

1 Soil contaminations in landfill: a case study of the landfill 2 in Czech Republic

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12 13 **Abstract**

14 Phytotoxicity test was determined to assess ecotoxicity of landfill soil. *Sinapis alba* L. was
15 used as heavy metals bioindicator. Soil samples 1 - 8, which were taken from the landfill
16 body, edge of the landfill body and its vicinity meet the limits for heavy metals Co, Cd, Pb,
17 and Zn specified in the applicable legislation. Hg and Mn threshold values are not established
18 in legislation, but values have been determined for the needs of the landfill operator. For
19 heavy metals Cr, Cu, and Ni sample 2 exceeded the threshold values, which attained the
20 highest values of all the samples tested for Cr, Cu and Ni. For Cr and Ni the values were
21 several times higher than values of the other samples. The second highest values for Cr, Cu,
22 and Ni showed sample 6 and 7. Both samples exceeded the set limits. An increase in plant
23 biomass was observed in plants growing on plates with soil samples, but no changes in
24 appearance, slow growth or necrotic lesions appeared. Ecotoxicity tests show that tested soils
25 (concentration of 50%) collected from the landfill body, edge of the landfill body and its
26 vicinity reach high percentage values of germination capacity of seeds of *Sinapis alba* L.
27 (101-137%). At a concentration of 25%, tested soil samples exhibit lower values of
28 germination capacity; in particular samples 3 to 8, yet the seed germination capacity in all 8
29 samples of tested soils range between 86 and 137%.

1 1 Introduction

2 Land degradation caused by human activities creates significant adverse effects on the
3 environments and ecosystems worldwide (Thomaz and Luiz, 2012; Bai et al., 2013; Li et al.,
4 2013; Chen et al., 2015) and solid waste is an important and emerging environmental
5 problem. It was estimated that 0.5–4.5 kg per person per day of solid waste is produced in
6 different regions of the world (Bakare et al., 2005; Swati et al., 2014). The most common
7 ways to manage such waste disposal are landfills and incinerators. Actually up to 95% total
8 municipal solid waste (MSW) collected is disposed of in landfills worldwide (El-Fadel et al.,
9 1997; Swati et al., 2014) and landfilling is the major MSW disposal method used in modern
10 cities (Wong et al., 2015). Landfills were thought to be the safe disposal method of MSW but
11 it is true only for properly engineered landfill sites. An engineered landfill site allows final
12 disposal of solid waste in a secure manner by minimizing the impacts on the environment as
13 modern landfills are often lined with layers of absorbent material and sheets of plastic to keep
14 pollutants from leaking into the soil and water (Swati et al., 2014).

15 The improper management of waste disposal raises public concern over potential harmful
16 effects to local communities and the environment. These concerns probably become more
17 pragmatic when recent intensive studies demonstrated increased human health risk caused by
18 exposure to toxic chemicals, such as dioxins and related compounds, and heavy metals in
19 these dumping sites (Agusa et al., 2003, Minh et al., 2003). Landfills containing hazardous
20 materials are under critical observation today for potential hazards, resulting in the need for
21 thorough risk analyses along with the soil and groundwater that have been contaminated with
22 chemicals leaching from landfills. Several reports have been published which are documented
23 on the leachate characterization and its effect on groundwater pollution (Boels and Fleming,
24 1993) but little information is available on the effect of landfills on the soil contamination
25 (Hernández et al., 1996) and its toxicological effects.

26 Soil is the key part of the Earth System as it control the hydrological, erosional, biological and
27 geochemical cycles. Soil System is also offering goods, services and resources to the
28 humankind (Keesstra et al., 2012, Mol and Keesstra, 2012, Berendse et al., 2015, Brevik et
29 al., 2015, Decock et al., 2015, Smith et al., 2015). This is why it is necessary to research how
30 the soils are affected by the use by the human societies. Pollution is one of those damaging
31 human activities and we need more information and assessment of the land pollution

1 [\(Kardanpour et al., 2015, Mahmoud and El-Kader, 2015, Riding et al., 2015, Roy and](#)
2 [Mcdonald, 2015, Sacristánet al., 2015, Wanget al., 2015\).](#)

3 Land and sSoil pollution by heavy metals has become a critical environmental concern due to
4 its potential adverse ecological effects. Heavy metals occur naturally at low concentrations in
5 soils. However, they are considered as soil contaminants due to their widespread occurrence,
6 acute and chronic toxicity (Youn-Joo, 2014).

7 More recently high concentrations of heavy metal(loid)s, such as As, Cd, Cu, Pb, and Zn in
8 soils have often been reported in number of countries. For example, significant adverse
9 impacts of As on human health have been recorded in Bangladesh, India, and China and it is
10 claimed that millions of people are potentially at risk from As poisoning (Bhattacharya et al.,
11 2012). Similarly, Cd accumulation in the offal of grazing animals in New Zealand and
12 Australia made it unsuitable for human consumption and affected access of meat products to
13 overseas markets (Loganathan et al., 2008). Similarly, there have been concerns about urban
14 development of horticultural sites which contained toxic levels of metal(loid)s such as As, Cu,
15 and Pb in soils resulting from excessive use of fungicides and herbicides that are rich in these
16 metal(loid)s (Pietrzak and Uren, 2011).

17 Plants can be used as bioindicators for toxicity assessment in aquatic and terrestrial
18 ecosystems (Gorsuch et al., 1991). The present research was aimed at assessing the soil
19 pollution at the landfill site (in operation) and in the vicinity of a MSW landfill site. The main
20 objective of this study was characterization of soil samples issued from a landfill located near
21 Klatovy, in south-western Czech Republic, in relation to their content of heavy metals. The
22 other objective was to recommend some sensitive plant to assess phytotoxicity effect on one
23 vegetal specie. White mustard (*Sinapis alba* L.) was selected as the test plant species due to
24 their sensitivity to a wide range of contaminants. To assess phytotoxicity of landfill soil a
25 laboratory study was conducted.

27 **2 Material and methods**

28 **2.1 Site description**

29 The investigated landfill (Štěpánovice, 49°26'15.934"N, 13°16'55.352"E, ca. 405 m asl) is
30 located in Pilsen Region, western part of the Czech Republic. It started operating during 1996

1 with an authorized volume of 569000 m³, at the moment, it is being used to dispose mixed
2 municipal waste. The landfill is formed by three sub-landfills: landfill A (closed in 2003, area
3 8750 m²); landfill B (working from 2003, area 26000 m²); landfill C (that will work after
4 closing part B). The total volume of both (A, B) parts of the landfill is 289000 m³. Planned
5 service life of the facility is up to year 2018 (Vaverková and Adamcová, 2014a).

