Transforming nano metal non selective particulates into chemoselective catalysts for hydrogenation of substituted nitrobenzenes

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ABSTRACT. The hydrogenation of nitro compounds is an industrial process that has experienced a renovated interest in the last ten years due to the discovery of highly selective and environmentally friendly solid catalysts. Particularly, the performance of chemoselective reactions in the presence of very sensitive groups such as double and triple CC bonds, with H₂ as reductant and no soluble additives needed, had been elusive for decades. The discovery that gold nanoparticles on solid supports could carry out such a reaction very selectively invigorated this area of research and claimed gold as an outstanding catalyst beyond oxidation processes. Subsequent work, devoted to understand how gold catalysts operate, established strong basis for the design of more efficient materials and the aperture of new routes for the synthesis of nitro derivatives. Here, we present three generations of materials that allowed improving the performance of the original gold catalysts. The relatively low activity of the initial Au/TiO₂ catalysts could be first boosted, without practical loss of selectivity, by the design of a material that incorporated two catalytic functions on the support: small amounts of platinum to enhance H₂ dissociation, and a greater amount of gold to activate the –NO₂ group. Later, we learnt how to control the catalytic structures and induce chemoselectivity to traditionally unselective metals such as platinum, ruthenium and nickel nanoparticles. Recently, Fe₃O₄ nanoparticles surrounded by a nitrogen-doped carbon layer have erupted as a promising alternative. A remarkable outcome from all that work is that the final pool of catalytic alternatives has been markedly expanded. Diversity is important because different solutions may open new gates to different catalytic processes, and we summarize here how the scope of new reactions and products could be expanded by means of properly designed metal catalysts in where support and metal work in a concerted way to direct the reaction toward the desired product. For example, whereas Au/TiO₂ is a chemoselective catalyst that drives the reaction efficiently to the fully reduced reaction product (aniline), the reaction could be tuned to obtain azocompounds in high yields by using nanoparticulated ceria to support the gold nanoparticles. On the other hand, whereas nitrobenzenes and aldehydes react in H₂ to afford imines in the presence of Au/TiO₂, the product distribution can be switched toward a more oxidized condensation product (a nitrone) using a chemoselective Pt/C catalyst, or to produce cyclohexanone oxime directly from nitrobenzene by means of supported Au and Pd metal catalysts. These and other examples represent some notable achievements, possibly, just the tip of the iceberg.

1. Introduction

Nitrogen is found in a variety of chemicals with industrial interest, within the fields of polymers, fertilizers, dyes, or biologically active compounds. Nitrogen is often integrated into these chemicals from NH₃ (ammonolysis) or HNO₃ (nitration) at the earliest reaction steps, and the resultant amines or nitro compounds, respectively, are primary building blocks in organic synthesis. The nitration route is particularly relevant to obtain nitroaromatic compounds and short nitroalkanes, because the selectivity toward the mono nitrated products is high following standard industrial protocols. From those, the production of amines by hydrogenation of the –NO₂ group is a key transformation that has focused the attention of researchers for more than a century.

In the last decade, a renewed interest in the hydrogenation of nitro compounds has awakened (Figure
i) due to the discovery of metal catalysts that overcome the following long-lasting problems:

- It is difficult to reduce selectively nitro groups when other sensitive functions are present (the latter must often be retained to preserve products with high synthetic value).
- The preceding holds particularly true when H₂ is the reducing agent (and H₂ is the preferred industrial choice).
- It is difficult to control the nitro reduction to afford selectively the fully hydrogenated amine or, otherwise, a particular reduction intermediate, as desired.
- Getting high activity and selectivity is challenging, since increasing the activity often causes the selectivity to decrease.
- It is not trivial to accomplish all the above through an environmentally sustainable solid catalyst that does not require the use of soluble, often toxic, additives.

We want to stress that reactions performed under hydrogen transfer conditions (using hydrazine, formic acid, or NaBH₄ for instance) often result in a greater chemoselectivity toward the nitro reduction than when H₂ is used directly. For instance, Pd/C, which is typically unselective for the hydrogenation of nitroalkenes with H₂ (leading to aminoalkanes), catalyzes the formation of oximes in high yields with ammonium formate. Conversely, catalysts that are selective with H₂ are generally also selective under hydrogen transfer conditions. Some of these reducing agents are still common in laboratory-scale organic syntheses, but industry prefers H₂ because it is readily available, cheap, and generates water as the reduction by-product.

