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**Agricultural Reuse of the Digestate from Anaerobic Co-Digestion of Organic Waste:
Microbiological Contamination, Metal Hazards and Fertilizing Performance**

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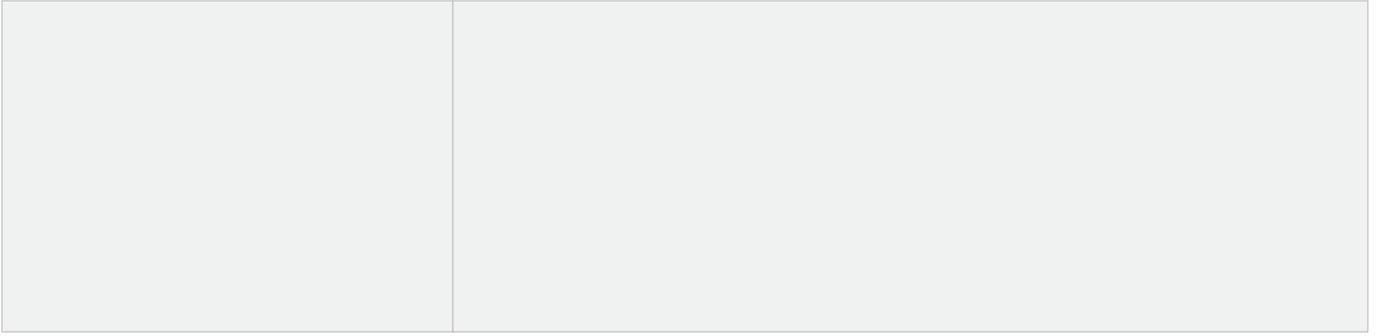
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TITLE PAGE

AGRICULTURAL REUSE OF THE DIGESTATE FROM ANAEROBIC CO-DIGESTION OF ORGANIC WASTE: MICROBIOLOGICAL CONTAMINATION, METAL HAZARDS AND FERTILIZING PERFORMANCE

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Keywords: anaerobic co-digestion, agricultural reuse, fertilizer, faecal indicator bacteria, pathogenic bacteria, heavy metal.

1 **ABSTRACT**

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The aim of this study was to evaluate the agricultural reuse of the digestate products (DPs) obtained from mesophilic anaerobic co-digestion of different organic wastes (sludge, cattle slurries and the organic fraction of municipal solid wastes). At this scope, the content of faecal indicators and pathogens as well as the heavy metal concentration of DPs was monitored. The fertilizing performance of the DPs was also investigated. Co-digestion trials were performed using laboratory-scale (LRs) and pilot-scale reactors (PRs). The microbiological analysis of DPs showed the common presence of *Salmonella* and an inadequate reduction of indicator organisms during the digestion process, both in the LRs and the PRs. Moreover, the presence of pathogens (e.g. *L. monocytogenes*) in some DP samples highlighted the importance of the microbiological quality evaluation of the DPs to study the possible health risks for consumer. In several samples of DPs, the Cu, Ni and Zn content exceeded the maximum admissible concentration for fertilizer, as specified by Italian law, suggesting possible environmental contamination if the DPs are used for agricultural purposes. Considering the fertilizing performance, significant differences of growth parameters were observed only for the DPs that were produced by LRs. In conclusion this work can be considered as a preliminary study to evaluate the possible agricultural reuse of the digestate obtained from different organic wastes.

1 1. INTRODUCTION

2

3 Millions of tons of solid waste are produced annually from municipal, industrial and agricultural sources. The
4 decomposition of these organic wastes results in large-scale contamination of land, water and air (Nasir et al.
5 2012). However, the European Commission has set the ambitious goal of increasing energy from renewable
6 sources to 20% in 2020, compared to 8.5% in 2005 (EREC 2008). To reach this goal, the use of all existing
7 renewable energy sources must increase. Anaerobic digestion is a suitable option for the production of renewable
8 energy in the form of biogas, which can be used to treat organic wastes such as manures, slurries, food
9 processing wastes, sewage sludge and the organic fraction of municipal solid waste (Rajeshwari et al. 2000;
10 Ward et al. 2008).

