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Quantifying saltmarsh vegetation and its effect on wave height dissipation: Results from a UK East coast saltmarsh

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Abstract

The degree to which incident wind waves are attenuated over intertidal surfaces is critical to the development of coastal wetlands, which are, amongst other processes, affected by the delivery, erosion, and/or resuspension of sediment due to wave action. Knowledge on wave attenuation over saltmarsh surfaces is also essential for accurate assessments of their natural sea-defence value to be made and incorporated into sea defence and management schemes. The aim of this paper is to evaluate the use of a digital photographic method for the quantification of marsh vegetation density and then to investigate the relative roles played by hydrodynamic controls and vegetation density/type in causing the attenuation of incident waves over a macro-tidal saltmarsh.

Results show that a significant statistical relationship exists between the density of vegetation measured in side-on photographs and the dry biomass of the photographed vegetation determined through direct harvesting. The potential of the digital photographic method for the spatial and temporal comparison of marsh surface vegetation biomass, density, and canopy structure is highlighted and the method was applied to assess spatial and seasonal differences in vegetation density and their effect on wave attenuation at three locations on a macro-tidal saltmarsh on Dengie Peninsula, Essex, UK. In this environmental setting, vegetation density/type did not have a significant direct effect on wave attenuation but modified the process of wave transformation under different hydrodynamic conditions. At the two locations, characterised by a relatively tall canopy (15–26 cm) with biomass values of 430–500 g m⁻², dominated by *Spartina* spp. (>70% of total dry biomass), relative incident wave height (wave height/water depth) is identified as a statistically significant dominant positive control on wave attenuation up to a threshold value of 0.55, beyond which wave attenuation showed no significant further increase. At the third location, characterised by only slightly less biomass (398 g m⁻²) but a shorter (6 cm) canopy of the annual *Salicornia* spp., no significant relationship existed between wave attenuation and relative wave height. Seasonally (between September and December) significant temporal increase/decrease in vegetation density occurred in one of the *Spartina* canopies and in the *Salicornia* canopy, respectively, and led to an expected (but not statistically significant) increase/decrease in wave attenuation. The wider implications of these findings in the context of form—process interactions on saltmarshes and their effect on marsh evolution are also discussed.

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1. Introduction

Recent concern over the possibility of increased flood risk (as a result of rising relative sea level and increased storminess) on many low-lying coasts requires a better understanding of the natural sea-defence capacity of intertidal areas such as mudflats and saltmarshes. Previous studies (e.g. Wayne, 1976; Brampton, 1992; Möller et al., 2001; Möller and Spencer, 2003) have shown that saltmarsh environments in a variety of physical settings can significantly increase attenuation of incident waves compared to unvegetated sand/mudflats. The most recent estimates for macro-tidal northwest European marshes (Möller et al., 1999, 2002) suggest rates of wave height reduction of up to 50% over the first 10–20 m of vegetated saltmarsh surface. Other studies have shown that vegetation can

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significantly reduce unidirectional tidal marsh surface flows (see e.g. Leonard et al., 1995a,b; Leonard and Reed, 2002).

The theory of wave energy dissipation due to viscous boundary layer friction, permeability and the viscous nature of the sea bed in shallow water has been developed in great detail (e.g. Sleath, 1984) since the early works of Putman and Johnson (1949), Bretschneider and Reid (1954) and others. It is known that, over any given surface, wave dissipation varies in response to hydrodynamic conditions, in particular, water depth and incident wave heights. Empirical studies by Le Hir et al. (2000), for example, suggest that relative wave height (i.e. wave height/water depth ratio) provides an overriding control on shallow water wave attenuation. An $H_{\rm rms}/h$ ratio of 0.15 was found to determine the limit of observed wave heights on the Brouage mudflat in France (Le Hir et al., 2000).

In addition to these hydrodynamic controls, however, wave attenuation over saltmarsh (as opposed to unvegetated mudflat surfaces) is increased by the effect of bed 'roughness' — the result of topographic variations in marsh surface geometry as well as vegetation canopy geometry and structure (see e.g. Hartnall, 1984; Dijkema, 1987; Pethick, 1992). It has proven difficult to incorporate these sea bed roughness effects into predictive energy dissipation models. A crude attempt was made by Möller et al. (1999) to quantify friction by comparing measured wave attenuation on a macro-tidal saltmarsh with modelled attenuation under identical hydrodynamic conditions but over unvegetated surfaces and smooth bottom topography. The resulting friction factors, however, do not allow the effects of topography and vegetation to be separately quantified.

