

Growth of saltwater algae on press water of champost (spent mushroom waste)

Hellen Elissen, Saskia Gaastra & Rommie van der Weide | Wageningen University & Research | December 2020





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Wageningen, December 2020

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Summary: For mushroom production usually a mixture of horse and chicken manure and straw is used. For every kg of mushrooms, 5 kgs of waste is produced, a total of 800.000 tonnes per year in the Netherlands. Processing options are often either costly or complex. As the champost contains high salt concentrations, washing results in a solid (less salty) fraction with better re-use options and a nutrient-rich liquid fraction. On the latter fraction growth of five species of commercially relevant micro algae was tested in small-scale well tests. In addition, the press water was also treated with H₂O₂/UV to reduce the brown colour of the solution. In the majority of the triplicate well tests, growth of three of the five common saltwater algae species (*Nannochloropsis gaditana, Porphyridium purpureum*, and *Tetraselmis chuii*) on press water fractions of spent mushroom waste (champost) was found to be comparable to or better than their growth on control media. *Dunaliella salina* growth was relatively low. *Arthrospira platensis* only grew on press water when treated with H₂O₂. Overall, the results indicate that washing of champost results in a mineral rich solution suitable for growth of several (saltwater) algae.

Keywords: champost, spent mushroom waste, washing, press water, micro algae growth, decolourization

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Preface

In the Netherlands, there are surpluses of manure and digestates/biogas slurries (digested manure and residual flows), which represent a negative value. At the same time, these residual flows contain valuable ingredients for the production of biomass (as raw material for food and feed products), for improving soil quality and for energy production. The number of feasible business cases in which the residual flow is upgraded has so far been limited. This is due both to the efficiency of the technologies used and the legislation and regulations related to the residual flows.

Recent information from research, scientific literature and companies provides new starting points for a biobased valorisation of manure/digestate streams and improving the efficiency of anaerobic digestion. The innovative aspect of our research is the cultivation of new types of biomass on the residual flows and the use of the conversion products to improve anaerobic digestion. This involves the use of separated manure and digestate products for the cultivation of mushrooms/fungi, worms, insects, specific bacteria and aquatic biomass. The resulting biomass can be further refined and marketed as food, feed and bio-based feedstock. There are also processed manure and digestate products that are valuable as fertilizer products for soil and plant growth, as substrate for improvement of anaerobic digestion or for export/use besides in agriculture. This gives a new interpretation to obligatory manure processing.

The aim of this project is to further explore and substantiate/test these ideas on lab and practical scale, leading to a proof of principles for new bio-based upgrading methods for manure and digestate that can be used in conjunction to better close cycles and/or sell outside regular agriculture. Bottlenecks in legislation and regulations are explored and put on the agenda. Key figures are also calculated that are necessary for assessing sustainability (e.g. costs, environmental effects) and for supporting legislation (e.g. minerals, food safety).

The livestock sector gains insight into the possibilities of biobased valorisation and better marketing of their most important residual flows. For the small and medium-sized enterprises (SMEs) involved, this research provides proof of principle for their technology and input in their business cases. The combined effects of the technologies provide new knowledge, methods and research directions for science. In a social context, the use and upgrading of manure and digestates in other ways also contributes to the transition to a circular bio-economy with an efficient and sustainable agri-food sector.

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Summary

For mushroom production usually a mixture of horse and chicken manure and straw is used. For every kg of mushrooms, 5 kgs of waste is produced, a total of 800.000 tonnes per year in the Netherlands. Processing options are often either costly or complex. As the champost contains high salt concentrations, washing results in a solid (less salty) fraction with better re-use options and a nutrient-rich liquid fraction. On the latter fraction growth of five species of commercially relevant micro algae was tested in small-scale well tests. In addition, the press water was also treated with H₂O₂/UV to reduce the brown colour of the solution.

