



An Antimicrobial Evaluation Of Tuttha Bhasma

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Abstract:

Tuttha Bhasma is a herbo-mineral preparation explained in Rasatarangini. The procedure includes shodhana of tuttha, gandhaka, tankana, which are mixed homogenously and triturated with Lakoocha swarasa and subjected to Kukkuta puta. This article explains about the antimicrobial activity of Tuttha Bhasma. 3 samples of tuttha bhasma was prepared and evaluated for its antimicrobial property in 4 strains. Antimicrobial study shows that, Tuttha Bhasma is having better antifungal properties than anti-bacterial properties.

Key words: *Tuttha Bhasma; Tuttha; Anti-Microbial; Anti-Fungal.*

Introduction:

Tuttha –copper Sulphate [CuSO₄7H₂O] is the artificially prepared and presently available form of Sasyaka (copper Sulphate). Tuttha is also considered as the upadhatu of Tamra and have the properties of Tamra and Sasyaka both¹. The Tuttha is also called by Tutathaka, Sikhigreeva, Hemarasa and Mayoorkam in Ayurveda. Tuttha is a chemical compound obtained though processing the Sulphuric acid over the copper. Tuttha reflects the color similar to the neck of a peacock and is heavy in weight. The Tuttha has an unpleasing taste of alkali and is bitter. This is capable to penetrate into the body and to remove ringworms. It is easy to digest and very effective in curing eye diseases, scabies and ailments generated by worms and poison².

Tuttha Bhasma is a herbo-mineral preparation explained in Rasatarangaini³. The procedure includes shodhana of tuttha, gandhaka, tankana, which are mixed homogenously and triturated with Lakoocha swarasa and subjected to Kukkuta Puta for 2 times. Tuttha Bhasma is chiefly indicated in Kushta, Svitra, Amlapitta, and Krimi⁴.

Antimicrobial activities of any therapeutic agent are understood by its degree of growth inhibition of microorganisms as well as bacterial property.

Usually different microbial species and strains have different degrees of susceptibility to therapeutic agents. The susceptibility of microorganisms can change with time even during therapy with a specific drug. Thus, it is essential for the physician to know the sensitivity of the pathogen before treatment.

Our ancient Indian scholars were aware of the existence of microorganisms or bacteria as well as causation of disease since Vedic period.

There are many references pertaining to Jivanuvada (bacteriology) in ancient literature such as Rigveda, Atharvaveda and Mahabharata etc which indicates familiarity of the subject in those days.

Materials and method:

Pharmaceutical study:

The ingredients are shuddha tuttha, shuddha gandhaka, shuddha tankana and Lakucha swarasa (*Atrocarpus lakoocha*). The powders of above said ingredients were mixed homogenously and triturated with Lakucha swarasa. It was subjected to Kukkuta Puta (a defined quantum of heat provided by cow-dung cakes). 3 samples of tuttha bhasma was prepared. Sample 1 was prepared by 3 kukkuta puta and sample 2 and 3 were prepared by 2 kukkuta puta.

Analytical study:

3 samples of tuttha bhasma was analysed as per classical parameters of bhasma pariksha and also using modern analytical parameters like organoleptic characteristics, physico chemical analysis, qualitative and quantitative analysis with the relevant instrumental methods like X-ray diffraction, laser diffraction method, SEM-EDAX.

Antimicrobial study:

Sample- Tuttha Bhasma 3 samples

Bacterial strains used

- **Gram negative strain** – Escherichia coli and Pseudomonas aeruginosa
- **Gram positive strain** – Staphylococcus aureus
 - a. **Methicillin sensitive(MSSA)**
 - b. **Methicillin resistant(MRSA)**
- **Fungus strain** – Candida albicans

Media used – Muller Hinton agar

Sabouraud's dextrose agar

Standard drug disc – Imipenem 10 microgram per disc

Ketaconazole 15 microgram per disc

Preparation of inoculum

For preparation of inoculum, identified clinical strain of all bacteria and fungi was inoculated in peptone water (Identification of the organism was done using standard procedure). It was kept undisturbed for 4 to 5 hours to multiplication. The obtained inoculum was compared with 0.5 McFarland standard optical density of 530 nm corresponding to 5×10^6 CFU/ml. The sterile swab containing organisms was squeezed to remove excess broth and swabbed on the culture plate with lawn culture.

Preparation of drug dilution

Each drug was suspended in distilled water at a concentration of 1mg/ml

Agar well preparation for the drug

Sterile petri dish was taken and Mueller Hinton agar was poured and left for solidification. Each plate was labelled. Standard cultures containing organisms were uniformly smeared all around the surface of the agar medium with sterile cotton swabs containing 10^5 CFU/ml of the bacterial suspension. Wells of 6 mm diameter were dug in the culture medium in the petri plates. 100 microlitres of drug suspension was loaded into each well. These petri plates were incubation at 37°C for 18-24 hours for bacterial growth. Sensitivity of the drug was measured by measuring the zone of inhibition. The same procedure was followed for fungus, Candida albicans using Sabouraud's dextrose agar as culture medium and incubating at 37°C for 48 hours and looked for zone of inhibition.

