these asymmetric cross-coupling has greatly enlarged the application scopes of transition-metal catalyzed reactions [174]. Although COFs are very novel materials with short history, the researches on COFs materials are evolving extremely rapid. More and more research articles have revealed that COFs materials possessed many unique properties, significantly differentiating from organic-inorganic hybrid materials such as MOFs [6]. Why chemists favor COFs materials for the specific catalytic C-C coupling reactions? The reasons can be summarized as the following three points. (i) From the prospect of industrial applications, Pd-based homogeneous catalyst has the intrinsic limitation. Especially for drug synthesis, the bio-toxicity of the noble-metal residue has long-time been worried and criticized by the public. And the efficiency of metal separation is unsatisfactory. Moreover, the cost for separation comprises a large portion of the total cost in drug production. Thus, developing heterogeneous catalysis, especially encapsulating the catalytically active species such as Pd, etc. into the pore space of porous materials like COFs with uncompromising catalytic efficiency will solve the issue of separating the toxic metal residues. (ii) From the prospect of catalytic efficiency, which is the core issue in catalysis, after binding metal active species, the metal catalytic center is confined by COFs pore. On one hand, this will increase the difficulty of substrate diffusing and approaching the catalytic sites and the products leaving the catalytic center, which would decrease the TOF (turn-over-frequency). However, on another hand, by the fine-tuning of COFs constituents and pore structures, the COFs will repulse solvent molecules and accelerate substrate adsorption and products desorption. Compared with homogeneous catalyst, this acceleration will greatly increase the efficiency and selectivity. Certainly, realizing this point is very challenging. However, this is just the most attractive property of COFs materials, that is, to search the optimal COFs constituent and pore structure. On one hand, this optimized COFs constituent and pore structure will settle and accommodate metal catalytic center to exert its catalytic ability of oxidative addition and reductive elimination to the maximum. On the other hand, this will fine-tune the selectivity of substrate adsorption and product desorption rate. Most of the following examples in this review are the breakthroughs and proceedings in these prospects. (iii) Compared with other 2D or 3D pore-structure materials, such as the most similar MOFs counterparts, the advantage of COFs in catalyzing C-C coupling reactions is the absence of another external transition-metal center (metal in MOF) to influence
the encapsulated Pd catalysis, which avoids as-induced side-reactions. (iv) As the above-mentioned, the stability to aqueous, acidic, basic and organic solvents renders COFs very inert to be solvated and decomposed. Certainly, due to the short history of COFs materials, its applications and adaptions for C-C coupling reactions require further development and improvement. Even so, the currently reported examples, i.e., the following recommended paragon ones, have already displayed the bright prospects for significant application and development.

We would divide the following examples and discussion according to the C-C coupling reaction types and recommend the following several typical COFs catalyst synthesis and catalytic performance.

2.1. Suzuki-Miyaura Reaction

In 2011, Wang et al. reported the first example of COF material for catalysis application [118]. They synthesized a imine-linked 2D COF material (COF-LZU1) by heating 1,3,5-triformylbenzene and 1,4-diaminobenzene in 1,4-dioxane/aqueous acetic acid solution in anaerobic condition after liquid nitrogen flash-frozen, evacuation and flame-sealing treatment (See Figure 2). The as-synthesized COF materials displayed a two-dimensional layered-sheet structure, with eclipsed nitrogen atoms in adjacent layers a distance of ~3.7 Å. This imine-linked COF-LZU1 material demonstrated high coordination affinity to Pd(OAc)$_2$ due to nitrogen-palladium strong interaction. Only by simple impregnation, Pd(OAc)$_2$ was effectively incorporated into COF-LZU1 channels and pores. This metal incorporation did not greatly eliminate COF material’s long-ordered structure although reduced the intensity of powder X-ray diffraction (PXRD) and Brunauer-Emmett-Teller (BET) surface area in a certain degree. The Pd-incorporated COF material demonstrated much enhanced catalytic activity, shorter reaction time and lower catalyst load than Pd(OAc)$_2$ and its Pd-MOF analogues in typical Suzuki-Miyaura cross-coupling reaction. Moreover, this COF material did not lose its catalytic activity at all even after four cycles reuse. This superior catalytic activity and stability after reutilization rendered it very promising catalyst for classical Suzuki-Miyaura reactions.

In 2015, Jiang et al. reported that a porphyrin-based H$_2$P-Bph-COF material could incorporate Pd(OAc)$_2$ species to efficiently catalyze a Suzuki reaction with excellent yields ranging from 97.1–98.5% [175]. This COF material was synthesized via the condensation reaction between 5,10,15,20-tetra(p-amino-phenyl) porphyrin and 4,4′-biphenyldialdehyde in a EtOH/mesitylene/acetic acid aqueous solution at 120 °C for 3 days in vacuum (See Figure 3). The as-formed COF material was nitrogen-rich due to the porphyrin unit’s tetrapyrole group and imine C=N bonds. These excess nitrogen groups aced as effective docking sites for Pd(OAc)$_2$ complexation. Solid State ($^13$C-NMR (nuclear-magnetic-resonance), FT-IR (Fourier Transform infrared spectroscopy), XPS (X-ray photoelectron spectroscopy) and ICP-AES (Inductively Coupled Plasma-Atomic Emission Spectroscopy) characterizations all confirmed the inclusion of Pd(OAc)$_2$. The porosity and crystallinity decreased in a certain degree after Pd incorporation. This Pd-H$_2$P-COF showed superior catalytic activity for Suzuki cross-coupling reactions for a variety of bromoarenes and phenylboronic acid, forming biphenyl derivatives with yields ranging from 97.1–98.5% surpassing its Pd-MOF and Pd/C counterparts (the yield of 4-methoxybiphenyl product was 65.0%, while the yield for Pd/MOF was 84.1% [176]). This report was inspiring and encouraging since the authors showed that porphyrin-based catalyst could not only promote the radical or carbene-based oxidation, but also accelerate the cross-coupling reaction by incorporating Pd complex in to a COF support.
Figure 2. Schematic representation for the synthesis of COF-LZU1 and Pd/COF-LZU1 materials. (a) Proposed structures of COF-LZU1; (b) and Pd/COF-LZU1; (c,d) possessing regular microporous channels (diameter of ~1.8 nm), simulated with a 2D eclipsed layered-sheet arrangement. C: blue, N: red, and brown spheres represent the incorporated Pd(OAc)$_2$. H atoms are omitted for clarity. (e) Catalytic activity test of Pd/COF-LZU1 in the Suzuki-Miyaura coupling reaction. Reprinted and adapted with permission from [118], Copyright 2011, American Chemical Society.

