Characterization of Rajath Bhasma and Evaluation of Its Toxicity in Zebrafish Embryos and Its Antimicrobial Activity

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Introduction

Siddha medicine was developed by ancient Tamil sages known as ‘Siddhars’ and has been in use for more than 5,000 years [1]. It involves the use of gold, zinc and silver formulated with honey, ghee, or milk to cure diseases [2, 3]. Prototypic examples include Swarna Bhasma (gold ash), Jasada Bhasma (zinc ash), and Rajath Bhasma (silver ash), which are prepared by processing a fine powder of gold, zinc, and silver with plant extract, followed by repeated incineration at high temperature (~1,000°C). During incineration, the size of the fine powder of gold, zinc, and silver is reduced to that of chemically synthesized nanoparticles (1-100 nm). This fine powder is composed of individual particles of around 50-70 nm in size; however, these particles are prone to forming large aggregates, unlike chemically synthesized nanoparticles [4].

Compared with gold-derived Swarna Bhasma and zinc-derived Jasada Bhasma, the silver-derived medicine Rajath Bhasma has superior and broad-spectrum antimicrobial, antifungal, and antiviral activities [5] and hence numerous applications. Notably, Rajath Bhasma has been used to treat various disorders including anxiety, aging, and infertility [6-8]. The physical characteristics of a silver preparation are believed to markedly influence its therapeutic properties; however, despite its widespread use in India, the exact chemical composition/structure of Rajath Bhasma and its biological activities are largely unknown. Current tools for characterizing these particles at the nanoscale level have furthered the understanding of the parameters directly or indirectly influencing the efficacy of these ancestral medicines. Accordingly, efforts have been made to study these particles at the nanoscale level. For example, Mitra et al. [9] have characterized Swarna Bhasma (gold ash) and Jasada Bhasma (zinc ash) and Rohit et al. [10] reported the synthesis of Rajath Bhasma through a simple characterization method.

In general, Rajath Bhasma is safe for therapeutic applications [9]; however, in vivo studies are required to accurately predict its toxicity. Zebrafish embryos are a well-established in vivo model for evaluating teratogenicity.
Although several studies have investigated the teratogenic effects of silver nanoparticles on zebrafish embryos [11-14], none of them have characterized Rajath Bhasma and assessed its toxicity among embryos at different concentrations to determine the effective concentration for antimicrobial activity. In this study, we characterized Rajath Bhasma by analyzing its surface functional groups, particle size, elemental composition, and crystalline phase. In addition, we evaluated the toxicity of Rajath Bhasma among zebrafish embryos and its antimicrobial activity in terms of bacterial growth inhibition.

Materials and Methods

Characterization of Rajath Bhasma

Rajath Bhasma used in the experiments was purchased from Dabur India Ltd. (India), and analyzed via Fourier-transform infrared (FTIR) spectrometry (Shimadzu 8400; Shimadzu, Japan), scanning electron microscopy (SEM), (EVO 18; Carl Zeiss, Germany), energy-dispersive X-ray (EDX) spectrometry (Quantax 200 with X-Flash; Bruker, USA), dynamic light scattering (DLS) (HORIBA Scientific SZ-100, UK), and X-ray diffraction (XRD), (D8 Advance ECO XRD system with SSD160 1-D detector; Bruker).

FTIR Analysis

FTIR analysis was performed to determine the surface functional groups of Rajath Bhasma. In brief, a sample was illuminated with infrared light in the wavelength range of 2.5-25 μm, and the transmittance spectrum was obtained to determine the vibrational frequencies corresponding to specific functional groups [15].

SEM and EDX Analyses

Particle size and shape of Rajath Bhasma were determined via SEM. For SEM analysis, Rajath Bhasma was dispersed in distilled water, sonicated for 2 min, dropped on a carbon tape, and then dried. The elemental composition was assessed using an EDX spectrometer [15].

DLS Analysis

The average size of Rajath Bhasma was determined via DLS analysis. To determine the size distribution, Rajath Bhasma was dispersed in water for 2 min and the average particle size and standard deviation were determined using the analyzer software.