In the majority of the triplicate well tests, growth of three of the five common saltwater algae species (*Nannochloropsis gaditana*, *Porphyridium purpureum*, and *Tetraselmis chuii*) on press water fractions of spent mushroom waste (champost) was found to be comparable to or better than their growth on control media. *Dunaliella salina* growth was relatively low. *Arthrospira platensis* only grew on press water when treated with H_2O_2 . Overall, the results indicate that washing of champost results in a mineral rich solution suitable for growth of several (saltwater) algae.

1 Introduction

For mushroom production usually a mixture of horse and chicken manure, gypsum and straw is used. After mushroom harvest a waste stream remains (~800.000 tonnes per year), called champost (or mushroom compost waste or spent mushroom substrate/waste/compost). For every kg of mushrooms, 5 kgs of waste is produced (Lin et al, 2014) with a dry matter content of 35 % (Schlatmann and Kosse, 2018). The waste can be used for biogas production, but yields are not very high (Oei and Albert, 2008). Other options such as pyrolysis and incineration of the champost seem interesting but also complex because of its high moisture content. Another limiting factor for certain re-uses is the high electrical conductivity of the waste due to high salt concentrations. High salt concentrations (NaCl, CaCl₂) result from the applied substrates and salt additions during the growth process. Therefore, washing of the waste leads to a better product that for example can be used as peat replacement (EC values < 1.0 dS/m) (Blok et al, 2015). The press water contains salts and nutrients and is possibly a good medium for algae production. In this report an initial test in well plates is described with five common commercially produced saltwater algae on different fractions of the press water. These saltwater algae are used for several applications (Table 1) (Enzing et al, 2014).

Table 1	Some commercial applications/products of algae species used in the experiments (From:
Enzing et al,	2014)

Species	Application
Arthrospira platensis	Food additive
	Phycocyanin
Dunaliella salina	β-carotene
	Xanthophyll
Porphyridium purpureum	Phycocyanin
	Food additive
Tetraselmis chuii	Aquaculture feed
	Carotenoids
Nannochloropsis gaditana	EPA
	Algae paste
	Aquaculture feed
	β-carotene

Production of the algae species on low cost safe waste streams is thus an interesting option. The press water of champost naturally has a dark brown colour, which may interfere with algal growth. DePraetere et al (2013) found that colour removal using different methods could result in a doubling of initial growth rate and a 50 % increase in final biomass yield of *Arthrospira platensis* grown on piggery wastewater. Treatment with hydrogen peroxide (H_2O_2) and UV resulted in 80 % colour removal and had no effect on phosphate concentrations, since it oxides coloured compounds. According to Vasconcelos Fernandes et al (2015) however, the translucent brown colour of anaerobically treated blackwater does not have to be a limiting factor for algal growth, as long as algal numbers and light intensity are high enough. Weide, van der, et al (2014) also found growth on different dilutions of chicken manure digestate. They observed that with increasing digestate concentrations, algae needed more time to start growing. In the test described in this report, one fraction has been treated with hydrogen peroxide (H_2O_2) and UV in order to decolour the medium and promote algal growth.

2 Materials and methods

2.1 Well test setup

At ACRRES a method was developed for screening quickly the suitability of waste streams for algal growth. This method enables simultaneous quick small-scale testing of several algal species and waste streams (Huurman and van der Weide, 2015). The method results in a '+/-' result for growth plus a quantification of algal numbers and/or OD measurements. For the screening 24-well plates are used. Figure 1 shows an example of such a test.

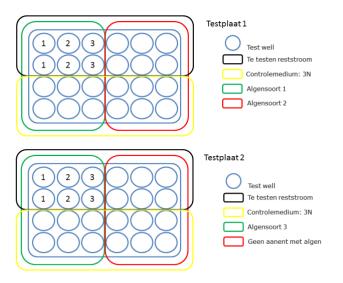


Figure 1 Example of well test for one test medium and three algae species (From: Huurman and van der Weide, 2015).