6 Every day, up to 37.5 tonnes of waste is authorized for landfilling after careful analysis: the
7 disposed waste includes municipal solid, non-hazardous wastes and the material for landfill
8 cover. Wastes may include scraps of paper, plastics and metals, packing, spent tires, textile
9 products, building materials, ashes from MSW incinerators, polluted terrain from
10 environment reclamation, etc. Particular details of waste composition, waste quantity stored
11 on landfill and landfill gas management are not presented in this article. Detailed information
12 and data were described in other articles (Vaverková and Adamcová, 2014a; Vaverková and
13 Adamcová, 2014b; Vaverková and Adamcová, 2014c).

14 The landfill site is located over an impermeable natural clay layer; bottom and side
15 boundaries may vary according to the period of cultivation, however they generally include
16 several protective layers, such as a compact clay layer (100 cm), geotextile membranes,
17 gravel (50 cm), geomembranes (2.5 mm) non-woven fabric (1200 g/m²), pulper products.

18 Landfill covers (top and side) are formed by a waste layer (terrain) to stabilize the surface,
19 drainage systems, compact clay (20 cm), soil bentonite and a vegetative soil layer (up to
20 100 cm). A grassy mantle and/or forestation with local vegetation will complete the recovery
21 of the environment after closing of each parcel. Systems for leachate treatment, and gas
22 recovery, collection and treatment are in operation. The landfill is situated in the north part of
23 widely opened valley directed towards W-E. The landfill is surrounded to the N and S by a
24 vegetation belt dominated by *Pinus sylvestris*. The hilly landscape in the western part of the
25 study area is used for agriculture, as well as the eastern lowland. The climate of the area is
26 typically inland, with mean annual rainfall over 582 mm and mean annual temperature of
27 8.0°C (Vaverková and Adamcová, 2015).

28

29 **2.2 Sample collection**

30 Soil samples were collected from landfill site at depth 10 cm (Fig. 1) [in 2014](#). They were
31 collected in sterilized plastic containers. Freeze and grounded soil samples were homogenized

1 by sieving through a stainless steel 0.2-mm sieve, and stored in sealed containers at -4 °C
2 until analysis. The materials were analyzed for the content of heavy metals (Hg, Cd, Pb, Cu,
3 Zn, Co, Ni, Cr, Mn). The examined samples were brought to the testing laboratory
4 (Department of Chemistry and Biochemistry, Faculty of Agronomy, Mendel University in
5 Brno) for analyses.

6 Fig. 1 provides sampling points where soil samples were collected. In total 8 sampling points
7 were determined. Samples collected from sampling points 6, 7 and 8 were used as blind
8 samples. Samples from sampling points 4 and 5 were collected directly from the landfill body
9 and samples 1, 2 and 3 were taken from the edge of the landfill body. The allocations of
10 sampling sites were chosen on the basis of the authors` decision and on the grounds of mutual
11 comparison of the landfill body and its borders with the nearest vicinity of the landfill
12 (agriculturally utilized soil and forests).

13

14 **2.3 Sample processing and chemical analysis**

15 A microwave digestion (Ethos SEL, Milestone, Italy) was used for isolation of analytes from
16 solid samples. Soil samples were air-dried and sieved. A fraction < 2 mm was used for the
17 analysis. 200 ± 0.1 mg of dried and homogenized soil samples was used for partial digestion
18 in the microwave oven using 3 ml of concentrated HNO₃ and 9 ml of concentrated HCl at 200
19 °C and 1000W for 30 min. The soil digests were adjusted to the final volume of 25 ml with
20 deionized water.

21 Electrothermal atomic-absorption spectrometer (AAS ZEE nit 60, Analytic Jena, Germany
22 equipped with Zeeman correction) was used under the recommended conditions specified by
23 the manufacturer for determination Cd (228.8 nm), Pb (283.3 nm), Co (240.7 nm), Cr (359.3
24 nm). The wavelengths are given in parentheses. 1% Pd/Mg(NO₃)₂ was used as modifier.

25 Flame atomic-absorption spectrometer (AAS ZEE nit 60, Analytic Jena, Germany equipped
26 with Zeeman correction) was used under the recommended conditions specified by the
27 manufacturer for determination Cu (324.7 nm), Zn (213.8 nm), Ni (232.0 nm), Mn (279.5
28 nm). Acetylene-air flame was used for determination of analytes. The wavelengths are given
29 in parentheses.

1 Total mercury content in soil samples was measured by one purpose atomic absorption
2 spectrometer AMA 254 (Advanced Mercury Analyzer) controlled by WinAMA software
3 (both Altec, Prague, Czech Republic). The homogenized solid samples were weighted ($100 \pm$
4 0.1 mg) into pre-cleaned combustion boats and inserted into the AMA254 analyzer. During
5 analysis the sample was dried at 120 °C for 90 s and thermally decomposed at 550 °C for 180
6 s under an oxygen flow. Selectively trapped mercury was subsequently released from the gold
7 amalgamator by a brief heat-up and finally quantified (measuring cycle, 60 s) as Hg⁰ by the
8 cold-vapor AAS technique at 253.65 nm.

9 LODs (limit of detection) of methods were 0.1 µg/kg for Hg, 0.02 µg/kg for Cd, 0.38 µg/kg
10 for Pb, 3.08 µg/kg for Cu, 3.70 µg/kg for Zn, 4.92 µg/kg for Co, 9.00 µg/kg for Ni, 0.70
11 µg/kg for Cr, and 12.10 µg/kg for Mn. The results were in good agreement with the certified
12 values.

13 **2.4 Test plant species**

14 The test species were white mustard (*Sinapis alba* L.). They were selected because they are
15 known to be sensitive to board range of chemicals. White mustard is ideal for studying soils
16 and soil extracts (Gerencsér et al, 2010; OECD Guideline 208 for the Testing of Chemicals,
17 2003). Seeds were surface-sterilized by soaking for 2 min. in a commercial sodium
18 hypochlorite (2%) solution to which a few drops of Tween-20 had been added. Then they
19 were rinsed twice in sterile distilled water. Damage or empty seeds hulls were discarded.

20 **2.5 Phytotoxicity test**

21 The earthen pot experiment was performed under laboratory conditions ~~(Fig. 2)~~. The earthen
22 pots (height of 10 cm and a diameter of 11 cm) were loosely filled with 200 g of medium,
23 than 100 seeds of white mustard were scattered on to the surface, covered with thin layer of
24 silica sand and covered with a glass plate (to avoid evaporation). The possible toxicological
25 effect was assessed according to CSN EN 13432 on growth of dicotyledonous plants. The
26 medium was specialized soil for germination and plant growth, enriched with soil samples
27 (25 %, 50 % w/w). Reference soil was composed from peat and silica sand. Plants were
28 grown under controlled conditions for 21 days. Humidity at level of 70 ± 25 % of water
29 absorption capacity was maintained to be constant. The toxicity tests were conducted at
30 ambient laboratory temperature of 22 ± 10 °C, continuous light was used. Values obtained

1 from two simultaneously conducted experiments were averaged and presented (germination
2 capacity, plant biomass).

