

Journal of Chemistry (JCHEM)



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Article history

Submitted 01.01.2024 Revised Version Received 04.03.2024 Accepted 04.04.2024

Abstract

Purpose: The aim of the study was to assess the impact of pH on the catalytic activity of metal nanoparticles in organic reactions in Ghana.

Methodology: This study adopted a desk methodology. A desk study research design is commonly known as secondary data collection. This is basically collecting data from existing resources preferably because of its low cost advantage as compared to a field research. Our current study looked into already published studies and reports as the data was easily accessed through online journals and libraries.

Findings: The pH of the reaction medium can significantly influence the surface charge, morphology, and composition of metal nanoparticles, consequently affecting their catalytic performance. At specific pH levels, the protonation or deprotonation of functional groups on the nanoparticle surface can occur, altering the electronic properties and reactivity of the catalyst. Additionally, pH can influence the stability of metal nanoparticles and their interaction with reactants, intermediates, and products. Studies have shown that optimizing pH conditions can enhance catalytic activity,

selectivity, and recyclability of metal nanoparticles in various organic transformations, including hydrogenation, oxidation, and coupling reactions. Understanding the pH dependence of metal nanoparticle catalysis is crucial for the rational design and optimization of efficient catalytic systems for organic synthesis.

Implications to Theory, Practice and Policy: Surface chemistry theory, electrochemical theory and colloid chemistry theory may be used to anchor future studies on assessing the impact of pH on the catalytic activity of metal nanoparticles in organic reactions in Ghana. Implementation of advanced experimental techniques for real-time monitoring of pH effects on nanoparticle catalysis. Techniques such as in situ spectroscopy and surface characterization methods can provide valuable insights into dynamic changes in nanoparticle structure and reactivity during catalytic reactions. Integration of pH considerations into regulatory frameworks and guidelines for sustainable catalysis.

Keywords: *pH, Catalytic Activity, Metal Nanoparticles, Organic Reactions*

INTRODUCTION

The catalytic activity of metal nanoparticles refers to their ability to accelerate chemical reactions by providing active sites for the formation of intermediate species and facilitating the conversion of reactants into products. This property has garnered significant interest due to its potential applications in various fields such as catalysis, energy conversion, and environmental remediation. Metal nanoparticles possess unique catalytic properties attributed to their high surface area-to-volume ratio, size-dependent electronic properties, and surface reactivity. For instance, gold nanoparticles have shown remarkable catalytic activity in oxidation reactions, hydrogenation, and carbon-carbon bond formation due to their tunable surface plasmon resonance and high surface energy (Hvolbæk, 2007). Similarly, silver nanoparticles have been employed as efficient catalysts in organic transformations, antimicrobial coatings, and pollutant degradation owing to their high surface-to-volume ratio and strong interaction with organic molecules (Xia, 2017).

In developed economies like the USA, Japan, and the UK, there has been a growing trend in research and development focusing on the catalytic activity of metal nanoparticles. For instance, in the USA, the National Nanotechnology Initiative (NNI) has been instrumental in advancing research in nanotechnology, including the synthesis and application of metal nanoparticles in catalysis (NNI, 2020). Japan, with its strong emphasis on technological innovation, has seen significant investment in nanotechnology research, leading to the development of advanced catalytic materials based on metal nanoparticles (Mizukoshi, 2018). Similarly, the UK has witnessed a surge in academic and industrial collaborations aimed at harnessing the catalytic potential of metal nanoparticles for sustainable chemical processes and environmental applications (Government Office for Science, 2016). These trends underscore the importance of metal nanoparticles in driving innovation and economic growth in developed economies.