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Figure 3. The preparation of H$_2$P-Bph-COF and Pd/H$_2$P-Bph-COF. Reprinted with permission from [175], Copyright 2015, Elsevier.
Apart from imine and porphyrin COF materials, a triazine-COF could also serve as efficient amphiphilic support for Pd(0) nanoparticles deposition [177]. The triazine-COF was prepared by condensing 4,4',4''-(1,3,5-triazine-2,4,6-triyl)tris(oxy)tribenzaldehyde and benzene-1,4-diamine in 1,4-dioxane/mesitylene/aqueous acetic acid by heating at 120 °C for 72 h. The as-prepared triazine COF contained both a long and flexible appendage and a nitrogen- and oxygen-rich skeleton. The nitrogen-rich moiety was responsible for the facile in-situ reduction of Pd²⁺ to Pd(0) without any external oxidants. The ether and imine units had strong interaction with Pd(0) nanoparticles by stabilizing and dispersing them on the COF material. This triazine-COF was very effective to catalyze the multi-fold Heck and Suzuki-Miyaura cross-couplings. The catalyst provided unprecedented high TON (turn-over number) and TOF (turn-over frequency) values for multi-fold Heck reactions compared with its homogeneous Pd, Pd-MOF and Pd/C counterparts. Extremely short reaction times 1.5 h and excellent isolated yield up to 99% was obtained. The recyclability test showed that after 3 cycles, the catalyst did not show apparent loss in activity. This report provided viable strategy to dock noble-metal nanoparticles into COF material without external reducing agents applying triazine monomer itself as reductant for efficient Heck and Suzuki-Miyaura cross-coupling reactions by appending long and flexible groups, forming an amphiphilic structure and incorporating more nitrogen and oxygen atoms.

Two different kinds of triazine-based imine and β-ketoenamine linked COFs TAT-DHBD and TAT-TFP could be synthesized from 1,3,5-tris-(4-aminophenyl)triazine (TAT) and 2,5-dihydroxybenzene-1,4-dicarboxaldehyde (DHBD) or 1,3,5-triformylphloroglucinol (TFP) under solvothermal conditions in dioxane/mesitylene mixture [178]. Upon Pd(OAc)₂ inclusion into the pore space and between interlayer region of the 2D sheets, Pd(II)-loaded TAT-DHBD and TAT-TFP COF materials were prepared. Sequentially, by the NaBH₄ reduction of Pd(II) to Pd(0) nanoparticles, the Pd(0)-loaded COFs were synthesized. Four Pd-loaded COF materials illustrated very good catalytic activity in Suzuki-Miyaura cross-coupling reaction between bromobenzene and phenylboronic acid. The best performance was obtained with Pd(0)-TAT-TFP COF catalyst which provided excellent conversion and yield for either electron-rich and electron-deficient bromobenzenes. The most appropriate substrate was 4-cyanobromobenzene with phenylboronic acid which showed almost quantitative formation of 4-cyanobiphenyl only after 4 h by this Pd(0)-TAT-TFP COF catalyst. These four Pd-COF catalysts all displayed very good stability and reusability without apparent leaching of Pd and loss of activity.

Very recently, thioether-containing COF was reported that it could be an excellent support for ultra-fine Pt and Pd nanoparticles providing very narrow size distribution and superior stability [179]. Inspired by their own designed shape-persistent thioether-containing organic cage which hosted ultra-fine Pt and Pd nanoparticles with very narrow size-distribution, the authors elaborately designed and synthesized the PtNPs@COF and PdNPs@COF by condensing a trialdehyde and a thioether-containing diamine in a dioxane/mesitylene/aqueous acetic acid solution at 120 °C for 3 days (See Figure 4). The as-formed Thio-COF material was further complexed with K₂PtCl₄ and K₂PdCl₄ in aqueous solution, and then reduced by a methanolic NaBH₄ solution to PtNPs@COF and PdNPs@COF. From a series of structural, morphological and compositional characterization, the authors demonstrated that ultra-fine Pd and Pt nanoparticles were uniformly incorporated into the pore space of the Thio-COF. The thioether functional group provided strong metal-sulfur interaction to stabilize the ultrafine noble-metal nanoparticles to prevent them from aggregation. Moreover, the long-range ordered pore-channel structure also provided assist for the stabilization of the residing external nanoparticles. The as-formed PtNPs@COF and PdNPs@COF illustrated very good catalytic activity towards 4-nitrophenol reduction and Suzuki-Miyaura cross-coupling between a variety of arylhalides and phenylboronic acid. PdNPs@COF provided excellent NMR yields up to 99% for cross-coupling of 4-methyl-iodobenzene and phenylboronic acid to form 4-methylbiphenyl. Moreover, these two COF-based catalysts showed excellent stability and recyclability after simple centrifugation or natural settling for cycling test and almost no decrease in conversion and yield in catalytic performance after 5th cycle tests, and no noticeable leaching of metal nanoparticles occurred. This example was
very significant because it was the first to use thioether functional group to stabilize the narrowly
distributed ultrafine noble-metal nanoparticles for effective cross-coupling reactions in a COF support.

![Figure 4.](image)

**Figure 4.** (a) Synthesis of Thio-COF and (b) schematic representation of the synthesis of Thio-COF supported PtNPs@COF and PdNPs@COF. Top and side views of the energy-minimized models of Thio-COF (yellow, S; blue, N; grey, C; red, O) are shown in (b). Reprinted with permission from [179]. Copyright 2017, American Chemical Society.

From these successful COFs heterogeneous catalysts for Suzuki-Miyaura cross-couplings, we can
deduce that suitable heteroatoms and their fine-tuned positions, COFs pore geometrical and chemical
properties, and the binding modes of Pd transition-metal center, will all determine whether COFs
catalysts can surpass their homogeneous analogues in catalytic efficiency and selectivity. Optimization
and combination of robust COFs catalysts for efficient Suzuki-Miyaura reactions would still have much
space for further development.

### 2.2. Heck, Sonogashira and Silane-Based Cross-Coupling Reaction

Heck cross-coupling reaction is the Pd-catalyzed coupling of aromatic or vinylic halides with
unsaturated olefinic C=C bond (See Figure 5). The catalytic cycle comprise oxidative addition of
aromatic halides, coordination with olefinic C=C bond, cis-insertion, cis-β-hydride elimination and
reductive elimination. This reaction require Pd(0) active species. The advantages of Heck reaction is
its high regio- and stereoselectivity. And the disadvantage is the costly Pd catalyst. The Sonogashira
cross-coupling is the Pd/Cu co-catalyzed coupling of terminal alkyne with aryl or vinylic halide.
It comprises three basic reactions. The first step is the oxidative addition of aryl halides to Pd(0)
center. Then a Cu-amine complex mediates the transmetallation reaction. The last step is the reductive
elimination, releasing the coupling product and regenerating the Pd(0) catalyst. Cu-amine complex
acts as cocatalyst to assist deprotonating the alkyne substrate.