3

4 **3 Results and discussion**

5 From the chemical analysis of solid samples with atomic absorption, the leaching values of
6 metals varied over a wide range as follows: Hg (0.0300 – 0.0663 mg/kg), Cd (0.0600 –
7 0.2044 mg/kg), Pb (2.5703 – 8.5287 mg/kg), Cu (32.43 – 51.24 mg/kg), Zn (25.67 – 41.97
8 mg/kg), Co (2.953 – 12.712 mg/kg), Ni (32.65 – 140.03 mg/kg), Cr (64.06 – 190.73 mg/kg) a
9 Mn (547.52 – 701.39 mg/kg). The **average** metal leaching values found in this study are
10 shown in Table 1.

11 Fig. **23** shows the graphically evaluated results of heavy metal content in individual soil
12 samples, with marked limit values for those heavy metals for which the limits are established
13 in the Decree of the Ministry of the Environment of the Czech Republic No. 13/1994 Coll., as
14 amended, laying down the details of agricultural land fund protection.

15 For heavy metals Cd, Pb, Zn, and Co, none of the eight examined soil samples did exceed the
16 limits specified in the Decree, as shown by the charts listed on Figure 3. For Cu, Ni, and Cr
17 some of the collected soil samples exceeded the limits established in the relevant legislation.
18 For Cu, the maximum limit for this heavy metal is 50 mg/kg. Samples that exceeded the limit
19 for Cu were as follows: sample 2 (58.62 mg/kg), sample 6 (51.24 mg/kg), and sample 8
20 (50.20 mg/kg). For Ni, the maximum limit is set at 25 mg/kg. This threshold value was
21 exceeded by all 8 soil samples; the highest value was measured in sample 2 (140.03 mg/kg).
22 The maximum allowed value for Cr (40 mg/kg) was exceeded by all 8 samples, with sample 2
23 showing the highest value (190.73 mg/kg). No limit values are established for Mn and Hg
24 presence in the soil. After the levels of heavy metals in collected soil samples were
25 determined, the phytotoxicity was tested.

26 Fourteen days from the establishment of the experiment, sprouts and the number of growing
27 plants occurring in the earthen pots were counted. The data were plotted into tables and
28 photographs were taken to document the course of the experiment. Germinating capacity and
29 growth of white mustard is shown in Fig. **34**. Twenty-one days from the establishment of the
30 experiment, the counting of sprouts and growing plants was repeated, the results were
31 recorded and photographs were taken.

1 Values were calculated from the obtained data (Table 2) and results were evaluated. The
2 number of sprouts (number of growing plants) occurring on samples of examined soil and on
3 the soil from the blank experiment was compared for all mixing ratios. Germinating capacity
4 was calculated as a percentage of the corresponding values obtained from soils in the blank
5 experiment.

6 Table 23 lists average values calculated from the results obtained after conducting the
7 experiment (see Table 2) as well as percentages of germination capacity for each sample of
8 examined soil.

9 Fig. 45 shows the percentage expression of germination capacity of seeds of white mustard
10 (25% share of soil of samples 1-8) after 14 days from the start of the experiment and after 21
11 days (end of the experiment).

12 Maximum germination capacity of seeds of white mustard at a concentration of 25% was
13 achieved for sample 1, both in the period of 14 days (139%) and after 21 days (137%). The
14 second highest value exhibited sample 2 (131 % after 14 days and 136% after 21 days). The
15 third highest values were measured for samples 3 and 8, where after 14 days the germination
16 capacity reached 106% and 111%, respectively; and after 21 days the germination capacity
17 was 110% and 107%, respectively.

18 Even sample 4 exhibits high values of germination capacity after 14 days (102%) and after 21
19 days (103%). The 100% germination capacity limit was approached also by sample 5 after the
20 period of 21 days when the seed germination capacity attained 100%. The lowest values of
21 germination capacity of white mustard seeds showed samples 6 and 7. Germination capacity
22 of sample 6 was 85% after 14 days and 91% after 21 days, and that of sample 7 was 80% after
23 14 days and 86% after 21 days.

24 Fig. 56 shows the percentage expression of germination capacity of seeds of white mustard
25 (50% share of soil of samples 1-8) after 14 days from the start of the experiment and after 21
26 days (end of the experiment).

27 At a concentration of 50%, all samples (Samples 1-8) reported seed germination capacity
28 values over 100%, once after 14 days and again after the 21 day period. The highest values of
29 germination capacity occurred in sample 1 after 14 days (138%) and after 21 days (133%).
30 The second highest germination capacity was observed in sample 5, where it reached 123%

1 after 14 days and 122% after 21 days. The third place in germination capacity of white
2 mustard seeds was occupied by sample 7 (122% after 14 days and 119% after 21 days).

3 An increase in plant biomass was observed in plants growing on plates with soil samples from
4 the landfill body and its vicinity, but no changes in appearance, slow growth or necrotic
5 lesions appeared. Ecotoxicity tests show that tested soils (at a concentration of 50%) collected
6 from the landfill body, edge of the landfill body and its vicinity reach high percentage values
7 of germination capacity of seeds of white mustard (101-137%) (Fig. 67). At a concentration
8 of 25%, tested soil samples exhibit lower values of germination capacity; in particular
9 samples 3 to 8, yet the seed germination capacity in all 8 samples of tested soils range
10 between 86 and 137%.

11 The analysis of the variance is listed in Table 34. P(ANOVA) was calculated using the *Maple*
12 software. P-value determines the significance level, where it is possible to reject the
13 hypothesis that both models used are equivalent. P-value is compared with a pre-chosen
14 constant (most commonly 0.05) and when it is smaller, the equivalence of the models is
15 rejected. Three cases where the assumption is that the behaviour of the samples is different
16 from the behaviour of the blanks by 5% significance are marked in Table 34. Four
17 measurements were provided for each sample – two concentrations and two germination
18 rates. 4 values of p are available for each sample. Not one sample can be discarded in most
19 cases, see Fig 78. The values of p factor (ANOVA) for germination after 14 days are plotted
20 on the x-axis of Fig 78, the values of p factor (ANOVA) for germination after 21 days are
21 plotted on the y-axis. The green area is the requirement for equivalent germination – H_0 for
22 samples and blanks positively satisfied on the standard range of significance 0,05 – 5%. The
23 pink areas indicate the failure to satisfy this condition for one of the germination rates (14 or
24 21) days. Sample 5 is located in this area, but it is just below the line for the 21-day
25 germination rate, it satisfies the 14-day germination rate. Sample 1 is also located in this area
26 for the germination rate of 21 days but only for 25% concentration. No samples are located in
27 the red area where hypothesis H_0 can be positively rejected. Due to this it is possible to
28 consider the assumption H_0 are not significantly affected by the landfill.