Recently, CO/H₂O mixtures could be used to reduce nitro compounds with gold catalysts. On the other hand, when referring to the chemoselective reduction of nitro compounds, the range of possibilities is broad, as it is the number of functional groups that may accompany the nitro group within the molecule. The accomplishment of high chemoselectivity is increasingly challenging as the reducibility of the competing functional group increases. For example, it is usually unnecessary to control the catalytic structures for achieving high selectivity with alky1, hydroxy or alkoxy substituents, and most common industrial catalysts can be used upon proper selection of the reaction conditions. Similarly, halonitrobenzenes, and specially monosubstituted chloronitrobenzenes, may fall short for the identification of catalysts with genuine selectivity features, since an extensive list of catalysts has been shown to work suitably with those compounds. While this is consistent with the fact that chlorine is a rather poor leaving group, one can surprisingly see publications showing “first time” high selectivity for the hydrogenation of chloronitrobenzenes. In contrast, the number of catalysts that can hydrogenate nitro compounds selectively in the presence of C=C, C≡C, C≡N or C=O bonds, and that fulfill the series of requirements depicted above, is very limited. C=C and C≡C bonds are considered to be the biggest challenge, and most catalysts that are selective for the hydrogenation of chloronitrobenzenes fail when the substrate incorporates an olefinic group. Au/SiO₂ catalysts are good examples, since they are able to hydrogenate successfully chlorinated nitrobenzenes, nitrophenols, nitrotoluenes and alkoxy nitrobenzenes, but they are unselective for the hydrogenation of 3-nitrostyrene. To reach chemoselectivity in the presence of C=C and C≡C bonds, Blaser et al used Pb or H₃PO₄ to tweak the performance of supported platinum nanoparticles, and cleverly added vanadium and iron salts to avoid the accumulation of partially reduced nitro compounds. Thinking in a more sustainable process, we developed a series of alternatives that do not require toxic additives in solution to achieve an optimal, broadband performance, and today rather general chemoselective catalysts can be accomplished with gold, platinum, ruthenium, nickel and, more recently, iron, silver, and rhodium. Here, we will summarize our contribution to the various generations of heterogeneous catalysts developed in the last ten years for the chemoselective hydrogenation of substituted nitrocompounds, and a series new reaction processes that were derived from them to synthesize useful organic precursors.

2. First Generation: Gold Catalysts

2.1. Discovery of Chemoselective Gold Catalysts.

We found that nitrobenzene was being reduced to aniline at 423 K in the presence of styrene, dimethylformamide (DMF) and Au/TiO₂. It turned out that, in the presence of
small amounts of water within the reactants, hydrolysis of DMF to formic acid (and dimethylamine) occurred, with the former acting as a hydrogen donor. Interestingly, styrene in the solution remained intact. When we used formic acid directly as the reducing agent, at a lower temperature (313 K), it was confirmed that Au/TiO2 can hydrogenate selectively nitrocompounds without altering C=C, C=O or C≡N substituents. In contrast, conventional platinum or palladium supported nanoparticles led to mixtures of products in where each functional group was indistinctly reduced. At that point, the question was if gold could perform that chemoselective hydrogenation with H2 instead of formic acid as the reducing agent. The designed isotopic and kinetic experiments clearly showed that Au/TiO2 (and Au/Fe2O3) were not only able to dissociate H2 for the –NO2 hydrogenation, but that anilines incorporating a variety of substituents (including C=C and C≡C bonds) could be obtained in high yields from the corresponding nitro compounds (Figure 2).

2.2. Elucidation of the Catalyst Specificity. To understand the mode of action of gold catalysts, kinetic measurements, IR spectroscopy, and DFT theory were combined to elucidate how molecules such as nitrobenzene, styrene (or mixtures of them), and nitrostyrene interact with the surface of various (supported and unsupported) gold catalysts. The in silico calculations suggested that both C=C and –NO2 groups interact similarly (weakly) with the surface of the (bare) gold nanoparticles, leading to unselective catalysis. In agreement with the theoretical calculations,
Figure 2. Scope of substituted anilines synthesized by hydrogenation of the corresponding nitro compounds with metal nanoparticles supported on TiO2. Yields (%) are displayed. To achieve optimal performance Pt/TiO2, Ru/TiO2, and Ni/TiO2 must be reduced in H2 at 723 K before reaction to promote a Strong Metal-Support Interaction. See experimental details in 14,27 [a] Low yield due to low catalytic activity. [b] Loss of yield due to hydrolysis of the nitrile group. [c] Unpublished results from our laboratory.

