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Introduction

Challenges and opportunities for the photo-(thermal) synthesis of ammonia

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For more than one century, the synthesis of ammonia (NH₃) through the Haber–Bosch route has allowed the industrial-scale production of fertilizers and other nitrogen-containing compounds. Despite its success, the development of alternative NH3 synthesis methods with a lower CO2 footprint is highly desirable. In this context, the use of sunlight as source of energy to perform the photo-catalytic nitrogen (N_2) reduction reaction represents a promising approach, although the reported efficiencies remain below those required to compete with the traditional HB route. To overcome these limitations, photo-thermal catalysis offers the opportunity to synergistically combine the effect of photo-induced charge carriers and localized heat to perform the synthesis of ammonia. In this review, we first provide an overview of the most recent advances in thermo- and photo-catalytic NH₃ production. Next, we describe the merits of photo-thermal catalysis to enhance the catalytic activity towards NH₃ synthesis. To conclude, we share our viewpoint on the future prospects and challenges ahead for photo-thermal catalysis to become a reality in the production of NH₃ and other industrially relevant chemicals.

Ammonia (NH₃) synthesis is considered one of the most relevant processes in the chemical industry. Together at BASF, German chemists Fritz Haber (1868-1934) and Carl Bosch (1874-1940) developed in 1913 the so-called Haber-Bosch (HB) process for the industrial-scale manufacturing of ammonia.^{1,2} This breakthrough contributed to the extensive production of NH₃ and its application to the large-scale fabrication of fertilizers, explosives and other nitrogen-containing chemicals of industrial interest. After more than one century, the actual ammonia synthesis has barely changed, and indeed circa 95% of the total ammonia production is still obtained through the Haber-Bosch route.

Despite the success of the traditional thermo-catalytic ammonia synthesis, the pursuit of higher efficiency, cost reduction, and environmental sustainability remains ongoing. In this regard, the synthesis of ammonia consumes 3-5% of the world's natural gas production, representing 1-2% of the annual energy supply and approximately 1% of all greenhouse gas generation.^{3,4} In view of this, the current focus of research aims to further optimize the process and explore novel approaches to meet the rising global demand for ammonia while minimizing energy consumption and greenhouse gas

emissions. One promising strategy in this vein is the use of blue hydrogen (H_2) for ammonia production.⁵ This method leverages existing fossil fuel infrastructure for exploitation and distribution while minimizing carbon dioxide (CO₂) emissions through carbon capture and storage (CCS). Recent reports indicate, indeed, that the use of blue ammonia can be a presentday convenient solution to mitigate the climate impacts associated with the conventional ammonia production route, at least until green ammonia becomes economically profitable.⁶ In fact, in September 2020 ARAMCO and the Institute of Energy Economics (Japan), in partnership with SABIC, successfully achieved the production and shipment of blue ammonia from Saudi Arabia to Japan, thus representing the world's first blue ammonia supply-chain demonstration.⁷ Unfortunately, this approach still relies on fossil gas, leading to a high concentration of HB facilities near the gas supply points and increasing transportation costs to remote or underdeveloped locations.8

To achieve decentralized ammonia production and move towards a carbon-free economy, alternative methods for the reduction of nitrogen (N_2) have been proposed. Among these methods, photo-catalysis represents an intelligent approach to drive chemical reactions by taking advantage of the solar energy. Assuming water or renewable H₂ as source of protons, photo-catalysis enables a CO2-free NH3 synthesis using sunlight as driving force. Besides, this strategy offers the additional advantage of allowing small on-site NH₃ synthesis plants powered by renewable energy, which essentially favors local production of ammonia close to the final consumption

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points. Nevertheless, and despite all these potential benefits, the reported efficiencies of most of the photo-catalytic systems for N_2 fixation described to date are still insufficient to meet industrial-scale requirements.⁹

In this context, the integration of both thermo-catalysis and photo-catalysis (from now on, "photo-thermal catalysis") has lately appeared as an excellent opportunity to both reduce the energy requirements and increase the catalytic efficiency of many chemical processes, including CO₂ conversion, H₂ generation or the degradation of pollutants, among others.¹⁰⁻¹⁵ Thanks to the combination of photo-induced localized heating and photo-excited charge carriers, many photo-thermal catalysts have exhibited comparable or even higher catalytic rates than those displayed under pure thermal conditions. For these reasons, it is quite surprising that, to date, only a few works have drawn attention to the synergistic combination of light and heat to perform the NH₃ synthesis reaction, thus remaining as an almost unexplored field of research. Given the emerging prospects of photo-thermal catalysis and the extraordinary relevance of ammonia in our modern societies, we herein present a review of the newest advances in thermo-catalytic, photo-catalytic and synergistic photo-thermo-catalytic NH₃ synthesis during the last five years. The goal is to give an overview of the up-to-date strategies of traditional thermo- and photocatalytic NH₃ production and, at the same time, to highlight the potential of photo-thermal catalysis as a hybrid tool to break the limitations of its individual thermochemical and photochemical counterparts. In this review, we will first briefly introduce the most well-accepted reaction pathways for NH₃ synthesis in thermo- and photo-catalysis. Then, we will describe recent advancements in thermo-catalytic and photocatalytic NH₃ synthesis, paying special attention to novel strategies to moderate reaction conditions and/or improve the catalytic performance. After that, we will outline a summary of the merits of photo-thermal catalysis to boost the catalytic activity towards NH₃ production, giving specific examples found in the literature. Lastly, we will share our opinion on the future directions and challenges ahead for photo-thermal catalysis to become a viable technology for the production of NH₃ and other commodities of industrial interest.

Reaction pathways for NH₃ synthesis

The reaction of N_2 and H_2 to produce NH_3 (eqn (1)) is an exothermic process thermodynamically favored by high pressures and low temperatures.

$$N_2 + 3H_2 \leftrightarrow 2NH_3 \quad \Delta H^\circ = -92.2 \text{ kJ mol}^{-1}$$
 (1)

Mechanistically, the catalytic N_2 reduction reaction is a multi-step process that involves: (i) N_2 adsorption, (ii) $N \equiv N$ bond activation and/or cleavage, (iii) $N_{2(ads)}$ or $N_{(ads)}$ hydrogenation and (iv) NH_3 desorption.^{16,17} Traditionally, two broadly accepted mechanisms have been proposed for the N_2 reduction reaction, namely the dissociative pathway and the associative pathway (Fig. 1).¹⁸



Fig. 1 Reaction pathways for the N_2 reduction reaction to ammonia. Adapted with permission from ref. 18. Copyright (2017) Elsevier.

In the dissociative pathway, the N \equiv N bond cleavage takes place before the hydrogenation of the nitrogen atom. In view of the high dissociation energy of the first N–N bond (410 kJ mol⁻¹), this route is extremely energy-intensive, and, indeed, traditional thermochemical HB systems are believed to follow this mechanism for the production of ammonia.¹⁹ On the contrary, the associative pathway is based on the sequential hydrogenation of nitrogen atoms while bonded to each other. Two different sub-types of associative pathways are possible depending on the hydrogenation site.

In the alternating associative pathway, nitrogen atoms of adsorbed N_2 are successively hydrogenated and, eventually, NH_3 is desorbed after the cleavage of the N–N bond. In the distal associative pathway, however, the hydrogenation occurs preferably on the distal nitrogen atom.⁹ Overall, the associative pathway does not require the cleavage of the first bond in N_2 molecule so in these cases the energy requisites are considerably lower. In consequence, the associative pathway is expected to be the preferential mechanism for electro- or photo-catalytic NH_3 production systems.

Thermo-catalytic NH₃ synthesis

As previously mentioned, NH_3 synthesis reaction is thermodynamically enhanced at high pressures and low temperatures. From a kinetic point of view, however, temperatures above 200 °C are necessary in order to achieve significant conversion values. For these reasons, as a compromise between kinetics

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and thermodynamics, thermo-catalytic NH_3 synthesis is generally conducted under medium temperatures and high pressures (400–450 °C and 150–250 atm), achieving conversion rates of 10–15% per pass.²⁰ These operating conditions represent the main drawback for the thermo-catalytic NH_3 synthesis, not only for the elevated consumption of energy but also for the limited stability of the catalysts. For these reasons, one of the main challenges in the field is the development of systems which enable milder reaction conditions, especially in terms of operating pressure, as compression cycles represent the major capital cost of the whole HB process.

Fe-based catalysts

Iron-based catalysts represent, undoubtedly, the most investigated and applied materials in the NH₃ synthesis reaction. Particularly, iron-fused catalyst derived from hematite (Fe₂O₃), magnetite (Fe₃O₄) and/or wüstite (Fe_{1-x}O) are the most commonly used in modern industries. Nevertheless, it is also true that early studies conducted with iron catalysts at BASF facilities showed quite disappointing activities and, in fact, this served as a spur for the study of the effect of additives on the catalytic performance. A comprehensive investigation was developed in between 1909 and 1911 by Carl Bosch's assistant Alwin Mittasch, who conducted a systematic testing of 2500 different materials in order to find an efficient and stable catalyst for the NH_3 synthesis.²¹ The scrutiny was prolonged until 1922 including about 5000 different candidates and eventually the best catalyst resulted to be a multi-component mixture consisting of magnetite-based iron catalyst with small amounts of alumina, calcium oxide, and potassium alkali.²¹ This exhaustive screening demonstrated the importance of the addition of promoters to the formulation of catalysts in order to provide beneficial structural and electronic effects, as we will later elaborate in this review.

Supported iron catalysts are also an appealing subfamily of ammonia synthesis catalysts. In these cases, carbon or metal oxides usually act as support materials for active iron species. When it comes to carbon-based catalysts, the one-step pyrolysis of metal-organic compounds has lately emerged as a straightforward method to obtain well-dispersed metal nanoparticles (NPs) encapsulated into a carbon matrix. For instance, Morlanés *et al.* developed a novel catalyst based on iron NPs embedded on carbon through the controlled pyrolysis at 735 °C of iron phthalocyanines.²² Upon promotion with Cs, the as-prepared catalyst (10%Cs-FePc) exhibited a stable production of NH₃ in the order of 14 000 µmol g⁻¹ h⁻¹ at 400 °C and 3 MPa, even surpassing an iron benchmark catalyst (Fe_KM1) under similar reaction conditions (Fig. 2A–C). Long-



Fig. 2 Catalytic performance of the phthalocyanine-derived catalysts for the ammonia synthesis reaction. (A) Pressure effect on the NH₃ synthesis rate at 400 °C. (B) Temperature effect on the NH₃ synthesis rate at 1 MPa. (C) Specific activity in μ mol NH₃ h⁻¹ g_{metal}⁻¹ and TOF (s⁻¹) (diagonal stripes) at 400 °C and 3 MPa. (D) Stability tests in the range of 400–520 °C and 3–7 MPa. Reproduced with permission from ref. 22. Copyright (2020) Royal Society of Chemistry.