All test combinations were tested in triplicate, and some were repeated as extra controls. Four plates in total were incubated on an orbital shaker at 100 rpm (continuously) in a climate cabinet at 20 °C on a glass surface with white LED lightning underneath (4.3 A, 12.3 V) covered with a white paper resulting in a light intensity of ~ 79 µmol/m²/s. 50 mL of each algae stock culture was previously centrifuged for 5 minutes at 3000 rpm, the supernatant (standard medium) discarded (to remove unwanted nutrients) and the algal pellet suspended in an equal amount of half-strength artificial seawater (15 g/L). Half-strength seawater was chosen not to disrupt the cell walls. This was repeated twice. An initial OD₄₄₀ nm of 0.3 was aimed for. In each well 2.25 mL of press water or control medium was added plus 0.25 mL of precultured algae. In controls only 2.5 mL of press water or control medium was added. For all cell counts Bürker Türk counting chambers of 0.0025, 0.04 and 0.1 mm² of Optik Labor (depth 0.1 mm) were used depending on estimated visual algal cell density. All counts were done under a 400x (40x10) Leitz Laborlux S and a Leica DMLS microscope. After 7 days the tests were terminated. OD measurements at the end of the test were not performed as the press water was often coloured and the resulting OD would not be an accurate representation of the algae numbers.

2.2 Press water

Mushroom spent waste was obtained from CNC (Gennep, the Netherlands) and kept at 4 °C until use. As an indication, the composition of this waste from several sources is shown in Annex I. The DM % was ~40 % and OM % was ~60 % of DM. Three different types of press water were produced from the waste:

- A: 1 kg of champost in a bucket was filled up to 5 L with demi water, stirred with a spoon and left for an hour to settle. The liquid was decanted and collected, the remaining solids pressed (by pressing a perforated bucket into a bucket with the soaked champost, thereby collecting the liquid in the upper bucket) and the press water was added to the decanted liquid. The mixed liquid was filtered over a woven mesh (Monodur PA 90n, average pore size 82.6 µm).
- B: The treatment for A was repeated with 1 kg of champost and subsequently the resulting liquid was filtered over a paper filter (Schleicher & Schuell No. 595, 180 mm).
- C: 180 mL of H₂O₂ (hydrogen peroxide solution 34.5-36.5%, Sigma-Aldrich) was added to 420 mL of liquid B and left for 1 hour. The solution was subsequently placed under UV light (Sylvania Blacklight blue, F15W BLB T8 in a 1L blue cap glass media bottle at 20 °C for 3 days.

All three fractions were analysed for their composition by Eurofins (Wageningen, the Netherlands) (Annex II). The coarse and fine filtered fractions were comparable in composition, but the fraction treated with H_2O_2 contained lower concentrations of most elements, had a lower EC and a lower pH. In addition, the colour of the H_2O_2 treated fraction was substantially lighter (Figure 2, Annex III). Figure 2 shows the resulting solutions and particles for A, B and C.

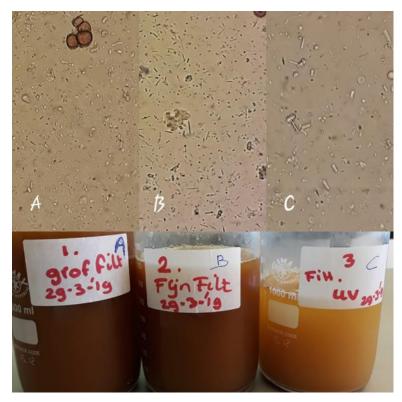


Figure 2 Solutions A, B and C with close-ups of the particles.

2.3 Saltwater algae

In a previous research growth of several saltwater algae strains was tested on salty waste streams (Elissen et al, 2018). The best growing/most common algae were selected for the current research (Table 2): *A. platensis*, *D. salina*, *N. gaditana*, *P. purpureum* and *T. chuii*.

Table 2Algal strains, sources and control media used in the well tests.

Strain & ACRRES code	Source	Control medium
Arthrospira platensis 1	SAG 21.99	Modified Spirulina 1)
Dunaliella salina 2	SAG 184.80	f/2 (CCAP) 2)
Nannochloropsis gaditana 3	ACRRES lab	f/2 (CCAP) 2)
Porphyridium purpureum 4	SAG 1380-1d	f/2 (CCAP) 2)
Tetraselmis chuii 8	AF&F (Almere, the Netherlands)	f/2 (CCAP) 2)

References: ¹⁾ Aiba & Ogawa (1977), Schlösser (1994), ²⁾ Guillard & Ryther (1962)

Previously purchased cultures that were kept in the ACRRES lab were used for the well tests.