2.3. The Nitro into Amine Transformation Path on Supported Gold Catalysts. Another interesting feature of Au/TiO2 catalysts is that aniline is practically the only product detected in the liquid phase during the hydrogenation of the nitrocompound, independently of the conversion.16 The Au/TiO2 catalyst, thus, avoids the accumulation of partially reduced intermediates, unlike other chemoselective alternatives that required soluble metal salts as co-catalysts for similar purposes.16 Notwithstanding, the nitro into amine transformation must still proceed through a series of elemental steps, as formerly proposed by Haber (Figure 3).

We used operando IR spectroscopy to characterize the species formed on the Au/TiO2 surface in the presence of H2 and nitrobenzene.16 The spectra evidenced the formation of nitrosobenzene, phenylhydroxylamine and aniline on the catalyst surface and excluded azobenzene, azoxybenzene and hydrazobenzene as potential reaction intermediates.16 Accordingly, it was inferred that the –NO2 into –NH2 transformation occurs on Au/TiO2 catalysts through the Direct Reaction Route depicted in Figure 3. Equivalent conclusions were later reached with Au/Al2O3 catalysts.23 Additional batch experiments with Au/TiO2 catalysts showed that whereas the accumulation of intermediates in the liquid phase is avoided when nitrobenzene is used as the only starting reactant, large amounts of condensation products are formed if the reaction feed includes both nitro and nitrosobenzene.16 From these results, it was concluded that the nitrosobenzene intermediate detected by IR spectroscopy must be strongly retained onto the catalyst surface during a batch reaction experiment because, if it was not so, one should observe an accumulation of condensation intermediates in the solution. Moreover, with TiO2 as support, the surface concentration of nitrosobenzene was found to be relatively low, minimizing the surface bimolecular nitroso/hydroxylamine interactions, and so the formation of condensation intermediates (Figure 3).

Figure 3. Reaction route proposed by Haber for the hydrogenation of nitroaromatic compounds. Names below each structure refer to the prefix that identifies the nitrogen functional group. Interestingly, recent results have demonstrated that the nitro hydrogenation reaction route on gold catalysts is support dependent. Ceria, in contrast to TiO2, favors the Condensation Route27,28, and when the catalyst was optimized it was possible to achieve very high yields of relevant azocompounds. This result has been explained by assuming that with Au/CeO2 a greater concentration of nitrosobenzene is generated on the catalyst surface.28 Thus, an efficient protocol for the synthesis of azocompounds was discovered,28 as depicted more in detail below.

2.4. An Activity Issue ... Gold was soon featured as a promising, versatile catalyst for the chemoselective hydrogenation of nitro compounds,29 but its activity should be improved for a potential industrial application. To do that, a thorough kinetic investigation with Au/TiO2 catalysts for the hydrogenation of nitrobenzene with H2 was performed to identify the rate-determining reaction step during the reduction of nitrocompounds. The reaction rate was observed to decrease with increasing nitrobenzene concentrations in the range 0.2-1.2 mmol/L at 393 K and 8 bar of H2 in toluene, whereas the dependence was direct, and linear, with the H2 pressure when the nitrobenzene concentration was held constant at ~ 0.42 mmol/L. Assuming Langmuir-Hinshelwood and Hougen-Watson hypotheses, the experimental data could be satisfactorily reproduced with a kinetic model that assumes the dissociation of H2 as the rate-determining step.