11 In anaerobic digestion, co-digestion is used to describe the combined treatment of several wastes with
12 complementary characteristics. The co-digestion of combined wastes results in a high methane yield compared to
13 single waste digestion, which is one of the main advantages of this anaerobic technology (Nasir et al. 2012).
14 There are several studies in the literature that address the utilization of co-digestion, such as co-digestion of the
15 organic fraction of municipal solid wastes (OFMSW), cattle manure and agricultural residues (Amon et al. 2007;
16 Macias-Corral et al. 2008), organic solid wastes and sewage sludge (Murto et al. 2004), or more specific wastes
17 (Demirel et al., 2013; Parawira et al. 2004; Traversi et al. 2013).

18 In addition to biogas, anaerobic digestion generates a digestate product (DP) that can be used as an agricultural
19 fertilizer because the nutrients present in the raw input material after the digestion process remain as accessible
20 compounds in the mineralized sludge (Alkanok et al., 2014; Diaz et al. 2011; Lethtomaki and Bjornsson, 2006).
21 The diverse origins of the input material used for biogas production indicate that biogas plants produce fertilizers
22 that vary in nutrient content. At present, considering the Italian laws, the DPs derived from the co-digestion of
23 OFMSW must be considered waste (D.Lgs. n. 205/2010). There is a lack of specific regulation on the use of
24 OFMSW or its derivatives in agriculture: only the agricultural use of wastewater digestion sludge (D.Lgs.
25 99/1992) and animal manure (D.M. n. 109/2006; D.M. n. 29819/2009) is regulated in Italy, and DPs derived
26 from animal by-products (including animal faeces) fall under the European regulation on animal by-products
27 (Commission Regulation n. 142/2011).

28 According to the literature, the physico-chemical properties of DPs have been widely investigated, whereas
29 fertilization studies are still scarce (Abubaker et al. 2012; Garfi et al. 2011; Ning et al. 2011; Nishikawa et al.
30 2012; Tambone et al. 2010). However, the DPs are not harmless products because they contain heavy metals and

1 may also contain organic pollutants, such as pesticides and pathogenic bacteria that are introduced to the soil
2 ecosystem by their application. Heavy metals can be present in the input material used for biogas production and
3 are not altered in the anaerobic digestion process (Sager, 2007); therefore, they may be concentrated due to mass
4 reduction during the process (Dabrowska and Rosinska 2012; Govasmark et al. 2011).

5 The application of digestate on fields can potentially spread pathogens from one farm to another, causing crop
6 contamination. The potential health risk of digested residues from biogas plants is partly dictated by the
7 substrates that are treated in the plants; for instance, organic wastes may contain pathogenic bacteria, depending
8 on the source and type of waste. In particular, wastes of animal and human origin can contain various pathogenic
9 bacteria (e.g., *Salmonellae*, *Enterobacter*, *Clostridia*, *Listeria*), parasites (e.g., *Ascaris*, *Giardia*,
10 *Cryptosporidium*), viruses (e.g., norovirus, enterovirus, rotavirus, Hepatitis A virus) and fungi (*Candida*,
11 *Aspergillus*, *Trichophyton*) (Sahlstrom 2003; Sidhu and Toze 2009; Venglovsky et al. 2006). The possible
12 presence of pathogens can be expected also in the OFMSW (Hassen et al. 2001), even if there is no specific
13 information about pathogen contamination of OFMSW in the literature. Some studies have posited that
14 pathogens can survive after anaerobic digestion (Sidhu and Toze 2009), and the growth of the remaining viable
15 bacteria after the application of DP to land has been demonstrated for some bacterial species (Bonetta et al.
16 2011a; Johansson et al. 2005).

17 The aim of this study was to investigate the content of faecal indicators and pathogens (*E. coli* O157:H7,
18 *Salmonella* spp., *Listeria monocytogenes*, *Giardia* spp. and *Cryptosporidium* spp.) as well as the heavy metal
19 concentration of DPs obtained from mesophilic anaerobic co-digestion of sludge (anaerobic and thickened),
20 cattle slurries and OFMSW. Moreover, this study includes an experiment conducted in a greenhouse that
21 investigated the fertilizing performance of the DPs on *Sorghum bicolor*.

22 This work is a portion of a larger multidisciplinary project DigestedEnergy concerning the improvement of
23 biomass anaerobic digestion process in order to produce biogas, the integration of the process in waste and
24 working refuse management and treatment cycle and the implementation of anaerobic digestion systems targeted
25 to medium-small sized urban, industrial and rural entities.

