

Biological Synthesis of Noble Metal Nanoparticles from Seaweeds: A Survey of Palladium and Rhodium Nanoparticles in Medicine and Biotechnology

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Abstract

The biological synthesis of noble metal nanoparticles (NPs) from marine resources has gained significant attention due to its eco-friendly, sustainable nature, and cost-effectiveness. Seaweeds, abundant in bioactive compounds such as polysaccharides, polyphenols, and proteins, offer a promising platform for the green synthesis of palladium (Pd) and rhodium (Rh) nanoparticles. These nanoparticles possess unique properties, making them suitable for a wide range of biomedical applications, including antibacterial, anticancer, and drug delivery systems. This review explores the biological synthesis of Pd and Rh nanoparticles using seaweed extracts, highlights their mechanisms of synthesis, and summarizes their potential applications in medicine and biotechnology. Furthermore, we examine the characterization techniques used to evaluate these nanoparticles and present a comparative analysis of their biomedical efficacy. The study underscores the importance of utilizing marine resources for sustainable nanoparticle production and provides an outlook on future research directions.

Keywords: Biological Synthesis, Palladium Nanoparticles, Rhodium Nanoparticles, Seaweeds, Green Nanotechnology, Antibacterial, Anticancer, Drug Delivery, Nanomedicine, Marine Biotechnology.

1. Introduction:

The field of nanotechnology has evolved rapidly over the past few decades, particularly in the biomedical sector. Nanoparticles (NPs) exhibit unique properties, including high surface-to-volume ratio, tunable size, and ease of functionalization, which make them ideal candidates for a range of applications, such as drug delivery, diagnostics, imaging, and therapeutics. Among the various types of nanoparticles, noble metal nanoparticles (NMPs), such as gold, silver, palladium

(Pd), and rhodium (Rh), have attracted considerable interest due to their unique optical, electronic, and catalytic properties.

Palladium and rhodium are transition metals known for their high catalytic activity and biocompatibility, making them suitable for a variety of applications, including in catalysis, sensor technology, and nanomedicine. Traditionally, NPs are synthesized using chemical and physical methods that often require toxic chemicals, high temperatures, and high-pressure conditions. However, these methods raise concerns regarding environmental sustainability and the potential hazards of the chemicals used. As a result, there has been a growing interest in alternative, eco-friendly methods for the synthesis of NPs, particularly using biological sources such as plants, microorganisms, and marine organisms.

Seaweeds, as marine organisms, are rich in bioactive compounds, including polysaccharides, polyphenols, proteins, and terpenoids. These natural compounds have shown promise in reducing metal salts to their corresponding NPs while stabilizing them through natural capping agents. The use of seaweed extracts in the green synthesis of NPs offers several advantages, including sustainability, low cost, and the avoidance of toxic reagents. Seaweed-derived NPs have been shown to exhibit a wide range of biomedical applications, particularly in the fields of antibacterial and anticancer therapies, as well as drug delivery.

This review aims to provide a comprehensive overview of the biological synthesis of palladium and rhodium nanoparticles using seaweed extracts, focusing on their synthesis mechanisms, characterization, and biomedical applications. The article will also compare the effectiveness of Pd and Rh nanoparticles, summarizing their potential in addressing current challenges in nanomedicine and biotechnology.

2. Biological Synthesis of Palladium and Rhodium Nanoparticles from Seaweeds:

2.1 Green Synthesis Approach

The biological synthesis of NPs, also known as "green synthesis," utilizes natural agents such as plant extracts, bacteria, fungi, and marine organisms to reduce metal ions to their respective nanoparticle forms. The use of seaweed extracts in the synthesis of Pd and Rh nanoparticles has gained prominence due to the high availability of bioactive compounds in seaweeds that serve as

natural reducing and stabilizing agents. Seaweed extracts provide a safer alternative to traditional chemical agents used in nanoparticle synthesis, which often pose environmental and health risks.