With regard to vegetation-induced roughness effects alone, the studies by Putman and Johnson (1949) and Bretschneider and Reid (1954) made significant attempts to derive a formula for calculating dissipation given known 'friction factors'. Subsequently, attempts have been made at defining vegetationinduced friction empirically, but such attempts have tended to be based on laboratory studies assessing the effect of vegetation on unidirectional flow rather than on the bi-directional flow of wave currents (e.g. Pethick et al., 1990; Fonseca and Cahalan, 1992; Shi et al., 1995; Coops et al., 1996). These studies have also suggested that the complexity of within-canopy flows results in species-specific flow structures related to canopy height, vegetation structure and plant spacing (Leonard et al., 1995a,b), thus exposing the traditional modelling analogy of regarding vegetation elements as cylinders of varying diameter, spacing and orientation within a flow field (e.g. Eckman et al., 1989) as inadequate.

With respect to wave attenuation, the potential effect of spatial and seasonal differences in marsh vegetation canopies has been addressed qualitatively by Möller and Spencer (2002), who observed a consistent decrease in attenuation at the marsh edge of the Tillingham marshes, Essex, UK, from autumn (September to December) through winter (December to February) and into spring (March to July). Variations in water depth, however, have also been shown to control wave attenuation at the saltmarsh edge, i.e. the transition from unvegetated sand/mudflat to vegetated marsh (Moeller et al., 1996; Möller and Spencer, 2002), making it difficult to isolate

the effect of vegetation-induced surface roughness changes. No study has thus, as yet, systematically investigated the relative importance of different types (and seasonal changes) of vegetation on wave attenuation in a field setting, under comparable hydrodynamic conditions. One of the reasons for this is the very time consuming and error-prone nature of traditional techniques for recording vegetation structure (e.g. the susceptibility of the point quadrat method described by Sutherland (1996) to wind disturbance — a common occurrence in dynamic coastal environments).

This paper thus aims to (a) test and develop a new, easily deployable technique for the quantification of marsh surface vegetation density, height, and structure and (b) investigate the relationship between such marsh surface vegetation characteristics and wave height dissipation (taking into account possible hydrodynamic controls, such as the incident wave height to water depth ratio).

2. Field setting

Field measurements were conducted on a macro-tidal saltmarsh near Tillingham on the Dengie Peninsula, Essex, UK (Fig. 1). The Dengie marshes lie between the estuaries of the Blackwater and Crouch on the northern margin of the Thames estuary, southern North Sea and form a generally narrow belt (700 m wide at their greatest extent) between lowlying, seawall-protected agricultural land and intertidal mudflats which extend for up to 4 km offshore. The saltmarshes have been morphologically dynamic over the past 100-150 years with phases of advance and retreat reported in the literature (e.g. Greensmith and Tucker, 1965; Harmsworth and Long, 1986; Pye, 2000). At Tillingham, this morphological history has led to mature marshes that grade into unvegetated mudflats through a distinct shore-normal mud-mound topography. Marsh surfaces on the mature marsh are near-horizontal at elevations of 2.4-2.7 m O.D. (Ordnance Datum Newlyn, which approximates to mean sea level). This coast is macro-tidal, with an offshore mean spring tidal range of 4.8 m (Reed, 1988). As other UK East coast marshes, the Dengie marshes are prone to the effect of North Sea storm surges and meteorological influences on tidal water surface elevations can be significant (Steers et al., 1979). Highest wave heights along this coast occur between February and April and between August and October, with minimum values between May and July (Herman, 2000). Further inshore, at the marsh edge at Tillingham, spring tide water depths of between 0.12 and 0.84 m and wave heights of around 20 cm are typically found (as observed over a series of spring tides during a 10-month period (October to August) by Möller and Spencer, 2002).

The vegetation is typical of the floristically diverse east coast saltmarshes of the British Isles (Adam, 1988). Midmarsh communities are typified by *Limonium vulgare*, *Puccinellia maritima* and *Atriplex portulacoides*. Near the seaward edge of the vegetated zone, a pioneer/seasonal marsh exists, characterised by *Salicornia* spp., *Spartina anglica*, *Aster tripolium* and *Suaeda maritima*. The vegetation exists in distinct patches of several square meters in size, dominated by

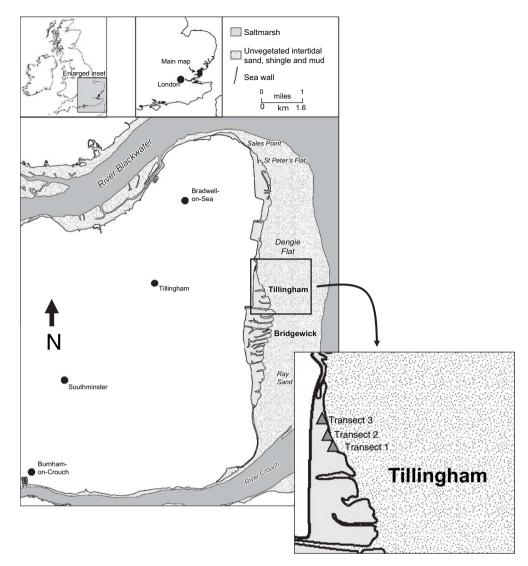


Fig. 1. Location of Tillingham on the Dengie Peninsula, Essex, UK (wave measurement transect locations 1, 2, and 3 are shown as triangles).

individual species, such that it is possible to identify approximately uniform vegetation types in close proximity to each other (i.e. experiencing similar hydrodynamic conditions).