26

27 **2. MATERIALS AND METHODS**

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29 **2.1 Anaerobic digestion reactors and sampling**

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1 commercial kit for immunomagnetic separation of *Giardia* and *Cryptosporidium* cysts and oocysts (Dynal), in
2 accordance with the manufacturer's instructions. After purification, the presence and number of cysts and
3 oocysts were determined by immunofluorescence with the *Cryptosporidium* Cell Test IF (Cellabs) and the
4 *Giardia* Cell Test IF (Cellabs) (ISS 2007). The oocysts were counted with an epifluorescent microscope (Zeiss),
5 taking into consideration the morphology, size and color of the particles. The results for cyst and oocyst
6 contamination were expressed as the presence/absence, and, when present, as the number of cysts and oocysts
7 per g of sample.

8

9 **2.3 Metal analyses**

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11 Metal content (Cd, Cr, Cu, Hg, Ni, Pb, Zn and Fe) was evaluated in DPs to investigate the chemical hazard
12 related to their reuse as fertilizers. Samples were pre-treated with HCl/NO₃ for digestion in a microwave vessel
13 at 200°C for 30 min, and metal concentration was measured by a Varian Series ICP 820-MS (Palo Alto USA)
14 equipped with a collision reaction interface (CRI) system.

15

16 **2.4 Plant growth test**

17

18 The plant growth test was performed with *Sorghum bicolor* to study the DPs fertilizing performance: two seeds
19 were sown in vases filled with 120 mL of substrate (zeolite:vermiculite:agriperlite:peat; 1:1:1:2) and 20 mL of
20 DP samples. Twelve replicate vases for each DP sample and a control sample (substrate without DP sample)
21 were prepared. Plants were grown in a greenhouse under controlled light, temperature and moisture conditions.
22 After 1 month of growth, plants were harvested and analysed for stem length and for the fresh and dry weights of
23 the stem and root.

24 The results were expressed as the percentage of stem length and as the fresh/dry weight of stem and root
25 compared to the control plants and statistically compared with ANOVA, test – Fisher.

26

27 **3. RESULTS AND DISCUSSION**

28

29 **3.1 Microbiological analyses**

30

1 The results of the analysis of the microbial indicator parameters obtained from the substrates entering the process
2 (input substrates) and from the DPs are reported in **Table 1**. Mesophilic bacterial counts and *E. coli* showed a
3 slightly lower mean value in the sludge (anaerobic and thickened) than in the other input substrates. On the other
4 hand, no relevant differences were observed among the various input substrates for enterococci count, except for
5 cattle slurry which had slightly lower mean value of this parameter compared to the other substrates. The
6 concentrations of the bacterial indicator parameters examined in this study for sludge (anaerobic and thickened),
7 cattle slurry and OFMSW were similar to, or sometimes lower than, those reported in other studies (Iwasaki et
8 al. 2011; Sahlstrom et al. 2004; Sidhu and Toze 2009; Soupir et al. 2006).

9 The DPs produced by LRs generally presented analogous levels of bacterial contamination to those of input
10 waste substrates, with the exception of mesophilic bacteria, which were present in the output samples at lower
11 levels. On the other hand, the digestion process in PRs seemed to cause a reduction of indicator bacteria. This
12 result is particularly important for enterococci, which are considered the best microbial indicators of vegetative
13 bacterial pathogen reduction during the digestion process (Larsen et al. 1994; Viau and Peccia 2009).

14 By contrast, the anaerobic digestion process did not seem to reduce the number of positive samples for *C.*
15 *perfringens*. The presence of *C. perfringens* contamination observed in this work in the DP samples was also
16 reported in earlier studies (Bagge et al. 2005; Bonetta et al. 2011a). The persistence of *C. perfringens* in the
17 digested residues could be an indicator of other pathogenic spore-forming bacteria.

18 The presence of helminth eggs was observed only in two samples of cattle slurry and in only one samples of DP
19 in the LRs. Although helminth infections are a major concern in the developing world, in agreement with results
20 obtained in this study, the occurrence of helminth eggs in digested biosolids has been also reported in other
21 industrialized countries (Rubio-Loza and Noyola 2010; Sidhu and Toze 2009).

22 The number of pathogen positive samples detected in the input wastes and in the DPs are presented in **Table 2**.
23 *Escherichia coli* O157:H7 was not found in any of the samples.