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term stability tests also demonstrated an outstanding stability of the catalyst after more than 100 hours of reaction (Fig. 2D).

The promotion with Cs was crucial to boost the catalytic activity and authors speculated that the presence of alkali both decreased the energetic barrier for nitrogen dissociation and increased the number of surface active sites by destabilization of adsorbed NH_x species. Interestingly, homologous materials prepared by conventional impregnation of carbon with iron/ alkali displayed much lower activity. The authors concluded, therefore, that the remarkable stability and catalytic activity of 10%Cs-FePc directly stemmed from the synthetic approach based on the pyrolysis of phthalocyanine precursor. A followup work from the same group demonstrated that it was also possible to obtain highly efficient bimetallic Fe-Co catalysts through the physical mixture of Fe and Co phthalocyanines followed by the pyrolysis treatment.²³ Under the optimal Fe:Co ratio, the bimetallic catalyst outperformed the monometallic Fe and Co counterparts, exhibiting a NH₃ production of 53 666 μ mol g⁻¹ h⁻¹ at 400 °C and 7 MPa. In a related precedent, the groups of Zou and Ma reported a K-promoted Fe/C catalyst derived from the pyrolysis under He of Fe-based metal-organic-framework (MOF) xerogels.²⁴ Upon addition of 1% K to the MOF-derived Fe catalyst, the catalytic activity reached 30 400 μ mol g⁻¹ h⁻¹ at 400 °C and 3 MPa, approximately 3 times higher compared to the sample without potassium. X-ray photoelectron spectroscopy (XPS) measurements revealed a shift to lower binding energies for the Fe 2p peak along with the increase of potassium concentration in the samples, thus suggesting a charge transfer from K ions to the surface of iron. Eventually, this electron-rich environment on the Fe surface favors nitrogen activation and improves the catalytic activity of promoted samples. Curiously, potassium not only enhanced the catalytic performance by modulating the electronic properties of the Fe active sites but also by increasing the thermal stability of the catalyst. Authors observed that unpromoted samples suffered from a severe deactivation after 24 h of reaction under high temperature conditions (475 °C). In situ temperature-programed reaction-mass spectrometry (TPR-MS) measurements demonstrated that the deactivation of the samples stemmed from the methanation of the catalyst under reaction conditions, which at the end led to carbon loss and agglomeration of Fe nanoparticles. However, under the same reaction conditions K-promoted samples retained more than 90% of their activity due to the inhibition of the methanation reaction by potassium.

Apart from carbon, oxyhydrides have been recently explored as potential supports in the formulation of new NH₃ synthesis catalysts. In these cases, a low content of iron in the order of 1–2% in weight is deposited on the inorganic support, in sharp contrast with the high iron loading of up to 50% that can be achieved through the controlled pyrolysis of Fe-containing carbon-based precursors. For example, the groups of Kobayashi and Kageyama reported the preparation of 1% Fe/ BaTiO_{2.35}H_{0.65} catalyst with a remarkable catalytic activity of 14 000 μ mol g⁻¹ h⁻¹ at 400 °C and 5 MPa.²⁵ Authors found that the oxyhydride support could successfully inhibit H₂ poisoning by removing the excess of adsorbed hydrogen through an effective spillover pathway. Furthermore, the oxyhydride served as electron donation source to the Fe metal sites, thus reducing the strength of the triple bond of adsorbed N₂ and, consequently, improving the catalytic activity. Further research from Kitano and co-workers demonstrated that the presence of other heteroanions like nitride (N³⁻) could enhance even more the performance of oxyhydrides as NH₃ catalysts.²⁶ In this regard, authors developed a new perovskite oxynitride-hydride BaCeO_{3-x}N_yH_z loaded with small amounts of Fe (1.2 wt%). Using this catalyst, the NH₃ synthesis rate at 400 °C and 0.9 MPa was 6800 μ mol g⁻¹ h⁻¹. In comparison, Fe-supported BaCeO₃ displayed a catalytic rate in the range of 30 μ mol g⁻¹ h^{-1} . Authors performed additional isotopic experiments in order to disentangle the promotion effect of Fe-BaCeO_{3-x}N_vH_z and found that, unexpectedly, the rate-determining step shifted from N2 dissociation to N-H bond formation due to the presence of N³⁻ and H⁻ anions. Ultimately, this change in the reaction mechanism translated into a lower activation energy for the ammonia synthesis (46–62 kJ mol⁻¹). In the case of Fe-BaCeO₃, however, the NH₃ formation followed the conventional mechanism characterized by high-energy barriers (85-121 kJ mol⁻¹).

Ru-based catalysts

In spite that iron-based catalysts have enjoyed a historical hegemony in the Haber–Bosch process, it was in the 1970s when a consortium between British Petroleum and M.W. Kellogg companies developed a novel carbon-supported ruthenium catalyst (Ru/C) for the NH_3 synthesis reaction.²⁷ Since then, the research on innovative Ru-based catalysts has gained considerable relevance and a vast number of supports such as perovskites, oxides or carbon-derived materials have been investigated.^{28–33}

For instance, the use of ceria (CeO_2) has been extensively reported due to its abundant oxygen vacancies that increase the electron density of Ru sites to promote N₂ activation.³⁴ Furthermore, it has been found that parameters like morphology of CeO2 or metal-support interaction are key to maximize the catalytic activity of the resultant materials.³⁵ In this regard, Ma and co-workers compared the catalytic activity for NH₃ synthesis of three different morphologies of ceria (nanocubes, nanorods, and nanoparticles) with a nominal Ru loading of 4%.36 At 400 °C and only 1 MPa, ceria nanorods displayed the highest catalytic activity (3830 μ mol g⁻¹ h⁻¹) followed by ceria nanocubes (1289 μ mol g⁻¹ h⁻¹) and ceria nanoparticles (529 μ mol g⁻¹ h⁻¹). XPS measurements also revealed the presence of CeH_{2+x} hydride on the samples and, curiously, authors identified a positive correlation between the amount of this hydride and the catalytic activity. Basically, the role of the CeH_{2+x} species seemed to be the electron donation to the Ru active sites, thus increasing the catalytic performance towards NH₃ synthesis. More recently in 2021, Li and collaborators described how the treatment of Ru-CeO₂ catalyst with hydrazine (N_2H_4) could modulate the electronic metal-support interaction and boost the catalytic activity in the NH₃ synthesis

reaction.³⁷ Authors found that the mild reduction of Ru-CeO₂ with hydrazine followed by H₂ treatment at 500 °C caused an increase in (i) the fraction of metallic Ru, (ii) the concentration of Ce³⁺ and (iii) the number of exposed species of Ru compared to the conventional reduction with only H₂. In the absence of N₂H₄ treatment, oxygen migration from the ceria support to the Ru species led to the lowering of exposed metallic Ru. However, the treatment with N₂H₄ improved the electronic metal-support interaction to create electron-rich states at the Ru sites and, eventually, increased the activity of Ru-CeO₂. Interestingly, a very recent work from Feng and collaborators revealed that ceria could act not only as a support but also as a promoter of Ru-based catalysts.³⁸ In their work, authors deposited small clusters of ceria on a Ru/Al₂O₃ catalyst and demonstrated that these sub-nanometric ceria domains could prevent the H₂ poisoning of Ru active sites. Furthermore, XPS data suggested that ceria clusters acted as electron donors to modulate the electronic structure of Ru, thus facilitating the N₂ adsorption and H₂ dissociation on its surface. Apart from ceria, other metal oxides such as MgO, La₂O₃ or Y₂O₃ have been recently explored as potential supports for Ru-based catalysts for the HB reaction.³⁹⁻⁴¹

Besides metal oxides, other materials like hydrides, nitrides, or amides have lately demonstrated to be excellent hosts for Ru in the preparation of new ammonia synthesis catalysts, even outperforming their Fe-based counterparts.^{42–45} For instance, Kitano *et al.* developed a Ru/Ca(NH₂)₂ catalyst promoted with Ba to create Ru–Ba core–shell nanoparticles supported on the calcium amide.⁴⁵ The as-prepared catalyst displayed an outstanding activity towards NH₃ synthesis of 60 400 µmol g⁻¹ h⁻¹ at 360 °C and pressure as low as 0.9 MPa, thus exceeding by 6 times the performance of the industrial benchmark Fe catalyst (Fig. 3). Furthermore, Ru–Ba/Ca(NH₂)₂



Fig. 3 (A) Temperature dependence on the catalytic performance of Ru/Ba-Ca(NH₂)₂ catalyst for the ammonia synthesis reaction at 0.9 MPa. (B) Particle size distribution and (C) HAADF-STEM image of Ru/Ba-Ca (NH₂)₂ catalyst. EDX elemental mapping images of (D) Ba (yellow) and (E) Ca (green), where Ru is displayed as red. Reproduced with permission from ref. 45. Copyright (2018) John Wiley and Sons.

could operate at relatively low temperatures below 300 °C, achieving NH_3 production rates of 7500 µmol g^{-1} h⁻¹ at only 260 °C.

