

A Brief Review on Nano-Catalysis and Its Applications

Kavita Rani*

M.Sc. Physical Chemistry, Group- 1, Department of Chemistry, University of Delhi

Abstract – In the time of nanoscience, when all gadgets and developments are decreasing and littler with improved properties; catalysis is a significant area of research. With this in mind, we present a summary of the applications of nanoscience in catalysis, energy conversion, and energy conservation. In the present study we are trying to condense in-writing knowledge that is beneficial to individuals to use nano-estimated inspiration in day-to-day life. Improvement of synergistic properties is discussed here due to reduced impetus size to nano scale.

Keywords: Catalysis, Nanocatalysis, Nano Science, Nano Material

-----X-----

INTRODUCTION

Catalysis assumes an emphasis on improvements in concoction and lies at the heart of countless material conventions, ranging from laboratory-level scholastic work to compound industry (Hemalatha et al 2013). By using catalytic reagents, the temperature of a process can be reduced, reagent-based waste reduced and a reaction selectivity improved that theoretically prevents unnecessary side reactions leading to green technology. In 1998, Anastas and Warner (1998) proposed a set of twelve principles which is Green Chemistry's main philosophy of reducing or eliminating chemicals and chemical processes which have negative environmental impacts. The design and development of perfect impulses is one of the important ideas of green science. As per these criteria, synergistic reagents are stronger than stoichiometric reagents (as specific as would be prudent). Stoichiometric reagents are used in mass and operate only once, while synergistic reagents are used in small amounts and can direct a solitary response many times over. The foundation of all the twelve principles of green chemistry is to function more like nature. Nature explicitly provided us with clues that micro-organisms and/or enzymes could be used to conduct environmentally benign reactions.

NANOPARTICLES CHEMISTRY

Particles which is in size 1-100 nano-meter (10⁻⁹ meter) are Nanoparticles (NPs). Nanochemistry's important action is the combination of stable nanoparticles, ranging between 1-100 nm. Nanoparticles can be synthesized in two main headings, such as I top-down technologies and (ii) bottom-up technologies, by different means types.

Numerous modern instruments have been used to represent nanomaterials to determine genuine size, form, surface structure, valence, synthetic arrangement, gap in the electron band, holding state, light outflow, retention, dissipation and diffraction properties like NMR, infrared spectroscopy (IR), bright and visible spectroscopies.

NANO-CATALYSIS

Since the late 1990s, nanocatalysis has risen as an area at the interface between homogeneous catalysis and heterogeneous catalysis and with nanoscience going ahead. The focus is on synthesizing, characterizing, exploring, and exploiting well-defined nanostructured catalysts, including nanoparticles (NPs) and nanomaterials. NPs, which are known to be the nanotechnology building blocks, refer to particles with at least one dimension that is < 100 nm. At the nanometer scale, metallic NPs are formed by clusters of atoms with intermediate properties between molecules and bulk metals. This characteristic defines new chemical and physical properties which are beneficial for different applications, especially for catalysis. The term nanomaterial refers to any solid with a dimension to the anometer. A specific definition of "nanomaterials" was created by the European Commission (EC) in 2011: Despite these differences in nomenclature, NPs are always implicated and "nanocatalysts" or "nanocatalysis" adequately summarizes these cases. Nano material strategies a trademark, coincidental or manufactured material containing particles in an unbound state or as an aggregate or agglomerate and in which at least one outer gage is within the size range of 1–100 nm for 50 percent or more of

the particles in the dissemination of the number size.

In recent decades, Because nanocatalysts are composed of small particles of acatalytically active content, usually with a range of 1–100 nm in diameter, they have attracted intense interest. We can be applied in various fields, such as catalysis, electrocatalysis, sensors, filters, electronic nanoscale, fuel cells, cosmetics, oil, climate, motors, water purification, and optoelectronics. The first scientific aim and challenge is associated with the synthesis of the separticles with maximum control over size and shape to tailor their physical and chemical properties and in a specific reaction to maximize their efficiency. The second challenge is to understand how optimum catalytic reaction performance is produced by the composition and atomic scale structures of NPs. Continuous developments in nanotechnology and material science can help explore new kinds of multifunctional nanocatalysts for the achievement of green and safe concoction types in the representation, judicious design, and development. This research provides new opportunities to understand the nature of the active sites, the metal-support interaction mechanism, and the origin of the structure–reactivity relationship by careful design and synthesis of specific size and shape catalyst particles at the nanoscale.

