



THE APPLICABILITY OF DISINTEGRATION TESTS FOR COHESIVE ORGANIC SOILS

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Abstract. The use of ripened fine-grained organic dredged materials as construction materials, e.g. as top soil on slopes such as landfills or dikes, is an important contribution to environmental engineering science. The materials are legally considered a waste and need to be beneficially re-used. Therefore, not only standard geotechnical parameters have to be determined but also their erosion resistance which is a particularly critical environmental parameter. There is a variety of different tests to determine the flow dependent erosion resistance of soils, such as the erosion function apparatus (Briaud *et al.* 2001). In this study, however, the focus lays on the aggregate stability as an indicator for the erosion resistance under static loading, which can be determined using wet sieving and disintegration tests. The disintegration tests after Weißmann (2003) and Endell (RPW 2006) have a similar setup; however, the specific boundary conditions for the tests as well as the evaluation procedures are different. Weißmann proposed his test to determine the erosion stability of dike cover materials while the Endell test should be used for mineral sealing liners in navigation channels. In this study both tests have been used to evaluate the aggregate stability of fine-grained organic dredged materials that have been installed in large-scale research dike facilities and in the recultivation layers of different landfills. The materials showed good visual performance with respect to rainfall induced erosion so far; however, problems in determining erosion and aggregate stability indices limit the value of the studies: both disintegration tests investigated have major limitations with respect to the organic soils tested. Particularly the evaluation methods are not suitable for the soils but also some boundary conditions are critical and are discussed in this paper. The gained knowledge is a valuable basis for the development of standard characterisation methods for dredged materials in environmental and geotechnical applications.

Keywords: soil erosion, disintegration tests, aggregate stability, waste management technologies, cohesive organic soils, dredged material, marsh clay, marl, environmental sustainability.

Introduction

Soil erosion plays an important role both in agricultural and geotechnical environments. In agriculture the workability and long-term availability of arable soils are very important issues while the erosion stability of slopes, (e.g. at dams and dikes) is of great interest in the geotechnical context. Most importantly, international soil protection legislation states that the top soil should always be protected against relocation by erosion. Coastal dikes are often constructed in environmentally sensitive areas and in spite of their flood protection function and the associated maintenance as a coastal protection structure dikes are large environmental earth constructions that characterise the

coastal landscape. They are usually made of a sand core, a cohesive top layer with high erosion resistance, and a grass cover. At the German Baltic Sea coast the cohesive layer is usually made of marl while at the North Sea coast it is mostly made of marsh clay which sometimes has considerable organic contents. During the last few years also dried fine-grained organic dredged materials have been investigated to be used in dike and landfill construction as protective and/or recultivation layers. These materials can be characterised as young cohesive organic soils, which is the focus of this paper.

Dredging means the removal of sediments from water bodies by excavators or hydraulic dredgers. It is a

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necessary operation in waterway and harbour maintenance but also in hydraulic construction works and ecological revitalisation projects. The major amount of the removed sediments is usually relocated within the water body which means the sediments are transported to another location and dumped to underwater reservoirs. However, due to environmental sustainability the relocation of contaminated and cohesive organic sediments is restricted in most areas of the world. In the Baltic Sea there is, for example, a general understanding under the HELCOM convention only to relocate clean sands (regarding precaution values usually provided by national law of the Baltic Sea bordering states).

Cohesive organic dredged sediments are usually taken ashore to be deposited on special land-fills (mainly if contaminated) or to be re-used beneficially according to the respective environmental regulations, e.g. the soil protection law, ground water protection law, and other soil-water related regulations (in Germany: BBodSchG 1998; BBodSchV 1999; WHG 2009). The re-use of fine-grained organic dredged materials in agriculture and landscaping has become an acknowledged method (Sigua *et al.* 2004; Canet *et al.* 2003). Usually the materials are processed before re-use. In most cases the materials are therefore pumped into containment areas where they are dewatered. After some time, the mud is taken out of the dredging polders and put on heaps on adjacent drying fields where the ripening process starts. The so ripened soil-like material is usually an in-homogenous mix of mineral and organic soil particles, depending on the dredging project. However, sometimes the dredged materials are also separated in so-called classification polders where the coarse and medium coarse sandy material settles around the inlet pipes, followed by different mixed soils with decreasing sand and increasing silt, clay and organic fractions down to the polder outlet where the very fine-grained organic sludge is deposited. Then particular batches can be classified and separately processed to receive better qualities for the re-use of the materials. The Hanseatic City of Rostock successfully runs two such containment areas as waste management technology with classification polders to process defined ripened soils for re-use. The mixed soils and fine-grained organic materials have been used in landscaping, agriculture and recultivation layers on landfills so far.

The project DredgDikes deals with the application of ripened dredged material in dike construction. Within the project three different cohesive organic dredged sediments have been installed in a full scale research dike facility (Cantré, Saathoff 2013). All materials are generally cohesive soils with an organic content of five to ten per cent. Visually, they seem to have considerable erosion resistance when installed and compacted with caterpillars. In contrast, laboratory flume tests showed rather low erosion resistance with respect to overflowing. Particularly for coastal and river dikes the

time dependent stability of the erosion protection layer with respect to a static water loading is of great interest because the soil aggregates may decompose due to aggregate explosion and swelling phenomena which may lead to dike failure. Therefore, two different disintegration tests have been investigated to study the aggregate stability of the materials as an index for the erosion resistance under static conditions. However, some important limitations regarding both the test setups and the evaluation methods disparage their conclusiveness which will be discussed in this paper. The results will be used to develop a set of characterisation tools for dredged materials in environmental and geotechnical engineering applications.

1. Basics

1.1. Aggregate and erosion stability

Erosion stability is defined as the stability against “the replacement and the transport of soil particles along the surface” (Blume *et al.* 2010) and can be characterized as a combination of the stability of the overall soil texture and the stability of the aggregates within the texture, which are agglomerations of fines of different mineral grains mixed with organic carbon, lime and other soil components (Le Bissonnais 1996).

Surface erosion can be triggered from water, wind, snow melting or gravitation and is influenced by different processes like rainfall or surface run-off. Here, only water induced erosion is investigated. The main processes of soil erosion by water are soil displacement, soil transport and soil deposition (Blume *et al.* 2010). The soil displacement is the release of erodible material and includes sub-processes like the destruction of soil aggregates and the siltation of bounded particles. The soil structure consists of single grains or aggregates which are characterized by their external shape and size or their internal structure (Blume *et al.* 2010). The stability of soil aggregates differs with respect to water and pressure which influence the mechanical capacity, the siltation and the erosion risk. An aggregate can be described as stable if the particle position does not change by voltage variation. Also, high organic contents improve the stability of aggregates against water immersion due to inter-particle cohesion with organic carbon (Chenu *et al.* 2000; Cantón *et al.* 2009; Dal Ferro *et al.* 2012).