In fact, one of the main advantages of Ru catalysts compared to their Fe-based equivalents lies in their improved catalytic activities under mild reaction conditions of temperature and pressure, therefore contributing to a considerable reduction of the capital cost of ammonia plants. Nevertheless, it is still hard to imagine a scenario in which Ru catalysts widely substitute traditional Fe catalysts, especially if one takes into account the high cost and limited availability of Ru. In the specific case of carbonaceous supports, the presence of Ru can also lead to the partial loss of the carbon due to its detrimental methanation, thus reducing the long-term stability of the catalyst.46 All these drawbacks still hamper the broad implementation of Ru catalysts in conventional Haber-Bosch plants and, indeed, by 2010 only sixteen ammonia plants around the globe operated using commercial ruthenium-based catalysts.21

Co-based catalysts

Alternative first-raw transition metals like cobalt or nickel have been explored in recent years as potential catalysts for NH₃ synthesis.47,48 Particularly for Co-based catalysts, notable catalytic activities at moderate operating conditions were achieved. For example, in a similar way to Fe catalysts derived from the pyrolysis of carbonaceous precursors, in 2020 Morlanés and collaborators reported the use of a barium-promoted Co/C catalyst obtained from the pyrolysis of Co phthalocyanine at 735 °C.²² The as-prepared materials displayed an ammonia production of 10 250 μ mol g⁻¹ h⁻¹ at 400 °C and 3 MPa. As in the case of Cs for Fe/C catalyst, the promotional effect of Ba on Co/C catalyst was uniquely ascribed to its electron donation activity but not to a protective effect against methanation of the carbon support. In 2015, the group of Pilecka evaluated the effect of barium as promoter of an activated carbon-supported Co catalyst for ammonia production.49 Under optimal conditions, Ba-Co/C catalyst achieved a NH₃ production of 88 235 μ mol g⁻¹ h⁻¹ at 400 °C and a high pressure of 9 MPa. However, in contradiction with the work of Morlanés et al., in this case authors found that even small amounts of Ba were able to enhance the catalytic activity by suppressing the undesirable methanation of the carbon matrix. All these results suggest that the nature and preparation protocol of the carbonaceous supports are decisive to determine both the activity and stability of the resultant catalysts.

The use of inorganic supports has been also described in the formulation of Co-based catalysts. Kitano and co-workers prepared a perovskite oxynitride-hydride, $BaCeO_{3-x}N_yH_z$, which was loaded with a 4.7 wt% of Co.²⁶ The as-prepared catalyst achieved an outstanding NH₃ production rate of 10 100 µmol g⁻¹ h⁻¹ at 400 °C and a very low pressure of 0.9 MPa. When authors decreased the temperature to 300 °C keeping constant the pressure, the catalytic activity significantly decreased to ~2000 µmol g⁻¹ h⁻¹. It is worth noticing that, under the same reaction conditions, an analogous cata-

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lyst with a 4.3 wt% of Ru displayed a NH₃ production rate close to 5000 μ mol g⁻¹ h⁻¹, thus highlighting the promising attributes of Co-based catalysts. In the last years, the group of Jiang has performed an extensive research on Co/CeO₂-based catalysts for the HB process. As we discussed in the previous section for CeO₂-supported Ru catalysts, authors found that the shape of CeO₂ (polyhedral, nanorods or hexagonal morphology) was critical to optimize the catalytic performance of a series of Co-based catalysts.⁵⁰ Authors suggested that the morphology had a strong influence of the reducibility of both Co oxides and CeO₂ support, thus modifying the metal-support interaction. Co NPs supported on CeO2 polyhedrons displayed the best catalytic activity achieving 86 000 μ mol g⁻¹ h⁻¹ at 430 °C and 10 MPa. In a follow-up work, the same authors studied the effect of alkali promoters (Ba and K) on the catalytic activity of Co/CeO₂ catalysts.⁵¹ Quite surprisingly, Ba showed a positive effect on the performance but the addition of K, even at small quantities, was detrimental to improve the catalytic activity. Later in 2019, authors made a last effort to strengthen the metal-support interaction of Co/CeO2 catalysts.⁵² To this end, they used dopamine during the synthesis to favor the stability of Co NPs and increase the interaction between Co and CeO₂ thanks to a strong interfacial bonding. This improved metal-support interaction was accompanied by a reduction of the particle size of Co, a better reducibility and a lower activation energy for N2 desorption. Under optimal conditions, the Co/CeO2 catalyst exhibited an outstanding NH3 production rate of 19120 µmol g⁻¹ h⁻¹ at 425 °C and a pressure as low as 1 MPa.

In general, most of the productivity values presented in this section are comparable or even outperform those achieved by many Ru-based catalysts. Taking into account that the main advantage of Ru-based catalysts is the possibility to operate under low pressure, Co-based catalysts have exhibited a huge potential to compete with Ru in the development of efficient yet affordable NH₃ catalysts operative under mild reaction conditions.

Photo-catalytic NH₃ synthesis

As discussed above, the traditional HB synthesis heavily relies on fossil energy sources (particularly the steam methane reformers for the production of H_2), thus contributing to estimated CO_2 emissions in the order of 1.5–1.6 tons of CO_2 per ton of NH₃.⁵³ For this reason, the introduction of water or renewable hydrogen (even blue hydrogen in the worst-case scenario) as reducing agents constitutes a key improvement to mitigate the carbon footprint of the NH₃ synthesis reaction. In addition to this, the implementation of renewable energy (either wind or solar) to drive the full process would further contribute to alleviate CO₂ emissions associated to the HB process.

With this in mind, photo-catalysis has emerged as a viable strategy to achieve a low-carbon emitting NH₃ synthesis route. On the one hand, photo-catalysis offers the possibility to integrate water as source of protons for the N₂ reduction reaction. On the other hand, photo-catalysis uses sunlight as unique driving force and operates under mild reaction conditions of pressure and temperature, therefore decreasing the global energy requirements of the process. Consequently, the photocatalytic NH₃ synthesis represents a promising strategy to overcome the limitations of the conventional thermo-catalytic pathway and, indeed, over the last years it has become a very dynamic field of research.54-57

Despite these encouraging features, the catalytic performance of most of the photo-catalytic NH₃ synthesis systems reported to date is still insufficient to meet the requirements for any industrial scale-up of this technology. This is mainly due to the limited utilization of sunlight by many photo-catalysts, the fast recombination rate between photo-generated electrons and holes and the slow reaction dynamics characteristic of photo-catalytic processes. These shortcomings are, obviously, shared by all photo-catalytic reactions but, particularly for the nitrogen reduction reaction, the process is even more complex considering the extremely high reduction potential (-4.2 eV vs. NHE) necessary to activate molecular nitrogen.¹⁹ Although it is possible to circumvent this thermodynamic limitation through alternative multiple-electron transfer processes with lower reduction potentials, this entails slower reaction kinetics that, inevitably, decrease the productivity of the photo-catalytic system. Furthermore, even if the photo-catalyst possesses a sufficiently high reduction potential at its conduction band to make the N2 reduction thermodynamically feasible, the selectivity can be compromised due to presence of the undesirable H₂ evolution reaction under aqueous media, thus further reducing the overall efficiency of the process.

All the above-mentioned drawbacks have motivated active lines of research devoted to improve the efficiency and the selectivity of photo-induced NH₃ synthesis. Therefore, in the following sections of this review we will describe the current advances in the design of novel photo-catalysts, paying special attention to different strategies to overcome the limitations that still impede photo-catalysis from becoming a viable route for NH₃ production.

Inorganic semiconductors

TiO₂-based catalysts. Among all inorganic semiconductor oxides, titanium oxide (TiO₂) has been the most studied material in photo-catalysis for a wide variety of applications including environmental remediation, production of solar fuels and, of more relevance for the scope of this review, NH₃ synthesis. In fact, it was in 1977 when Schrauzer and Guth reported for the first time the use of a series of doped-TiO₂ for the photo-catalytic reduction of N_2 in water to produce NH_3 .⁵⁸ Since then, many works have explored TiO₂-based photo-catalysts for the production of ammonia, although the reported efficiencies are still far from satisfactory.^{59–62}

Defect engineering has recently gained considerable relevance as a tool to enhance the catalytic activity in semiconductor photo-catalysts. Thus, for example, the introduction of oxygen vacancies, heteroatom doping or lattice distortions can modify the surface, chemical and/or light absorption properties.^{63,64} When it comes to light-induced NH₃ synthesis,

TiO₂-based catalysts have also benefited from this strategy to improve their catalytic performance. For instance, Zhang and co-workers developed ultrathin defective TiO₂ nanosheets through doping with copper ions.⁶⁵ Upon doping with Cu, authors increased the oxygen vacancy (VO) defect states and this translated into an enhancement in the light absorption in the range of 400-800 nm (Fig. 4A). These new absorption features derived not only from the presence of oxygen vacancies but also from $2E_g \rightarrow 2T_{2g}$ transitions from O to Cu atoms and d-d transitions from copper ions. TiO₂ nanosheets samples containing a 6 mol% of Cu displayed the best catalytic results, achieving a NH_3 production rate of 78.9 µmol g⁻¹ h⁻¹ and quantum yields in the range of 0.05-0.25% when using different monochromatic lights from 700 to 400 nm (Fig. 4B and C). Additional photocurrent and time-resolved photoluminescence (PL) decay measurements together with DFT calculations demonstrated that doped TiO2 nanosheets exhibited an improved charge separation of electron-hole pairs owing to the presence of VO acting as electron-trapping sites.

In a related precedent, Tang and collaborators reported the synthesis of porous carbon-doped anatase TiO_x (C-TiO_x) nanosheets obtained from the thermal-oxidation etching of Ti_3SiC_2 .⁶⁶ As a result of the carbon doping, the obtained C-TiO_x material displayed a high concentration of Ti^{3+} species that could be easily adjusted by modifying the synthetic con-

ditions. According to both steady-state and time-resolved PL analysis, authors found that the Ti³⁺ sites could act as electron traps to extend the lifetime of charge carriers, compared to the un-doped TiO2. N2 chemisorption measurements also indicated that $C-TiO_x$ samples exhibited a high amount of active sites for N₂ activation. All these features, together with an extended light absorption in the visible region thanks to the carbon doping, contributed to the enhancement of the catalytic activity of C-TiO_x nanosheets towards the photo-catalytic N_2 reduction. At the optimal Ti^{3+} concentration, the as-prepared material showed a remarkable NH₃ production of 109.3 μ mol g⁻¹ h⁻¹ under visible light and an apparent quantum efficiency of 1.1% at 400 nm. All these results suggest that the introduction of both oxygen vacancies and/or Ti³⁺ sites is a powerful strategy to improve the efficiency of TiO₂-based catalysts for photo-catalytic N₂ reduction.

Besides defect engineering, surface functionalization represents an alternative route to enhance the catalytic activity of TiO_2 -based photo-catalysts. Traditionally, noble metals like platinum, palladium or ruthenium have been proposed as potential co-catalysts in combination with TiO_2 . However, the rise of plasmonic photo-chemistry has recently shifted the attention to alternative metals like Au or Ag that present intense plasmon bands in the visible-NIR range. Apart from the extended light absorption, the creation of a metal/semi-



Fig. 4 (a) UV-vis diffuse reflectance spectra (DRS) of TiO_2 nanosheets with different percentages of Cu doping. (b) NH₃ yield of different TiO_2 nanosheets samples under UV-vis radiation (c) UV-vis diffuse reflectance spectrum and quantum efficiency for NH₃ evolution by 6%-TiO₂ sample under monochromatic light of different wavelengths. Reproduced with permission from ref. 65. Copyright (2019) John Wiley and Sons.