Highly--Active Nanocatalysts

The best example to demonstrate the exceptional catalytic activity of nanomaterials is a catalyst with gold nanoparticles distributed on a titania support in the 5 nm regime. This catalyst exhibits high activities at room temperatures for epoxidizing hydrocarbons and CO oxidation. (Bond and Thompson, 1999) It has been suggested that the quantum confinement effects change the electronic structure of this noble metal and lead to the unusual catalytic activities observed. This discovery has spurred extensive research efforts in searching novel nanocatalysts for the important catalytic reactions with low reactivity, such as activation of saturated hydrocarbons in reforming reactions, oxygen reduction reactions in fuel cells, and lingo---cellulose biomass hydrolysis. Tiny, charged metal clusters have been documented to exhibit easy activation and dehydrogenation of methane molecules for example.

Highly--Selective Nanocatalysts

Highly— selective catalysts can help to reduce energy consumption in chemical industries required for product separation and waste disposal processes. Alternative energy resource development is also based on highly selective catalysts. A key step in the conversion of biomass is the selective conversion of biomass-derived carbohydrates to liquid fuels and useful chemicals. In our laboratory, numerous molecular level factors affecting catalytic selectivity have been identified using surface science studies

over the past few decades that allow nanotechnology to design and engineer highly selective catalysts (samoljae and Kliever, 2009).

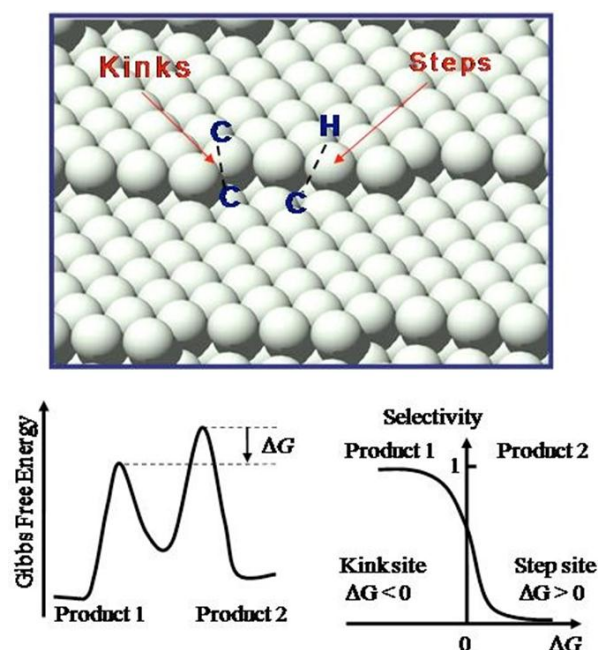


Fig1: The upper panel: the model kinked Pt surface (the upper panel).

At the kink and step site, respectively, the C—C bond and the C—H bond are dissociated. The lower left panel: the potential surface of schematic free energy for a two-way reaction. Product 1 is formed by breaking the bond C—C and Product 2 is formed by breaking the bond C—H. At the kink site, the activation barrier for product 1 is lowered, which contributes to the disparity in selectivity between phase and kink sites, as shown in the panel below right.

The rate— determining steps for various products usually occur at different active sites on the catalyst surface for a multiple—, path-catalytic reaction. Consider a catalytic reaction involving a cyclic hydrocarbon reactant (Figure 2), the C---C bond split leads to the ring opening product (Product 1); while the C---H bond dissociation gives a dehydrogenation product (Product 2). For the two things, the overall statures of the Gibbs free energy hindrances regulate the proportion of item 1 to item 2 generated at a given surface location. At the progression destinations on platinum surfaces, the splitting of C---H bond occurs all the more promptly than that of the C---C bond, which prompts a higher probability of framing the dehydrogenation element. The breaking of the C—C bond is less difficult at the wrinkle locations, and the ring— opening item is relying on increment. From this basic picture the selectivity of heterogeneous synergistic responses is finally decided by the aggregate groupings of the diverse destinations for different response

pathways. For catalysts with nanoparticles, the size and shape of the nanoparticles determines the concentration of active sites. Advances in the synthesis of nanoparticles enable the precise control of surface active sites through the production of mono-, disperse and shape- controlled catalysts. An example is shown in Figure 3 where pyrrole hydrogenation selectivity over Pt nanoparticles exhibits dependency on both size and shape. (Kuhn and Tsung, 2008). The change in selectivity is induced by the nanocatalysts change in surface structure with different sizes and shapes.

