First cultivation of *Agaricus flocculosipes* and a novel Thai strain of *A. subrufescens*

Thongklang N¹,², Sysouphanthong P³, Callac P⁴ and Hyde KD¹,²

¹School of Science, Mae Fah Luang University, Chiang Rai 57100, Thailand  
²Institute of Excellence in Fungal Research, and School of Science, Mae Fah Luang University, Chiang Rai 57100, Thailand  
³Key Laboratory for Plant Diversity and Biogeography of East Asia, Kunming Institute of Botany, Chinese Academy of Science, Kunming 650201, Yunnan, China  
⁴UR 1264, Mycologie et Sécurité des Aliments, 33883 Villenave d’ Ornon, France


Abstract

*Agaricus flocculosipes* and *A. subrufescens* are edible species that belong to section *Arveses* of the genus *Agaricus*. *Agaricus subrufescens* (almond mushroom) is known to produce bioactive compounds with medicinal properties, such as anti-cancer and anti-tumor activity and fruiting bodies are also edible and nutritious. *Agaricus subrufescens* is presently cultivated in Brazil, China, Japan, Taiwan and some European countries for use as foods and nutraceuticals. *Agaricus flocculosipes* is a newly described species currently known only from Thailand, Mayotte Island and China. Species of *Agaricus* have high potential for cultivation as many species are edible and have medicinal properties. Herein we report the first cultivation of *A. flocculosipes* and a Thai strain of *A. subrufescens*. The strains were isolated from wild sporocarps and subcultured on compost extract agar medium. Standard compost based on wheat straw and horse manure was used as the cultivation substrate. A comparative study of the cultivation of one strain of *A. flocculosipes* and two strains of *A. subrufescens* from France and Thailand was carried out with four replicates trays. The mushroom spawn was inoculated into the compost and incubated at 25 °C. The mycelia fully covered and grew throughout the media after incubation for two weeks. A casing layer made up of sand, peat and lime (1:1:1) was applied. Fruiting phase was performed at 25 °C and 95% humidity. The first primordia of the French and Thai strains of *A. subrufescens* occurred after the casing layer was added at days 12 and 24, respectively. Yields were obtained from both strains of *A. subrufescens*. The yield of the Thai strain (85.90 ± 27.06 g kg⁻¹) was lower than that of the French strain (178.56 ± 68.28 g kg⁻¹) in the first production cycle lasting 34 days. The first primordia of *A. flocculosipes* formed at day 32 after casing, and the strain produced only a few fruit bodies (1.04 ± 0.95 g kg⁻¹). Further research is needed to develop suitable agriculture wastes and regimes for growing these mushrooms and for increasing production yields so that these strains and species can be industrialized.

**Keywords** – *Agaricus* – edible mushroom – fruiting test – mushroom growing
**Introduction**

Most of edible mushrooms are macrofungi belonging to the phylum basidiomycota. Since ancient times mushrooms have been consumed by humans, not only as part of the normal diet but also as a delicacy because they have a high source of protein, desirable taste, aroma and also have medicinal properties (Zhong & Tang 2004, Firenzuoli et al. 2008, De Silva et al. 2012a, 2012b, 2013).

_Agaricus_ is probably the most important and potentially cultivatable genus, as most species are edible, some are delicious and widely consumed and others have considerable medicinal properties. _Agaricus bisporus_ (the button mushroom) was first domesticated in France in 1650 (Atkins 1978, Patel 2013). _Agaricus bisporus_ is very popular worldwide and is the most widely cultivated species of edible mushroom, accounting for 31.8% of the world market. Species of _Lentinus_ are second ranking and species of _Pleurotus_ are third ranking in the world market (Chang 1999).

_Agaricus subrufescens_ is also a cultivated mushroom (Firenzuoli et al. 2008) with significant medicinal properties and belonging to the section _Arvenses_ of the genus _Agaricus_, while _Agaricus bisporus_ belongs to _Agaricus_ section _Bivelares_ (Kerrigan et al. 2008, Parra 2013). This mushroom is cultivated commercially in Brazil, China, Japan and Taiwan (Kerrigan 2005). Main synonyms of _A. subrufescens_ are _A. blazei_ sensu Heinemann (misapplied), _A. braziliensis_ (illegitimate), and _A. rufotegulis_. This species has a broad climatic and geographical distribution range (Asia, Europe and Oceania). Recently Thongklang et al. (2014) showed that samples from Brazil, France and Thailand are amphithallic and interfertile.