In developing economies, the exploration of metal nanoparticles' catalytic activity has been gaining momentum, driven by the need for sustainable and cost-effective solutions to various societal challenges. For example, in China, the world's largest emitter of greenhouse gases, there has been a notable increase in research focusing on the catalytic conversion of carbon dioxide into value-added chemicals using metal nanoparticles as catalysts (Zhang, 2018). This research aligns with China's commitment to reducing carbon emissions and transitioning towards a more sustainable economy (National Development and Reform Commission, 2020). Similarly, India, with its burgeoning population and rapid industrialization, has been investing in nanotechnology research to address environmental pollution and enhance energy efficiency through the development of efficient catalytic systems based on metal nanoparticles (Department of Science and Technology, 2019). These efforts underscore the pivotal role of metal nanoparticles in driving innovation and addressing pressing societal and environmental challenges in developing economies.

In Brazil, a country known for its abundant natural resources, there has been a growing interest in utilizing metal nanoparticles as catalysts for biomass conversion and biofuel production (de Souza, 2015). The Brazilian government has been supporting research initiatives aimed at harnessing the catalytic potential of metal nanoparticles to valorize agricultural residues and mitigate dependence on fossil fuels (Ministério da Ciência, Tecnologia, Inovações e Comunicações, 2017). Similarly, in South Africa, efforts are underway to leverage metal nanoparticles in catalytic converters for automotive applications to reduce emissions and comply with stringent environmental regulations (Department of Science and Innovation, 2020). These examples highlight the diverse applications

of metal nanoparticles in addressing unique challenges faced by developing economies and underscore the importance of continued investment in nanotechnology research and development.

In Nigeria, a country with a growing population and expanding industrial sector, there is increasing interest in leveraging metal nanoparticles for catalytic applications to address environmental pollution and enhance industrial processes. Research efforts have focused on utilizing metal nanoparticles as catalysts for wastewater treatment, air pollution control, and the production of clean energy sources (Ogunsona, 2019). The Nigerian government, recognizing the potential of nanotechnology in driving sustainable development, has been promoting collaborations between academia, industry, and government agencies to advance research and innovation in this field (National Agency for Science and Engineering Infrastructure, 2018). Furthermore, initiatives such as the National Nanotechnology Development Agency (NNDA) are actively supporting capacity building and technology transfer in nanoscience and nanotechnology to stimulate economic growth and improve quality of life (NNDA, 2020). These efforts underscore the role of metal nanoparticles as catalysts for sustainable development in Sub-Saharan Africa.

In Kenya, a country with a growing economy and increasing urbanization, there is a burgeoning interest in utilizing metal nanoparticles for catalytic applications in various sectors such as agriculture, healthcare, and environmental management. Research initiatives have focused on developing nanoparticle-based catalysts for water purification, crop protection, and medical diagnostics (Ong'uti, 2017). The Kenyan government, in collaboration with international partners and research institutions, has been supporting projects aimed at harnessing the potential of nanotechnology to address local challenges and stimulate economic growth (Ministry of Education, Science and Technology, 2014). Additionally, initiatives such as the Kenya Industrial Research and Development Institute (KIRDI) are facilitating technology transfer and commercialization of nanoparticle-based catalytic solutions to enhance industrial competitiveness and promote sustainable development (KIRDI, 2020).

In South Africa, a leading economy in the African continent, there has been significant interest in the catalytic applications of metal nanoparticles, particularly in the fields of renewable energy and environmental sustainability. Research endeavors have focused on utilizing metal nanoparticles as catalysts for the production of clean fuels, such as hydrogen and biodiesel, from renewable feedstocks (Mamba, 2018). Additionally, metal nanoparticles have shown promise in catalyzing chemical reactions for wastewater treatment, air pollution control, and carbon dioxide capture and utilization (Wu, 2019). The South African government, through initiatives like the Department of Science and Innovation (DSI), has been investing in research and development to harness the catalytic potential of metal nanoparticles for addressing pressing societal and environmental challenges while fostering economic growth and job creation (DSI, 2020).

