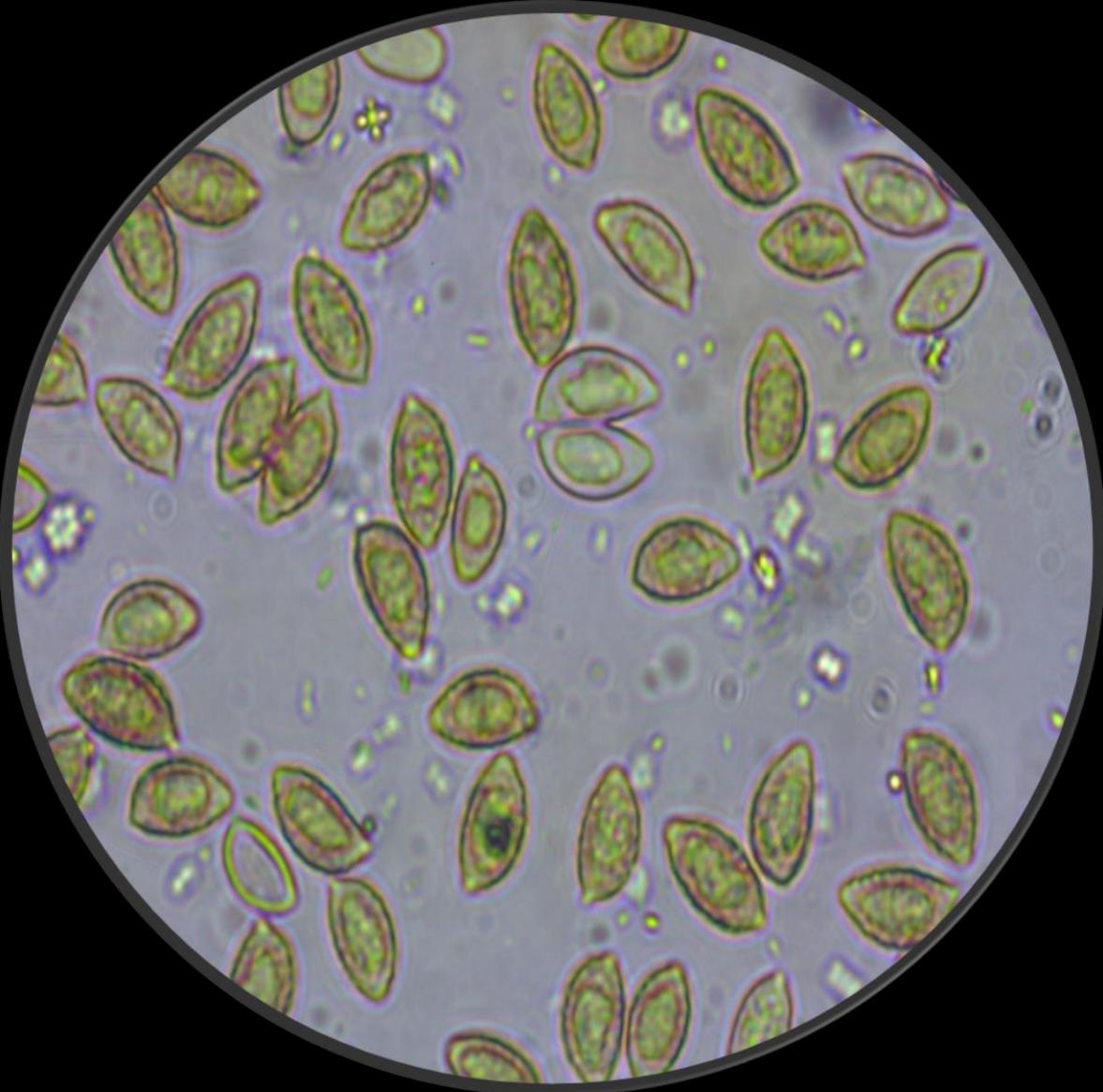


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Mantarcılık
Uygulama ve Araştırma Merkezi
KONYA-TÜRKİYE***



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ADINA SAHİBİ

PROF.DR. GIYASETTİN KAŞIK

YAZI İŞLERİ MÜDÜRÜ

PROF.DR. GIYASETTİN KAŞIK

Haberleşme/Correspondence

S.Ü.

Mantarcılık Uygulama ve Araştırma Merkezi Müdürlüğü
Alaaddin Keykubat Yerleşkesi, Fen Fakültesi B Blok,
Zemin Kat-42079/Selçuklu-KONYA

Tel:(+90)0 332 2233998/ Fax: (+90)0 332 241 24 99

Web: <https://dergipark.org.tr/tr/pub/mantar>

E-Posta:mantarcilik@gmail.com

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Bu sayıımızda yer alan eserler hakkında aşağıda isimleri yazılı hakemlerimize yaptıkları değerlendirmeler için teşekkür ederiz.

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Ağrı Merkez/Hamur İlçesinde Belirlenen Makromantarlar

Zeki ÇELİK¹, Mehmet Hakkı ALMA²

*Sorumlu yazar: zeki.celik@tarimorman.gov.tr

¹Ağrı İl Tarım ve Orman Müdürlüğü, Ağrı, Türkiye / zeki.celik@tarimorman.gov.tr

²Biyosistem Mühendisliği Bölümü, İğdır Üniversitesi, İğdır, Türkiye / mhakki.alma@igdir.edu.tr

Öz: Bu çalışma, 2018-2019 yılları arasında Ağrı ili Merkez ve Hamur ilçesi sınırları içerisinde yer alan bölgede doğal olarak yetişen ve toplanan makromantar örnekleri üzerinde yapılmıştır. Arazi ve laboratuvar çalışmaları sonucu teşhisleri yapılan mantar örneklerinin *Pezizomycetes* ve *Agaricomycetes* sınıflarına ait 5 takım ve 20 familya içerisinde dağılım gösteren toplam 57 makromantar taksonu olduğu tespit edilmiştir. Bunların 31'i yenen, 20'si yenmeyen ve 6 tanesi ise zehirli özellikleştir. Bu çalışma ile ülkemiz mikobiotasının belirlenmesine katkı sağlanması yanısıra mantarların yörede etnomikolojik amaçlı olarak yararlanma durumlarının belirlenmesi amaçlanmıştır.

Anahtar kelimeler: Basidiomycota, Ascomycota, Ağrı Merkez/Hamur, Makromantar

Macrofungi Determined in Ağrı Central/Hamur District

Abstract: This study was carried out on macrofungus specimens naturally grown and collected in the region located within the borders of central and Hamur district of Ağrı province between 2018-2019. It has been determined that the mushroom samples identified as a result of field and laboratory studies have a total of 57 macrofungi taxa, which are distributed in 5 orders and 20 families belonging to the *Pezizomycetes* and *Agaricomycetes* classes. Of these, 31 are edible, 20 are inedible, and 6 are poisonous. In this study, it was aimed to contribute to the determination of our country's mycobiota, as well as to determine the ethnomicological use of fungi in the region.

Key words: Basidiomycota, Ascomycota, Agri Central/Hamur, Macrofungus

Giriş

Genel anlamda mantarlar, ökaryotik, eşeyle ve eşeyiz çoğalabilen, spor üreten, hif diye bilinen tipik olarak hücre duvarıyla kuşatılmış dallanan ve ipliksel somatik yapıya sahip, saprofitik, parazitik veya simbiyotik bir yaşam sürdürken, hücre çeperi kitinden oluşan, absorbsiyonla beslenen, klorofilsiz organizmalar olarak tanımlanırlar. Makromantarlar Fungi âleminin, Ascomycota ve Basidiomycota bölgümlerine ait, gözle görüldükelle tutulan, saplı ve sapsız, şapkalı veya top şeklinde başta olmak üzere farklı ve birçok çeşitli şekillerde üreme yapıları olan canlılardır. Mantarlar, ekolojik olarak büyük öneme sahip canlılardır. Ayrıca, çürükçül olanlar besinlerini bitkisel atıklar, cansız ağaç parçaları ve hayvan gübresi gibi ölü organik maddelerden temin ederek doğadaki besin maddelerinin ve azot, fosfor, potasyum, demir, gibi organik yapıdaki elementlerin geri dönüşümünde önemli katkıları bulunmaktadır (Chang and Miles, 2004; Vargas and Zardoya, 2014; Chang and Wasser, 2018).

Ülkemiz sınırları içerisinde yetişen makromantarlarla ilgili yapılmış olan çalışmalar Sesli ve ark.,(2020) ve Solak ve ark., (2015) tarafından liste halinde sunulmuştur. Genel anlamda bu çalışmalarda belirtildiği üzere ülkemizde konu ile ilgili yapılan ilk çalışma 1915'te başlamış olup 2015 yılına kadar toplam olarak 2422 makromantar taksonu belirlenmiştir. Ayrıca verilen listede yapılan çalışmalardan sonra (Sesli ve ark.,2020; Akçay ve ark., 2022) sonucunda verilmiş olan bu sayı artmaktadır. Bu çalışmada, Ağrı Merkez ve Hamur ilçesi sınırları içerisinde doğal olarak yetişen makromantarlar tespit edilerek, Doğu Anadolu Bölgesi ve ülkemiz mikobiotasının zenginleştirilmesine katkı sağlanması ve bunula birlikte belirlenen bu mantarların habitatları, mevsimsel dağılımları ve yörede etnomikolojik amaçlı olarak yararlanma durumlarının belirlenmesi amaçlanmıştır.

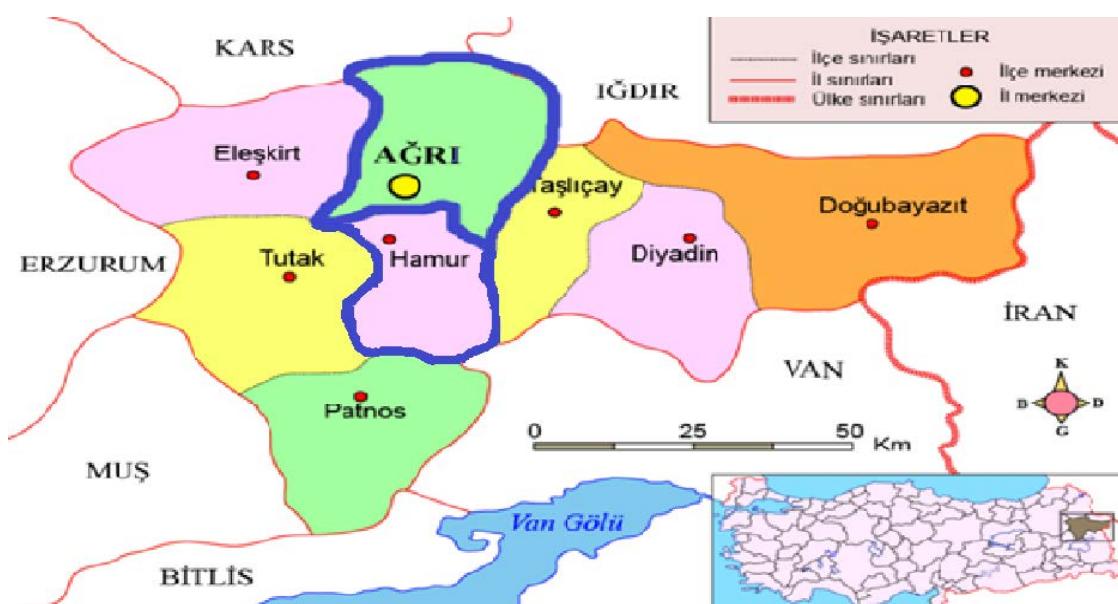
Materyal ve Metot

Mevcut çalışmanın materyalini oluşturan makromantar örnekleri 2018-2019 yılları arasında Ağrı ili



Merkez ve Hamur ilçesi sınırları içerisinde yer alan bölgede doğal olarak yetişen ve toplanan örnekler üzerinde yapılmıştır. Bununla birlikte mantar örneklerinin yetişmelerinin uygun olduğu bilinen İlkbahar ve sonbahar aylarında toplanmıştır. Çalışma alanında toplanan makromantar örneklerinin morfolojik ve etnomikolojik özellikleri not edilerek örneklerin teşhisinde veri olarak kullanılmıştır. Aynı zamanda araziden laboratuvara taşınan mantarın belirlenmiş olan mikolojik teknikler ve teşhis anahtarı uygulanması sonucunda fungaryum

materyalleri haline getirilmiştir. Yapılmış arazi ve laboratuvar çalışmaları sonrasında elde edilen veriler (Buczacki, 1989; Bresinsky and Besl, 1990; Phillips, 1981; Moser, 1983; Breitenbach and Kranzlin, 1986, 1991, 1995; 2000; Jordan, 1995; Ellis and Ellis, 1990; Kränzlin, 2005; Dähncke, 2004; Jordan, 2004) ilgili literatürle karşılaştırılarak mantar örneklerinin teşhisleri yapılmış olup kayıtları yapılan örnekler İğdır Üniversitesi Ziraat Fakültesinde saklanmaktadır.



Şekil 1. Araştırma alanının haritası (Kocaman ve ark., 2011).



Şekil 2. Araştırma alanından görünüm.

Bulgular



İlgili çalışma sonucunda, 2018-2019 yılları arasında Ağrı ili Merkez ve Hamur ilçesi sınırları içerisinde yer alan bölgede doğal olarak yetişen makromantarların tespit edilmesi amacı ile yapılmıştır. Belirlenen taksonlar; habitat, substrat, toplama yeri, ilçe, köy, coğrafi konum, yükseklik, toplama tarihi, şahsi fungaryum numarası ve yenilebilirlik durumları (örn.: Ağaç altı ve kalıntıları üzeri, merkez, dere kenarı, kütük üzeri, çayırlık alan, gübre üzeri, 05° 79'620"K, 43° 86'959"D, 1730 m, 05.04.2018, ÇELİK. 32), kendilerine ait fungaryum numaraları ile birlikte verilmiştir. Teşhis edilen örneklerin Pezizomyces ve Agaricomycetes sınıflarına ait olup 5 ordo ve 20 familya olarak dağılım gösteren 57 makromantar taksonu olduğu tespit edilmiştir. Ayrıca bunların 31'i yenen, 20'si yenmeyen ve 6 tanesi ise zehirli olan makromantar tespit edilmiştir. Buna ek olarak teşhis yapılan mantar örneklerinin türleri Sesli ve ark., (2020) ve [http://indexfungorum.org.](http://indexfungorum.org/), [http://www.mycobank.org.](http://www.mycobank.org) veri tabanları baz alınarak sistematik sıraya dizilmiştir.

Ascomycota

Pezizales

Helvellaceae Fr.

1- **Paxina queletii** Bres, **Papaztası;** Kavak (*Populus* sp.) ağaçları altında, Hamur-Ceylanlı, 05° 79'622"K, 43° 82'956"D, 1723 m, 05.04.2018, ÇELİK. 12. Yenir.

2- **Helvella leucopus** Pers., **Top semermantarı;** Kavak (*Populus* sp.) ağaçları altı, Hamur-Ceylanlı, 05° 79'612"K, 43° 82'950"D, 1720 m, 05.04.2018, ÇELİK. 18. Yenir.

Morchellaceae Rchb.

3- **Morchella elata** Fr., **Siyah göbek;** Kavak (*Populus* sp.) ağaçları altı, Hamur, 05° 89'613"K, 43° 74'876"D, 1773 m, 05.04.2018, ÇELİK. 14. Yenir.

4- **Morchella esculenta**(L.) Pers., **Kuzugöbeği;** Kavak (*Populus* sp.) ağaçlar altı, Merkez Yukarı yol düzü, 05° 93'433"K, 44° 10'421"D, 1733 m, 01.05.2018, ÇELİK. 13. Yenir.

Pezizaceae Dumort.

5- **Peziza vesiculosa** Bull., **Yığışanak;** Hayvan gübresi üzerinde, Merkez Yukarı yol düzü, 05° 97'533"K, 44° 10'471"D, 2180 m, 20.05.2018, ÇELİK. 15. Yenmez.

Basidiomycota

Agaricales

Agaricaceae Chevall.

6- **Agaricus arvensis** Schaeff., **Atmantarı;** Çayırlık alanda ve Kavak (*Populus* sp.) ağaçları altı, Merkez Yukarı yol düzü, 05° 93'567"K, 44° 10'578"D, 1880 m, 20.05.2018-12.06.2018, ÇELİK. 9. Yenir.

7- **Agaricus bisporus** (J.E. Lange) Imbach, **Kültür mantarı;** Çayırlık, Hamur ve Merkez Yukarı yol düzü, 05° 97'533"K, 44° 10'471"D, 2180 m, 20.05.2018-15.10.2018, ÇELİK. 34, 44. Yenir.

8- **Agaricus campestris** L., **İçi kıızı;** Çayırlık alan, Merkez Yukarı yol düzü, 05° 96'093"K, 43° 71'091"D, 2172 m, 20.05.2018, ÇELİK. 36. Yenir.

9-*Agaricus xanthodermus* Genev., Ağulu kıızı;

Çayırlık, Hamur-Ceylanlı ve Merkez Yukarı yol düzü, 05° 96'062"K, 43° 69'913"D, 1960 m, 20.05.2018, ÇELİK. 7. Zehirli.

10- **Bovista plumbea** Pers., **Puf mantarı;** Çayırlık alan, Hamur-Ceylanlı, 05° 77'910"K, 43° 86'298"D, 1970 m, 05.04.2018, ÇELİK. 6. Yenir.

11- **Bovistella utriformis** (Bull.) Demoulin & Rebrev, **Yan poslak;** Genellikle dağlık alanda, güneşli yüksek meralarda, çayırlıklarda, otlaklıarda kumlu topraklarda, kısa bodur ağaçların bulunduğu alanda; Hamur-Ceylanlı, 05° 77'910"K, 43° 86'298"D, 1995 m, 30.05.2018, ÇELİK. 50. Yenir.

12- **Lycoperdon perlatum** Pers. **Fissakuri;** Kumlu toprak ve Çayırlık alanda; Hamur-Ceylanlı, 05° 75'934"K, 43° 86'278"D, 1985 m, 30.05.2018, 05° 67'678"K, 43° 84'256"D, 1895 m, 12.10.2018, ÇELİK. 11, 17. Yenir.

13- **Coprinus comatus** (O.F. Müll.) Pers., **Söbelen;** Çayırlık alanda, Söğüt (*Salix* sp.), ağaçları altı, Hamur-Ceylanlı ve Merkez Yukarı yol düzü, 05° 95'869"K, 43° 70'105"D, 2160 m, 20.05.2018, ÇELİK. 10, 19. Yenir.

14- **Lepiota erminea** (Fr.) P. Kumm., **Süt pullu;** Kavak (*Populus* sp.) ağaçlar altı, çayırlık alanda, Hamur-Özdirek ve Merkez, 05° 95'869"K, 43° 70'105"D, 2160 m, 20.05.2018, 16.10.2018, ÇELİK. 43, 51. Yenir.

15- **Leucoagaricus leucothites**(Vittad.) Wasser, **Ak etlice;** Çayırlık alan ve Kavak (*Populus* sp.) ağaçlar altı, Hamur-Özdirek, 05° 85'829"K, 43° 76'225"D, 2120 m, 16.10.2018, ÇELİK. 42. Yenmez.

Bolbitiaceae Singer

16- **Panaeolus papilionaceus** (Bull.)Quél., **Süslü terscanı;** Hayvan gübresi üzeri, çayırlıklarda Hamur-Özdirek, Merkez Yukarı yol düzü, 05° 55'679"K, 43° 72'303"D, 2060 m, 20.05.2018, 24.10.2018, ÇELİK. 16, 31. Yenmez.

17- **Conocybe apala** (Fr.) Arnolds, **Ak yalinetek;** Çayırlık, Hamur-Özdirek, Merkez Yukarı yol düzü, 05° 58'676"K, 43° 71'345"D, 2043 m, 24.10.2018, ÇELİK. 21. Yenmez.

18- **Conocybe semiglobata** Kühner & Watling, **Kör yalinetek;** Çayırlık, park ve bahçelerde, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 56'587"K, 43° 71'323"D, 1978 m, 24.05.2018, ÇELİK. 22. Yenmez.

19- Conocybe subovalis Kühner & Watling

Piç yalinetek; Çayırlık, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 56'587"K, 43° 71'323"D, 1975 m, 24.05.2018, ÇELİK. 37. Yenmez.

Cortinariaceae R. Heim ex Pouzar

20- **Cortinarius purpurascens** Fr., **Ekli örümcekmantarı;** Kavak (*Populus* sp.), ağaçları altı ve asidik topraklarda, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 56'587"K, 43° 71'323"D, 1775 m, 01.05.2018, ÇELİK. 38. Yenir.

21- **Hebeloma mesophaeum** (Pers.) Quél., **Ala turpkokan;** Kavak (*Populus* sp.) ağaçları altı, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 56'587"K, 43°



71°323"D, 1775 m, 01.05.2018, 24.10.2018, ÇELİK, 2,23.
Yenmez.

Entolomataceae Kotl. & Pouzar

22- **Entoloma sericeoides** (J.E. Lange) Noordel.,

Seri kıvrıkbaba; Kavak (*Populus sp.*) ve Söğüt (*Salix sp.*), ağaçları altı, Merkez Yukarı yol düzü, 05° 56'587"K, 43° 71'323"D, 1732 m, 01.05.2018, ÇELİK, 24. Zehirli.

Inocybaceae Jülich

23- **Inocybe flocculosa** Sacc, **Aykümbet**; Kavak (*Populus sp.*) ve konifer ağaçları altı, Hamur-Özdilek, Merkez Yukarı yol düzü, 05° 55'477"K, 43° 70'463"D, 1898 m, 05.05.2018, 24.10.2018, ÇELİK, 1,39. Zehirli.

24-**Inocyb flocculosa** Sacc, **Aykümbet**; Kavak (*Populus sp.*), ağaçları altı, kalkerli topraklar, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 56'467"K, 43° 71'453"D, 1798 m, 24.10.2018, ÇELİK, 2,23. Zehirli.

25- **Pseudosperma rimosum** (Bull.) Matheny & Esteve-Rav., **Uysalkümbet**; Konifer ve Kavak (*Populus sp.*), ağaçları altı, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 55'679"K, 43° 72'303"D, 2060 m, 20.05.2018, 24.10.2018, ÇELİK, 3, 33. Zehirli.

Pleurotaceae Kühner

26- **Pleurotus eryngii** (DC.) Quél., **Çakşır mantarı**; Heliz (Çakşır) bitkisi üzeri, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 79'620"K, 43° 86'959"D, 2163 m, 05.04.2018, 20.05.2018, ÇELİK, 8. Yenir.

27- **Pleurotus ostreatus** (Jacq.) P. Kumm., **İstiridye mantarı**; Kavak (*Populus sp.*) ve Söğüt (*Salix sp.*) kütüğü üzeri, Hamur, Merkez Yukarı yol düzü, 05° 79'612"K, 43° 86'933"D, 2154 m, 20.05.2018, ÇELİK, 5. Yenir.

Pluteaceae Kotl. & Pouzar

28- **Pluteus romellii** (Britzelm.) Sacc., **Eli çitkirdi**; Kavak (*Populus sp.*) ve Söğüt (*Salix sp.*), kütüğü üzeri, Hamur, Merkez Yukarı yol düzü, 05° 79'612"K, 43° 86'933"D, 2154 m, 05.04.2018, 20.05.2018, ÇELİK, 40. Yenir.

29-**Pluteus salicinus** (Pers.) P. Kumm. **Söğüt çitkirdi**;

Odyn kalıntısı üzeri, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 79'412"K, 43° 86'833"D, 2154 m, 30.05.2018, 20.10.2018, ÇELİK, 41. Yenir.

30-**Volvopluteus gloiocephalus** (DC.) Vizzini, Contu & Justo, **Kakilvik**; Söğüt (*Salix sp.*) ve Kavak (*Populus sp.*) altı, Hamur, Merkez Yukarı yol düzü, 05° 80'612"K, 43° 83'953"D, 2178 m, 20.05.2018, ÇELİK, 45. Yenir.

Psathyrellaceae Vilgalys, Moncalvo & Redhead

31-**Coprinellus disseminatus** (Pers.) J.E. Lange, **Minikmürekkep**; Söğüt (*Salix sp.*) kütüğü üzeri, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 79'620"K, 43° 86'959"D, 2063 m, 05.10.2018, ÇELİK, 4, 20. Yenir.

32- **Coprinellus impatiens** (Fr.) J.E. Lange, **Cammürekkebi**; Söğüt (*Salix sp.*) kütüğü üzeri, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 79'620"K, 43° 86'959"D, 2063 m, 05.06.2018, ÇELİK, 26. Yenmez.

33- **Coprinellus micaceus** (Bull.) Vilgalys, Hopple & Jacq. Johnson, **Pullumürekkep**; Kavak (*Populus sp.*)

ve Söğüt (*Salix sp.*) kütüğü üzeri, Merkez Yukarı yol düzü, 05° 79'531"K, 44° 84'659"D, 2073 m, 05.05.2018, ÇELİK, 28. Yenmez.

34-**Coprinopsis nivea** (Pers.) Redhead, Vilgalys & Moncalvo, **Ak döbelen**; Hayvan gübresi üzeri, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 79'640"K, 43° 86'859"D, 2163 m, 05.06.2018, ÇELİK, 47. Yenmez.

35-**Coprinopsis atramentaria** (Bull.) Redhead, Vilgalys & Moncalvo, **Kütük döbeleni**; Kavak (*Populus sp.*) kütüğü üzeri, Merkez Yukarı yol düzü, 05° 79'531"K, 44° 84'659"D, 2073 m, 05.10.2018, ÇELİK, 48. Zehirli.

36-**Parasola plicatilis** (Curtis) Redhead, Vilgalys & Hopple, **Açık sevelen**; Çayırlık, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 80'660"K, 43° 85'759"D, 2103 m, 05.04.2018, 20.05.2018, ÇELİK, 29. Yenmez.