Wild edible mushrooms in the section _Arvenses_ include _Agaricus arvensis_, _A. augustus_ and _A. silvicola_, all of which are collected from the wild for human consumption in Europe and Americas (Kalač & Svoboda 2000, Wisitrassameewong et al. 2012). Zhao et al. (2012) described a new species in the section _Arvenses_ of the genus _Agaricus_ from Thailand as _A. flocculosipes_; this new mushroom is potentially cultivatable and is expected to be a good edible species that may have potential commercial value for Thailand and other countries.

Approximately 30,000 mushrooms throughout the world have been described (Kirk et al. 2008), and in Thailand 22 species are cultivated commercially for the Thai market (Thawthong et al. 2014). Our studies of basidiomycetes from northern of Thailand have reported several new species for Thailand. A new wild edible mushroom has been cultivated successfully; _Pleurotus giganteus_ can form fruiting bodies at 25°C and has the possibility of being grown in Thailand (Klomklung et al. 2012).

_Agaricus flocculosipes_ and _Agaricus subrufescens_ are newly recorded or described species in Thailand. _A. flocculosipes_ is also recorded from Mayotte Island (Zhao et al. 2012) and China (Gui et al. 2014, in press). In Thailand, we questioned local people and found that these wild species are not commonly consumed and not many people are aware that these mushrooms are likely edible. These mushrooms could be introduced to Thai market for cultivation and medicinal use.

The objective of this study was to determine ways to cultivate the Thai mushroom strains of section _Arvenses_ ( _A. flocculosipes_ and _A. subrufescens_) on a laboratory scale using compost. Research was directed towards the cultivation of these wild strains for fruiting competence, and we discuss the possible introduction to the local market.

**Materials & Methods**

**Mushroom strains**

We collected numerous species of _Agaricus_ between 2004 and 2013; some are potentially cultivatable and have medicinal properties. We isolated several Thai strains of _A. flocculosipes_ and _A. subrufescens_ from northern Thailand for potential cultivation. These species belong to section _Arvenses_, whose species are generally edible. _Agaricus subrufescens_ (CA918= MFLUCC 11-0653) was found growing in a grassland at Mae Fah Luang University, Thailand by S.C. Karunarathna, P.
Callac and S. Rapior in 2011. *Agaricus flocculosipes* (CA917, MFLUCC 11-0652) was also found along with CA918 on the campus of Mae Fah Luang University, Thailand by S.C. Karunaratna, P. Callac and S. Rapior in 2011. Mycelia were isolated and subcultured on compost agar medium. The strains were incubated at 25°C for two or three weeks.

**Spawn production**

Spawn is the media for transfer of the mushroom mycelium to the growing substrate for upscale production of mycelia for cultivation. Rye grain spawn prepared by Euromycèl, France was used for spawn production (Navarro & Savoie 2013, Llarena-Hernández et al. 2014). The spawn boxes containing 100 g of rye grain were inoculated with a half colony from actively growing mycelia of a 9 mm diameter Petri dish.

**Compost and casing layer**

Commercial compost based on horse manure mixed with wheat straw and with various additives as the main substrate for *A. bisporus* cultivation was provided by SA Renault, Pons, France. The substrate, compost (8 kg) was inoculated with spawn (2% weight of compost) and incubated at 25 °C and 85 % humidity for 15 days. After the mycelia fully covered and grew throughout the compost, a casing layer was applied. The casing layer, made up of a mixture of limestone: peat: thin sand (1:1:1), was added (about 4 cm deep) above the colonized mycelium in the compost, then incubated at 25 °C and 95% humidity and low CO₂ concentration (Llarena-Hernández et al. 2011, 2014). After casing, the trays were watered once a day. The fruiting test was carried out with four replicates trays (Royse 2010, Llarena-Hernández et al. 2011).