In Ghana, a country with a growing economy and increasing industrialization, there is a growing interest in utilizing metal nanoparticles for catalytic applications to enhance industrial processes and address environmental issues. Research efforts have focused on developing nanoparticle-based catalysts for sustainable agriculture, pollution abatement, and renewable energy production (Opoku-Damoah, 2020). The Ghanaian government, recognizing the potential of nanotechnology in driving economic development and improving quality of life, has been supporting research initiatives and collaborations aimed at harnessing the catalytic properties of metal nanoparticles (Ministry of Environment, Science, Technology and Innovation, 2016). Additionally, institutions such as the Council for Scientific and Industrial Research (CSIR) are actively engaged in research

and technology development to promote the adoption of nanoparticle-based catalytic solutions in various sectors of the economy (CSIR, 2020).

pH level is a crucial parameter in chemistry that measures the acidity or alkalinity of a solution on a scale from 0 to 14, where pH 7 represents neutrality, pH below 7 indicates acidity, and pH above 7 indicates alkalinity. The pH of a solution influences the surface charge and reactivity of metal nanoparticles, thereby impacting their catalytic activity. At lower pH levels, metal nanoparticles may undergo surface oxidation, leading to the formation of metal oxides or hydroxides, which can significantly alter their catalytic properties (Hvolbæk, 2007). Conversely, at higher pH levels, the surface of metal nanoparticles may become more negatively charged, affecting their adsorption capacity and interaction with reactant molecules, thus influencing catalytic activity (Xia et al., 2017).

Considering four likely pH levels ranging from acidic to alkaline, each level can have distinct effects on the catalytic activity of metal nanoparticles. At pH levels below 7 (acidic conditions), metal nanoparticles may exhibit enhanced catalytic activity due to increased surface reactivity and protonation of functional groups on the nanoparticle surface, facilitating adsorption and activation of reactant molecules (Hvolbæk, 2007). However, excessively low pH levels may also lead to surface corrosion and aggregation of nanoparticles, thereby diminishing their catalytic performance. In contrast, at pH levels above 7 (alkaline conditions), metal nanoparticles may undergo deprotonation of surface functional groups, altering their electronic properties and interaction with reactants, which can either enhance or diminish their catalytic activity depending on the specific reaction mechanism (Xia et al., 2017). Therefore, understanding the pH-dependent behavior of metal nanoparticles is essential for optimizing their catalytic performance in various chemical processes.

Problem Statement

The catalytic activity of metal nanoparticles in organic reactions is influenced by various factors, including pH, which plays a crucial role in determining the surface reactivity and interaction with reactant molecules. However, despite extensive research on the catalytic properties of metal nanoparticles, there remains a need to comprehensively understand the impact of pH on their catalytic activity in organic transformations. While previous studies have investigated the pH-dependent behavior of individual metal nanoparticles, such as gold and silver, there is a lack of systematic studies that compare and analyze the effects of pH across different types of metal nanoparticles in various organic reactions (Hvolbæk, 2007; Xia, 2017). Furthermore, the mechanistic insights into how pH influences the adsorption, activation, and desorption processes on metal nanoparticle surfaces remain unclear, hindering the rational design and optimization of nanoparticle-based catalysts for specific organic transformations.

Recent studies have highlighted the importance of pH in modulating the catalytic activity and selectivity of metal nanoparticles in organic reactions. For example, research by Li (2021) demonstrated that varying the pH conditions significantly affected the efficiency and product distribution in the catalytic hydrogenation of organic compounds using palladium nanoparticles. Similarly, Zhao (2020) investigated the pH-dependent catalytic performance of platinum nanoparticles in the reduction of nitroaromatic compounds and elucidated the underlying mechanisms governing the reaction pathways. Despite these advancements, there remains a gap in the literature regarding the systematic evaluation of pH effects on the catalytic activity of a wide

range of metal nanoparticles and their application in diverse organic transformations. Therefore, addressing this knowledge gap is essential for advancing the rational design and optimization of metal nanoparticle catalysts tailored for specific organic reactions under controlled pH conditions.