3. Field instrumentation and measurement

Vegetation and wave measurements were carried out at three 10-m long cross-shore transects during two field visits, in September and December 2004. Measurements of hydrodynamic conditions (water depths and waves) were made over three and nine consecutive spring tides, respectively. The timing of the field visits was chosen to coincide with times of the year of contrasting seasonal vegetation cover/structure. Transects differed with respect to vegetation type with Transect 1 containing a dense *Spartina* cover, Transect 2 a less dense combination of *Spartina*, *Salicornia*, and *Suaeda*, and Transect 3 a uniform *Salicornia* cover. All transects were located the same distance from the marsh edge and in close vicinity to each other (see Fig. 1) to minimise potential spatial differences

in incident wave conditions. All transects were characterised by topographically smooth marsh sections such that the effect of topographic variation on wave attenuation across the transects could be assumed to be negligible.

3.1. Vegetation measurements

Field measurements of vegetation were collected at 18 locations (six locations near each of the three transects: three between the wave recording instruments and three in the immediate vicinity (but outside) of the transect) during the September and December field visits. At all 18 locations, digital side-on photographs of a 0.6 m wide and 0.2 m deep strip of vegetation were taken against a red background plate (to provide maximum colour contrast against the brown to green tones of the vegetation). The field methodology was initially developed for saltmarsh environments by Möller (2000) and, in parallel, for grasslands by Zehm et al. (2003). Further development has led to the methodology adopted for the purpose of this study, as illustrated in Fig. 2.

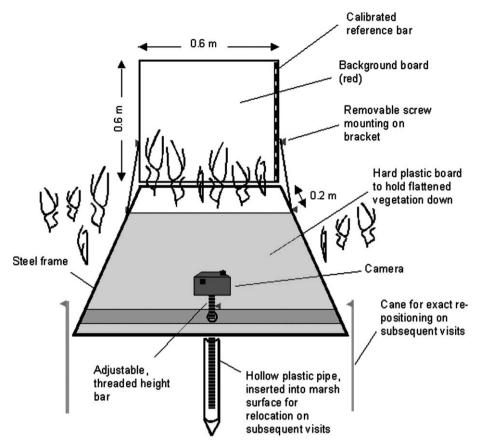


Fig. 2. Portable digital photograph frame, first developed by Möller (2000) and Zehm et al. (2003), as used in this study to capture side-on photographs of marsh vegetation.

The method used in this study differed from that of Zehm et al. (2003) in a number of important respects in order to adapt it to use in saltmarsh environments:

- (1) a rigid plastic (slightly textured) board (matt red colour finish, to reduce reflection and maximise contrast against vegetation) was used as a background to the photograph instead of an opaque cloth (optionally white or black), which was thought to be too sensitive to wind disturbance on exposed saltmarsh environments and would not give sufficient colour contrast in dim light conditions;
- (2) a smaller area (0.6 m wide and 0.2 m deep) was photographed, as the area used by Zehm et al. (2003) (1.0 m wide and 0.3 m deep) was thought to be too large (and too deep) for dense saltmarsh vegetation (it would not allow subtle differences in vegetation density to be distinguished); and
- (3) the vegetation in front of the photographed plot was pressed to the ground with a plastic sheet held within the photographic frame, rather than a tarpaulin (which was found to be inadequate for the more rigid and dense saltmarsh vegetation).

Digital images were then classified into binary black (vegetation) and white (background) images using the Erdas Imagine 8.5 image processing software and output as binary matrix files for further quantitative analysis using program routines

written in Matlab software code. Image classification included the following key steps:

- (1) cropping of images to standard size of background board and checking resolution of image (1 pixel approximated 0.25 mm²) with reference to object of known size (a scale bar of white and black squares of 1 × 1 cm clipped to the background board);
- (2) performing a supervised classification of each image into a binary image of black vegetation and white background (at least 15 training areas of the 'vegetation' and 'background' category were selected for this purpose from one or more images collected during each field visit and the resulting 'signature' file used to classify all images collected during that field visit);
- (3) checking the performance of step (2) for each classified image and, if necessary (i.e. if misclassification of the 'vegetation' or 'background' class occurred), modifying the signature file generated under step (2) by selecting additional training areas and reapplying it to all images collected during the field visit;
- (4) saving each binary image file in matrix (ASCII text) format ('0' values for 'background', '1' values for 'vegetation' class).

Following this classification procedure, the binary image analysis in Matlab used a series of programming routines,

providing detail on overall image vegetation density (% black versus white pixel cover within the 0.6×0.6 m photographed area) and structure (e.g. canopy height variation across the image). Fig. 3 illustrates the digital image analysis process. Note that the method for determining vegetation canopy height differs from the more traditional methods of determining the height of individual plants in that it provides an average of the height of any vegetation component in the image, measured across the width of the image.