37-**Psathyrella candolleana** (Fr.) Maire, **Güzel pulcuklu**; Söğüt (*Salix sp.*), ağaçları altı, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 55'584"K, 43° 71'433"D, 1995 m, 24.05.2018; 06.10.2018, ÇELİK, 30, 35. Yenmez.

38- **Psathyrella lutensis** (Romagn.) Bon, **Üç pulcuklu**; Kavak (*Populus sp.*) ve Söğüt (*Salix sp.*), ağaçları altı, Hamur, 05° 56'687"K, 43° 71'523"D, 1992 m, 20.05.2018, ÇELİK, 32. Yenmez.

39- **Psathyrella prona** (Fr.) Gillet, **Yel pulcuklu**; Çayırlık alan, Hamur özdirek köyü, 05° 56'867"K, 43° 71'323"D, 2175 m, 06.05.2018, 10.10.2018, ÇELİK, 27, 66. Yenmez.

40-**Psathyrella spadiceogrisea** (Schäff.) Maire, **Yol pulcuklu**; Ağaç kalıntıları üzeri, Merkez Yukarı yol düzü, 05° 55'654"K, 43° 71'476"D, 1895 m, 26.05.2018; 13.10.2018, ÇELİK, 30, 35. Yenmez.

Strophariaceae Singer & A.H. Sm.

41- **Agrocybe dura** (Bolton) Singer, **Yaz meteliği**; Çayırlık alanlarda, Hamur, 05° 56'664"K, 43° 72'523"D, 2085 m, 06-08.05.2018, ÇELİK, 57. Yenir.

42-**Agrocybe pediades** (Fr.) Fayod, **Kollu metelik**; Çayırlık alanlarda, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 55'474"K, 43° 71'513"D, 2195 m, 24.05.2018; 06.10.2018, ÇELİK, 52, 60. Yenir.

43- **Agrocybe paludosa** (J.E. Lange) Kühner & Romagn. ex Bon, **Yaş metelik**; Çayırlık alanda, Hamur, Merkez Yukarı yol düzü, 05° 55'474"K, 43° 71'513"D, 2165 m, 22.05.2018, ÇELİK, 53. Yenmez.

44- **Agrocybe praecox** (Pers.) Fayod, **Bahar meteliği**; Kavak (*Populus sp.*), ağaçları altı, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 55'574"K, 43° 71'534"D, 2095 m, 06.05-10.2018, ÇELİK, 55. Yenir.

45-**Pholiota aurivella** (Batsch) P. Kumm., **Sarı pulbaş**; Söğüt (*Salix sp.*), ağaçları üzeri, Merkez Yukarı yol düzü, 05° 55'632"K, 43° 71'513"D, 2095 m, 06.10.2018, ÇELİK, 54. Yenmez.

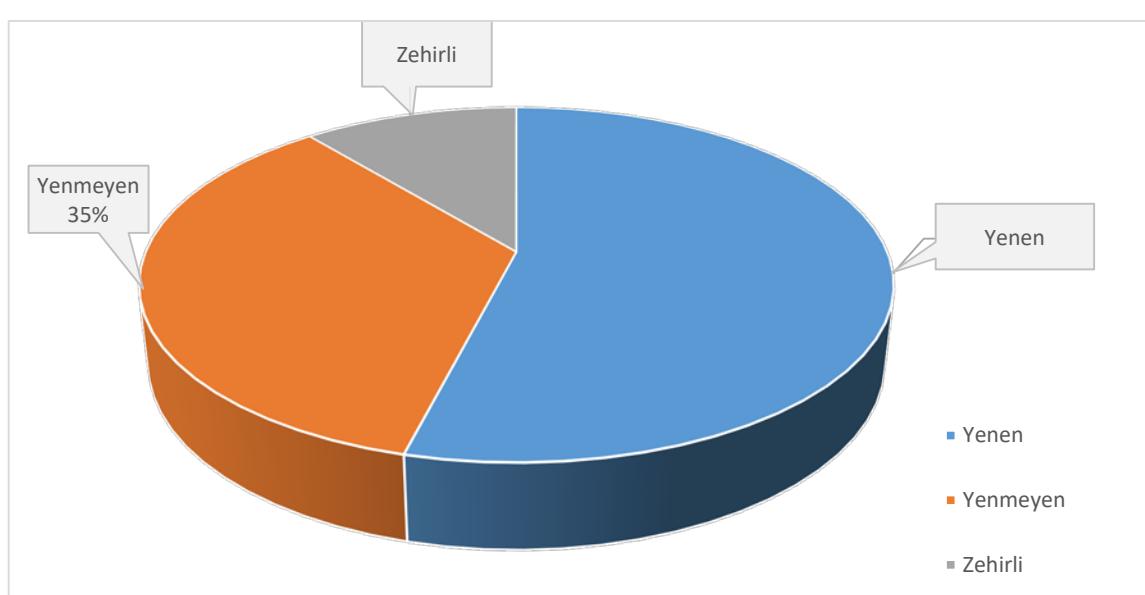
46-**Pholiota gummosa** (Lasch) Singer, **Yapışkan pulbaş**; Kavak (*Populus sp.*), ağaçları altı, Hamur-Ceylanlı, 05° 55'474"K, 43° 71'513"D, 1995 m, 16.10.2018, ÇELİK, 65. Yenmez.

**Hymenogastraceae** Vittad.47-*Psilocybe coronilla* (Bull.) Noordel., **Kefgarık**;

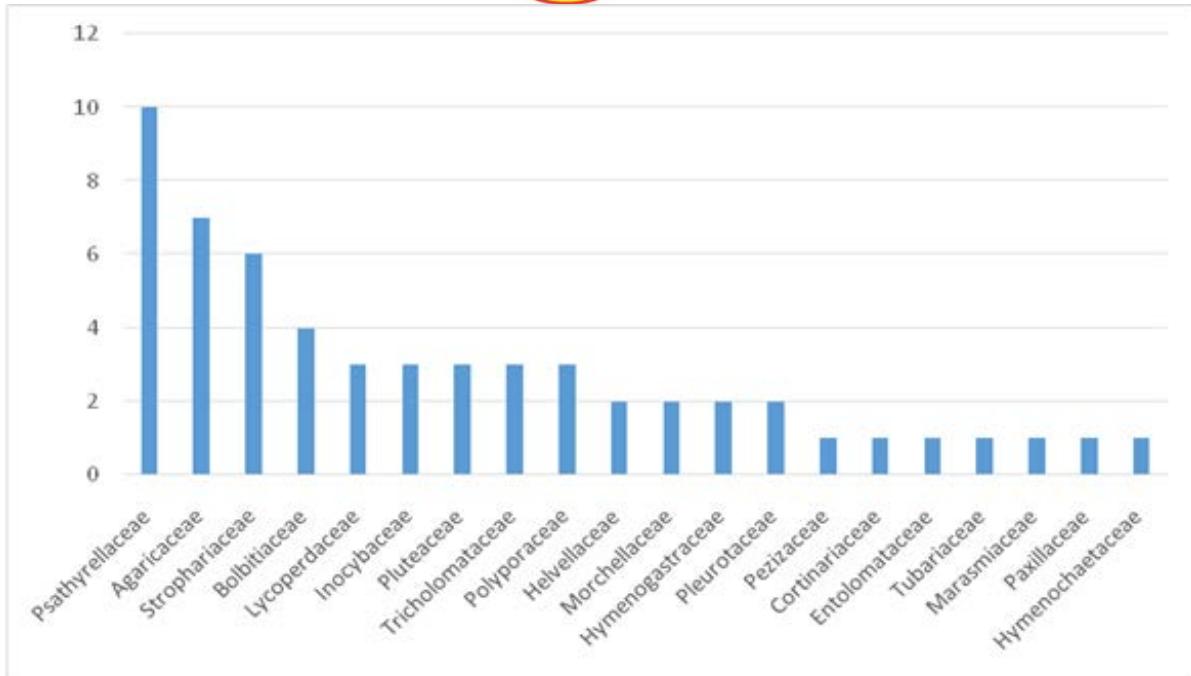
Çayırlıkk, Hamur-Ceylanlı, 05° 55'674"K, 43° 71'713"D, 2165 m, 06.05.2018, ÇELİK, 67. Yenir. Yenir.

Tubariaceae Vizzini48- *Tubaria furfuracea* (Pers.) Gillet, **Fırkırlı****tubarya**; Söğüt (*Salix* sp.), ağaçları üzeri, Merkez Yukarı yol düzü, 05° 55'867"K, 43° 71'583"D, 2105 m, 22.05.2018, ÇELİK, 62. Yenmez.**Marasmiaceae** Roze ex Kühner49- *Marasmius oreades* (Bolton) Fr. **Mıhbaşı**;Çayırlıkk alan ve Kavak (*Populus* sp.), ağaçları altı, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 55'387"K, 43° 71'513"D, 2155 m, 06-22.05.2018, ÇELİK, 61, 56. Yenir.**Tricholomataceae** R. Heim ex Pouzar50- *Lepista irina* (Fr.) H.E. Bigelow, **Süslü cincile**;Kavak (*Populus* sp.), ağaçları altında, Hamur-Ceylanlı, Merkez Yukarı yol düzü, 05° 55'387"K, 43° 71'513"D, 2130 m, 22.05.2018, ÇELİK, 63. Yenir.51- *Lepista personata* (Fr.) Cooke, **Diken mantarı**;Kavak (*Populus* sp.), ağaçları altında, Hamur-özdirek, Merkez Yukarı yol düzü, 05° 55'387"K, 43° 71'513"D, 2155 m, 06-22.05.2018, ÇELİK, 58, 70. Yenir.52-*Tricholoma populinum* J.E. Lange,**Kavakkarakız**; Kavak (*Populus* sp.), ağaçları altı, Hamur ve Merkez Yukarı yol düzü, 05° 55'457"K, 43° 71'554"D, 1978 m, 24.10.2018, ÇELİK, 59, 69. Yenir.**Boletales****Paxillaceae** Lotsy53- *Paxillus involutus* (Batsch) Fr., **Pax mantarı**;Kavak (*Populus* sp.), ağaçları altı, Hamur-Ceylanlı, 05° 55'387"K, 43° 71'513"D, 2145 m, 06-16.05.2018, ÇELİK, 68. Zehirli.**Hymenochaetales****Hymenochaetaceae** Donk54- *Phellinus igniarius* (L.) Quél., **Kara toynak**;Huş (*Betula* sp.), ağaç gövdesi üzeri, Hamur ve Merkez Yukarı yol düzü, 05° 55'457"K, 43° 71'554"D, 2078 m, 24.05.2018, ÇELİK, 73, 79. Yenmez.**Polyporales****Polyporaceae** Fr. ex Corda55-*Fomes fomentarius* (L.) Fr., **Kavmantarı**;Söğüt (*Salix* sp.), ağaçları üzeri, Hamur-Özdirek köyü, 05° 55'776"K, 43° 71'654"D, 1988 m, 24.10.2018, ÇELİK, 74, 76. Yenmez.56- *Lentinus tigrinus* (Bull.) Fr., **Kaplanmatarı**;Söğüt (*Salix* sp.), kütüğü üzeri, Hamur-ceylanlı ve Merkez Yukarı yol düzü, 05° 55'457"K, 43° 71'554"D, 2089 m, 22.05.2018, ÇELİK, 87, 96. Yenir.57- *Polyporus squamosus* (Huds.) Fr.,**Görkemli**; Söğüt (*Salix* sp.), kütüğü üzeri, Hamur-ceylanlı-özdirek köyü ve Merkez Yukarı yol düzü, 05° 55'487"K, 43° 71'384"D, 2119 m, 04-24.10.2018, ÇELİK, 85, 93. Yenir.**Tartışma**

Bu çalışma sonunda, Ağrı ili Merkez ve Hamur ilçesi sınırları içerisinde yer alan bölgede 2018-2019 yılları arasında doğal olarak yetişen Pezizomyces ve Agaricomycetes sınıflarına ait olan 20 familya'ya ait toplam 57 makromantar taksonu tespit edilmiştir. Bu mantar örneklerinde 31 yenen, 20 yenmeyen, 6 zehirli özellikleştir (Şekil 3). Yenen türler toplam türlerin %54, yenmeyenler %35, zehirli türler ise %11'ni oluşturmaktadır. Ayrıca bölge genelinde tanıtan ve besin olarak tüketilen mantar takson sayısı oldukça azdır. Bunun en önemli sebebi yerel halkın çoğunuğunun tüm mantarların zehirli olduğunu düşündüğü öngörmektedir.



Şekil 3. Yörede tespit edilen makromantarların yenilebilirlik durumuna göre dağılımı.



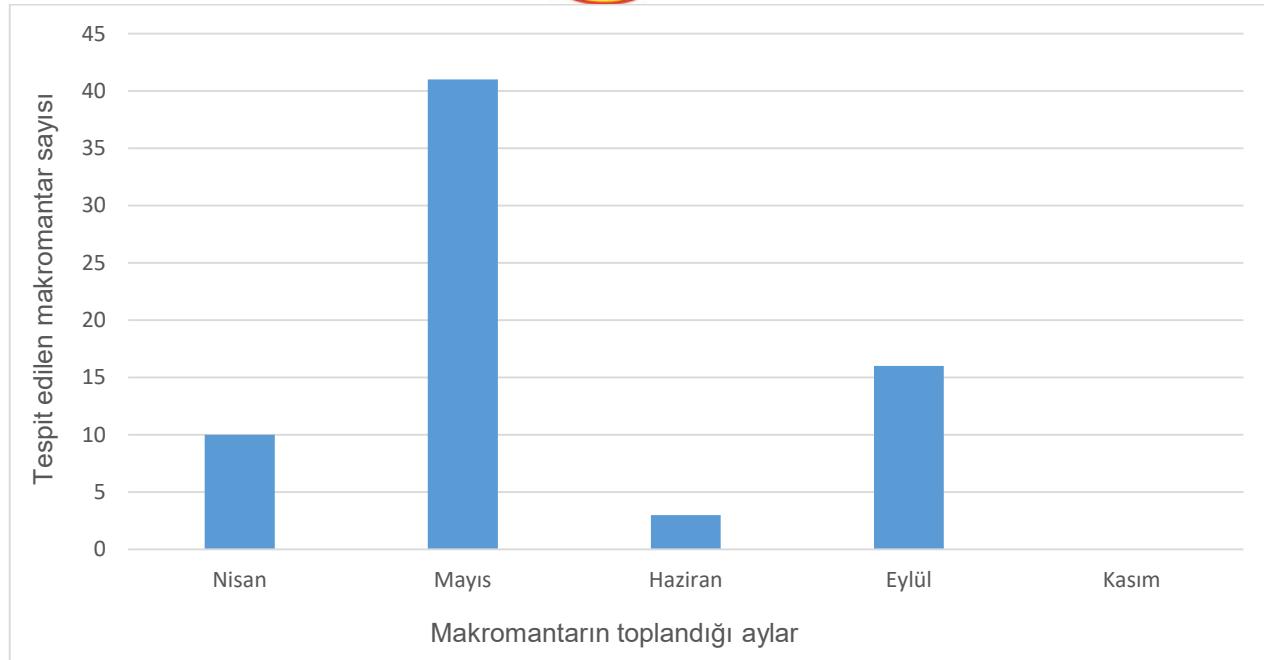
Şekil 4. Tespit edilen makrofungal taksonlarının familyalara göre dağılımı.

Tablo 1. Bazı yeniden türlerin yöresel isimleri

Yenen Türler	Yenen Türlerin Yöresel İsimleri	Türlerin Kullanım Amaçları
<i>Agaricus bisporus</i>	Çayır mantarı (Demirel ve Koçak, 2016)	Yemeklik olarak kullanılmaktadır.
<i>Agrocybe dura</i>	Çayır mantarı (Demirel ve ark., 2017)	Yemeklik olarak kullanılmaktadır.
<i>Lycoperdon perlatum</i>	Fıssakuri (Demirel ve ark., 2017)	Halk arasında yara üzerine sürüllererek kullanılmaktadır.
<i>Pleurotus ostreatus</i>	Ağaç mantarı (Çağlı ve Öztürk, 2020)	Yemeklik olarak kullanılmaktadır.
<i>Pleurotus eryngii</i>	Çakşır mantarı (Demirel ve Koçak, 2016)	Yemeklik olarak kullanılmaktadır.

Çalışma bölgesinde en fazla makromantar örneği Mayıs ve Eylül aylarında toplandığı tespit edilmiştir. Bunun sebebi ilkbahar yağışlarının ve sıcaklığın mantar yetişmesi için uygun olmasından kaynaklanmaktadır. Haziran ayında daha az örnek toplandığı tespit edilmiştir. Temmuz, Ağustos ve Kasım aylarında ise ekolojik

faktörler makromantaların yayılışına ve yetişmesine uygun olmadığından; ayrıca Aralık, Ocak, Şubat ve Mart aylarında ise düşük sıcaklık ve toprağın karla kaplı olmasından dolayı herhangi bir örnek toplanamamıştır (Şekil 5).



Şekil 5. Araştırma yöresinde toplanan makromantoların aylara göre dağılımı.

Tablo 2. Araştırma yöresine yakın bölgelerde yapılmış olan çalışmalarla benzerlik durumu

Aştırma Yoresi	Tespit edilen makromantar sayısı	toplam takson sayısı	Benzerlik oranı (%)
Ağrı (Demirel ve ark., 2003)	47	27	57.44
Malazgirt (Muş) (Akçay ve ark., 2012)	50	24	48
Karz Dağı (Bitlis-Tatvan) (Sadullahoglu, 2013)	79	28	35.44
Zilan Vadisi (Van-Erciş) (Koçak, 2014)	98	51	52.04
Van (Demirel ve ark., 2015)	122	41	33.60
Kağızman (Kars) (Uzun ve ark., 2020)	86	41	47.67
Muradiye (Van) (Çağlı ve Öztürk, 2020)	86	26	30.23

Aynı zamanda, Tablo 1'de görüldüğü üzere çalışmada teşhis edilen makromantoların çalışma alanına yakın olan bölgelerde yapılmış benzer çalışmalar; Malazgirt (Muş) (Akçay ve ark., 2012); Karz Dağı (Bitlis-Tatvan) (Sadullahoglu, 2013); Zilan Vadisi (Van-Erciş) (Koçak, 2014); Van (Demirel ve ark., 2015) ve Kağızman (Kars) (Uzun ve ark., 2020) ile karşılaştırılmıştır. Bununla birlikte, Malazgirt (Muş) %48; Bitlis-Tatvan %35.44; Van-Erciş %52.04; Van %33.60 ve Kağızman (Kars) %47.67 gibi benzerlik oranlarının olduğu görülmektedir. Mevcut sonuçlara göre bu benzerlik ve farklılıkların olması belirtilmiş olan çalışma alanlarının sahip oldukları kendilerine ait bitki örtüsü ve iklim özelliklerinden kaynaklandığı düşünülmektedir. İlgili araştırma bölgemizde yaygın makromantar taksonu *Psathyrellaceae* familyasına ait olan türlerin olduğu belirlenmiştir. Ayrıca bu çalışma ile tespit edilen türler, ülkemizde Türkiye makromantoları olarak kaydedilmiştir.

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Ecological Factors Influencing the Occurrence of *Armillaria mellea* (Basidiomycota, Agaricales, Physalacriaceae) in Yuvacık Dam Watershed in Kocaeli, Türkiye

Sabiha ACER¹, Ersel YILMAZ^{2*}
Ayhan KARAKAYA³

*Corresponding author: ersel@iuc.edu.tr

¹ İstanbul Üniversitesi – Cerrahpaşa, Orman Fakültesi, Orman Mühendisliği Bölümü, İstanbul / sacer@iuc.edu.tr

² İstanbul Üniversitesi – Cerrahpaşa, Orman Fakültesi, Orman Mühendisliği Bölümü, İstanbul / ersel@iuc.edu.tr

³ TC Orman Genel Müdürlüğü, Kavak ve Hızlı Gelişen Orman Ağaçları Araştırma Enstitüsü Müdürlüğü / ayhankarakaya@ogm.gov.tr

Abstract: The occurrence of *Armillaria mellea* (Vahl) P. Kumm. and the ecological characteristics of this fungus were studied in Kocaeli, Yuvacık dam basin mixed-broad leaved forests. During the surveys, we analyzed the sporocarps (fruiting bodies) of *A. mellea* growing up on woody plants in plots selected by cluster sampling in the Yuvacık dam watershed dominated by broad-leaved forests. The Runs test results showed that randomness rules complied in the selection of the plots, and there was no tendency ($p= 0.109 > 0.05$, $z= -1.603$). The presence/absence of *A. mellea* and environmental variables were tested with Chi-square analysis, and the temperature differed among these environmental variables. To the dendrogram, *A. mellea* was mainly seen in the south of the study area and preferred western aspects. It is understood that this macrofungus prefers the south of the study area because of the altitude. Our data showed that sporocarps of *A. mellea* generally occurred in the western aspect, at temperatures of 15–20°C, >80% humidity and 800–1000 m altitude. Our logistic regression analysis model ($z=-9.508+0.307 \times \text{temperature}+0.081 \times \text{humidity}$) showed that if the temperature and humidity change by 1 unit in the region, sporocarp formation is affected by 36% and 8.4%, respectively.

Key words: *Armillaria mellea*, Sporocarp, Chi-square analysis, Logistic regression, Temperature, Humidity, Altitude

Yuvacık Barajı Havzasında (Kocaeli, Türkiye) *Armillaria mellea* (Basidiomycota, Agaricales, Physalacriaceae)'nın Oluşumunu Etkileyen Ekolojik Faktörler

Öz: Kocaeli, Yuvacık baraj havzası karışık-geniş yapraklı ormanlarında *Armillaria mellea* (Vahl) P. Kumm.'nın oluşumu ve bu mantarın ekolojik özellikleri incelenmiştir. Araştırmalar sırasında, geniş yapraklı ormanların hâkim olduğu Yuvacık baraj havzasında küme örneklemesi ile seçilen parsellerde odunsu bitkiler üzerinde yetişen *A. mellea* sporokarpları (fruktifikasyon organları) analiz edilmiştir. Runs testi sonuçları, parsellerin seçiminde rastgelelik kurallarına uyulduğunu ve herhangi bir eğilimin olmadığını göstermektedir ($p= 0.109 > 0.05$, $z= -1.603$). *A. mellea*'nın varlığı/yokluğu ve çevresel değişkenler Ki-kare analizi ile test edilmiş ve sıcaklık bu çevresel değişkenler arasında farklılık göstermiştir. Dendrograma göre *A. mellea* daha çok çalışma alanının güneyinde görülmekte ve batı bakını tercih etmektedir. Bu makrofungsun çalışma alanın güneyini tercih etmesinin sebebinin ise yükseklik olduğu anlaşılmaktadır.



Verilerimiz, *A. mellea* sporokarplarının genel olarak batı bakıda, 15-20°C, >80 % nemde ve 800-1000 m yüksekliklerde bulunduğu göstermektedir. Elde ettiğimiz lojistik regresyon analizi modeli ($z = -9.508 + 0.307 \times \text{temperature} + 0.081 \times \text{humidity}$), bölgede sıcaklık ve nem 1 birim değişirse sporokarp oluşumunun sırasıyla %36 ve %8.4 oranında etkilendiğini göstermektedir.

Anahtar kelimeler: *Armillaria mellea*, Sporokarp, Ki – kare analizi, Lojistik regresyon, Sıcaklık, Nem, Yükseklik

Introduction

The genus *Armillaria* (Basidiomycota, Agaricales, Physalacriaceae) are natural components of the mycoflora in many forests worldwide. They occur worldwide in boreal, temperate, and tropical forests, and through diverse parasitic activities, they influence many host species. The genus *Armillaria* is, therefore, significantly considered in the ecology and management of many natural forests (Kile et al., 1991). It also has a long and controversial taxonomic history. This discussion goes back to the Danish botanist Martin Vahl's description of *Agaricus melleus* (syn: *Armillaria mellea*) in 1787. Many taxonomic revisions have subsequently been published addressing the controversial genus name. *Armillaria* Fr. Staude is now the broadly accepted generic name with *Armillaria mellea* (Vahl) P. Kumm. (honey mushroom) (Balmantarı) representing the type of the genera. (Volk & Burdsall, 1995; Watling et al., 1982; Coetze et al., 2018, Sesli et al., 2020). *A. mellea* was recognized as a mixture of at least 10 different "biological species" in the late 1970s (Hagle, 2010; Lushaj et al., 2010) and *A. mellea* sensu stricto is described as a pathogen in broadleaf and conifers (Kile et al., 1991; Guillaumin et al., 1993; Lushaj, 2008).

Among the seven species of *Armillaria* spread in Europe, six species are wood decay fungi with wide distribution and of great ecological and economical importance (Kile et al., 1991; Guillaumin et al., 1993; Lushaj, 2008). In the Balkan countries, seven species have been reported from Albania (Lushaj, 2008), six species from Slovenia and the Czech Republic (Munda, 1997; Jankovský, 2003), and five from Greece and Serbia (Tsopelas, 1999; Lushaj et al., 2010). There have been identified eleven species in Türkiye; *A. borealis* Marxm. & Korhonen (Kuzey balmantarı), *A. bulbosa* (Romagn.) Kile & Watling, *A. cepistipes* Velen.(Narin balmantarı), *A. gallica* Marxm. & Romagn. (Şiş balmantarı), *A. mellea*, *A. obscura* (Schaeff.) Herink (Top balmantarı) *A. ostoyae* (Romagn.) Herink (Külahlı balmantarı), *A. socialis* (DC.) Fayod (Yığın balmantarı), *A. solidipes* Peck, *Desarmillaria ectypa* (Fr.) R.A. Koch & Aime (Sivri balmantarı), and *D. tabescens* (Scop.) R.A. Koch & Aime (Sesli and Denchev, 2014, Sesli et al., 2020, Solak and Türkoğlu, 2022). *A. mellea* has been recorded on deciduous species such as *Quercus*, *Fagus*, *Castanea*, *Carpinus*, *Populus*, *Salix*, *Corylus*, and *Juglans* and coniferous species such as *Pinus*, *Abies*, and *Cedrus* in

Türkiye (Selik, 1973; Sümer, 1977; Doğan and Öztürk, 2006; Türkkul, 2008).