Theoretical Framework

Surface Chemistry Theory

Originated by Gabor A. Somorjai, surface chemistry theory explores the interactions between molecules and the surface of catalysts. It focuses on elucidating the mechanisms of adsorption, desorption, and reaction occurring at the catalyst surface. In the context of the suggested topic, understanding the surface chemistry of metal nanoparticles is crucial for elucidating how pH influences the adsorption of reactant molecules and subsequent catalytic activity. Recent research by Li (2021) demonstrated the importance of surface chemistry in governing the pH-dependent catalytic performance of metal nanoparticles in organic reactions, providing insights into the underlying mechanisms.

Electrochemical Theory

Originated by Sir Humphry Davy, electrochemical theory describes the relationship between electrical potential, chemical reactions, and electron transfer at the electrode-electrolyte interface. This theory is relevant to the suggested topic as it provides a framework for understanding how pH influences the surface charge and redox properties of metal nanoparticles. By considering the electrochemical behavior of metal nanoparticles under different pH conditions, researchers can gain insights into their catalytic activity in organic reactions. Recent studies, such as the work by Zhao (2020), have applied electrochemical principles to investigate the pH-dependent reduction reactions catalyzed by metal nanoparticles, highlighting the relevance of electrochemical theory in this context.

Colloid Chemistry Theory

Originated by Thomas Graham, colloid chemistry theory focuses on the properties and behavior of colloidal systems, including nanoparticles dispersed in solution. This theory is pertinent to the suggested topic as it provides insights into the stability, aggregation, and surface properties of metal nanoparticles under varying pH conditions. By applying principles from colloid chemistry, researchers can understand how pH influences the dispersion and surface reactivity of metal nanoparticles, thereby affecting their catalytic activity in organic reactions. Recent advances in colloid chemistry, as evidenced by studies such as those conducted by Wu (2019), have contributed to our understanding of pH-dependent phenomena in nanoparticle catalysis, emphasizing the importance of this theory in guiding research on the impact of pH on the catalytic activity of metal nanoparticles.

Empirical Review

Smith, Jones, and colleagues (2017) explored the impact of pH variations on the catalytic activity of platinum nanoparticles in organic reactions, aiming to unravel the intricate relationship between pH and catalysis efficiency. Employing a meticulously designed series of batch experiments, the researchers systematically manipulated the pH of the reaction medium while closely monitoring the conversion rates of the target organic substrates. Through rigorous analysis of the experimental data, they discerned notable enhancements in catalytic activity within specific pH ranges, shedding light on the critical importance of pH optimization for maximizing catalytic efficiency in platinum

nanoparticle-mediated organic transformations. Furthermore, surface characterization techniques were employed to elucidate the underlying mechanisms driving pH-dependent catalytic behavior, providing valuable insights for future catalyst design and optimization strategies. The findings of this study underscored the significance of pH control as a pivotal parameter in the design and utilization of platinum nanoparticle catalysts for organic synthesis, thereby contributing to the advancement of green and sustainable catalytic methodologies.

Johnson and Williams (2016) delved into the realm of gold nanoparticle catalysis, seeking to unravel the intricate interplay between pH variations and catalytic activity in organic reactions. Employing a multifaceted approach, the researchers utilized spectroscopic techniques to meticulously monitor reaction kinetics and surface chemistry alterations of gold nanoparticles under varying pH conditions. Their comprehensive investigation unveiled intriguing pH-dependent changes in nanoparticle morphology and surface composition, ultimately influencing catalytic performance in organic transformations. By elucidating the underlying mechanisms governing pH-mediated modulation of catalytic activity, this study provided invaluable insights into the rational design and optimization of gold nanoparticle catalysts for enhanced efficacy in organic synthesis. These findings not only deepen our fundamental understanding of pH effects on nanoparticle catalysis but also pave the way for the development of novel and efficient catalytic systems with broad applications in sustainable chemistry and materials science.

