



SHORT COMMUNICATION

BIOTECHNOLOG

**COMPARATIVE STUDY OF AGITATION RATE AND STATIONARY PHASE FOR  
THE REMOVAL OF  $\text{Cu}^{2+}$  BY A.LENTULUS**

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**ABSTRACT**

The aim of this research work was to find out the effect of agitation rate for the removal of  $\text{Cu}^{2+}$ . Experiments were performed under shaking and stationary conditions towards optimizing the rate for efficient  $\text{Cu}^{2+}$  removal by the immobilized fungal strain. Increase in agitation rate from 100-180 rpm resulted into higher copper removal.



## KEY WORDS

A.lentulus, Agitation rate,  $\text{Cu}^{2+}$  removal

## INTERDICTION

Heavy metal pollution is one of the most important environmental problems today. One of the processes used for metal removal utilizes natural material<sup>1,2</sup>. In this study, adsorption ability of immobilized *A.lentulus* was investigated under stationary phase and agitation rate for the removal of metal. The effect of metal toxicity has been largely reflected through decrease in specific growth rates, extended lag phase, and decrease in radial colony expansion and respirations rates in presence of metallic pollutants. Less attention has been paid to concomitant changes in colony and hyphal morphology. In the present study, an interesting observation was made about the change in morphology of the pellets induced by metal stress. The pellets in absence of metal were smaller as compared to those in presence of metal above certain concentration (55mg/l). Size of pellet was found to be 0.4 mm in absence of  $\text{Cu}^{2+}$ . A gradual increase in pellet size was observed when  $\text{Cu}^{2+}$  concentrations were increased from 50mg/l to 100mg/l. Above 100mg/g, size of fungal pellets remains almost same. This indicates that change in pellet size may be due to the fact that the pellets get aggregated in response to metal toxicity at higher concentrations. *A. niger* produces larger pellets in response to nickel toxicity as compared to smaller pellets in control (absence of nickel)<sup>3</sup>.

### General Procedure:

1 Biomass preparation Fungal strain was grown in 250 ml Erlenmeyer flasks containing 100 ml of growth media. After exponential phase of growth, cells were harvested by centrifugation at 4000 rpm and 6 C for 10 min. The harvested cells were washed by double distilled water. This biomass was termed as resting biomass.

*A. lentulus* was immobilized by mixing sodium alginate, gelatin with above resting biomass in the ratio of 1:5. This mixture was dropped in the form of beads of sodium alginate, gelatin covering the biomass. These beads were dipped into calcium chloride solution overnight. Finally, beads of calcium alginate, gelatin entrapping the fungal biomass were obtained.

2 Copper removal studies using synthetic solution.

A weigh amount of treated/ untreated fungal biomass (4.5 g/l) was added in an Erlenmeyer flask containing 100 ml solution of known concentration of  $\text{Cu}^{2+}$ .

Initially, fungal growth and  $\text{Cu}^{2+}$  removal studies were conducted under shaking (100rpm) and stationary conditions (Fig 1). As compared to pellet morphology in shaking conditions, the biomass growth in stationary mode was in the form of a thick mat. The growth in stationary conditions was delayed as compared to shaking conditions, even in the absence of any copper stress, which can be attributed to reduced mass transfer and oxygen availability. The presence of  $\text{Cu}^{2+}$  further reduced the growth rates in stationary condition. It shows that the toxicity of  $\text{Cu}^{2+}$  was also reduced under stationary mode due to limited contact with the contaminant. For further verification of these results, one of the batches was run in shaking mode till 48 h and then shifted to stationary mode (Fig 2).

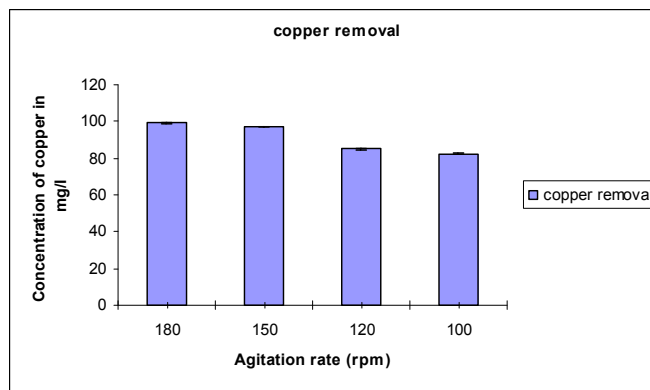
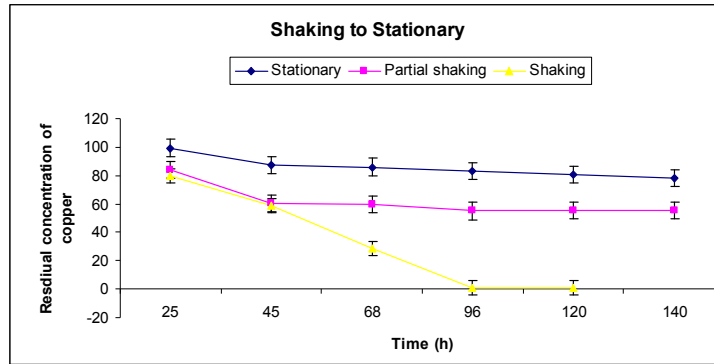
It was observed that in the stationary mode leading to the development of thick mat only 40%  $\text{Cu}^{2+}$  removal was observed. Hence, shaking conditions are necessary to ensure  $\text{Cu}^{2+}$  removal during growth of *A. lentulus*. Further experiments were directed towards optimizing the agitation rate for efficient  $\text{Cu}^{2+}$  removal. Increase in agitation speed (100-180 rpm) resulted into slightly higher biomass production and  $\text{Cu}^{2+}$  removal changed from 85% to 99.7%. (Fig. 2 and Fig 3)<sup>4</sup>. The oxygen transfer often influences the fungal pellet



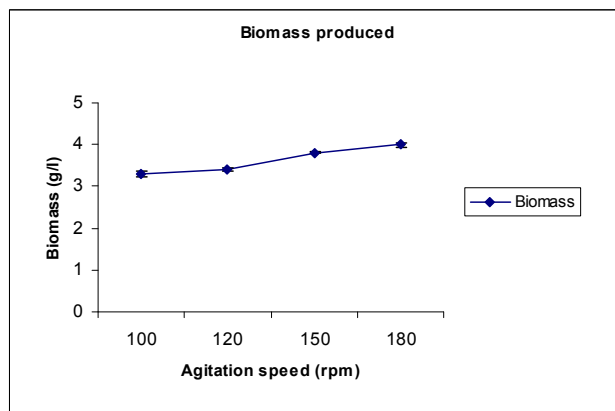
growth, and molecular diffusion is assumed to be the determinant mechanism for oxygen transport in pellet if they are larger than a

certain size, because of lack of oxygen in the pellet centre<sup>5</sup>.

**Fig 1**  
**Removal of Cu<sup>2+</sup> by A.lentulus in shaking and stationary conditions at 100mg/l initial Cu<sup>2</sup> concentration .**



**Fig 2**  
**Removal of Cu<sup>2+</sup> by A.lentulus in shaking and stationary conditions at 100mg/l initial Cu<sup>2</sup> concentration**



**Fig 3**  
**Biomass production in shaking and stationary conditions at 100mg/l initial Cu<sup>2</sup> concentration.**



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