To calibrate and validate the digital photographic record described above, vegetation contained in the photographs (i.e. from a surface area of $0.6 \times 0.2 = 0.12 \,\mathrm{m}^2$) at the locations immediately outside each of the recording transects was also harvested for determination of wet and dry biomass and species composition/structure during the first (i.e. September 2004) field visit. In addition, a separate experiment involved the placement of the photo frame in an area of dense, tall Spartina, which was photographed once with the vegetation cover intact and three subsequent times with the vegetation successively thinned through harvesting individual stems at the base of the frame. Each time the vegetation cover was thinned by partial harvesting, cuttings were placed in a separate bag for return to the laboratory, where samples were dried and weighed to determine the dry biomass that had been removed in the successive clipping stages.

3.2. Wave measurements

Bottom mounted Druck 'PTX1830' pressure transmitters were deployed at the seaward and landward ends of the three 10-m long cross-shore transects (Fig. 1). Marsh surface topography between each pair of sensors was smooth and the surface elevations of the marsh surface at either end of each transect are comparable (1.06/1.08 m, 1.07/ 1.06 m, and 1.04/1.01 m O.D. (Ordnance Datum: approximates to Mean Sea Level) at the landward/seaward end of Transect 1, 2, and 3, respectively). Low (1 Hz) and high (4 Hz) frequency pressure measurements were triggered upon submergence of the sensors and stored as (a) 5 min averaged pressure (for water depth measurements) and (b) continuous 4 Hz readings (for computation of wave spectra and summary statistics). Measurements were controlled and stored, together with a record of datalogger battery voltage and date/time references, via a Campbell Scientific CR10 datalogger. The measurement and data processing method followed the procedure outlined in Möller and Spencer (2002).

Upon downloading of the pressure data, tidal stage records were derived from the 5 min averaged pressure readings. The high frequency (4 Hz) pressure time series were then split into series of 2¹¹ (2048) values (i.e. 8.5 min duration) when water

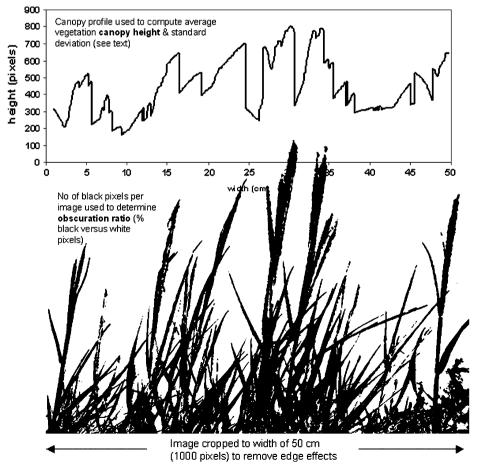


Fig. 3. Schematic diagram illustrating the derivation of obscuration ratio (black pixel coverage (%) of image) and canopy height from classified binary black and white images.

depths reached 20, 30, 40, 50, 60, and 70 cm on the rising tide. These high frequency records were processed according to the following sequence:

- (1) de-trending of pressure time series (using a polynomial fit) to remove any low-frequency tidal component present;
- (2) computation of pressure spectra using a Fast Fourier Transform (FFT) algorithm followed by adjustment of the spectra to allow for frequency-dependent attenuation of the wave-induced pressure signal with water depth (Lee and Wang, 1984); and
- (3) computation of summary parameters such as water depth (h) and zero-upcrossing period (T_z) from the corrected free surface record, and computation of total wave energy (in J m⁻²) (used for the computation of root-mean-square wave height (H_{rms}) and significant wave height (H_s)) from its frequency spectrum $(S(f_n)\Delta f)$ (see Tucker, 1991), where:

$$S(f_n)\Delta f = 0.5a_n^2$$

$$E_{\text{tot}} = \sum_{n=1}^{N} S(f_n) \Delta f g \rho = \sum_{n=1}^{N} 0.5 a_n^2 g \rho$$

$$H_{\rm s} = 4\sqrt{E_{\rm tot}\frac{1}{g\rho}}$$

$$H_{
m rms} = 2\sqrt{2}\sqrt{E_{
m tot}rac{1}{g
ho}}$$

and a_n = amplitudes of the component waves of the spectrum, N = total number of measurements (time series length), g = gravitational acceleration, and ρ = water density (1.02 g cm⁻³).

A detailed calibration of the above method for wave recording against a video record of water level fluctuations using a calibrated staff is described in Moeller et al. (1996).

This processing sequence thus resulted in wave parameters for the seaward and landward sensors of each of the transects in water depths at 10 cm intervals between 20 and 70 cm depths.