To further check this model, we designed a series of Au/TiO2 catalysts with variable sizes of the gold nanoparticles,30 taking into account the following observations: a) the activity of gold for H2 dissociation is correlated with the number of low coordination sites in the metal nanoparticle;31 and b) the number of low coordinated sites depends, at least in part, on the gold particle size.32 We quantified the number of low coordination gold species in this
series of catalysts using IR spectroscopy and CO as a probe molecule. A direct correlation between the total number of low coordinated zero-valent gold centers and the activity for H₂ dissociation, measured by H₂/D₂ exchange, was found. Furthermore, and more importantly, catalysts activating H₂ fastest were also the most active in the hydrogenation of nitrocompounds, in agreement with the kinetic model inferred. Thus, to increase the catalytic activity of Au/TiO₂, one should control the gold nanoparticles architecture to increase the number of low coordination metal centers. We noted, however, that the increase in activity that can be achieved through this synthetic strategy is limited. Thus, we had to find other ways to increase the activity of the catalyst.


Prompted by the realization that a) the hydrogenation of nitrocompounds by gold is limited by H₂ dissociation and b) other metals such as platinum dissociate H₂ readily, we envisioned the use of TiO₂-supported bimetallic platinum-gold formulations as a promising strategy to improve previous results. Since it was found that the reaction controlling step was H₂ dissociation, we thought that by adding small amounts of Pt to the Au/TiO₂ catalyst, the rate of the reaction, i.e. of H₂ dissociation, could be increased. Meanwhile, for such a small Pt content, no effect on selectivity might occur. Then, we investigated the influence of increasing amounts of platinum, from 50 to 2000 ppm on the Au/TiO₂ catalyst, on the activity and selectivity for the hydrogenation of 3-nitrostyrene and found that the addition of 100 ppm of Pt is sufficient to promote a ~ 8-fold increase in activity of our most active Au/TiO₂ catalysts at 358 K, with little formation of undesired by-products (93.4 % selectivity to 3-vinylaniline at ~95 % conversion). Higher Pt contents led to gradually higher activity, but with a gradually lower chemoselectivity. Meanwhile, catalysts with no gold and 100 ppm of Pt on TiO₂ gave negligible conversion, stressing the bifunctional character of the Au-Pt formulation. The bifunctional Au-Pt/TiO₂ catalyst offers additional advantages over the original Au/TiO₂ catalysts such as lower separation costs and greater productivity through reactions performed in the absence of a solvent (nitro/Au mol ratio = 6738). Interestingly, the conversion vs time curve with the bimetallic material show a nearly constant slope to conversions up to 90 %, suggesting that the deactivation of the catalyst after almost 7000 turnovers is insignificant. Au/TiO₂ catalysts in the absence of trace amounts of Pt, in contrast, only achieve ~ 1350 turnovers and suffer from evident deactivation when the reaction is performed free of a solvent.

Recently, similar bimetallic Au-Pt formulations have been proposed for the hydrogenation of less demanding, from the chemoselectivity standpoint, nitroaromatics that incorporate –Cl and –OH. Whether or not these investigations bring any advantage is difficult to evaluate, because of the lack of cross-references to earlier work. With monochlorinated nitrocompounds, other metals, such as Pd, can also be selective (Pd-Au/Al₂O₃ with a 1:20 mol ratio).

4. Third Generation: Platinum, Ruthenium and Nickel Catalysts

4.1. The Role of the Metal Nanoparticle Architecture. To expand the scope of chemoselective catalysts for the hydrogenation of nitrocompounds, we next focused on the design of methods that could transform highly active, but traditionally unselective metals into highly chemoselective catalysts.

The effect of the active sites architecture on the hydrogenation of either styrene or nitrobenzene was firstly investigated through Pt/Al₂O₃ catalysts with variable metal content (0.2-5 wt %). On average, particle size increased when the metal content was increased. Complimentary, IR spectra of the catalysts probed with CO showed that the fraction of atoms in extended platinum terraces increases with the metal loading, consistent with nanoparticles that are, on average, larger. In the catalytic experiments, styrene reduced faster (mol converted × total mol of Pt' × h⁻¹) as the average platinum particle size and the number of terrace platinum sites increased. The particle size effect was, in contrast, minor on the nitrobenzene hydrogenation rate. In agreement, Pt/Al₂O₃ catalysts with the lowest platinum contents provided the greatest selectivities in the hydrogenation of 3-nitrostyrene (65 %, at almost complete conversions), whereas catalysts with the highest contents (5 wt %) produced 3-vinylaniline with a selectivity <25 %. The selectivity to 3-vinylaniline could be increased (~ 91 % at 95 % conversion) by the design of a catalyst that virtually lacked platinum terrace sites accessible to the reactants, using a high area mesoporous active carbon as the support. A similar effect is achieved by capping the surface of the metal nanoparticles with strongly bonded organic thiol ligands. Interestingly, the performance/structure dependence just depicted for nitro groups competing with C=C bonds in nitrostyrenes is reversed, and much less intense, for chloronitrobenzenes, where the selectivity decreases from 96.4 % to 84.5 % when the particle size decreases from ~12.2 nm to ~1.9 nm. The advantage of Pt with respect to gold is that owing to the high activity for H₂ dissociation, the reactions could be performed under conditions where gold is virtually inactive (313 K and 2 bar, in toluene). Recently, impressive molecular engineering has been accomplished to synthesize Pt single-atom centers and very tiny Pt clusters on iron oxide supports, which are
shown highly active, selective and of general use in the hydrogenation of nitroaromatics.\(^48\)