![Figure 5.](image)

**Figure 5.** Scheme of Pd-catalyzed Heck reaction and its catalytic cycle. Adapted with permission
from [171]. Copyright 2000, ACS.
Banerjee showed that introduction of a large number of nitrogen and oxygen atoms to the skeleton of COF material would reinforce its stability when metal nanoparticles or complexes were deposited. They showed that by the condensation of 1,3,5-triformylphloroglucinol and paraphenylenediamine in mesitylene/dioxane/aqueous acetic acid solution at 120 °C for 3 days in inert atmosphere would generate a nitrogen- and oxygen-rich COF material (See Figure 6) [180]. The as-synthesized imine COF material was further deposited with Pd(II) complexes by immersing in a methanol solution containing Pd(OAc)$_2$. The Pd(0) nanoparticles were generated from in-situ reduction of Pd (II)-COF with NaBH$_4$. The authors demonstrated that Pd(0) and Pd(II) inclusion did not greatly change the crystallinity and flower-like morphology of the COF material. Moreover, the Pd(0)-COF showed superior catalytic activity towards Heck and Sonogashira-type reaction. The Pd(II)-COF demonstrated considerably robust catalytic ability for an intramolecular C-H activation and further C-C coupling reaction synthesizing 9H-carbazole from diphenylaniline. This report manifested its significance in incorporating Pd(0) and Pd(II) into the same COF material and applied the metal-COF composite in highly selective C-C coupling and C-H activation transformations. Moreover, the fact that this COF material demonstrated greatly enhanced stability towards aqueous, acidic and alkaline environment was very inspiring, which illustrated nitrogen and oxygen atoms would considerably increase interactions with metal nanoparticles or metal complexes.

Figure 6. Synthesis of Pd(II) and Pd(0)-doped COFs (i.e., Pd(II)@TpPa-1, Pd(0)@TpPa-1) and summary of their catalytic activity towards Sonogashira, Heck and oxidative biaryl couplings. The doped Pd(0) nanoparticles are probably situated on TpPa-1 surface. (The scheme is to represent the synthesis and the organization of the Pd nanoparticles on COF (TpPa-1) and it is not exactly fit to scale). Reprinted with permission [180], Copyright 2014, Royal Society of Chemistry.

Chai et al. reported that two different nitrogen ligands bipyridine and imine could be incorporated into a single COF skeleton to provide differentiated Pd coordinating sites [181]. They designed and synthesized X% bpy-COF by condensing X% 2,2-bipyridine-5,5′-dicarbaldehyde, 100-X% 4,4′-biphenyl dialdehyde and 4,4′,4′′,4′′′-(pyrene-1,3,6,8-tetrayl)tetraaniline (PyTTA) building blocks in a mesitylene/dioxane/3M acetic acid solution at 120 °C for 3 days (See Figure 7). The as-formed X% bpy-COF contained two different kinds of nitrogen ligands, namely, bipyridine and imine. The further Pd(OAc)$_2$ complexation was furnished by a simple solution-infiltration procedure. From a variety of characterization experiments results, the authors demonstrated that Pd(OAc)$_2$ coordinated with...
both bipyridine and imine unit, but in different region. Pd(OAc)$_2$ combining with bipyridine mainly dwelled in the pore space, while Pd(OAc)$_2$ jointing with imine resided between adjacent layer of the 2D COF. Furthermore, the authors demonstrated that these Pd@bpy-COF displayed very good catalytic performance towards classical Pd-catalyzed Heck reaction between a series of arylhalides and styrene. Pd(II)@75% bpy COF showed the best catalytic ability providing >90% yield after four consecutive runs. The superior activity for Heck reaction of these Pd@bpy-COF catalysts was attributed to the uniform dispersion and the ultra-high loading of Pd(OAc)$_2$.

In other occasions, Pd(0) nanoparticles could be generated in-situ by choosing a predesigned metal-anchored building block. Initially, a 2,2-bipyridine-5,5$'_{-}$diamine palladium chloride(bpy-PdCl$_2$) complex was formed by the coordination reaction between PdCl$_2$ and 2,2$'_{-}$bipyridine-5,5$'_{-}$diamine [182]. Then, via the Schiff-base condensation between Bpy-PdCl$_2$ and 1,3,5-triformylphloroglucinol (Tp), a Pd@Tp-bpy COF was synthesized. This in-situ generated Pd@Tp-bpy COF did not require any external reducing agents for Pd(II) reduction to Pd(0) nanoparticles. The as-formed Pd@Tp-bpy COF showed excellent catalytic performance towards...
a tandem C-C and C-O bond formation reaction between 2-bromophenol and phenylacetylene. The Pd@Tp-bpy COF heterogeneous catalyst promoted tandem cyclization provided various 2-arylbenzofurans in good to excellent yields not regarding electron-rich or electron-deficient substituents on phenylacetylene or 2-bromophenol. Moreover, this keto-enamine and bipyridine anchored Pd@TpBpy COF showed very good stability and recyclability.

A triazine-based COF-SDU1-palladium hybrid could be an active catalyst for the cross-coupling between silanes and aryl iodides [183]. The COF-SDU1 material was synthesized via the imine condensation reaction from tri-(4-formacylphenoxy)-1,3,5-triazine (trif) and p-phenylenediamine in o-dichlorobenzene/n-butanol/6M AcOH heating at 85 °C for 7 days (See Figure 8). The as-obtained COF solid contained two kinds nitrogen ligand azine and imine which are both suitable coordinating sites for Pd(II). After a simple solution-infiltration, mono-dispersing Pd(II) ion was docked in the 2D COF material. This Pd(II)-COF-SDU1 material showed excellent catalytic activity towards a one-pot silane oxidation-cross-coupling reaction. The transformation of phenylsilanes with aryl iodides to biphenyls was effectively catalyzed in methanolic solution. A variety of electron-rich and electron-deficient aryl iodides were cross-coupled with phenylsilanes in good to excellent yields. This COF material also displayed good recyclability and reusability without detectable Pd leaching and loss of activity.

Figure 8. A triazine-based covalent organic framework-palladium hybrid for one-pot silicon-based cross-coupling of silanes and aryl iodides. C: blue, N: orange, O: purple and red spheres represent the incorporated Pd(II). H atoms are omitted for clarity. Reprinted with permission from [183]. Copyright 2015, Royal Society of Chemistry.

The key points of COFs successful application in Heck, Sonogashira cross-couplings are summarized as follows: (i) very low Pd-loading (ii) very high catalytic efficiency (iii) very facile desorption of halide ions.

