THERMAL AND MECHANICAL SHOCKS AFFECTING THE FIRST FLUSH OF PRODUCTION OF 
LENTINULA EDODES ON EUCALYPTUS SALIGNA LOGS*

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ABSTRACT

The aim of this work was to evaluate the effect of thermal and mechanical shocks on the productivity of Lentinula edodes colonized on 140 Eucalyptus saligna logs at different immersion times in water and in the first flush of production. The logs were immersed in water at low temperature (16ºC) or environmental temperature (22ºC), for 6, 10, 14, 18, 22, 26 and 38 hours. The mechanical shock treatment was accomplished by dropping the logs onto the floor three consecutive times, in a vertical position. Water temperature and immersion time affected L. edodes yield, which increased two to four times when the logs were immersed in cool water for the shortest times (6 and 10 hours). The mechanical shock treatment did not increase the sporophore yield.

Key words: shiitake, stress factors, Lentinula edodes, sporophore yield

Shiitake, or Lentinula edodes [(Berk.) Pegler], is one of the most important edible mushrooms in Asian countries, and the second-most cultivated in the world (3).

Traditionally, Shiitake has been produced on recently felled logs measuring 1.0 m in length by 10 to 15 cm in diameter. Among the plant species more frequently used in cultivation, eucalyptus logs are particularly important and are predominantly used in Brazil, since they are inexpensive and can be easily purchased (6).

The production of sporophores starts when the logs are completely colonized by the fungus. Under natural conditions, production occurs right after the opening rains, causing complete log wetting and a sudden temperature decrease in the environment (5, 16). Under business growing conditions, these changes are provoked and managed, and include a reduction in temperature and increases in moisture, aeration, and luminosity, causing the fungus metabolism to change and forcing it to reallocate its energy sources to the production of mushrooms (15). The management intended to induce the production of sporophores is usually made by immersing the logs in water, which is frequently cooled to create an additional stress factor for the mycelium (5). Generally, most Shiitake strains respond better to induction when submerged in water at a temperature between 5ºC and 20ºC (5, 10, 16), with a water immersion time between 8 and 72 hours (12), with longer immersion times required for logs with greater diameters or densities (16).

The mechanical shock consists in exposition of the logs to strong impacts at one end, after they are removed from the immersion tank. Its use dates from the eleventh century in China, where it was observed that, under natural conditions, when the logs were vigorously stricken, after a few days the Shiitake mushrooms would appear abundantly on them (2). This practice was widely used years ago, however, without experimental data (14). No detailed studies were found on the effects of water 


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temperature and log immersion time, or yet on the effect of mechanical shock on the induction of *L. edodes* primordia formation on eucalyptus logs.

The experiment was conducted under greenhouse conditions, in Jaboticabal, São Paulo, Brazil. The treatments were arranged in a completely randomized design, with five replicates, in a 7×2×2 factorial arrangement (seven times of immersion in water; two immersion water temperatures; with or without mechanical shock). The production of inoculum was based on the JAB-K1 *L. edodes* strain, cultivated in PDA medium for 10 days at 25°C. The inoculum was grown on a substrate previously sterilized consisting of *Eucalyptus* sp. sawdust, rice bran, and water (4:1:1 v/v), placed inside 1-liter capacity polypropylene bags and incubated in an oven at 25°C for 30 days.

One hundred and forty *Eucalyptus saligna* logs 10 to 15 cm in diameter and 1 m in length, were inoculated with the substrate colonized by the fungus two days after the trees were cut. The logs were then inoculated with *L. edodes*, using a mechanical inoculator, after drilling the logs with a high-rotation (7000 rpm) power drill; a spacing of 15 cm between holes and 5 cm between hole rows was adopted, totaling 50 ± 10 holes per log. The holes were drilled at a 2 cm depth with a diameter of 1.2 cm, according to the methodology described by Przybylowicz and Donoghue (5). After inoculation, the logs were disposed at random, stacked at an angle, with a height of 120 ± 10 cm, separated at 3.0 cm to 5.0 cm from one another, in a shed protected from direct sunlight, at a temperature near 25 ± 2°C and relative humidity from 60% to 75% (15,23), for a period of 170 days until the formation of sporophores was induced.