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conductor heterojunction favors the formation of a Schottky barrier that facilitates the injection of plasmon-induced energetic electrons from the metal to the semiconductor and, at the same time, prevents the return of those electrons back to the metal, thereby improving the charge separation at the heterojunction interface. An appealing example of this approach can be found in the work of Yang and collaborators, where they describe the decoration of oxygen vacancy-containing TiO₂ nanosheets with Au nanocrystals (Au/TiO₂-OV) and its application in the N₂ photo-reduction reaction.⁶⁷ Authors claimed a tandem pathway for the N2 reduction by Au/TiO2-OV catalyst, where both oxygen vacancies and Au nanocrystals contributed to improve the catalytic performance. In fact, the catalytic activity increased not only with the number of oxygen vacancies in the TiO₂ but also with the loading of Au, achieving a NH₃ production rate close to 100 μ mol g⁻¹h⁻¹ using a 2.53 wt% Au. It is worth mentioning that authors employed a 420 nm cut-off filter during the experiments, so the catalytic performance entirely derived from the excitation of Au nanocrystals. Indeed, the analysis of the action spectrum revealed a clear match with the absorption spectrum of Au/TiO₂-OV, therefore indicating that the photo-induced hot electrons from plasmonic Au crystals were responsible for the N₂ reduction. Mechanistic studies including N2 chemisorption, transient absorption spectroscopy (TAS) and photo-voltage measurements suggested that, on the one hand, oxygen vacancies acted as active sites for the activation of N2 molecules while; on the other hand, Au nanoparticles provided plasmoninduced hot electrons to drive the photo-reduction towards NH₃.

Feng et al. also explored the functionalization of oxygen vacancy-enriched TiO_{2-x} with plasmonic Ag nanoparticles and PW₁₀V₂ to create a ternary composite for N₂ photo-reduction.⁶⁸ In this case, a Z-scheme heterojunction between TiO₂ and $PW_{10}V_2$ was responsible of the N₂ reduction, where electrons and holes were located at the TiO₂ conduction band (CB) and PW₁₀V₂ valence band (VB), respectively. Upon plasmon excitation, Ag nanoparticles injected hot electrons to the CB of TiO₂, thus improving the charge separation and increasing the number of available electrons for the N₂ reduction, as demonstrated by transient photo-current and PL measurements. For a broader analysis on the state-of-the-art for the photo-catalytic N₂ reduction reaction by plasmonic structures, readers are referred to an excellent review from Puertolas and collaborators.56

Bi-based catalysts. A pioneer work from Zhang and collaborators in which authors demonstrated the photo-reduction of N_2 using BiOBr nanosheets set the basis for a broad field of study on Bi-based photo-catalysts.⁶⁹ Since then, different strategies have been adopted in order to improve the catalytic performance of Bi-based catalysts such as defect engineering, surface functionalization or the assembly of heterostructures, among others.⁷⁰

For example, Di *et al.* prepared Bi₃O₄Br layers with surfaceconfined defects based on Bi and O vacancies.⁷¹ Authors found a clear positive relationship between the degree of defects of the Bi₃O₄Br materials and the amount of NH₃ generated under light radiation. Both femtosecond transient absorption spectroscopy (fs-TAS) and time-resolved transient PL measurements indicated that the presence of defects was responsible of an enhancement in the charge carrier separation owing to the capacity of defects to act as electron traps. In spite of these encouraging results, a serious weakness of this work relies on the catalytic stability tests, which were limited to only 4 consecutive cycles. Unfortunately, this reduced number of catalytic cycles is insufficient to properly assess the stability of the catalyst under study. Xu and collaborators induced defects on pristine bismuth subcarbonate (Bi₂O₂CO₃) through hydrothermal treatment of NaBiO₃ with graphitic carbon nitride.⁷² Importantly, authors found that the introduction of defect levels improved the light absorption properties of the samples, showing and extended absorption towards the NIR region. Consequently, defective Bi2O2CO3 displayed catalytic activity towards NH₃ production under visible light, in sharp contrast with the pristine counterpart. It is worth mentioning that the defect density also determined the position of the defect level, so there was an optimum degree of defects to maximize the catalytic performance.

An alternative approach for the formulation of new Bibased photo-catalysts is the design of heterojunctions combining two or more elements. The group of Wang recently reported the synthesis of a Bi₂MoO₆/BiOBr composite containing both bismuth and oxygen vacancies.⁷³ Interestingly, bismuth vacancies not only modulated the band position of the composite but also induced the formation of additional oxygen vacancies. The synergy between bismuth and oxygen vacancies led to an improvement in the light absorption properties of the resulting heterostructure. Furthermore, bismuth vacancies provided adequate adsorption sites for N2 activation. As a result, the composite displayed a \sim 4-fold NH₃ production compared to the samples with a poor content of defects. Stability tests were performed for 5 consecutive cycles, showing a slightly decrease in the catalytic performance that authors attributed to the partial loss of oxygen vacancies under reaction conditions. The same group expanded this approach through the design of a ternary heterostructure, namely a Bi@BiOBr-Bi2MoO6 composite fabricated by two consecutive solvothermal methods.⁷⁴ Both semiconductors, BiOBr and Bi₂MoO₆, created a type-II heterojunction in which, under light radiation, photo-excited electrons migrated to the CB of Bi₂MoO₆ and holes accumulated at the VB of BiOBr. Owing to the surface plasmon resonance of metallic Bi and the presence of oxygen vacancies derived from the solvothermal treatments, the light absorption properties of the composite extended towards the IR region of the spectrum. All in all, the combination of the three elements had a positive effect on the catalytic performance, achieving a NH₃ photo-reduction rate as high as 167.2 μ mol g⁻¹ h⁻¹. In this regard, other Bi-based ternary heterojunctions containing plasmonic particles have been proposed, for instance a combination of graphitic carbon nitride, particles and Bi₂WO₆ Bi to create а g-C₃N₄(a)Bi(a)Bi₂WO₆ composite or a heterostructure made of

 TiO_2 , Au particles and $(TiO_2/Au/BiO_4I)$.^{75,76} Furthermore, besides classical semiconductors other potential candidates to fabricate heterojunctions with Bi-based materials have recently included MOFs, MXenes or even graphenes.^{77–79}

Other semiconductor-based catalysts

In addition to TiO₂ and Bi-based catalysts, other families of semiconductors have drawn the attention of researchers as promising candidates to drive the photo-catalytic reduction of N₂. For instance, niobates are a well-known class of materials that have been extensively explored as catalysts for photo-catalytic H₂ evolution and, in the last years, they have shown a great potential for the N₂ photo-fixation.^{80,81} In 2019, the groups of Zhao and He described a Ag/KNbO3 nanocomposite prepared by a hydrothermal method followed by photo-deposition of Ag nanoparticles.⁸² Authors found that it was possible to modulate the morphology, crystallinity and textural properties of KNbO3 by adjusting both the amount of KOH and the hydrothermal temperature during the material synthesis. Under optimal conditions, KNbO3 nanorods decorated with a 0.5% Ag molar ratio displayed a remarkable NH₃ production rate of 385 μ mol L⁻¹ g_{cat}⁻¹ h⁻¹ under simulated sunlight using ethanol as hole scavenger. Given the wide bandgap of KNbO₃, the photo-activity under visible light was exclusively attributed to Ag nanoparticles, which injected plasmon-induced hot electrons to the CB of KNbO3 to drive the N2 reduction to NH3. In a follow-up work from the same group, authors tried to substitute the precious Ag nanoparticles by a more affordable cocatalyst. Hence, authors photo-deposited NiO nanoparticles on KNbO3 and the as-prepared NiO/KNbO3 composite reached a NH₃ productivity as high as 470 μ mol L⁻¹ g_{cat}⁻¹ h⁻¹ under simulated sunlight.⁸³ In this case, both NiO and KNbO₃ semiconductors formed a type-II heterojunction that favored charge migration across the heterostructure, with electrons and holes concentrating at the CB of KNbO3 and VB of NiO, respectively.

Metal sulphides are also a long-established family of materials for photo-catalytic applications. With respect to N₂ photo-fixation, a pioneer work from Miyama and collaborators in 1980 proposed for the first time the use of Pt-CdS as photocatalyst for NH₃ production.⁸⁴ Since then, different examples of sulphide-based photo-catalysts for N2 fixation have been reported.⁸⁵⁻⁸⁸ One of the main advantages of metal sulphides is their low bandgap that allows for the exploitation of lowenergy wavelengths in the visible range ($\lambda > 400$ nm).⁸⁹ However, metal sulphides usually suffer from low stability and fast charge carrier recombination, thus hampering their broad applicability.⁹⁰⁻⁹² These drawbacks have motivated intense lines of research to improve the stability and charge separation efficiency of metal sulphides. One of these strategies consists in the fabrication of heterojunctions between metal sulphides and other semiconductors. For instance, in 2020 Sun et al. prepared CdS nanorods functionalized with sulfur vacancy-rich 1T MoS₂ nanosheets (SV-1T-MoS₂).⁹³ The as-prepared SV-1T-MoS₂/CdS composite with optimal MoS₂ loading (30%) showed a photo-catalytic NH₃ production of 8221 µmol L⁻¹ g_{cat}^{-1} h⁻¹ with an AQE of 4.4% using methanol as sacrificial

agent. This value represented a ~4-fold enhancement compared to bare CdS, thus evidencing the positive effect of MoS_2 nanosheets as co-catalyst. In this case, the introduction of MoS_2 containing sulfur vacancies improved the electron mobility and charge carrier separation, as demonstrated by both steady-state and time-resolved PL, transient photo-current and electrochemical impedance spectroscopy (EIS) measurements (Fig. 5).