2.3. Cross-Dehydrogenative-Coupling Reactions

In 2016, a hydrazone-based COF material was reported to be used as an outstanding photocatalyst for a series of cross-dehydrogenative-coupling reaction (CDC) under visible-light illumination [184]. The hydrazone-based TFB-COF was synthesized from the Schiff-base condensation reaction of 2,5-dimethoxyterephthalohydrazide (DMTH) and 1,3,5-triformylbenzene (TFB) under solvothermal conditions (See Figure 9). The as-formed TFB-COF showed good catalytic activity towards a variety of CDC reactions between N-phenyltetrahydroisoquinoline and nucleophilic reagents CH3NO2, CH3COCH3 and PhCOCH3 in O2 atmosphere under visible-light illumination. Good to excellent yields up to 87% were obtained by this hydrazone-based TFB-COF photocatalyst. Moreover, the conversion and selectivity did not show apparent loss after three runs of recycling test. The morphology and XRD
peaks did not display great change. To our best knowledge, this example was the first recorded COF material without transition metal for CDC applications.

Likewise, Liu et al. reported that another hydrazone-based 2D COF material COF-JLU5 could promote the photocatalytic oxidative transformation of N-phenyltetrahydroisoquinoline with a variety of C-centered or P-centered nucleophiles [185]. This 2D COF-JLU5 was synthesized via the Schiff-base condensation reaction of 1,3,5-tris-(4-aminophenyl) triazine and 2,5-dimethoxyterephthaldehyde in the presence of 6 M aqueous acetic acid using mesitylene/1,4-dioxane as the solvent at 120 °C for 3 days (See Figure 10). The as-synthesized COF-JLU5 demonstrated good crystallinity and porosity. Moreover, this hydrazone-based COF showed excellent catalytic performance towards the CDC oxidative transformation of N-phenyltetrahydroisoquinoline with various nucleophiles. The aza-Henry, Mannich-type and C-P bond formation reactions were effectively catalyzed by this COF-JLU5 under visible-light illumination in an O2 atmosphere. Moreover, from the results of the electron-paramagnetic-resonance (EPR) characterization of singlet oxygen ¹⁰O₂ and superoxide anion in different conditions, the authors proposed a plausible mechanism for this COF photocatalytic CDC reaction. Initially, the visible-light activated COF-JLU5 abstracted an electron from N-phenyltetrahydroisoquinoline forming the corresponding radical cation. The radical cation was further transformed to the radical via deprotonation by superoxide anion. The further single-electron-transfer (SET) oxidation of N-phenyltetrahydroisoquinoline radical to its cation was facilitated by superoxide anion or activated-state COF-JLU5. The finally nucleophilic attack between nitroalkane, ketones, malonates and dialkylphosphates with tetrahydroisoquinoline cation yielded the final C-C or C-P bond formation products.
Figure 10. (a) Synthesis of COF-JLU5 by imine condensation reaction. (b) Photocatalytic C-C/C-P coupling reaction of N-phenyltetrahydroisoquinolines with various nucleophiles using COF-JLU5 as photocatalyst. (c) Schematics of plausible mechanism of the COF-JLU5 photocatalyzed CDC reactions. Reprinted and adapted with permission from [185], Copyright 2017, Royal Society of Chemistry.
Despite quite rare examples yet, these COFs catalyzed cross-dehydrogenative-coupling reactions clearly show the most valuable feature of COF materials as C-C coupling catalysts, that is, without transition metal participation, their pore and channel render the formation of carbocation formation and stable enough to allow another nucleophilic reagent attacking effectively. These examples significantly differentiate homogeneous dye-sensitized singlet $O_2$ reaction. The dye molecules cannot provide such a pore structure and catalytic platform to realize the cross-coupling reactions. The unique skeleton and independence of transition-metal catalytic center is the most attractive trait of COFs for cross-coupling reactions.

### 2.4. Chiral Asymmetric C-C Bond Formation Reaction

In 2014, Jiang et al. described an example of a chiral-organocatalytic COF material synthesized through pore surface engineering [186]. This organocatalytic COF material was prepared by integrating a chiral pyrrolidine unit into the main-chain of porphyrin-imine COF material (See Figure 11). They first introduced ethynyl group into the imine moiety. And by the facie alkyne-azide click reaction, they grafted the triazole-substituted pyrrolidine ring to imine by a post-treatment catalyzed by CuI catalyst. The as-prepared COF demonstrated its activity for organocatalysis and displayed a variety of advantages in catalyzing an enantioselective asymmetric Michael-addition reaction. Although moderate diastereoselectivity (d.r.) and $ee$ values were obtained, the conversion of the reactants was much accelerated by this organocatalytic COF material (See Figure 12). And the most important was that this COF material the first time COFs materials realized enantioselectivity control in catalytic organic synthetic reaction.

**Figure 11.** (a) The general strategy for the pore surface engineering of imine-linked COFs via a condensation reaction and click chemistries (the case for $X = 50$ was exemplified). (b) A graphical representation of [Pyr]$_x$-$H_2$P-COF with different densities of catalytic sites on the pore walls (gray: carbon, red: nitrogen, green: oxygen, purple: carbon atoms of the pyrrolidine unit; hydrogen is omitted for clarity). Reprinted with permission from [186], Copyright 2014, Royal Society of Chemistry.

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**Figure 12.** Comparison of the pyrrolidine control, amorphous nonporous polymers, and COFs as catalysts for a Michael addition reaction. Reprinted with permission from [186], Copyright 2014, Royal Society of Chemistry.
In 2015, Jiang et al. reported the design and synthesis of another organocatalytic COF material on *Nature Chemistry* [187]. This was the milestone event for COF materials applied for organic synthetic purpose. The authors discovered that introduction of methoxy group in edge unit would greatly increase the stability of COF material against humidity, acidity and basicity since the methoxy group reinforced the interlayer interaction. The COF material was synthesized by condensing triphenylbenzenetriamine (TPB), 2,5-bis(2-propynoxy)terephthalaldehyde (BPTA) and 2,5-dimethoxyterephthalaldehyde (DMTA) (See Figure 13). The alkynyl-containing [HC≡C]x-TPB-DMTA-COFs were further transformed to [(S)-Py]x-TPB-DMTP-COFs by a post-synthetic click reaction with (S)-2-(azidomethyl)pyrrolidine catalyzed by CuI catalyst. The as-formed [(S)-Py]x-TPB-DMTP-COFs demonstrated extremely strong stability against boiling water, 12 M HCl and 14 M NaOH solution soaking. The COF material displayed very little reduction in crystallinity and porosity after these harsh condition treatments. The XRD diffraction peaks intensities show no apparent decrease and the Brunauer-Emmett-Teller (BET) and Langmuir surface area almost remained unchanged. Besides extraordinary stability and uncompromised crystallinity and porosity, the most significant point of this COF material was its functionality to catalyze chiral asymmetric organic reactions. For a typical organocatalytic asymmetric Michael-addition between unactivated cyclohexanones and nitrostyrenes, this COF material displayed superior catalytic activity compared with its homogeneous counterpart. The COF material required only half the time to complete the transformation with similar excellent ee and d.r. values in contrast with the homogeneous organocatalyst. After 5-cycle reusability test, this COF material did not show any apparent loss in catalytic activity. The shining point of this COF material was its combination of unprecedented stability, good crystallinity, and highly developed mesoporous structure with very powerful catalytic ability, accelerated reaction kinetics, excellent yields, d.r. and ee value for chiral asymmetric Michael addition in aqueous solution (See Figure 13d.)