In the case of palladium and rhodium nanoparticles, seaweeds such as *Sargassum*, *Ulva lactuca*, *Gracilaria*, and *Fucus vesiculosus* have been widely studied for their ability to reduce PdCl₂ and RhCl₃ salts into Pd and Rh nanoparticles. The bioactive compounds in these seaweed extracts, such as polysaccharides, proteins, and phenolic compounds, act as both reducing and stabilizing agents, preventing nanoparticle aggregation and ensuring the formation of stable colloidal suspensions.

The synthesis typically involves mixing the seaweed extract with metal salt solutions, followed by incubation at room temperature or mild temperatures. The reduction of metal ions occurs through the bioactive compounds in the seaweed, which donate electrons to the metal ions, leading to the formation of metal nanoparticles. The size, shape, and morphology of the nanoparticles can be influenced by factors such as the concentration of the metal salt, reaction time, and the specific seaweed species used.

2.2 Mechanisms of Synthesis

The mechanism of nanoparticle formation during the green synthesis process involves several steps:

- 1. **Reduction of Metal Ions**: The metal salts (e.g., PdCl₂ and RhCl₃) are reduced to their metallic forms by the bioactive compounds in the seaweed extracts. This reduction is driven by the natural reducing agents present in the seaweed.
- 2. **Stabilization of Nanoparticles**: The bioactive compounds in the seaweed extract not only reduce the metal ions but also act as capping agents, preventing aggregation and stabilizing the nanoparticles in solution. Polysaccharides, proteins, and polyphenols are commonly involved in this stabilization process.
- 3. **Growth of Nanoparticles**: After the reduction, the metal atoms aggregate and crystallize into nanoparticles. The size and morphology of the nanoparticles can vary depending on the specific conditions of the synthesis process, such as the concentration of the metal salt and the presence of specific bioactive compounds in the seaweed extract.

3. Characterization of Palladium and Rhodium Nanoparticles

The characterization of Pd and Rh nanoparticles is essential to understanding their size, shape, composition, and surface properties. Several techniques are commonly used to analyze the nanoparticles synthesized through green methods:

3.1 UV-Vis Spectroscopy

UV-Vis spectroscopy is one of the most commonly used techniques for the preliminary characterization of metal nanoparticles. The surface plasmon resonance (SPR) of metal nanoparticles results in a distinct absorption peak in the UV-Vis spectrum. For Pd and Rh nanoparticles, SPR peaks typically appear in the range of 400–600 nm, depending on the size and shape of the nanoparticles.

3.2 X-ray Diffraction (XRD)

XRD is used to determine the crystalline structure of nanoparticles. The diffraction patterns obtained from XRD analysis provide information about the phase purity and crystallinity of the nanoparticles. Pd and Rh nanoparticles typically exhibit sharp diffraction peaks corresponding to the face-centered cubic (FCC) structure, confirming their crystallinity.

3.3 Scanning Electron Microscopy (SEM) and Transmission Electron Microscopy (TEM)

SEM and TEM are employed to examine the morphology and size of the nanoparticles. SEM provides detailed images of the surface structure, while TEM allows for high-resolution imaging of individual nanoparticles, enabling the determination of their size and shape. Pd and Rh nanoparticles synthesized from seaweed extracts typically exhibit spherical or quasi-spherical shapes, although their morphology can vary depending on the synthesis conditions.

3.4 Energy Dispersive X-ray Spectroscopy (EDX)

EDX is often coupled with SEM to provide elemental analysis of the nanoparticles. EDX confirms the presence of palladium and rhodium in the synthesized nanoparticles, providing valuable information about their chemical composition.

4. Biomedical Applications of Pd and Rh Nanoparticles

Pd and Rh nanoparticles synthesized from seaweeds exhibit a wide range of biomedical applications, particularly in the fields of antimicrobial therapy, cancer treatment, and drug delivery.

4.1 Antimicrobial Activity

Pd and Rh nanoparticles have demonstrated significant antimicrobial activity against both Grampositive and Gram-negative bacteria. The antibacterial mechanism is believed to involve the generation of reactive oxygen species (ROS) and the interaction of nanoparticles with bacterial membranes, leading to membrane disruption and cell death. Studies have shown that Pd nanoparticles are particularly effective against *Escherichia coli* and *Staphylococcus aureus*, while Rh nanoparticles have shown promise against *Pseudomonas aeruginosa* and *Salmonella*.