3 Results and discussion

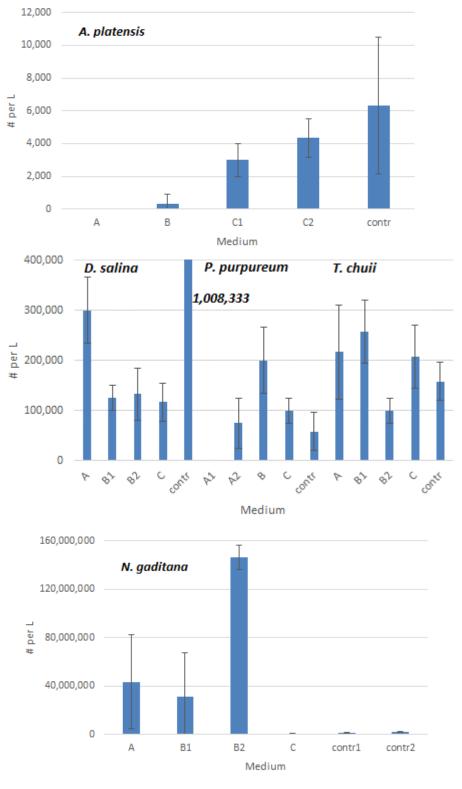
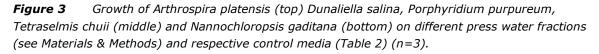


Figure 3 shows the growth results of different saltwater algae with press water fractions.



Numbers behind the mediums refer to repetitions of the triplicates. Separate graphs were created because of large differences in algal numbers for the different species.

Highest algae numbers were found for *N. gaditana*, followed by *D. salina*, *P. purpureum* and *T. chuii* with comparable numbers, followed by *A. platensis* with the lowest algae numbers. Several combinations of press water fractions and saltwater algae resulted in growth in numbers comparable to growth on control media.

- *A. platensis*: Growth on the control medium was the best (but with a large standard deviation), followed by growth om media C, B and A. Growth in each repeated C triplicate was comparable. No algae survived in A.
- *D. salina*: Growth on the control medium was by far the best and was followed by media A, B and C, the latter two being similar. Growth in each repeated B triplicate was also comparable.
- *P. purpureum*: Growth on medium B was the best, followed by C, A and the control, the latter three all being very similar. Growth in the repeated A triplicates was different, as no algae survived in A1. It is not clear why there were sometimes differences in growth between two triplicates.
- *T. chuii*: Growth was best in B1, but this was very similar to growth in A, C and the control. Only growth in one of the repeated B triplicates (B2) was lower.
- *N. gaditana*: Growth was best in one of the repeated B triplicates (B2), followed by A and the repeated control triplicates. Relatively very low numbers were found in the controls and in medium C. The low growth in the controls could be due to the medium choice.

Table 3 shows the combinations which resulted in similar or better algae growth to control media.

Table 3Combinations of algae with different media (see Materials & Methods) that resulted insimilar or higher growth compared to control media (+) and combinations that resulted in lowernumbers (-). +/- denotes triplicates resulting in different algal numbers for the same medium/algaespecies combination

Medium	A. platensis	D. salina	P. purpureum	T. chui	i N. gaditana
Α	-	-	+/-	+	+
В	-	-	+	+	+
С	+	-	+	+	-

Press water from champost thus seems a promising medium for growth of several saltwater algae species. From the results it's not clear that (the method in this experiment used for) decolorization of these media resulted in better algae growth, except for *A. platensis*. As mentioned, decolorization has also been found to promote growth of *A. platensis* on piggery wastewater (DePraetere et al, 2013). This species may be more susceptible to lower light conditions than the other algae species that grew well on the original brownish press water fractions. Mixotrophic growth on C components in the press water in the absence of sufficient light could also have taken place in the tests were good algae growth was found.