4.2. The Role of Strong Metal Support Interactions. We observed that properly prepared Pt/C catalysts can catalyze the reduction of nitroaromatic compounds selectively, but they suffer from a main drawback: up to ~25 % phenylhydroxylamine intermediates form during the reaction\(^7\) — and these species are associated with a loss in product quality.\(^7\) The problem was circumvented by selection of TiO\(_2\) as the support, considering that: a) TiO\(_2\) in Au/TiO\(_2\) catalysts is key for avoiding the accumulation of reaction intermediates;\(^46\) b) TiO\(_2\) in Au/TiO\(_2\) is key to boost the chemoselectivity;\(^5\) and c) terrace platinum sites, detrimental to selectivity,\(^7\) can be blocked by TiO\(_2\) species from the support due to strong metal support interaction phenomena at high activation temperatures.\(^4\) Strong metal-surface interactions are often invoked as critical in catalysis.\(^4\)

HRTEM and EELS evidenced the presence of TiO\(_x\) patches on top of TiO\(_2\)-supported platinum nanoparticles reduced at 723 K in H\(_2\)], and IR spectroscopy demonstrated the absence of terrace sites accessible to small reactants, such as CO.\(^7\) The selectivity to 3-vinylaniline (from 3-nitrostyrene) was then 93 % at 95 % conversion, a result that drastically differs from the low selectivities observed from the support due to strong metal support interaction phenomena at high activation temperatures.\(^4\) Strong metal-surface interactions are often invoked as critical in catalysis.\(^4\)

The effect of the TiO\(_x\) decoration on the nitro group activation with regard to a C=C bond is overwhelming: the former makes the styrene hydrogenation rate to fall two orders of magnitude when present together with nitrobenzene in a 1:1 mol ratio.\(^45\) Recent investigations have shown that the decoration of Pt nanoparticles with Zn species in Pt/ZnO catalysts leads to a similar increase in the catalyst selectivity.\(^44\)

In a subsequent series of experiments, the generality of the decoration concept was demonstrated through catalysts that included other metals such as ruthenium and nickel. With this battery of alternatives, a broad range of substrates containing C=C, C≡C, iodine, chlorine, bromide, cinnamate, nitrite, and carbonyl substituents could be satisfactorily tackled (Figure 1). More recently, silver-cerium oxide core-shell nanoparticles have been also shown highly active and selective for the hydrogenation of a number of nitrostyrenes.\(^4\)

5. New Synthetic Opportunities

The discovery of new chemoselective catalysts brought back some old processes that were left behind due to unsatisfactory performance, while opened the door to new synthetic reaction routes. A series of illustrative examples will be described below.

5.1. Synthesis of Oximes. Oximes may be obtained from \(\alpha,\beta\)-unsaturated nitro compounds provided that the catalyst is able to avoid a parallel hydrogenation of the C=C bond and a consecutive hydrogenation of the desired –C=NOH group. Au/TiO\(_2\) was such a catalyst that could produce oximes in high yields from conjugated nitroalkenes, which may also include functionalities such as Br, -OCH\(_3\) or –NO\(_2\). For example, 2-nitrophenyl acetaldehyde oxime was formed with 96 % selectivity from \(\beta,2\)-dinitrostyrene at 95 % conversion on Au/TiO\(_2\), whereas commercial Pd/C and Pt/C catalysts provided yields < 1 %.\(^7\) The method could be applied to produce cyclohexanone oxime, a 5 million metric-ton product used to manufacture Nylon 6, in high yields from 1-nitro-1-cyclohexene (~ 91 % at 383 K and 15 bar of H\(_2\), see Figure 4B). Industrially, cyclohexanone oxime is obtained by ammoximation of cyclohexanone, whereby cyclohexanone is obtained by air oxidation of cyclohexane at ~8% conversion per pass, and hydroxylamine has to be used or, otherwise, synthesized in situ with H\(_2\)O\(_2\) in NH\(_3\) (aq.) as it is done in the Sumitomo-ENI process (Figure 4A).