The definition of aggregate stability and soil structure stability is not standardised and both definitions are often used synonymously (Ametzketta 1999). The German DIN standard 19683-16 (2009) defines aggregate stability as an important parameter for the siltation of soils and provides a comparison of the stability of different agricultural farmlands. Here, aggregate stability is defined as the resistance of soil particles to their structure (Blume *et al.* 2010). Hartge and Horn (2009) describe aggregate stability as the

resistance of aggregates against destructive triggers which can be expressed by a shear parameter. The determination of the aggregate stability against water is not trivial because it depends on the surface binding energy of the soil particle, the water cohesion and the adhesion forces. Adhesion is caused by suspended solids that may deposit at the contact points of the soil particles and thus glue the particles together. Both the building and reversibility of aggregates depend on the degree of drying and the type of rehydration and thus on the aggregate size, aggregate density and the pathways through intra aggregate pores are important parameters (Hartge, Horn 2009). Le Bissonnais (1996) also describes a high dependence of water infiltration and soil erosion by aggregate stability. In his opinion the erosion rates are the results of the aggregate disintegration divided into disintegration of macrostructure and dispersion. The aggregate disintegration through water can be the result of a variety of mechanisms, four of which are particularly important (Le Bissonnais 1996):

I. Slaking occurs with overpressure through enclosed air during moisture infiltration and appears when dry aggregates are put fast into water. The disintegration decreases with increasing clay content.

II. Swelling and slaking during wetting and dehydration lead to aggregate blasting. The disintegration increases with increasing clay content.

III. Soil fragments are dissolved by the splash effect if the kinetic energy of rain drops is high enough to cause disintegration. This phenomenon occurs most often with wet soils because the aggregates are weaker and can be dissolved more easily.

IV. Physico-chemical dispersion means the reduction of attraction between single colloid particles during drainage of soils and is facilitated with slaking and swelling. It is one of the most effective processes of aggregate destruction. Auerswald (1993) on the other hand focuses on the influence of the initial soil moisture and the resulting erosion process to explain aggregate stability. Investigations about the shear strength from soils or aggregates show lower strengths with increasing moisture. Investigations with a sudden wetting, however, show a lower aggregate blasting at increasing moisture content. Hereby, the initial moisture seems to have an influence on the mechanisms of temperature weakening, impact hardness, air blasting and swelling pressure with respect to aggregate disintegration (Auerswald 1993):

I. With penetrating water a heat front proceeds into the aggregates due to an energy release that weakens the aggregate bounding substances. This mechanism is called temperature weakening and is important for soils with moistures below 10% or above 30% (gravimetric).

II. The soil stiffness exponentially increases with decreasing soil moisture, thus the impulse recurrence from a rain drop to the soil increases.

III. The decrease of soil moisture causes an increase of air filled pores. If water penetrates quickly into an aggregate without air escaping, an overpressure occurs inside the aggregate, leading to air blasting. This phenomenon increases with decreasing moisture content.

IV. Quick and irregular wetting of aggregates causes irregular swelling. The resulting drag stress causes aggregates to break-up.

1.2. Methods to determine erosion stability

To determine the soil erosion stability a variety of methods have been proposed (Table 1) which can be divided into tests to determine the erosion resistance of the whole soil texture (e.g. laboratory flumes or erosion function apparatus after Briaud) and tests to determine the aggregate stability (e.g. wet-sieving methods, flow-rate methods, and water-immersion methods). In the following, different tests to determine the aggregate stability will be briefly reviewed.

For the wet-sieving tests air-dried soil aggregates are put into a sieve-diving apparatus with different sieve sizes to be moved vertically through a liquid medium (water or ethanol). Thereby, the instable aggregates are resolved in the liquid and the remaining grains are automatically sieved. The mass left on the sieve is measured and related to the initial mass.

Table 1. Overview of tests to determine aggregate stability

Method	Test
Wet-sieving	German DIN-standard 19683-16
	Hartge/Horn
	Le Bissonnais
Flow-tests	Pinhole test after ASTM D 4647-06
	Modified pinhole test after BAW
Dispersion-tests	Dispersion test after ASTM D 4221-99
	Crumb-test after ASTM D 6572-06
	Modified crumb test after Haghghi
	Disintegration test after Weißmann
	Disintegration test after Endell (BAW)

The pinhole test after ASTM standard D 4647-06 and the modified pinhole test (RPW 2006) represent direct qualitative measurements of the dispersibility of compacted clay. A defined hole is put into a compacted soil sample and a water flow is realised through the hole. After the test the increase in hole diameter is measured and related to a degree of aggregate stability.

The disintegration of aggregates or compacted samples in water immersion is analysed using dispersion tests. They can be performed using the crumb test after ASTM standard D 6572-06, the modified crumb test after Haghghi *et al.* (2012), the dispersion test after ASTM

standard D 4221-99 or different disintegration tests. In this paper the disintegration tests after Endell (enhanced by the Federal Waterways Engineering and Research Institute, BAW) and Weißmann have been studied.

2. Investigated soil materials

For the investigations five different soils were used: three different fine-grained, organic dredged material batches from a containment area near Rostock, Germany and two standard dike cover materials (marl and marsh clay). Apart from their composition the dredged materials differ in their ripening time: materials M1 and M3 had been ripened for 5 years, while material M2 had been ripened for



Fig. 1. Locations from where the soil samples were taken

Table 2. Reliable results of geotechnical characterisation

Material	M1	M2	M3	Marl	Marsh clay
Clay [%]	25–28	22–25	15	5	17
Sand [%]	29–34	40–47	54	89	46
W [%]	61–68	55–73	46	13	21
OM [%]	10–11	9–10	6	0.9	1.7
LC [%]	9–10	8	10	12	0.9
c_u [kPa]	53–132	19–34	120	24	83
φ [°]	28–30	28–31	30	42	66
c [%]	35–47	13–19	59	15	3
k_f [m/s]	5E-08	8E-10	5E-09	4E-08	4E-10

Table 3. Preliminary results of geotechnical characterisation

Material	M1	M2	M3	Marl	Marsh clay
LL [%]	80–98	64–88	52–57	17	36
PL [%]	75–81	54–67	49–54	13	22
SL [%]	58	42–47	51	16	22
PI [%]	4–22	11–24	3–4	4	14
CI [-]	2–5	0–1	2–4	1	1
w_{opt} [%]	40–43	32–35	31	9.2	14
OD [g/cm ³]	1.1–1.2	1.3	1.4	2.1	1.8

two years at the time of the investigation. For materials M1 and M2 three subsamples were taken respectively. The marsh clay batch was sampled from a dismantled dike cover layer in Hamburg Port while the marl was originally extracted from the Baltic Sea off the coast of the Island of Rügen (Fig. 1).