4. Results

4.1. Vegetation measurement

The results from the calibration of the digital photograph technique against biomass measurements determined by harvesting the photographed vegetation in the vicinity of the wave recording transects are shown in Fig. 4. The relationship between vegetation pixel density (black pixel density in the classified binary image, i.e. obscuration ratio of the background plate) and dry biomass was positive when all data were included in a regression analysis (p = 0.05, $r^2 > 0.6$). The data in Fig. 4a, however, show that the relationship

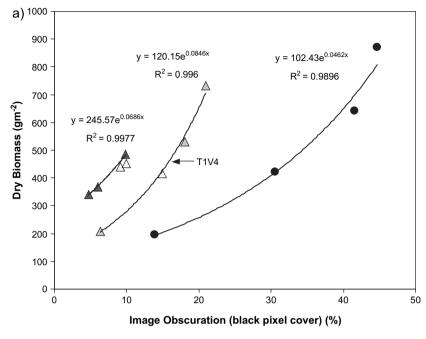
between the vegetation pixel density (black pixel density in the classified binary image) and dry biomass varied between three distinct groups of measurement, namely those made on Transect 2, those on Transect 3, and the separate *Spartina* thinning experiment. With the exception of the measurements obtained from location T1V4, the data from Transect 1 seem to fit into the relationship established for Transect 3. The best fit to the data was achieved with an exponential regression fit. Table 1 summarises the results from the regression analysis performed on these data.

The different canopy heights and species composition of the vegetation are shown in Fig. 4b, c. Transect 3 is distinguished by a significantly lower canopy height (across all three locations on Transect 3, the average height was 5.9 ± 2.6 cm) and different species composition (>60% of dry biomass was Salicornia) than Transect 2 (average height across all three locations: 22.5 ± 8.0 cm (although canopy heights on T2V4 were more comparable with Transect 3), >65% dry biomass Spartina). Of the three locations measured on Transect 1, the canopy height on T1V4 was greatest (20.1 \pm 7.7 cm) and, given its vegetation composition (74% Spartina by dry weight), this might have put this site more in line with Transect 2 than Transect 3. The vegetation at locations T1V5 and T1V6 on Transect 1 (although consisting mainly of Spartina (95 and 87% of dry biomass, respectively)) was much lower in height $(11.6 \pm 5.6 \text{ and } 13.6 \pm 7.9 \text{ cm}, \text{ respectively}), \text{ and}$ this appears to have put these measurements more in line with the measurements obtained from the low Salicornia canopy of Transect 3. The Spartina thinning experiment was characterised by the tallest canopy $(45.1 \pm 4.1 \text{ cm})$ and plots furthest on the right in Fig. 4a, providing evidence for an overriding importance of canopy height on the relationship between %image obscuration ratio and dry biomass.

In addition to the overall positive correlation between %image obscuration and dry biomass, the scatterplot presented in Fig. 4a suggests that this relationship may not be linear. It suggests, rather, that at already high biomass values, a further increase in biomass does not lead to a corresponding increase in %image obscuration.

Table 2 shows that, with the exception of Transect 2, the vegetation within the wave recording transects was generally representative of the overall vegetation characteristics determined by measurements at all six locations within each vegetation type. This table also illustrates that the highest variability in canopy height (a potential measure of vegetation canopy 'roughness') on the wave recording transects themselves was recorded on Transect 2 (8.0 cm), followed by Transect 1 (7.1 cm) and Transect 3 (2.6 cm).

Seasonal changes in image vegetation density (%image obscuration ratio) and canopy height recorded on the three vegetation measurement locations on the wave recording transects themselves are illustrated in Fig. 5. While Transect 1 experienced no significant change in image vegetation density between September and December, Transects 2 and 3 experienced a significant increase (from 6.1 ± 1.1 to 13.7 ± 5.8) and decrease (from 9.5 ± 0.2 to 6.4 ± 1.6) in image vegetation density, respectively (Fig. 5a). On all three



△ Sep 04: Trans 1 (h = 15 cm; 86% Spartina, 11% Salicornia)

△ Sep 04: Trans 2 (h = 23 cm; 73% Spartina, 25% Salicornia)

▲ Sep 04: Trans 3 (h = 6 cm; 98% Salicornia)

● Dec 04: h = 45 cm; Spartina Clip Experiment

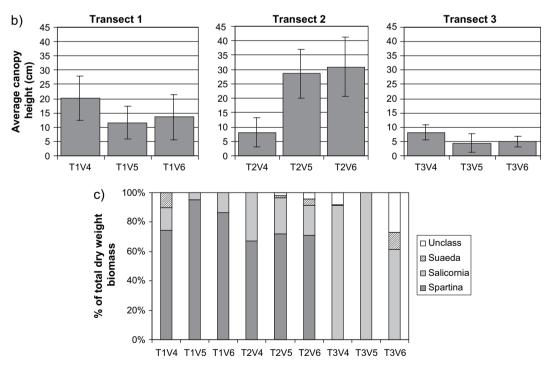


Fig. 4. (a) Relationship between aboveground biomass (dry weight) of photographed vegetation and image obscuration ratio for three vegetation monitoring locations on each of the three cross-shore transects and for the *Spartina* thinning experiment; (b) mean (± 1 standard deviation indicated by error bars) of canopy height as determined from classified binary side-on image for each measurement location in the immediate vicinity of Transect 1 (T1V4 to T1V6), Transect 2 (T2V4 to T2V6), and Transect 3 (T3V4 to T3V6); (c) species composition (% of total dry weight biomass) of vegetation harvested (same locations as shown in (b)).