When compared to algae numbers found in experiments with the same algae species on several 'salty' waste streams by Elissen et al (2018) only *A. platensis* showed overall lower numbers in the fractions and control medium used in the current test. The other algae species grew in comparable numbers. Annex III shows the well plates at the start and end of the experiment. Superficially looking, colour of the H_2O_2 treated fraction C changed from yellow to colourless, while wells with fractions A en B did not change (remained brown), except in the wells with *N. gaditana*, where the colour became more dark brown, possibly as a result of relatively high algal numbers.

Conclusions and recommendations

4

In the majority of the triplicate well tests, growth of four common saltwater algae species (*N. gaditana*, *P. purpureum*, *D. salina* and *T. chuii*) on press water fractions of spent mushroom waste (champost) was found to be comparable to or better than their growth on control media. *A. platensis* only grew on press water when treated with H_2O_2 . This was probably the result of higher light penetration in the liquid. The other well-growing species seemingly were not hindered by the dark colour of the press water or grew in mixotrophic mode on C sources in the press water. Overall the results indicate that washing of champost could lead to

1. A mineral rich solution for (saltwater) algae growth, still containing some particles, that may influence light absorbance.

2. A 'less salty' solid fraction that may be more appropriate for subsequent use of the waste as for example fertilizer or as feedstock for terrestrial insects or worms

Press water of champost could be a cheap nutrient solution for the production of several commercial algae species. It is however recommended to repeat experiments with the combinations that resulted in similar or better growth than growth on the control medium in larger test volumes. In addition, a business case should be formulated in which the extra costs for washing the champost should be included.

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Annex 1 Compositions of champost

Kwaliteitsonderzoek Compost champost zonder dekaarde

Uw klantnummer: 2818922

CNC Grondstoffen (RHP) BV T. v.d. Venne Postbus 13 6590 AA GENNEP

Analyse-/ordernummer: 2014903193/003317897 Onderzoek

Datum verslag: 18-04-2014

Datum monstername:

02-04-2014

Monster genomen bij: CNC P.H Vervoort, postbus 13 6590 AA GENNEP Datum ontvangst: 07-04-2014

champost analyse cns pve

Type monster: Champost

Resultaat		Eenheid	Resultaat	Toetswaarde	Conclusie	Resultaat in produkt (g/kg)
bepaald in het monster volgens de op de hieronder vermelde normer	Droge stof	g/kg product	440			
	Ruw as	g/kg ds	454			
	nOrg. stof	% van de ds	54,6	10,0		
	Stikstof (N)	g/kg ds	21,5			9,5
	Fosfor (P)	g/kg ds	5			
	Fosfaat (P2O5)	g/kg ds	11,5			5,06
	Kalium (K)	g/kg ds	29			
	Kali (K ₂ O)	g/kg ds	35			15
	Zwavel (S)	g/kg ds	21,9			9,6
	Magnesium (Mg)	g/kg ds	4,5			
	Magnesia (MgO)	g/kg ds	7,5			3,3
	Chloride	g/kg ds	10			
	Zuurgraad (pH)		5,7			
	C-anorganisch	96	0,20			
	Koolzure kalk	%	1,1			
	Geleidingsvermogen	mS/cm 25°C	17,82			
	Cadmium (Cd)	mg/kg ds	0,31	1,00		
	Chroom (Cr)	mg/kg ds	7,5	50		
	Koper (Cu)	mg/kg ds	34	90		
	Kwik (Hg)	mg/kg ds	0,05	0,30		
	Nikkel (Ni)	mg/kg ds	4,2	20		
	Lood (Pb)	mg/kg ds	6,4	100		
	Zink (Zn)	mg/kg ds	192	290		
	Arseen (As)	mg/kg ds	1,4	15		

Toelichting

De onderzochte parameters voldoen aan de samenstellingseisen van compost zoals vermeld in de Uitvoeringsregeling Meststoffenwet.