29

30 **4. Conclusions**

31 Phytotoxicity test was determined to assess ecotoxicity of landfill soil. Fourteen days after the
32 establishment of the experiment, sprouts and the number of growing plants occurring in the

1 earthen pots were counted. The data were plotted into tables and photographs were taken to
2 document the course of the experiment. Twenty-one days from the establishment of the
3 experiment, the counting of sprouts and growing plants was repeated, the results were
4 recorded and photographs were taken. Results were evaluated from the acquired data. The
5 number of sprouts (number of growing plants) on the soil samples and on the soil from the
6 blank experiment was compared for all mixing ratios. Germinating capacity was calculated as
7 a percentage share of corresponding values obtained from the soil in the blank experiment.
8 Results in the tables (germinating capacity of seeds) are mean values obtained from the
9 conducted experiment.

10 Plant growth test can be good protocol to assess the phytotoxicity of soil contaminated by
11 heavy metals. White mustard is sensitive plant that can be used as heavy metals bioindicator.
12 Soil samples 1 to 8, which were taken from the landfill body, edge of the landfill body and its
13 vicinity meet the limits for heavy metals Co, Cd, Pb, and Zn specified in the applicable
14 legislation. Hg and Mn threshold values are not established in legislation, but values have
15 been determined for the needs of the landfill operator. For heavy metals Cr, Cu, and Ni some
16 samples exceeded the threshold values, namely sample 2, which attained the highest values of
17 all the samples tested for Cr, Cu and Ni. For Cr and Ni the values were several times higher
18 than values of the other samples.

19 After sample 2, the second highest values for Cr, Cu, and Ni showed sample 6 and also
20 sample 7, this one particularly for Cr and Ni. Both of these samples exceeded the set limits,
21 but their measured values were not as high as in the case of sample 2.

22 An increase in plant biomass was observed in plants growing on plates with soil samples from
23 the landfill body and its vicinity, but no changes in appearance, slow growth or necrotic
24 lesions appeared. Ecotoxicity tests show that tested soils (at a concentration of 50%) collected
25 from the landfill body, edge of the landfill body and its vicinity reach high percentage values
26 of germination capacity of seeds of white mustard (101-137%). At a concentration of 25%,
27 tested soil samples exhibit lower values of germination capacity; in particular samples 3 to 8,
28 yet the seed germination capacity in all 8 samples of tested soils range between 86 and 137%.

29

30 **Author contribution**

1 D. Adamcová, M.D. Vaverková, Z. Havlíček and E. Břoušková designed the experiments and
2 D. Adamcová and M.D. Vaverková carried them out. S. Bartoň performed the analysis of the
3 variance. M.D. Vaverková prepared the manuscript with contributions from all co-authors.

4

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12

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9

10

11 Table 1. Content of heavy metals in examined soil samples.

Sample	Hg (mg/kg)	Cd (mg/kg)	Pb (mg/kg)	Cu (mg/kg)	Zn (mg/kg)	Co (mg/kg)	Ni (mg/kg)	Cr (mg/kg)	Mn (mg/kg)
1	0.0300	0.0670	6.5554	34.07	25.67	4.888	39.13	69.99	689.55
2	0.0311	0.1446	2.5703	58.62	34.20	12.712	140.03	190.73	608.67
3	0.0343	0.1446	5.1769	39.37	31.51	2.953	32.65	65.92	547.52
4	0.0663	0.1576	8.5287	34.25	39.29	5.825	36.94	64.06	610.10
5	0.0403	0.1343	5.1095	32.43	33.93	10.284	33.11	70.10	584.58
6	0.0386	0.2044	5.4088	51.24	41.97	6.874	44.05	86.69	625.12
7	0.0459	0.0600	5.0800	43.80	32.10	5.375	42.76	74.85	661.00
8	0.0312	0.1471	4.1255	50.20	31.68	5.469	37.59	69.94	701.39

12

1 ~~Table 2. Results for germination capacity of seeds of white mustard for examined samples.~~

Sample	Summary—germination test	
	14 days	21 days
-		
1A-25	79	82
1B-25	91	95
1A-50	99	99
1B-50	100	100
2A-25	88	95
2B-25	72	81
2A-50	83	88
2B-50	86	88
3A-25	61	68
3B-25	68	74
3A-50	81	85
3B-50	91	93
4A-25	66	70
4B-25	59	63
4A-50	63	65
4B-50	92	96
5A-25	53	54
5B-25	68	75
5A-50	91	94
5B-50	86	89
6A-25	48	55

6B-25	56	62
6A-50	82	87
6B-50	62	65
7A-25	56	61
7B-25	41	50
7A-50	84	87
7B-50	91	92
8A-25	57	59
8B-25	78	79
8A-50	84	86
8B-50	85	89
Blank-I	69	72
Blank-II	75	78
Blank-III	70	73
Blank-IV	52	56

1 *A ——— 1 ——— performance, ——— B ——— second ——— performance

- 1 Table 23. Average values and percentages of germination capacity of seeds of white mustard
 2 for examined samples.

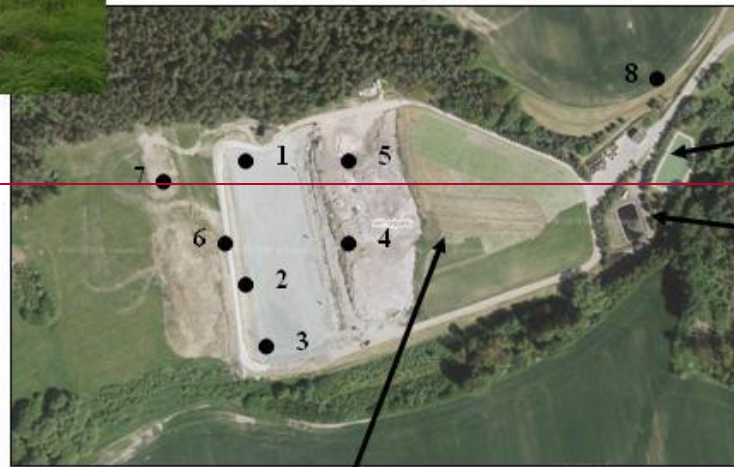
Sample - Mean	Summary - germination test		% number of seeds germinated	
25%	14 days	21 days	14 days	21 days
1	85	88.5	139	137
2	80	88	131	136
3	64.5	71	106	110
4	62.5	66.5	102	103
5	60.5	64.5	99	100
6	52	58.5	85	91
7	48.5	55.5	80	86
8	67.5	69	111	107
Blank	61	64.5	100	100
50%	14 days	21 days	14 days	21 days
1	99.5	99.5	138	133
2	84.5	88	117	117
3	86	89	119	119
4	77.5	80.5	108	107
5	88.5	91.5	123	122
6	72	76	100	101
7	87.5	89.5	122	119
8	84.5	87.5	117	117
Blank	72	75	100	100