Later, it was found that Pt/TiO\(_2\) catalysts could efficiently afford cyclohexanone oxime from nitrocyclohexane with the TiO\(_2\)-decorated platinum catalysts.\(^4\) This process was previously performed by DuPont and 70 % yields were achieved with a PbO-modified Pd catalyst, at 413 K and 35 bar.\(^46\) Yields up to 80 % were achieved under solvent-free conditions (>13000 turnovers) at 353 K and 4 bar of H\(_2\) with the decorated Pt/TiO\(_2\) catalyst doped with Na to avoid Lewis catalyzed side reactions.\(^45\) Au/Al\(_2\)O\(_3\) was later shown to work for this reaction with yields to the oxime of ~ 83 % at 373 K and 6 bar in ethanol (0.33 mol nitrocyclohexane/L).\(^47\)
5.2. Direct Synthesis of Cyclohexanone Oxime From Nitrobenzene. The limitation of the procedures described above to synthesize cyclohexanone oxime is the requirement to prepare 1-nitro-1-cyclohexene or nitrocyclohexane. Then, we have recently developed a single-step procedure, involving Au/C and Pd/catalysts in the presence of NH$_2$OH·HCl and H$_2$ at 333 K, where cyclohexanone oxime can be obtained directly from nitrobenzene (Figure 4C). In this process, cyclohexylamine and aniline are formed first by hydrogenation on either Pd or Au centers. The former two subsequently condense into an imine catalyzed by Pd, with gold next acting as a Lewis acid to catalyze the imine hydrolysis to cyclohexanone. Finally, in the presence of NH$_2$OH·HCl, cyclohexanone oxime is formed in high yields (97 %) through an efficient cascade-type reaction.

5.3. Synthesis of Aromatic Azocompounds in One Step. Azocompounds could recently be synthesized directly from nitroaromatics using Au/CeO$_2$ catalysts. These compounds are widely used as dyes, food additives and drugs. Only up to 15 % selectivity to azobenzene was achieved with a low area ceria, which was increased to > 95 % with nanoparticulated CeO$_2$. Kinetic and IR data evidenced that the conversion of nitrosobenzene to the corresponding hydroxylamine is with this catalyst relatively slow, leading to high concentration of nitrosobenzene on the catalyst surface. In this scenario, nitrosobenzene and the corresponding hydroxylamine condense before further hydrogenation to aniline occur, forming azoxybenzene and, subsequently, azobenzene.

These results contrast to those obtained when the support was TiO$_2$ (Figure 5). Azocompounds can be obtained in high yields not only because the Condensation Route is favored when CeO$_2$ is the support, but also because the undesired hydrogenation of the azo group to aniline is inhibited in the presence of even traces of azoxybenzene. Therefore, by using a gold catalyst on a high surface area CeO$_2$ it is possible now to produce azocompounds in one step with very high yields.

5.4. Cascade Type Reactions: One-Step Synthesis of Imines, Secondary Amines, Nitrones, and $\beta$-Amino Carbonyls. Amines are ubiquitous precursors to lend new N–C bonds by condensation with a second functionality. Thanks to the discovery of chemoselective catalysts for the hydrogenation of nitrocompounds, we can now conceive single-step processes that combine several cascade reactions (among others, a full or partial reduction of the nitro group). Cascade-type reactions are interesting because they allow the design of more intensive process, increase the atom-economy and are, thus, more ecofriendly.