Apart from above-mentioned chiral organocatalyst incorporated COF materials for asymmetric organic transformations and noble-metal incorporated COF materials for achiral organic transformations, homochiral organocatalytic COF skeleton could also be a perfect support for Pd nanoparticle to catalyze asymmetric reductive Heck reaction which was conventionally catalyzed by organometallic Pd species with chiral ligands [188]. The Pd nanoparticle dispersed in a chiral COF (CCOF) skeleton could effectively be a heterogeneous catalyst for asymmetric Henry and reductive Heck reaction providing excellent isolated yields and ee values. The CCOF was synthesized by a condensing reaction between cyanuric chloride and S-(+)2-methylpipperazine with K2CO3 in a dioxane solution, Furthermore, Pd(0) nanoparticles were included into the CCOF material by in-situ reduction of Pd(NO3)2 methanolic solution with NaBH4 in CCOF aqueous suspending solution (see Figure 14). The as-formed Pd@CCOF was uniformly dispersed between CCOF 2D layer, not residing in the pore space due to PdNPs large size (2–5 nm) compared with the CCOF micropore (1.5 nm). The incorporation of Pd nanoparticles greatly influenced the porous structure of CCOF, enlarging its BET surface area and pore size in certain amount. Furthermore, the authors demonstrated the synergistic catalytic activity by subjecting the Pd@CCOF catalyst in typical Henry and reductive Heck reaction conditions. To the authors delight, the Pd@CCOF catalyst displayed extremely superior catalytic ability towards these two reactions providing excellent yield up to 99% and perfect ee value up to 97%. To our best knowledge, this was the first example of a heterogeneous Pd-catalyzed chiral asymmetric reductive Heck reaction. The Pd@CCOF material illustrated satisfactory recyclability and reusability after 5 cycles reuse almost without apparent loss of catalytic activity providing isolated yield up to 93% and ee value 91% for the 5th cycle test. This report of Pd@CCOF material manifested its importance due to that it was the first COF material combining noble-metal catalyst with chiral organocatalyst in a single COF carrier to fulfill asymmetric transformations previously catalyzed by homogeneous organometallic compounds with complex and elaborately designed chiral ligands.
Figure 13. Synthesis and structure of stable crystalline porous COFs. (a) Synthesis of TPB-DMTP-COF through the condensation of 2,5-dimethoxyterephthalaldehyde (DMTA, blue) and TPB (black). Inset: The structure of the edge units of TPB-DMTP-COF and the resonance effect of the oxygen lone pairs that weaken the polarization of the C=N bonds and soften the interlayer repulsion in the COF. (b) Graphic view of TPB-DMTP-COF (red, O; blue, N; grey, C; hydrogen is omitted for clarity). (c) Synthesis of chiral COFs ([([S]-Py)x]-TPB-DMTP-COFs, x = 0.17, 0.34 and 0.50; blue, DMTA; black, TAPB; red, BPTA; green, ([S]-Py sites) via channel-wall engineering using a three-component condensation followed by a click reaction. (d) Scope of reactants. Different β-nitrostyrene derivatives investigated for the Michael reactions catalyzed with chiral COFs, their products, e.e. yields and d.r. values (red, cyclohexanone; green, newly formed C–C bond; blue, nitrostyrene derivatives). R, substituent H, Cl, Br, Me or OMe. Reprinted with permission from Reference [187], Copyright 2015, Springer.
Recently Cui et al. adopted a multivariate strategy to design and synthesize a family of two-component and three-component 2D CCOF for the purpose of asymmetric catalytic organic synthesis [189]. Three binary and four tertiary 2D CCOFs were prepared via the imine condensation reaction between 2,5-dimethoxy-1,4-terephthalaldehyde (DMTA), 1,3,5-tris(4-aminophenyl) benzene (TPB1) and four L-proline- and L-imidazolidine-based TPB derivatives. The crystalline CCOFs were prepared via two-step condensation and deprotection procedure (see Figure 15). The as-prepared CCOFs demonstrated very high catalytic activity towards three kinds of meaningful asymmetric organic transformations, namely, α-aminooxylation, aldol and Diels-Alder reactions. The CCOF materials provided excellent yields, diastereoselectivity and enantioselectivity. The catalytic performance of the CCOF surpassed the homogeneous chiral proline and imidazolidinone catalyst and other heterogeneous support catalyst such as imidazolidinone-grafted mesoporous silica and proline-MOFs in both diastereoselectivity and enantioselectivity. This report manifested its importance in that it built a toolbox of CCOFs for asymmetric organic transformations by a multivariate strategy.

Cui et al. reported two imine-based chiral TADDOL (2,2-dimethyl-α,α,α′,α′-tetraaryl-1,3-dioxolane-4,5-dimethanol) COF materials CCOF-1 and CCOF-2, which could catalyze the asymmetric addition reaction between Et₂Zn and a variety of aromatic aldehyde providing excellent conversions and ee values [190]. These two CCOFs were synthesized via the imine condensation reaction between tetraaryl-1,3-dioxolane-4,5-dimethanols (TADDOLs) derivatives TTA and TTPA with 4,4′-diaminodiphenylmethane (4,4′-DADPM) (Figure 16). The as-formed TADDOL CCOF-1 and CCOF-2 acted as excellent heterogeneous catalysts for the asymmetric addition reaction between Et₂Zn and aromatic aldehydes after treatment with Ti(OiPr)₄. Up to 99% conversion and >90% ee value was realized when the heterogeneous catalytic reaction was conducted at −30 °C. The CCOF-1/CCOF-2 chiral catalyst also displayed very good recyclability and reusability after consecutive four runs providing comparable conversion and ee value with the fresh one. This example was a milestone discovery in COF application to catalysis due to that it was among the first main-chain chiral COFs to fulfill asymmetric organic transformations.