24 *Salmonella* was present in all of the sludge samples (anaerobic and thickened) and in one sample of OFMSW,
25 but was never found in cattle slurry. The identification of species revealed in the sludge samples the presence of
26 *Salmonella enterica* (67%) and *Salmonella enterica* subsp. *arizonae* (33%). *Salmonella* isolated from OFMSW
27 sample was identified as *S. enterica*. In general, the presence of *Salmonella* spp. was reported in almost all DPs
28 collected from the LRs and in some samples collected from the PRs; in all cases, it belonged to the species
29 *enterica*. The presence of *Salmonella* in wastewater sludge such as in anaerobically digested sludge has also
30 been reported in other studies (Dahab and Surampalli 2002; Sahlstrom et al. 2004; Sidhu and Toze 2009). The

1 contamination of OFMSW samples could be ascribed to the use of thickened sludge as a diluent for OFMSW
2 pre-treatment.

3 *Listeria monocytogenes* was present in some samples of sludge and in one samples of OFMSW, but was never
4 found in cattle slurry. This pathogen was found in a sole sample of DP obtained from the PRs. *Listeria monocytogenes*
5 is a common contaminant of organic wastes and, in general, of biomasses. This bacterium is wide spread in the
6 environment (Colleran 2000), but it is normally present in low numbers (Sidhu and Toze 2009). Different studies
7 showed the *Listeria monocytogenes* contamination in wastewater sludge (Sahlstrom et al. 2004; Sidhu and Toze
8 2009). As reported for *Salmonella* the presence of *Listeria monocytogenes* in samples of OFMSW could be
9 related to the mixing of OFMSW with thickened sludge in the pre-treatment. Many studies have reported that
10 animal manure is generally contaminated by different pathogens (Sahlstrom 2003; Venglovsky et al. 2006)
11 contrary to that observed in this study. The effect of the mesophilic digestion process on *Listeria* has not been
12 investigated in depth; according to the results obtained in this study, a reduction, but not a complete elimination,
13 was observed in the study conducted by Horan et al. (2004) using a laboratory-scale plant.

14 *Giardia* was revealed in all samples of sludge (mean 132 cysts g⁻¹), 1 sample of OFMSW (296 cysts g⁻¹) and all
15 DPs (mean 11 and 67 cysts g⁻¹ in LR and PRs respectively); *Cryptosporidium* was observed only in 1 sample of
16 LR and PRs at a low concentration (6 oocysts g⁻¹). *Giardia* and *Cryptosporidium* (oo)cysts are frequently
17 isolated from wastewater, although *Giardia* cysts are more frequently present in these samples compared with
18 *Cryptosporidium* oocysts, which have a more seasonal distribution. Anaerobically digested biosolids can also be
19 contaminated by these protozoa but there are only limited information. Some previous studies reported that DPs
20 may contain up to 10 g⁻¹ *Cryptosporidium* oocysts and 10² g⁻¹ of *Giardia* cysts (Chauret et al. 1999; Hu et al.
21 1996).

22 The results of the microbiological analyses of the DPs obtained from the LR and PRs did not always coincide.
23 A direct comparison between the data obtained in the two types of reactors cannot be made because, as observed
24 in other studies, both the volumes of input substrates involved in the digestion process and the microbial
25 dynamics were different (Wagner et al. 2008).

26 Because no specific Italian law has been established to regulate the hygienic quality of DPs, the results of the
27 microbiological analyses obtained in this study were compared with the maximum admissible concentration (*E.*
28 *coli* < 1000 CFU g⁻¹) specified in the Italian law for fertilizers (D.M. n. 29819/2009). From this comparison, the
29 microbiological analyses of the DPs reveal values for *E. coli* that are higher than the allowed limit in almost all
30 LR digestates and in some PR samples. Moreover, considering the European regulations for animal by-products,

1 all of the DP samples exceeded the standard for enterococci ($m = 1 \times 10^3$ and $M = 5 \times 10^3$ CFU g^{-1})
2 (Commission Regulation n. 142/2011).

3 The presence of *Salmonella* in almost all of the DP samples could represent a hygienic problem because the
4 absence of *Salmonella* in 25 g of material serves as an indicator of bacterial pathogen absence and is considered
5 the standard for the use of DP as a fertilizer. This standard is reported in the Italian laws for fertilizer (D.M. n.
6 29819/2009) and wastewater sludge (D.Lgs. n. 99/1992), as well as in European regulations of animal by-
7 products (Commission Regulation n. 142/2011).