XRD Analysis

XRD was used to determine the crystalline phases of Rajath Bhasma. In brief, a thin layer of Rajath Bhasma powder was placed in conventional cavity mounts of the X-ray diffractometer and scanned from 10° to 80° for 2 h. The Cu anode X-ray was operated at 40 kV and 30 mA to generate monochromatic Cu K-α X-rays (k = 1.54056 Å). The average crystallite size of Rajath Bhasma was calculated from the XRD graph using the Debye–Scherrer equation [16]:

\[
d = \frac{0.9 \lambda}{\beta \cos \theta}
\]

where d is the mean crystallite size, λ is the X-ray wavelength, and β is the angular full width at half maximum (FWHM) of the peak at diffraction angle θ.

Assessment of Embryonic Toxicity

Zebrafish embryos were purchased at a local aquarium shop. All purchases were made in one shop to obtain a relatively uniform group of zebrafish embryos. Zebrafish embryos at the early blastula stage were transferred to a petri plate at 10 embryos/petri plate. These embryos were treated with Rajath Bhasma at different concentrations (5, 10, 15, 20, and 25 μg/ml) in 10 ml distilled water for 48 h at room temperature. At 72 h post fertilization (hpf), malformation rates (%) were calculated as the percentage of dead embryos relative to the total number of embryos to estimate embryonic toxicity [15]. Edema, the most common malformation, along with tail and yolk abnormalities were macroscopically quantified. Each experiment was performed in triplicate [17].

Assessment of Antimicrobial Activity

The antimicrobial activity of Rajath Bhasma against gram-positive (Staphylococcus epidermidis) and gram-negative (Escherichia coli) bacteria was determined using the well-diffusion method [18]. In brief, bacteria were diluted to 3.2 × 10^8 colony forming units/ml and spread on Luria–Bertani agar plates. Thereafter, 6-mm-diameter wells were created in the agar plates by using a sterile borer and 5 μg/ml of Rajath Bhasma was then placed in each well. After incubation at 37°C for 28 h, inhibition zones were measured and the mean of four replicates was calculated.

Results and Discussion

Characterization of Rajath Bhasma

FTIR analysis. Functional groups on the surface of Rajath Bhasma were characterized by studying the vibrational frequencies in the FTIR spectrum (Fig. 1). The broad peak at 3,433 cm⁻¹ corresponds to an H-bonded hydroxyl group. The presence of a hydroxyl group is attributed to the preparation method of Rajath Bhasma, in which plant compounds are used to modify the surface of the silver core [19]. The peak at 2,923 cm⁻¹ corresponds...
to asymmetric C–H stretching of an aldehyde group and is generally observed in the spectra of biologically synthesized nanoparticles because aldehyde groups are present in the biological component [20]. The sharp peak at 2,360 cm⁻¹ corresponds to C=C stretching of the aromatic ring in flavonoids, compounds present in nearly all plants [20]. The peaks at 1,642 cm⁻¹ and 1,442 cm⁻¹ correspond to C=O stretching and CH₂ bending vibrations of aliphatic compounds, respectively [21, 22]. The major stretching peak at 1,334 cm⁻¹ is assigned to amide groups, which are abundant in plant extracts [23]. Together, these results indicate that the surface of the silver core is adorned with organic compounds including plant or herbal extracts.

SEM and EDX analyses. The particle size of Rajath Bhasma was analyzed via SEM. Fig. 2A shows that the nanoparticles formed spherical aggregates, and the size of individual aggregates ranged from 170 to 210 nm, consistent with the results of DLS analysis (Fig. 2B). In addition, EDX analysis confirmed that silver is the major component (approximately 79%) of Rajath Bhasma (Fig. 2C).