Contact & info

Monster genomen door: Derden Contactpersoon monstername: Patrick Bens: 0652002106

Na verzending van dit verslag wordt, indien de aard en de onderzoeksmethode van het monster dit toelaat, het monster nog twee weken bij BLGG AgroXpertus voor u bewaard. Binnen deze tijd kunt u eventueel reclameren en/of aanvullend onderzoek aanvragen.

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onze vormen van dienstverlening zijn o izoek worden deze ervlof de specificat AgroXpertus stelt zich niet aarspreke gebruik van door of namene BLOG # Op al o Op ver BLGG uit het ning zijn orze Algemene Voorwaarden van t specificaties van de analysemethoden toege ansprakalijk voor eventuele schadelijke gev BLGG AgroXpertus verstreide onderzoeken



BLGG AgroXpertus is ingeschreven in het RvA-register voor testaboratoria zoals nader omschreven in de erkenning onder m. L122 voor uitsluitend de monsternemings- en/of de analysemethoden.

BLGG AGROXPERTUS

T monstername: Patrick Bens: 0652002106 T klantenservice: +31 (0)88 876 1010 E klantenservice@blgg.agroxpertus.nl I blgg.agroxpertus.nl

Postbus 170 NL - 6700 AD Wageningen

	Eenheid	Resultaat
Droge stof	G/kg product	333
Org. Stof	% van de ds	62,0
Fosfaat (P ₂ O ₅)	g/kg ds	11,9
Stikstof (N)	g/kg ds	21,0
Cadmium (Cd)	mg/kg ds	0,29
Chroom (Cr)	mg/kg ds	7,7
Koper (Cu)	mg/kg ds	34
Kwik (Hg)	mg/kg ds	0,04
Nikkel (Ni)	mg/kg ds	4,3
Lood (Pb)	mg/kg ds	6,8
Zink (Zn)	mg/kg ds	145
Arseen (As)	mg/kg ds	1,1
Kali (K ₂ O)	g/kg	21,5
Geleidingsvermogen	mS/cm 25°C	12,4

Name	compost	champost	kolen	bruinkool	kippenmest	B-hout	RWZI slib	palmpit	diermeel
PROXANAL									
Moisture (a.r.)	61,20	63,88	9,95	58,69	9,29	9,08	75	11,01	2,50
Fixed Carbon			44,12		14,40	21,62	28.4	17,59	12,71
Volatile Matter			33,62		47,82	76,53		77,28	63,31
Ash	39,28	36,52	12,30	8,70	37,79	1,85	43	5,14	23,95
ULTANAL									
Ash	39,28	36,52	12,30	8,70	37,79	1,85	43	5,14	23,95
C (d.b.)			71,05	61,75	37,38	50,26	28.4	48,34	43,07
H (d.b.)	3,77	3,82	4,61	4,50	4,19	6,91	4.6	6,20	6,04
N (d.b.)	2,29	1,90	1,50	0,60	3,76	1,03	3.5	2,62	9,16
CI (d.b.)	0,77	0,63	0,01	0,05	0,13	0,07	.07	0,21	0,87
S (d.b.)	3,35	2,70	0,69	0,39	0,74	0,00	0.09	0,26	1,27
O (by diff.)			9,83	24,02	15,64	39,88	19.6	37,23	15,64
Stookwaarde (a.r.)	2.66	2.51	27,49	8,10	13,99	18,54	2 a 3	19,36	17,27
Asanalyse									
macro- elements									
AI	0,31	0,32	1,71	0,20	0,18	0,04	2.7	0,15	0,29
Ca	4,95	6,93	0,35	1,34	13,93	0,32	11.8	0,31	6,88
CI	0,77	0,63	0,01	0,85	0,13	0,07	0.07	0,21	0,87
Fe	0,22	0,20	0,54	0,02	0,11	0,05	7.8	0,30	0,04
K	3,59	2,62	0,12	0,23	3,47	0,16	0.5	0,63	0,61
Mg	0,55	0,45	0,13	0,02	0,85	0,06	1.0	0,18	0,19
Na	0,32	0,26	0,04	0,01	0,15	0,04	1.8	0,00	1,11
P	0,57	0,47	0,04	1,84	2,30	0,08	5.2	0,62	4,17
Si	4,09	3,55	3,02	0,07	0,93	0,20	10.1	0,39	0,00
Ti	0,02	0,01	0,10	0.00	0,01	0,02	0.3	0,00	0,00