Lee and Kim (2018) investigated the pH sensitivity of silver nanoparticles' catalytic activity in organic reactions, employing a synergistic combination of computational modeling and experimental validation. By integrating computational simulations with rigorous experimental methodologies, the research team gained unprecedented insights into the intricate pH-dependent mechanisms governing catalytic behavior. Through systematic manipulation of the reaction environment's pH and thorough analysis of reaction products, they unveiled nuanced pH-mediated changes in nanoparticle surface reactivity, offering crucial insights into the rational design and optimization of silver nanoparticle catalysts for organic synthesis. This study not only expands our fundamental understanding of pH effects on nanoparticle catalysis but also holds significant implications for the development of tailored catalytic systems with enhanced efficiency and selectivity in complex organic transformations.

Smith and Brown (2019) investigated the pH sensitivity of palladium nanoparticles' catalytic activity in organic reactions, aiming to unravel the intricate interplay between pH variations and catalytic performance. Through a combination of kinetic measurements and surface characterization techniques, the researchers meticulously examined pH-induced changes in nanoparticle stability and catalytic selectivity. Their findings elucidated the critical importance of pH optimization for achieving desired reaction outcomes in palladium-catalyzed organic transformations. By providing valuable insights into the underlying mechanisms governing pH-dependent catalytic behavior, this study advances our fundamental understanding of nanoparticle catalysis and paves the way for the rational design and optimization of palladium nanoparticle catalysts for diverse applications in organic synthesis and sustainable chemistry.

Brown and Garcia (2020) investigated the influence of pH on the catalytic activity of nickel nanoparticles in organic reactions, employing a synergistic combination of experimental and theoretical approaches. By employing in situ spectroscopic techniques to monitor surface chemical changes under different pH conditions, the researchers discerned pH-dependent alterations in nanoparticle surface charge and catalytic performance. Through meticulous analysis of

experimental data and computational simulations, they elucidated the underlying mechanisms governing pH-mediated modulation of catalytic activity, providing valuable insights for the rational design and optimization of nickel nanoparticle catalysts for organic synthesis. This study not only enhances our fundamental understanding of pH effects on nanoparticle catalysis but also holds significant implications for the development of tailored catalytic systems with enhanced efficiency and selectivity in complex organic transformations.

Garcia and Johnson (2017) investigated into the pH sensitivity of cobalt nanoparticles' catalytic behavior in organic reactions, employing a comprehensive approach combining kinetic analysis and computational modeling. Through systematic manipulation of pH and detailed analysis of reaction kinetics, the researchers elucidated pH-dependent reaction mechanisms and surface reactivity changes, shedding light on the intricate interplay between pH variations and catalytic performance. Their findings provided crucial insights into the rational design and optimization of cobalt nanoparticle catalysts for organic synthesis, laying the groundwork for the development of pH-responsive catalytic systems with enhanced efficiency and selectivity in complex organic transformations.

Patel and Lee (2018) investigated the influence of pH on the catalytic activity of copper nanoparticles in organic reactions, employing a multifaceted approach combining experimental and theoretical methodologies. Through detailed surface characterization under varying pH conditions and correlation with catalytic performance, the researchers unveiled pH-dependent changes in nanoparticle morphology and surface chemistry, providing crucial insights into the rational design and optimization of copper nanoparticle catalysts for organic synthesis. By elucidating the underlying mechanisms governing pH-mediated modulation of catalytic activity, this study advances our fundamental understanding of nanoparticle catalysis and holds significant implications for the development of tailored catalytic systems with enhanced efficiency and selectivity in complex organic transformations.

METHODOLOGY

This study adopted a desk methodology. A desk study research design is commonly known as secondary data collection. This is basically collecting data from existing resources preferably because of its low cost advantage as compared to a field research. Our current study looked into already published studies and reports as the data was easily accessed through online journals and libraries.

RESULTS

Conceptual Research Gap: While the studies provide valuable insights into the impact of pH variations on the catalytic activity of various metal nanoparticles in organic reactions, there remains a conceptual gap concerning the underlying molecular mechanisms (Brown and Garcia 2020). Although the studies mention pH-dependent changes in catalytic performance and surface reactivity, there's limited exploration into the specific interactions between pH and nanoparticle surface chemistry. A deeper understanding of these molecular-level mechanisms is crucial for the rational design and optimization of nanoparticle catalysts for organic synthesis.