In the life cycle of *Armillaria* species, basidiospores germinate on the woody substrate, and formed mycelium colonizes this wood tissue. Healthy trees are infected either by root contacts with infected woody tissue or by rhizomorphs growing out from the infected centre. *Armillaria* invades the root system and the lower stem of the infected trees, killing the cambium and causing heart rot. Sporocarps develop on dead/moribund wood tissue and release basidiospores into the environment (Heinzelmann et al., 2019). *Armillaria* overwinters as a mycelium or rhizomorph on the stems or roots of infected or dead trees. *Armillaria* overwinters as a mycelium or rhizomorph on the stems or roots of infected or dead trees. Rhizomorphs are black, branched, and filamentous structures that develop from spores and supply nutrient and air transmission (Selik, 1986; Baumgartner et al., Hagle, 2010; 2011;).

Most species of *Armillaria* are considered to be facultative necrotrophs that have parasitic and saprophytic phases. First, *Armillaria* colonizes the cambium of living roots (parasitic phase). Then, the fungus kills the cambium, causing a necrotic lesion beneath the root bark. Lastly, the fungus feeds on the dead tissue (saprophytic phase). This saprophytic capability makes *Armillaria* root disease so difficult to prevent (Baumgartner et al., 2011). Relatively few studies have focused on the distribution of the species, physical space factors, or stand type characteristics of the forest (Bruhn et al., 2000). *A. mellea* was associated with high pathogenicity in deciduous and coniferous except for pines in alkaline soils in England and observed that it grows rhizomorphs under soil over short distances (Rishbeth, 1982). Termorshuizen and Arnolds (1994) pointed out that host preferences of *Armillaria* species may be affected by soil type and their virulence may be related to tree vitality affected by soil type. It was detected on broad-leaved hosts growing on clay and loess soils in The Netherlands. It was recognized as a dominant parasite on broadleaved forests and fruit trees, especially in warm regions in the Czech Republic (Jankovský, 2003). This species was identified in deciduous and coniferous forests below 1100 m altitude in Greece (Tsopelas, 1999). *A. mellea* was determined on oaks between 700–1050 m altitude in Serbia (Keča et al., 2009), while on a few broadleaved and conifers species,



below 1200 m in Albania (Lushaj et al., 2010). It was a species with low frequency in Poland (Łakomy, 2006) and the Ukrainian Carpathians (Tsykun et al., 2012).

Macrofungi studies in Turkey focused on its biodiversity rather than the ecological characteristics of fungi. There are a few papers based on this framework in our research area (Akata et al. 2018; Doğan et al., 2021). Yuvacık dam watershed, is located east of the Marmara region and in the transition zones of the three phytogeographic regions. The study area consists predominantly of pure and mixed broadleaved trees and mixed stands of deciduous and coniferous trees. *A. mellea*, which may attack deciduous and coniferous species, has been identified in previous studies in the research area. The present study aims to find out the ecological characteristics of *A. mellea* in the Yuvacık dam watershed area. We compared the presence/absence status of *A. mellea* to various environmental parameters in the sampled plots and analysed them with logistic regression.

Material and Method

Study Area

Yuvacık dam watershed is located in the east of the Marmara region and the south of the Izmit district of Kocaeli province. It is coordinated at 40°32' N and 29° 58' E and covers 25.759 ha (Fig. 1 (Google Earth, 2022)).

The Yuvacık Dam is fed by three main streams, numerous creeks and the Hüseyinli pond, which is used for irrigation. The average altitude of the area varies from 170-1300 m (Beşkardeş, 2012; Efe et al., 2013).

The study area is in the transition zones of the three different phytogeographic regions including the Euro-Siberian (Sub-Euxine, underside section), the Mediterranean (the Aegean, underside section), and Irano-Turanian (the side of the inner Anatolia). In the watershed, the predominant forest tree species are *Fagus sylvatica* subsp. *orientalis* Lipsky, *Quercus* species, *Carpinus betulus* L., *Pinus nigra* Aiton, *P. sylvestris* L. and *Abies nordmanniana* subsp. *equi-trojani* (Asch. & Sint. ex Boiss.) Coode & Cullen). In addition, some other species are also found, such as *Castanea sativa* Mill., *Corylus avellana* L., *Alnus glutinosa* (L.) Gaertn., *Populus tremula* L., and *Juglans regia* L. in the area. Forests are made up of pure stands as well as mixed stands of these particular species (Beşkardeş, 2012; Efe et al., 2013). To the Thornthwaite methodology, the annual average precipitation and temperature are 1038.7 mm and 9.5 °C, respectively. The study area has a climate model that reflects similar oceanic climate parameters such as humidity, mild temperature and no water shortage (Zengin et al., 2005). The prevailing soil type is eutric cambisol and orthic luvisol in the FAO system (FAO 2022).

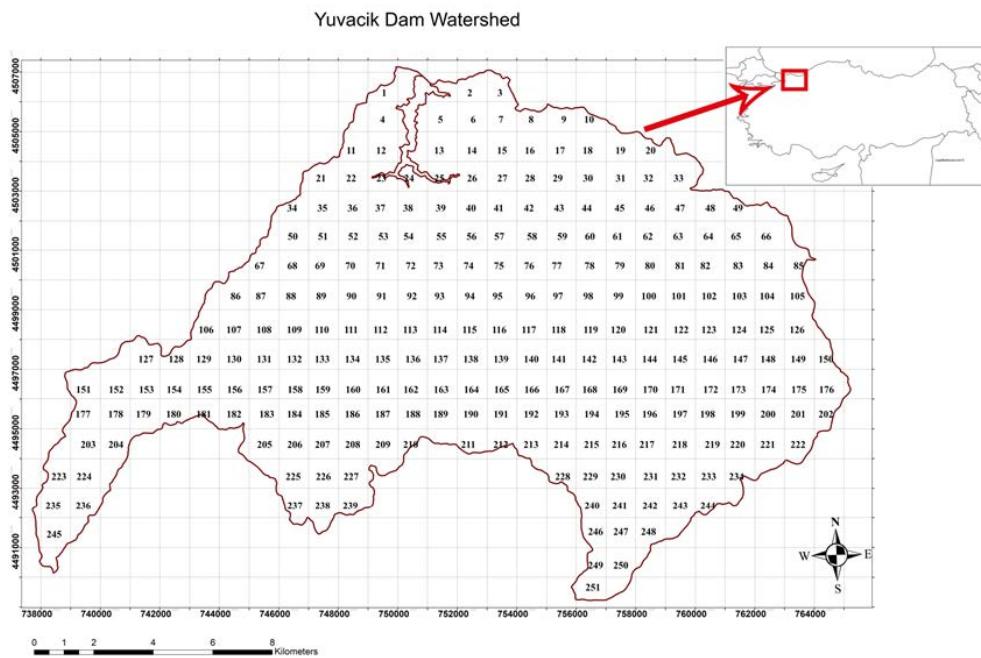


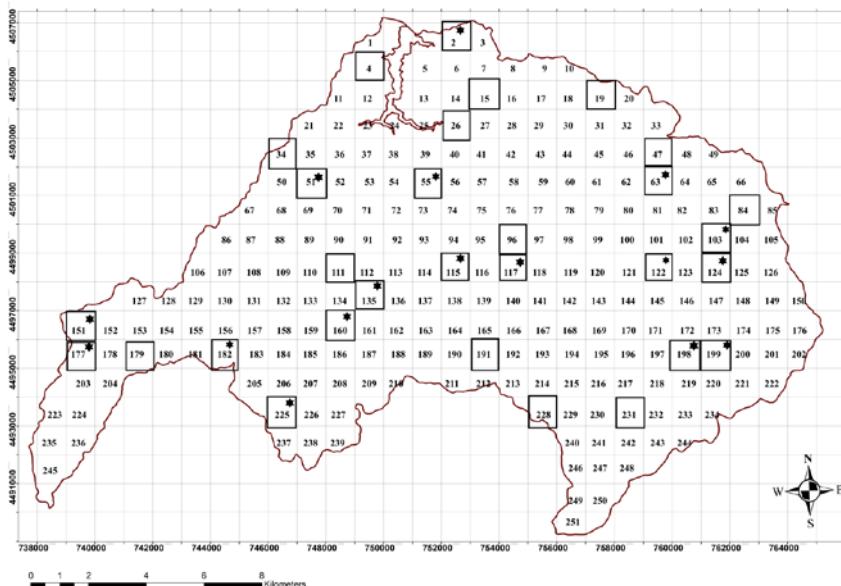
Figure 1. Location of Yuvacık dam watershed in the Marmara Region

The cluster sampling method was used in the Yuvacık dam watershed, spread over a 25759 hectar wide geographical area. The study area was divided into plots of 1 km² (grids) and 251 plots were obtained. 30 of 251 plots which correspond to approximately 12% of the study

area were selected by simple random sampling (Fig 2). In our research, the transect method used 4 parallel transect sections running across plots of 1 km², 200 m apart as sampling patterns.



Yuvacik Dam Watershed

Figure 2. 30 sampled plots in the Yuvacik dam watershed (*where *Armillaria mellea* occurs)

Fungal Survey

Field surveys were systematically conducted during the vegetation season for symptomatic hosts and *A. mellea* sporocarps. Fallen and living trees, shrubs and stumps were examined in four parallel transects, each 1 km in length. Whenever sporocarps were encountered on a plot, at least one individual mushroom was collected from genets representing each morphology type. The photographs were taken in their habitat. During the field studies, environmental variables of the sampled plots such as stand type, age class, tree species, crown closure, aspect, altitude, temperature, and air humidity were noted (Table 1). Microscopic characteristics of sporocarps specimens were examined using a Bresser light microscope. The measurements of at least 25 spores per specimen were taken. Identification was made from the relevant literature (Breitenbach and Kränzlin, 1991). Collected specimens turned into fungarium specimens and preserved in the cabinets by İZT-391 (3828) numbering at Poplar and Fast-Growing Forest Trees Research Institute, Kocaeli.

Statistical analysis

The Runs Test was performed to control whether the field study was on randomly selected plots. Chi-square analysis was used to determine whether there was a relationship among environmental variables in plots where *A. mellea* was observed. Hierarchical cluster analysis was performed considering ecological similarities and differences among plots. Besides, a dendrogram was created to show the similarity of the plots according to the environmental variables. In this

paper, to find out the correlation between a set of independent variables such as temperature (°C), humidity (%), altitude (m), age class, and crown closure and the binary dependent variable with two possible values (presence=1/absence=0) logistic regression analysis was performed (Küçüker et al., 2010).

$$P(Y) = \frac{e^z}{1 + e^z} \rightarrow P(Y) = \frac{1}{1 + e^{-z}}$$

z is a linear combination of independent variables.

$$Z = \beta_0 + \beta_1 X_1 + \beta_2 X_2 + \dots + \beta_n X_n$$

$\beta_0, \beta_1, \beta_2, \dots, \beta_n$ are regression coefficients.

The calculation of the regression coefficients is as follows:

$$\ln \left(\frac{P(Y)}{Q(Y)} \right) = \beta_0 + \beta_1 X_1 + \beta_2 X_2 + \dots + \beta_n X_n$$

$$\frac{P(Y)}{Q(Y)} = e^{\beta_0 + \beta_1 X_1 + \beta_2 X_2 + \dots + \beta_n X_n} = e^{\beta_n X_n}$$

$$Q(Y) = 1 - P(Y)$$

The odds of an event occurring are calculated as the ratio of the probability of the property being present compared to the probability of it being absent. Odds= $(P(Y))/(1-P(Y))$ Which means, the Exp (β) value of each parameter is considered as the odds ratio. Exp (β_i) values indicate to what extent the dependent variable Y is more likely to be observed with the effect of the variable X_i (Akarp, 2016).

Table 1. Sampling plots, hosts and environmental variables of *Armillaria mellea*

Plot No	Pre/Abs (1/0)	Coordinates	Altitude (m)	Slope (°)	Temperature (°C)	Humidity (%)	Aspect*	Host**	Colonization Site	Age Class***	Crown Closure****
2	1	36T/4506955-754968	537	31	13	81	NE	OK	Collar	BC	2
4	0	35T/4505440-749692	341	46	9	79	N	-	-	BC	2
15	0	35T/4504656-753915	612	24	8	84	S	-	-	BC	2
19	0	36T/4504596-757176	789	26	10	78	SW	-	-	BC	2
26	0	35T/4503521-752445	187	17	5	68	SE	-	-	BC	2
34	0	35T/4502608-746524	645	10	6	81	SW	-	-	BC	2
47	0	36T/4502324-759452	1012	23	6	77	SW	-	-	BC	2
51	1	35T/4501076-747851	459	26	12	77	N	OK	Collar	BC	2
55	1	36T/4501924-751604	587	10	8	74	NE	OK	Collar	BC	2
63	1	36T/4501672-759596	898	29	6	78	SW	OK	Stump	BC	2
84	0	36T/4500908-762885	1167	19	7	85	W	-	-	BC	2
96	0	36T/4499536-754539	1119	9	7	75	SW	-	-	CD	3
103	1	36T/4499040-761978	632	19	8	79	NW	OK	Stump	BC	2
111	0	35T/4498488-748296	592	44	15	86	SW	-	-	BC	2
115	1	35T/4498827-752720	886	25	17	83	W	BE	Stump	CD	3
117	1	36T/4498396-754532	1120	10	18	86	NW	BE	Collar	CD	3
122	1	36T/4498164-759912	1261	15	17	77	SE	OK	Stump	BC	2
124	1	36T/4498496-761687	743	29	11	77	NE	BE	Stump	BC	3
135	1	35T/4497228-749676	1061	27	16	82	SW	BE	Stump	CD	3
151	1	35T/4496392-739216	1041	28	20	72	SE	BE	Stump	CD	3
160	1	35T/4496768-748156	843	26	16	81	W	BE	Stump	CD	3
177	1	35T/4495642-739516	903	16	17	78	SW	HBM	Stump	CD	3
179	0	35T/4495080-741356	842	27	15	67	W	-	-	BC	2
182	1	35T/4495496-744556	987	7	17	62	NE	HBM	Stump	CD	2
191	0	35T/4495584-753917	1186	7	14	55	E	-	-	CD	3
198	1	36T/4495726-760196	991	15	23	76	SE	OK	Stump	BC	2
199	1	36T/4495612-761875	900	23	15	81	E	OK	Stump	BC	2
225	1	35T/4493436-746768	1042	19	15	57	NW	OK	Stump	BC	2
228	0	36T/4493800-755480	975	18	12	60	SE	-	-	BC	2
231	0	36T/4493560-758760	850	8	16	56	SW	-	-	BC	2

*N: North, S: South, W: West, E: East, SW: Southwest, NW: Northwest, SE: Southeast

**OK: Oak, HBM: Hornbeam, BE: Beech

***a (0 – 8 cm), b (9 – 20 cm), c (21 – 36 cm), d (36 - 52 cm) Diameter at Breast Height (Age class)

****1 (11 – 4 %), 2 (41 – 70%), 3 (71 – 100%) Crown closure



To investigate whether environmental factors affect the presence of sporocarps of *A. mellea* in the plots, we analysed all variables together and removed the least expressive independent variable stepwise with Binary Logistic and Backward LR method on IBM SPSS Statistics 21 (IBM CR, 2012).

Results

According to the Runs test, there was no relationship between the presence of *A. mellea* and the selection order of the plots ($p= 0.109 > 0.05$, $z= -1.603$). Our results showed that randomness rules complied in

the selection of the plots, and there was no tendency. *A. mellea* was detected in 17 of the 30 plots. This count represents 56% of the study area. All of the sporocarps of *A. mellea* were found on deciduous trees and particularly on stumps. 71% of plots, where *A. mellea* sporocarps occur were at an altitude of >800 m. Our determinations showed that they grew up mostly in the western aspect, "bc" age class and "2" crown closure. In addition, its sporocarps occurred in a high frequency at a temperature of 15–20 °C and >80 humidity (Table 1; Fig 3).

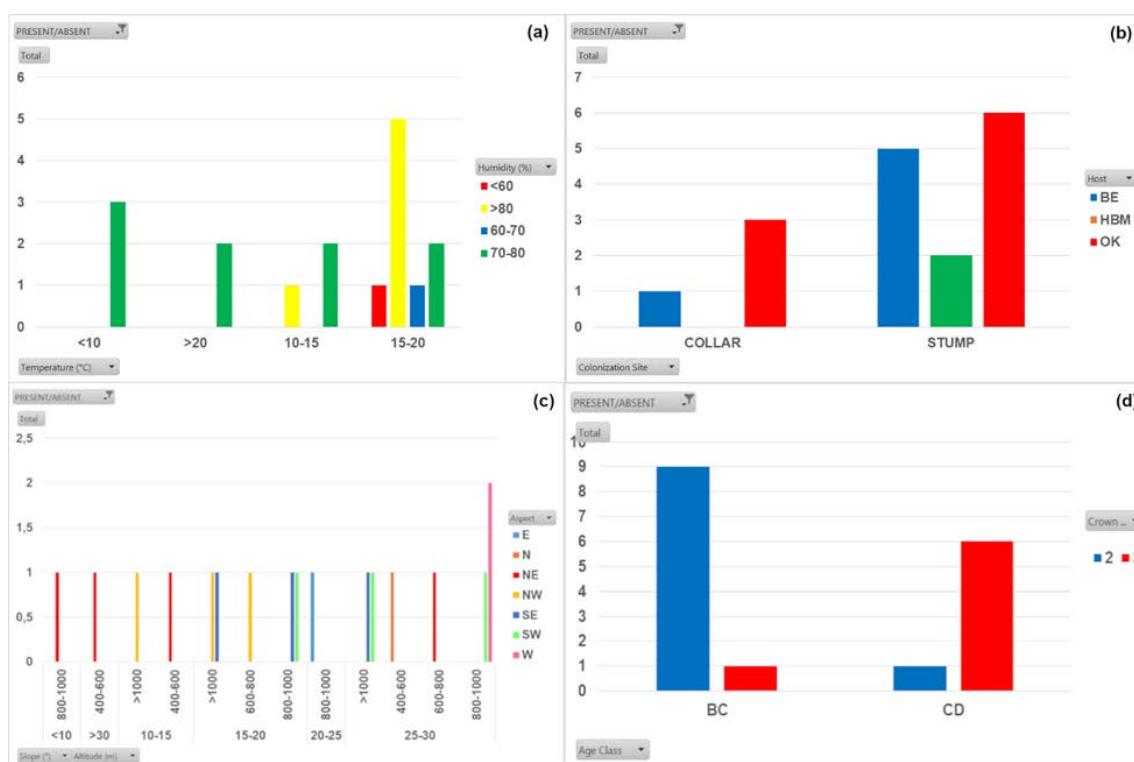


Figure 3. The effect of environmental variables on the occurrence of *Armillaria mellea* (a)Temperature-humidity, (b)colonization site–host preference, (c)altitude–slope–aspect, (d)age class–crown closure

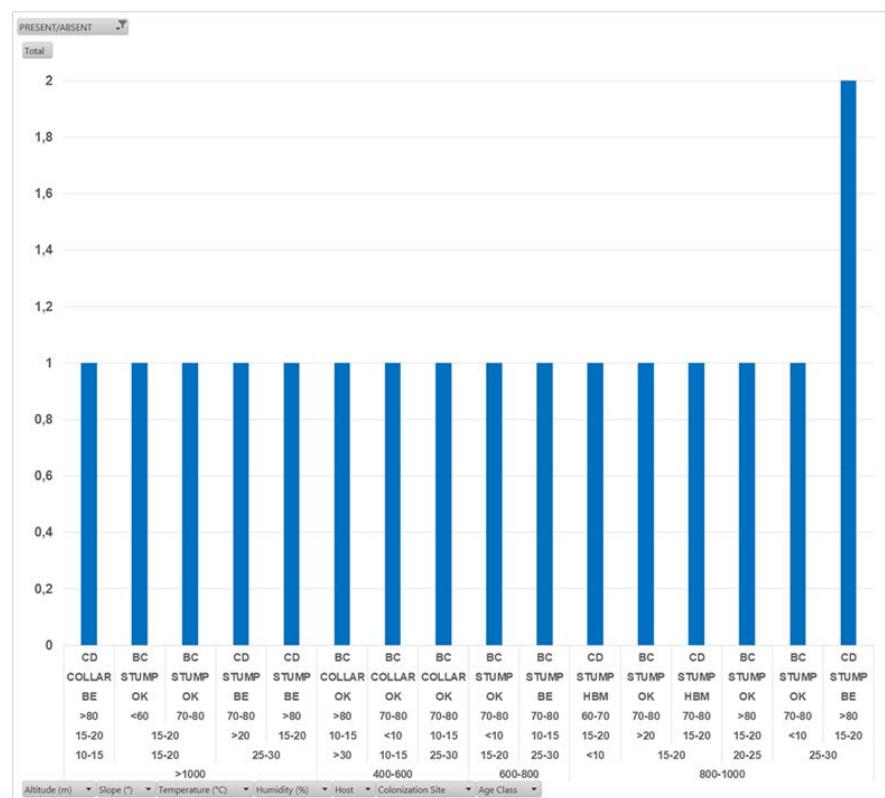
The presence/absence of *A. mellea* and environmental variables were tested with Chi-square analysis, and the results are given in Table 2. The temperature was different among these environmental variables, except for the host and colonization area. In addition, the graphical representations of these environmental variables were given in Figure 4. Environmental similarities and differences among plots where *A. mellea* was present and absence was determined by Hierarchical clustering analysis. In addition, a dendrogram showing the similarities of the plots was created for environmental variable groups in Figure 5.

The dendrogram consisted of 3 clusters including in order of 9, 12 and 9 plots from top to bottom and *A. mellea* was found on 6, 7, and 4 plots, respectively. To the dendrogram, *A. mellea* was mainly seen in the south of the study area and preferred western aspects. It is understood that this macrofungus prefers the south of the study area because of the altitude. In addition, *A. mellea* needed optimum conditions in terms of temperature and humidity; in other words, the probability of its occurrence decreases at high humidity and low temperature, as well as at low humidity and high temperature.

Table 2. Chi-square analysis of environmental variables, effect on the occurrence of *Armillaria mellea*.

Environmental Variables	df	χ^2	p-Value	Cramer's V	Effect Size
Slope (°)	5	4.830	0.348>0.05 NS	0.401	Relatively Strong
Altitude (m)	4	4.457	0.437>0.05 NS	0.385	Moderate
Temperature (°C)	1	5.954	0.015<0.05 *	0.454	Relatively Strong
Humidity (%)	3	3.180	0.365>0.05 NS	0.326	Moderate
Aspect	7	8.824	0.366>0.05 NS	0.542	Relatively Strong
Age Class	1	2.334	0.127>0.05 NS	0.279	Moderate
Crown Closure	1	2.334	0.127>0.05 NS	0.279	Moderate
Host	4	10.168	0.038<0.05 *	0.550	Relatively Strong
Colonization Site	3	22.961	0.000<0.001 ***	0.875	Very Strong

NS: Non-significant, *correlation is significant at the 0.05 level

Figure 4. The relationship between environmental variables and *Armillaria mellea* in Yuvacık dam watershed.

OK: Oak, HBM: Hornbeam, BE: Beech, a (0–8 cm), b (9–20 cm), c (21–36 cm), d (36–52 cm)
Diameter at Breast Height (Age class); 1 (11–4 %), 2 (41–70%), 3 (71–100%) Crown closure

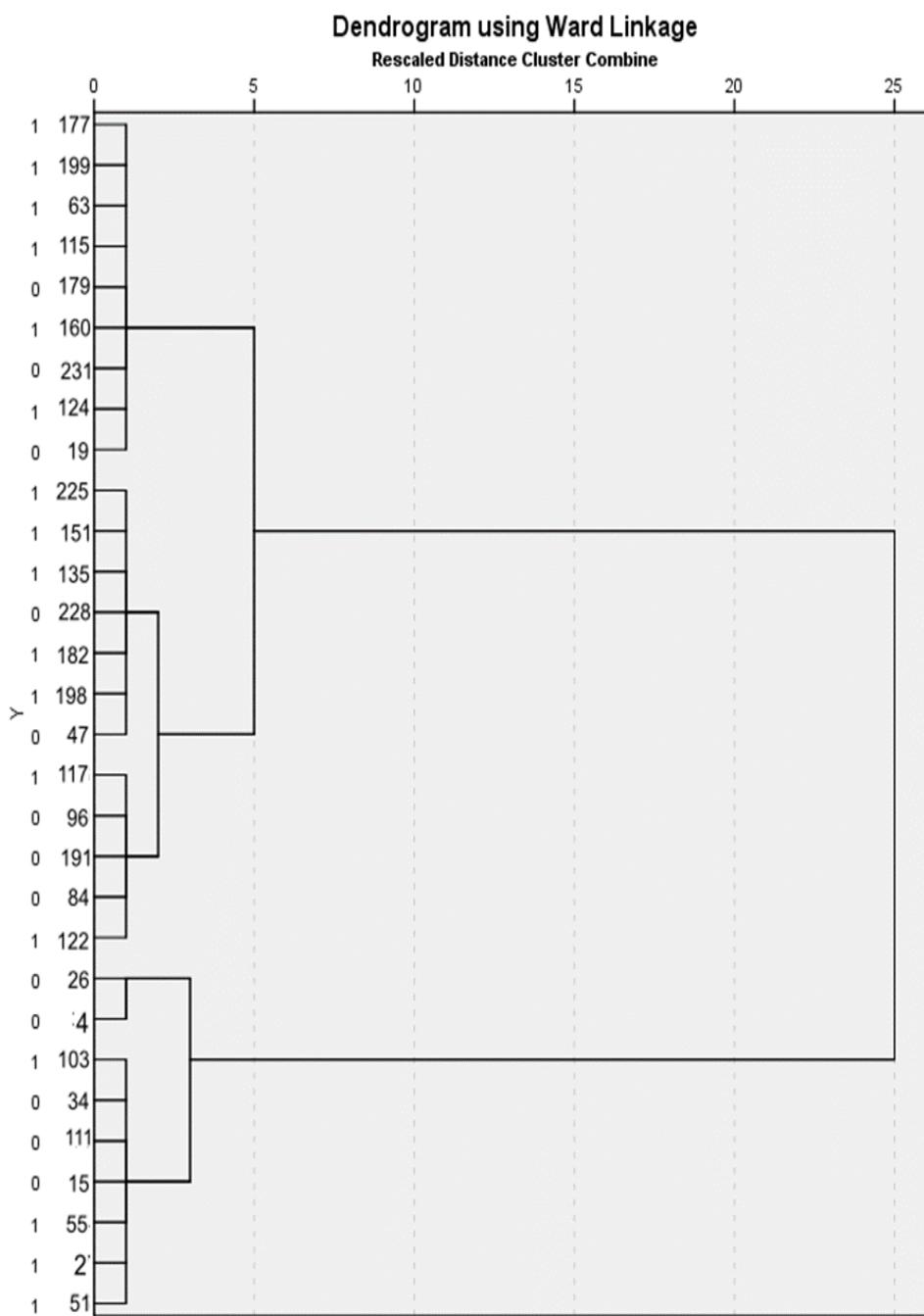


Figure 5. The clustering of the plots according to the presence/absence of *Armillaria mellea* for environmental variable groups in the study area.