Table 1 Results of exponential regression analysis — image obscuration (% of $0.6 \times 0.6 \times 0.2 \,\mathrm{m}$ area) versus dry biomass of clipped vegetation (g m⁻²) (nb: $p \leq 0.05$ throughout): Biomass = $a \, \mathrm{e}^{b \, \mathrm{obscuration}}$

	R^2	a-Variable	b-Exponent
Transect 2 (including T1V4)	0.996	120.15	0.0846
Transect 3 (including T1V5 and T1V6)	0.998	245.57	0.0686
Spartina thinning experiment	0.990	102.43	0.0462

transects, vegetation canopy height decreased from September to December, but average heights for December remained within ± 1 standard deviation relative to the September averages (Fig. 5b).

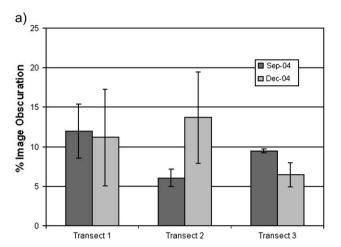
4.2. Wave attenuation

Wave measurements at the seaward and landward stations on each of the three wave recording transects were obtained during three and nine tidal cycles in September and December 2004, respectively. Water depths during the September tides were considerably larger (maximum depths of 70 cm recorded on two out of the three tides) than in December (maximum depths of 50 cm recorded on only three out of the nine tides). Table 3 summarises the wave conditions encountered at the seaward transect stations in September and December 2004. This shows that maximum significant wave heights recorded at the seaward transect stations were greater in December (max 32 cm) than September (max 19 cm). Relative wave heights (i.e. wave height/water depth ratios) were thus larger in December (0.12–0.86) than in September (0.08–0.54).

Table 3 also illustrates that incident wave conditions were comparable between Transects 1 and 2 but significant wave heights (H_s) at Transect 3 were generally 2–3 cm lower.

Fig. 6a shows the average significant wave height (H_s) attenuation (± 1 standard deviation) observed over the three transects in September and December, in any water depth. Average H_s attenuation increased over Transects 1 and 2 between these two field visits, but decreased over Transect 3 — although all these changes remained within the limits of the standard deviations recorded in September.

Fig. 6b resolves these data further to show average $H_{\rm s}$ attenuation (± 1 standard deviation) for the three transects in different water depths during the September and December field campaigns. Wave attenuation ($H_{\rm s}$ reduction) over the 10-m transect distance varied from 0.08% (in 40 cm water depth



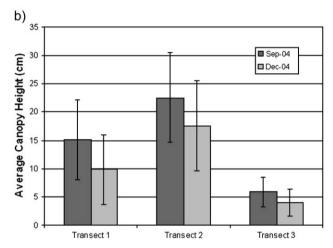


Fig. 5. Seasonal changes (September versus December) in (a) image vegetation density (%image obscuration ratio) and (b) canopy height at the three locations on each of the three cross-shore transects.

over Transect 3 on tide 8 in December (14th)) to 33% (in 20 cm water depth over Transect 1 on the same tide). At any water depth, in September and December, highest average wave attenuation was observed over Transect 1, although attenuation was highly variable (Fig. 6b).

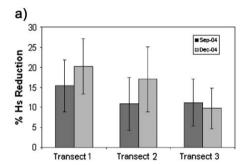
Wave attenuation across any surface is known to vary with relative wave height (as discussed above) (see e.g. Le Hir et al., 2000). Fig. 7 shows that a statistically significant relationship between H_s/h (where h = water depth) and wave height attenuation (% H_s reduction) existed on Transects 1 and 2 ($r^2 > 0.6$, p < 0.05), but not on Transect 3 ($r^2 = 0.2$),

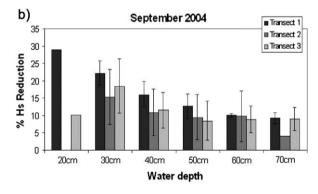
Table 2
Average vegetation canopy profile height and standard deviation within (three location) and overall (all six locations) at the three wave recording transects, September 2004

	Vegetation type (% mix by dry weight)	Vegetation profile height (cm)				Obscuration ratio (% of 0.6×0.6 m area covered by vegetation in image)			
		Average		Std. deviation		Average		Std. deviation	
		Within	Overall	Within	Overall	Within	Overall	Within	Overall
Transect 1	86% Spartina, 11% Salicornia	15.1	15.3	7.1	6.9	11.9	11.6	3.4	3.0
Transect 2	73% Spartina, 25% Salicornia	22.5	15.3	8.0	6.4	6.1	10.6	1.1	7.0
Transect 3	98% Salicornia	5.9	7.0	2.6	2.8	9.5	8.2	0.2	2.2
Clip exper.	100% Spartina		45.1		4.1		44.7		

