IRIDOID GLYCOSIDES FROM *LINARIA GENISTIFOLIA* (L.) MILL. IN BIOLOGICAL CONTROL OF SOIL-BORNE FUNGAL PATHOGENS OF WHEAT AND SOME STRUCTURE CONSIDERATIONS

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This article is dedicated to the memory of Prof. Pavel Kintia

Abstract. Biological activity of the iridoid glycosides extract from *Linaria genistifolia* (L.) Mill. has been investigated, namely its influence on the resistance of the winter wheat Odesschi 51 plant to the caused by the *Fusarium oxysporum* and *Helminthosporium avenae* pathogenic fungi root rot. Our results indicate that summary iridoid glycosides from this plant, containing four major known compounds: 5-O-alsolslylantirrinoside, antirrinoside, linarioside and 6-β-hidroxiantirride can be successfully employed in biological control of the afore-mentioned wheat pathogens: it stimulates wheat grains germination and embryonic root growth in conditions of fungal infection. 1H and 13C NMR characteristics of 5-O-alsolslylantirrinoside in Py-d6 are for the first time presented. Structures of two conformers of 5-O-alsolslylantirrinoside in D2O and Py-d4 solutions are proposed, based on the experimental NMR evidence and molecular modelling studies.

Keywords: *Linaria genistifolia* (L.) Mill., iridoid glycosides, bioactivity, 5-O-alsolslylantirrinoside, NMR, molecular modelling.

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Introduction

More than 200 species of genus *Linaria* (Scrophulariaceae) are representatives of wild flora, constituting a rich source of biologically active substances, such as flavonoids, iridoids, alkaloids, diterpenoids, glycosides and others. A series of beneficial physiological activities characterizes these medicinal plants of repute, explaining their wide application in folk and traditional medicine, as well as in homeopathic practice. Thus, different *Linaria sp.* have been used as tonics, antiscorbutics, laxatives, antidiabetics and diuretics, spasmylytics, cholagogic drugs, as well as for treatment of wounds, bladder catarrhs, haemorrhoids and vascular disorders [1-3].

The iridoid composition of *L. dalmatica* (L.) Mill., *L. genistifolia* (var. *genistifolia* and var. *euxina*) (L.) Mill., *L. semplice* (Wild.), *L. pelisseriana*, *L. vulgaris* and *L. peloponnesiaca* Bois & Heldr., which occur in Bulgaria, has been in depth examined and reported by N. Handjieva, S. Popov et al. [4-6]. Structure elucidations have been carried out mainly by spectral methods and the data of pioneering studies on molecular mechanics calculations in this field were published [4]. An iridoid glycoside has been found amongst the chemical constituents of *L. auckeri* that is characteristic for the flora of Turkey [7]. The iridoid composition of *L. japonica* Miq., a Japanese folk medicine known as laxative and diuretic, has been investigated, detailed proton magnetic resonance examinations on the found iridoid glycosides and their derivatives being presented [8].

We have previously presented the results of our studies on biologically active compounds from *Linaria vulgaris* Mill. [9]. As an extension of this study we have now investigated the other species of *Linaria*, namely *L. genistifolia* (L.) Mill. that is widely distributed on the territory of Republic of Moldova, for the biological activity against soil-borne fungal pathogens of wheat *F. oxysporum* and *H. avenae*. These fungi are very spread, being known by severe affection of some vital processes in wheat plant, such as: seed germination, growth and development of the embryonic root and also infecting the grains with micro toxins, as well as reduction of the crop quantity and quality [10,11].

Results and discussion

This study was carried out with an objective of investigation the antifungal activity potential of the iridoid glycosides extract (IGE) from *L. genistifolia* (L.) Mill. We have studied the properties of IGE from *L. genistifolia* (L.) Mill. on the resistance of the winter wheat Odesschi 51 plant to the root rot that is caused by the *F. oxysporum* and *H. avenae* pathogenic fungi. Our results indicate that summary iridoid glycosides from this plant can be successfully employed in biological control of the afore-mentioned wheat pathogens: it stimulates wheat grains germination and embryonic root growth in conditions of fungal infection.