Results and discussion:

Table no: 1 showing antimicrobial activity of Tuttha Bhasma

Name of organism	Zone diameter				
	Test drug			Standard drug	
	Sample 1	Sample 2	Sample 3		
Escherichia coli	15 mm	15 mm	17 mm	25 mm	
Pseudomonas aeruginosa	17 mm	19 mm	21 mm	25mm	
Staphylococcus aureus	MRSA	25 mm	25 mm	24 mm	25 mm
	MSSA	24 mm	26 mm	28 mm	
Candida albicans	30 mm	35 mm	40 mm	25 mm	



Figure no:1 antimicrobial activity in E.coli strain



Figure no: 2 showing antimicrobial activity in pseudomonas aeruginosa

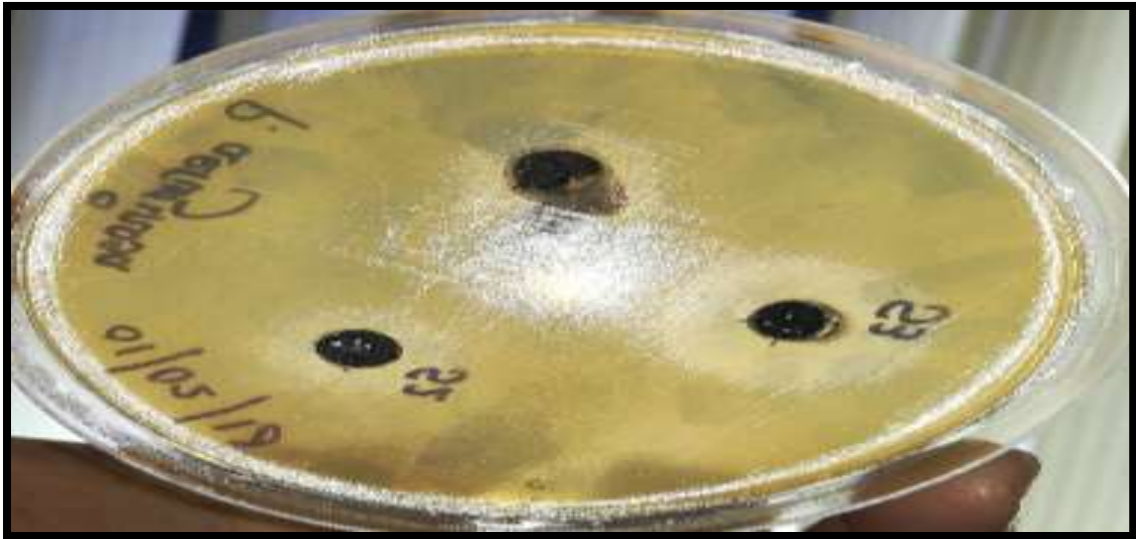


Figure no: 3 showing antimicrobial activity in staphylococcus aureus(MRSA)



Figure no: 4 showing antimicrobial activity in staphylococcus aureus(MSSA)



Figure no: 5 showing antimicrobial activity in Candida albicans

The antimicrobial activities of any therapeutic agents are understood by the degree of growth inhibition of microorganisms it produces as well as its bacterial side of property using different microbial species or even

strains have different degree of susceptibility to therapeutic agents. The susceptibility of microorganisms can change with time, even during therapy to a specific drug⁵. Thus it is essential to the physician to know the efficacy of drug before treatment. Krimighna property of Tuttha bhasma evaluated by antibacterial, antifungal effect.

Bacterial strains used includes Gram negative staphylococcus aerues, E.coli, Pseudomonas aeruginosa and gram positive strains staphylococcus aerues and also methicillin sensitive and resistant strain and Candida albicans as fungal strain. All the samples shown mild anti-bacterial activity against E.coli where the zone of inhibition were 15 and 17 mm compared to 25 mm of standard drug. Moderate activity was shown against pseudomonas aeruginosa (zone of inhibition up to 21 mm).

Good antibacterial activity was observed against staphylococcus aerues (both methicillin sensitive and resistant). Excellent antifungal activity was observed in Tuttha bhasma samples where the maximum zone of inhibition up to 40 mm was observed which was better than all standard drug. Hence the sample of Tuttha bhasma found to be effective in having antibacterial and antifungal property.

Comparatively Tuttha Bhasma found to be more effective in fungus rather than bacteria. CuS was found to be present in Tuttha Bhasma and Tamra Bhasma. Previous study has established that Tamra Bhasma has good in-vitro antibacterial activity⁶.

A study has shown that tuttha has moderate antimicrobial activity. According to this study, antimicrobial efficacy increased after Shodhana⁷. Present study has confirmed that Tuttha Bhasma is even more potent as antibacterial and antifungal agents when compared with Shodhita Tuttha. More potent antimicrobial activity was observed even at a lesser dosage than that of Shodhita tuttha.

Conclusion:

Comparatively Tuttha Bhasma are found to be more effective in fungus rather than bacteria

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