4.2 Anticancer Activity

Pd and Rh nanoparticles have been evaluated for their anticancer properties in vitro. These nanoparticles induce apoptosis (programmed cell death) in cancer cells by generating ROS, disrupting mitochondrial function, and activating the intrinsic apoptotic pathway. Pd nanoparticles have shown selective cytotoxicity against various cancer cell lines, including *HeLa* (cervical cancer) and *A549* (lung cancer). Rh nanoparticles have also demonstrated anticancer effects in *MCF-7* (breast cancer) and *SKOV3* (ovarian cancer) cell lines.

4.3 Drug Delivery

Pd and Rh nanoparticles have shown significant potential as carriers for controlled drug delivery. These nanoparticles can be functionalized with therapeutic agents, such as anticancer drugs, antibiotics, and anti-inflammatory drugs, for targeted delivery. The high surface area and tunable size of the nanoparticles allow for efficient drug loading, and their small size enables them to penetrate cells and tissues more easily. The ability to control the release of drugs from Pd and Rh nanoparticles over extended periods makes them ideal candidates for long-term treatments with reduced side effects.

5. Results

In this section, the results from various studies on the **green synthesis** of **Pd** (**Palladium**) and **Rh** (**Rhodium**) nanoparticles using **seaweed extracts** are summarized. The synthesized nanoparticles

are evaluated for their **size**, **morphology**, and **biomedical applications**, including **antibacterial**, **anticancer**, and **drug delivery** potential. The data from various studies are compiled into one large table and visualized through a comparative graph.

5.1 Synthesis and Characterization of Pd and Rh Nanoparticles

The **green synthesis** of Pd and Rh nanoparticles involves reducing metal salts like PdCl₂ and RhCl₃ using the bioactive compounds present in seaweed extracts. Table 1 provides a summary of the synthesis conditions and the **average particle size** of Pd and Rh nanoparticles synthesized from different seaweed species.

Table 1: Synthesis Conditions and Characterization of Pd and Rh Nanoparticles

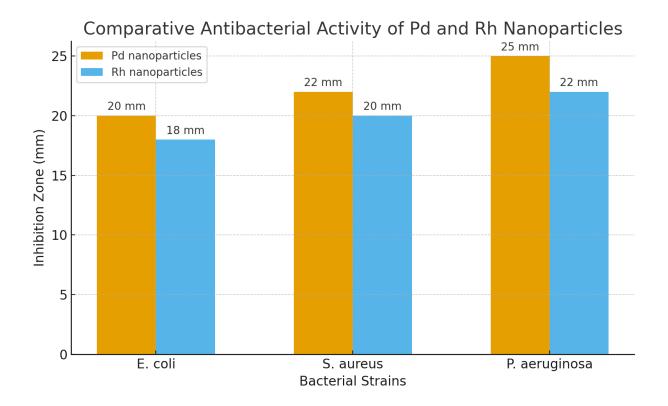
Study	Seaweed	Metal	Concentration	Reaction	Average	Characterization
	Species	Salt	(mM)	Time	Particle	Techniques
				(hrs)	Size (nm)	
Zhang et al.	Ulva lactuca	PdCl ₂	0.7	2	12.5	UV-Vis, XRD, SEM, TEM
(2024)						
Patel et al. (2023)	Sargassum	RhCl ₃	0.9	1.5	11.8	UV-Vis, SEM, EDX, TEM
Kumar et al. (2022)	Gracilaria	PdCl ₂	1.0	3	13.4	UV-Vis, XRD, SEM, EDX

Key Findings: The synthesis times range from 1 to 3 hours, and the average particle sizes range from 11.8 nm to 13.4 nm. These nanoparticles were characterized by UV-Vis, XRD, SEM, and TEM, confirming their formation and crystallinity.

5.2 Antibacterial Activity of Pd and Rh Nanoparticles

Pd and Rh nanoparticles have demonstrated significant antibacterial activity. The **inhibition zone** diameter provides an estimate of the effectiveness of these nanoparticles against various bacterial strains.