1 Table 34. Analysis of the variance.

		25%			
Sample A, B	14 days		21 days		
	Germ.	p(ANOVA)	Germ.	p(ANOVA)	
1	75, 91	0.091	82, 95	0.012	
2	88, 72	0.654	95, 81	0.076	
3	66, 68	0.811	68, 74	0.074	
4	66, 59	0.636	70, 63	0.398	
5	53, 68	0.533	54, 75	0.045	
6	48, 56	0.140	55, 62	0.601	
7	56, 41	0.110	61, 50	0.055	
8	57, 78	0.924	59, 79	0.075	
		50%			
Sample A, B	14 days		21 days		
	Germ.	p(ANOVA)	Germ	p(ANOVA)	
1	99, 100	0.084	99, 100	0.617	
2	83, 86	0.094	88, 88	0.063	
3	81, 91	0.874	85, 93	0.064	
4	63, 92	0.686	65, 96	0.417	
5	91, 86	0.535	94, 89	0.041	
6	82, 62	0.206	87, 65	0.559	
7	84, 91	0.146	87, 92	0.054	
8	84, 85	0.940	86, 89	0.070	
Blank	Germ. 14 days		Germ. 21 days		
I,II, III, IV	69, 75, 70, 52		72, 78, 73, 56		



Detail of the landfill



Rainwater reservoir

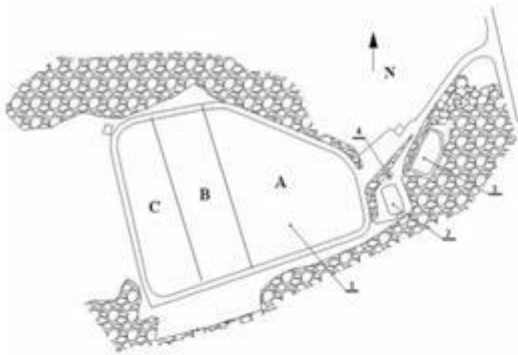
Leachate pond

● Sampling points

Landfill

1

2



1 - landfill, 2 - detention receiver of leachate, 3 - rainwater reservoir, 4 - entrance gate