Fiquur 42: Samenstelling compost, champost en andere gangbare brandstoffen

Annex 2 Composition of champost press water fractions

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Rapport

	ongineer				
Monster	Onderzoek-/ordernr: 820490/004683869	Datum monstername 29-03-2019		Datum verslag: 18-04-2019	
	Code onderzoek: 485	Datum ontvan 18-04-2019	gst:	Monster genomen door: Derden	Contactpersoon monstername:
Resultaat	EC		m 25°C		
	pН	6,1			
		mmol/l	ppm (=mg/l)	
	Nitraat (NO ₃)	< 1,0	< 63		
	Ammonium (NH ₄)	3,5	63		
	Fosfor (P)	1,95	60		
	Kalium (K)	57,5	2248		
	Calcium (Ca)	12,2	489		
	Magnesium (Mg)	6,6	160		
	Zwavel (S)	31,1	997		
	Natrium (Na)	10,9	251		
	Chloride (CI)	28,4	1007		
	Bicarbonaat (HCO3)	< 1,0	< 62		
	Silicium (Si)	0,50	14,04		
		µmol/l	ppm (=mg/l)	
	Borium (B)	21	0,23		
	Koper (Cu)	6,1	0,39		
	IJzer (Fe)	22	1,23		
	Mangaan (Mn)	56	3,08		
	Molybdeen (Mo)	< 1.0	< 0,10		
	Zink (Zn)	10	0,65		
	De resultaten zijn uitge Zij kunnen niet dienen			re fractie van het aangeboden r offenwet.	monster.

de Alle bepalingen

Em: MENGMS en DAHFD

 Q
 Methode geaccrediteerd door RvA

 Em:
 Eigen methode, Gw: Gelijkwaardig aan, Cf: Conform

 Bij deze verrichting is de gestelde houdbaarheidstermijn tussen monstername en analyse overschreden. Dit heet mooglik de betwurbaarheid van het resultaat behivioed.

 De resultaten hebben uitsluitend betrekking op het in behandeling genomen materiaal.

Pagina: 1 Totaal aantal pagina's: 1 820490, 18-04-2019

onder verantwoording van it. T.G.M. Aanhane, Buainess Unit Manager. Ierestretering zijn orze Algemene Voorwaarden van bepeseing. Op verzoek endoarlew van de aansjeventhoden to sugeordaan. Euroffan Agro Teating niet aansonkelijk voor eventuele schadelijke gevolgen voorhivierind niet aansonkelijk voor eventuele schadelijke gevolgen voorhivierind e en/of de s





Agro

Meststoffenonderzoek Anorg. vaste enkelvoudige GROF FILT

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	Origineel				
Monster	Onderzoek-/ordernr: Datum monster 820489/004683869 29-03-2019		ername:	Datum verslag: 18-04-2019	
	Code onderzoek: 485	Datum ontvan 18-04-2019	gst:	Monster genomen door: Derden	Contactpersoon monstername:
Resultaat	EC pH	9,3 mS/a 6,2	m 25°C		
		mmol/l	ppm	(=mg/l)	
	Nitraat (NO3)	< 1,0	< 63		
	Ammonium (NH ₄)	3,6	65		
	Fosfor (P)	2,03	63		
	Kalium (K)	57,6	2252		
	Calcium (Ca)	12,3	493		
	Magnesium (Mg)	6,7	163		
	Zwavel (S)	31,2	1000		
	Natrium (Na)	10,8	248		
	Chloride (CI)	29,5	1046		
	Bicarbonaat (HCO3)	< 1,0	< 62		
	Silicium (Si)	0,49	13,76		
		µmol/l	ppm	(=mg/l)	
	Borium (B)	23	0,25		
	Koper (Cu)	6,2	0,39		
	IJzer (Fe)	22	1,23		
	Mangaan (Mn)	56	3,08		
	Molybdeen (Mo)	< 1,0	< 0,10	0	
	Zink (Zn)	10	0,65		

De resultaten zijn uitgedrukt als de wateroplosbare fractie van het aangeboden monster. Zij kunnen niet dienen als toetsing van de meststoffenwet.