All materials were investigated with respect to their soil mechanical characteristics such as water content, grain-size distribution, compaction parameters, shear strength, water permeability and shear parameters, principally according to the German DIN-standards for soil mechanical analyses (DIN... 1998). Because of the high organic content of the dredged materials the grain-size distribution was determined according to DIN ISO 11277 for fine-grained, organic soils. For the organic content the TOC (Total Organic Carbon) value was determined after DIN ISO 10694 with an elemental analyser at temperatures above 1000 °C, because the marine dredged sediments possess high lime contents which influence the results from a 550 °C muffle furnace.

The results of the geotechnical characterization are shown in Table 2 and Table 3. Because of different challenges using the standardized DIN characterization for the specific dredged materials the tables are divided into reliable and preliminary results.

3. Disintegration tests

For both disintegration tests in this study cylindrical compacted samples with different water contents are put into a wire mesh basket which is connected to an electric scale. The disintegration is measured by recording the weight loss caused by the dropping of aggregates at water immersion. The dropping of aggregates is described as “crumbling” in the following while disintegration describes the status of the overall sample. Both disintegration tests use the weight loss as a function of time for further analysis. The principle of the test procedure is shown in Figure 2. In spite of the similar setup of the disintegration tests and similar processes studied the tests differ in several important issues which will be discussed in the following.

3.1. Disintegration test after Endell

The disintegration test after Endell (enhanced by the Federal Waterways Engineering and Research Institute, BAW) is used to estimate the erosion susceptibility of clay to be used in waterways construction. Sealing compounds with higher disintegration are more sensitive to erosion which is especially interesting while installing mineral sealing liners while they are still uncovered (RPW 2006).

For the analysis at least five proctor compacted samples (proctor power of 0.6 MNm/m³) with a diameter of 2 cm and a height of 4 cm are used. The samples

are each put into a wire mesh basket which is not specified with respect to its dimensions. It is connected to an electric scale and put into a water basin with distilled water (Fig. 2).

After the RPW (2006) the weight has to be recorded until a constant weight is reached, but not less than 24 hours. The result of this test is the disintegration number $Z(t)$ as a function of time (Eqn (1)):

$$Z(t) = \frac{A1 - A(t)}{A1 - A2}, \quad (1)$$

where: $Z(t)$ – disintegration number as function of time; $A1$ – uplift sample weight including mesh basket weight [g] at the beginning; $A(t)$ – uplift sample weight including mesh basket weight [g] at the time t of the test; $A2$ – uplift mesh basket weight [g] without sample.

To compare different samples the disintegration number after 8 hours $Z(8)$ is proposed. Therefore, it must be ensured that all dropped particles drop out of the wire mesh basket; otherwise, the test has to be repeated. According to Endell there is one specific moisture content for every soil at which the most significant disintegration occurs. Therefore, the test has to be carried out with at least five different water contents resulting in different undrained shear strengths (e.g. $c_u = 10 \dots 70$ kPa). The lowest disintegration should be expected at a water content near the liquid limit because of a reduced water immersion in saturated samples.

RPW (2006) specifies a limit value for the disintegration number of $Z(8) = 0.05$. Above this value the soils are defined as susceptible to erosion while soils with a value $Z(8) < 0.05$ are defined as erosion resistant. However, there is no standardised limit value for the disintegration number yet.

3.2. Disintegration test after Weißmann

The test equipment of Weißmann's disintegration test was developed independently to that of Endell to determine the erosion resistance of marsh clay embankments on sea dikes. The test facility is similar to that of Endell, except little differences (Fig. 3). The wire mesh basket is defined with meshes of 8×8 mm² and an overall size of $10 \times 10 \times 12$ cm³. The soil samples are bigger than those of the Endell test (5 cm for both diameter and height). The water temperature shall have 20 °C during the entire test and tap water should be used. Nine cylindrical samples have to be prepared in a Proctor apparatus with optimum water content (standard proctor power of 0.6 MNm/m³) to guarantee good compaction.

From the nine samples three test series with different water contents are prepared (three repetitions):

- (1) Dry sample – dried at 50 °C to a constant mass;
- (2) Standard sample – optimum water content;

(3) Wet sample – lagged with filter paper and put into a water basin for ten minutes, then kept in a desiccator for 24 hours.

The result of the Weißmann test is the time t_{30} when a sample lost 30% of its initial weight. The maximum testing time is 24 hours, even if the 30% weight loss cannot be reached; then the material is classified as erosion resistant. According to Weißmann the disintegration time increases with increasing water content and can be described approximately with an exponential curve, shown in