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Figure 14. (a) Scope of different β-nitrostyrene derivatives with cyclohexanone investigated for the Michael-addition reactions catalyzed with Pd@CCOF (b) Pd NPs-Loaded homochiral COF for heterogeneous asymmetric catalysis. Reprinted and adapted with permission from [188], Copyright 2017, American Chemical Society.
Figure 15. (a) Synthesis of the CCOFs. Blue and red color represents TPB units, and black color represents DMTA units. (b) Multivariate chiral COFs with controlled crystallinity and stability for asymmetric catalysis. Blue represents TPB units, grey represents imine units and yellow represents R and R' units in subfigure a. Reprinted with permission from [189], Copyright 2017, American Chemical Society.
variety of chiral asymmetric organic synthetic reactions, such as V-salen CCOF catalyzed cyanation and diastereoselectivity. Moreover, the good recyclability and reusability was proved in the case of reactions not only provided satisfactory yields, but also realized excellent controls of enantioselectivity by a bimetallic Cr(salen)-Mn(salen)-CCOF heterogeneous catalyst. These M(salen)-CCOF catalyzed Cr-salen CCOF and the tandem one-pot epoxidation and aminolysis opening of alkene catalyzed by Fe-salen CCOF and Mn-salen CCOF, aminolysis opening of epoxides catalyzed by Fe, Mn, Cr, V and Co. The M(salen)-based CCOFs demonstrated very good catalytic activity to a stability. And these two CCOFs were further complexed with a variety of metal ion such as Zn, Fe, Mn, Cr, V and Co. 

Cui et al. reported that two Zn(salen)-based 2D CCOFs were synthesized by condensation of chiral 1,2-diaminocyclohexane and trisalicylaldehyde or tributyltrisalicylaldehyde (Figure 17) [191].

They found that the bulky tributyltrisalicylaldehyde containing CCOF possessed superior stability. And these two CCOFs were further complexed with a variety of metal ion such as Zn, Fe, Mn, Cr, V and Co. The M(salen)-based CCOFs demonstrated very good catalytic activity to a variety of chiral asymmetric organic synthetic reactions, such as V-salen CCOF catalyzed cyanation reaction of aldehydes, Diels-Alder cycloaddition reaction catalyzed by Co-salen CCOF, epoxidation catalyzed by Fe-salen CCOF and Mn-salen CCOF, aminolysis opening of epoxides catalyzed by Cr-salen CCOF and the tandem one-pot epoxidation and aminolysis opening of alkene catalyzed by a bimetallic Cr(salen)-Mn(salen)-CCOF heterogeneous catalyst. These M(salen)-CCOF catalyzed reactions not only provided satisfactory yields, but also realized excellent controls of enantioselectivity and diastereoselectivity. Moreover, the good recyclability and reusability was proved in the case of
V-salen CCOF catalyzed cyanation of aldehydes, which showed almost no loss of enantioselectivity, and conversion after 5 runs of cycle tests.

Two chiral COFs LZU-72 and LZU-76 acted as efficient heterogeneous organocatalyst and effectively catalyzed the asymmetric aldol reaction providing excellent yield and enantioselectivity [192]. The chiral COFs were synthesized by the bottom-up approach from condensing the chiral block units other than a post-synthesis procedure. The authors discovered that the linear-structured 4,4'-(2-(pyrrolidin-2-yl)-1H-benzimidazole-4,7-diyl)dianiline was incorporated into the achiral skeleton. The further condensation reaction between 1,3,5-triformylbenzene and 1,3,5-triformylphloroglucinol with the chiral diamine formed the final chiral COFs LZU-72 and LZU-76 (see Figure 18). These two organocatalytic COFs showed outstanding catalytic performance towards a value-added C-C bond forming reaction, asymmetric aldol reaction. LZU-76 displayed the best performance providing 94.0: 6.0 e.r. value, which was comparable with the benchmark homogeneous analogue (S)-4,7-diphenyl-2-(pyrrolidin-2-yl)-1H-benzimidazole (DPBIP) catalyst for the asymmetric aldol reaction. The authors proposed that the direct construction of chiral COFs from chiral building blocks provided a promising pathway to design and synthesize novel chiral COF structures which could accommodate a large amount of uniform chiral catalyst moiety for more meaningful value-added organic transformations.

Figure 18. (a) Schematic representation for the direct construction of chiral COFs, LZU-72 and LZU-76. (b) Catalytic activity test of LZU-76 in asymmetric aldol reaction. Reprinted and adapted with permission from [192], Copyright 2016, American Chemical Society.

From the above-mentioned examples, it is feasible to incorporate chiral units or moieties into COFs structure for asymmetric enantioselective catalysis. COFs pore and tunnel structure would be further self-assembled into helical structure to fulfill more challenging chiral synthesis.

2.5. Bimetallic or Bifunctional COF-Catalyzed C-C Bonds Formations

The first bifunctional organocatalytic COF material was designed and realized by Banerjee et al., which was stable in aqueous and acidic condition [193]. This COF material was prepared via the Schiff-base condensation between 2,3-dihydroxynaphthalaldehyde (2,3-Dha) and 5,10,15,20-tetrakis (4-aminophenyl)-21H,23H-porphine unit (Tph) in dichlorobenzene (o-DCB,) and dimethylacetamide (DMAc) with a catalytic amount of 6.0 M acetic acid. The as-synthesized COF material displayed unprecedented stability in water and acidic solution due to the catechol group, the presence of trans conformation of imine bonds and intramolecular hydrogen bonding [-OH·N=C; D = 2.579, d = 1.858 Å,
was effectively accelerated by the basic sites of the Dha-Tph-COFs. This discovery manifested its significance in that it was the first stable bifunctional COF catalyst in water and acidic solution.

![Figure 19. (a) The synthesis of 2,3-DhaTph and 2,3-DmaTph by the condensation of Tph and 2,3-Dha/2,3-Dma. The catalytically active porphyrin and catecholic –OH groups are shown in coral and cyan colors, respectively. An ORTEP diagram of 2,3-DhaTph and 2,3-DmaTph monomer units. (b) The catalytic activity towards acid–base catalyzed reaction with various reactants. Reprinted with permission from [193], Copyright 2015, Royal Society of Chemistry.](image)

Besides Mn and Pd bimetallic docking for bimetallic COF catalyst, Rh and Pd bimetallic docking could also be realized through this 2D BPy COF (see Figure 21) [195]. By condensing 4,4′,4″-(pyrene-1,3,6,8-tetrayl)tetraaniline (PyTTA) and various ratio of X% 2,2-bipyridyl-5,5-dialdehyde and 100-X% 4,4′-biphenyldialdehyde in a three-component solvent, a series of structurally tunable 2D COF were prepared. And the authors demonstrated that by a further solution-infiltration method, Pd(OAc)₂ and Rh(COD)Cl was sequentially incorporated into the COF structure in a programmed synthetic procedure. Pd(OAc)₂ dispersed in the interlayer space and y = 146.11], which have been confirmed from the monomer crystal structure (see Figure 19).