It was investigated whether there is an effect of temperature (°C), humidity (%), altitude (meters), age class and crown closure on the presence of sporocarps of *A. mellea* in the plots. To create a new model using the Binary Logistic and Backward LR method, the age class, crown closure and altitude variables were removed from the model at the last step (Step 5). In terms of significance levels, positive advantages and Exp (B) values, the temperature (°C) and humidity (%) variables remained in the model, and the following statistically significant model was created.

$$z = -9.508 + 0.307 \times \text{temperature} \\ + 0.081 \times \text{humidity}$$

In step 5 of Table 3, we found that the temperature affects the occurrence of sporocarps of *A. mellea* 1.36 times (36%). In other words, when the temperature is altered by 1 unit, sporocarps' probability of occurrence is affected by 36% ($e^{0.307}$). Furthermore, the humidity alteration affects



the presence of sporocarps of *A. mellea* by 8.4% ($e^{0.081}$). In the new model, when the ratio of the relevant independent variables is included in the model, the probability ratio of

the dependent variables ($P(Y)$) is obtained. If $P(Y)$ is >0.5 , we estimate that *A. mellea* is found in the area.

Table 3. Logistic regression analysis and estimated logit coefficients, the effects of the environmental variables on the occurrence of *Armillaria mellea* sporocarps.

Variables in the Equation							
Steps- Variable		Estimated Logit Coefficient	Standard Error	Wald	df	Significance Level	Exp(B)/Odds Ratio
Step 1 ^a	altitude	-.001	.002	.352	1	.553	.999
	slope	-.084	.071	1.388	1	.239	.919
	temperature	.382	.164	5.413	1	.020	1.465
	humidity	.121	.069	3.060	1	.080	1.128
	age class	.830	1.980	.176	1	.675	2.294
	crown closure	-1.153	1.725	.447	1	.504	.316
	constant	-10.140	6.273	2.613	1	.106	.000
Step 2 ^a	altitude	-.002	.002	.411	1	.521	.998
	slope	-.072	.064	1.295	1	.255	.930
	temperature	.358	.149	5.765	1	.016	1.430
	humidity	.116	.067	3.037	1	.081	1.124
	crown closure	-.646	1.220	.280	1	.597	.524
	constant	-9.459	5.804	2.656	1	.103	.000
Step 3 ^a	altitude	-.001	.002	.243	1	.622	.999
	slope	-.073	.063	1.318	1	.251	.930
	temperature	.366	.146	6.320	1	.012	1.442
	humidity	.120	.066	3.331	1	.068	1.127
	constant	-10.635	5.429	3.837	1	.050	.000
Step 4 ^a	slope	-.060	.058	1.076	1	.300	.942
	temperature	.344	.138	6.192	1	.013	1.410
	humidity	.119	.066	3.264	1	.071	1.127
	constant	-11.478	5.285	4.716	1	.030	.000
Step 5 ^a	temperature	.307	.121	6.429	1	.011	1.360
	humidity	.081	.051	2.511	1	.113	1.084
	Constant	-9.508	4.570	4.329	1	.037	.000

a. Variable(s) entered on step 1: altitude. slope. temperature. humidity. age. crown.

Discussions

To chi-square results, the presence of *A. mellea* is independent of the slope, altitude, humidity and aspect, age class and crown closure variables, but it is dependent on temperature, host and colony location variables. *A. mellea* was specified mostly at an altitude of >800 m while it was not seen above 1200 m in the present study. This finding is consistent with previous studies conducted in

Europe (Guillaumin et al., 1993; Tsopelas, 1999; Keča et al., 2009; Lushaj et al., 2010; Mesanza et al., 2017).

In this survey, all of the sporocarps were found on broad-leaved trees. It was recorded on deciduous trees by previous studies in the Netherlands, Greece, Czech Republic, Serbia, Albania, and Spain (Termorshuizen and Arnolds, 1994; Tsopelas, 1999; Jankovský, 2003; Keča et al., 2009; Lushaj et al., 2010; Mesanza et al., 2017).



In our study, sporocarps were mostly found on the root collars of living trees of oak and beech, and stumps in the western aspect. Mesanza et al. (2017) observed *Armillaria* species mostly in the western aspect of Spain. Also, they mainly found sporocarps on the root collars of dead and living trees and stumps. Our determinations showed that *A. mellea* grew up in the "bc" age class and "2" crown closure. Dálya et al. (2019) evaluated the presence and distribution of root rot fungi in the Vallombrosa (Italy) forest after severe storm damage. They noted that *A. mellea* grew between 20–60, according to the age classes of forest stands. Our findings are similar to the study results of Dálya et al. (2019).

A. mellea is widely spread in the presence of an Atlantic climate as in England, part of France and the Mediterranean climate in Italy, Portugal and Spain (Guillaumin et al., 1993). In our study, we found out that the optimum temperature was 15–20 °C and humidity >80 humidity (%) for the growth of the sporocarps of the species. Guillaumin et al., (1993) reported that *A. mellea* is a thermophilic species of Atlantico-Mediterranean distribution. Keča (2005) reported that the fastest growth was at 22 °C in his study of mycelium development on five *Armillaria* species, including *A. mellea*. Hasegawa et al., (2011) showed that *A. mellea* was thermophilic according to Kira's warmth index (WI) and indicated a very similar thermal preference between Europe and Japan.

In addition, the dendrogram showing the similarities of the plots was created for environmental variable groups by hierarchical clustering analysis. We also provided the knowledge that the occurrence of *A. mellea* decreased both at high humidity and low temperature or low humidity and high temperature. In other words, the sporocarps of *A. mellea* prefer optimum conditions of temperature and humidity.

The results of logistic regression analysis revealed that temperature affects the growth of the sporocarps of *A. mellea* by 36%. When the temperature varies by 1 degree, the growth of the sporocarps is affected 1.36 times. This rate is 8.4% for humidity. This logistic regression equation can be used to predict *Armillaria*'s prevalence in this study area.

Armillaria has not been reported as a major epidemic or disease agent for the study area so far. However, there are significant losses in the world, especially in terms of oak species. It is required to observe *A. mellea*, which can be a substantial threat to deciduous and coniferous species. Our results show that

the sporocarps of *A. mellea* grow on stumps by 80%. This detection indicates that the species is currently saprophytic in the area and reveals that treatment of stumps should be attentive to avoid an epidemic caused by *A. mellea* in the future.

A. mellea is the most virulent species of broad-leaved forests and has caused significant economic losses in western Europe. However, *Armillaria* species are thought to be beneficial for the carbon cycle in natural ecosystems as a saprophytic, especially white-rot. It is also used in the food and pharmaceutical industry. It is a requirement to realize regular surveys in the field for a species that is considered to have such different effects. There is little knowledge of *Armillaria*'s diversity, distribution, pathogenicity, pandemic, and virulence in the forest of Türkiye. The species diversity of the *Armillaria* complex should be investigated and supported by molecular methods. It is required to give importance to the pathogenicity, virulence, and pandemic risk analysis studies of *Armillaria* species in Türkiye forests. Since the incidence of *A. mellea* is higher in the "bc" age class and "2" crown closures in the study area, it is revealed that more attention should be paid to sites with this feature in terms of epidemics in forestry studies. In addition, our findings support that this fungus commonly affects younger individuals. Studies on assessment as a non-wood forest product (in the food and pharmaceutical industry) maybe provide significant knowledge of the environmental requirements of these species.

Author contributions

Sabiha Acer coordinated the research, planned and carried out the identifications and discussed the results. Ersel Yilmaz developed the theory and analytical methods, conducted the field experiment, and run the data analysis. Ayhan Karakaya supervised the work, designed and conducted the field experiment, and discussed the results. All authors helped in writing, read, and approved the final manuscript.

Conflicts of interest

The authors declare no competing interests.

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Domestication of Wild Edible Mushrooms in Eastern Africa: A Review of Research Advances and Future Prospects

Susan Njuguini KABACIA^{1*}, Mary Nyawira MUCHANE²

*Corresponding author: susannjuguini@gmail.com

¹ National Museums of Kenya, Nairobi, Kenya / susannjuguini@gmail.com

² Nairobi, Kenya / mmurethi@yahoo.com

Abstract: Mushroom farming is an emerging industry in Africa with great potential to improve livelihoods. However, the industry is still dominated by a few exotic varieties faced with challenges of regional adaptability. Also, information concerning their domestication status, challenges and future prospects is scattered and unclear. The purpose of this paper was to review and detail the diversity of mushrooms occurring in Eastern Africa (EA), reveal the wild edible/medicinal species (WEMs) that have already been domesticated, their domestication status, the nutritional composition status of the edible species and availability of their germplasm (mother cultures). To achieve this, a detailed review of published research articles, books and reports from EA was conducted. Data was collected from articles focusing on the diversity of WEMs, nutritional composition analysis and domestication methods and status. From the review, 306 WEMs are shown to have desirable characteristics for utilization as food /medicine, with Tanzania documenting the highest number (147) followed by Malawi (90). Among these, 82 species are edible ectomycorrhizal species with great potential to support the mushroom industry if sustainably harvested and managed. The rest are saprophytic fungi species. Only 14 species among the saprobe group have been tissue cultured, tested for spawn production and cultivation. 51 species have been analyzed for nutritional composition. However, none of these have been commercially introduced to cultivation and the availability of their germplasm for research and propagation purpose is uncertain. The result from this study clearly shows research on the domestication of WEMs in Africa is still in its infancy stages.

Key words: Cultivation, Distribution, Species, Substrates, Symbiosis

Doğu Afrika'da Yenilebilir Yabani Mantarların Kültüre Alınması: Araştırmadaki İlerlemeler ve Gelecekteki Beklentiler Üzerine Bir İnceleme

Öz: Mantar yetiştirciliği, geçim kaynaklarını iyileştirme konusunda büyük potansiyele sahip, Afrika'da gelişmekte olan bir endüstridir. Bununla birlikte, sektörde hala bölgesel uyum sağlama zorlukları karşı karşıya kalan birkaç egzotik tür hakimdir. Ayrıca, mantarların kültüre alma durumları, karşılaşlıklarla zorluklar ve gelecek beklenimleri ile ilgili bilgiler dağıtık ve belirsizdir. Bu makalının amacı, Doğu Afrika'da bulunan mantarların çeşitliliğini incelemek ve detaylandırmak, halihazırda kültüre alınmış olan yabani yenilebilir/tıbbi mantar türlerinin (WEM'ler), kültüre alma durumlarını, yenilebilir türlerin beslenme kompozisyon durumunu ve anaç kültürlerinin (germplazmlarının) mevcudiyetini ortaya çıkarmaktır. Bunu başarmak için, yayınlanan araştırma makaleleri, kitaplar ve Doğu Afrika raporlarının ayrıntılı bir incelemesi yapılmıştır. Veriler, yabani yenilebilir/tıbbi türlerin çeşitliliğine, besin bileşimi analizine ve kültüre alma yöntemlerine ve statüsüne odaklanan makalelerden toplanmıştır. Araştırmada 306 yabani yenilebilir/tıbbi türün gıda/ilac olarak kullanım için arzu edilen özelliklere sahip olduğu gösterilmiştir, en yüksek sayımı ortaya koyan Tanzanya (147) ve onu Malavi (90) takip etmektedir. Bunların arasında 82 tür, sürdürülebilir bir şekilde hasat edilir ve yönetilirse mantar endüstrisini destekleme potansiyeli yüksek, yenilebilir ektomikorizal türlerdir. Geri kalanı saprofitik mantar türleridir. Saprofitik grubundan sadece 14 türün doku kültürü yapılmış ve tohumlu misel üretimi ve yetiştirmeye için test edilmiştir. 51 tür besin bileşimi açısından analiz edilmiştir. Bununla birlikte,



bunların hiçbirini ticari olarak kültüre alınmaya çalışılmamıştır ve anaç kültürlerin araştırılması ve çoğaltılması amacıyla mevcut durum belirsizdir. Bu çalışmadan elde edilen sonuç, Afrika'da yabani yenilebilir/tıbbi türlerin kültüre alınmasına ilişkin araştırmaların henüz başlangıç aşamasında olduğunu açıkça göstermektedir.

Anahtar kelimeler: Kültür mantarcılığı, Dağılım, Türler, Substratlar, Simbiyoz

Introduction

Wild edible mushrooms are valuable non-wood forest resources, cherished by many communities in the world (Boa, 2004; Dejene et al., 2017a; Degreef et al., 2016; Degreef et al., 2020; Ngom et al., 2022). In Africa, wild mushroom gathering and hunting is a common practice mostly among the rural populace, especially during the rainy season (Dejene et al., 2017b; Kashiki et al., 2021; Ngom et al., 2022), which is suitable fructification of fungi. They are the source of food, health promoting properties and economic value (Boa, 2004; Dejene et al., 2017a; Sande, 2019; Niazi and Ghafoor, 2021). Ethnomycological knowledge obtained from different parts of the world reveals the use of mushrooms in various ways including in traditional medicine (Atri and Chadha, 2018). In Tanzania for example, the wild mushroom collection is a socio-economic activity among the Hehe and Benna communities. It is estimated that the collectors earn approximately US \$500 to 1000 for 750 - 1500 kg of mushrooms per season (Atri and Chadha, 2018). In other regions of Africa, wild edible mushrooms are collected to serve various nutritional needs, especially protein which in most cases is lacking or it is expensive (Atri and Chadha, 2018). Among the most cherished mushrooms in Africa are *Termitomyces* which are preferred due to their unique taste, flavor and texture (Degreef et al., 2020). Despite the role mushrooms play to improve livelihoods and the well being of many communities in Africa, there is increasing unsustainable harvesting which raises concerns about their depletion from the wild which will consequently increase food insecurity among local communities in the near future (Waiganjo et al., 2008; Dejene et al., 2017a; Degreef et al., 2020). To protect the planet and its biodiversity while reducing food insecurity and maintaining peace and human prosperity in line with the 2015 Sustainable Development Goals (SDGs), local mushroom cultivation and expansion of the mushroom industry have been proposed as a viable way of sustaining mushroom supply and protecting WEMs in nature (Niazi and Ghafoor, 2021).

Mushroom cultivation has increased tremendously in many parts of the world. It is carried out in more than 100 countries with a growth rate of 6-7% annual production (Niazi and Ghafoor, 2021). China leads with over 80% in the production of edible and medicinal mushrooms. The mushroom industry is the second largest in China and has created over 25 million job opportunities (Willis, 2018). However, mushroom cultivation is low in Africa and the region contributes less than 1% of the annual worldwide production of mushrooms (Adejumo and Awosanya, 2005; Onyango et al., 2011; Dejene et al., 2017a). This is majorly attributed to over-reliance on wild harvesting and low uptake of mushroom cultivation when compared to the cultivation of other food crops such as maize and beans. Partly, it is dependent on imported exotic mushroom mother cultures which tend to be expensive and sometimes difficult to access making mushroom farming expensive and unattractive. In East Africa, only cultivated mushrooms of exotic origin (*Agaricus*, *Pleurotus*, *Lentinula* and *Ganoderma* sp) cultured from imported mother cultures (Musieba et al., 2012) dominates the mushroom industry. These imported strains are associated with low yields and susceptibility to pests and diseases, the assumption being that the species lack regional adaptability (Otieno et al., 2015). Over the past years, there has been increased interest to exploit and domesticate wild edible and medicinal saprophytic mushrooms with desirable characteristics such as good regional adaptability, ability to grow on readily available substrates, low susceptibility to pests and diseases and good flavor for sustainable nutritional source in Eastern Africa (EA) (Mdachi et al., 2004; Magingo et al., 2004; Oriyo et al., 2004; Mshadete and Cuff, 2008; Mwita et al., 2010; Musieba et al., 2012; Juma et al., 2015; Wendiro, Wacoo and Wise, 2019).

Eastern Africa is a tropical region with 10 countries (Tanzania, Kenya, Uganda, Rwanda, Burundi, South Sudan, Djibouti, Eritrea, Ethiopia, and Somalia). The region is known for its unique forested ecosystems hosting valuable mycodiversity (Pegler and Rayner, 1969; Pegler, 1977; Boa, 2004). Although research on the diversity and taxonomy of mushroom resources in this



region is increasing tremendously with new species being reported (Boa, 2004; Tibuhwa, 2011; Njuguini et al., 2018; Nteziryayo et al., 2019; Muchane et al., 2021; Ngom et al., 2022) information concerning domestication of these wild edible varieties is still scanty. Unlike many parts of the world where more than 100 species have been domesticated and 60% commercialized (Chang and Miles, 2004; Dejene et al., 2017a; Willis, 2018) in EA available information points only to the successful cultivation trials, but reports of adopted species by farmers for cultivation is lacking (Magindo et al., 2004, Mshadete and Cuff, 2008, Musieba et al., 2012, Juma et al., 2015). Additionally, the diversity of WEMs in EA is also poorly investigated. The potential of WEMs contribution to the mushroom industry remains less known and explored, with information concerning their domestication status, challenges and future prospects still scattered and not clear. There is still no clear distinction between edible and medical mushrooms.

Domestication of WEMs in EA provides a great opportunity of producing mushroom strains adapted to EA environmental and climate conditions as well as strains less susceptible to diseases. Since most of these WEMs are adapted to high temperatures and humid conditions, their production can be faster compared to exotic strains from temperate regions (Musieba et al., 2012). Optimization and simulation of growth conditions to match that of natural habitats is also possible since habitats and growth conditions of domesticated strains are known and accessible. Such information will enable spawn producers and mushroom farmers to use only strains ideal for conditions in the growing facility and use of right substrate. Domestication of WEMs will also increase the production of traditionally preferred and high-nutritional indigenous mushrooms. This will increase levels of mushroom farming adoption and production. This will also minimize losses, increase value for money, and lower the cost of production thus expanding the mushroom industry

in EA. Although many of the common edible species have therapeutic properties and are used for medical purposes, information regarding their unique medicinal properties is also required for proper characterization.

The purpose of this paper is to review and detail the diversity of WEMs with special reference to EA, its potential in the mushroom industry, domestication status, the nutritional value in comparison to introduced exotic species and availability of their germ-plasm (mother cultures).

Material and Method

Literature was obtained through Internet-based scientific literature search engines (ISI web of Science, Research gate and Google scholar) and Libraries. Research materials that fulfilled the following criteria were selected; research materials and articles in eight selected countries focusing on the diversity of WEMs, nutritional composition analysis, articles detailing domestication, experimental methods and status. Data collected included; wild edible mushroom species (WEMs), nutrient content of WEMs, domesticated mushroom species, status of domestication (tissue culture, spawn or cultivated). Methods & materials used during domestication, and the yield of cultivated mushrooms was considered. Data were extracted from the results section and appendices with the list of WEMs, tables of means, graphs, and figures. Graphs and tables were used to summarize WEMs species in different countries.

Results

Diversity of edible (medicinal) mushrooms

This study has revealed a diverse community of wild edible mushroom (WEMs) species in East Africa with a total of 306 species of WEMs distributed within the division *Basidiomycota* (82 genera within 39 families) and *Ascomycota* (4 genera within 2 Phylum) (Table 1).

Table 1: list of wild edible / Medicinal mushrooms in East Africa Region

Species	Habitat association	Country	Reference
<i>Afroboletus luteolus</i> (Heinem.) Pegler & T.W.K. Young	Ectomycorrhizal	Tanzania, Burundi, Malawi	Harkonen et al., 2003; Boa, 2004; Tibuhwa, 2013; Degreef et al., 2016
<i>Afroboletus costatisporus</i> (Beeli) Watling.	Ectomycorrhizal	Malawi	Boa, 2004
<i>Afrocantharellus fistulosus</i> Tibuhwa & Buyck	Ectomycorrhizal	Tanzania	Tibuhwa, 2013
<i>Afrocantharellus splendens</i> Buyck	Ectomycorrhizal	Tanzania	Tibuhwa, 2013



<i>Afrocantharellus symoensi</i> (Heinemann)	Ectomycorrhizal	Tanzania	Tibuhwa, 2013
<i>Agaricus campestris</i> (L. ex Fr.)	Saprobe	Tanzania, Somalia, Ethiopia	Pegler, 1977; Rammeloo and Wallyne, 1993; Boa, 2004; Mdachi <i>et al.</i> , 2004; Tibuhwa, 2011; Woldegiorgis, 2015; Dejene <i>et al.</i> , 2017a; Dejene <i>et al.</i> , 2017b ; Njugini <i>et al.</i> , 2018
<i>Agaricus campestroides</i> (Heinem & Gooss. - Font)	Saprobe	Ethiopia	Dejene <i>et al.</i> , 2017b
<i>Agaricus amboensis</i> (Fayod) Sacc.	Saprobe	Somalia, Uganda	Boa, 2004
<i>Agaricus arvensis</i> Schaeff.	Saprobe	Ethiopia, Tanzania, Uganda	Harkonen <i>et al.</i> , 2003; Degreef <i>et al.</i> , 2016; Dejene <i>et al.</i> , 2017a; Dejene <i>et al.</i> , 2017b; Ngom <i>et al.</i> , 2022
<i>Agaricus augustus</i> Fr.	Saprobe	Kenya, Tanzania	Pegler, 1977; Tibuhwa, 2011; Njuguni <i>et al.</i> , 2018
<i>Agaricus bimbaumii</i> (Corda) Singer	Saprobe	Tanzania	Tibuhwa, 2013
<i>Agaricus bingensis</i> Heinem.	Saprobe	Malawi, Uganda	Rammeloo and Wallyne 1993; Boa, 2004
<i>Agaricus bisporus</i> (Lange) Imbach	Saprobe	Tanzania, Ethiopia, Kenya, Malawi	Rammeloo and Wallyne, 1993; Tibuhwa, 2011; Tibuhwa, 2013; Woldegiorgis, 2015
<i>Agaricus croceolutescens</i> (Heinemann & Gooss.-Font)	Saprobe	Malawi	Rammeloo and Wallyne, 1993; Boa, 2004
<i>Agaricus placomyces</i> Perk.	Saprobe	Tanzania	Tibuhwa 2013
<i>Agaricus silvaticus</i> Schaeff.	Saprobe	Kenya	Pegler, 1977; Njuguni <i>et al.</i> , 2018
<i>Agaricus</i> sp. L.: Fr. Emend. Karst	Saprobe	Tanzania, Malawi	Nakalembe <i>et al.</i> , 2009
<i>Agaricus subedulis</i> Heinem.	Saprobe	Ethiopia	Dejene <i>et al.</i> , 2017a.
<i>Agaricus sylvicola</i> (Vitt.) Lév.	Saprobe	Rwanda	Degreeef <i>et al.</i> , 2016
<i>Agaricus volvatus</i> Heinem. & Gooss.	Saprobe	Kenya	Tibuhwa 2011; Njuguni <i>et al.</i> , 2018
<i>Agrocybe pediades</i> (Fr.) Fayod	Saprobe	Ethiopia	Dejene <i>et al.</i> , 2017a
<i>Amanita aff. calopus</i> (Beeli)	Ectomycorrhizal	Malawi	Rammeloo and Wallyne, 1993; Boa, 2004
<i>Amanita aurea</i> (Beeli) E.J. Gilbert	Ectomycorrhizal	Tanzania	Harkonen <i>et al.</i> , 2003
<i>Amanita bingensis</i> (Beeli) Heinem.	Ectomycorrhizal	Malawi	Rammeloo and Wallyne, 1993; Boa, 2004
<i>Amanita cf. robusta</i> (Beeli)	Ectomycorrhizal	Malawi	Rammeloo and Wallyne 1993
<i>Amanita flammeola</i> (Pegler & Pearce)	Ectomycorrhizal	Malawi	Rammeloo and Wallyne, 1993; Boa 2004
<i>Amanita fulva</i> (Schaeff) Fr.	Ectomycorrhizal	Malawi	Boa, 2004; Rammeloo and Wallyne, 1993
<i>Amanita goossensiae</i> (Beeli)	Ectomycorrhizal	Malawi	Boa, 2004; Rammeloo and Wallyne, 1993
<i>Amanita hemibapha</i> (Berk. & Br.)	Ectomycorrhizal	Malawi	Boa, 2004; Rammeloo and Wallyne 1993
<i>Amanita loosii</i> Beeli	Ectomycorrhizal	Burundi, Tanzania	Buyck and Nzigidahera, 1995; Harkonen <i>et al.</i> , 2003; Boa, 2004; Tibuhwa, 2013; Degreeef <i>et al.</i> , 2016
<i>Amanita mafingensis</i> Härk. & Saarim.	Ectomycorrhizal	Burundi	Harkonen <i>et al.</i> , 2003; Tibuhwa, 2013; Degreeef <i>et al.</i> , 2016