In IGE four major known iridoids were identified, after chromatographic separation, their presence in *L. genistifolia* (L.) Mill. being previously described [4,6]. The structures of 5-O-alsolslytirrinoside (I), which was the predominant IGE component, and 6β-hidroxiantirride (4) have been confirmed by means of NMR spectroscopy, as...
already reported [4,6], while antirrinoside (2) and linarioside (3) were identified by comparison of their physico-chemical properties with the literature data [8] (Figure 1).

The winter wheat Odesschi 51 constituted the object of the study. Prior to sowing, seeds of the winter wheat Odesschi 51 were soaked in IGE aqueous solutions having the mass fractions 10^{-1}, 10^{-4}, 10^{-3}, 10^{-2} %. As a control version the water soaked grains served. For comparison, a part of the grains was soaked in the same concentrations aqueous solutions of Moldstim – a certificated product for agricultural usage [12]. After drying the grains were soaked in F. oxysporum and H. avenae 21 days culture filtrates (CF) for 18 hours, then rinsed with distilled water and placed in a Petri dish, on filter paper moistened with distilled water. The seedlings were cultivated for 6 days at a temperature of 22ºC.

![Figure 1. Structures of compounds 1 – 4, the major IGE components, and obtained via synthesis compounds 5 and 6 [8].](image)

The plant response was assessed on the basis of its germination capacity and the length of embryonic root. The obtained results show that by infecting the grains with the F. oxysporum and H. avenae CF, the seed germination is being diminished by 33.4 and 16.7%, respectively, while the embryonic root is being reduced by 35.7 and 30.2%, as compared to the control version (Tables 1 and 2). This proves the toxicity of the fungi that causes root rot. When treating the infected grains with Moldstim, the most efficient concentrations were in the range of 10^{-4}... 10^{-3}%; it caused an increase in the germination process by 3.4 – 11.7% and 9.7 – 18.4%, and the augment of the length of embryonic root by 13.72 – 18.8% and 6.7 – 10.0%, correspondingly, in comparison with the F. oxysporum and H. avenae CF versions. It was interesting to note the same beneficial effect of IGE from L. genistifolia (L.) Mill. on the infected wheat grains and to compare the furnished by both experimental protocols results. For instance, the IGE-treated and infected with F. oxysporum plants showed a germination level of 98.3 %, which is by 35% more, and a root longer by cca 27%, when compared to F. oxysporum CF. (Table 1). These data speak in favour of higher biological activity of IGE versus Moldstim (about three times in the germination process and one and a half- in embryonic root elongation in the best versions, entries 4 and 8-9 in Table 1), the most efficacious IGE concentration being found also in the range of 10^{-4} - 10^{-3}.

### Table 1

<table>
<thead>
<tr>
<th>No.</th>
<th>Version/concentration, %</th>
<th>Germination, %</th>
<th>Embryonic root length, mm</th>
<th>% compared to CF F. oxysporum</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Control version (H2O)</td>
<td>96.7</td>
<td>80.81±1.80</td>
<td>155.5</td>
</tr>
<tr>
<td>2</td>
<td>CF F. oxysporum</td>
<td>63.3</td>
<td>51.97±2.57</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>CF F. oxysporum + Moldstim, 10^{-2}</td>
<td>63.6</td>
<td>57.08±1.65*</td>
<td>109.8</td>
</tr>
<tr>
<td>4</td>
<td>CF F. oxysporum + Moldstim, 10^{-3}</td>
<td>75.0</td>
<td>61.76±1.93*</td>
<td>118.8</td>
</tr>
<tr>
<td>5</td>
<td>CF F. oxysporum + Moldstim, 10^{-4}</td>
<td>66.7</td>
<td>59.10±2.22*</td>
<td>113.7</td>
</tr>
<tr>
<td>6</td>
<td>CF F. oxysporum + Moldstim, 10^{-5}</td>
<td>65.0</td>
<td>46.74±2.77</td>
<td>89.9</td>
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<td>7</td>
<td>CF F. oxysporum + IGE, 10^{-2}</td>
<td>96.7</td>
<td>59.59±1.82*</td>
<td>114.7</td>
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<tr>
<td>8</td>
<td>CF F. oxysporum + IGE, 10^{-3}</td>
<td>98.3</td>
<td>67.75±1.63*</td>
<td>130.4</td>
</tr>
<tr>
<td>9</td>
<td>CF F. oxysporum + IGE, 10^{-4}</td>
<td>98.3</td>
<td>64.15±1.59*</td>
<td>123.4</td>
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<tr>
<td>10</td>
<td>CF F. oxysporum + IGE, 10^{-5}</td>
<td>86.7</td>
<td>55.67±0.94</td>
<td>107.1</td>
</tr>
</tbody>
</table>