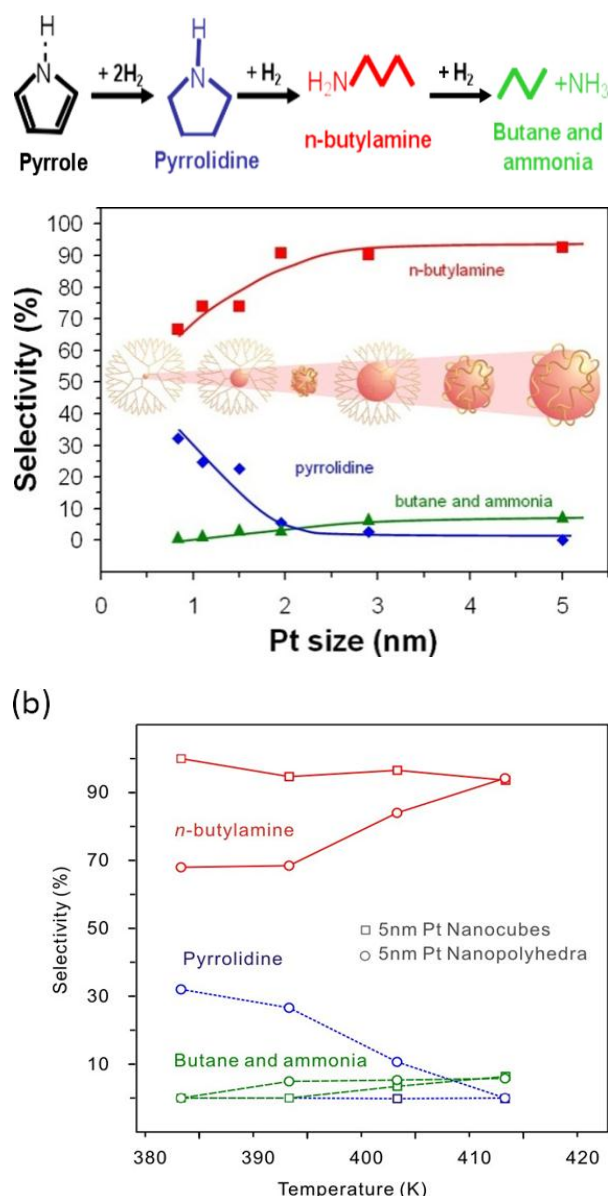


Fig2: (A) Nanoparticulate size dependence of pyrrole hydrogenation selectivity under reaction condition: 4 Torr pyrroleum, 400 Torr H, 413 K. Small nanoparticles show high pyrrolidine selectivity. (B) Nanoparticles form the selectivity dependency of pyrroleum hydrogenation under the following conditions: 4 Torr pyrroleum, 400 Torr H. At relatively low temperatures, particles of the nanopolyhedra have a higher pyrrolidine selectivity than nanocubes.

A significant class of selective catalysts are bimetallic alloys which provide special atomic arrangements of metal components for surface-active sites (Norskov, 2008). This field has recently been revolutionized by the emergence of nano-scale rational catalyst design. High-throughput, computer-based screening provides alloy candidates with the elements and atomic arrangement that may have optimum catalytic properties. Nanoalloys are then synthesized and put to the test. The successful synthesis of alloy catalysts with intended composition is based on the fact that reducing the size of alloy particles usually leads to a reduction of the immiscible gap. Following this method a Ni₁Zn₃ alloy catalyst for partial hydrogenation of acetylene was discovered with a higher selectivity.

In a more recent study, photoelectron spectroscopy was applied in-situ X-ray to monitor the surface segregation of bimetallic nanoparticles under reaction conditions. It was observed that the bimetallic composition on surfaces of nanoparticles alternates between oxidizing and reducing conditions when the chemical gas environment is switched between. This new insight can lead to the development of "smart" catalysts whose surface structure depends advantageously on the reaction environment.