**Yield data and Statistical analysis**

The fruiting bodies, including those with open and closed caps, were manually harvested, and counted and weighed daily. The mushroom yields were recorded for 34 days. Yield data means total weight of fresh mushroom in one crop per kilogram of substrate (Royse 2010, Llarena-Hernández et al. 2011). The data set was analyzed statistically for variance and mean by one-way ANOVA analysis using Duncan’s multiple range tests. Differences were considered significant for P < 0.05.

**Results**

*Agaricus subrufescens*

A comparative study of the cultivation of the Thai and French strains of *A. subrufescens* was carried out with four replicates trays. Fruiting bodies of the Thai strain were produced at 25 °C and 95% humidity. The first primordia of the French *A. subrufescens* strain appeared on day 12 (Fig 1 e-f), while that of the Thai *A. subrufescens* strain appeared on day 24 (Fig 1 a-b), after casing was applied. Although yields were obtained for both strains, the yield of the Thai strain was lower than that of the French control strain, with the average wet yield of French and Thai strains 1,428.50 g and 687.17 g, respectively (Table 1). We found that the *A. subrufescens* French strain produced 323 fruiting bodies in production cycle (34 days), while Thai strain produced 158 fruiting bodies in the production cycle (34 days). Comparison of Thai *A. subrufescens* and French strain production are given in Table 1.

*Agaricus flocculosipes*

Cultivation of the Thai strain of *Agaricus flocculosipes* was achieved, but with low yield and in some trays, no yield. The mycelia of *A. flocculosipes* fully covered and grew throughout the culture substrate after 2 weeks incubation. The first primordia of *A. flocculosipes* formed on day 32 after casing (Fig 1 c-d). The average wet yield of *A. flocculosipes* was 8.33 g, with two fruiting bodies produced in 34 days (Table1).
Fig. 1 – Cultivation of species of Agaricus. a-b: A. subrufescens Thai strain (CA918) produced at 25 °C and 95% humidity, c-d: A. flocculosipes (CA917) developing on compost media at 25 °C and 95% humidity, e-f: French strain of A. subrufescens (CA487).
**Confirmation of cultivated species**

The Thai strain of *A. subrufescens* that fruited at 25 °C and 95% humidity was checked to confirm the species. The macromorphological characters traits of this species are described in detail; the pileus surface covered with silk-like fibres and small scales, a reddish-brown cap, the odour of almond, a two-layered and floccose annulus, yellow staining when cut and a positive Schäffer’s reaction all are typical of *A. subrufescens*.

The fruiting bodies of *A. flocculosipes* that grew on compost were confirmed as being the same species as the inoculated strain. The floccose stipe, almond odour, well-developed squamules on the pileus, two-layered annulus, and catenulate cheilocystidia were typical of the species.

**Table 1** Comparison first cycle yields (34 days) Thai and French strains of *Agaricus subrufescens*.

<table>
<thead>
<tr>
<th>Contents</th>
<th><em>Agaricus flocculosipes</em> Thai strain (CA917)</th>
<th><em>Agaricus subrufescens</em> Thai strain (CA918)</th>
<th><em>Agaricus subrufescens</em> French strain (CA487)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Occurrence of primordia after soil casing</td>
<td>30-32 days</td>
<td>21-24 days</td>
<td>7-14 days</td>
</tr>
<tr>
<td>Average wet weight</td>
<td>8.33 g</td>
<td>687.17 g</td>
<td>1428.50 g</td>
</tr>
<tr>
<td>Yield data*</td>
<td>1.04 g kg⁻¹</td>
<td>85.90 g kg⁻¹</td>
<td>178.56 g kg⁻¹</td>
</tr>
<tr>
<td>Number of fruiting bodies</td>
<td>2</td>
<td>158</td>
<td>323</td>
</tr>
</tbody>
</table>