DFT studies also revealed that the presence of both MoS₂ and sulfur vacancies was critical to promote N2 adsorption. In spite of this, the study on the stability of the SV-1T-MoS₂/CdS composite was limited to only three consecutive uses, showing a remarkable 20% decrease compared to the fresh sample. This highlights again the importance of performing longer stability tests to demonstrate the reusability of the photo-catalysts under study, especially on those affected by photo-corrosion events. In a related precedent, the group of Yang fabricated nanoporous films of a WS2@TiO2 type II heterojunction.⁹⁴ Compared to bare TiO_2 , the presence of WS₂ improved the charge carrier separation and transfer, as demonstrated by both the increase in the fluorescence time and the reduced semicircle in the Nyquist plots. Upon optimal WS₂ loading, the WS₂@TiO₂ photo-catalyst rendered a maximum NH₃ production rate of 1390 μ mol g_{cat}^{-1} h⁻¹ in the absence of any sign of deactivation after 4 consecutive cycles. The proposed reaction mechanism involved the charge carrier generation at WS₂ and subsequent electron transfer to the CB of TiO₂, where they were finally injected into anti-bonding orbitals of N₂. Photogenerated holes were accumulated at the VB of WS₂ and then consumed by the hole scavenger. Other composites including In₂O₃/In₂S₃ microspheres, MoS₂/n-MgIn₂S₄ sheets or g-C₃N₄/ ZnMoCdS have been also reported although in all these cases the catalytic performance was far from adequate.95-97



Fig. 5 (a) Steady state and (b) time-resolved PL spectra of CdS nanorods and 30 wt% SV-1T-MoS₂/CdS composites at λ_{ex} = 405 nm. (c) Transient photo-current measurements and (d) EIS Nyquist plots of CdS nanorods, 30 wt% SV-1T-MoS₂/CdS and 30 wt% 1T-MoS₂/CdS composites. Reproduced with permission from ref. 93. Copyright (2020) American Chemical Society.

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Carbon-based materials

In addition to inorganic semiconductors, metal-free carbonbased materials have attracted a great deal of attention as affordable and sustainable candidates for applications in catalysis, especially when these materials are obtained from biomass or other renewable sources.¹⁰⁵ In this regard, graphitic carbon nitride (g-C₃N₄) is one of the most relevant examples of this family of materials and its applications span from heterogeneous organic synthesis to photo- and electrocatalysis, among others.^{106,107} When it comes to N₂ photo-fixation, a cutting-edge work of 2015 from Dong and co-workers demonstrated for the first time that g-C₃N₄ containing nitrogen vacancies could perform the photo-reduction of N₂ gas under visible light.¹⁰⁸ Not surprisingly, the study demonstrated that nitrogen vacancies are crucial to endow g-C₃N₄ with the capability to photo-reduce N2 thanks to the improved nitrogen adsorption and enhanced charge transfer efficiency.

Apart from the introduction of vacancies, doping with heteroatoms is a straightforward approach to expand the light absorption properties and modulate the redox potentials of $g-C_3N_4$.¹⁰⁹ In 2020, Liang and collaborators explored the possibility to merge the positive effects of both nitrogen vacancies and boron-doping in a graphitic carbon nitride.¹¹⁰ With that purpose, authors first obtained a defective carbon nitride (CN_r) through calcination of melamine followed by an annealing treatment under N₂ in presence of NaBH₄ (Fig. 6A). The asprepared boron-doped defective carbon nitride (BNUCN_x) displayed a remarkable NH₃ production rate of 435 μ mol g⁻¹ h⁻¹ under visible light. Both steady-state PL and EIS spectra and photo-current measurements revealed that the defective boron-doped carbon nitride featured a good charge separation efficiency and a low charge-transfer resistance that ultimately improved the catalytic activity (Fig. 6B and C). Furthermore,



Fig. 6 (a) Schematic illustration of the synthesis protocol for the preparation of boron-doped defective ultrathin carbon nitride (BNUCN_x). (b) Steady state PL and (c) EIS spectra of different samples. Reproduced with permission from ref. 110. Copyright (2020) Elsevier.

photo-catalysts, the use of tungsten oxides has recently sparked the interest of the scientific community. Although pristine WO₃ has shown a slight activity towards the N₂ photo-This article is licensed under a Creative Commons Attribution-NonCommercial 3.0 Unported Licence. reduction, the doping with Fe (0.25%) increased by 4 times the productivity up to 477 μ g gcat⁻¹ h⁻¹ thanks to the improved charge mobility and charge separation efficiency.98 As for the case of TiO₂-based catalysts, the introduction of oxygen vacancies or dopants to tungsten oxides is a straightforward method to enhance their catalytic activity. In 2022, Yang and pen Access Article. Published on 07 December 2023. Downloaded on 8/23/2025 6:12:03 AM. collaborators compared the catalytic activity of both 2D WO_{3-x} nanosheets and WO_{3-x} nanoparticles containing a high and

low content of oxygen vacancies, respectively.99 The introduction of oxygen vacancies in the tungsten oxide lattice featured a series of advantages including extended light absorption, improved charge carrier separation and enhanced N2 adsorption. Consequently, 2D WO_{3-x} nanosheets with abundant oxygen vacancies displayed a NH3 production of 82.41 µmol g_{cat}^{-1} h⁻¹ under irradiation with a visible LED (λ = 420 nm), which was almost 4 times higher compared to the sample with a low content of oxygen vacancies. The stability tests, however, were limited to only 5 cycles and authors did not provide any explanation to the observed deactivation of the catalyst upon consecutive uses. Zhang et al. studied the effect of Mo doping on the photo-catalytic activity of W18O49 nanowires and it was found that the Mo-W centers acted as active sites for N2 activation thanks to the polarization of chemisorbed nitrogen molecules.¹⁰⁰ Consequently, Mo-doped W₁₈O₄₉ nanowires displayed a 7-fold higher NH₃ production compared to pristine $W_{18}O_{49}$, thus achieving 195 µmol NH₃ g_{cat}^{-1} h⁻¹. In a related precedent, Zhang and co-workers demonstrated that oxygen vacancies displayed a dual effect to both induce the hydrogen spillover and promote N2 adsorption on Ru-decorated W₁₈O₄₉.¹⁰¹ Under irradiation, photo-generated electrons migrated to Ru NPs to activate and dissociate H2O in the form of H* while oxygen vacancies facilitated the migration of the H* species to W18O49. Eventually, H* species reacted with adsorbed N₂ to produce NH₃. Recycling experiments showed, however, that the catalyst lost its activity upon consecutive reuses. Interestingly, apart from tungsten, other oxides from the same group like molybdenum ones have been investigated as active materials for N₂ photo-fixation. Thanks to the presence of oxygen vacancies, defective molybdenum oxides featured plasmonic behavior that provided extended light absorption along the visible and NIR regions, thus exhibiting remarkable catalytic activities for N2 photo-reduction.^{102,103} More recently in 2023, Fu et al. also described the use of defective ZrO_{2-x} in combination with Ru NPs to create a metal/semiconductor heterojunction.¹⁰⁴ In this case, the presence of oxygen vacancies in the zirconia lattice induced a higher concentration of carriers, therefore providing more electrons for the N₂ reduction reaction. Furthermore, the formation of a Schottky barrier at the interface between the defective ZrO_{2-x} and the Ru NPs improved the charge separation and the effective injection of electrons from the semiconductor to the metal.

In addition to the above-discussed semiconductor-based

the introduction of N vacancies and B dopants led to a narrowing in the bandgap that enhanced the optical absorption properties of the final material. In the same spirit, other researchers have investigated the effect of the doping of $g-C_3N_4$ with diverse heteroatoms such as S or P.^{111,112}

As in the case of inorganic semiconductors, the construction of heterojunctions containing g-C₃N₄ can be a useful tool to refine its electronic and catalytic properties. Among all of the different types of heterojunctions, type-II heterojunctions are one of the most investigated systems. For instance, the groups of Zhang and Wu studied the combination of p-type Cu₂O and n-type C₃N₄ semiconductors to build a composite for N₂ photo-fixation.¹¹³ The strong electric field at the interface between both elements greatly enhanced the charge separation efficiency, with electrons and holes accumulating at g-C₃N₄ and Cu₂O, respectively. Another example of type-II heterojunction between g-C₃N₄ and ZrO₂ was reported recently by Mou et al.¹¹⁴ Upon light radiation, photo-excited electrons migrated from the CB of $g-C_3N_4$ to the CB of ZrO_2 , remaining the holes at g-C₃N₄. Thanks to the effective charge separation, the g-C₃N₄/ZrO₂ heterostructure displayed a NH₃ production rate as high as 1446 $\mu mol \; L^{-1} \; h^{-1},$ compared to the negligible activity of both elements separately. Apart from type-II heterojunctions, Z-scheme heterojunctions are also a very representative exponent of composite materials with improved catalytic efficiencies. In spite that Z-scheme heterostructures feature a distinct design compared to type-II heterojunctions, both architectures provide a spatial separation of electron and holes and the creation of well-defined redox active sites.¹¹⁵ Several examples of Z-scheme heterostructures for N2 photo-reduction include g-C₃N₄/Fe₂O₃, g-C₃N₄/Ga₂O₃ or B-doped g-C₃N₄/ WO₃.¹¹⁶⁻¹¹⁸ For a more comprehensive revision on g-C₃N₄based materials and heterostructures for photo-catalytic N2 reduction, readers are referred to excellent reviews covering this topic.115,119

Although g-C₃N₄ has been, by far, the most widely explored metal-free material, other carbon-based materials such as doped graphenes, reduced graphene oxides or carbon nanotubes have been investigated, alone or in the form of heterostructures, potential photo-catalysts for as NH_3 production.¹²⁰⁻¹²⁴ It is worth reminding that, especially for N containing photo-catalysts such as g-C₃N₄ or N-doped carbons, the incorporation of a set of control experiments including the use of isotopically labelled ¹⁵N₂, blank experiments working under N2-free atmosphere and/or in the absence of photo-catalyst is crucial to ensure the reliability of the reported results. Despite it is out of the scope of the present review, readers will find excellent guidelines for the accurate evaluation of the catalytic activity photo-fixation in N_2 experiments elsewhere.125-127

Metal-organic frameworks (MOFs)

In addition to commonly studied inorganic semiconductors and carbon-based materials, metal–organic frameworks are a novel class of materials that can be effectively used as photocatalysts due to their tunability, wide light harvesting properties and large surface area. Applications include photocatalytic water splitting, CO₂ reduction and organic photosynthesis, among others.^{128–130} Unfortunately, the study of metal–organic frameworks as photo-catalysts for ammonia synthesis is still in its initial stage.

To date, the greatest number of publications is devoted to transition metal-based metal-organic frameworks. Quite remarkable NH₃ production rates were achieved using iron-based MOFs.¹³¹ In the case of MIL-101(Fe), MIL-88(Fe) and MIL-100 (Fe), the ammonia yield reached 1007.1 μ mol g⁻¹ h⁻¹, 800.7 μ mol g⁻¹ h⁻¹ and 930.64 μ mol g⁻¹ h⁻¹, respectively (Fig. 7). The difference between isoreticular iron-based and chromium-based MIL-101, which exhibited no activity, was explored in the same study. Lower activation energy and higher electron density due to two additional filled d-orbitals were identified as fundamental reasons for such a different behavior.