Section of the landfill



Rainwater reservoir

Leachate pond

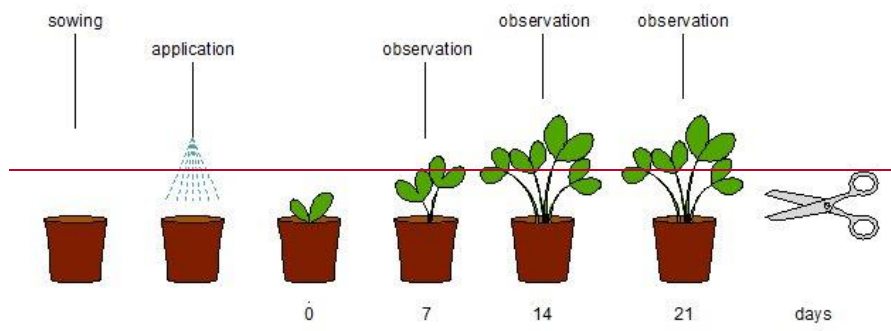
● Sampling points

Landfill

3

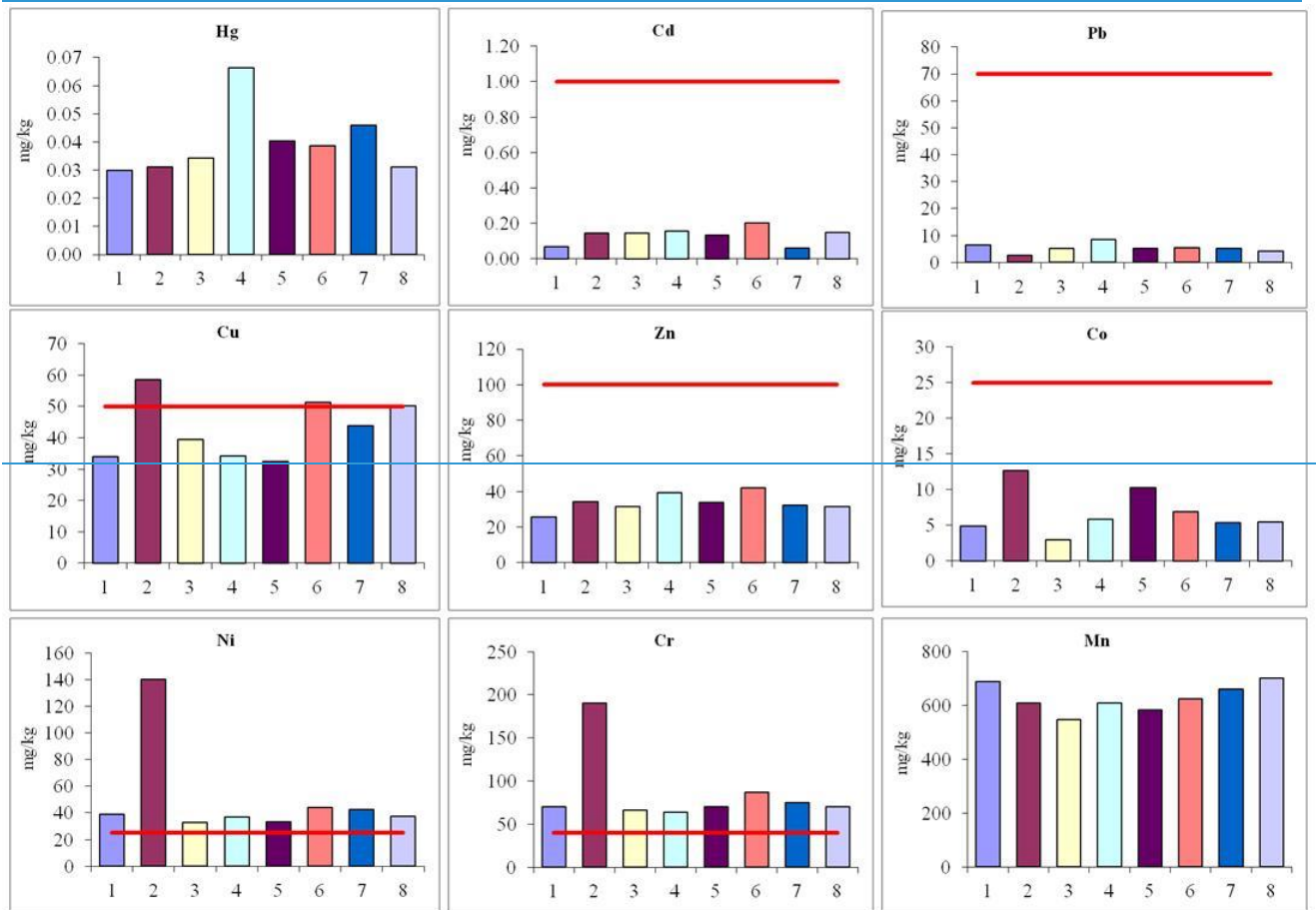
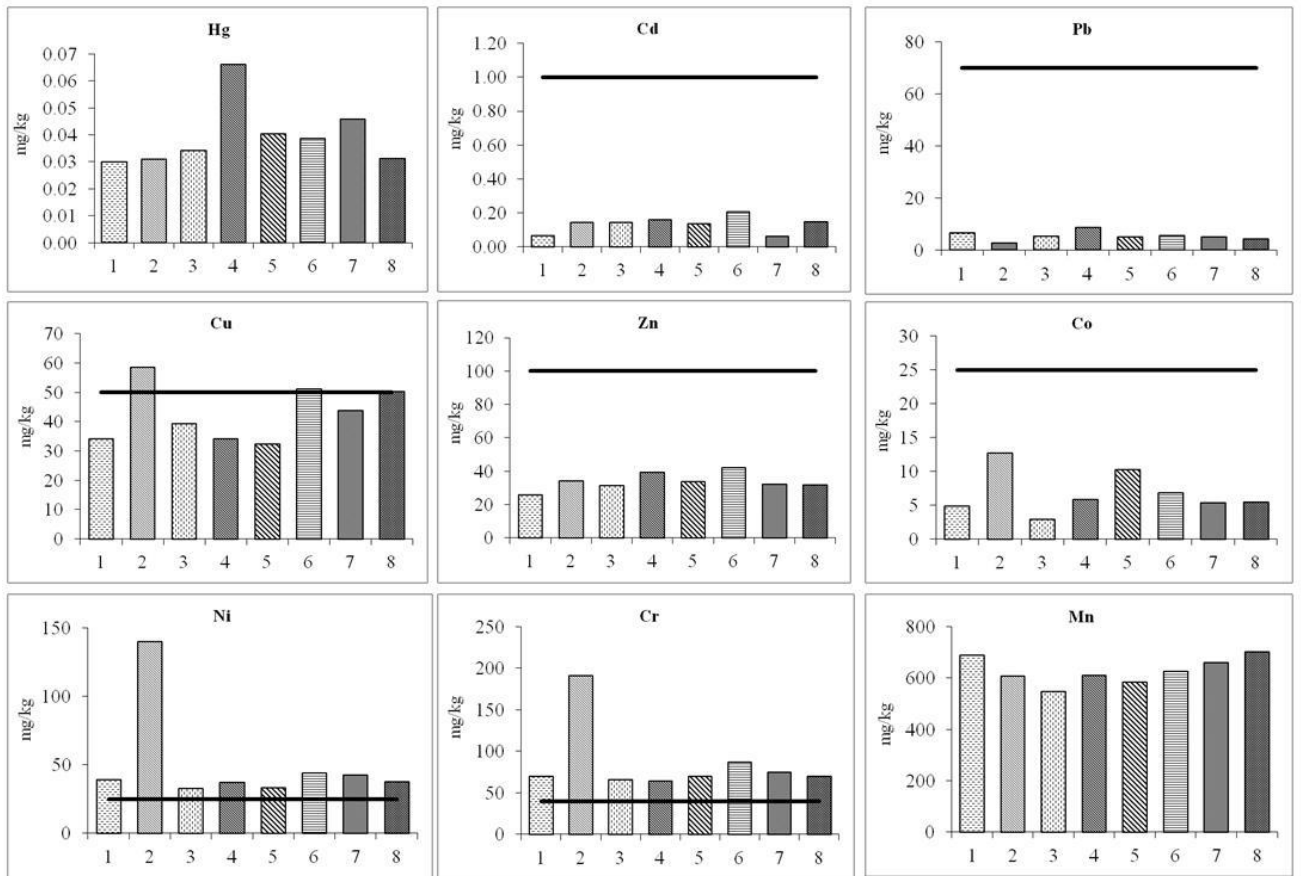
1 Figure 1. Map of Štěpánovice landfill and sampling points.

2



- 1
- 2
- 3
- 4

Figure 2. Layout of the phytotoxicity test.



1 Mn, Hg – no threshold values are set in the Decree No. 13/1994 Coll.

2

3 Figure 23. Content of heavy metals in examined soil samples with marked limit values set in

4 the Decree No. 13/1994 Coll.:-

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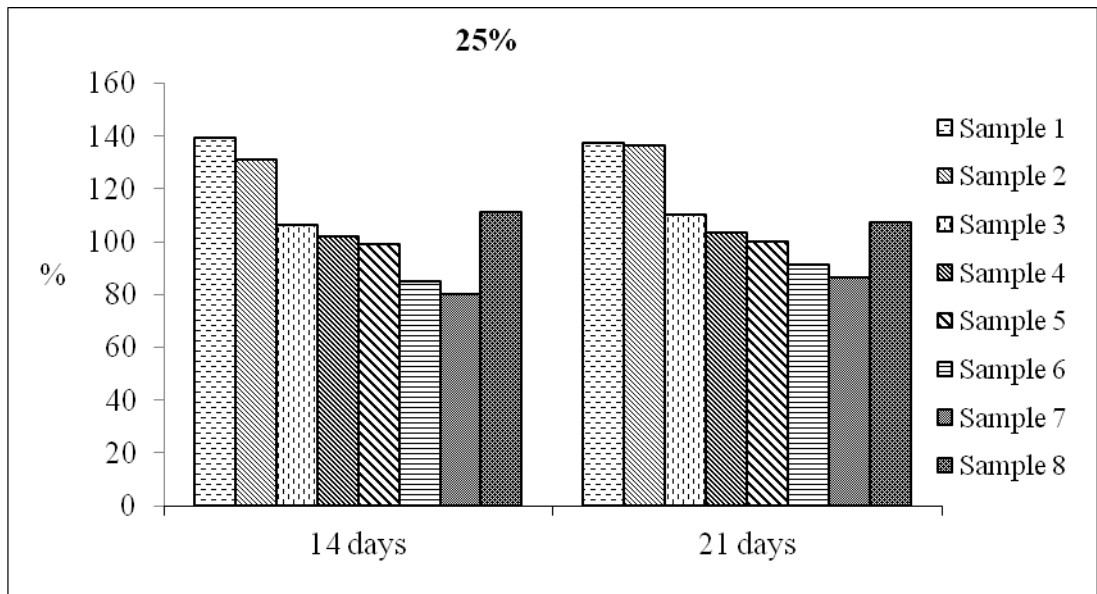