The 140 logs were arranged horizontally inside two empty masonry tanks and tied to the bottom of the tank with a metal chain. In one tank, seventy logs were immersed in water at a temperature of 16°C, achieved by adding block ice. This temperature was 10°C below room temperature (26°C) (12,13), and was maintained cool throughout the treatments by adding ice blocks whenever needed. The other 70 logs were immersed in water at a temperature of 22°C, using the same procedures, but without cooling the water. In both tanks, the water immersion times were 6, 10, 14, 18, 22, 26, and 38 hours.

After each immersion period, the logs were removed from the tanks and submitted or not to mechanical shock. The mechanical shock consisted of strong impacts at one end of the log, obtained by dropping the log three consecutive times, in the vertical position, against a steady shield on the floor, from a height of 30 cm. After the treatments were carried out, the logs were transferred to an environment favorable for the formation of sporophores (90 ± 5% RH, 20 ± 2°C, and 500 ± 200 lux) (5,16). The harvest period was from 5 to 8 days after induction. Harvest was accomplished by hand and the standard adopted for the sporophores was the same required by the consumer market, i.e., sporophores with the pileus completely expanded and with the edges still facing down (13). After harvest, the sporophores fresh mass per log, number of sporophores per log, and fresh mass per sporophore were determined. The data were submitted to analysis of variance and the F test at 5% probability, and regression equations were fitted.

The coefficient of variation oscillated between 66.36% (fresh mass per sporophore) and 102.60% (sporophore fresh mass per log), revealing high variability between replicates of the same treatment (Table 1). The irregular sporophores yield distribution and the inclusion of data from unproductive or little productive logs in the statistical analysis explain these results. According to Royse (7), this irregular yield distribution is a common fact in the cultivation of *L. edodes* on logs. Montini and Eira (4) also adopted this procedure, because the exclusion of unproductive log data in order to decrease the experimental error would result in an incorrect representation of reality.

With respect to the production of sporophores fresh mass per log, number of sporophores per log, and fresh mass per sporophore, there were no interactions between the factors immersion time (IT), water temperature (WT), and mechanical shock (MS), demonstrating that the behavior of one factor is not influenced by changes in another factor. The immersion water temperature (WT) effect was significant in relation to the production of *L. edodes* sporophores fresh mass per log, number of sporophores per log, and fresh mass per sporophore (Table 1). In cooled water (16°C), the sporophores fresh mass per log, number of sporophores per log, and fresh mass per sporophore were, respectively, about four, four, and two times higher than those obtained in water without cooling (22°C), demonstrating that temperature reduction favors the development of *L. edodes*, and therefore is an adequate technique to increase yield. Similar results were obtained by Song *et al.* (10), who observed *L. edodes* dry weight and sporophores production velocity increases on solid and semi-solid artificial substrates, with water cooled to 5°C in relation to the control (25°C) for 24 hours. These authors verified an alteration in the metabolism of the fungus, caused by a change in the profile of mycelium lipids, mainly due to an increase in the quantity of unsaturated fatty acids, such as linoleic acid, which would be the correct substrate for primordia initiation (8,11); this change seems to have favored the formation of sporophores. Studies have demonstrated the capacity of the low temperature stress factor in changing the composition of fatty acids in other microorganisms (17), causing an increase in the amount of unsaturated fatty acids, such as palmitoleic, linoleic, and oleic acids, which are presumed to act maintaining the fluidity of the lipid on the *L. edodes* membrane (8) and favoring the production of sporophores. The mechanical shock effect was non-significant, indicating that it does not change sporophores fresh mass per log, number of sporophores per log, and fresh mass per sporophore. Therefore, it did not prove to be an effective technique to increase sporophores productivity (Table 1).
The *L. edodes* sporophores fresh mass per log, number of sporophores per log, and fresh mass per sporophore on eucalyptus logs showed a decreasing exponential behavior as immersion time in water increased. These results demonstrate that a yield reduction occurred with longer times of immersion in water. However, it must be taken into consideration that primordia formation is the most sensitive stage to environmental influences, where extreme log moisture, temperature, and relative humidity values may change or inhibit the production of sporophores (5,16). Thus, the excess moisture contained in the logs, resulting from prolonged exposures to water, could be the reason for the systematic drop in sporophores yield, as immersion times became longer.