Table 3
Summary of wave conditions encountered at the seaward stations of the three transects in September and December 2004

	$T_{\rm z}$ (s)		$E_{\rm tot} ({\rm J m}^{-2})$		$H_{\rm s}$ (cm)	
	September	December	September	December	September	December
Transect 1						
Mean	2.6	2.0	8.6	12.0	11.1	12.3
Std. Dev.	0.7	0.4	5.5	13.3	3.9	6.4
Min.	1.5	1.3	2.4	2.2	6.2	5.9
Max.	3.3	2.8	17.3	54.9	16.6	29.6
Transect 2						
Mean	2.6	2.1	9.8	13.7	11.7	12.9
Std. Dev.	0.6	0.4	7.0	16.3	4.6	7.4
Min.	1.6	1.3	2.3	2.3	6.0	6.0
Max.	3.3	2.8	22.0	63.3	18.8	31.8
Transect 3						
Mean	2.6	2.1	4.6	8.5	8.0	9.7
Std. Dev.	0.7	0.5	3.6	12.6	3.3	6.7
Min.	1.2	1.1	0.9	0.9	3.7	3.8
Max.	3.2	2.8	12.5	51.7	14.1	28.7





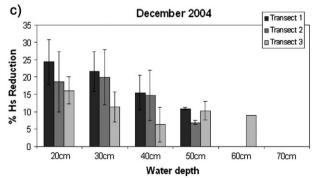


Fig. 6. Average significant wave height (H_s) attenuation (± 1 standard deviation) across the three transects in September and December for (a) all water depths, and (b) divided into the different water depth categories (note: standard deviations are not shown if less than three observations are present).

in September, when H_s/h measurements were consistently <0.55. In December, however, when H_s/h measurements reached values as high as 0.86, no significant relationship was found to exist between H_s/h and H_s attenuation $(r^2 < 0.3)$. When the data set was restricted to records for which $H_s/h < 0.55$, however, a positive relationship between H_s/h and H_s attenuation was present in December $(r^2 > 0.5, p < 0.05)$, although, again, only for Transects 1 and 2 (as in September). See Table 4 for details of the regression equations that apply to the regression lines shown in Fig. 7.

5. Discussion

5.1. Quantifying vegetation density, height and structure by digital imaging

The results presented above suggest that the digital photograph method developed as a prototype by Möller (2000) for saltmarsh environments and in more detail by Zehm et al. (2003) for grasslands, is applicable to a range of relatively simple saltmarsh vegetation canopies. The modifications made to the method as outlined above proved successful and allowed its application to coastal environments, where dim light conditions and/or adverse weather (e.g. wind) can be problematic when using the method as configured for grasslands by Zehm et al. (2003).

It is also evident, from the varying regression relationships presented in Fig. 4a, that it is possible to estimate aboveground biomass from image obscuration ratio, although this statistical relationship varies with species type/composition and is affected by canopy height. Similar results were obtained by Zehm et al. (2003) when comparing overall image obscuration (pixel count) with harvested aboveground biomass (dry weight) over a gradient from pioneer grassland vegetation to tall, mature stands, although the photographed plot area was deeper (30 cm) and wider (100 cm) than in the saltmarsh vegetation photographs of this study (20 cm \times 60 cm). The

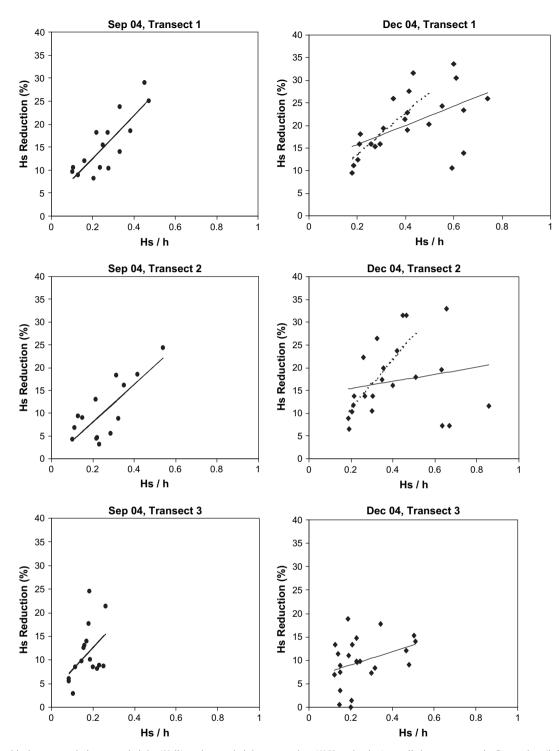


Fig. 7. Relationship between relative wave height (H_s/h) and wave height attenuation (% H_s reduction) on all three transects in September (left) and December (right) (note: solid regression lines apply to all data, stippled lines apply to conditions where $H_s/h < 0.55$; on Transect 3, all H_s/h values were <0.55; see Table 4 for regression equations).