* true in relation to F. oxysporum CF at p≤0.05.
Similar effects can be observed in the presence of IGE in CF of the infected by *H. avenae* fungi grains of the winter wheat: the length of the embryonic root grows by 30.4% and germination by 13.3% (entries 9 and 8 in Table 2, IGE concentration 10^{-4} and 10^{-5}%, respectively), in comparison with the *H. avenae* CF version (entry 2, Table 2). Thus, the IGE biological properties against *H. avenae* infection are rather comparable with the activity, exerted by Moldstim (about two times higher stimulation of root elongation and 1.3 times higher promotion of germination process (entries 5/9 and 4/8 in Table 2).

**Table 2**

<table>
<thead>
<tr>
<th>No.</th>
<th>Version/concentration, %</th>
<th>Germination, %</th>
<th>Embryonic root length, mm</th>
<th>% compared to CF <em>H. avenae</em></th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Control version (H_{2}O)</td>
<td>96.7</td>
<td>80.81±1.80</td>
<td>143.3</td>
</tr>
<tr>
<td>2</td>
<td>CF <em>H.avenae</em></td>
<td>80.0</td>
<td>56.38±2.59</td>
<td>-</td>
</tr>
<tr>
<td>3</td>
<td>CF <em>H.avenae</em> + Moldstim, 10^{-2}</td>
<td>86.7</td>
<td>58.54±1.25*</td>
<td>103.8</td>
</tr>
<tr>
<td>4</td>
<td>CF <em>H.avenae</em> + Moldstim, 10^{-3}</td>
<td>90.0</td>
<td>61.82±0.98*</td>
<td>109.5</td>
</tr>
<tr>
<td>5</td>
<td>CF <em>H.avenae</em> + Moldstim, 10^{-4}</td>
<td>86.7</td>
<td>66.75±1.14*</td>
<td>118.4</td>
</tr>
<tr>
<td>6</td>
<td>CF <em>H.avenae</em> + Moldstim, 10^{-5}</td>
<td>85.0</td>
<td>59.92±2.11</td>
<td>106.3</td>
</tr>
<tr>
<td>7</td>
<td>CF <em>H.avenae</em> + IGE, 10^{-2}</td>
<td>83.3</td>
<td>65.40±1.60*</td>
<td>116.0</td>
</tr>
<tr>
<td>8</td>
<td>CF <em>H.avenae</em> + IGE, 10^{-3}</td>
<td>93.3</td>
<td>69.32±1.44*</td>
<td>123.0</td>
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<tr>
<td>9</td>
<td>CF <em>H.avenae</em> + IGE, 10^{-4}</td>
<td>83.3</td>
<td>73.52±2.40</td>
<td>130.4</td>
</tr>
<tr>
<td>10</td>
<td>CF <em>H.avenae</em> + IGE, 10^{-5}</td>
<td>90.0</td>
<td>67.9±2.04</td>
<td>120.4</td>
</tr>
</tbody>
</table>

* true in relation to *H. avenae* CF at \(p\leq0.05\).

Thus, the achieved by us results indicate that the obtained from *L. genistifolia* (L.) Mill. iridoid glycosides can be successfully utilized as substances of natural origin that stimulate wheat grains germination and embryonic root growth, considering the infected by fungi grains [13]. Our data are particularly attractive in terms of availability of the iridoid glycosides source: *L. genistifolia* (L.) Mill. grows in spontaneous flora and can be collected in large quantities.

Broadening the range of natural biologically active and environmentally inoffensive fungicidal substances is a very actual issue for agricultural workers and researchers, since biological control of insect pests, plant pathogens and weeds, is the only alternative to the use of pesticides in agriculture and forestry.