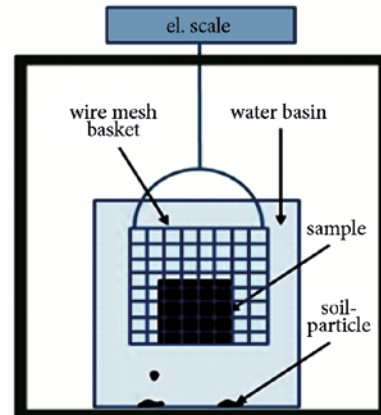


Fig. 2. Schematic of disintegration tests

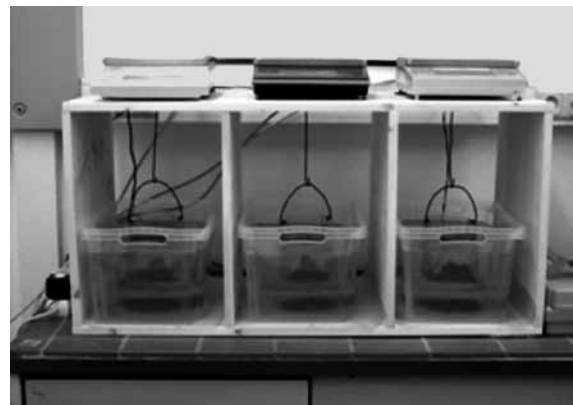


Fig. 3. Disintegration equipment after Weißmann

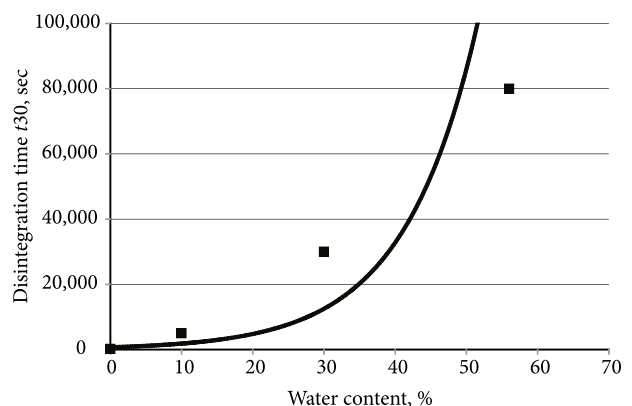


Fig. 4. Diagram of disintegration time depending on water content after Weißmann

Figure 4. After entering the results of the three test series into this diagram the disintegration number $t_{30,V}$ can be calculated with Equations (2)–(5).

For determining the auxiliary parameters A and B (Eqns (3) and (4)) two not defined water contents have to be chosen and the associated disintegration times have to be taken out from the diagram. The reference value $t_{30,V}$ is used at a Consistency Index $I_c = 0.8$ depending on the associated water content w_v , which can be calculated from the liquid and the plastic limit (Eqn (5)). Finally the disintegration time $t_{30,V}$ can be calculated at w_v with Equation (2):

$$t_{30}(w) = t_{30,0} + A \cdot w \cdot e^{B \cdot w}; \quad (2)$$

$$B = \frac{\ln((t_{30(w1)} - t_{30,0}) \cdot w_2) - \ln((t_{30(w2)} - t_{30,0}) \cdot w_1)}{w_1 - w_2}; \quad (3)$$

$$A = \frac{t_{30(w1)} - t_{30,0}}{w_1 \cdot e^{B \cdot w_1}}; \quad (4)$$

$$w_V = 0.2 \cdot w_L + 0.8 \cdot w_P, \quad (5)$$

where: $t_{30,0}$ – disintegration time t_{30} at $w = 0\%$; A, B – auxiliary parameter; w_v – water content at $I_c = 0.8$.

3.3. Modification of the test procedure for the dredged materials

Initial tests showed difficulties in evaluating the results of the dredged materials with both tests. Therefore, some modifications were necessary. For a better understanding of the modifications, the original attributes of the Endell and Weißmann tests are compared in Table 4.

For the Endell test one sample at natural water content w_n and two samples with water contents $w > w_n$ and $w < w_n$ were chosen respectively. Samples with $w < w_n$ were dried step by step until the selected water content was achieved. To produce samples with $w > w_n$ water was added and the material was mixed. Both water addition and sample drying were restricted to values in an interval in which good compaction could be ensured for all samples. The wire mesh basket was made cylindrical having a mesh size of 8 mm and a diameter and height of 3.2 and 5 cm respectively. In this way the small samples would not fall through the mesh and could be installed upright inside the basket.

The experiments after Weißmann were modified after initial tests with the dredged materials. Instead of the optimum water content the natural water content was used for sample preparation. The proposed optimal water content at Proctor's density of the inhomogeneous dredged materials could not be determined reliably. Also, all samples compacted at natural water content showed little spread in the disintegration curve and so ensured comparability of all three repetitions.

For both tests the recording interval with respect to weight loss was chosen to 5 s to ensure a precise measurement of the disintegration during the first few minutes of testing. Every subsample was tested for at least 24 hours.

Table 4. Differences between the tests of Endell and Weißmann

	Endell	Weißmann
Water type	Distilled water	Tap water
Basket	Not defined. Chosen: Size: 3.2×5.0 cm ² , 8 mm mesh	Defined: Size: 10×10×12 cm ³ , 8 mm mesh
Sample size	H = 4 cm, D = 2 cm	H = D = 5 cm
Result	Disintegration number Z(t)	Disintegration time t_{30}

4. Results

4.1. Disintegration test after Endell

4.1.1. Dredged materials M1, M2 and M3

The results of the disintegration test are the process of disintegration as a function of the time and computed a disintegration number after eight hours. The functions of material M1 and M2 have similar shapes. Figure 5 shows the measured relative sample weight of material M1 relating to the disintegration time.

Already after a short time all samples start to crumble and large aggregates fall off to the bottom of the water basin which can be seen from the recorded scale data. This process generally dominates in a time interval of 10 to 1,000 s. As soon as the samples have lost about 50–70% of their initial weight this process seems to stop; for the rest of the time the recorded weight remains nearly constant. However, there seems to be an exception for samples with low water contents: shortly after the start the dry material absorbs water and the weight increases to about 120% of the initial weight. After some time the increase stops and the recorded weight shows both the largest and fastest decrease among all samples.

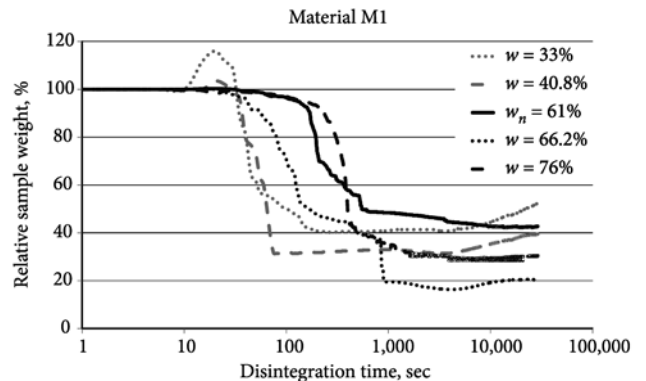


Fig. 5. Disintegration curves (Endell) of material M1

Table 5. Disintegration number $Z(8)$ of material M1 and M2

Material	w [%]	$Z(8)$ after 8 h
M1	33.0	0.7273
	40.8	0.6056
	$w_n = 61.0$	0.5728
	66.2	0.6954
	76.0	0.6056
M2	30.0	0.7273
	42.0	0.3991
	$w_n = 54.0$	0.4714
	78.0	0.4784
	79.0	0.6348

Table 6. Disintegration number $Z(8)$ of material M3

Material	w [%]	$Z(8)$ after 8 h
M3	21.9	0.9206
	36.0	0.9958
	36.9	0.7273
	49.1	0.8033
	$w_n = 49.7$	0.5425
	54.6	0.9524
	57.8	0.9003

Later during the test the initially dry samples seem to start to absorb water again, resulting in a second weight gain. The analysis of the disintegration numbers after eight hours show the lowest disintegration (the best erosion stability) for the natural water content w_n for material M1 and for a water content below w_n for material M2. Table 5 gives an overview about the disintegration numbers $Z(8)$ of the materials M1 and M2.