Nanocatalysis in Green Chemistry

Green chemistry is commonly recognized as "planning, developing and upgrading compound procedures and products designed to diminish or destroy substances that are harmful to human well-being and the environment." The concept of performing this kind of chemistry is gaining popularity among the players in the chemical industry, as the major challenges of today are achieving sustainable production processes, with lower energy use and less environmental impact. Furthermore, green methods of reproduction could also provide economic benefits for businesses. For example, platinum prices have increased significantly since 2000, so it is advisable to cover and reuse such expensive catalysts of precious metals (Norskov, 2008). Many examples of commercialised green chemistry processes demonstrate the promise of this innovative theory. (Sinfelt, 1984) However, the overall impact of this new way of doing chemistry is still marginal compared to conventional industrial processes (Studt, 2008) This is because businesses optimize productivity within the current policy limits while keeping an eye on social acceptability.

APPLICATIONS OF NANOCATALYSTS

Carbon nanotubes

Due to their unique physical and mechanical properties, significant attention has been given to

nano-sized carbon materials (NCMs) in particular CNTs or carbon nano-tubes. CNTs have been used as field discharge sources in various fields (Trans et al., 1998), electrical nano-conducts (Mintmire et al., 1992), optional Li particle batteries (Maurin et al., 1999), double layer electrical condensers (Frac-kowiak et al., 2000), subatomic sifters (Wang et al., 1999) and control modules (Bessel et al., 2001). Likewise, CNTs were later used for adsorption of hydrogen (Dillon et al., 1997), as they are exceptionally permeable, solid, stable, and minimal effort. Hydrogen is considered as an ideal energy carrier due to its high energy content and non-polluting nature, which can be widely used in the near future (Züttel, 2004; Shi and Hwang, 2007; Saito et al., 1993; Ruoff and Lorents, 1995).

Water refining

Hydrogen is the most recent in the development of energy suppliers, with its credit having numerous social, monetary and ecological advantages reminiscent of its use for material firms, which constitutes 40 percent of its use. It infers the pre-emptive importance of H₂ for the world of the present and of tomorrow. Impressive measurements of hydrogen are outlined in the various forms of decrease, in particular the decrease of metal impurities used for different hydrogenation and different reactions isolated from their use in the hydrogenation responses themselves. Arrangement techniques that can genuinely generate impurities in metallic structure without using any hydrogen to put oxidic or different types of impetus into their metallic structures can therefore be deeply invaluable in reducing hydrogen use.

Shashikala et al. (2007) said that combining nano-metallic silver particles using a novel electro-synthetic statement technique on carbon-shrouded alumina leads the way to a continuing economy of hydrogen. This strategy gives an exceptionally powerful silver impetus in regulating water organisms. Additionally, it is widely recognized that the impurities that Ag has assisted are reusable. The combined qualities of Al₂O₃ and carbon, for example, low acidity, high mechanical strength and closeness of meso pores in carbon coverage in alumina (CCA), are equally useful for the program of extremely active AgCCA impurities. Nanocatalysts processing biodiesel have extremely specific surface and high catalysis exercises that can unravel the above problems.

Wen and so on (2010) analyzed the possibility of using the strong KF / CaO base nanocatalyst for the production of bio-diesel with a yield of over 96%. The impetus is well utilized to transform the higher corrosive substance oil into biodiesel. It is permeable, with molecule sizes of 30–100 nm. XRD examination found the impetus to have a new KCaF₃ gem that increases reactant action and solidity. The high clear surface area and enormous pore size are

ideal for interaction with impetus substrates, which increased the performance of transesterification accordingly. The production of Chinese fat seed oil biodiesel positively impacts the use of agricultural and ranger service products.

In sedate conveyance CNTs may be ideal for bio-applications in biorecognition and drug conveyance systems along these lines. By the way, biocompatibility of CNTs is still uncertain, and the prolonged functionalization process of CNT surface remains a significant obstacle to earth applications (Magrez et al., 2006; Cheng et al., 2009).