However, Min Luo et al. showed that the incorporation of silver particles into the MIL-101(Cr) cavity allowed reaching an activity of 138.81 μ mol NH₃ g⁻¹ h⁻¹, which they attributed to the synergy of ligand-metal charge transfer (LMCT) and localized surface plasmon resonance effect (LSPR).¹³² Inspired by naturally occurring multi-metal cluster nitrogenases, Zhao et al. recently synthesized MIL-53(Fe²⁺/Fe³⁺), which displayed high nitrogen fixation activity as high as 306 μ mol NH₃ g⁻¹ h^{-1} .¹³³ It is important to note that the sample was irradiated only in the visible range above 420 nm, which means that it can potentially be used to convert solar energy. In a similar vein, Li and co-workers adopted a strategy to mimic the functionality of natural enzymes in synthetic materials.¹³⁴ In this case, the approach involved modifying the MIL-88A iron MOF with polyoxometalates, which acted as charge regulators and simulated the π^* -back-donating mechanism of nitrogenase. This modification resulted in a remarkable six-fold increase in the photocatalytic activity of the MOF.

Some Ti-based MOFs have been explored in photocatalytic reactions and have shown good nitrogen sorption capacity, thus standing for potential catalysts in the N₂ photo-fixation reaction. Huang *et al.* showed that a decrease in the optical band gap of NH₂-MIL-125(Ti) led to an increase in the photo-



Fig. 7 (a) Photo-catalytic nitrogen fixation activity of different MIL MOFs. Charge density difference of a N₂ molecule adsorbed at (b) MIL-101(Cr) and (c) MIL-101(Fe). Reproduced with permission from ref. 131. Copyright (2020) Elsevier.

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catalytic performance, as in the case of hydrogen evolution reaction.¹³⁵ However, this Ti-MOF, which exhibits one of the most impressive H₂ evolution activities, reached a very modest NH₃ production of 12.3 μ mol g⁻¹ h⁻¹. It should be mentioned, however, that the photocatalytic nitrogen reduction tests were tested under visible light and without a sacrificial agent. Interestingly, hafnium and zirconium-containing MOFs (UiO-66-Vis and U(0.5 Hf)) turned out to be ten times more active (256.6 μ mol NH₃ g⁻¹ h⁻¹ and 351 μ mol NH₃ g⁻¹ h⁻¹, respectively).^{136,137} In fact, a recent work indicated that Zrbased MOFs activity could be increased via the incorporation of Ca²⁺ dopants, resulting in a noteworthy increase in photocatalytic activity from 105 μ mol g⁻¹ h⁻¹ in the unmodified MOF to 222.2 µmol g⁻¹ h⁻¹ in the modified counterpart.¹³⁸ Authors suggested that the introduction of Ca²⁺ dopants promoted the redistribution of electrons around active Zr sites, thus reducing the activation energy of N2 triple bonds as elucidated through DFT calculations. Furthermore, the utilization of porphyrins and metalloporphyrins as ligands for stable Zrbased MOF represents an alternative approach to formulate new photocatalysts for NH₃ production.¹³⁹ Hence, using porphyrin ligands for MOF construction led to NH₃ production rates of 776.24 μ mol g⁻¹ h⁻¹, while replacing the ligands with metalloporphyrin resulted in a significant boost in photocatalytic activity, reaching 1502.51 μ mol g⁻¹ h⁻¹.

Besides transition metal-based, attention should be paid to lanthanide-based MOFs since Hu et al. developed two radicalcontaining gadolinium-based MOFs.¹⁴⁰ Purple crystals of Gd-IHEP-7 could be obtained using a solvothermal technique. The procedure for obtaining Gd-IHEP-8 included heating Gd-IHEP-7 to 120 °C in the air when single-crystal-to-single-crystal transformation occurred. The rearrangement of the Gd³⁺ coordination environment was accompanied by an increase in photocatalytic nitrogen fixation activity (up to 220 µmol NH₃ g^{-1} h⁻¹). Authors suggested that the factor responsible for the higher activity of Gd-IHEP-7 was the calculated change in the stability of the intermediates. Interestingly, Gd-IHEP-7 and Gd-IHEP-8, unlike most other viologen-based compounds, were stable in air and water due to radical-radical interaction. Compared to Gd-based MOFs, deficient activity (34.2 µmol $NH_3 g^{-1} h^{-1}$) was shown in the case of cerium-based MOF-76 (Ce) even using full-spectrum light of Xe-lamp.¹⁴¹

Photo-thermal catalytic NH₃ synthesis

As we have earlier discussed in this review, the thermo-catalytic NH_3 synthesis route is constrained by its harsh operative conditions whereas the photo-catalytic NH_3 production is extremely hampered by its poor catalytic efficiency and low selectivity. In order to overcome these limitations, a smart approach would be the combination of these two strategies into a single process. In this spirit, photo-thermal catalysis arises from the synergistic incorporation of thermal and nonthermal contributions of light and features exceptional advantages compared by its individual counterparts. In fact, this cooperative effect has implications in both reaction thermodynamics and kinetics. For instance, from the thermodynamic point of view, non-thermal effects can reduce the energetic barriers of certain reaction steps. The reduction in the activation energy can derive from charge transfer events to the surface of the catalyst that alter the acidity/basicity of the active sites. Besides, the energy transfer to certain vibrational modes of the transition state can also favour bond elongation, thus decreasing the energy requirements of the process.¹⁴² In this regard, Song and collaborators recently demonstrated the potential of non-thermal effects to adjust the thermodynamics of dry reforming reaction.¹⁴³ Authors demonstrated that photo-induced hot electrons from plasmonic Pt and Au nanoparticles could be injected to adsorbed CO₂ and CH₄ molecules, thereby weakening the C=O and C-H bonds and facilitating bond dissociation. In this case, the apparent activation energy under radiation decreased from 241 to 171 kJ mol⁻¹, suggesting a change in the energetics of the reaction. In addition to this, the presence of non-thermal effects opens up the possibility to activate specific chemical bonds that unlock certain reaction pathways that are inaccessible under pure thermal conditions, thus facilitating tailored selectivity patterns or alternative low-energy reaction intermediates. For instance, the groups of Liu and Everitt demonstrated that it was possible to adjust product selectivity in the photo-thermal CO₂ hydrogenation by plasmonic Rh nanoparticles.¹⁴⁴ The use of blue or UV LEDs to perform the reaction facilitated the CH₄ production with selectivities up to 98%, while the reaction under dark conditions displayed comparable amounts of CH₄ and CO. Theoretical calculations demonstrated that the injection of photo-induced hot electrons from Rh nanoparticles to anti-bonding orbitals of adsorbed intermediates played a crucial role to determine the selectivity patterns. Curiously, the presence of hot carriers can not only tune the reaction selectivity towards the desired products, but also prevent the formation of detrimental surface species. Indeed, several recent studies have shown the hot carriers' potential in mitigating the formation of unwanted byproducts, especially in the context of photo-thermal dry reforming of methane.145,146

When it comes to thermal effects, the photo-induced heat at the nanoscale provides high temperatures at the active sites in a more efficient way compared to traditional thermo-catalysis. These elevated temperatures translate into catalytic activities that are several orders of magnitude above those obtained with conventional photo-catalysis, mainly due to both the improved reaction kinetics and energy-mass transfer. In addition to this, thermal effects favor the desorption of products or poisonous species from the catalyst surface, thereby enhancing the activity and stability of the catalysts.¹⁴⁷ Last but not least, while in photo-catalytic systems the use of light is limited by the band gap and the position of both conduction and valence bands of the photocatalysts, photo-thermal catalysis takes advantage of the full solar spectrum, even the lowenergy photons from the visible and IR that are typically unable to trigger photo-chemical reactions. A comprehensive discussion on the fundamentals of photo-thermal catalysis and the benefits from the synergies underlying the photothermal phenomenon lies beyond the scope of this review, so readers are referred to specific works on this topic.^{148–151}

Photo-thermal catalysis can be considered a relatively recent sub-field of catalysis and, in fact, only a few research groups have explored the combination of light and heat to perform the HB process under mild reaction conditions. For instance, García and co-workers lately reported the use of Cspromoted ruthenium NPs supported on strontium titanate (Cs_vRu_x(a)ST) as photo-catalyst for the NH₃ synthesis reaction under 350 °C and ambient pressure.¹⁵² All the tests were performed using a continuous-flow set up equipped with a heating ribbon and a thermocouple to control the temperature. Under optimal Cs and Ru loading, the photo-catalyst achieved an impressive NH₃ production rate in the order of 3.6 mmol g^{-1} h⁻¹, outperforming by a 68% a control experiment in the dark at the same temperature. Authors suggested that the remarkable catalytic activity derived from the combination of both photo-induced hot carrier and local thermal effect mechanisms taking place at the irradiated Ru NPs. Altogether, both contributions facilitated the activation of N₂ and its further hydrogenation (Fig. 8A). Interestingly, this work also showed that the choice of the support for Ru NPs was not trivial at all, as authors demonstrated that both the basicity and low thermal conductivity of strontium titanate were key to improve the catalytic performance (Fig. 8B). In 2023, the same group explored the MOF-derived approach to obtain Cs-promoted Ru clusters supported on ZrO2 as photocatalyst for the photo-thermal NH3 synthesis.153 In this case, the authors impregnated the UiO-66 MOF with Ru and Cs precursors and calcined the resulting material in air at 550 °C. Under these conditions, the MOF served as sacrificial template to obtain ZrO2 decorated with Cs and Ru clusters. The as-obtained catalyst exhibited a notable activity under 350 °C and 1 sun radiation, achieving a NH₃ production rate of 5.1 mmol g^{-1} h⁻¹. Analogous materials prepared by impregnation of commercial ZrO₂ displayed negligible catalytic activity, thus stressing the potential of the MOF-derived strategy to formulate efficient catalysts that are not achievable by conventional synthetic routes. Mechanistic experiments revealed that both thermal and nonthermal effects operated simultaneously to enhance the catalytic activity. On the one hand, Ru sites absorbed visible and NIR light and locally increased the temperature of the catalyst owing to plasmon excitation. On the other hand, plasmon excitation could also generate hot electrons that could be injected into anti-bonding orbitals of adsorbed N_2 .