<i>Amanita masasiensis</i>	Ectomycorrhizal	Tanzania	Harkonen et al., 2003; Tibuhwa, 2013
Hark. & Saarim.			
<i>Amanita Pers.</i>	Ectomycorrhizal	Burundi	Buyck and Nzigidahera, 1995; Dejene et al., 2017a
<i>Amanita pudica</i> (Beeli) Walleyn	Ectomycorrhizal	Burundi	Degreeef et al., 2016
<i>Amanita rhodophylla</i> (Beeli)	Ectomycorrhizal	Malawi	Rammeloo and Wallyne, 1993; Boa, 2004
<i>Amanita robusta</i> (Beeli)	Ectomycorrhizal	Malawi	Boa, 2004
<i>Amanita rubescens</i> (Pers.: Fr.)	Ectomycorrhizal	Burundi, Malawi	Rammeloo and Wallyne, 1993; Buyck and Nzigidahera, 1995; Boa, 2004; Degreeef et al., 2016
<i>Amanita tanzanica</i> Hark. Saarim.	Ectomycorrhizal	Tanzania, Burundi	Harkonen et al., 2003; Tibuhwa, 2013; Degreeef et al., 2016
<i>Amanita vaginata</i> (Bull.: Fr)	Ectomycorrhizal	Malawi	Rammeloo and Wallyne, 1993; Boa, 2004
<i>Amanita verna</i> (Bull.) Lam.	Ectomycorrhizal	Burundi	Nteziryayo, et al., 2019
<i>Amanita zambiana</i> (Pegler & Pierce)	Ectomycorrhizal	Kenya, Burundi	Rammeloo and Wallyne, 1993; Boa, 2004; Wandati et al., 2013; Nteziryayo et al., 2019
<i>Amylosporus IJ-2014</i>	Saprobe	Tanzania	Hussein et al., 2016
<i>Armillaria borealis</i> Marxmüller & Korhonen	Saprobe	Rwanda	Degreeef et al., 2016
<i>Armillaria cepistipes</i> Velen.	Saprobe	Rwanda	Degreeef et al., 2016
<i>Armillaria heimii</i> Pegler	Saprobe	Tanzania, Ethiopia	Tibuhwa, 2013; Degreeef et al., 2016; Dejene et al., 2017a
<i>Armillaria lutea</i> Gillet	Saprobe		Degreeef et al., 2016
<i>Armillaria mellea</i> (Vahl) P. Kumm.	Saprobe	Kenya, Uganda	Pegler, 1977; Harkonen et al., 2003; Boa, 2004; Tibuhwa, 2013; Njuguini et al., 2018
<i>Armillaria ostoyae</i> (Romagn.) Herink	Saprobe	Rwanda	Degreeef et al., 2016
<i>Armillaria tabescens</i> (Scop.) Emel	Saprobe	Rwanda	Degreeef et al., 2016
<i>Auricularia auricula - judae</i> (Bull.: Fr.) Wettst.	Saprobe	Kenya, Malawi, Rwanda, Uganda	Rammeloo and Wallyne, 1993; Boa, 2004; Onyango et al., 2012; Degreeef et al., 2016; Hussein et al., 2016; Njuguini et al., 2018
<i>Auricularia Bull.ex</i> Juss	Saprobe	Ethiopia; Kenya	Wandati et al., 2014; Dejene et al., 2017; Njuguini et al., 2018
<i>Auricularia cornea</i> Ehrenb.	Saprobe	Tanzania, Burundi, Rwanda	Tibuhwa et al., 2013; Degreeef et al., 2016
<i>Auricularia delicata</i> (Mont. ex Fr.) Henn.	Saprobe	Kenya, Tanzania, Uganda	Rammeloo and Wallyne, 1993; Harkonen et al., 2003; Boa, 2004; Tibuhwa, 2011; Tibuhwa, 2013; Degreeef et al., 2016; Njuguini et al., 2018; Ngom , et al., 2022
<i>Auricularia fuscosuccinea</i> (Mont.) Henn.	Saprobe	Malawi	
<i>Auricularia fuscosuccinea</i> (Mont.) Henn.	Saprobe	Tanzania	Boa, 2004
<i>Auricularia polytricha</i> (Mont.) Sacc.	Saprobe	Tanzania, Kenya	Harkonen et al., 2003; Boa, 2004; Tibuhwa, 2011; Hussein et al., 2016; Njuguini et al., 2018;
<i>Bolbitius vitellinus</i> (Pers.:Fr.)	Ectomycorrhizal	Malawi	Rammeloo and Wallyne, 1993
<i>Boletus clavipes</i> (Perk) Pilat & Dermek	Ectomycorrhizal	Tanzania	Mdachi et al., 2004
<i>Boletus loosii</i> Heinem.	Ectomycorrhizal	Burundi	Degreeef et al., 2016
<i>Boletus pallidissimus</i> Watling	Ectomycorrhizal	Tanzania	Tibuhwa, 2013
<i>Boletus pruinatus/</i> <i>xerocomellus</i>	Ectomycorrhizal	Tanzania	Mdachi et al., 2004



<i>pruinatus</i> Fr. & Hok Sutara			
<i>Boletus spectabilissimus</i> Watling	Ectomycorrhizal	Tanzania	Harkonen et al., 2003; Tibuhwa, 2013
Buntarantara	Saprobe	Uganda	Nakalembe et al., 2009
Buntarantara	Saprobe	Uganda	Nakalembe et al., 2003
Bunyabikandaigo	Saprobe	Ethiopia	Dejene et al., 2017a
<i>Calvatia rubroflava</i> (Cragin) Lloyd	Saprobe	Malawi	Rammeloo and Wallyne, 1993
<i>Calvatia utiliformis</i> (Bull.:Pers)	Ectomycorrhizal	Burundi	Boa, 2004
<i>Cantharellus rufopunctatus</i> (Beeli) Heinem.	Ectomycorrhizal	Malawi, Tanzania	Harkonen et al., 2003; Boa, 2004; Mdachi et al., 2004; Tibuhwa, 2013; Gateri et al., 2014;
<i>Cantharellus cibarius</i> Fr.	Ectomycorrhizal	Malawi	Rammeloo and Wallyne, 1993; Boa, 2004
<i>Cantharellus cibarius</i> var. <i>defibulatus</i> Heinem.	Ectomycorrhizal	Burundi	Boa, 2004
<i>Cantharellus congolensis</i> Beeli	Ectomycorrhizal	Burundi, Tanzania, Malwi	Rammeloo and Wallyne, 1993; Buyck and Nzigidahera, 1995; Harkonen et al., 2003; Boa, 2004; Degreef et al., 2016
<i>Cantharellus cyanescens</i> Buyck.	Ectomycorrhizal	Burundi	Buyck and Nzigidahera, 1995; Boa, 2004
<i>Cantharellus cyanoxanthus</i> R. Heim ex Heinem.	Ectomycorrhizal	Burundi, Tanzania	Buyck and Nzigidahera, 1995; Tibuhwa, 2013; Degreef et al., 2016
<i>Cantharellus defibulatus</i> (Heinem.) Eyssart. & Buyck	Ectomycorrhizal	Burundi	Degreeef et al., 2016
<i>Cantharellus densifolius</i> Heinem.	Ectomycorrhizal	Burundi, Tanzania, Malawi	Rammeloo and Wallyne, 1993; Buyck and Nzigidahera 1995; Boa, 2004; Tibuhwa, 2013; Degreef et al., 2016
<i>Cantharellus floridula</i> (Pegler)	Ectomycorrhizal	Tanzania	Harkonen et al., 2004; Tibuhwa, 2013
<i>Cantharellus Fr. var. latifolius</i> Heinem.	Ectomycorrhizal	Burundi	Buyck and Nzigidahera, 1995
<i>Cantharellus isabellinus</i> Heinem.	Ectomycorrhizal	Tanzania	Harkonen et al., 2003; Boa, 2004; Tibuhwa, 2013
<i>Cantharellus longisporus</i> Heinem.	Ectomycorrhizal	Malawi	Rammeloo and Wallyne, 1993; Boa, 2004
<i>Cantharellus luteopunctatus</i> (Beeli) Heinem.	Ectomycorrhizal	Tanzania	Tibuhwa, 2013
<i>Cantharellus miomboensis</i> Buyck & V. Hofst.	Ectomycorrhizal		Degreeef et al., 2016
<i>Cantharellus parvisporus</i> (Eyssart. & Buyck)	Ectomycorrhizal	Tanzania	Tibuhwa, 2013
<i>Cantharellus platyphyllus</i> Heinem. var. <i>cyanescens</i> (Buyck) Eyssart. & Buyck	Ectomycorrhizal	Tanzania, Burundi	Harkonen et al., 2003; Boa, 2004; Degreef et al., 2016
<i>Cantharellus pseudocibarius</i> Henn.	Ectomycorrhizal	Burundi	Boa, 2004



<i>Cantharellus rhodophyllus</i> Heinem.	Ectomycorrhizal	Tanzania	Tibuhwa, 2013
<i>Cantharellus ruber</i> Heinem.	Ectomycorrhizal	Burundi, Tanzania	Buyck and Nzigidahera, 1995; Harkonen <i>et al.</i> , 2003; Boa, 2004; Tibuhwa, 2013; Degreef <i>et al.</i> , 2016
<i>Cantharellus rufopunctatus</i> (Beeli) Heneim.	Ectomycorrhizal	Burundi, Tanzania	Buyck and Nzigidahera, 1995; Tibuhwa, 2013
<i>Cantharellus splendens</i> Buyck	Ectomycorrhizal	Burundi	Buyck and Nzigidahera, 1995; Degreef <i>et al.</i> , 2016
<i>Cantharellus subincarnatus</i> Eyssart. & Buyck	Ectomycorrhizal	Tanzania	Tibuhwa, 2013
<i>Cantharellus symoensii</i> Heinem.	Ectomycorrhizal	Burundi , Tanzania	Buyck and Nzigidahera, 1995; Harkonen <i>et al.</i> , 2003; Boa, 2004; Degreef <i>et al.</i> , 2016
<i>Cantharellus tenuis</i> Heinem.	Ectomycorrhizal	Malawi	Rammeloo and Wallyne, 1993; Boa, 2004
<i>Cantharellus tomentosus</i> Eyssart. & Buyck	Ectomycorrhizal	Tanzania	Tibuhwa, 2013
<i>Clavaria albiramea</i> (Corner) Buyck & Duhem	Saprobe	Malawi	Boa, 2004
<i>Clavatia rubroflava</i> (Cragin) Lloyd	Saprobe	Ethiopia	Dejene <i>et al.</i> , 2017a
<i>Clavulina wisoli</i> R.H. Petersen	Saprobe	Tanzania	Harkonen <i>et al.</i> , 2003
<i>Collybia aurea</i> (Beeli) Pegler	Saprobe	Burundi, Rwanda	Buyck and Nzigidahera 1995; Degreef <i>et al.</i> , 2016
<i>Collybia confluens</i> (Pers.) Kumm.	Saprobe	Malawi	Rammeloo and Wallyne, 1993; Boa, 2004
<i>Collybia dryophila</i> (Bull.) Kumm.	Saprobe	Malawi	Rammeloo and Wallyne, 1993; Boa, 2004
<i>Coprinellus domesticus</i> (Bolton)Vilgalys, Hopple & Jacq. Johnson	Saprobe	Ethiopia	Dejene <i>et al.</i> , 2017a
<i>Coprinellus niveus</i> Fr.	Saprobe	Ethiopia	Dejene <i>et al.</i> , 2017a
<i>Coprinus cf. molestus</i> Bouriquet	Saprobe	Burundi	Buyck and Nzigidahera, 1995
<i>Coprinus cinereus</i> (Schaeff.)	Saprobe	Tanzania	Harkonen <i>et al.</i> 2003; Boa, 2004, Mshandete and Cuff 2007 and 2008, Raymond <i>et al.</i> , 2012
<i>Coprinus comatus</i> (O.F Murill.) Pers	Saprobe	Malawi	Rammeloo and Wallyne, 1993; Tibuhwa, 2013
<i>Coprinus disseminatus</i> (Pers.) J. E Lange	Saprobe	Tanzania, Malawi, Uganda , Kenya	Pegler, 1977; Boa, 2004; Tibuhwa, 2013; Njuguini <i>et al.</i> 2018
<i>Coprinus micaceus</i> (Bull.) Fr.	Saprobe	Kenya , Tanzania	Pegler 1977; Njuguini <i>et al.</i> , 2018
<i>Coprinus</i> sp Pers.	Saprobe	Ethiopia	Dejene <i>et al.</i> , 2017a
<i>Coprinus pseudoplicatilis</i> Voglina	Saprobe	Ethiopia	Dejene <i>et al.</i> , 2017a



<i>Coprinus sterquilinus</i> (Fries)	Saprobe	Kenya	Boa, 2004
Fries			
<i>Coprinopsis nivea</i> (Pers.) Redhead, Vilgalays & Moncalvo	Saprobe	Ethiopia	Dejene et al., 2017
<i>Cotylidia aurantiaca</i> (Pat.) A.L. Welden	Saprobe	Burundi , Rwanda	Degreeef et al., 2016
<i>Craterellus</i> Pers.	Ectomycorrhizal	Ethiopia	Dejene et al., 2017a
<i>Cymatoderma dendriticum</i> (Pers) D.A Raid	Saprobe	Malawi	Rammeloo and Wallyne, 1993; Boa, 2004
<i>Cystodermella elegans</i> (Beeli)	Saprobe	Rwanda	Degreeef et al., 2016
Harmaja			
<i>Dacryopinax spathularia</i> (Schwein.) G.W. Martin	Saprobe	Rwanda	Degreeef et al., 2016
<i>Entoloma argypus</i> P. Karst	Saprobe	Tanzania	Rammeloo and Wallyne, 1993; Boa, 2004
<i>Favolus brasiliensis</i> (Fr) Fr.	Saprobe	Malawi	Rammeloo and Wallyne, 1993
<i>Ganoderma appplanatum</i> (Pers.) Pat.	Saprobe	Ethiopia, Uganda	Dejene et al., 2017b; Ngom et al., 2022
<i>Ganoderma lucidum</i> (Curtis) P. Karst	Saprobe	Tanzania	Mdachi et al., 2004
<i>Ganoderma</i> sp P. Karst sp	Saprophytic	Kenya	Njuguini et al., 2018
<i>Gyroporus castaneus</i> (Bull Quel.)	Saprobe	Malawi	Rammeloo and Wallyne, 1993
<i>Hygrophoropsis aurantiaca</i> (Wurfen) Maire	Saprobe	Ethiopia	Dejene et al., 2017a
<i>Hymenagaricus</i> sp Heinem.	Saprobe	Ethiopia	Dejene et al., 2017a
<i>Hypholoma fasciculare</i> (Huds: Fr.) P.Kumm.	Saprobe	Burundi	Nteziryayo et al., 2019
<i>Hypholoma subviride</i> (Berk. & M.A. Curtis) Dennis	Saprobe		Boa, 2004; Degreeef et al., 2016
<i>Inonotus</i> P.Karst.	Saprobe	Tanzania	Mdachi et al., 2004
<i>Joga kadzonzo</i>	Ectomycorrhizal	Kenya	Wandati et al., 2013
<i>Joga muhama</i>	Ectomycorrhizal	Kenya	Wandati et al., 2013
<i>Kuehneromyces mutabilis</i> Schaeff. Singer & A.H.Sm	Saprobe	Tanzania	Rammeloo and Wallyne, 1993; Boa, 2004
<i>Lactarius</i> aff. <i>gymnocarpus</i> R. Heim ex Singer	Ectomycorrhizal	Burundi	Buyck and Nzigidahera, 1995
<i>Lactarius deliciosus</i> (L.ex Fr.) Gray	Ectomycorrhizal	Malawi, Tanzania , Burundi	Rammeloo and Wallyne, 1993; Harkonen et al., 2003; Tibuhwa, 2013
<i>Lactarius edulis</i> (Verbeken & Buyck)	Ectomycorrhizal	Burundi , Tanzania	Buyck and Nzigidahera, 1995
<i>Lactarius</i> <i>gymnocarpoides</i> Verbeken	Ectomycorrhizal	Tanzania, Malawi	Harkonen et al., 2003; Boa, 2004



<i>Lactarius Pers.</i>	Ectomycorrhizal	Tanzania	Harkonen et al., 2003; Boa, 2004; Mdachi et al. 2004; Tibuhwa, 2013
<i>Lactarius inversus</i> (Gooss.-Font &r. Heim.) Verbeken	Ectomycorrhizal	Burundi	Buyck and Nzigidahera, 1995
<i>Lactarius kabansus</i> Pegler & Pearce	Ectomycorrhizal	Burundi , Tanzania	Buyck and Nzigidahera, 1995; Harkonen et al., 2003; Boa, 2004; Degreef et al., 2016; Ntezirayayo et al., 2019
<i>Lactarius luteolus</i> Peck.	Ectomycorrhizal	Tanzania	Tibuhwa, 2013
<i>Lactarius medusae</i> Verbeken	Ectomycorrhizal	Tanzania	Harkonen et al., 2003; Tibuhwa, 2013
<i>Lactarius pelliculatus</i> (Beeli) Buyck.	Ectomycorrhizal	Tanzania	Boa, 2004
<i>Lactarius phlebophyllus</i> Heim. L. Roussel	Ectomycorrhizal	Tanzania	Boa, 2004
<i>Lactarius piperatus</i> Verbeken	Ectomycorrhizal	Malawi	Boa, 2004
<i>Lactarius pumilus</i> Verbeken	Ectomycorrhizal	Tanzania	Harkonen et al., 2003; Tibuhwa, 2013
<i>Lactarius rubroviolascens</i> R. Heim.	Ectomycorrhizal	Tanzania	Boa, 2004
<i>Lactarius tanzanicus</i> Karhula & Verbeken	Ectomycorrhizal	Tanzania	Tibuhwa, 2013
<i>Lactarius vellereus</i> (Fr.) Fr.	Ectomycorrhizal	Malawi	Boa, 2004
<i>Lactarius volvoides</i> Kurhula	Ectomycorrhizal	Tanzania	Harkonen et al., 2003; Tibuhwa, 2013
<i>Lactifluus densifolius</i> Verbeken & Karhula	Ectomycorrhizal	Burundi	Degreeef et al., 2016
<i>Lactifluus edulis</i> (Verbeken & Buyck) Buyck	Ectomycorrhizal	Burundi	Degreeef et al., 2016
<i>Lactifluus gymnocarpoides</i> (Verbeken) Verbeken	Ectomycorrhizal	Burundi	Degreeef et al., 2016
<i>Lactifluus Kigomaensis</i> sp. nov. De Crop & Verbeken	Ectomycorrhizal	Tanzania	Decrop et al., 2012
<i>Lactifluus longisporus</i> (Verbeken) Verbeken	Ectomycorrhizal	Burundi	Degreeef et al., 2016
<i>Lactifluus luteopus</i> Verbeken	Ectomycorrhizal	Burundi	Degreeef et al., 2016
<i>Laetiporus Murr.</i> (IJ- 2014)	Saprobe	Tanzania	Juma et al., 2016
<i>Laetiporus sulphureus</i> (Bull.) Murill	Saprobe	Burundi , Ethiopia, Tanzania	Boa, 2004; Tibuhwa, 2011; Woldegiorgis et al., 2015; Dejene et al. 2017a; Ntezirayayo et al., 2019
<i>Langemannia gigantea</i> (Batsch) Rostk.	Saprobe	Burundi	Boa, 2004
<i>Lentinus cladopus</i> (Lev.)	Saprobe	Malawi	Rammeloo and Wallyne, 1993; Boa, 2004; Degreeef et al., 2016
<i>Lentinus edodes</i> (Berk.) Pegler	Saprobe	Ethiopia	Woldegiorgis, 2015
<i>Lentinus Fr.</i>	Saprobe	Ethiopia	Dejene et al., 2017
<i>Lentinus prolifer</i> (Pat. & Har.) Pegler	Saprobe	Uganda	Boa, 2004



<i>Lentinus sajor caju</i> (Fr.; Fr.)	Saprobe	Tanzania	Boa, 2004; Rammeloo and Wallyne, 1993; Hussein et al., 2015; Degreef et al., 2016
<i>Lentinus squarrosulus</i> Mont.	Saprobe	Tanzania	Rammeloo and Wallyne, 1993, Boa, 2004; Hussein, et al., 2015, Nteziryayo et al. 2019
<i>Lenzite elegans</i> (Spreng.) Pat.	Saprobe	Tanzania	Boa, 2004; Rammeloo and Wallyne, 1993
<i>Lepista cafrorum</i> (Kalchbr. & Mc Owan)	Saprobe	Malawi	Boa, 2004; Rammeloo and Wallyne, 1993
<i>Lepista nuda</i> (Bull.) Cooke	Saprobe	Burundi	Boa, 2004
<i>Lepista sordida</i> (Schumach.) Singer	Saprobe	Rwanda	Degreef et al., 2016
<i>Leucoagaricus holosericeus</i> (Gillet) M.M Moser	Saprobe	Ethiopia	Dejene et al., 2017a
<i>Leucoagaricus leucothites</i> (Vittad.) Wasser	Saprobe	Tanzania	Boa, 2004; Dejene et al., 2017a
<i>Leucoagaricus rhodocephalus</i> (Berk.) Pegler	Saprobe	Tanzania	Boa, 2004
<i>Leucoagaricus rubrotinctus</i> (Peck.) Singer	Saprobe	Ethiopia	Dejene et al., 2017a
<i>Leucocoprinus birnbaumii</i> (Corda) Singer	Saprobe	Ethiopia	Dejene et al., 2017a
<i>Leucocoprinus cepistipes</i> (Sowerby) Pats.	Saprobe	Ethiopia	Dejene et al., 2017a
<i>Inocybe</i> (Fr.) Fr.	Saprobe	Malawi	Boa, 2004
<i>Lycoperdon</i> sp Pers.	Saprobe	Ethiopia	Dejene et al., 2017a
<i>Lycoperdon perlatum</i> Pers.	Saprobe	Burundi	Boa, 2004
<i>Macrocybe lobayensis</i> (R. Heim)) Pegler & Lodge	Saprobe	Malawi	Boa, 2004
<i>Macrolepiota aberdarense</i> Mbaluto	Saprobe	Kenya	Mbaluto, 2015
<i>Macrolepiota dolichaula</i> (Berk. & Br.) Pegler & Rayner	Saprobe	Malawi, Tanzania, Uganda	Pegler, 1977; Rammeloo and Wallyne, 1993; Harkonen et al. 2003; Boa, 2004; Njuguini et al. 2018; Nteziryayo et al. 2019; Ngom et al., 2022
<i>Macrolepiota mastoidea</i> (Fr.) Singer	Saprobe	Kenya	Pegler, 1977
<i>Macrolepiota procera</i> (Scop.) Singer	Saprobe	Kenya, Tanzania, Ethiopia, Malawi, Uganda	Pegler, 1977; Rammeloo and Wallyne, 1993; Boa, 2004; Tibuhwa et al., 2011; Tibuhwa, 2013; Hussein et al., 2015; Dejene et al., 2017b; Njuguini et al., 2018; Ngom et al., 2022
<i>Macrolepiota rhacodes</i> (Vittad.) Singer	Saprobe	Burundi	Boa, 2004
<i>Macrolepiota</i> Singer	Saprobe	Ethiopia	Dejene et al., 2017a; Njuguini et al., 2018
<i>Malombo</i>	Saprobe	Kenya	Wandati et al., 2013
<i>Marasmiellus inoderma</i> (Berk.) Singer	Saprobe	Rwanda	Degreef et al., 2016
<i>Marasmius arborescens</i> (Henn.) Beeli	Saprobe	Rwanda	Degreef et al., 2016



<i>Marasmius bekolacongoli</i> Beeli	Saprobe	Burundi, Rwanda	Degreeef et al., 2016
<i>Marasmius oreades</i> (Bolton) Fr.	Saprobe	Burundi	Boa, 2004
<i>Micropsalliota brunneosperma</i> (Singer) Pegler	Saprobe	Malawi	Boa, 2004
<i>Morchella elata</i> Fr.	Saprobe	Burundi	Boa, 2004
<i>Morchella esculenta</i> Fr.	Saprobe	Ethiopia	Dejene et al., 2016
<i>Mycoamaranthus congolensis</i> (Dissing & M. Lange)	Ectomycorrhizal		Degreeef et al., 2016
<i>Castellano & Walleyn Pholiota nameko</i> (T.Ito) S. Ito & S. Imai	Saprobe	Ethiopia	Gizaw, 2017
<i>Obulando</i>	Saprobe	Kenya	Wandati et al., 2013
<i>Obumpokompoko</i>	Saprobe	Uganda	Nakalembe et al., 2009.
<i>Obutoosa</i>	Saprobe	Uganda	Nakalembe et al., 2009
<i>Olando</i>	Saprobe	Kenya	Wandati et al., 2014
<i>Oudemansiella tanzanica</i> nom. prov. Magingo	Saprobe	Tanzania	Magingo et al., 2004
<i>Panus conchatus</i> (Bull.) Fr.	Saprobe	Tanzania	Hussein et al., 2015
<i>Parmelia sulcata</i> Taylor	Saprobe	Burundi	Boa, 2004
<i>Paxillus brunneotomentosus</i> Heinem. & Rammeloo	Saprobe	Rwanda	Degreeef et al., 2016
<i>Perenniporia mundula</i> (Wakef) Ryvarden	Saprobe	Malawi	Boa, 2004
<i>Phlebopus colossus</i> (R.Heim) Singer	Saprobe	Burundi, Malawi	Boa, 2004; Degreeef et al., 2016
<i>Pholiota</i> sp (Fr.) P Kumm.	Saprobe	Ethiopia	Dejene et al., 2017a
<i>Plebopus sudanicus</i> (Har. & Pat.)	Saprobe	Malawi	Rammeloo and Wallyne, 1993; Boa, 2004
<i>Pleurocybella porrigens</i> (Pers.) Singer	Saprobe	Burundi	Boa, 2004
<i>Pleurotus</i> (HK-37)	Saprobe	Tanzania	Raymond et al., 2013
<i>Pleurotus</i> (Jack. Ex. (Fr.) P. Kumm.	Saprobe	Uganda, Kenya,	Nakalembe et al., 2009; Gateri et al., 2014
<i>Pleurotus</i> aff. <i>cystidiosos</i> O.K. Mill.	Saprobe	Burundi	Buyck and Nzigidahera, 1995
<i>Pleurotus citrinopileatus</i> Singer	Saprobe	Kenya	Musieba et al., 2011; Musieba et al., 2012; Okoth 2013; Nteziryayo et al., 2019
<i>Pleurotus contrarius</i> Sacc.	Saprobe	Tanzania	Pegler, 1977
<i>Pleurotus</i> <i>cystidiosos</i> O.K. Mill.	Saprobe	Burundi, Tanzania	Huseein et al., 2016; Degreeef et al., 2016
<i>Pleurotus djamor</i> (Rumph. ex Fr.) Boedijn	Saprobe	Tanzania, Kenya, Uganda, Rwanda	Boa, 2004; Harkonen et al., 2003; Tibuhwa, 2013; Nakalembe et al., 2015; Degreeef et al., 2016; Njuguini et al., 2018;



