ABSTRACT

The present study deals with the molecular characterization of some mushrooms collected frem Miandam, Swat Valley, KPK, Pakistan. Collected mushrooms were characterized based on molecular and phylogentic techniques using universal primers (ITS1 & ITS4) for amplification of 5.8S rRNA gene of LSU (larger subunit) after extracting DNA using most reliable methods. Eleven (11) taxa belonging to six different Genera viz. Amanita, Crepidotus, Inocybe, Mycena, Russula and Turbinellus were characterized through molecular and phylogenetic analyses of amplified target region. In this analyses their genetic variability and phylogenetic relatedness was studied. These include Amanita franchetii (Boud.) Fayod, Crepidotus sp. MD32, Mycena pura (Pers.) P. kumm., Inocybe sp. MD19, Inocybe sp. MD34, Russula sp. MD03, Russula sp. MD13, Russula sp. MD14, Russula nigricans (Bull.) Fr., Turbinellus sp. MD29, and Turbinellus sp. MD31. Out of these eleven taxa, Crepidotus sp. MD32, Inocybe sp. MD19, Inocybe sp. MD34, Turbinellus sp. MD29, and Turbinellus sp. MD31 may be new to science after complete morpho-anatomical characterization. Russula sp. MD03, Russula sp. MD13, both closely resemble to each other and with some unidentified specie sequences already present in GenBank. Thus these may also be new to science if their morpho-anatomic characterization is done. Whereas Amanita franchetii (Boud.) Fayod, Mycena pura (Pers.) P. Kumm., Russula nigricans (Bull.) Fr., and Russula sp. MD14, seem to be never described earlier from Swat Valley, KPK, Pakistan. After phylogenetic data analyses maximum intraspecific variations were observed within Turbinellus sp. MD29 (39.9%), and Turbinellus sp. MD31 (37.3%). Minimum intraspecific variations were found within the sequences of Amanita franchetii (Boud.) Fayod. (5.97%), and Mycena pura (Pers.) P. kumm. i.e (-5.17%). This study may serve as baseline for further research to explore the factors responsible for such variations and their effects on the natural adaptability of Mushrooms to our changing environment. Further research should also be taken up to find out how climatic conditions and genetic variability can influence fungal phenology and diversity.