Thoughtful et al. (2010) recompleted the combination of carbonated polypyrrole nanoparticles (CPyNs) with controlled estimates and their textural properties, and investigated the potential capacity of CPyNs as imaging tests and medication transporters depending on their porosity, alluring properties and biocompatibility.

Malachite green color photocatalytic decoloration

Every day large quantities of unconsumed colors made by materials and printing firms are released into the water. The closeness of hues and shades in water significantly damages the oceanic condition (Robinson et al., 2001; Pearce et al., 2003; Talarposhti et al., 2001). These contaminants make concoction oxygen (COD) common, biochemical oxygen (BOD) appeal, poisonous consistency, terrible stench and, most importantly, wastewater shading. Even at low focuses, the shade of such contaminants can be seen in light of the fact that the closeness of colors in water is deeply noticeable. This impact is unwanted as the shading squares have daylight access to oceanic greenery, and decrease photosynthetic activity within the biological system (Srivastava et al., 2004; Cheng et al., 2008).

Methylene blue photodegradation

Soni et al. contemplated photocatalytic action in the obvious piece of the sun powered range (442 nm) for deeply organized slender, mesoporous titanium-doped thiourea films. This property, which is critical for the further manufacture of photovoltaic gadgets that operate in daylight, is uncovered as an aspect of illumination time and film thickness by testing methylene blue photodegradation upon contact with a N-doped TiO₂ film (Soni et al. 2008).

TiO₂ is known as the best and earth-neighborly photocatalyst and most widely used to photograph various pollutants (Fujishima and Honda, 1972; Fujishima et al., 2000; Graetzel, 2001; Hay and Raval, 1998). As with E, TiO₂ photocatalysts can

also be used to kill microorganisms. Coli suspension (Kikuchi et al., 1997);

Squander water treatment

Halogenated natural mixes (HOCs) are among the most common water contaminations in industrialized nations. For example, pharmaceutical assembly or the IT division, these natural atoms take on a significant job as solvents and added substances in various companies. For the most part, HOCs are dangerous and hurtful synthetic compounds that are often extremely industrious and can cause serious medical problems, such as malignant growth or mutagenic injury. In this way a complete annihilation of these mixes is desired, or even required.

The traditional work of handling wastewater cannot adapt to this issue. In this way, it still does not seem feasible to use costly and energy-concentrated methodologies to take care of this problem. The normal act of discarding indus-process wastewater contaminated with a limit heap of natural solvents is cremation and only hints of HOCs. For example, at a wastewater treatment plant, alcohols could be extracted by bacte-rial treatment, as it may be, organic mixes.

The HOC is often the critical component which leads to cost-intensive disposal. Current detoxification techniques such as adsorption on activated carbon or wastewater com-ponents oxidation do not result in an environmentally friendly and cost-effectively priced solution. For example, it would take huge quantities of activated carbon to extract HOC traces from water loaded with other com-ponents that are involved in sorption. In the case of oxidation, excess quantities of oxidant are needed because it would oxidize all reduced water components. The risk arises when insufficient quantities of oxi-dant are added, that if the oxidation is not complete, even more toxic components can occur. Overpriced incineration is the last and often performed result. However, aqueous wastage incineration is a waste of energy. Hence a decentralized selective dehalo-generation treatment of wastewater can give a significant eco-nomic advantage for a medium-sized enterprise.

Hildebrand et al. (2008) proposed a consideration of detoxifying water using the HDH technique on palladium-containing nano-impulses by severe destruction of HOCs. Detoxification ensures that steady HOCs are transformed into natural aggravates which can be effectively removed by biodegradation at a wastewater treatment plant. They are viable and specific in diminishing responses to hydrodehalogenation.

CONCLUSION

In the last decade, the field of nanocatalysis (use of nanoparticles to catalyze responses) has experienced an explosive improvement in both homogeneous and heterogeneous catalysis. Nanoparticles have an overwhelming proportion of surface-to-volume compared with mass materials, they are alluring to be used as impulses. Cata lysts accelerate and boost thousands of different chemical reactions daily, and thus form the basis for the worldwide multibillion dollar chemical industry and indispensable technologies for environmental protection. Nanotechnology and nanoscience work is expected to have a major impact on new catalyst production. The detailed understanding of chemistry of nanostructures and the ability to control materials on the nanometer scale will ensure a rational and cost efficient devel-opment of new and more capable catalysts for a chemical process.