Nevertheless, for most of the photo-thermal systems, it is still extremely challenging to provide an accurate evaluation of the specific role of each of the contributions to the overall catalytic activity. Zhang and collaborators recently addressed this issue employing a model catalyst based on Ru on carbon.¹⁵⁴ Authors first estimated a local temperature of the catalyst (Treaction) based on the thermodynamic equilibrium conversion. By introducing this T_{reaction} into the Arrhenius equation under thermo-catalytic conditions, authors calculated the actual reaction rate derived from pure thermal effects $(R_{\text{temperature}})$. Once the thermal component was isolated, it was possible to extract the non-thermal contribution (R_{electron}) by subtracting the thermal component to the total photo-thermal rate (R_{PTC}). Interestingly, authors found that $R_{electron}$ transitioned from a linear to superlinear relationship with the light intensity, thus confirming the presence of non-thermal effects in the system. In fact, under 5.3 W cm^{-2} light intensity and 350 °C the hot-electron contribution accounted for a 73.6% of the total NH₃ production rate. In situ diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS) measurements also revealed the presence of *NH_r species under light radiation at 230 °C, while no intermediates were detected under dark conditions at temperatures below 260 °C. Based on these observations and additional DFT calculations, authors concluded that light could promote the NH3 synthesis reaction at lower temperatures thanks to the injection of photo-induced hot-electrons into unoccupied anti-bonding orbitals of adsorbed N_2 , thus facilitating the cleavage of the N \equiv N triple bond. Overall, this study demonstrated that it is possible to differentiate and quantify both the thermal and non-thermal contributions of photo-thermal systems using Le Chatelier's principle to determine the actual reaction temperature.

As commented before, local heat generation at the nanoscale is considered one of the main advantages of photo-



Fig. 8 (a) Proposed mechanism for the photo-thermal NH₃ synthesis by $C_{S_y}Ru_x@ST$ catalyst: (i) N₂ adsorption and activation, (ii) N₂ dissociation and formation of intermediate species, (iii) hydrogenation of activated nitrogen species, and (iv) NH₃ formation and desorption. (b) NH₃ production rate using different supports. Dark bars correspond to experiments under light irradiation and red bars correspond to dark conditions. Reproduced with permission from ref. 152. Copyright (2022) American Chemical Society.

thermal catalysis. In 2019, Mao and co-workers proposed a dual-temperature-zone catalyst to tentatively trick the thermodynamic equilibrium limit while using sunlight to promote the NH₃ synthesis.¹⁵⁵ The idea here was to locate the N₂ dissociation in a "local high-temperature" center, where the nitrogen dissociation takes place, and a local low-temperature center for the N-hydrogenation in order to physically separate these two elemental steps of ammonia synthesis. Thanks to the afore-mentioned capability of photo-thermal catalysis to create hot spots, authors succeeded in designing a catalytic system to provide this non-uniform heat distribution, something otherwise extremely complex to achieve under pure thermal conditions. In this case, plasmonic Fe nanostructures acted as nano-heaters to perform the N2 dissociation step while subsequent N-H hydrogenation took place at the "cooler" surface of defective $TiO_{2-x}H_{y}$ owing to its low thermal conductivity. When irradiated at 102 sun, the catalyst reached a temperature as high as 495 °C, thus achieving a NH₃ concentration of 1939 ppm. This concentration was above the thermodynamic equilibrium value of NH₃ synthesis at 495 °C and 1 atm (1249 ppm) and even outperformed the value obtained by commercial-wüstite-based Fe catalyst. The presence of the dual-temperature-zone between Fe and TiO2-xHy was key to allocate the elementary steps of NH₃ synthesis into two spatially separated catalytic sites, thus surpassing the thermodynamic limit of the reaction (Fig. 9).

Despite the potential of the aforementioned dual-temperature-zone catalyst, some authors have called the attention to the existence of important thermal gradients due to the shallow penetration of light, especially in the case of thick catalyst beds. This situation would, inevitably, confine the effect of photo-thermal catalysis to the illuminated surface region. In their work of 2019, Li et al. made the most of this light-induced thermal gradient to boost the NH₃ production rate using a conventional cesium-promoted Ru on magnesium

oxide (Cs-Ru/MgO) catalyst.¹⁵⁶ Authors demonstrated that the non-isothermal gradient between the surface and the bottom of the catalyst bed favored high temperatures on the surface, where the N₂ scission took place, while the cooler temperatures from the bottom preserved the NH₃ yield. In addition to this, authors found that not only the presence but also the sign of the thermal gradient was critical to enhance the catalytic activity. Under a negative gradient, thermophoretic forces place parallel to the flow of reactants, thus removing NH₃ products from the hottest region and circumventing the undesirable reverse reaction of NH₃ decomposition. However, under the same absolute value, a positive gradient displayed much lower reaction rates and product yields. Overall, these results evidenced the potential of light-induced thermal gradients to act as thermodynamic pumps to shift the equilibrium and improve the catalytic activity of photo-thermal systems (Fig. 10).

An appealing example of the potential of photo-thermal catalysis to unlock new reaction pathways was reported in 2018 by the group of Zhang. In their work, authors described the use of K-promoted Ru NPs supported on disordered $TiO_{2-x}H_x$ (K/ $Ru/TiO_{2-x}H_x$) as photo-thermal catalyst for the solar-driven NH_3 synthesis.¹⁵⁷ The K/Ru/TiO_{2-x}H_x catalyst featured a series of advantages that have been described along previous sections of this review such as the introduction of oxygen vacancies to expand the light absorption properties of TiO_2 or the use of alkaline promoters to tune the electronic environment of Ru NPs and promote N₂ activation. However, the most striking observation in this work was the presence of two simultaneous reaction mechanisms within the $K/Ru/TiO_{2-x}H_x$ catalyst. Hence, activated N2 could react not only with H atoms on the Ru surface, but also with the H atoms incorporated in the $TiO_{2-x}H_x$ support. Remarkably, this latter pathway was notably different from those reported to date and it could even occur under room temperature, thus suggesting a mechanism



Fig. 9 (A) NH₃ concentration under combined solar irradiation and external heating by TiO_{2-x}H_y/Fe at a constant temperature of 495 °C. (B) NH₃ concentration as a function of the local temperature difference (LTD). (C) Comparison of equilibrium limits at the temperature of "cool" $TiO_{2-x}H_{y}$ and "hot" Fe for photo-thermal ammonia synthesis on $TiO_{2-x}H_v/Fe$ (1 atm). Reproduced with permission from ref. 155. Copyright (2019) Elsevier.

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Fig. 10 (a) Illustration of the thermal gradient generated by the photo-thermal heating of the irradiated catalyst surface. Effect of light-induced thermal gradients on NH₃ synthesis. (b) Measured dark thermal (red squares) and heated white light illumination NH₃ synthesis rates as a function of equivalent temperature. Circled data point indicates the light-only condition with no external heating applied. (c) NH₃ synthesis rates under dark and additionally illuminated conditions using a blue LED. Reproduced with permission from ref. 156. Copyright (2019) American Chemical Society.

with a considerably low energy barrier. In fact, authors calculated the activation energies of both thermal and photothermal reactions, achieving 86 and 65 kJ mol⁻¹, respectively. This decrease in the activation energy was explained by the presence of both plasmon-induced confined magnetic fields and heat generation at the vicinity of Ru NPs. In general, this study opened a new venue for the development of alternative NH₃ synthesis routes driven by solar light thanks to the combination of low-energy reaction pathways and the confined energy of plasmonic NPs.

Catalyst engineering and optimization

As mentioned before, photo-thermal catalysts benefit from a synergistic contribution of both photo and thermal components of light. To further intensify the photo-thermal effect and enhance production rates, researchers have focused on improving the light-to-heat conversion by optimizing size and shape of active sites, adjusting the porosity and nature of support or through structural engineering, among other strategies.148

For plasmonic materials, size and shape of nanoparticles affect not only the capability of charge carrier generation, but also the thermal properties.¹⁵⁸ Theoretical calculations indicate that smaller nanoparticles produce electron-hole pairs with higher energy in comparison to larger nanoparticles.¹⁵⁹ This characteristic holds significant importance in the context of photo-thermal catalysis, as it enhances the likelihood of populating acceptor states and overcoming Schottky barriers. However, a small particle size may lead to inefficient charge carrier separation due to spatial confinement. In addition, larger particles also show a red-shift on the absorption capability of the catalyst, unlocking the use of visible and IR light ranges.¹⁶⁰ Another point is that the plasmon-heating effect in metallic nanoparticles scales quadratically with their radius.¹⁶¹ Therefore, the engineering of the ideal catalyst relies on the perfect balance of the size. As for the morphology, geometries that display strong and inhomogeneous electric fields associated with confinement effects contribute to high-energy hot electrons and holes.¹⁶² The opposite is also true, geometries without defects and homogeneous electric fields, like nanospheres, exhibit a less efficient production of hot carriers.

Another possibility is to use secondary light-absorbing elements, whereby a catalytically active metal is coupled with another material able to absorb light in a wider range and generate photo-carriers and/or heat.¹⁶³⁻¹⁶⁵ Core-shell structures have also received great attention, as the idea is to mimic a nanoreactor, confining the heat in the core, increasing the temperature precisely where the active sites are located.¹⁶⁶ Still in this context, MOFs can also be used as sacrificial templates to obtain carbon-based materials featuring broadband light absorption in the visible and IR light ranges.¹¹ The combination of carbonaceous supports with highly dispersed metal nanoparticles makes MOF-derived materials a very attractive family of catalysts for photo-thermal applications.¹⁶⁷

Conclusions and outlook

Due to its widespread use as building block for the production of fertilizers, NH₃ is considered one of the key chemicals in our society. Despite the technological maturity of the Haber-Bosch process, there is still room for improvement. With this objective in mind, ongoing research is being devoted not only to find catalysts able to operate under milder conditions without losing productivity, but also to the development of alternative synthetic routes. One of the possibilities in this regard is the integration of light into the NH₃ synthesis process to achieve photo-catalytic N2 fixation routes powered by sunlight. However, to date the reported efficiencies remain far from those required to compete with the conventional HB approach. In an effort to overcome the limitations of both thermo- and photo-catalytic approaches, photo-thermal catalysis has demonstrated potential to achieve high activities at moderate reaction conditions through the synergistic combination of thermal and non-thermal contributions of sunlight.