For example, Au/TiO$_2$ catalysts were used to obtain imines with good yields (80-87 %) directly from nitroaromatic compounds, aldehydes and H$_2$ in a single reaction step (Figure 6), even when sensitive substituents such as C=C bonds or Br were present. Substituted imines are important building blocks in the synthesis of fine chemicals, pharmaceutical compounds, and
agrochemicals. The results lie on a catalyst that facilitates full reduction of the nitro group into an amine without reduction of the –C=O bond and the rest of functionalities. Secondary amines could also be obtained in high yields by simple prolongation of the reaction time; and aldehydes incorporation conjugated C=C bonds led to α,β-unsaturated imines, whereas α,β-unsaturated ketones yielded β-amino carbonyl compounds. We could drastically tune the distribution of reaction products in mixtures of nitrocompounds, aldehydes and H₂ by using a different chemoselective catalyst, mesoporous carbon-supported platinum nanoparticles, opening the possibility to obtain another valuable class of building blocks for organic syntheses: nitrones. This cascade reaction requires not only a chemoselective catalyst, but that the aldehyde condenses with the hydroxylamine intermediate before the nitro group is fully reduced to an amine. The Pt/C catalyst depicted herein above, which lacks platinum terrace sites accessible to the reactants, can do this reaction very efficiently, with yields to a variety of substituted nitrones that can exceed 90 % (Figure 6). Common supported platinum catalysts, in contrast, were found unselective.

![Figure 5: Influence of the support in the distribution of products during the hydrogenation of nitrobenzene. On TiO₂, gold nanoparticles catalyze a full reduction of the nitro group, whereas azocompounds form in high yields when the support is nanoparticulated CeO₂.](image)

![Figure 6: A diversity of chemicals can be obtained from nitro compounds via single-step reactions thanks to the development of a variety of catalysts that hydrogenate the –NO₂ chemoselectively in the presence of other reactive functionalities.](image)

On the other hand, α,β- Unsaturated ketones could also be engaged in cascade-type processes with anilines synthesized in situ from nitro compounds. In this case, the addition reaction takes place through the conjugated C=C bond to produce β-amino carbonyls in good yields (83-91%) in the presence of Au/TiO₂. The support plays a
dual role here: a) induces chemoselectivity and b) supplies Lewis acid sites for the addition reaction, and the tolerance to ligands such as ethyl cinnamates or nitriles opens the door to important fragrances and biocides. The Au/TiO₂ sample, moreover, catalyzes the three-component reaction between nitrobenzenes, phenylacetylenes, and H₂ to methyl substituted imines. Gold acts in this case as a catalytically active species both for reduction of the –NO₂ group to –NH₂ and for the subsequent coupling between the amine and the C≡C group. Au/Fe₂O₃ has also been used in inter and intramolecular tandem reactions between nitro compounds and C≡C to form indoles.

N-alkylated anilines could be also obtained from nitrobenzenes and alcohols on Au/Fe₂O₃, in a process where gold catalyzes oxidation of the hydroxyl group into aldehyde via formation of Au–H species that cause reduction of the –NO₂ group into –NH₂, with final formation of the imine upon amine/aldehyde condensation. Recently, MgO, a solid base, and a chemoselective Pt/TiO₂, catalyst were coupled to obtain α'-aminochalcones from nitroacetophenone and benzaldehyde via a one-pot reaction, with yields and selectivities higher than those in typical multistep processes.

6. Conclusions

We have shown how supported gold nanoparticles have opened new possibilities for chemo-selective hydrogenation of substituted nitroaromatics and this has allowed to discover new reaction processes for producing substituted anilines, imines, oximes, nitrones, azocompounds, and β-amino carbonyl compounds. Moreover, the research on the reaction mechanism and how gold and the support interact and collaborate during the reaction have taught us how to modify other metal catalysts to expand their scope as catalysts for a number of reactions of fundamental and applied interest.

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REFERENCES

1 Blaser, H.-U.; Steiner, H.; Studer, M. ChemCatChem, 2009, 1, 210–221
4 Corma, A.; Serna, P. Science 2006, 313, 332–334
Platinum nanoparticles are embedded on the mesoporous carbon matrix, with terrace sites blocked by carbonaceous species.


44 Yarulin, A; Berguerand, C; Alonso, A.O.; Yuranov, I.; Kiwi-Minsker, L. Cat. Today, 256, 241–249


46 GB Patent 860340, 1961, to DuPont