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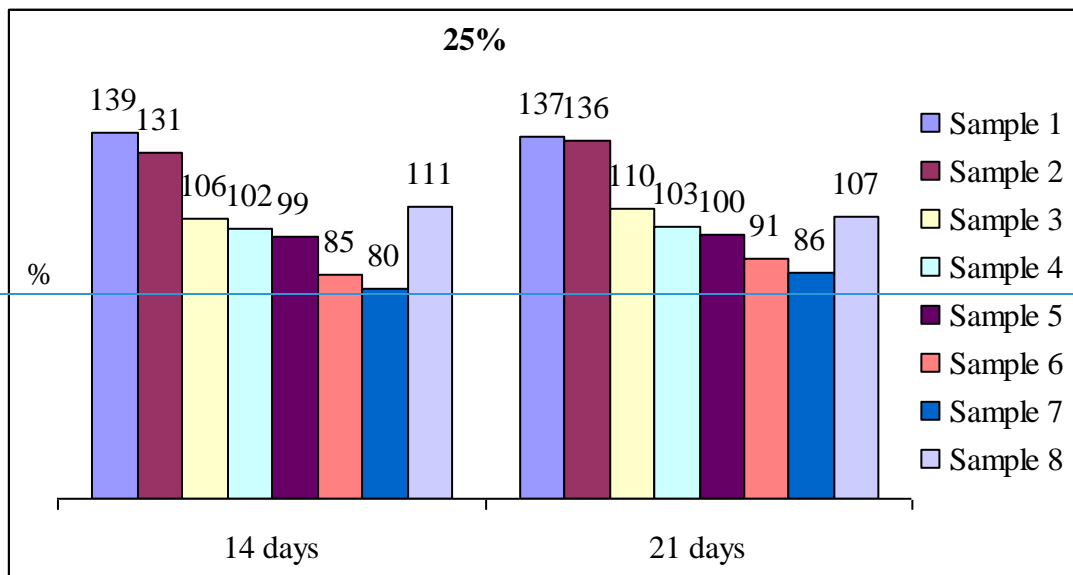
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4 Figure 34. Samples of white mustard.

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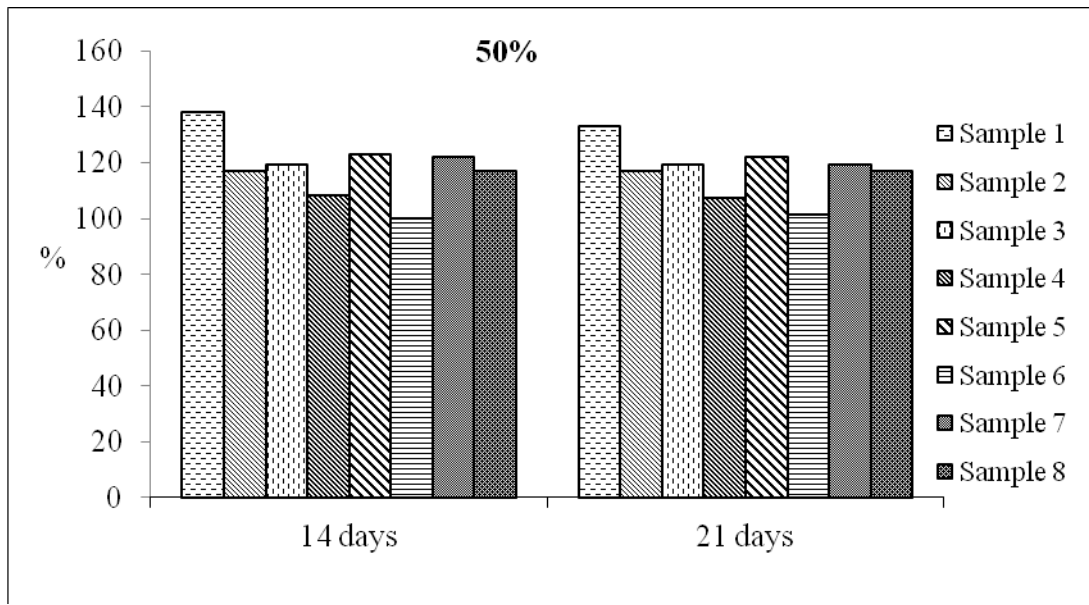
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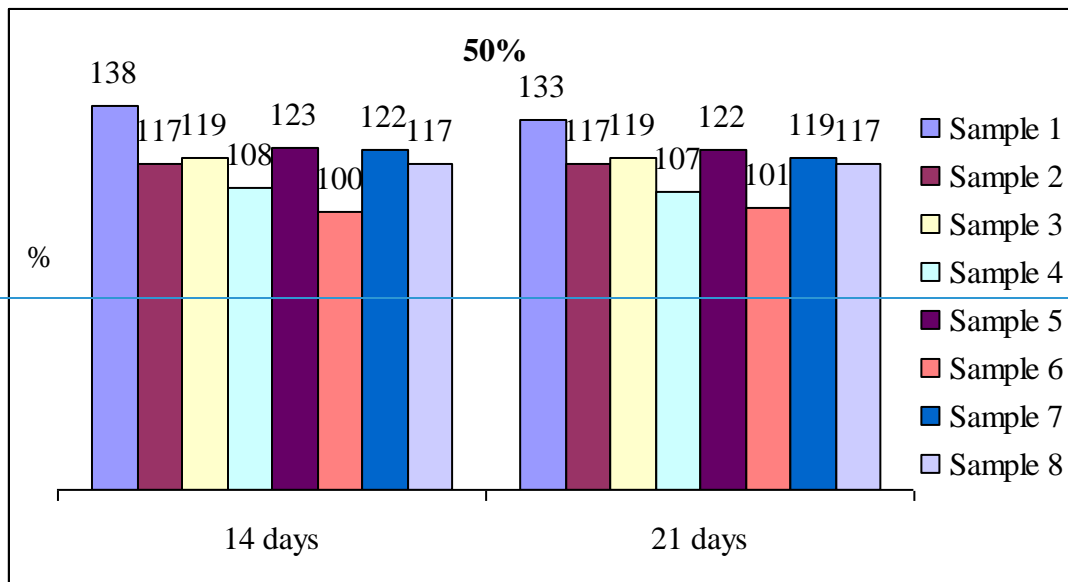
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Figure 45. Comparison of the germination capacity at a concentration of 25%.