The more sandy dredged material M3 shows a different set of disintegration curves. Almost all samples show a high disintegration rate. The sample weight reduces to nearly zero within approximately 100 s with the exceptions of the samples with $w_n = 49.7\%$ and $w = 36.9\%$. The dry samples also show an initial weight increase to 110% comparable to the materials M1 and M2. Interestingly, there are samples with nearly the same water content which show very different disintegration behaviour: the sample with $w = 49.1\%$ shows a significantly higher disintegration number than that with $w_n = 49.7\%$. A similar observation was made for $w = 36.0\%$ and $w = 36.9\%$ (Fig. 6 and Table 6).

4.1.2. Conventional dike construction material marl and marsh clay

The development of the marl disintegration partly differs from that of the dredged materials, in particular regarding materials M1 and M2. Compared to the dredged materials, the tested marl can only absorb little amounts of water

because of its large sand and small clay fraction. The maximum possible water content to produce proctor samples was $w = 14.1\%$. The crumbling of the different marl samples proceeded very fast and in steps (Fig. 7).

The sample with the natural water content shows the lowest disintegration number, a result that was also found for the dredged materials M1 and M3 (Table 7). Some samples even show a full disintegration like material M3.

The investigated marsh clay shows an explicit distinction of the disintegration curves with respect to the water contents of the subsamples (Fig. 8).

The best result was again achieved with the w_n sample. The driest sample crumbled fastest with a total weight loss of 98% and a very high disintegration number $Z(8) = 0.9874$ (Table 8). The disintegration numbers with $Z(8) = 0.2589$ and $Z(8) = 0.2834$ for the samples with w_n and $w = 23\%$ are the best results of all materials tested.

Table 7. Disintegration number $Z(8)$ of marl

Material	w [%]	$Z(8)$ after 8 h
Marl	5.6	1.0
	7.8	0.8823
	9.2	0.9821
	$w_n = 11.8$	0.6838
	14.8	0.807

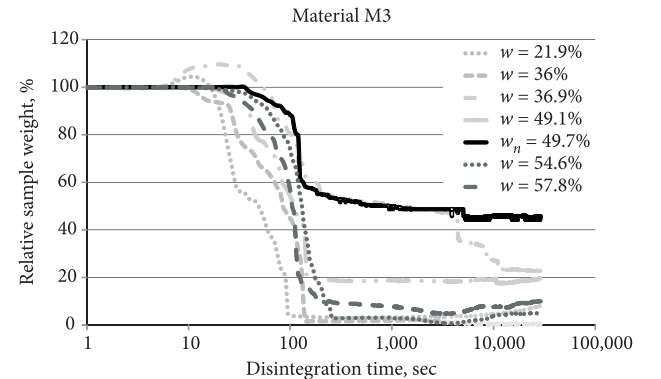


Fig. 6. Disintegration curves (Endell) of material M3

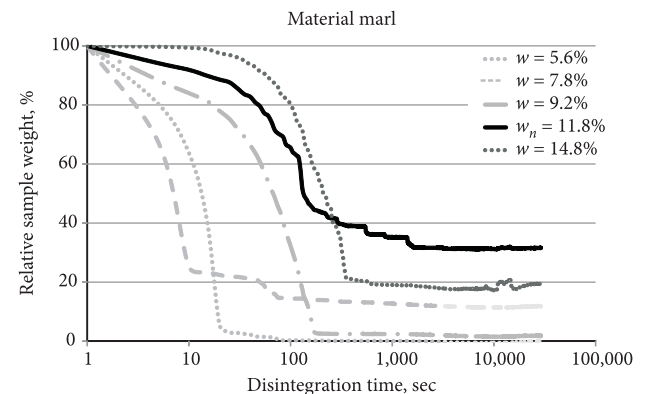


Fig. 7. Disintegration curves (Endell) of marl

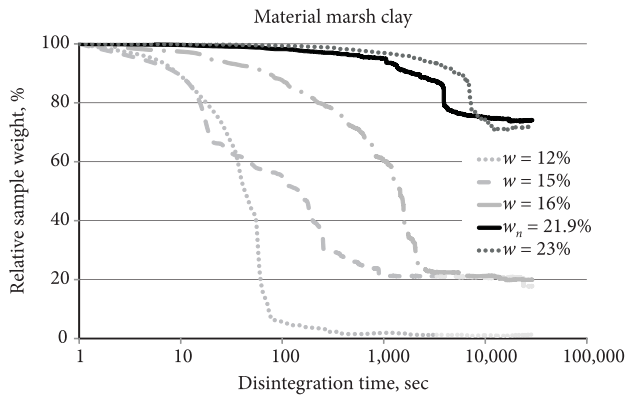


Fig. 8. Disintegration curves (Endell) of marsh clay

Table 8. Disintegration number $Z(8)$ of marsh clay

Material	w [%]	$Z(8)$ after 8 h
Marsh clay	12.0	0.9874
	15.0	0.8216
	16.0	0.8008
	$w_n = 21.9$	0.2589
	23.0	0.2834