<i>Pleurotus eryngii</i> (D C.) Quel.	Saprobe	Tanzania	Tibuhwa, 2011and 2013
<i>Pleurotus flabellatus</i> Sacc.	Saprobe	Tanzania, Kenya , Rwanda	Pegler, 1977; Mshandete and Cuff, 2008; Degreef et al., 2016
<i>Pleurotus florida</i> (Mont.) 0 (Singer)	Saprobe	Kenya	Wandati et al., 2014
<i>Pleurotus lignatilis</i> (Pers. Ex Fr.) Kummer	Saprobe	Uganda	Pegler, 1977
<i>Pleurotus limpidus</i> (Fr.) Sacc.	Saprobe	Tanzania	Pegler, 1977
<i>Pleurotus luteoalbus</i> Beeli	Saprobe	Kenya	Pegler, 1977
<i>Pleurotus opuntiae</i> (Durieu & Lev.) Sacc.	Saprobe	Kenya	Pegler, 1977
<i>Pleurotus ostreatus</i> (Jacq. ex Fr.) Kumm.	Saprobe	Burundi, Ethiopia, Tanzania , Uganda	Pegler, 1977; Woldegiorgis, 2015; Ngom et al., 2022
<i>Pleurotus sajor- caju</i> (Fr.) Sing.	Saprobe	Tanzania	Harkonen et al., 2003; Mdachi et al. 2004; Tibuhwa, 2011; Tibuhwa, 2013
<i>Pleurotus sapidus</i> (Quel.)	Saprobe	Kenya	Musieba, 2013; Otieno et al., 2015
<i>Pleurotus tuber-regium</i> (Rump. Ex Fr.) Singer	Saprobe	Burundi, Tanzania	Rammeloo and Wallyne 1993; Boa, 2004; Harkonen et al., 2003; Tibuhwa, 2011 and 2013; Degreef et al., 2016
<i>Pluteus umborosus</i> (Pers.) P. Kumm.	Saprobe	Tanzania	Hussein et al., 2016
<i>Polyozellus multiplex</i> (Underw.) Murill	Saprobe	Burundi	Boa, 2004
<i>Polyporales</i> Gaum.	Saprobe	Tanzania	Hussein et al., 2016
<i>Polyporus P.Micheli</i> ex Adans	Saprobe	Uganda, Tanzania, Ethiopia,	Nakalembe et al., 2015; Harkonen et al., 2003; Tibuhwa 2013., Dejene et al. 2017b;
<i>Polyporus brasiliensis</i> Speg.	Saprobe	Malawi	Boa, 2004
<i>Polyporus cinnabarinus</i> (Jacq.) Fr.	Saprobe	Burundi	Ntezirayayo et al., 2019
<i>Polyporus moluccensis</i> (Mont.) Ryvarden	Saprobe	Malawi, Tanzania	Boa 2004
<i>Polyporus tenuiculus</i> (P. Beauv.) Fr.	Saprobe	Kenya , Rwanda, Tanzania,Uganda,	Nakalembe et al., 2009; Tibuhwa, 2013; Hussein et al., 2016; Njuguini et al., 2018; Ngom et al., 2022
<i>Psathyrella atroumboonata</i> Pegler	Saprobe	Rwanda, Malawi,	Boa 2004; Rammeloo and Wallyne, 1993; Degreef et al., 2016
<i>Psathyrella candolleana</i> (Fr.) Maire	Saprobe	Malawi	Rammeloo and Wallyne, 1993
<i>Psathyrella tuberculata</i> (Path.) A.H. Sm.	Saprobe	Rwanda	Degreeef et al., 2016
<i>Ptychoverpa bohemica</i> (Krombh) J. Schrot.	Saprobe	Burundi	Boa, 2004
<i>Pulveroboletus aberrans</i> (Heinemann & Goos.- Font)	Ectomycorrhizal	Malawi	Boa, 2004; Rammeloo and Wallyne, 1993
<i>Pulveroboletus</i> Spec.	Ectomycorrhizal	Malawi	Rammeloo and Wallyne, 1993
<i>Pycnoporus sanguineus</i> (L.;Fr.)	Saprobe	Malawi	Rammeloo and Wallyne, 1993



<i>Rubinoboletus balloui</i> (Peck) Heinem. & Rammeloo	Ectomycorrhizal	Degreeef et al., 2016	
<i>Rubinoboletus luteopurpureus</i> (Beeli) Hein. & Rammeloo	Ectomycorrhizal	Malawi	Rammeloo and Wallyne, 1993; Boa, 2004
<i>Russula Pers.</i>	Ectomycorrhizal	Ethiopia	Gateri et al., 2014; Dejene, et al. 2017
<i>Russula afronigricans</i> Buyck	Ectomycorrhizal	Malawi	Boa, 2004
<i>Russula albofloccosa</i> Buyck.	Ectomycorrhizal	Tanzania	Harkonen et al., 2003
<i>Russula atropurpurea</i> (Krombh.) Britelm.	Ectomycorrhizal	Malawi	Rammeloo and Wallyne 1993
<i>Russula congoana</i> Pat.	Ectomycorrhizal	Burundi, Tanzania	Harkonen et al., 2003; Boa, 2004; Tibuhwa, 2013; Degreeef, et al., 2016
<i>Russula cyanoxantha</i> (Schaeff.) Fr.	Ectomycorrhizal	Malawi	Boa, 2004; Rammeloo and Wallyne 1993
<i>Russula delica</i> Fr.	Ectomycorrhizal	Malawi	Boa, 2004; Rammeloo and Wallyne 1993
<i>Russula hiemisilvae</i> Buyck	Ectomycorrhizal	Burundi, Tanzania	Harkonen et al., 2003; Mdachi et al., 2003; Boa, 2004; Degreeef, et al., 2016
<i>Russula liberiensis</i> Singer	Ectomycorrhizal	Tanzania	Boa, 2004
<i>Russula ochroleuca</i> (Pers.) Fr.	Ectomycorrhizal	Malawi	Boa, 2004; Rammeloo and Wallyne, 1993
<i>Russula phaeocephala</i> Buyck	Ectomycorrhizal	Burundi, Tanzania	Buyck and Nzigidahera 1995; Boa, 2004; Degreeef et al., 2016
<i>Russula rosea</i> Pers.	Ectomycorrhizal	Malawi	Boa, 2004; Rammeloo and Wallyne, 1993
<i>Russula roseoviolacea</i> Buyck	Ectomycorrhizal	Burundi	Degreeef et al., 2016
<i>Russula schizoderma</i> Pat	Ectomycorrhizal	Malawi	Rammeloo and Wallyne, 1993; Boa 2004
<i>Russula sejuncta</i> Buyck	Ectomycorrhizal	Burundi	Degreeef et al., 2016
<i>Russula sublaevis</i> (Buyck) Buyck	Ectomycorrhizal	Tanzania	Boa, 2004
<i>Russula tanzaniae</i> Buyck.	Ectomycorrhizal	Tanzania	Boa, 2004
<i>Russula xerampelina</i> (Schaeff.) Fr.	Ectomycorrhizal	Burundi	Boa, 2004
<i>Rusulla cellulata</i> Buyck	Ectomycorrhizal	Burundi, Tanzania	Buyck and Nzigidahera 1995, Harkonen et al., 2003, Boa, 2004; Tibuhwa, 2013; Degreeef et al., 2016
<i>Rusulla ciliata</i> Buyck	Ectomycorrhizal	Tanzania	Harkonen et al., 2003; Tibuhwa 2013
<i>Rusulla compressa</i> Buyck	Ectomycorrhizal	Tanzania, Kenya	Harkonen et al., 2003; Boa, 2004; Tibuhwa, 2013; Wandati et al., 2013
<i>Schizophyllum commune</i> Fr.	Saprobe	Malawi, Kenya, Tanzania, Ethiopia, Burundi	Pegler, 1977; Rammeloo and Wallyne 1993; Harkonen et al., 2003; Boa, 2004; Degreeef et al., 2016; Dejene et al., 2017a; Muchane et al., 2021
<i>Sparassis crispa</i> (Wulfen) Fr.	Saprobe	Burundi	Boa, 2004
<i>Stereopsis hiscens</i> (Berk. & Rav.) D. A. Reid	Saprobe	Malawi	Boa, 2004; Rammeloo and Wallyne 1993
<i>Stropharia rugosoannulata</i> Farlow ex Murill	Saprobe	Kenya	Njuguini et al., 2018



<i>Suillus cavipes</i> (Opat.) A.H.Sm. & Thiers	Ectomycorrhizal	Burundi	Boa, 2004
<i>Suillus granulatus</i> (L.) Roussel	Ectomycorrhizal	Tanzania , Kenya, Malawi	Rammeloo & Wallyne 1993, Harkonen et al., 2003; Boa 2004; Tibuhwa 2013, Degreef et al., 2016, Njuguini et al., 2018; Muchane et al., 2021
<i>Suillus luteus</i> (L.) Roussel	Ectomycorrhizal	Burundi , Ethiopia , Tazania , Kenya , Malawi	Buyck and Nzigidahera, 1995; Mdachi et al., 2003; Boa, 2004; Degreef et al., 2016; Dejene et al., 2017a; Njuguini et al., 2018
<i>Termitomyces (Beeli) R. Heim</i>	Termitophilic	Burundi	Buyck and Nzigidahera, 1995; Harkonen et al., 2003; Nakalembe et al., 2009; Gateri et al., 2014; Wandati, 2014; Woldegiorgis et al., 2015
<i>Termitomyces (Oruka-stipe)</i>	Termitophilic	Kenya	Wandati et al., 2014
<i>Termitomyces aurantiacus</i> (Heim)	Termitophilic	Uganda, Ethiopia, Kenya, Malawi	Rammeloo and Wallyne, 1993; Buyck, 1995; Harkonen et al., 2003; Boa, 2004; Nakalembe et al., 2009; Tibuhwa, 2012 ; Tibuhwa, 2013; Dejene et al. 2017; Woldegiorgis et al., 2015
<i>Termitomyces clypeatus</i> Heim	Termitophilic	Kenya, Tanzania, Ethiopia, Malawi	Pegler, 1977; Rammeloo and Wallyne 1993; Harkonen et al., 2003; Boa, 2004; Tibuhwa, 2011, Tibuhwa, 2012; Tibuhwa, 2013; Nakalembe and Kabasa, 2009; Ashagriel et al., 2014; Woldegiorgis et al., 2015; Dejene et al., 2017b; Ngom et al., 2022
<i>Termitomyces eurhizus</i> (Berk.) Heim	Termitophilic	Kenya, Tanzania, Uganda, Malawi	Pegler, 1977; Harkonen et al., 2003; Boa, 2004; Nakalembe et al., 2009, Dejene et al., 2017b
<i>Termitomyces globulus</i> R. Heim & Gooss.- Font	Termitophilic	Kenya, Uganda	Pegler, 1977; Nakalembe et al., 2009
<i>Termitomyces le-testui</i> (Pat.) Heim	Termitophilic	Tanzania, Burundi, Ethiopia, Kenya, Uganda 2004	Pegler, 1977; Harkonen et al., 2003; Boa, 2004; Tibuhwa, 2012; Ashagriel et al., 2014; Woldegiorgis et al., 2015; Degreef et al., 2016; Dejene et al., 2017a
<i>Termitomyces mammiformis</i> R. Heim	Termitophilic	Burundi, Tanzania	Pegler, 1977, Tibuhwa, 2012; Tibuhwa, 2013, Degreef et al., 2016
<i>Termitomyces microcarpus</i> (Berk. & Broome) R. Heim	Termitophilic	Tanzania, Ethiopia, Uganda, Kenya, Rwanda; Uganda	Pegler, 1977; Rammeloo and Wallyne 1993; Harkonen., et al., 2003; Boa 2004; Nakalembe and Kabasa 2009; Olila et al., 2007; Tibuhwa, 2012; Tibuhwa, 2013; Woldegiorgis et al., 2015; Degreef et al., 2016 ; Dejene et al., 2017a ; Muchane et al., 2021; Ngom et al., 2022
<i>Termitomyces robustus</i> (Beeli) R. Heim	Termitophilic	Burundi, Tanzania, Ethiopia, Uganda, Malawi	Pegler 1977; Rammeloo and Wallyne 1993; Boa, 2004; Tibuhwa, 2013; Degreef et al., 2016; Dejene et al., 2017a; Ngom et al., 2022
<i>Termitomyces saggitiformis</i> (Kalchbr. & Cooke) D. A Reid	Termitophilic	Tanzania	Tibuhwa, 2012; Tibuhwa, 2013
<i>Termitomyces singidensis</i> Saarim & Hark.	Termitophilic	Tanzania	Harkonen et al., 2003; Boa, 2004; Tibuhwa, 2013
<i>Termitomyces sp</i> (Joga utuwe)	Termitophilic	Kenya	Wandati et al., 2013
<i>Termitomyces (Mariondonik)</i>	Termitophilic	Kenya	Wandati et al., 2013
<i>Termitomyces striatus</i> (Beeli) R. Heim	Termitophilic	Malawi, Tanzania, Uganda	Pegler, 1977; Rammeloo and Wallyne, 1993; Buyck and Nzigidahera, 1995 ; Boa, 2004; Tibuhwa 2013;Degreef et al., 2016
<i>Termitomyces titanicus</i> Pegler & Pearce	Termitophilic	Burundi, Malawi	Boa, 2004; Tibuhwa, 2012; Degreef et al., 2016
<i>Termitomyces tylerianus</i> Otieno	Termitophilic	Uganda, Kenya	Pegler, 1977; Rammeloo and Wallyne, 1993; Harkonen et al., 2003; Nakalembe et al., 2009 ; Tibuhwa, 2013; Degreef et al., 2016
<i>Termitomyces umkowaani</i> (Cooke & Massee) D.A Reid	Termitophilic	Tanzania	Tibuhwa, 2011; 2012 & 2013; Muchane et al., 2021



<i>Termitomyces eurhizus</i> (Berk) Heim	Termitophilic	Tanzania	Tibuhwa, 2012
<i>Termitomyces le-testui</i> (Pat.) R. Heim	Termitophilic	Ethiopia, Tanzania	Boa, 2004; Tibuhwa, 2013; Kabede, 2017
<i>Termitomyces rabuori</i> Otieno	Termitophilic	Kenya	Pegler, 1977
<i>Termitomyces schimperi</i> (Pat.) R. Heim	Termitophilic	Burundi, Rwanda, Tanzania, Malawi	Rammeloo and Wallyne, 1993; Pegler, 1977; Boa, 2004; Kabede, 2017; Dejene et al., 2017a; Degreef et al., 2016
<i>Trametes polyzona</i> Pers.	Saprobe	Burundi	Ntezirayayo et al., 2019
<i>Trametes suaveolens</i> (L.) Fries	Saprobe	Burundi	Boa, 2004
<i>Tremella fuciformis</i> Berk.	Saprobe	Tanzania, Uganda	Tibuhwa, 2013; Ngom et al., 2022
<i>Tricholoma caligatum</i> (Viv.) Ricken	Saprobe	Burundi	Boa, 2004
<i>Tricholoma lobayense</i> Heim	Saprobe	Malawi	Rammeloo and Wallyne, 1993
<i>Tricholoma magnivelare</i> (Peck) Redhead	Ectomycorrhizal	Burundi	Boa, 2004
<i>Tricholoma spectabilis</i> Peerally & Sutra	Saprobe	Burundi	Buyck et al., 1995
<i>Troglia infundibuliformis</i> Trogia; Fr.	Saprobe	Malawi	Boa, 2004
<i>Tubosaeta brunneosetosa</i> (Sing.) E. Horak	Saprobe	Malawi	Boa, 2004; Rammeloo and Wallyne, 1993
<i>Tylopilus niger</i> (Heinem. & Gooss.-Font.) Wolfe	Saprobe	Ethiopia	Dejene et al., 2017a
<i>Vascellum</i> F. Smarda	Saprobe	Ethiopia	Dejene et al., 2017a
<i>Vasellum pratense</i> (Pers.) Kreisel	Saprobe	Malawi	Boa, 2004; Rammeloo and Wallyne 1993
<i>Volvariella bobycina</i> (Scheaeff.) Singer	Saprobe	Malawi, Tanzania	Boa, 2004; Pegler, 1977; Rammeloo and Wallyne, 1993
<i>Volvariella speciosa</i> (Fr.: Fr.) Sing.	Saprobe	Uganda	Nakalembe et al., 2009
<i>Volvariella volvacea</i> (Bull.;Fr.)Sing.	Saprobe	Tanzania, Malawi, Kenya, Uganda	Pegler,1977, Rammeloo and Wallyne, 1993, Mshandete and Cuff, 2008, Harkonen et al., 2003, Boa, 2004, Tibuhwa et al., 2011, Tibuhwa, 2013
<i>Xerocomus pallidosporus</i> Heinem.	Saprobe	Malawi	Boa, 2004
<i>Xerocomus soyeri</i> Heinem.	Saprobe	Malawi	Boa, 2004, Rammeloo and Wallyne, 1993
<i>Xerocomus subspinulosus</i> Heinem.	Saprobe	Burundi	Buyck et al., 1995, Degreef et al., 2016
<i>Xerula radicata</i> (Relhan: Fr.) Dorfelt	Saprobe	Malawi	Boa, 2004

Distribution of the wild edible Mushrooms in different Phyla and families

In the division Ascomycota, the macrofungi species belonged to the class Lecanormycetes represented by 1

species *Parmelia sulcata* Taylor from the Parmeliaceae family and class Pezizomycetes represented by 3 species (*Morchella elata* Fr. , *Morchella esculenta* Fr. and *Ptychoverpa bohemica* (Krombh) J. Schrot.) from



Morchellaceae family. In the division *Basidiomycota*, the macrofungi species belonged to the class *Agaricomycetes* represented by species from *Agaricales* (47%), *Russulales* (16%), *Polyporales* (10%), *Cantharellales* (10%) and *Boletales* (8%). Species from other orders (*Auriculariales*, *Dacrymycetales*, *Hymenochaetales*, *Incertae sedis*, *Stereopsidales*, *Thelephorales*, and *Tremellales*) had less than 2% presentation.

Overall, the *Russulaceae* family had the highest number of species (51 species), followed by *Agaricaceae* (41 species), *Cantharellaceae* (30 species), *Lyophyllaceae* (26 species), *Polyporaceae* (21 species), *Amanitaceae* (20 species), *Pleurotaceae* (20 species), *Boletaceae* (18 species) and *Tricholomataceae* (11 species). The other families had less than 10 species (Figure 1). One species from order *Polyporales* was only identified up to order level while 9 species were described by their local names.

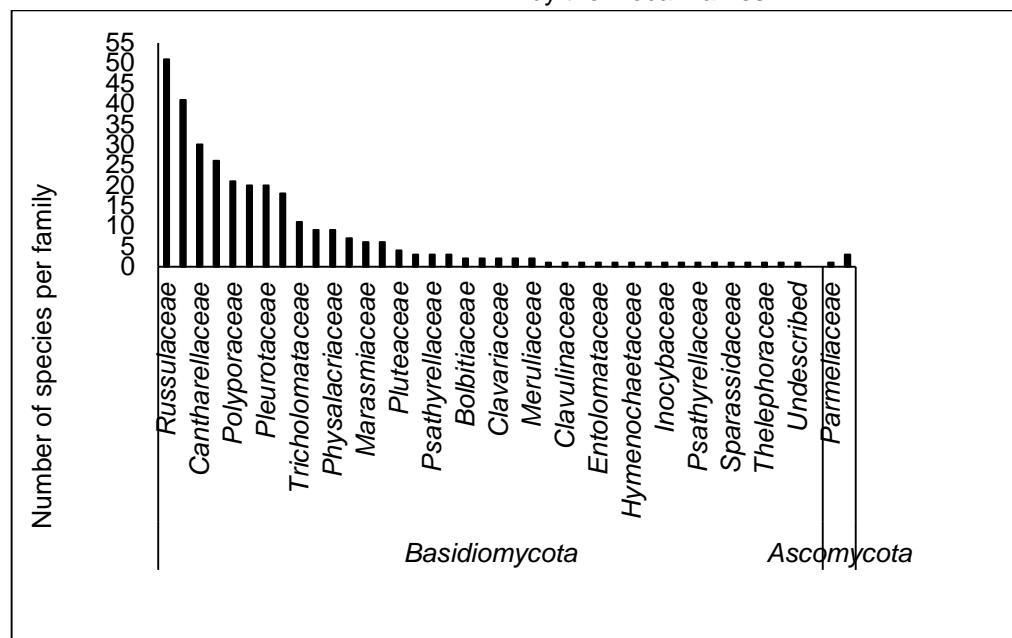


Figure 1: Distribution of WEMS into phyla and families

Distribution of Wild Edible Mushrooms in East African Countries

The WEMs occurred in all 8 East African countries namely Burundi, Kenya, Ethiopia, Somalia, Rwanda, Tanzania, Uganda and Malawi. Tanzania (143 species), Malawi (90 species) and Burundi (89) had the highest number while Somalia had the least with only 2 species (*Agaricus campestris* (L. ex Fr.), *Agaricus amboensis*

(Fayod) Sacc.) from *Agaricaceae* family (Figure 2). The number of WEM species in other countries declined in order, Kenya (53 species), Ethiopia (46 species) Rwanda (28) and Uganda (28 species). WEMs communities in Tanzania, Malawi, and Burundi were dominated by ectomycorrhiza species while in other countries, the highest proportion of WEMs communities comprised saprophytic macrofungi species (Table 2).

Table 2: Wild edible / medicinal mushrooms species in East African countries

Families	Countries								
	Kenya	Uganda	Tanzania	Burundi	Rwanda	Malawi	Ethiopia	Somalia	Total
Agaricaceae	11	5	13	3	3	12	16	2	65
Amanitaceae	1	–	4	8	–	13	–	–	26
Auriculariaceae	4	–	4	1	2	2	1	–	14
Bolbitiaceae	–	–	–	–	–	1	–	–	1
Boletaceae	–	–	5	4	1	7	–	–	17
Boletinellaceae	–	–	–	1	–	2	–	–	3
Bondarzewiaceae	–	–	1	–	–	–	–	–	1



Cantharellaceae	-	-	21	15		6	1	-	43
Clavuriaceae	-	-	-	-	-	1	-	-	1
Clavulinaceae	-	-	1	-	-	-	-	-	1
Dacrymycetaceae	-	-	-	-	1	-	-	-	1
Entolomataceae	-	-	1	-	-	-	-	-	1
Fomitopsidaceae	-	-	1	1	-	-	1	-	3
Ganodermataceae	1	-	1	-	-	-	1	-	3
Gyroporaceae	-	-	-	-	-	1	-	-	1
Hygrophoropsidiaceae	-	-	-	-	-	-	1	-	1
Hymenochaetaceae	-	-	1	-	-	-	-	-	1
Inocybaceae	-	-	-	-	-	1	-	-	1
Lyophyllaceae	12	10	15	9	3	7	10	-	66
Marasmiaceae	-	-	-	3	3	-	-	-	6
Meruliaceae	-	-	-	-	-	1	-	-	1
Morchellaceae	-	-	-	2	-	-	1	-	3
Parmeliaceae	-	-	-	1	-	-	-	-	1
Paxillaceae	-	-	-	1	-	-	-	-	1
Physalaciaceae	1	1	3	-	6	2	1	-	14
Pleurotaceae	9	3	10	4	3	1	1	-	31
Pluteaceae	1	2	2	-	-	2	-	-	7
Polyporaceae	2	3	8	3	1	7	3	-	27
Psathyrellaceae	-	-	-	-	1	2	3	-	6
Repetobasidiaceae	-	-	-	1	1	-	-	-	2
Russullaceae	1		32	19	-	12	1	-	65
Schizophyllaceae	1	-	1	1	-	1	1	-	5
Sparassidaceae	-	-	-	1	-	-	-	-	1
Stereopsidaceae	-	-	-	-	-	-	1	-	1
Strophariaceae	1	-	1	2	1	-	2	-	7
Suillaceae	2	-	2	2	-	2	1	-	9
Therephoraceae	-	-	-	1	-	-	-	-	1
Tremellaceae	-	-	1	-	-	-	-	-	1
Tricholomataceae	-	-	-	5	2	6	-	-	13
unidentified	4	4	-	-	-	-	-	-	9
Total	51	28	128	88	28	89	46	2	

Groups of Wild mushrooms in East Africa and substrate utilization

(a) Saprophytic WEMs

In East Africa, saprophytic WEMs diversity comprised 147 species (46% of total) within the division Basidiomycota (51 genera within 26 families) and Ascomycota (3 genera within 2 families) (Fig. 2). WEMs

in Ascomycota belonged to the *Parmeliaceae* family (*Parmelia sulcata* Taylor) and *Morchellaceae* family (*Morchella elata* Fr., *Morchella esculenta* Fr. Basidiomycota division, *Agaricaceae* family (40 species) had the highest number of species, followed by *Polyporaceae* (21 species), *Pleurotaceae* (20 species), *Physalaciaceae* (9 species), *Tricholomataceae* (7



species), Auriculariaceae (7 species), Marasmiaeae (6 species), Strophariaceae (6 species) and Psathyrellaceae (4 species) families. The other families had less than 4 species (Fig. 2).