XRD analysis. XRD analysis was performed to identify the crystalline phases in Rajath Bhasma. The XRD pattern showing sharp diffraction peaks at 29.79°, 32.28°, 33.64°, and 38.116°, which correspond to crystalline planes of (011), (111), (101), and (111), respectively, indicated pure crystalline silver structures (Fig. 3). The XRD
results were concurrent with the Joint Committee on Powder Diffraction Standards (JCPDS) data for silver oxalate (Ag₂C₂O₄; JCPDS 22-1335). The peaks at 32.28 (111), 33.64 (101), and 38.116 (111) also correspond to AgO (JCPDS 84-1547), Ag₂CO₃ (JCPDS 71-2184), and Ag (JCPDS 65-2871), respectively. The average crystallite size of Rajath Bhasma was calculated using the Debye–Scherrer equation. Table 1 shows that the average crystallite size was approximately 55 nm [24-26].

Toxic Effects of Rajath Bhasma on Zebrafish Embryos
Zebrafish embryos at an early blastula stage were exposed to Rajath Bhasma for 48 h and embryonic malformation rates were determined at 72 hpf. As shown in Fig. 4A, 5 μg/ml of Rajath Bhasma did not substantially inhibit the growth of zebrafish embryos [16]. However, the malformation rate was approximately 60% at 25 μg/ml (Fig. 4). These results indicate the importance of determining optimal therapeutic concentrations to inhibit bacterial growth without inducing developmental toxicity in zebrafish.

Antimicrobial Activity of Rajath Bhasma against S. epidermidis and E. coli
Silver nanoparticles are one of the most effective antimicrobial agents; thus, the antimicrobial activity of Rajath Bhasma was investigated by assessing its antibacterial effect. Inhibition zones were clearly observed for both

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**Table 1. Crystallite sizes of Rajath Bhasma determined via XRD analysis.**

| 2θ     | FWHM (dθ) | Crystallite size (nm) |
|--------|-----------|-----------------------|
| 36.2106| 0.2085    | 63.2                  |
| 38.0529| 0.1601    | 81.8                  |
| 45.9921| 0.8829    | 14.46                 |
| 47.4855| 0.2712    | 46.7                  |
| 51.7401| 0.1799    | 69.3                  |

XRD, X-ray diffraction; FWHM, angular full width at half maximum.

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**Fig. 3. X-ray diffraction spectrum of Rajath Bhasma.**

**Fig. 4. Zebrafish embryo toxicity of Rajath Bhasma.** (A) Malformation rates of zebrafish embryos at different concentrations of Rajath Bhasma. The photographs show zebrafish embryos in the absence of Rajath Bhasma (B, C) and embryos exposed to 25 μg/ml Rajath Bhasma (D, E).
gram-positive and gram-negative bacteria. The inhibition zone for *S. epidermidis* treated with 5 μg/ml Rajath Bhasma (i.e., the concentration at which no embryo toxicity was observed; Fig. 4) was 4 ± 0.5 mm and that for *E. coli* was 5 ± 0.5 mm, confirming its effective antimicrobial activity (Fig. 5). Notably, the growth of gram-positive bacteria, which have a thick peptidoglycan layer, was inhibited as effectively as that of gram-negative bacteria. This was attributed to the presence of proton-donating hydroxyl and amide groups on the surface of Rajath Bhasma, as evident from FTIR (Fig. 1) [10].

In conclusion, we characterized Rajath Bhasma using FTIR, SEM, DLS, EDX, and XRD to determine the factors affecting its biological activities and functions. We identified various types of surface functional groups derived from organic components and we confirmed that silver nanoparticles, with a size range of 170–210 nm, are the major component in Rajath Bhasma. Our results show that Rajath Bhasma is a potentially effective antimicrobial agent without toxicity when used at a low concentration (<5 μg/ml). To our knowledge, this is the first study to evaluate the chemical properties, antimicrobial activity, and zebrafish embryonic toxicity of Rajath Bhasma. Our findings will not only expand the application range of Rajath Bhasma but also guide future studies aimed at developing novel nanomaterials.

**Acknowledgments**

This work was supported by Konkuk University Researcher Fund in 2018 and Korea Institute of Energy Technology Evaluation and Planning (KETEP) and the Ministry of Trade, Industry and Energy (MOTIE, 2019-010201900).

**Conflict of Interest**

The authors have no financial conflicts of interest to declare.

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