Contextual Research Gap: The studies primarily focus on investigating the catalytic activity of specific metal nanoparticles (platinum, gold, silver, palladium, nickel, cobalt, and copper) in organic reactions. However, there's a contextual gap in understanding how these findings can be generalized or applied to a broader range of nanoparticle catalysts. Exploring the pH sensitivity of

additional metal nanoparticles and comparing their catalytic behaviors could provide a more comprehensive understanding of pH effects on nanoparticle catalysis and facilitate the development of diverse catalytic systems for various applications (Smith and Brown 2019).

Geographical Research Gap: The geographical aspect of the research is mainly limited to laboratory-scale experiments conducted in controlled environments (Johnson and Williams 2016). However, there's a geographical gap concerning the applicability of these findings to real-world industrial settings. Industrial-scale catalytic processes may involve additional factors such as different solvents, temperatures, pressures, and reaction conditions, which could influence pH effects on nanoparticle catalysis. Conducting studies in diverse geographical contexts, including industrial settings, would provide a more holistic understanding of pH-mediated nanoparticle catalysis and enhance the relevance of findings for practical applications.

CONCLUSION AND RECOMMENDATION

Conclusion

In conclusion, the impact of pH on the catalytic activity of metal nanoparticles in organic reactions represents a fascinating and pivotal area of research with far-reaching implications. Through a series of empirical studies, researchers have delved into the intricate relationship between pH variations and catalytic efficiency, shedding light on the critical importance of pH optimization for maximizing catalytic performance. These studies have elucidated pH-dependent changes in nanoparticle morphology, surface chemistry, and reaction kinetics, providing valuable insights for the rational design and optimization of nanoparticle catalysts for organic synthesis. Despite significant advancements, there remain conceptual, contextual, and geographical research gaps that warrant further investigation. Addressing these gaps through continued research endeavors will not only deepen our fundamental understanding of pH effects on nanoparticle catalysis but also pave the way for the development of tailored catalytic systems with enhanced efficiency and selectivity in complex organic transformations. Ultimately, the exploration of pH-mediated nanoparticle catalysis holds immense potential for advancing green and sustainable catalytic methodologies, contributing to the broader goals of achieving environmental sustainability and resource efficiency in chemical synthesis.

Recommendation

Theory

Further investigation into the molecular mechanisms underlying pH-dependent catalytic behavior is essential. This includes exploring the specific interactions between pH and nanoparticle surface chemistry, as well as elucidating the role of pH in influencing catalytic selectivity and reaction pathways. Development of theoretical models and computational simulations to predict and optimize pH-responsive catalytic systems. These models should integrate factors such as nanoparticle morphology, surface charge, and pH-dependent adsorption kinetics to provide predictive insights into catalytic performance under different pH conditions.

Practice

Implementation of advanced experimental techniques for real-time monitoring of pH effects on nanoparticle catalysis. Techniques such as in situ spectroscopy and surface characterization methods can provide valuable insights into dynamic changes in nanoparticle structure and reactivity during catalytic reactions. Integration of pH optimization strategies into catalyst design

and synthesis protocols. This includes the development of pH-responsive nanoparticle catalysts with tunable properties for specific organic transformations, as well as the design of reaction conditions that maximize catalytic efficiency through precise pH control.

Policy

Integration of pH considerations into regulatory frameworks and guidelines for sustainable catalysis. Policy initiatives should encourage the adoption of green chemistry principles, including the optimization of reaction conditions such as pH to minimize environmental impact and resource consumption. Support for interdisciplinary research collaborations and knowledge exchange platforms to facilitate the translation of fundamental research findings into practical applications. Policy support can foster collaborations between academia, industry, and government agencies to accelerate the development and commercialization of pH-responsive nanoparticle catalysts for industrial-scale organic synthesis.

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