(b) Ectomycorrhizal WEMs

In Eastern Africa, ectomycorrhizal WEMs comprised 131 species (41% of the total) within the division Basidiomycota (23 genera within 12 families) (Fig. 4). *Russulaceae* family (51 species) had the highest

number of species, followed by *Cantharellaceae* (30 species), *Amanitaceae* (19 species), *Boletaceae* (18 species) and *Suillaceae* (3 species). The other families had less than 2 species (Fig. 4). The highest documented ectomycorrhizal species belongs to the genus *Cantharellus* and *Russula* sp (26 species), followed by *Amanita* (20 species) and *Lactarius* (16 species) respectively (Table 1, Fig. 2).

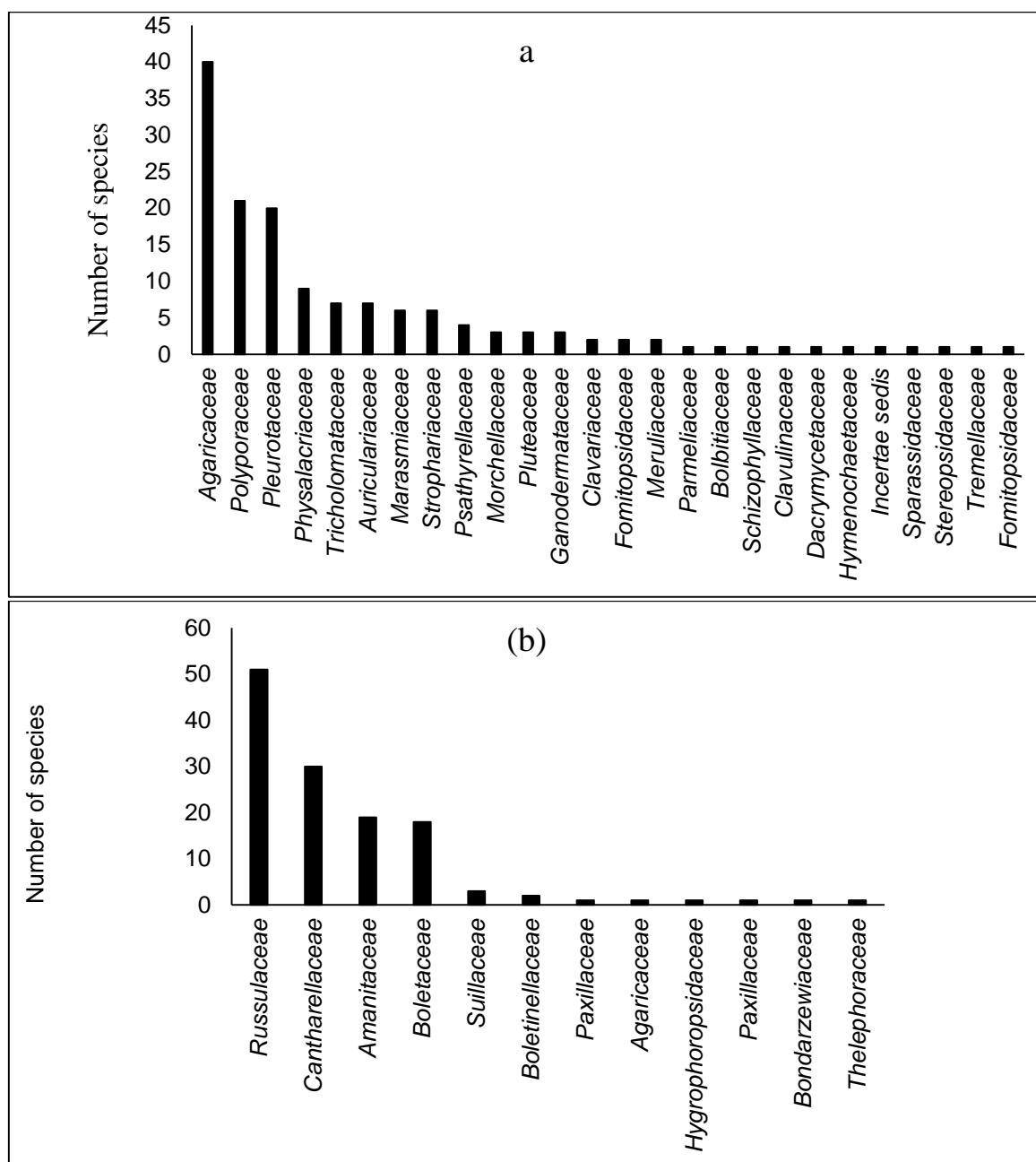


Figure 2: Number of Saprobes(a) and Ectomycorrhizal (b) wild edible mushroom species per family in Eastern Africa



(c) Termitophilic WEMs mushrooms

In East Africa, termitophilic species documented are 25 species (7.8% of the total). The species were reported in all the countries of East Africa with the exception of Somalia (Table 1).

Domestication Status of WEMs in East Africa

Domestication of wild edible mushrooms is gaining popularity in the East African region. Domestication is the process of bringing WEMs into cultivations. There are about 2000 WEMs in the world, but only around 30 species are in cultivation. In East Africa 14 saprophytic species (*Amylosporus* IJ-2014, *Coprinus cinereus* (Schaeff.) Gray, *Lentinus sajor caju* (Fr.; Fr.), *Lentinus squarrosus* Mont., *Hypholoma fasciculare* (Huds: Fr.) P.Kumm., *Nameko pholiota* (T.Ito) S. Ito & S. Imai, *Oudemansiella tanzanica* Magindo, *Pleurotus citrinopileatus* Singer, *Pleurotus flabellatus* Sacc., *Polyporales* sp Gaum., *Polyporus cinnabarinus*, *Trametes polyzona* Pers., *Volvariella volvacea* (Bull.; Fr.) Sing.) have been tissue cultured, and tested for spawn production and mushroom production under different growth conditions (Table 3). However, the cultivation of WEMs in this region is still in its early stages with only 24 species under experimental trials (Table 3).

The species *Auricularia polytricha* (Mont.) Sacc., *Laetiporus* sp Murr., *Polyporus tenuiculus* (P. Beauv.)

Fr., *Pleurotus cystidiosus* O.K. Mill. and *Termitomyces* sp (Beeli) R. Heim. are still at spawn and tissue culture stages (Table 3). *Auricularia auricula* (Bull.: Fr.) did not fruit despite successful growth in spawn run and pinning stages (Table 3).

Process of cultivating wild edible mushrooms in East Africa

(a) Tissue culture and mycelia colonization rates of WEMs

The pure mycelial culture was established from fourteen mushroom species (Table 3). The main culture media used were Potato Dextrose Agar (PDA) and Malt Extract Agar (Malt) (Table 3). However, to culture *Pluteus umbrosus* (Pers.) P. Kumm., *Hypholoma fasciculare* (Huds: Fr.) P. Kumm., *Trametes polyzona* Pers., *Pleurotus citrinopileatus* Singer and *Lentinus squarrosus* Mont. AVOINE media was used (Table 3). SDYA media was also used for obtaining *Pholiota Nameko* mycelial culture (Table 3). The species were incubated upside down at 25-30 °C depending on the species' temperature requirements for 4 up to 48 days (Table 3). Pure cultures were subcultured for one to three months and cultures were stored at 4 °C to –30 °C and some in liquid nitrogen at 196 °C..

Table 3: Domestication status of edible and medicinal Mushrooms in Eastern Africa

Species	Tissue culture			Spawn production			Mushroom cultivation				References
	Media	Incubation	Substrat e	Incubation	Incubation	Subst rate	Spawn	Pinning	Fruiting	Yield	
Temperature											
Period			period			run					
--days-			°C			-----days-----					-g/kg-
<i>Coprinus cinereus</i>	PDA	4	a	-	10	g	7 (12)	9 (15)	11 (17)	238	Mshandete and Cuff, 2008; Raymond et al., 2013
<i>Volvariella volvacea</i>	PDA	5	a	-	12	g	9	11	13	114	Mshandete and Cuff, 2008
<i>Pleurotus flabellatus</i>	PDA	6	a	-	13	g	12	14	17	371	Mshandete and Cuff, 2008
<i>Oudemansiella tanzanica</i>	MALT	7	b	-	21	i, h, g	18 (19)	20 (21)	19 (22)	-	Magindo et al., 2004
<i>Auricularia auricula</i>	MALT	5 (7)	a, b	-	12	j, h, l, j	8 (14)	13 (23)	20 (35)	96- (266)	Onyango et al/2011
<i>Pleurotus citrinopileatus</i>	PDA, AVOINE	7 (10)	a, c	25	-	h, i, k, m, n, o, p, q, v, x	8 (21)	13 (28)	-	28.3 (397.7)	Ntezirayo et al., 2019
<i>Pleurotus</i> sp.2 (HK-37)	MALT	5 (7)	-	28	-	g	12 (14)	18 (43)	20 (46)	119.2	Raymond et al/2013



<i>Pleurotus cystidiosus</i>	PDA	27	a	-	-	-	-	-	-	-	-	Juma et al., 2015
<i>Pleurotus djamor</i>	PDA	-	-	-	-	-	-	-	-	-	123 (238)	Nakalembe et al., 2015
<i>Polyporales sp.</i>	PDA	48	a	25	-	t	67	-	105	23.29	Juma et al., 2015	
<i>Polyporus tenuiculus</i>	PDA	20	a	-	-	-	-	-	-	-	-	Juma et al., 2015
<i>Laetiporus sp. IJ-2014</i>	PDA	24	a	-	-	-	-	-	-	-	-	Juma et al., 2015
<i>Lentinus squarrosus</i>	AVOINE	-	a	25	-	u	25(27)	-	-	212.6	Ntezirayo et al., 2019	
<i>Amylospora sp. IJ-2014</i>	PDA	23	a	25	-	t	31	-	43	15.5	Juma et al., 2015	
<i>Auricularia polytricha</i>	PDA	27	a	-	-	-	-	-	-	-	-	Juma et al., 2015
<i>Nameko pholiota</i>	PDA, MALT & SDYA	18(28)	a,b,c ,d,e, f	-	-	-	60	-	-	55 (797)	Gizaw, 2015	
<i>Termitomyces microcarpus</i>	PDA	10	b	-	3 (31)	i, u	-	-	-	-	-	Olila et al., 2007
<i>Lentinus saju caju</i>	PDA & MALT	4	a	25	10	n, v	24 (28)	49 (52)	1 (2)	52	Hussein et al., 2015	
<i>Panus conchatus</i>	PDA & MALT	4	a	-	17	n, v	-	-	-	-	-	Hussein et al., 2016
<i>Pluteus umbrosus</i>	AVOINE	7	a	25	17	n, v	-	-	-	-	-	Hussein et al., 2016
<i>Hypholoma fasciculare</i>	AVOINE	-	a	25	-	u	24(26)	-	-	153.5	Ntezirayo et al., 2020	
<i>Trametes polystylos</i>	AVOINE	-	a	25	-	u	28(30)	-	-	208.7	Ntezirayo et al., 2020	
* <i>Pleurotus ostreatus</i>	MEA	-	a	-	14	v, j, w, x, y	15(20)	20(29)	22(31)	803 (2170)	Tekeste et al., 2020	
* <i>Agaricus subreflexus</i>	-	-	f2	-	-	z	14(21)	7(29)	22(31)	684 (1428)	Thongklang et al., 2014	

Key: Spawn substrate included (a) Sorghum grains, (b) Millet grains and (c) Wheat grains (d) Barley grain (e) rice (f) shredded maize (f2) Rye grain. Cultivation substrate included (g) Sisal waste (h) Rice straw, (i) Saw dust, (j) Sugarcane bagasse (k) Wheat straw, (l) Grass straw, (m) Maize cobs, (n) Banana leaves, (o) Ground nut, (p) Soya bean, (q) Coffee husk, (r) Coconut waste (s) Bean straw (t) Dried sugarcane tops (u) Cotton waste (v) Barley straw (w) Sesame stalks (x) Teff straw (y) Commercial compost. The values in parenthesis are maximum number of days taken or yield produced. The words in asterisks are for species commonly cultivated in the world.

(a) Spawn preparation

Spawn preparation started by obtaining good quality grains. Sorghum was the most used grain in spawn production in the EA region (Table 4). The grains were first soaked in water overnight followed by boiling for 10-20 min. Excess water was drained and 1-2% calcium carbonate (agricultural lime) was added and mixed thoroughly with the grains (Table 4). The grains were allowed to drain off excess water through a sieve or by spreading on a clean plastic sheet to air dry. Once drained, the grains were filled into 330-750 ml bottles up to $\frac{3}{4}$ of the jar, which are closed with lids and autoclaved for 1 hour at 121°C to kill the contaminants (Table 4). The bottles were allowed to cool and then inoculated aseptically with 2 pieces of 1cm culture of mushroom mycelia. The inoculated bottles were fitted with lids which were then shaken thoroughly by hand to spread the mycelia. The bottles were incubated at 25-28°C with caps

loosely fitted in a well ventilated incubator for 10 days- 21 days depending on the species (Table 3).

(a) Substrate preparation requirements and growth conditions for wild edible mushrooms

Different types of agricultural and industrial waste have proved potential in the cultivation of fourteen species of domesticated mushrooms in the EA region (Table 4). This is evident by the ability of the species to colonize the substrates and form fruitbodies. The preparation of substrates in this study employed two methods of substrate preparation which were compost and non-compost methods. The compost preparation method was used to cultivate *Coprinus cinereus* (Schaeff.), *Pleurotus flabellatus* (Sacc.) and *Volvariella volvacea* (Bull.;Fr.) Sing. on sisal waste and manure. The outdoor composting outdoor method took 21 days. For the non-composted substrates, the materials are first soaked in water in the ratio of 1:2 (substrate: water) overnight for 4 days (Table 4). Prepared substrates were



subjected to sterilization and pasteurization at 100-121°C for 1-6 hours and pasteurization at 70°C for two hours. The sterilized bags are allowed to cool to prevent the mycelium from heat destruction. Thereafter, 1-6% (three-six teaspoonfuls) of spawn were inoculated into each bag of the substrate (Table 4). The bags were transferred to the incubation room with temperatures between 23-30°C and humidity ranging between 69-81%.

The mycelial colonization progress was monitored daily until the bags were fully colonized (Table 4). Contaminated bags were removed once spotted and taken outside the incubation room for disposal. Once fully colonized the bags were transferred to the fruiting room with lower temperatures 18-30°C and humidity 50-95% (Table 4).

Table 4: Mycelial run requirements for Wild Edible Mushrooms in East Africa (EA)

Species	Culture	Media treatment		Incubation		Culture preservation	References
	Source	Reagent	Antibiotic	Temperature (°C)	Days		
<i>Oudemansiella tanzanica</i>	Tissue	-	-	25	7	-	Magingo et al., 2004
<i>Pleurotus HK-37</i>	Pure culture	Distilled water, 70% Ethanol	-	-	8	-	Raymond et al., 2013
<i>Coprinus cinereus</i>	Tissue	3% hydrogen peroxide, 70% ethanol	250mg/l ampicillin	28	4	Malt at 4 °C	Mshadete and Cuff, 2008
<i>Pleurotus flabellatus</i>	Tissue	-	250mg/l ampicillin	28	6	Malt at 4 °C	Mshadete and Cuff, 2008
<i>Volvariella volvaceae</i>	Tissue	-	250mg/l ampicillin	28	5	Malt at 30 °C	Mshadete and Cuff, 2008
<i>Lentinus saju caju</i>	Tissue	Distilled water, 70% ethanol, MEA	-	28	4	MEA at 4 °C & liquid nitrogen	Hussein et al., 2016
<i>Lentinus squarrosus</i>	Tissue	Distilled water, 70% Ethanol	1 capsule chloramphenical	25		-	Ntezirayo et al., 2019
<i>Amylospora</i> sp IJ -2014	Tissue	PDA		25	-	-	Juma et al., 2015
<i>Nameko pholiota</i>	Pure culture	-	0.025g chloramphenicol/250	25	30	-	Gizaw, 2015
<i>Hypholoma fasciculare</i>	Tissue	Distilled water, 70% ethanol	-	25		AVOINE at 4 °C (monthly subculturing)	Ntezirayo et al., 2019
<i>Trametes polyzona</i>	Tissue	Distilled water, 70% ethanol	-	25		-	Ntezirayo et al., 2019
<i>Pleurotus citrinopileatus</i>	Tissue	Distilled water, & 70% ethanol	-	25		-	Musieba et al., 2012; Musieba et al., 2013, Ntezirayo et al., 2019
<i>Auricularia auricula</i>	-	-	-	-	-	-	Onyango et al., 2011

From the study, 47 edible species were analyzed for nutritional and medicinal benefits. Among the most commonly utilized species are the *Termitomyces*,

Pleurotus and *Ganoderma* species. Species belonging to *Termitomyces* were the most (21%) analyzed in this study followed by *Pleurotus* species (12%) (Table 5)



Table 5: Nutritional status of wild edible mushrooms of East Africa

Species	Proximate Analysis	Mineral Content	Vitamins	Amino Acids	Anti-oxidants	Reference
<i>Afrocantharellus</i>						
<i>splendens</i>	-	-	-	-	+	Tibuhwa et al., 2014
<i>Agaricus</i>						
<i>bisporus</i>	+	+	+	+	+	Wandati et al., 2013
<i>Agaricus</i>						
<i>campestris</i>	-	+	-	-	+	Woldegiorgis et al., 2015
<i>Agaricus</i> sp	-	-	-	+	-	Mdachi et al., 2004
<i>Amanita</i>						
<i>zambiana</i>	+	+	-	-	+	Wandati et al., 2013
<i>Auricularia</i>						
<i>judae</i>	-	-	-	-	+	Hussein et al., 2015
<i>Boletus clavipes</i>	-	-	-	+	-	Mdachi et al., 2004
<i>Boletus</i>						
<i>pruinatus</i>	-	-	-	+	-	Mdachi et al., 2004
<i>Cantharellus</i>						
<i>Rufopunctatus</i>	-	-	-	-	+	Tibuhwa et al., 2014
<i>Cantharellus</i>	-	-	-	+	-	Mdachi et al., 2004
<i>Ganoderma</i>						
<i>lucidum</i>	-	-	-	+	-	Mdachi et al., 2004
<i>Inonotus</i> sp	-	-	-	+	-	Mdachi et al., 2004
<i>Joga kadzonzo</i>	+	+	-	-	+	Wandati, 2014
<i>Joga muhamma</i>	+	+	-	-	+	Wandati, 2014
<i>Lactarius</i> sp	-	-	-	+	-	Mdachi et al., 2004
<i>Lentinus</i>						
<i>edodes</i>	-	+	-	-	+	Woldegiorgis et al., 2015
<i>Lentinus</i> sajor						
<i>caju 1</i>	-	-	-	-	+	Hussein et al., 2015
<i>Lentinus</i> sajor						
<i>caju 2</i>	-	-	-	-	+	Hussein et al., 2015
<i>Lentinus</i>						
<i>squarrosulus</i>	-	-	-	-	+	Hussein et al., 2015
<i>Lentinus</i>						
<i>sulpureus</i>	-	+	-	-	+	Woldegiorgis et al., 2015
<i>Macrolepiota</i>						
<i>procera</i>	-	-	-	-	+	Hussein et al., 2015
<i>Malombo</i>	+	+	-	-	+	Wandati et al., 2013
<i>Auricularia</i> sp	+	+	-	-	+	Wandati et al., 2013
<i>Obulando</i>	+	+	-	-	+	Wandati et al., 2013
<i>Oando</i>	+	+	+	+	+	Wandati et al., 2013
<i>Panus cochatus</i>	-	-	-	-	+	Hussein et al., 2015
<i>Pleurotus caju</i>	-	-	-	+	-	Woldegiorgis et al., 2015
<i>Pleurotus florida</i>	-	+	+	+	+	Wandati et al., 2013



<i>Pleurotus ostreatus</i>	-	-	-	-	+	Woldegiorgis et al., 2015
<i>Pleurotus tenuiculus</i>	-	-	-	-	+	Hussein et al., 2015 Nakalembe, 2013; Nakalembe et al., 2015
<i>Polyporus tenuiculus</i>	+	+	+	+	-	Woldegiorgis, et al., 2015
<i>Russula hiemisilvae</i>	-	-	-	+	-	Wandati, 2014
<i>Rusulla compressa</i>	+	+	-	-	+	Woldegiorgis et al., 2015
<i>Termitomyces (oruka-stipe)</i>	+	+	+	+	+	Wandati et al., 2013
<i>Termitomyces aurantiacus</i>	-	+	-	-	+	Woldegiorgis et al., 2015
<i>Termitomyces clypeatus</i>	-	+	-	-	+	Woldegiorgis et al., 2015
<i>Termitomyces eurhizus</i>	+	-	-	+	-	Nakalembe et al., 2013
<i>Termitomyces globulus</i>	+	-	-	+	-	Nakalembe et al., 2013
<i>Termitomyces letestui</i>	-	+	-	-	+	Woldegiorgis et al., 2015 Woldegiorgis et al., 2015; Nakalembe et al., 2015
<i>Termitomyces microcarpus</i>	-	+	+	-	+	Nakalembe et al., 2015
<i>Termitomyces tyleranus</i>	-	+	+	-	-	Woldegiorgis et al., 2015
<i>Termitomyces sp</i>	-	+	-	-	-	Wandati et al., 2013
<i>Termitomyces sp</i>	+	+	+	+	+	Wandati et al., 2013
<i>Termitomyces sp (joga utuwe)</i>	+	+	-	-	+	Wandati et al., 2013
<i>Termitomyces sp (mariondonik)</i>	+	+	-	-	+	Wandati et al., 2013
<i>Volvariella speciosa</i>	-	+	+	-	-	Nakalembe et al., 2015

Key: analyzed (+), Not analyzed (-)

Discussions

The numbers of WEMs observed in this review corresponds to the number of identified and described wild edible mushroom species (300 species) reported in sub-Saharan Africa (Soro et al., 2019). This is a high number considering we have about 2000-2166 edible mushroom species which have been reported from all

over the world (Rai et al., 2005; Nakalembe et al., 2015; Li et al., 2021). From the results, EA is endowed with a wide range of WEMs whose benefits can be applied to support mushroom industry as well as tree restoration programs, especially with tree species growing in association with ectomycorrhizal fungi.



There are over six thousand (6000) ECM fungi species mainly in division *Basidiomycota* and into lesser extent in division *Ascomycota* (Smith and Read, 2008). ECM fungi are characterized by presence of a fungal mantle that envelops host roots and a Hartig net that surrounds root epidermal. In addition, the ECM fungi form soil-borne mycelia network important in uptake and translocation of nutrients and water to host tree species as well as movement of nutrients between individual tree hosts (Simard and Durall, 2004). These ECM fungi develop a dense mycelia network of about 200 m in a gram of dry soil. This mycelia network in soil is very important in the formation of soil aggregates, which in return facilitate carbon sequestrations in soil (Walland, 2006). ECM macrofungi are also the most expensive macrofungi, with a total market of billions of US dollars. The highly traded ECM species include *Tuber melanosporum*, *Tuber magnatum* (white truffle), *Tricholoma matsutake*, *Boletus edulis*, *Cantharellus cibarius* and *Amanita caesarea*. The majority of these species were from Miombo woodland and associated with *Brachystegia*, *Julbernardia*, *Isoberlinia* and *Uapaca* tree species known to symbiotically associate with ectomycorrhizal mushrooms species (Degreef et al., 2020). Miombo woodland makes up approximately 10% of the ecosystem in Kenya, Tanzania, Uganda and Burundi extending to DRC Congo, Malawi, Zimbabwe, Angola, Zambia and Mozambique explaining the high number of ectomycorrhiza mushroom species found in Tanzania, Burundi and Malawi (Degreef et al., 2020). The Ectomycorrhiza mushrooms form part of the most appreciated wild edible mushrooms (Kamalebo and De Kesel, 2020). All chanterelles are edible most and appreciated owing to large quantities harvested during the rainy seasons and the long shelf life. Apart from local consumption, the sale of the produce takes place in the markets and along the roads in many African countries (Degreef et al., 2016). Besides, *Amanita* comprises highly toxic species such as *Amanita phalloides* and *Amanita muscaria*, it's a genus with valuable members such as *Amanita losii* which has been proposed as the most productive (Degreef et al., 2020). The high number of *Russula* species could be attributed to its associations with different tree hosts. Other ectomycorrhiza species were documented in exotic plantation forests such as *Pinus*, Cypress and *Eucalyptus*. A few are from *Acacia* species plantations.

Ectomycorrhiza mushrooms are valuable WEMs widely collected for food and are also an important

income source, generating US 70 per month for families during the rainy season (Degreef et al., 2020). Despite the key role of Miombo mushrooms in communities, the species are threatened by the practice of felling host trees that form symbiotic relationships with mushrooms to create farmlands and fuel sources (Degreef et al., 2020). As well, ectomycorrhizal mushrooms cannot be cultivated, due to their symbiotic nature with specific tree species. Thus, sustainable harvesting remains the most appropriate strategy for the conservation of beneficial mushrooms and their habitat. The other alternative is to cultivate ectomycorrhizal seedlings which have been inoculated with specific fungus for continued mushroom supply when conditions are conducive for fructification. It is important to note that native fungi fail to form symbiotic relationships with exotic tree species and therefore the right fungus should be identified for appropriate indigenous tree species (Ducouso et al., 2012). Therefore, sustainable land management practices and cheap fuel alternatives have been proposed to secure and sustain ectomycorrhiza mushrooms (Degreef, et al., 2020).