Moreover, since the COF material was consisted from the Dha unit which contains weak acidic catechol groups and Tph group containing basic pyrrole groups and imine C=N bonds, this COF possessed acidic and basic sites providing it as a promising bifunctional heterogeneous catalyst. In a model cascade deacetalization-Knoevenagel reaction, this COF material demonstrated excellent isolated yield up to 96%. The deacetalization of benzaldehydedimethylacetal was catalyzed by the acidic sites of this COF catalyst, while the further Knoevenagel reaction between benzaldehyde and malonitrile was effectively accelerated by the basic sites of the Dha-Tph-COFs. This discovery manifested its significance in that it was the first stable bifunctional COF catalyst in water and acidic solution.

Apart from single metal deposited COF materials, bimetallic docked COF materials was designed, synthesized and applied as effective catalysts for a Heck-epoxidation tandem reaction. Mn and Pd bimetallic docking to a bipyridine-imine COF could be realized by a programmed synthetic procedure [194] (see Figure 20). Firstly, a Py-2,2′-BPypPh COF skeleton was constructed via the Schiff-base condensation reaction between 4,4′,4″-(pyrene-1,3,6,8-tetrayl)tetraaniline (PyTTA) and 2,2-bipyridyl-5,5-dialdehyde (2,2′-BPypDCA). The as-formed COF contained two differentiated nitrogen-complexing sites, the bipyridyl and imine units. MnCl₂ was docked with bipyridine ligands first, and then Pd(OAc)₂ was combined with both bipyridine and imine units. The as-synthesized bimetallic docked COF material acted as effective bifunctional catalyst for two different types of organic transformations, i.e., Pd-catalyzed Heck cross-coupling reaction and Mn-catalyzed epoxidation reaction. The COF material transformed iodobenzene and styrene to trans-stilbene oxide in a tandem reaction. Initially, Pd(OAc) incorporated in COF transformed iodobenzene and styrene into trans-stilbene with excellent yield up to 95%, while the Mn in COF catalyzed the epoxidation reaction of trans-stilbene to trans-stilbene oxide in almost quantitative yield (99%). The control group experiments proved that Mn@Py-2,2′-BPypPh COF and Pd@Py-2,2′-BPypPh COF alone could only catalyze the separate epoxidation and Heck reaction. This finding was important since it demonstrated that by elaborately design and choose the ligands, different metal species could be incorporated into a single COF skeleton to fulfill different genres of organic transformation by the certain metal.
coordinated with both imine units and bipyridine ligands, while the more structurally rigid Rh(COD)Cl deposited in the pore space and complexed with the bipyridine ligands. This Rh/Pd bimetallic docked BPh COF demonstrated superior catalytic activity towards a tandem addition-oxidation reaction between phenylboronic acid and benzaldehyde to form initially the intermediate diphenylmethanol and further be oxidized to the final product benzophenone. The authors demonstrated that Rh(COD)Cl moiety in COF was accountable for the addition reaction between phenylboronic acid and benzaldehyde surpassing its homogeneous Rh(COD)Cl analogue in catalytic activity. Pd(OAc)$_2$ was responsible for the oxidation from diphenylmethanol to benzophenone. The as-synthesized Rh/Pd-BPh COF showed excellent recyclability and reusability providing isolated benzophenone products up to 85% yield even after 5 cycles reuse without noticeable leaching of the metal and apparent loss of activity. This report manifested its significance in that it illustrated that two different kind of organometallic compounds could be docked in a structurally tunable COF material by different coordinating groups to render the COF material different catalytic ability towards totally differentiated reaction types in the first time.

Figure 20. Left, (a) Schematic representation of Mn/Pd bimetallic docked COFs prepared via a programmed synthetic procedure; (b) top view and (c) side view of Mn/Pd@py-2,2'-bpy-Ph COF; (d) appearances and (e) PXRD (powder X-ray diffraction) patterns of the COFs before and after metallic loading treatment. Right, Schematic illustration of the tandem reaction catalyzed by Mn/Pd@py-2,2'-bpy-Ph COF.

Figure 21. Left, (a) Use of a three-component condensation system to modulate the nitrogen content of the 2D imine-type COFs. (b) Designed strategies for the monometallic (Route 1) and bimetallic docking (Route 2). (c) Open channels of the COFs. (d) Open channels of the metal loaded COFs. Right, One-pot cascade reactions using different homogeneous/heterogeneous catalysts. Reprinted and adapted with permission from [195], Copyright 2016, Wiley.
2.6. 3D COF-Catalyzed C-C Bonds Formations

Wang et al. described that an interpenetrating dynamic 3D COF LZU-301 could be a Lewis-base catalyst for the Knoevenagel condensation between malonitrile and three aromatic aldehydes [196] (see Figure 22). The authors discovered that for the small size benzaldehyde, the 3D COF LZU-301 provided excellent yield up to 72% in 4 h and 99% in 10 h. However, for the larger 2-naphthalenealdehyde and 9-anthracenealdehyde, the yield notably decreased to 21% and 5% due to a size-effect. The larger fuse-ring aromatic aldehyde could not be accommodated into the pore space of LZU-301, thus did not have enough interaction with its pyridine Lewis-base catalytic site, leading to inferior catalytic performance. This example manifested its significance in that it was the only few 3D COFs which demonstrated considerable catalytic activities in meaningful organic synthesis.

![Solvothermal synthesis of a three-dimensional (3D) COF material, LZU-301, via imine condensation. For clarity, only the single framework of LZU-301 is shown.](image)

**Figure 22.** Left, (a) Solvothermal synthesis of a three-dimensional (3D) COF material, LZU-301, via imine condensation. For clarity, only the single framework of LZU-301 is shown. (b) Side and top views of porous crystalline structure of LZU-301, which features with a 9-fold interpenetration of the underlying diamond net. Different color represents different penetrating frameworks from a side-view. Right, Knoevenagel condensation catalytic experiments. *Only Ph-CHO reacted with malonitrile under the reaction conditions. Reprinted and adapted with permission from [196], Copyright 2017, American Chemical Society.