Under the conditions the experiment was developed, on the first flush of production, the cooling of the immersion water significantly increases the production of *Lentinula edodes* on *Eucalyptus saligna* logs (2 to 4 times); mechanical shock does not influence the production of *Lentinula edodes* on *Eucalyptus saligna* logs and the prolonged exposure of *Eucalyptus saligna* logs in water decreases the production of *Lentinula edodes* sporophores.

### Table 1. Values for the F Test statistic of causes of variation in the analysis of variance with regard to sporophores fresh mass per log, number of sporophores per log, and fresh mass per sporophore, in *L. edodes* mycelia submitted to seven immersion times, two water temperatures, and two mechanical shocks (five logs per treatment).

<table>
<thead>
<tr>
<th>Causes of variation</th>
<th>Sporophores fresh mass per log (g log⁻¹)</th>
<th>Number of sporophores per log (units log⁻¹)</th>
<th>Fresh mass per sporophore (g sporophore⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Immersion Time (IT)</td>
<td>1.16 NS</td>
<td>0.71 NS</td>
<td>2.56*</td>
</tr>
<tr>
<td>Water temperature (WT)</td>
<td>20.80**</td>
<td>19.58**</td>
<td>9.72**</td>
</tr>
<tr>
<td>Mechanical Shock (MS)</td>
<td>0.06 NS</td>
<td>0.03 NS</td>
<td>0.02 NS</td>
</tr>
<tr>
<td>IT × WT interaction</td>
<td>0.65 NS</td>
<td>0.73 NS</td>
<td>0.53 NS</td>
</tr>
<tr>
<td>IT × MS interaction</td>
<td>0.22 NS</td>
<td>0.21 NS</td>
<td>0.69 NS</td>
</tr>
<tr>
<td>WT × MS interaction</td>
<td>0.35 NS</td>
<td>0.25 NS</td>
<td>0.67 NS</td>
</tr>
<tr>
<td>IT × WT × MS interaction</td>
<td>0.48 NS</td>
<td>0.52 NS</td>
<td>0.54 NS</td>
</tr>
<tr>
<td>Mean</td>
<td>55.71</td>
<td>5.14</td>
<td>6.63</td>
</tr>
<tr>
<td>Coefficient of variation</td>
<td>102.60%</td>
<td>82.05%</td>
<td>66.36%</td>
</tr>
</tbody>
</table>

* Significant at the 5% probability level; ** Significant at the 1% probability level; NS Non significant.

The *L. edodes* sporophores fresh mass per log, number of sporophores per log, and fresh mass per sporophore on eucalyptus logs showed a decreasing exponential behavior as immersion time in water increased. These results demonstrate that a yield reduction occurred with longer times of immersion in water. However, it must be taken into consideration that primordia formation is the most sensitive stage to environmental influences, where extreme log moisture, temperature, and relative humidity values may change or inhibit the production of sporophores (5,16). Thus, the excess moisture contained in the logs, resulting from prolonged exposures to water, could be the reason for the systematic drop in sporophores yield, as immersion times became longer.

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**RESUMO**

**Coque térmico e mecânico afetando o primeiro fluxo de produção de *Lentinula edodes* em toras de *Eucalyptus saligna***

O objetivo deste trabalho foi avaliar o efeito do choque térmico e mecânico na produtividade de *Lentinula edodes* em 140 toras de *Eucalyptus saligna*, completamente colonizadas pelo fungo, em diferentes tempos de imersão em água e no primeiro fluxo de produção. As toras foram imersas em água resfriada (16°C) ou à temperatura ambiente (22°C); os períodos de imersão corresponderam a 6, 10, 14, 18, 22, 26 e 38 horas; o choque mecânico foi acompanhado por três quedas consecutivas da tora, em posição vertical, no chão. A temperatura da água e o tempo de imersão afetaram a produção de *L. edodes*, resultando em aumentos significativos (2 a 4 vezes) nos tratamentos em que as toras foram submetidas à água resfriada e nos tempos de imersão mais curtos (6 e 10 horas). O choque mecânico não resultou em aumento na produção de basidiomas.

**Palavras-chave:** Shiitake, fatores de estresse, *Lentinula edodes*, produção de basidiomas

**REFERENCES**