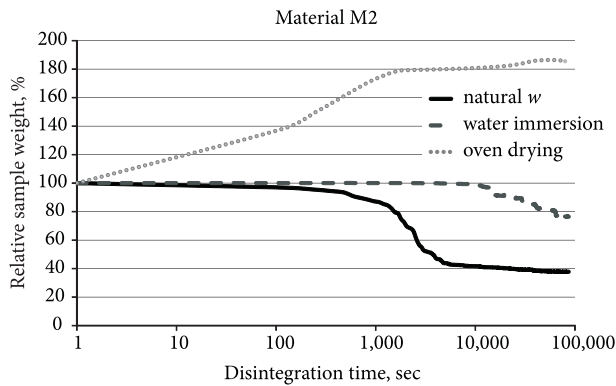


Fig. 9. Disintegration curves (Weißmann) of material M2

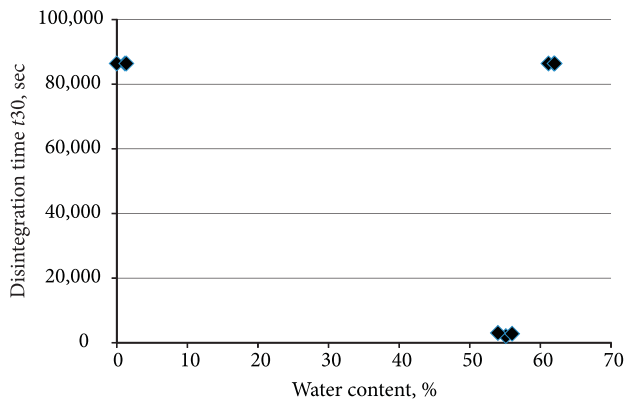


Fig. 10. Disintegration time t_{30} depending on water content for material M2

4.2. Disintegration test after Weißmann

4.2.1. Dredged materials M1, M2 and M3

The results of the disintegration tests after Weißmann are the disintegration curves as functions of time as well as the disintegration time t_{30} for three defined water contents. Depending on the available amount of material two subsamples were chosen for the investigation of materials M1 and M2 and only one for material M3. Since all dredged materials show similar curve shapes for the different sample preparation modes the curves of M2 are presented exemplary (Fig. 9).

The weight of the oven-dried samples started to increase constantly as soon as they were submerged. The weight gain approached up to 260% of the initial weight although aggregates had fallen off the sample. During the test large air bubbles were observed to come out of the soil sample.

The “water immersion” samples which were initially wrapped in filter paper show the highest stability. For a considerable period of time these samples did not lose significant weight before they started to crumble after about 10,000 s. However, there are also intervals in which no weight change was recorded although particles fell off the sample. At the end of the test the water immersion samples started to disintegrate to a relative weight loss of about 20% with no further change.

The w_n samples show the highest disintegration of the three conditions. They usually started to crumble after about 1,000 s and show a final weight loss of about 60%.

Because both the oven-drying and the water-immersion samples show no weight loss of 30% the disintegration time has to be defined to 86,400 s which is why no exponential disintegration curve can be determined (Fig. 10).

The dependency of disintegration and water content cannot be described with this data and consequently no disintegration time $t_{30,V}$ can be calculated. Only the disintegration time after 30% mass loss of the samples with natural water content can be determined and compared (Table 9).

Table 9. Disintegration time $t_{30}(w_n)$ of M1, M2 and M3

Material	Disintegration time $t_{30}(w_n)$ [s]
M1	1.600 and 1.320
M2	2.600 and 4.000
M3	1.120

4.2.2. Conventional dike construction material marl and marsh clay

The marl samples show a different behaviour. After few seconds all samples started to crumble and lost a large

amount of mass within a short period of time (Fig. 11). After 2,000 s the disintegration was already finished with a final weight of 20 to 30% of the initial mass.

As distinguished from the investigations with dredged material the optimal water content w_{opt} was used instead of the oven-dried because the oven-dried samples showed only weight increase instead of mass loss. For further results of the disintegration of oven-dried marsh clay samples compare Beyer *et al.* (2012). For the installation of w_{opt} the material was oven dried at 55 °C and rewetted afterwards. The result of the disintegration time at w_v is $t_{30,V} = 731$ s (Table 10).

Table 10. Disintegration time t_{30} of marl

Parameter	w [%]	t_{30} [s]
w_n	11.5	482
w_{im}	12.43	672
w_{opt}	12.0	178
w_v	13.72	731

Table 11. Disintegration time t_{30} of marsh clay

Parameter	w [%]	t_{30} [s]
w_n	24.0	60,098
w_{im}	43.71	86,400
w_{opt}	13.4	373
w_v	25.08	62,438

The disintegration curves of marsh clay show that the crumbling of samples with w_n started late and the disintegration of 30% was reached very late: $t_{30} = 60,000$ s. The “water immersion” samples show hardly any mass loss during the whole investigation (maximum 2%). The samples with w_{opt} on the contrary showed a result comparable to that of the marl. Here, crumbling started already after 100 s and resulted in an almost complete mass loss (Fig. 12). The disintegration time at w_v is $t_{30,V} = 62,438$ s (Table 11).

5. Discussion

During the investigation both benefits and major limitations of the different disintegration tests could be observed. There is a general problem of both tests with respect to the testing procedure – a sample in a wire basket submerged in water. However, the two experimental setups show test specific limitations also. Finally, the results are discussed in comparison to test results from a dynamic erosion test using a small scale laboratory flume and to data taken from the literature.

5.1. Testing procedure

The testing procedure of both disintegration tests is based on a sample inside a wire mesh basket which is submerged

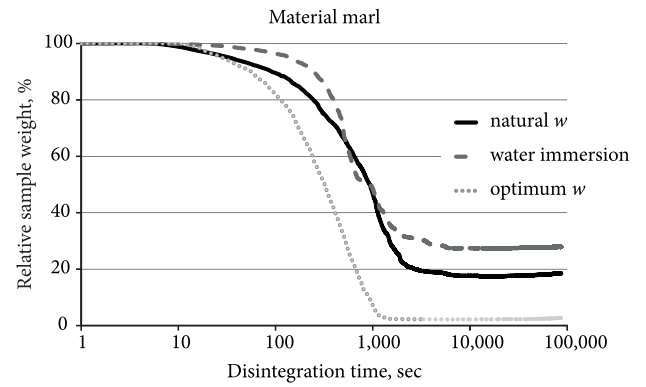


Fig. 11. Disintegration curves (Weißmann) of marl

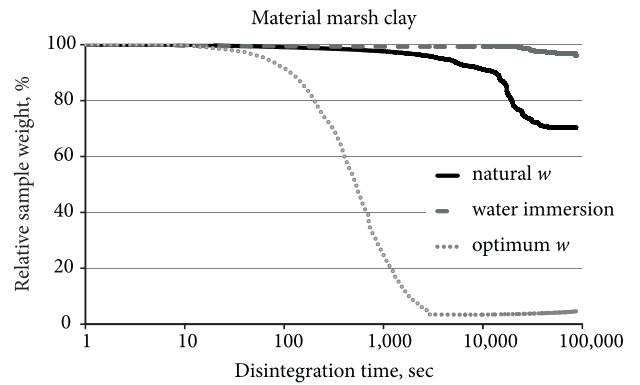


Fig. 12. Disintegration curves (Weißmann) of marsh clay

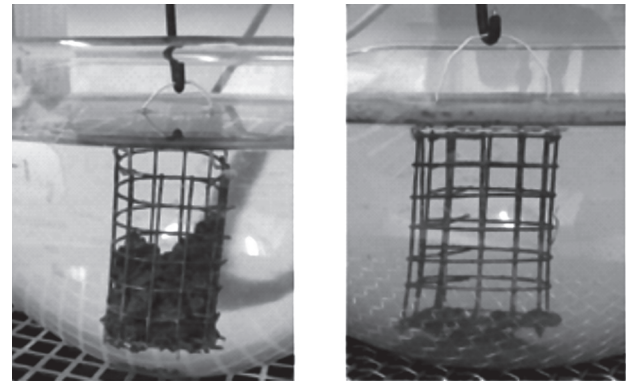


Fig. 13. Total disintegration with different mass loss

in water. The major limitation of this procedure which was observed in a large variety of test is, that soil aggregates that fall off the sample during disintegration stay in the basket and are not detected as disintegration by weight loss (Fig. 13 left).