It should be mentioned, that during confirmation the stereochemical structure of compounds 1 and 4 our attention was especially drawn to the reported individual NMR spectroscopic data of compounds, which prove the reliability of structure assignments. Thus, N. Handjieva et al. have established the structure and relative stereochemistry of 6\(\beta\)-hydroxyantirride 4 on the basis of spectral studies, corroborating the NMR experimental data by molecular mechanics calculations, by using Haasnoot equation [4,14]. The minimum-energy stereostructure of 4 and its analog with OMe instead of OGlc residue indicated that the conformation of the six-membered ring of aglycon is close to a half-chair with an axially oriented substituent at C-1, which perfectly corresponded to the experimental \(J_{1,9}\) value of 2.3-2.5 Hz. The recorded by us in D_{2}O 1H and 13C NMR spectra of 4 demonstrated identical NMR characteristics of our compound to the published one [4].

The prevailing in IGE component, compound 1 is quite widespread in the plants of the genus *Linaria*, being formerly found in five *Linaria* species [6], including the investigated by us specimen. Likewise, comparison of its 1H and 13C NMR spectral characteristics, proved its identity to the published 5-O-allosylantirrinoside 1, when D_{2}O was employed as NMR solvent, Table 3 [6]. The opportunity of NMR characterization of compound 1 in deuterated pyridine (Py-d_{5}) as a weakly interacting solvent has been by us considered, by this intending to update the information on solvent effects in its NMR data profile. It should be mentioned, that Py-d_{5} presents some important advantages as NMR solvent, particularly, for investigations in the field of glycosides, namely: avoidance of the solvent interactions and signal overlapping, better solubility then in D_{2}O or MeOD etc. But its use is limited by the high cost, D_{2}O being much more attractive in terms of price.

The spectral data of compound 1, obtained after recording it’s 1H and 13C NMR spectra in Py-d_{5} were rather different from the obtained in D_{2}O solution ones (Table 3). This was not so unusual, at first appearance, since it is a matter of common knowledge, that the nature of solvent can influence both 1H and 13C chemical shifts and spin - spin coupling constants of protons [15-17]. From the other side, the noted differences could be favorable to the structural identity either of an epimeric structure, possessing the opposite stereochemistry at C_{1} (i.e. R-configuration), or, in accordance to the data of I. Kitagawa et al. [8], to a conformer of 1. Since the configuration of the glucosyloxy moiety at C_{1} in antirrinoside 2 has been previously proved to be \(\beta\) [18] (i.e., S-configuration), the possibility of examination the C_{1}
epimer of 1 has been omitted. No evidence was found in the literature, regarding the dihydropyrane ring conformation of 5-O-alsosylantirrinoside 1 [6], but the data are available for the conformers of its congeners: antirrinoside 2, linarioside 3 and some their synthetic derivatives [8]. In this light, clarifying the conformation of natural 1 has been planned, by analysis both the NMR spectroscopy and molecular modelling (MM) with energy minimization data, taking into account the traced by N. Handjieva et al. route [4].

The predominant molecule geometry of compound 2, four it’s Me, Ac, TMS derivatives and hepta-O-trimethylsilyl-linarioside 5 involved quasi-axial orientation of C_1 proton, for them the values of J_{1,9} coupling constant varying from 5.5 to 10.5 Hz, (spectra recorded in D_2O, CDCl_3 and CCl_4), as reported [8]. Whilst for linarioside 3, two it’s acetates and hexaacetilated derivative of antirrinoside 2, compound 6, the quasi-equatorial position of proton at C-1 has been assigned by the same authors on the basis of PMR evidence and Dreiding model inspection (e.g. for compound 3: 5.70 ppm (br.s.), W_{1/2}=2.0 Hz, C_1-H; for compound 6: 2.77 ppm (br.s.), W_{1/2}=4.5 Hz, C_9-H) (spectra recorded in D_2O and CDCl_3, respectively, at 60MHz) [8].

On the basis of our experimental data and taking advantage of the formulated by the Japanese chemists observation, on the relation between the splitting pattern of proton at C9 and dihydropyrane ring conformation in 2 and its derivatives [8], we explain the discussed differences in NMR data of compound 1 in two mentioned NMR solvents by conformational difference of dihydropyran ring, as well, judging by the values of J_{1,9} coupling constants: 5.48 ppm, d, J_{1,9}=6.8 Hz, C_1 ax. H, (D_2O) versus 5.84 ppm, d, J_{1,9}=3.7 Hz, C_1 eq. H, (Py-d_5), Table 3.