Termitomyces is a paleotropical genus of agarics growing in association with termites and their nests. The species belong to *Lyophyllaceae* (division *Basidiomycota*) with 30-40 species estimated globally (Kirk et al., 2008). Although species in this genus are saprobic, decomposing plant-derived material (e.g., wood, dry grass, and leaf litter) organic matter, the species are completely dependent on termites to survive. They are the most preferred species that form a favorite delicacy for most African and Asian communities. Their good flavor, taste and texture make them acceptable among these communities (Sathiya et al., 2020). Out of 27 that have been documented in Africa and Southeast Asia, 23 species are edible and 3 species are medicinal. *Termitomyces* mushrooms live in a mutualistic obligate relationship with termites, also found in Miombo woodland, although widely distributed in montane areas (Harkonen et al., 2003). Unlike many saprophytic mushrooms, termitomyces species are difficult to cultivate artificially due to the complexity of the symbiotic relationship that exists between the termites and the mushroom forming fungi (Hsieh and Ju, 2018). However, attempts to culture mycelia on artificial media have been fruitful. In order to maintain sustainable utilization of *termitomyces* in their natural habitats, efforts are needed to limit the destruction of ecosystems that are known to harbor the species in East Africa. Even as



communities harvest the species for home consumption, surplus can be sold to buy other types of foods and for improved livelihoods. Communities that shy off from consuming wild mushrooms should also be educated on which species are edible and sustainable methods of harvesting.

Cultivation of WEMs in this region is however still in its early stages with only 24 species under experimental trials. This is in comparison with 100 under artificial cultivation and 60 % of the species at commercial level in China (Willis, 2018). Of this, about 20 species are currently being cultivated at an industrial level (Gizaw, 2015). The advantage of cultivating mushrooms is the diversification of livelihoods and strengthening the resilience of farmers, the crop has a short cycle that takes only 2 months. Mushrooms are grown at a very low cost since cultivation is mostly done indoors with low requirements for water compared to other crops (FAO, 2017). Cultivation of wild edible mushrooms is also an environmentally friendly technological process of recycling organic waste (Girmay et al., 2016; Hussein et al., 2016). Since small-scale cultivation of over 20 species is undertaken across China, the model used can be used for technology transfer to researchers and farmers in East African countries and Africa at large. The demand for mushrooms is also growing in East Africa (Kenya, Tanzania, Burundi, DR Congo, Uganda) though partly met despite importation and wild gathering efforts (Degreef et al., 2016).

Domestication of WEMs in the region is an enormous opportunity to expand the mushroom industry since most WEMs have regional adaptability, grow rapidly at higher temperatures ($>25^{\circ}\text{C}$) and their growth conditions can easily be replicated using locally available agricultural and industrial waste (Table 3). Types of agricultural and industrial wastes include wheat straw, rice straw, beans and barley, leaves of various trees, maize stalks, millets, cotton husks and banana leaves (Table 3) (Magingo et al., 2004)

The attempt to domesticate species would also offer year round valuable protein sources and broadening of income avenues for families in East Africa. However, none of these species have been commercially introduced into cultivation and the availability of their mother cultures (germplasm) for research and propagation purpose is uncertain. Despite the limited effort to successfully cultivate these species in East Africa, the attempts made elsewhere have been successful (Thawthong, et al., 2014; Rizal et al., 2015;

Bandara et al., 2020). Thus, further studies are required in the region to determine optimal conditions (temperature, humidity, pH, aeration, suitable substrates) for each species in order to successfully grow them invitro.

The process of domesticating WEMs begins with the identification and selection of edible mushroom species whose cultivation potential is known. The cultivability of a species is established by determining if species in the same genus are cultivatable. The edibility of wild mushrooms is best obtained through local knowledge from herbalists, local collectors, and local communities. The knowledge can also be combined with scientific information (Thawthong et al., 2014). The choice of the best substrate for any mushroom species is also a very important step in the successful cultivation of mushrooms. Thus, the process is followed by selection of suitable artificial media and agricultural or industrial substrates. The process of cultivating mushrooms addresses the problem of agricultural and industrial waste by converting it into rich protein food sources (Mshandete and Cuff, 2008, Raymond, et al., 2013, Degreef, et al., 2020). Culturing the species on artificial culture helps to obtain pure mycelia of a specific strain to be cultivated (Mshandete and Cuff, 2008). The step is followed by choice of spawn with high quality viability which is the ability of the mycelia to produce fruiting bodies under different treatments of substrates, suitable conditions such as pH, temperatures and humidity (Thawthong, et al., 2014). Two methods are used in substrate preparation which mainly varies with species requirements. The two main methods are composting and non-composting. During the compost preparation, the substrate is made into piles that are turned after every three days from the 5th day of substrate preparation to the 21st day. According to Mshandete and Cuff (2008), the composting method during substrate preparation manipulates the natural succession of microorganisms which involves preparing compost using agricultural or industrial wastes, organic and inorganic manures, and calcium sulfate (gypsum). The non-composting method involves soaking the substrate overnight to soften it. The aim of soaking is to moisten the substrate which is followed by draining off excess water by spreading it on a surface that allows excess water to drain off (Nteziryayo et al., 2019). To ensure that the right moisture is retained, a palm squeeze test is done to ensure that water does not drip between the fingers (Hussein et al., 2016). Pasteurization of the substrate is meant to kill the microorganisms that would



compete with mushroom mycelium and hinder production. In some instances, supplements are added to the substrates as a nitrogen source such as chicken manure, cow dung, or rice bran at a rate of 1 to 30% depending on the type of supplement used. The aim is to increase mushroom yield and overall production (Mshadete, 2013).

Among the species analyzed for nutritional composition, *Ganoderma lucidum* is known for immunity enhancement, especially among HIV & AIDS patients, *Volvariella volvacea* for lowering high blood pressure, and *Schizophyllum commune* for fighting cancer (Chandrawanshi et al., 2017). Attempts to domesticate the species in EA have only been made for *Volvariella volvacea* which was successful. Medicinal mushrooms owe their potential to the secondary metabolites they produce which enhance their immunomodulation, antibacterial, antifungal, antioxidant, antidiabetic, anticancer, antiallergic, anticholesterolemic, cardiovascular protector, antiparasitic, antiviral, hepatoprotective and detoxification effects (Valverde et al., 2015).

Species belonging to *Termitomyces* were the most (21%) analyzed in this study followed by *Pleurotus* species (12%) (Table 5). This is probably because *Termitomyces* mushrooms are the most consumed edible mushrooms due to their good taste and flavor and ease of availability during the rainy seasons. *Pleurotus* mushrooms grow easily on dead logs immediately after an adequate rainy season and it's also the 2nd most cultivated species in the world making it a species of interest to researchers. The composition of edible mushrooms meaty taste and immunity enhancement properties make them acceptable as nutritional sources worldwide (Teklit, 2015). Also, the protein content in wild edible mushrooms is almost equal to that of milk (Lister, 2015) which makes them a good alternative to expensive and unavailable meat, especially among rural poor populations (Musieba et al, 2012) Additionally, wild mushrooms comprise carbohydrates, vitamins, minerals, antioxidants, phytochemicals and fiber essential for human health (Lister, 2015, Juma et al., 2016, Bandara et al., 2017). The valuable vitamins which are also lacking in most food sources such as Vit D, E, B1, ,B2 and B12 form part of the nutritional composition of most mushrooms (Valverde et al., 2015). Vitamin D in

mushrooms is the only non-animal source and thus an important diet, especially, for vegetarians. As a result, mushrooms are gaining popularity recently promising to be potential functional food and medicines that would prevent and treat diseases such as diabetes, cancer and malnutrition (Zhang et al., 2012). It's worth noting that some of the medicinal mushrooms documented in East Africa are cultivatable such as *Ganoderma lucidum*, a means through which communities can exploit to generate income.

Conclusion

There is a high diversity of cultivatable wild edible saprophytic mushrooms in Eastern

- Africa with the potential to expand the mushroom industry.
- The yield of wild edible mushrooms is almost comparable to that of exotic species though domestication of the species is still in infancy stages.
- The East Africa region is endowed with many agricultural and industrial wastes suitable for the cultivation of most saprophytic mushrooms in the region.
- Efforts made to analyze Wild edible mushrooms in the region are an indication of the growing interest to explore natural resources for improved health and nutrition.
- Availability of the WEMs mother cultures (germ-plasm) for research and propagation is uncertain.

Recommendation

- Further research is needed for more detailed information on diversity, domestication, nutrient content analysis of cultivatable wild mushrooms and their suitable growth requirements for the best yields.
- More research is required to determine nutritional and phytochemical composition of over 80% wild edible/ medicinal mushrooms in East Africa region.
- Regional gene bank is needed for storage and conservation of wild mushrooms germplasm.

Author Contributions

All authors have equal contribution.

Conflict of Interest

The authors declare no conflict of interest.



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***Hebeloma limbatum*: Türkiye Mikotası İçin Yeni Bir Kayıt**

Ertuğrul SESLİ

*Sorumlu yazar: ertugrulsesli@trabzon.edu.tr

Trabzon Üniversitesi, Fatih Eğitim Fakültesi, Biyoloji Eğitimi Ana Bilim Dalı, Söğütlü, Trabzon,
Türkiye / ertugrulsesli@trabzon.edu.tr

Öz: *Hebeloma limbatum* Beker, Vesterh. & U. Eberh. (Hymenogastraceae)'a ait meyveşmiler Türkiye'den ilk kez toplanmış ve burada arazi resimleri ve kısa bir tartışma ile birlikte sunulmuştur. Koleksiyonun teşhisini araziden elde edilen veriler ve mikroskopik inceleme sonuçlarına göre yapılmıştır. Yeni kayıt nispeten küçük, yapışkan, turuncumsu kahverengi veya beyazımsı şapka; krem rengi veya açık kahverengi lameller; beyazımsı sap; badem veya limon şeklinde, süslü, soluk sarı, 10–14 × 5–7.5 µm boyutlarında bazidiyosporlar; çomak veya şişe biçiminde, bazen başlıklı ve 30–100 × 5–14 µm büyülüklüğünde sistytyumlar ile teşhis edilir.

Anahtar kelimeler: Etli mantar, *Hebeloma*, Trabzon, Türkiye, Yeni kayıt

***Hebeloma limbatum*: A New Record For the Turkish Mycota**

Abstract: Fruit bodies belonging to *Hebeloma limbatum* Beker, Vesterh. & U. Eberh. (Hymenogastraceae) were collected for the first time from Turkey and presented herein with field photos and a short discussion. The identification of the collection was made according to the data obtained from the field and the results of microscopic examination. The new record is identified with relatively small, sticky, orangey brown or whitish pileus; cream-colored or light brown lamellae; whitish stipe; almond- or lemon-shaped, ornamented, pale yellow, 10–14 × 5–7.5 µm sized basidiospores; club- or bottle-shaped, sometimes capped, and 30–100 × 5–14 µm sized cystidia.

Key words: Fleshy fungi, *Hebeloma*, Trabzon, Türkiye, New record

Giriş

Hebeloma (Fr.) P. Kumm. cinsi mantarlar genellikle keskin ve bazen kötü ve itici kokuları, öbekler halinde toprakta veya çürümekte olan ağaçlar üzerinde yayılış göstergeleri ile arazide; ilginç ve çok sayıdaki sistytyumları, badem veya limon biçiminde, bol ve genellikle süslü bazidiyosporları ile laboratuvara kolaylıkla yakın cinslerden ayırt edilirler. Ancak cins içerisindeki birçok tür birbirine yakın özelliklere sahip olduğundan teşhis edilmeleri kolay değildir. Bir bölümü zehirli olup genellikle beslenme amaçlı kullanılmazlar. Mevcut çalışmadan önce Doğu Karadeniz bölümünden uzun süreli arazi çalışmalarında genellikle sonbaharda, ladin-kayın ormanlarında, çürümekte olan döküntüler ve ağaçlar üzerinde, çimenlerde ve diğer tip arazilerde *Hebeloma leucosarx* P.D. Orton, *H. populinum* Romagn., *H. laterinum* (Batsch) Vesterh., *H. radicosum* (Bull. : Fr.) Ricken (Sesli ve Baydar, 1996), *Hebeloma aff. aestivale* Vesterh. (Sesli ve ark., 2015), *H. sacchariolens* Quél.

(Akata ve Sesli, 2017) ve *H. avellaneum* Kauffman (Sesli ve ark., 2018) türleri tespit edilmiştir. Literatür araştırmalarına göre bu cins günümüzde Türkiye'de yaklaşık 40 civarında türle (Sesli ve ark., 2020) ve dünyada ise yaklaşık 690 kayıtla temsil edilmektedir (Kirk ve ark., 2008). Elbette bu sayının tamamı farklı tür olmayıabilir. Diğer öbekler üzerinde olduğu gibi *Hebeloma* konusunda da Türkiye'de ve dünyada yapılacak yeni çalışmalar araştırmacıları beklemektedir.

Mevcut çalışmanın amacı Türkiye için yeni olan *Hebeloma limbatum* türünün morfolojik özelliklerini tanıtmaktır.

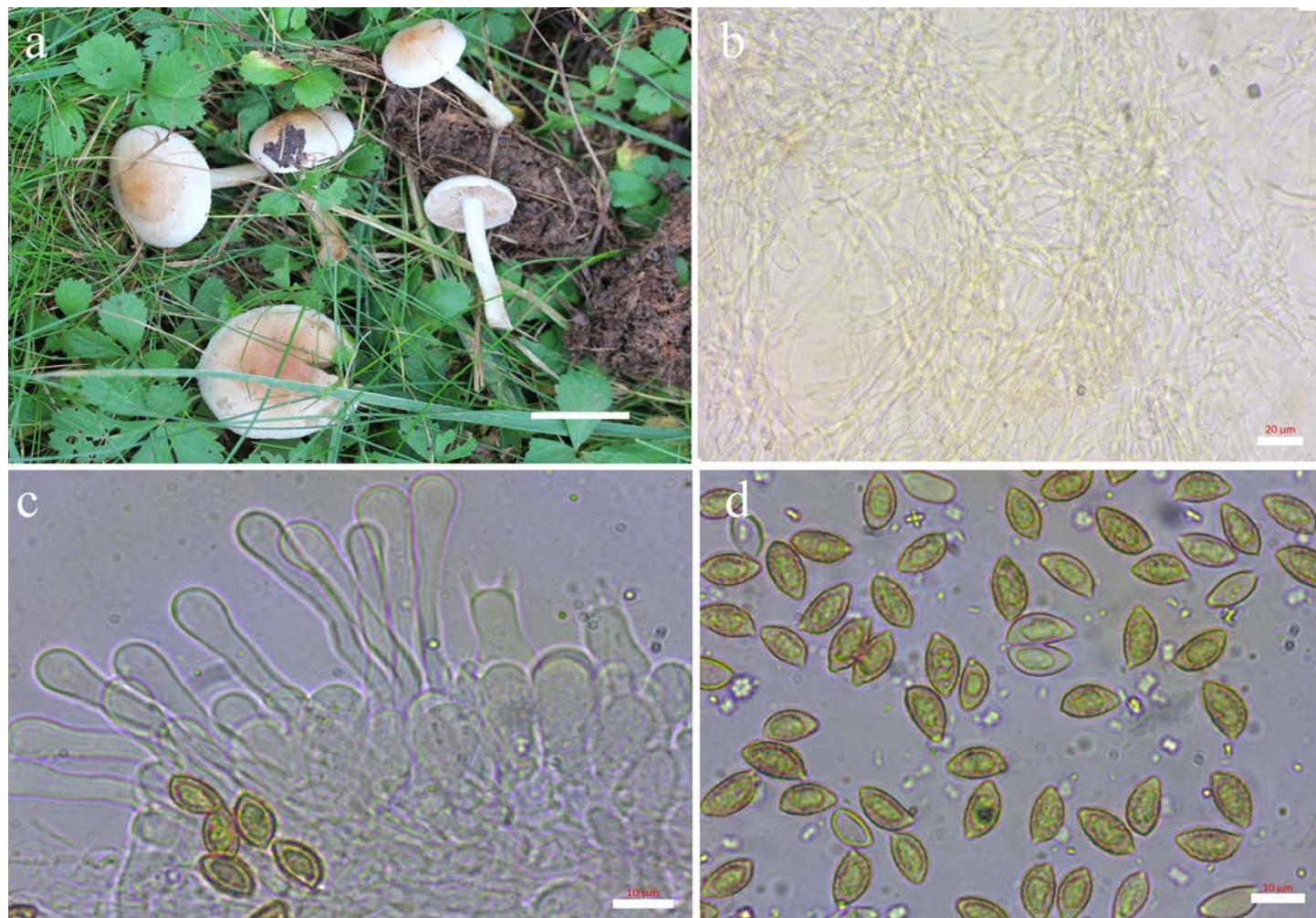
Materyal ve Metot

Araştırmancın materyali olan meyveşmiler (Şekil 1) 16.11.2018 tarihinde Karadeniz Teknik Üniversitesi park alanında saptanmış, fotoğrafları çekilmiş, önemli morfolojik özellikleri not edilmiş, standart yöntemlerle toplanarak laboratuvara getirilmiş, kurutulup etiketlenerek



fungaryum dolabına yerleştirilmiştir. Bazidiyosporların görüntülenebilmesi için şapkadan bir parça kesilerek 2 dakika %5'lik amonyak çözeltisi içerisinde tutulmuş, daha sonra bir pens yardımı ile lam üzerine alınmış ve sporlar lam üzerine dökülünceye kadar aralıklı basınç uygulanmıştır. Bazidiyum, sistityum ve şapka derisinin hifal yapısını görüntülemek için keskin jiletle binoküler mikroskop altında çok sayıda ince kesitler alınmış, %5'lik amonyak çözeltisi ile işlemen sonra Axio Imager A2 araştırma mikroskopu altında inceleme yapılmıştır.

İnceleme sırasında hücresel yapıların en ve boy ölçümleri yapılmış ve fotoğrafları çekilmiştir. Boyutların doğru olarak belirlenebilmesi için şapka derisi hücrelerinden, bazidiyumlardan ve de sistityumlardan 25'er ölçüm yapılmıştır. Teşhisler elde edilen verilerin ilgili literatür ile karşılaştırılması sonucunda yapılmıştır (Eberhardt, 2016). Koleksiyon materyali Trabzon Üniversitesi Fatih Eğitim Fakültesi'ndeki kişisel fungaryumda saklanmaktadır.



Şekil 1. *Hebeloma limbatum*: a- meyvemeler, b- şapka derisi kesiti, c-sistityumlar, bazidiyumlар ve bazidiyoller, d- bazidiyosporlar (ölçek çubukları: a: 30 mm, b: 20 µm, c ve d: 10 µm)

Bulgular

Hymenogastraceae / Papazküregiller

Hebeloma limbatum Beker, Vesterh. & U. Eberh.
[Şu eserde: Beker, Eberhardt, Vesterholt & Schütz, Fungal Biology 120(1): 83 (2016)] / Dönkturpkokan

Şapka dışbükey veya yayvan, 25–60 mm, merkeze doğru beyaz rengi, sarımsı veya turuncumsu kahverengi, bal rengi, koyu pembemsi kahverengi, soluk krem rengi, kenara doğru beyazımsı; kenarı dişli veya yivli, uzun süre içeriye doğru kıvrık, yüzeyi düz ve yapışkandır. Lameller sapa çentikli olarak bağlı veya genişliği ölçüsünde birleşik, krem, beyaz veya soluk kahverengi renkte, geniş,

kenarları damlacıklı ve kahverengi beneklidir. Dokusu dayanıklı, acı, turp kokulu, beyazımsı veya açık kahverengidir. Sap silindirik veya çomak biçiminde ve tabanda hafif soğansı, dolu, kırılgan, 15–80 × 4–15 mm; yüzeyi beyaz tozsu yapılarla kaplı, pullu veya lifli ve tabanda soluk kahverengimsidir. Bazidiyumlар çomak biçiminde, 4 sporlu, kancalı ve 20–40 × 5–15 µm'dır. Bazidiyosporlar badem veya limon şeklinde, süslü, açık sarı ve 10–14 × 5–7.5 µm'dır. Sistityumlar çomak, ıspatula veya şipe şeklinde, bazen başlıklı ve 30–100 × 5–15 µm'dır. Şapka derisi hifleri jelatinli, şeffaf, 2–9 µm,



kancalı ve düzensiz yerleşimlidir. Yapısı jelatinli üst tabaka ile kısa silindirik ve sarımsı hücrelerden oluşmuş alt tabakadan meydana gelmiştir. Yenmez.

İncelenen örnekler: Türkiye, Trabzon, Karadeniz Teknik Üniversitesi yerleşkesi, $40^{\circ}59'38.67''\text{K}$ / $39^{\circ}46'16.72''\text{D}$, hem geniş ve hem de iğne yapraklıların bulunduğu park alanında, otlar ve döküntüler arasında, çimenlik alanda, öbekler halinde, 16.11.2018, E. Sesli 4023.

Tartışma

Yeni kaydın bağlı olduğu *Hebeloma* cinsinde bazidiyokarplar *Tricholoma* veya *Collybia* tipinde, yüzeyleri yapışkan, tarçın rengi, soluk kahverengimsi, toprak veya kil rengi, beyazımsı ve kokuları genellikle iticidir. Bazı türlerin lamelleri damlacıklar oluşturur. Sapları üzerinde tozsu, pulsu veya tanecikli miselyum bulunur. İçeriği bazı türlerde turp kokusunu andırır, bazları tatsız, bazları acı ve bir kısmının kokusu pek belli değildir. *Hebeloma limbatum* ilk olarak İtalya'dan toplanarak tanımlanmış olup (Eberhardt ve ark., 2016) yeni koleksiyon 25–60 mm, merkeze doğru sarımsı, turuncumsu veya pembemsi kahverengi, kenarlarda beyaz rengi, soluk krem rengi veya beyazımsı şapka; krem veya beyaz rengi, soluk kahverengi ve damlacıklı lameller; badem veya limon şeklinde ve $10\text{--}14 \times 5\text{--}7.5 \mu\text{m}$ bazidiyosporlar; çomak veya şşe şeklinde ve $30\text{--}100 \times 5\text{--}15 \mu\text{m}$ sistityumlar ile teşhis edilir. Dış görünüş olarak benzer bir tür, *H. leucosarx*, benzer büyüklüğe ve renk tonuna sahip olmakla birlikte daha küçük bazidiyosporlara ve sistityumlara sahiptir. Ayrıca bu türde sistityumlara şişe şeklinde olmayıp her zaman çomak biçimindedir. Ayrıca bu tür söğüt ve huş altında yetişir. Bir diğer tür, *H. longicaudum* (Pers.: Fr.) Kumm. biraz daha küçük şapkaya, benzer renklere, daha uzun (60–110 mm) sapa, daha küçük ($10\text{--}13 \times 5\text{--}7 \mu\text{m}$) bazidiyosporlara, bazları dallanmış, silindir biçiminde sistityumlara sahip olmasına yeni kayıttan farklılık gösterir. Şapka rengi ve sap yapısı ile yakın gözüken bir diğer tür, *H. sinapizans* (Paul.: Fr.) Gill. iki kat büyük (50–120 mm) şapkalı, soluk kırmızımsı veya beyaz renkli, ayrık lamelli ve daha küçük ($35\text{--}70 \times 7\text{--}12 \mu\text{m}$) sistityumludur. *Hebeloma velutipes* Bruchet huş veya fındık altında yetişir, küt tepe çıkışlıdır,

bazidiyosporlarının kenar kısımları dalgalıdır. *Hebeloma senescens* (Batsch) Berk. & Mr. daha az benzerlik gösterir, genellikle iğne yapraklılar altında yetişir, yüzeyi krem rengi, soluk kırmızımsı veya benzeri renklerdedir, daha küçük bazidiyosporlar ($8.5\text{--}11.5 \times 5\text{--}7 \mu\text{m}$) üretir. Ayrıca bu türün silindirik sistityumları kıvrımlıdır. Söğüt ile mikorizal yaşayan *H. pusillum* Lge. daha küçük (10–30 mm), konik veya çan şeklinde ve belirgin tepe çıkışlı şapkaya; pembemsi tonlu beyaz lamellere; badem şeklinde fakat daha küçük ($9\text{--}12.5 \times 5\text{--}6.5 \mu\text{m}$) bazidiyosporlara sahiptir. *Hebeloma crustuliniforme* (Bull.: Fr.) Quel. konik veya çan şeklinde, tipik tepe çıkışlı, soluk turuncumsu, beyazımsı şapkaya; çentikli ve krem rengi lamellere; eliptik veya badem biçiminde ve biraz daha küçük bazidiyosporlara sahiptir. Laden altında, iğne yapraklı veya yaprak döken ağaç ormanlarında yayılış gösteren *H. cavipes* Huijsman yakın büyülüklükte olmakla birlikte, krem rengi veya soluk sarımsı şapkası, sapa tüm genişliği ile bağlı lamelleri ve daha küçük sistityumları ile *H. limbatum*dan farklılık gösterir. *Hebeloma candidipes* Bruchet genellikle ladin altında yayılış gösterir, daha küçük, kırmızımsı veya gri kahverengi şapkaya; beyazımsı veya beyaz Kahvesi ve sapa genişliği ölçüsünde bağlı lamellere; daha küçük bazidiyosporlara ve sistityumlara sahiptir. *Hebeloma bruchetii* Bon aynı büyülüklükte bazidiyokarplara sahip olmakla birlikte söğüt altında yetişir, bazidiyosporları ($8\text{--}10 \times 5\text{--}7 \mu\text{m}$) ve sistityumları ($40\text{--}60 \times 8\text{--}11 \mu\text{m}$) çok daha küçüktür (Breitenbach ve Kränzlin, 2000; Knudsen ve Vesterholt, 2008; Eberhardt ve ark., 2016).

Yazar Katkıları

Tüm yazarlar eşit katkıya sahiptir.

Çıkar Çatışması

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