Another 3D COF DL-COF-1 and DL-COF-2 was synthesized via the dual linking reaction between 1,3,5,7-tetraaminoadamantane (TAA) and 4-formylnaphthaleneboronic acid (FPBA) or 2-fluoro-4-formylphenylboronic acid (FFPBA) forming two kinds of linkage in COF skeleton, namely, boroxine and imine bonds (see Figure 23) [197]. The as-synthesized 3D COFs displayed large specific surface area and incorporated both acidic boroxine sites and basic imine site. These two different sites rendered these 3D COFs as versatile bifunctional heterogeneous catalyst. To demonstrate its catalytic activity, a one-pot deacetalization-Knoevenagel reaction was applied. DL-COF-1 exhibited excellent yields towards both the acid-catalyzed deacetalization (yield up to 100%) and base-catalyzed Knoevenagel condensation reaction (yield up to 98%). The COF crystals can be recycled and reused for three times with almost no loss of activity and no identical change in PXRD and N$_2$ uptake characterization. This example was very encouraging since it was the first bifunctional 3D COF with large specific surface area to fulfill heterogeneous catalytic applications.

Qiu et al. reported that 3D imine COF BF-COF-1 and BF-COF-2 could act as effective basic catalyst for C-C formation Knoevenagel reaction (see Figure 24) [135]. These 3D BF-COF-1 and BF-COF-2 materials were synthesized via the imine condensation between 1,3,5,7-tetraaminoadamantane (TAA) and 1,3,5-triformylbenzene (TFB) or triformylphloroglucinol (TFP). The as-formed 3D BF-COF-1 and BF-COF-2 demonstrated excellent size-selection catalytic activity. For small-size reactants, such as benzaldehyde and malonitrile, these 3D COFs exhibited very good conversion up to 99%. However, for the large reactants, which could not be accommodated into the pore space for efficient catalytic interaction, the reaction proceeded much sluggishly with poor yield. To our best knowledge, this was the first catalytic 3D COF materials, which showed excellent size-selection catalytic mode towards base-promoted Knoevenagel condensation.
**Figure 23.** Left, Strategy for preparing 3D COFs with dual linkages (DL-COFs). (a) Model reaction of 1-adamantanamine (AA) with 4-formylphenylboronic acid (FPBA) to form the triangular molecule with dual linkages. (b) Condensation of tetrahedral 1,3,5,7-tetraaminoadamantane (TAA) and FPBA or 2-fluoro-4-formylphenylboronic acid (FFPBA) to give 3D COFs with dual linkages, DL-COF-1 or DL-COF-2. (c) On the basis of triangular and tetrahedral building units, both DL-COFs show 3D networks with ctn topology. Right, Three-dimensional COFs with dual linkages for bifunctional cascade catalysis. Reprinted with permission from [197], Copyright 2017, American Chemical Society.

**Figure 24.** Left, schematic representation of the strategy for preparing 3D microporous base-functionalized COFs. (a) Model reaction of 1-adamantanamine with benzaldehyde to form the molecular N-(1-adamantyl) benzaldehyde imine. (b) Structure of 1,3,5,7-tetraaminoadamantane (TAA) as a tetrahedral building unit. (c) Structure of 1,3,5-triformylbenzene (R=H, TFB) or triformylphloroglucinol (R=OH, TFP) as a triangular building unit. (d) Condensation of tetrahedral and triangular building units to give a 3D network with the symbol ctn (BF-COF-1 or BF-COF-2). Right, Catalytic activity of BF-COFs in the size-selective Knoevenagel condensation reaction. Reprinted with permission from [135], Copyright 2014, Wiley.
In several C-C coupling reaction such as Knoevenagel condensation, 3D COFs displayed similar high catalytic efficiency as 2D COFs. However, they still differentiate greatly in that 3D COFs catalysis is more sensitive to substrate steric factor. This renders specific 3D COFs the ideal heterogeneous catalyst for size- and shape-dependent catalytic applications.

3. Conclusions

We have overviewed the paragon examples of covalent organic frameworks (COFs) applied as heterogeneous catalyst for C-C bonds formation reactions. Although in its infancy stage compared with other common catalysts used for C-C bonds formation, such as homogeneous transition-metal catalysts, organocatalysts and enzyme catalysts, COF materials still displayed its multi-faceted advantages. First, metal-loaded COF materials showed parallel or even superior catalytic activity in a series of C-C bond formation reactions in contrast with their homogeneous analogues including Heck, Suzuki-Miyaura, Sonogashira cross-coupling, carbon-centered nucleophilic addition and cross-dehydrogenative couplings (CDC). Moreover, COF materials usually demonstrated excellent recyclability and reusability for consecutive runs of catalytic transformations without apparent loss of activity and selectivity. The last but not the least, there appeared some chiral COF (CCOF) materials which showcased powerful control on enantioselectivity in asymmetric organic C-C formation events such as asymmetric Michael addition, Henry reaction, Diels-Alder cyclo-addition, α-aminooxylation, epoxidation and opening of epoxides. Apart from continuously emerging novel COF structures acting as effective heterogeneous catalysts for C-C bond formation reaction in organic synthesis, the new strategy to apply bimetallic and bifunctional COF in catalysis broadens the application fields of COF-based catalysis in more organic reaction types. The new approaches to in-situ loading noble-metal nanoparticles on COF skeleton without requiring external reductants realized the minimal influence and impacts on metal-loading avoiding the decrease of crystallinity and porosity to the least limit. Comparing to homogeneous catalysts, COF-based heterogeneous catalysts still have great space to improve for more challenging reaction types and wider substrate scope for C-C bond formation in organic synthesis. The main current limitations and challenges for COFs materials for C-C bond formation reaction concentrate on the direct activation of inert C-H bonds especially sp\(^3\)C-H bonds to construct sp\(^3\)C-sp\(^2\)C or sp\(^3\)C-sp\(^3\)C bonds. And the activation of energetically important and inert small molecules such as methane, methanol and carbon dioxide for C-C bond formation in highly value-added chemicals such as first-line drugs, natural products and organic functional compounds is another developing direction for COFs materials as heterogeneous catalysts. The third one is to developing more metal-free COFs materials acting as highly efficient versatile heterogeneous catalysts for a number of organic synthetic transformations. Comparing to current common homogeneous catalyst, COFs materials demonstrated better recyclability and reusability and lower cost in separating catalyst and the products. However, COFs materials still need to improve their efficiency, selectivity and substrate scopes. Especially, the current catalytic reaction pathways and mechanisms by COFs materials are not clear in comparison with their homogenous analogue. More extensive and deep mechanistic study in COFs materials catalyzed C-C formation reaction need to be conducted in order to further design more efficient and selective catalytic process and more effective COFs catalyst. As the focus frontier of current materials and catalysts research, COF materials would have brighter prospects for more challenging, meaningful C-C bond formation reaction and more novel principles on COF catalysis would be uncovered along with this promising application field.

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