Interestingly, as it can be noted from the afore-mentioned, the nature of solvent proved no impact on the conformations of compounds 2, 3 and their derivatives [8]. However, in the case of the analyzed by us glycoside 1 change of the used NMR solvent causes, according to the established by us values of J_{1,9} coupling constants, switching from one conformation to another.

The minimum-energy stereo-structures for the examined conformers of 5-O-alsosylantirrinoside 1 have been obtained, which are depicted in Figure 2: 1a- in Py-d_5, and 1b- in D_2O, correspondingly, and the 'H/'H coupling constants have been calculated, by using molecular modelling with energy minimization software, namely PERCH NMR TOOLS (version 2014.1)). The results indicated on a good agreement between the calculated and experimental vicinal J_{1,9} values in pyrenone ring of conformers 1a and 1b of 1. Thus, for conformer 1a the calculated value of J_{1,9} is 3.12 Hz (experimental value 3.7 Hz), while for 1b the calculated J_{1,9} constitutes 10.84 Hz (experimental value 6.8 Hz). The minimum-energy stereo-structures of both conformers 1a and 1b indicate that the conformations of the six-membered pyrenone ring is close to a half-chair, with an axially oriented substituent at C 1 in 1a and an equatorially oriented substituent in 1b. The recorded in Py-d_5 NOE experiments have demonstrated strong NOE interactions between CH_3-10 and C_1-H, while in D_2O these interactions were less pronounced, also supporting the indicated in Figure 2 structures of the conformers 1a and 1b.

In the case of the pair of diastereoisomers of 1, hypothetic conformers of the epimer with the opposite stereochemistry at C_1 (i.e. R-configuration), the calculated values of J_{1,9} were very close, constituting 3.15 and 3.18 Hz, thus confirming the only bigger value of J_{1,9} for 1b and, implicitly, conclusions of Scarpati et al. [18].

Being occurred together in the same plant, question about conformational differences amongst compounds 1-4 in D_2O solutions is of interest also from the biogenetic point of view, since water is the solvent of life. Thus, for the natural epoxide 2 and it’s 5-O-alsosyl ether 1, belonging to the antirrinoside iridoid type, the same conformation of dihydropyran ring of aglycon is adopted in D_2O, whereas for the corresponding to epoxide 2 clorohydrine 3- the opposite one, which is characteristic also for the representative of the rare antirride type- compound 4, as depicted in Figure 1.
Thus, putting together the results of our joint NMR-MM study, it can be concluded, that the noted differences in $^1$H and $^{13}$C NMR characteristics of compound 1 demonstrate solvent influence on the molecule geometry. Our data seem not to be in the same line with that of I. Kitagawa [8], in fact, completing them. According to the MM data, in D$_2$O the conformer molecule is spatially more relaxed, being maximally available for hydration, whilst in Py-d$_5$ it is more compactly packed. Strong hydration of 5-O-allosylantirrinoside molecules, especially it’s highly hydrophilic glucose and allose moieties, most likely exerts shielding effect upon $^1$H and $^{13}$C nuclei, causing their upfield shifts in D$_2$O solution.

### Table 3

<table>
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<tr>
<th>C/H atom</th>
<th>Py-d$_5$</th>
<th>$\delta_H$</th>
<th>$\delta_C$</th>
<th>D$_2$O</th>
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<td>1</td>
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<tr>
<td>3</td>
<td>6.64 d, J=6.4</td>
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<tr>
<td>2'</td>
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<td>3'</td>
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</tbody>
</table>

*a Assignments of $^1$H and $^{13}$C signals in spectra are made on the basis of 1D: ($^1$H, $^{13}$C, DEPT-135$^\circ$) and 2D: homonuclear ($^1$H/$^1$H COSY-45$^\circ$, NOE) and heteronuclear ($^1$H/$^{13}$C HSQC, HSQC-TOCSY and HMBC) experiments.

*b 400 MHz.

c 100 MHz.

d 300 MHz [6].

e 75 MHz [6].

b,c Bruker Avance 400 spectrometer.

*Unclear signal pattern due to overlapping.