*Yield data = total weight of mushroom per kilogram of substrate.*

**Discussion**

Although there are well over 300 genera of mushrooms, only a few species are cultivated commercially due to the fact that some are mycorrhizal and apparently will not fruit without the host roots being present (Egli 2011). However, there are many saprobic genera, and these have the possibility of being cultivated (Klomklung et al. 2012, Thawthong et al. 2014). A simple fruiting test using wild strains is essential for testing cultivars that might be introduced to the market. The general methodology for assessing fruiting ability have been described, firstly the strains were isolated and inoculated to grain spawn and finally compost inoculation and casing with soil (Carrera et al. 2001, Coello-Castillo et al. 2009, Llarena-Hernández et al. 2011). Preparation of substrates suitable for *Agaricus* fruiting involves using standard compost and success in fruiting tests to predict whether it is possible to grow the wild strains (Llarena-Hernández et al. 2011, Navarro & Savoie 2013, Thawthong et al. 2014).

*Agaricus* has high potential for cultivation as many of the species in this genus are edible and some have medicinal properties (Bernardshaw et al. 2005, Adams et al. 2008, De Silva et al. 2012a, 2012b, Wisitrassameewong et al. 2012). In this study we tested a new Thai strain of *A. subrufescens* and the new species, *A. flocculosipes* (section *Arvenses*) for fruiting competence. The two strains were isolated by tissue culture and as tropical strains, incubated at 25 °C (Llarena-Hernández et al. 2011, 2014, Navarro & Savoie 2013). Colony morphology on agar plates depended on the species of mushroom. The mycelia of all strains are white and have an almond odour during both spawn production and on compost media agar.

In fruiting trials of the Thai and French *A. subrufescens* strains, yields of the Thai strain were lower than the French strains; the productivity of the Thai strain was low (85.90 ± 27.06 g kg⁻¹), while the French strain was 178.56 ± 68.28 g kg⁻¹. *Agaricus subrufescens* cultivation trials by Llarena-Hernández et al. (2011) showed that wild strains [of *A. subrufescens*] produced a wide range of mushroom yields, depending on the mushroom species and strains. The Brazilian wild strain of *A. subrufescens* (WC837) that grew with wheat and horse manure as main substrate had a yield of 41.9 ± 12.2 g kg⁻¹, which is lower than our Thai strain corrected, while a French strain of *A. subrufescens* (CA487) had a yield of 207.9 ± 47.9 g kg⁻¹. The yield of the mushroom is dependent on the substrate used for cultivation. Eira (2003) used various local substrates for *Agaricus subrufescens* (as *A. blazei*) cultivation in Brazil; the yields of this mushroom ranged from 3 to 25 kg of fresh mushroom /100 kg of substrate (Eira 2003), depending on the substrate used.
Agaricus flocculosipes is a new potential cultivable species in the section Arvenses described by Zhao et al. (2012). This species produced fruiting bodies but yields were low, although the mycelium has the ability to colonize well throughout the compost media, and the fruiting period was shorter when compared with the French and Thai strains of A. subrufescens. Further work will be carried out to develop suitable media for culturing these taxa, and genetic improvement will be attempted by breeding for increasing yield (Thongklang et al. 2014).

Normally, the time to fruiting and thus mushroom harvesting begins 15-20 days after casing. In this study, we found that French strain of A. subrufescens began to fruit 12 days after the casing layer was applied, while fruiting of the Thai wild A. subrufescens strain occurred 24 days after casing. Agaricus flocculosipes began to produce fruiting bodies at 32 days. However, the time of first fruiting is dependent on the material and method of casing. Llarena-Hernández et al. (2014) reported that increasing the depth of the casing layer applied to compost significantly improved yield and fruiting time in A. subrufescens. However further work will be carried out to develop suitable conditions for improving time to fruiting.

Acknowledgements

This work was supported by the Royal Golden Jubilee Ph.D. Program-RGJ–I (PHD / 0061 / 2552, 4. S. MF/52/A.1), and the Thailand Research Fund grant (BRG 5580009) under the research grant entitled "Taxonomy, Phylogeny and Biochemistry of Thai Basidiomycetes" for financial support. We thank to Institut National de la Recherche Agronomique (INRA), France and School of Science, Mae Fah Luang University is acknowledged for providing facilities and training.

References