REFERENCES

- K. Hemalatha, G. Madhumitha, A. Kajbafvala, N. Anupama, R. Sompalle and S. M. Roopan (2013). "Function of nanocatalyst in chemistry of organic compounds revolution: An overview," Hindawi Publishing Corporation, Journal of Nanomaterials, Volume, Article ID 341015, 23 pages, <http://dx.doi.org/10.1155/2013/341015>
- P. T. Anastas and J. C. Warner (1998). Green Chemistry: Theory and Practice, Oxford University Press: New York.
- Bond, G.C. and D.T. Thompson (1999). Catalysis by gold. Catalysis Reviews---Science and Engineering, 1999. 41(3---4): p. 319-388.
- Somorjai, G.A. and C.J. Klier (2009). Reaction selectivity in heterogeneous catalysis. Reaction Kinetics and Catalysis Letters. 96(2): pp. 191---208.
- Heiz, U. and U. Landman (2007). Nanocatalysis. Nanoscience and technology, Berlin ; New York: Springer. xvi, 503 p.
- Kuhn, J.N., et. al. (2008). Structure Sensitivity of Carbon---Nitrogen Ring Opening: Impact of Platinum Particle Size from below 1 to 5 nm upon Pyrrole Hydrogenation Product Selectivity over Monodisperse Platinum Nanoparticles Loaded onto Mesoporous Silica. Journal of the American Chemical Society, 130(43): p. 14026---+.
- Tsung, C.K., et. al. (2009). Sub---10 nm Platinum Nanocrystals with Size and Shape Control: Catalytic Study for Ethylene and Pyrrole

- Hydrogenation. Journal of the American Chemical Society, 131(16): pp. 5816---5822.
- Norskov, J.K., et. al. (2008). The nature of the active site in heterogeneous metal catalysis. Chemical Society Reviews. 37(10): pp. 2163--2171.
- Sinfelt, J.H., G.H. Via, and F.W. Lytle (1984). APPLICATION OF EXAFS IN CATALYSIS - -- STRUCTURE OF BIMETALLIC CLUSTER CATALYSTS. Catalysis Reviews---Science and Engineering, 26(1): pp. 81-140.
- Studt, F., et. al. (2008). Identification of non--precious metal alloy catalysts for selective hydrogenation of acetylene. Science. 320(5881): pp. 1320-1322.
- Saito, R., Fujita, M., Dresselhaus, G., Dresselhaus, M.S. (1993). Appl. Phys. Lett. 60, 2204.
- Mintmire, J.W., Dunlap, B.I., White, C.T. (1992). Phys. Rev. Lett. 68, 631.
- Ruoff, R.S., Lorents, D.C. (1995). Carbon 33, pp. 925.
- Maurin, G., Bousquet, C., Henn, F., Bernier, P., Almairac, R., Simon, B. (1999). Chem. Phys. Lett. 312, 14.
- Wang, J., Pan, Z.J., Zhang, Z.H., Zhang, X.D., Wen, F.Y., Ma, T., Jiang, Y.F., Wang, L., Xu, L., Kang, P.L. (2006). Ultrason.
- Dillon, A.C., Jones, K.M., Bekkedahl, T.A., Kiang, C.H., Bethune, D.S., Heben, M.J. (1997). Nature 386, 377.
- Shashikala, V., Kumar, Siva, Padmasri, A.H., David Raju, B., Venkata Mohan, S., Nageswara Sarma, P., Rama Rao, K.S. (2007). J. Mol. Catal. A: Chem. 268, 95.
- Wen, L., Wang, Y., Lu, D., Shengyang, H., Heyou, H. (2010). Fuel 89 (9), pp. 2267.
- Magrez, A., Kasas, S., Salicio, V., Pasquier, N., Seo, J.W., Celio, M. (2006). Nano Lett. 6, 1121.
- Oh, W.-K., Yoon, H., Jang, J. (2010). Biomaterials 31, pp. 1342.

Corresponding Author

Kavita Rani*

M.Sc. Physical Chemistry, Group- 1, Department of Chemistry, University of Delhi

Kavita Rani*