Methode Alle bepalingen

Em: MENGMS en DAHFD

Methode geaccrediteerd door RvA
 Em: Eigen methode, Gw: Gelijkwaardig aan, Cf: Conform
 Bij deze verrichting is de gestelde houdbaarheidstermijn tussen monstername en analyse
 overschreden. Dit heeft mogelijk de betrouwbaarheid van het resultaat beinvioed.
 De resultaten hebben uitsluitend betrekking op het in behandeling genomen materiaal.

Pagina: 1 Totaal aantal pagina's: 1

ort is vrijgegeven onder venntwoording van Ir. T.G.M. Aanhane, Business Unit Manager. ze vormen van diemstverlening zijn erze Algemene Voorwaarden van toepassing. Op verzoek doez entrid ne specificatie van die antisjementhoon toepaconden. Euroffna Ago Testing ngen BV statt zich niet aamsprakelijk voor eventuele schadelijke gevolgen voorhioelend abuit oor, dovr of aamsen zwa vestetate naterizzeisereeutsten en off ad viceen.



820489, 18-04-2019

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Rapport

Agro

Meststoffenonderzoek Anorg. vaste enkelvoudige FILT + UV

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F +31 (0)88 876 1011 E klantenservice.glastuinbouw@eurofins-agro.com www.eurofins-agro.com

Origineel Monster Onderzoek-/ordernr: Datum monstername: Datum verslag: 820491/004683869 29-03-2019 18-04-2019 Code onderzoek: Datum ontvangst: Monster genomen door: Contactpersoon monstername: 485 18-04-2019 Derden Resultaat EC 6,9 mS/cm 25°C pН 4.7 mmol/l ppm (=mg/l) Nitraat (NO₃) 2.3 143 Ammonium (NH₄) 1.8 32 Fosfor (P) 2,65 82 Kalium (K) 41,4 1619 Calcium (Ca) 9,4 377 119 Magnesium (Mg) 4.9 22,3 715 Zwavel (S) Natrium (Na) 8,4 193 Chloride (CI) 20,2 716 Bicarbonaat (HCO₃) < 1,0 < 62 Silicium (Si) 0 47 13 20 ppm (=mg/l) µmol/l 19 0,21 Borium (B) Koper (Cu) 5,1 0,32 IJzer (Fe) 43 2,40 Mangaan (Mn) 47 2,58 Molybdeen (Mo) < 1.0 < 0.10 0,72 Zink (Zn) 11

De resultaten zijn uitgedrukt als de wateroplosbare fractie van het aangeboden monster. Zij kunnen niet dienen als toetsing van de meststoffenwet.

Methode Alle bepalingen

Em: MENGMS en DAHFD

 Q
 Methode geaccrediteerd door RvA

 Em:
 Eigen methode, Gw: Gelijkwaardig aan, Cf: Conform

 *
 Bij deze verrichting is de gestelde houdbaarheidstermijn tussen monstername en analyse overschreden. Dit heeft mogelijk de betrouwbaarheid van het resultaat beinvioed.

 De resultaten hebben uitsluitend betrekking op het in behandeling genomen materiaal.

Pagina: 1 Totaal aantal pagina's: 1

820491, 18-04-2019

Is vrijgegeven onder verantwoording van Ir. T.G.M. Aanhane, Busitess Unt menagen. vomen van diemskreitering zijn onze Algemene Voorwaarden van bepossing. Op verzoek ze enrid de specificatie van de analysemethichet negozoedhe. Eurotisk ago Teeling in BV stelt zich niet aansprakelijk voor eventuele achabelijk gevolgen voortvoelend Martin and erdozen.



Annex 3 Pictures of well plates before and after the test (7 days)



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