This may have different reasons such as the defined small mesh size in case of the Weißmann test together with varying sizes of agglomerates that evolve during disintegration. In Figure 13 both samples are completely disintegrated with the difference that the disintegration is recorded correctly for the right sample and no weight loss is recorded for the left sample. For the Endell test the RPW (2006)

proposes to repeat the test until there is one test in which all aggregates fall down. There may be soil samples, however, which always decompose into larger aggregates where this solution will not solve the problem. Weißmann, on the other hand, gives no attention to this problem. The problem may be solved by using baskets with different mesh sizes; this may not affect the Endell method (no definitions with respect to the basket) but an adjustment to the Weißmann method would be needed. To compare different disintegration tests the same mesh size should be used, thus this parameter needs to be recorded for every test.

5.2. Disintegration test after Endell

In RPW (2006) the disintegration number after eight hours $Z(8)$ is used to specify a soil's susceptibility to erosion. The value of $Z(8) = 0.05$ above which a soil is considered erodible has been developed for mineral sealing liners. In this investigation dredged material, marl and marsh clay were tested which may perform differently with respect to the disintegration number and the actual erosion resistance. This may be caused by the materials' lower clay content compared to that of mineral sealing liners.

The lowest disintegration number was determined at w_n for the majority of samples. The most substantive explanation seems to be connected to the sample preparation which was necessary to install the different water contents. Some of the samples needed to be dried, which seems to have a considerable influence on the aggregate stability, particularly for the fine-grained, organic materials. The influence of the drying temperature in soil sample preparation on the aggregation has already been investigated by different researchers (Basma *et al.* 1994; Sunil, Krishnappa 2012). These investigations show that a higher drying temperature leads to an accumulation of fine particles with organic and to larger grains with decreasing specific surface area (Mikutta *et al.* 2005). There is also an irreversible dehydration process of the clay particles causing cementation of fine particles which are then not dispersed (Sunil, Krishnappa 2012). These effects can even be detected in a standard grain-size analysis. The "reduction" of fine particles also reduces their cementing ability and the samples start to crumble faster.

A second explanation may be found in the fast infiltration of the dried materials once they are submerged which leads to a quick displacement of air confined in the soil sample (Rohoskova, Valla 2004). Large air bubbles escape the samples due to over-pressure, usually causing "aggregate blasting" (Auerswald 1993; Le Bissonnais 1996)

For dredged materials the preparation with water addition leads to a higher disintegration. At the beginning higher water contents lead to less crumbling because more

micro-pores are pre-filled with water. Then the behaviour changes and more aggregates fall off, possibly because the aggregate cohesion is weakened through the initial water immersion. No dependences of disintegration and water content could be found for the dredged materials (compare Rudat 2012).

According to Dal Ferro *et al.* (2012) and Cantón *et al.* (2009) there is a stabilisation of aggregates by organic carbon. In the investigation no correlation between organic content and disintegration could be validated. Both fine-grained dredged materials M1 and M2 have high organic and clay contents but show lower erosion resistance than marsh clay with less organic content.

To compare the erosion resistance of the different materials the disintegration number $Z(8)$ w_n was chosen (Table 12). The conventional dike construction material marsh clay has the lowest disintegration number $Z(8) = 0.2589$, followed by the dredged material M2 with $Z(8) = 0.4714$. The maximum disintegration could be observed for dredged material M1 and marl.

All materials – including the conventional dike construction materials – did not even closely achieve a value of $Z(8) < 0.05$ which is the proposed erosion resistance limit for mineral liners. If the disintegration test after Endell shall be used for dike construction materials the disintegration value needs modification.

Table 12. Disintegration number $Z(8)$, w_n in comparison

Sample	$Z(8)$, w_n
M1	0.5728
M2	0.4714
M3	0.5425
Marl	0.6838
Marsh clay	0.2589

5.3. Disintegration test after Weißmann

In the disintegration tests no dependency of disintegration time t_{30} and water content could be found. All oven-dried samples showed a considerable weight increase of up to 260% of the initial weight instead of a mass loss, although moderate crumbling could be recognized for all samples. There are two reasons for this phenomenon: the high amount of water which is absorbed and the escaping air bubbles which reduce the buoyancy. According to Beyer *et al.* (2012) the influence of buoyancy cannot be eliminated in this test procedure. This may lead to questionable results for unsaturated samples.

Due to the missing dependency of water content and disintegration the disintegration time at $I_p = 0.8$ cannot be determined. Even the modification to use samples with w_{opt} instead of oven-dried samples does not show any improvement. All samples crumbled in a very short time

because of the preparation with oven-drying, rewetting and compaction (Lindh, Winter 2003).

To compare all materials the disintegration time at w_n was chosen again (Table 13). For materials M1 and M2 two subsamples were investigated respectively, for material M3, marsh clay and marl only one each.

Table 13. Disintegration time t_{30} at natural water content

Material	$t_{30,wn}$ [sec]
M1	1,320 and 1,600
M2	2,600 and 4,000
M3	1,120
Marl	482
Clay	60,098

Like in the Endell test the conventional dike construction material marsh clay shows the best disintegration behaviour among the tested materials. The best result among the dredged materials was obtained with material M2 with $t_{30,wn} = 2,600$ s and $t_{30,wn} = 4,000$ s. The sandy materials M3 and marl show the poorest results.