8 Considering that in this study the suitability of *Salmonella* presence as an indicator of bacterial pathogens has not
9 been demonstrated, other indicators, such as viral contaminants, should be introduced and studied. In particular,
10 an evaluation of the effectiveness of the anaerobic process and determination of the most effective post-treatment
11 of DP, based on different input substrates and the type of anaerobic process used, could be useful for
12 identifying specific indicators for each biogas plant.

13

14 3.2 Metal analyses

15

16 The metal concentrations of the DPs are reported in **Table 3**. In several DP samples, the metal content exceeded
17 the maximum admissible concentration under the Italian law for fertilizers (D.M. n. 29819/2009). In particular,
18 the levels of Cu, Ni and Zn were found to be higher than the allowed levels in all of the DP samples from the LR
19 and in a large number of the PR samples. The Cd concentration showed values higher than the limits in two of
20 the DP samples from the LR and in a sole PR sample. On the other hand, the concentrations of Hg and Pb were
21 lower than the limits specified by law.

22 If the DPs are considered as sludge, then it is possible to compare the metal concentrations with the Italian law
23 for wastewater treatment sludge (D.Lgs. n. 99/1992). In this case some DP samples produced by the LRs
24 exceeded the allowed limits for Zn and Ni.

25 The high levels of some heavy metals observed in the DP samples are ascribed to contamination of the input
26 substrates that were used for anaerobic digestion in this study. In fact, different authors have reported that
27 wastewater sludge, cattle manure and OFMSW can contain hazardous substances such as heavy metals (Dong et
28 al. 2010; Tulayakul et al. 2011; Uysal et al. 2010). Moreover, it is important to note that during anaerobic
29 digestion, the composition of organic substances results in an increase of heavy metal concentration in the dry
30 matter of sludge (Dabrowska and Rosinska 2012).

1 The heavy metal content lower than that showed in the DPs analysed in this study was reported in other works
2 focusing on DPs from food and garden waste (Govasmark et al., 2011), slurry (Jin et al., 2011) or OFMSW
3 (Dong et al. 2010; Gracia et al., 2012). A similar contamination was observed in the study of Dabrowska et al.
4 (2012) that analysed the heavy metal content in DPs from wastewater sludge highlighting the role of this input
5 substrate in the heavy metal contamination of the digestate.

6 The presence of significant amount of Cu, Ni and Zn in DPs suggests that there is a possibility of environmental
7 contamination if the DPs are used for agricultural purposes. In addition to environmental concerns, the release of
8 heavy metals (e.g., Cu, Zn, Pb, Cd) into soils, water and plants, through the use of DPs as fertilizers, could also
9 pose public health risks throughout the food chain.

11 3.3 Plant growth test

12
13 The general effects of DPs used as fertilizers on relative biomass fractions (root, stem) and plant stem lengths are
14 reported in **Table 4**. As shown for the DPs produced by PRs (compared to the control), neither the weight of the
15 stem and root nor the stem length were affected by treatment with DPs, which produced parameter values that
16 were similar to those in the control plant. Significant differences ($p < 0.05$, ANOVA test – Fisher) of growth
17 parameters with respect to the control were observed only for the DPs produced by LRs. The different fertilizing
18 performance of DPs produced by the two types of reactors (LRs vs. PRs) could be ascribed to differences in the
19 management practices and operating conditions of the digestion process, which lead to the production of
20 fertilizers with varying contents of plant macro- and micro-nutrients, as well as chemical contaminants
21 (Abubaker et al. 2012; Garfi et al. 2011).

22 Unfortunately, it is difficult to compare the results of this study to those obtained in previous studies because the
23 fertilizing performance of DPs depends on the origin and composition of the feedstock and has been investigated
24 under experimental conditions (pot vs. field trials) that vary in the plant tested (e.g., tomato, rice, potato) and the
25 fertilization rate (Abubaker et al. 2012; Garfi et al. 2011; Ning et al. 2011; Nishikawa et al. 2012).

27 4. CONCLUSIONS

28
29 In this work, the possible agricultural reuse of DPs produced by the anaerobic co-digestion of sludge (anaerobic
30 and thickened), cattle slurry and OFMSW was investigated from several different points of view.