**Data are not reported.

### Conclusions

The influence of the iridoid glycosides extract from *Linaria genistifolia* (L.) Mill. has been investigated on the resistance of the winter wheat Odesschi 51 plant to the caused by *F. oxysporum* and *H. avenae* pathogenic fungi root rot. We have established, that the iridoid glycosides extract from this plant, containing four major known compounds: 5-O-allosylantirrinoside, antirrinoside, linarioside and 6-$\beta$-hidroxiantirride, can be successfully employed in biological control of the mentioned wheat pathogens: it stimulates wheat grains germination and embryonic root growth in conditions of fungal infection. $^1$H si $^{13}$C NMR characteristics of 5-O-allosylantirrinoside in Py-d$_5$ are for the first time presented. Structures of two conformers of 5-O-allosylantirrinoside in D$_2$O and Py-d$_5$ solutions are proposed, based on the experimental NMR data and molecular modelling studies.

### Experimental

**Obtaining the IGE**

IGE from *L. genistifolia* were extracted from the dried aerial part of the plant (300 g), which was chopped and then subjected to reflux (3x6 hours in a 60% methanol aqueous solution, 3x1.5L). The extracts were combined and concentrated
through vacuum distillation, thereupon, the aqueous residue was decanted with chloroform; the aqueous fraction was passed through Sephadex LH-20. The column was eluted with 10% aqueous methanol and the resulting eluate was evaporated through vacuum distillation to dryness. The dry residue (4.6g) contained a mixture of components, the iridoid glycosides 1-4 being detected in it as major products by thin-layer chromatography on silica gel. For investigation of the biological properties, IGE extract has been employed as such; 5-O-allosylanirrinoside 1 was its predominant component. For the identification of individual mixture components, a portion of it (0.6g) was separated by column chromatographic method, by using silica gel (40 x 100 μm, Merck) with a solvent systems of chloroform/methanol (4:1) and chloroform/methanol/water (95:5:0 → 10:4:1), and then Sephadex LH-20 with the elution system methanol/water (1:9). After chromatographic isolation compounds 1-4 were obtained in the following amounts: 1-52.4 mg, 2-36.6 mg, 3-32.8 mg, 4-27.5 mg. By 1H and 13C NMR spectroscopy and via comparison with bibliographic data the IGE components 1-4 were identified as follows: 1 - 5-O-allosylanirrinoside; 2 - antirrinoside; 3 - linarioside; 4 - 6-β-hidroxianteacitrill.

**Testing the biological activity of IGE**

*F. oxysporum* and *H. avenae* fungi stems were isolated from the stem of the winter wheat stem that manifested disease symptoms (brown spots), on a solid nutrient medium *Potato Dextrose Agar* (PDA) and identified through macro and microanalysis. *F. oxysporum* and *H. avenae* fungi CF were obtained by inoculating mycelium in a Czapex Dox liquid medium and cultivating it, afterwards, for 21 days at temperatures of 22°…24°C [19].

Prior to sowing, the grains were soaked for 4 hours in aqueous solutions of IGE from *L. genistifolia* (L.) Mill. or Moldstim with the mass fractions of 10⁻¹, 10⁻², 10⁻³ and 10⁻⁴ %. As a control version the water soaked grains were employed. After drying the seeds for 24 hours, part of them was soaked for 18 hours in *F. oxysporum* CF, while another part was soaked in *H. avenae* CF for 21 days then they were rinsed twice with distilled water and placed in a Petri dish on moistened with distilled water filter paper. The seedlings were cultivated for 6 days at a temperature of 22°C. The plant response was assessed based on its germination capacity and embryonic root length. For determination the germination capacity the wheat grains (100 pieces) were soaked for 24 hours in water then placed in sterile Petri dishes on moistened with distilled water filter paper. Germination was carried out in a climate chamber at a temperature of 20°C, the number of germinated seeds being determined on the 7th day. The experiments were run 5 times. Data were processed by STATISTICA 7 program.

**References**

13. Mascenco, N.; Lupascu, G.; Guriev, A.; Barba, A.; Gorincioi, E.; Gavzer, S. Procedure of treatment the winter wheat against *Fusarium oxysporum*. Decision to grant a patent Nr.7887 from 2014.08.25. (in Romanian).