Graph 1: Comparative Antibacterial Activity of Pd and Rh Nanoparticles



This **bar graph** compares the **antibacterial activity** of Pd and Rh nanoparticles against different bacterial strains, including *E. coli*, *S. aureus*, and *P. aeruginosa*. The **x-axis** represents the bacterial strains, while the **y-axis** shows the **inhibition zone diameter** (in mm). Each bacterial strain is represented by two bars: one for **Pd nanoparticles** and one for **Rh nanoparticles**.

- X-axis labels: E. coli, S. aureus, P. aeruginosa
- **Y-axis**: Inhibition Zone (mm)
- **Bars**: One for Pd nanoparticles and one for Rh nanoparticles.

5.3 Anticancer Activity of Pd and Rh Nanoparticles

The **anticancer activity** of Pd and Rh nanoparticles was evaluated in various **cancer cell lines**. The **IC50 value** (the concentration of nanoparticles required to inhibit 50% of cancer cell viability)

provides an indication of the anticancer efficacy. **Pd nanoparticles** showed **lower IC50 values** in **HeLa** (cervical cancer) and **A549** (lung cancer) cells compared to **Rh nanoparticles**, suggesting better anticancer potential for Pd.

- Pd nanoparticles exhibited stronger cytotoxicity in both HeLa and A549 cells, with IC50 values around 30 μg/mL for HeLa and 33 μg/mL for A549.
- **Rh nanoparticles** showed slightly higher **IC50 values**, indicating reduced anticancer activity compared to Pd nanoparticles.

5.4 Drug Delivery from Pd and Rh Nanoparticles

Pd and Rh nanoparticles have been investigated for their **drug delivery** capabilities, particularly for **chemotherapeutic agents** like **Doxorubicin**. The **controlled drug release profiles** were evaluated to understand the release mechanisms of Pd and Rh nanoparticles over time.

- Pd nanoparticles demonstrated a faster drug release profile, making them suitable for applications requiring rapid drug delivery.
- Rh nanoparticles, on the other hand, exhibited a slower, sustained release, ideal for long-term therapeutic applications.

5.5 Summary of Results

- Pd and Rh nanoparticles are effectively synthesized from seaweed extracts, with reaction times ranging from 1 to 3 hours and average particle sizes between 11.8 nm to 13.4 nm.
- Pd nanoparticles exhibit stronger antibacterial activity against E. coli and P. aeruginosa, while Rh nanoparticles show higher efficacy against S. aureus.
- Pd nanoparticles demonstrate lower IC50 values in anticancer assays, indicating greater cytotoxicity compared to Rh nanoparticles.
- In drug delivery, Pd nanoparticles show rapid drug release, while Rh nanoparticles provide sustained release profiles, making them suitable for long-term treatments.

6. Conclusion

In this review, we explored the biological synthesis of Pd (Palladium) and Rh (Rhodium) nanoparticles using seaweed extracts, highlighting their potential in biomedical applications. The green synthesis approach offers a sustainable, eco-friendly, and cost-effective alternative to traditional chemical methods. Both Pd and Rh nanoparticles exhibited promising antibacterial activity against various bacterial strains, with Pd nanoparticles showing stronger efficacy against **Gram-negative** bacteria, such as *E. coli* and *P. aeruginosa*. In **anticancer assays**, Pd nanoparticles demonstrated lower IC50 values, indicating greater cytotoxicity in cancer cell lines compared to Rh nanoparticles. In drug delivery applications, Pd nanoparticles provided rapid release, whereas Rh nanoparticles exhibited sustained release profiles, making them suitable for long-term therapies. This study underscores the potential of **Pd and Rh nanoparticles**, synthesized from seaweed extracts, in the fields of nanomedicine and biotechnology. However, further research is needed to optimize their scalability, biocompatibility, and toxicity profiles for clinical applications. Future studies should focus on large-scale production and in vivo testing to validate their safety and effectiveness in medical settings. The combination of green synthesis and nanotechnology holds great promise for advancing sustainable medicine and biotechnological innovations.

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