1 The microbiological analysis of DPs performed in this study revealed the presence of *Salmonella* and an
2 inadequate reduction of indicator organisms during the digestion process, both in the laboratory and pilot-scale
3 reactors. Therefore, this contamination could make the DP unsuitable as an agriculture fertilizer. Moreover,
4 the presence of pathogens (e.g. *L. monocytogenes*) in some DP samples highlights the importance of the
5 microbiological quality evaluation of the DPs to study the possible health risks for consumer.
6 Considering the metal content of the DPs analysed the high levels of Cu, Ni and Zn may cause environmental
7 contamination. Thus, heavy metal pollution should be a concern when we apply DPs to soil, particularly in
8 relation to the possible health risks for humans that are caused by some heavy metals (e.g., Cd, Cr and Pb).
9 The significant positive effect on plant growth observed with the DPs obtained from LRs demonstrates the need
10 for further investigations. In particular, it could be interesting to study fertilizing performance on different plants
11 using field trials.
12 In conclusion this work can be considered as a preliminary study to evaluate the possible agricultural reuse of
13 the digestate obtained from different organic wastes.

14

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16

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24

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29

Table 1. Mean, minimum and maximum values (expressed as log₁₀ CFU g⁻¹) or percentage of positive samples of bacterial indicator parameters in input substrates and DPs.

	N	Mesophilic count			<i>E.coli</i>			Enterococci			<i>C.perfringens</i>	Helmint eggs
		mean	m	M	mean	m	M	mean	m	M	(%)	(%)
Organic waste												
Anaerobic sludge	4	6.3	5.3	8.7	3.5	2.9	3.8	4.2	3.1	4.9	75	0
Thickened sludge	4	7.1	6.7	7.9	4.2	2.9	5.7	4.3	4.2	4.5	75	0
Cattle slurry	4	7.8	7.3	8.5	5.3	4.9	5.5	3.5	2.8	4.2	50	50
OFMSW	4	7.9	7.4	8.7	4.9	4.3	5.0	4.6	3.8	5.2	100	n.a.
DPs												
LRs	8	6.6	5.4	7.8	4.4	<2	5.6	4.8	4.6	5.0	71	13
PRs	8	6.1	5.1	7.0	3.6	<2	4.4	3.8	3.1	5.2	100	0

N: number of samples; n.a.: not analysed; m: min; M: max

Table 2. Number of pathogen positive samples in input substrates used for anaerobic co-digestion and DPs.

	<i>E. coli</i> O157:H7	<i>Salmonella</i> spp.	<i>L.</i> <i>monocytogenes</i>	<i>Cryptosporidium</i> spp. <input type="checkbox"/>	<i>Giardia</i> spp. <input type="checkbox"/>
Organic waste					
Anaerobic sludge	0/4	4/4	1/4	0/2	2/2
Thickened sludge	0/4	4/4	3/4	0/2	2/2
Cattle slurry	0/4	0/4	0/4	0/2	0/2
OFMSW	0/4	1/4	1/4	0/2	1/2
DPs					
LRs	0/8	7/8	0/8	1/2	2/2
PRs	0/8	3/8	1/8	1/2	2/2

Only two samples were analysed

Table 3. Metal content and Italian law Limits measured in DP samples expressed as mg/kg s.s.

	LR		PR		Law limits	
	Mean±DS	Max	Mean	Max	LV fertilizer ^a	LV sludge ^b
Cd	1.4±0.3	1.6	0.6±0.7	1.6	1.5	20
Cr	250.6±187.0	560.3	96.9±58.1	240.0	/	/
Cu	447.6±111.2	561.0	159.9±102.3	420.0	230	1000
Hg	0.7±0.1	0.8	0.5±0.5	1.1	1.5	10
Ni	239.0±89.2	355.9	107.8±61.4	220.0	100	300
Pb	104.1±17.2	126.0	52.6±23.6	91.0	140	750
Zn	2818.4±560.1	3753.7	655.0±234.5	1200.0	500	2500
Fe	21129.3±6171.0	29837.7	13700.0±2022.7	16.000.0	/	/

^a Limit value (LV) D.M. n. 29819/2009; ^b Limit value (LV) D.lgs. n. 99/1992

Table 4. Plant growth parameters measured after application of DP as fertilizer. Results were expressed as percentage respect to control (100%).

DPs	Stem lenght	Fresh weight stem	Fresh weight root	Dry weight stem	Dry weight root
LRs	149.5±10.5	194.8±56.4	235.9±43.8	225.4±39.9	221.1±40.2
PRs	123.8±10.3	171.6±39.6	156.8±27.7	164.8±37.6	135.2±16.2