In this review, we have first provided an overview of the main reaction pathways for NH₃ synthesis. Next, we have reviewed recent strategies for the formulation of novel catalysts for the thermo-catalytic NH₃ production able to operate at

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milder conditions. In this regard, catalysts based on Ru or Co are considered promising candidates although their higher costs compared to inexpensive Fe-based catalysts still represent a major pitfall for their wide implementation. Later, we have discussed potential methodologies to improve the catalytic performance of photo-catalytic systems for N₂ photo-fixation, mainly the introduction of vacancies, defects or the construction of heterojunctions. In the last section of the review, we highlighted the significance of photo-thermal catalysis to surpass the limitations of its individual counterparts thanks to the synergistic integration of thermal and non-thermal effects. This cooperative effect can improve reaction rates, create positive thermal gradients or even trigger alternative lower-energy reaction pathways.

Although the technological advantages are evident, to date there has been little discussion on the benefits of photothermal catalysis in terms of sustainability. A rigorous analysis in this regard would require appropriate metrics to evaluate the environmental impact of photo-thermal catalysis but, unfortunately, these figures of merit have not been perfectly established in the field. Ozin and collaborators recently tried to address this issue and proposed a series of equations to assess the CO₂ footprint of thermal and photo-thermal CO₂ hydrogenation reactions.¹⁶⁸ These equations considered multiple parameters related to the photo-thermal process such as CO2 emissions per kWh of electricity, CO2 produced per mol of H₂, total irradiation time or the power consumption of lamps, pumps, etc., among others. Based on their model, authors found that the source of radiation was critical to reduce the CO₂ emissions associated with the photo-thermal process. In other words, energy-saving lamps with low power are preferred, as the net CO₂ emission rate scales up with the power of the lamp. In the best-case scenario, the direct use of sunlight instead of electricity would further reduce the carbon footprint or even achieve negative CO₂ emissions, although this strategy would require additional technological development. Equations also indicated that reactor configuration plays an important role, as flow reactors displayed a higher potential to reduce CO2 emissions compared to their batch counterparts. Assuming a CO2-free source of H2, authors concluded that photo-thermal catalysis has the potential to achieve net CO₂ emission rates lower than traditional thermal catalysis, especially in the case of flow reactors coupled with energy-efficient lamps or directly powered with solar radiation. Overall, this pioneer work reinforced the prospects of the photo-thermal approach in terms of its CO₂ footprint and set the basis for subsequent studies focused on other chemistries of interest beyond CO₂ hydrogenation. In addition to this, assessing the economic feasibility of the entire process is imperative to ensure a realistic scale-up of this technology. While performing an accurate economic analysis of the photothermal HB process is highly challenging due to the immaturity of the field and the lack of actual data from large-scale prototypes, we would like to share some considerations for future estimations. Based on available techno-economic analyses, the main contributors to the total energy consumption in blue

ammonia production include the ammonia separation refrigerant, cooling water and gas used to provide heat to the reactors.⁶ In the case of green ammonia, electricity for compressors, water for general cooling and refrigerant for separation purposes are the major contributors to the energy requirements of the operation. With this in mind, the photothermal approach could potentially reduce the energy demand (and consequently, the cost) of the process by harnessing natural sunlight instead of relying on electricity or gas to power the HB reactors. Furthermore, considering that photothermal processes can operate at lower temperatures than pure thermal systems, the energy costs could be further reduced. The techno-economic analysis should also include the cost of the photo-thermal reactor based on parameters such as the average solar intensity, illumination hours, irradiated area, type of solar collector, amount and lifetime of the photocatalyst, etc. Nevertheless, previous works have revealed that the major capital costs for NH₃ synthesis primarily stem from feed costs rather than energy costs. This implies that, ultimately, the hydrogen or gas market prices will play a pivotal role to determine the profitability of any potential route for NH₃ production.⁶

Despite its promising future, if photo-thermal catalysis is to lead a paradigm shift in the field of heterogeneous catalysis and compete with long-established thermo-catalytic processes, a series of challenges need to be addressed. For instance, a deeper understanding of the synergies between hot carriers and thermal effects at the fundamental level would be desirable in order to fully comprehend the mechanisms underlying photo-thermal reactions. To this end, in situ/in operando techniques (X-ray diffraction, electron microscopy, XPS, Fouriertransformed IR, Raman, UV-vis absorption, etc) could help not only to assess the contribution of thermal and nonthermal effects on the catalytic performance, but also to provide full information on the catalytic sites and transitional species present under reaction conditions.169-172 In this regard, computational methods can be also interesting tools to disentangle not only the reaction pathways but also the intermediate states involved.

An accurate measurement of the local temperature is crucial in photo-thermal catalysis as imprecise temperature records can lead to misinterpretations of the thermal contribution.¹⁷³ For this reason, and especially in the case of thick catalyst beds, one or more sets of thermocouples should be utilized to ensure a correct monitoring of the temperature across the catalyst. Ideally, these measurements should be complemented with additional temperature registrations using IR cameras or temperature-responsive photoluminescence probes.

From the point of view of scalability, photo-thermal catalysis requires adequate photo-reactors capable of harvesting sunlight and channeling it directly to the catalyst. Based on lessons learned from a mature technology like the solar thermal energy, solar concentrators or condenser lenses could serve as elements to both harness and concentrate sunlight.^{174–176} For instance, the groups of Steinfeld and

Furler recently developed a solar fuel system that enables the production of syngas from water and CO₂ using concentrated solar energy.¹⁷⁷ The solar redox unit consisted of a sun-tracking parabolic concentrator coupled with a secondary reflector. Under normal solar radiation conditions, the system could reach a peak flux concentration above 5000 suns and an average flux of approximately 2700 suns. Many reaction systems, however, require less severe solar flux concentrations in the order of 50 to 100 suns. In these cases, the use of parabolic troughs to reflect and concentrate sunlight can be a more affordable and practical option. These systems use sets of concave mirrors to concentrate light into its focal line, achieving temperatures in the range of 500-700 K.178 Alternatively, high-power LEDs or incandescent light bulbs could also act as irradiation sources in conjunction or not with solar light radiation.¹⁷⁹ In addition to that, further considerations on heat management should be taken into account when designing new photo-reactors, as thermal management has proved to be critical if one wants to maximize the benefits of the photothermal effect. Moreover, the use of batch reactors at the laboratory scale should be progressively shifted to continuousflow reactors that are better suited for potentially scalable systems at the industrial scale. In any case, it is necessary to perform extensive research in reactor design and engineering to piece together all these considerations and develop suitable photo-reactors for applications in photo-thermal catalysis.

As we have discussed in detail along this review, the HB process is one of the best examples of energy integration in the chemical industry. In this particular case, for photothermal catalysis to be successful, catalysts with excellent light absorption and light-to-heat conversion properties are crucial. Examples of this kind of materials include carbon-based solids, defective semiconductors or transition metal carbides and nitrides. These materials, alone or in combination with adequate metallic active sites, can provide exceptional electronic and thermal effects to synergistically boost the production of NH₃ from N₂ and H₂. Along with absorption properties, stability is equally important when assessing the potential application of a catalyst. Conventional stability cycles usually cover only a few hours of activity, however, this limited timespan is insufficient to gain any insight on the real robustness of the system, so long-term stability tests in the order of tens of hours are imperative.

The present review has described alternative NH₃ synthesis routes based on the use of solar light as source of energy, namely photo-catalysis and photo-thermal catalysis. However, it is worth reminding that the use of electro-catalysis to drive the N₂ reduction reaction has recently gained considerable interest. Compared to the traditional HB process, the possibility to produce NH₃ using renewable electricity would allow a decentralized route for N₂ fixation with lower CO₂ emissions and improved energy efficiency. However, in spite of these promising features, the viability and reliability of most of the available reports on electrocatalytic N₂ reduction is still under debate, given the low amount of NH₃ produced (in the orders of nmol to μ mol) and the presence of multiple impurities that may lead to false positives.^{180–182} For instance, a recent work from Chen and collaborators demonstrated the presence of nitrogen-containing compounds such as NO_3^- or NO_2^- in many commercial catalysts for ammonia electro-synthesis, thus stressing the necessity to develop specific protocols to ensure reliable and reproducible results.¹⁸³ In any case, electrochemical NH₃ synthesis has become a very dynamic field of research and we certainly believe it will play a decisive role in the future NH₃ production systems beyond the HB process. For these reasons, and despite being out of the scope of this work, those readers interested in the topic are referred to specific reviews available in the literature.^{181,184,185}

In summary, photo-thermal catalysis has proved to hold great promise as a viable route to drive the N₂ hydrogenation to NH₃ using solar light as energy source. Apart from its advantages in terms of energy efficiency and operational costs, a potential niche for the photo-thermal NH₃ synthesis would be the distributed production of fertilizers. Conventional HB facilities tend to be localized close to the natural gas supply points and require daily productions in the range of thousands of metric tons of NH₃ to be economically feasible.¹⁸⁶ These facts illustrate some of the constraints of the HB process, particularly the heavy dependence on fossil fuels and a certain inflexibility in the scale.⁴ On the contrary, photo-thermal NH₃ synthesis represents an opportunity for the deployment of small-scale delocalized NH₃ plants powered by renewable energy. This on-site production of ammonia would increase the access to fertilizers, especially in remote or isolated areas, promoting their economic and social growth. Interestingly, besides the well-known application of NH₃ as precursor for the manufacture of fertilizers, in recent years NH3 has been considered as an excellent H₂ carrier owing to its high volumetric and gravimetric energy density and well-established distribution infrastructure.¹⁸⁷ Nevertheless, the decomposition of NH₃ to obtain N₂ and H₂ is a highly endothermic reaction that requires elevated temperatures in the range of 500-600 °C to achieve significant conversion rates, thus posing a considerable penalty to the overall efficiency. Given its ability to reach elevated local temperatures, photothermal catalysis has the potential to drive the light-induced cracking of NH₃ under milder operating conditions, thus lowering the cost and improving the efficiency of the whole process, in a similar way to the NH3 synthesis reaction. Within the context of hydrogen economy, the development of such a light-mediated NH3 synthesis-decomposition cycle would be considered a major breakthrough with multiple implications in the decarbonization of a wide variety of areas including transportation or the industrial sector. All in all, we hope that this review will stimulate the progress of new photo-thermal systems focused not only on the chemistry of NH3 but also on other commodities of industrial interest.

Author contributions

Conceptualization: D.M. and J.G. Writing – original draft: D. M., A.S., M. Z. Writing – review and editing: J.G.

Conflicts of interest

There are no conflicts to declare.

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