EVALUATION OF ARMILLARIA MELEA SPECIES FEATURES CONCERNING SOME HEAVY METALS UP TAKE IN NATURAL CONDITIONS OF DIFFERENT FORESTRY ECOSYSTEMS

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Abstract

It was necessary to identify the metals (heavy, noble and rare metals) bioabsorption characteristics to the edible mushroom species Armillaria melea, in the view to relieve its quality as indicator. The samples were collected from some forestry ecosystems: Adanca, Gorgota, Manesti, Bucegi-Paduchiosu and Dambovita River valley from Dambovita County.

All over the world today are developing large studies concerning biochemical and physiological features of mushrooms in the view to promote them as biological tools in different types of environmental biotechnologies grace of their bioabsorption capacity for heavy, rare or noble metals.

This paper is about the elemental content of Armillaria melea in cap and stipe, bioabsorption factor, correlated to the mineral content of substrate and its natural pH value. The elemental content of biological and environmental samples was determined by spectrometry of fluorescence (EDXRF) using ELVA_X apparatus. Biological samples and their substrate samples have been dried at 60°C some hours first. After drying the solid samples have been grindend until to fine powder and weighed.

For the evaluation of EDXRF results was used a certified reference sample NIST SRM 1571- Orchard leaves.

So, it was determinate the presence of some heavy metals as copper, manganese, lead.

Key words: Armillaria melea, bioabsorption factor, translocation factor, pH.

MATERIAL AND METHOD

Biological samples consisted in Armillariella mellea species samples and their substrate fresh harvested from some forestry ecosystems of Dambovita County: Adanca, Gorgota, Manesti, Bucegi - Paduchiosu and Dambovita River valley. Mushrooms species where identifying using some book guides very known in Europe published after 2000 (4, 5, 14). The samples were weighted first and then dried at 60°C for some hours. After drying operation the samples were weighted again. The method for analysing was chosen carefully and not at random. The analyses were made by EDXRF because this method is not destructive and one can use the same samples to be analyzed by others methods (1, 5, 13). The elemental content of biological and environmental samples was determined using Elva-X spectrometer having a X-ray tube with Rh anode. The samples were excited for 300s and the characteristic X-rays were detected by a multichannel spectrometer based on a solid state Si-pin diode X-ray detector with a 140 mm Be window and a energy resolution of 200eV at 5.9 KeV 91,20 (2,19). Elva-X software was used to...
interpret the EDXRF spectra. The accuracy and precision of results were evaluated by measuring a certified reference sample (NIST SRM 1571-Orchard biological samples). By this method were obtained the concentration of studied metals. By these methods were registered all the quantities which were in a concentration higher than 1ppm. Every result represents the average of minimum 5 and maximum 10 determinations.

Bioabsorption factor was calculated after the following equation:

\[
Fb\% = \frac{C_m}{C_s} \times 100
\]

where: 
- \( Fb\% \) = bioabsorption factor
- \( C_m \) = metal concentration in mushroom;
- \( C_s \) = metal content of substrate.

\[
Tf = \frac{C_c}{C_s}
\]

where: 
- \( Tf \) = translocation factor;
- \( C_c \) = metal concentration in cap;
- \( C_s \) = metal concentration in stipe/soil.

RESULTS AND DISCUSSIONS

On can see from figure 1 that the pH of soil had a very important effect on the uptake of copper all over the area studied, that is releaved by the value of regression factor (0,868). pH value of soil was 6,03, easy acid in Gorgota forest and the lowest was 4,29, powerfull acide in Adanca. The greatest content of copper was found in samples of Armillariella mellea harvested from Gorgota forest (35,11ppm in cap and 21,5ppm in stipe). The lowest concentrations of copper were determined at exemplars growing on Adanca forest very closed as values from cap to stipe (13,88ppm respectively 13,4ppm). At Armillariella mellea fruitbodies from Manesti forest copper content in cap was 16,89ppm and in stipe was 14,47ppm.

On can observe a important difference between the mushrooms growing in Gorgota and Adanca.

Concerning manganese content of Armillariella mellea on can see an important accumulation in caps and stipes in all cases (figure 4). The maximum concentrations was obtained in caps of mushrooms growing in Adanca forest (292ppm) and the lowest one in exemplars from Gorgota (158ppm). Speaking about stipe content of manganese, maximum of 410ppm was in mushrooms stipes from Gorgota and minimum of 385ppm in those from Adanca. In case of manganese storage pH values were stricktly involved in, this thing beeing releavated by regression factor very closed to 1 (meaning 0,977).
Bioabsorption factor calculated for manganese was in all the others cases between 36 and 133% (figure 5). By expection, in case of mushrooms harvested from Adanca where the values were very higher, 1460% in cap and 1925% in stipe. A impressive value of translocation factor was 19 in case of mushrooms harvested from Adanca. The lowest one (0.77) was obtained at the exemplars from Manesti forest. A good translocation of 1.33 presented the mushrooms growing in Gorgota forest. So, absorption of manganese content was very different from a forest to other in all studied cases. Both pH of soil and manganese content influenced the accumulation of this metal species in high quantities inside fruiting body of these mushrooms.

Speaking about the influence of soil pH on translocation factor for manganese, on can see that it is powerfull in all studied cases (figure 6). This is sow by the values of regression factor which is over 0.800. Blue line inside graphics reprezent translocation factor for manganese.

The concentration of lead in soil was highest in Gorgota forest (2450ppm) and the minimum was determinated in mountains forest (Paduchiosu-440ppm) (figure 7). Maximum of lead content was determinated in fruiting body of mushrooms growing in mountains forest (640ppm) followed by those from Dambovita valey (630ppm) and minimum in A. mellea harvested from Gorgota forest (460ppm). Regression factor of 0.791 indicates a great influence of pH of soil on lead accumulation in mushrooms.

Bioabsorption factor for lead was under 50% at mushrooms harvested from Gorgota forest and Dambovita valey and over 100% in samples prelevated from mountains forest (figure 8).

The values of regression factor less than 0.5 shows that pH had no influence on translocation of lead into fruiting body of A. mellea, situation available in all studied cases (figure 9).

It is at least strange this uncorrelation of pH with translocation factor, because its influence on accumulation of lead is certain if we think at figure 7 which put in evidence a clear influence of pH of soil on accumulation of lead in fruiting body also in this case.
CONCLUSIONS

Copper accumulation in A. mellea was moderate, a little higher in cap than in stipe.
Manganese was stored in higher quantities, but much more in stipe than in cap.
Lead was determined in higher and very higher concentration in fruiting body.
Bioabsorption factor for copper was over 100% only in mushrooms growing in Adanca forest.
Bioabsorption factor was over 100% at mushrooms from Gorgota and over 1000% at the exemplars harvested from Adanca forest.
Translocation factor for copper was over 1 in case of mushrooms from Adanca forest.
Translocation factor was over 1 at the mushrooms from Gorgota and even 19 at those from Adanca forest.

pH had a clear influence on accumulation of copper, manganese and lead, but with differences.
In case of lead accumulation inspite of pH influence on accumulation in fruiting body, concerning translocation factor it was find any correlations.

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