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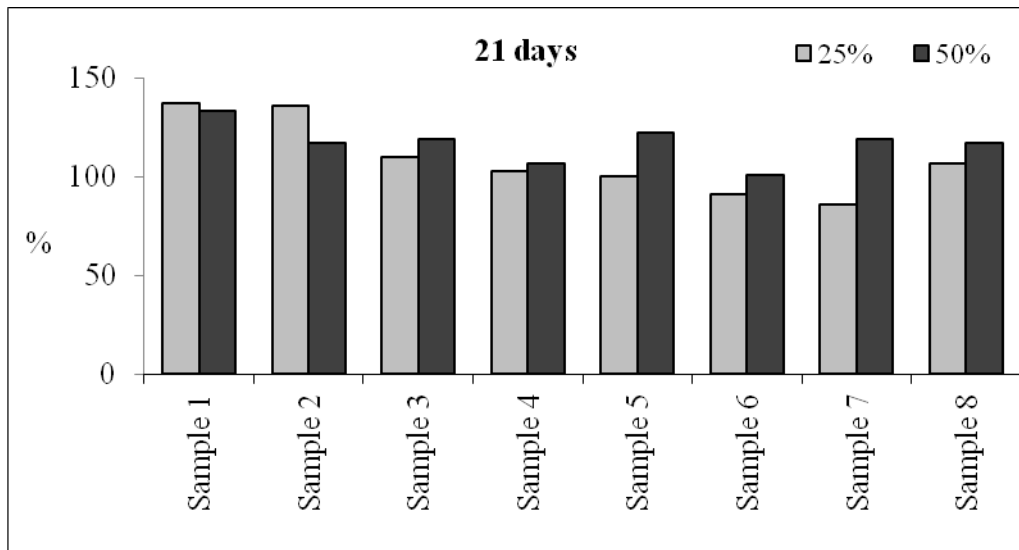


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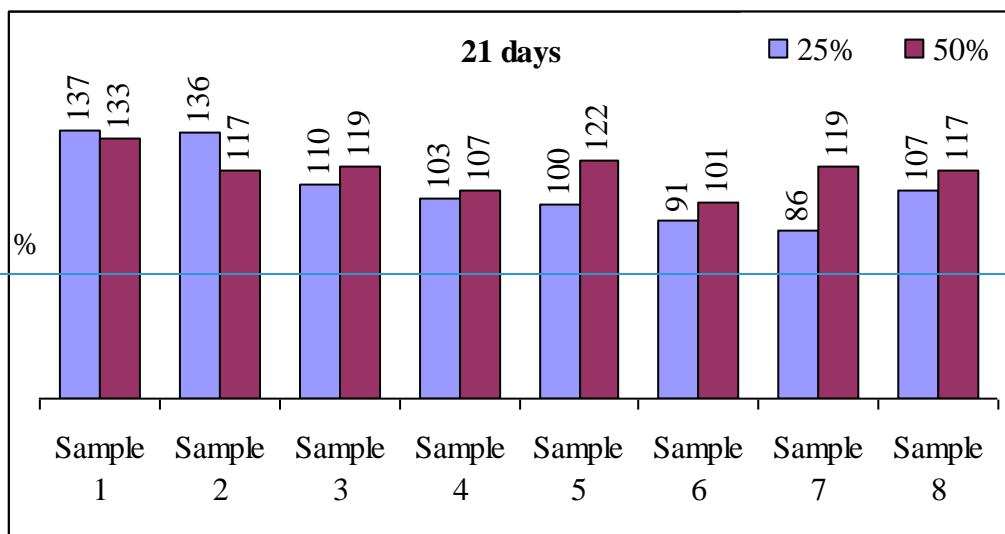
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4 Figure 56. Comparison of the germination capacity of soil samples at a concentration of 50%.

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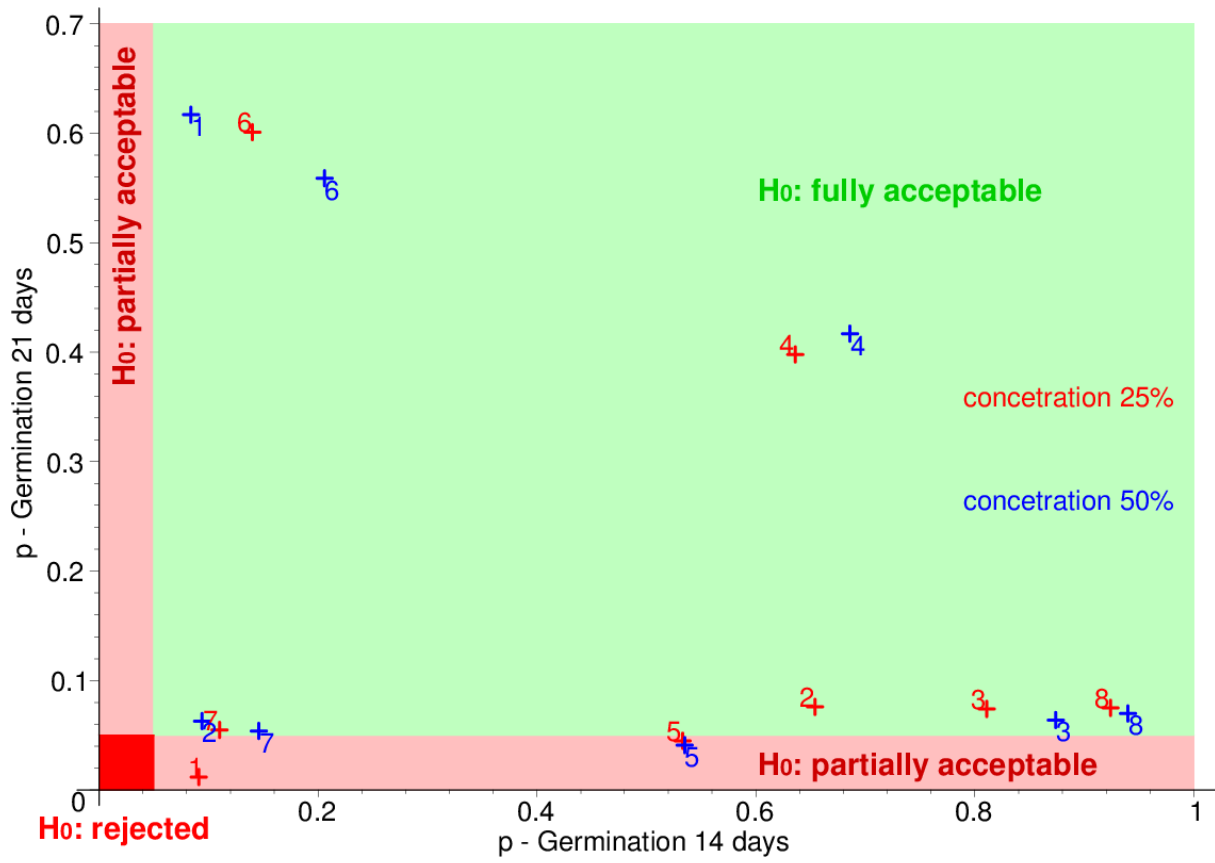
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3

4 Figure 67. Results of germination capacity of white mustard seeds (at concentrations of 25%
5 and 50%).



1

2

3 Figure 78. The values of p factor (ANOVA)