Based on his disintegration test Weißmann developed an evaluation method to classify the application of marsh clay as dike construction material. Additional evaluation criteria are water permeability (B1), degree of shrinkage (B3), and plasticity index (B4). As a result an evaluation number N is determined (Eqns (6)–(10)):

$$N = \sqrt[4]{B1 \cdot B2 \cdot B3 \cdot B4}; \quad (6)$$

$$B1 = 0.7 - (\log(kf) + 4) / 20; \quad (7)$$

$$B2 = 0.2 \cdot \log(t_{30,v}); \quad (8)$$

$$B3 = 1.0 - 1.25 \cdot (V_s - 0.05); \quad (9)$$

$$B4 = 0.3 + 2 \cdot I_p. \quad (10)$$

The evaluation number N can be divided into five suitability classes. Materials with suitability class 1 are very well suited as dike cover layer, while materials with suitability class 5 are not advisable (Table 14). For all materials investigated in this study the evaluation number and suitability class was determined (Table 16). In this case the disintegration time $t_{30,wn}$ was used. The parameters used to compute the evaluation numbers are presented in Table 15, partly determined using the “not reliable” results from Table 3.

All investigated materials can be classified at least as class 4 “limited suitability” and are therefore applicable as dike construction materials. The best suitability was determined for the marsh clay (suitability class 2 “well suitable”) closely followed by the dredged material M2 (suitability class 3, nearly “well suitable”).

Table 14. Suitability classes with respect to the evaluation numbers N (Weißmann 2003)

Valuation number N	Degree of suitability	Suitability class
$1.00 \geq N \geq 0.85$	Very well suited	1
$0.85 > N \geq 0.75$	Well suited	2
$0.75 > N \geq 0.65$	Suited	3
$0.65 > N \geq 0.50$	Limited suitability	4
$N < 0.50$	Not advisable	5

Table 15. Calculation parameters to determine the evaluation number

Parameter	M1	M2	M3	Marl	Marsh clay
k_f [m/s]	5E-08	8E-10	2E-08	3E-08	5E-10
t_{30} [sec]	1,320	2,600	1,120	482	60,098
V_s [-]	0.33	0.26	0.176	0.026	0.1886
I_p [-]	0.051	0.157	0.028	0.041	0.139

Table 16. Evaluation numbers N and suitability classes (SC)

	M1	M2	M3	Marl	Marsh clay
B1	0.865	0.955	0.885	0.876	0.965
B2	0.624	0.683	0.610	0.537	0.956
B3	0.750	0.750	0.843	1.000	0.827
B4	0.200	0.614	0.200	0.200	0.578
N	0.533	0.740	0.549	0.554	0.815
SC	4	3(+)	4	4	2

5.4. Comparison of disintegration tests with investigations in a laboratory flume

In the project DredgDikes full scale overflowing tests on dike slopes are planned to determine the surface erosion stability of the dredged materials M1–M3. For preliminary tests a small scale laboratory flume was used with dimensions of 2.90 m length and 0.25 m width and a variable slope inclination with a maximum of 3:1. With the associated pumps a flow velocity of 2.3 m/s and an associated shear stress of 210 N/m² can be realised. For the experimental tests 20 samples (ten vegetated and ten non-vegetated samples) with a thickness of 7 cm prepared from the dredged materials M1, M2 and M3 and tested in the flume. The degree of compaction ranged from 0.6–0.85. For reference the conventional dike construction material marsh clay was used. For the non-vegetated samples an erosion rate was determined using laser scan data as the quotient of the eroded soil volume and the volume of water.

The marsh clay showed by far the lowest erosion rate, while material M2 showed the best results among the

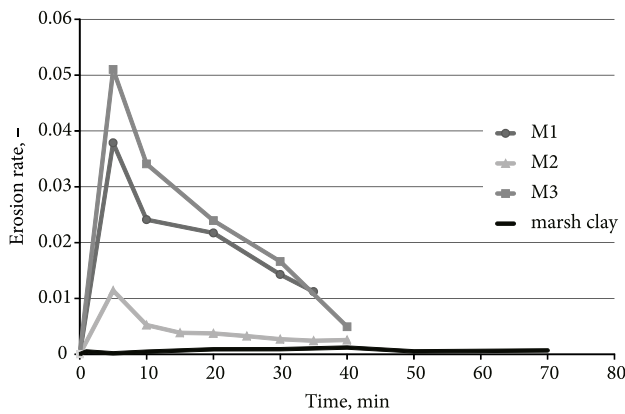


Fig. 14. Erosion rates of dredged material and marsh clay (modified after Lesch 2012)

dredged materials (Fig. 14). The samples of materials M1 and M3 in comparison experienced four to five times the erosion of M2.

When comparing the results of the flume test with those of the disintegration tests the single value of each test is related to the marsh clay values, because the lowest erosion in all investigations have been obtained at these samples. Altogether, similar orders of magnitudes can be recognized especially between the Weißmann test and the laboratory flume (Table 17), whereas the Endell test results show a smaller difference between the disintegration of marsh clay and the dredged materials. The reasons for this may be the small Endell test samples which are quickly wetted, leading to a comparably fast disintegration of all samples.

Table 17. Comparison of orders of magnitudes of disintegration tests and laboratory flume

	M1	M2	M3	Marsh clay
Lab flume	31	9	45	1
Endell	2.2	1.8	2.1	1
Weißmann	38	15	54	1

Conclusions

The use of dredged materials in dike construction is an important contribution to environmental protection engineering since a legal waste is beneficially re-used in environmental constructions. For the characterisation of the materials the erosion stability has to be investigated. During the investigations both benefits and major limitations of disintegration tests could be observed. The following conclusions can be drawn:

1. There is a major problem with the small mesh size in the Weißmann test facility and there are limitations with respect to the varying agglomerate sizes. In case

of the Endell test the proposal of RPW to repeat the test until there is one in which all particles fall through the mesh is problematic with respect to the agglomerations that characterise a fine-grained organic dredged material. For the small mesh size proposed there may not be one such result.

2. The sample preparation with drying and rewetting of cohesive organic soils has a considerable influence on the aggregate stability and complete drying should be avoided.
3. The sample preparation methods for disintegration tests need to be modified for cohesive organic soils.
4. In both disintegration test setups no dependency between disintegration time and water content could be found. All oven-dried samples show an increase of the initial weight instead of a mass loss, although moderate crumbling could be recognized. Thus the modification of different test conditions should be considered.
5. Disintegration tests can be used to determine qualitative disintegration differences between materials using samples prepared at natural water content w_n .
6. The disintegration tests after Endell and Weißmann cannot be used to compute the quantitative aggregate stability for cohesive organic soils yet.
7. The results of the study on disintegration tests shows that there is still work needed to establish a laboratory test that allows to determine the erosion stability of fine-grained organic soils to be used in geotechnical constructions in environmentally sensitive areas such as coastal lowlands. The presented determination of benefits and limitations of the existing tests together with an analysis of necessary adaptations and boundary conditions pave the way for environmental engineers to discuss this development.

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