relationship between image obscuration (pixel count) and dry weight of aboveground biomass established in this study is in agreement with the non-linear trend suggested by Zehm et al. (2003) for grasslands, i.e. for dense stands of vegetation, an increase in aboveground biomass results in only minor increases in image obscuration. The threshold biomass value beyond which no further image obscuration would occur does not appear to have been reached in any of the images analysed as

part of this study (Fig. 4a), suggesting that the plot area of 0.2×0.6 m is adequate for these types of saltmarsh vegetation communities. For a larger plot area vegetation density would become too large for differences in aboveground biomass to manifest themselves as differences in image obscuration. Further studies using a larger sample size to further increase confidence in the results as summarised in Fig. 4a are, however, needed.

Table 4
Details of regression analysis presented in Fig. 7

	X-variable		Y-intercept		R^2	
	September	December	September	December	September	December
Transect 1	46.47	21.27	3.23	11.49	0.69	0.28
Transect 1 ^a		45.96		4.44		0.61
Transect 2	41.32	7.80	-0.24	13.87	0.61	0.32
Transect 2 ^a		53.93		0.23		0.53
Transect 3	47.43	14.31	3.10	6.15	0.20	0.13

^a Using data for H_s/h values < 0.55 only (on Transect 3, all H_s/h values were < 0.55).

Although further studies are needed to establish the accuracy of the digital imaging technique in different saltmarsh settings (and especially mixed marsh vegetation communities), this method provides a series of key advantages over more traditional vegetation survey methods (see e.g. Sutherland, 1996) as it:

- (1) minimises time in the field (can be deployed quickly at pre-defined sites);
- (2) is a non-destructive method (although areas around the photographed plots are temporarily disturbed);
- (3) allows detailed, objective, and quantitative measurement of a series of vegetation structure parameters (e.g. canopy 'roughness' and density) without measurement errors frequently encountered in direct field measurements (e.g. wind affecting point quadrat measurements (Sutherland, 1996); and
- (4) is insensitive to operator error, optical distortions and technical problems (see detailed validation conducted by Zehm et al., 2003).

5.2. Seasonal vegetation density and canopy structure variations

The seasonal change in vegetation density (from September to December) recorded with the digital imaging technique in this study was highly variable between the three transects. When comparing vegetation densities at Tillingham between November, March, and June surveys, previous studies (Möller and Spencer, 2002) observed the highest vegetation densities in November. The observations from the three marsh edge transects presented in this study, however, suggest both increases and decreases in vegetation density from September to December. Such spatial variability is to be expected within marsh edge communities, given the differences in the relative dominance of annual versus perennial species. While annual Salicornia was dominant on Transect 3, where vegetation density decreased significantly between September and December, Spartina dominated Transect 2, where continued growth into the autumn months led to an increase in vegetation density between September and December. While Transect 1 was also dominated by Spartina, it appears that it reached its peak density earlier than Transect 2 and thus no significant change in vegetation density was observed between September and December.

With respect to canopy structure, the results from this study suggest that it is the mean canopy height, rather than its standard deviation (i.e. canopy 'roughness'), that changes seasonally at the three transects studied: while the mean canopy height decreased from September to December at all three transects, standard deviations (canopy 'roughness') remained constant. In terms of the effect on wave attenuation, it is thus more likely to be the height and density variations of the vegetation layer that explain observed decreases in attenuation in the winter months (as observed by Möller and Spencer, 2002) than changes in the 'roughness' or variability of the top of the canopy.

5.3. Relative importance of hydrodynamic conditions and vegetation on wave attenuation

Over the short-term individual tidal inundations, a link exists between the surface roughness characteristics of the marsh and the hydrodynamic process of wave transformation across the surface through the effects of friction, percolation, and viscous damping (see e.g. Putman and Johnson, 1949; Bretschneider and Reid, 1954; Sleath, 1984; Le Hir et al., 2000). In addition, however, the wave attenuation process itself is also a function of water depth and incident wave conditions (as higher and longer waves are affected more strongly by the mudflat/saltmarsh surface). In particular, wave heights over shallow mudflat environments have been shown to be limited by critical relative wave height (wave height/water depth) ratios (Le Hir et al., 2000).

Previous data obtained at the Tillingham site, however, clearly contained a seasonal signal, not only in terms of both overall energy input (incident wave energy) but also in terms of attenuation patterns (Möller and Spencer, 2002). Seasonally changing surface friction (due to changing vegetation density, structure, or composition) especially in marsh edge environments affected by seasonal vegetation growth (e.g. *Salicornia europaea* or *Suaeda maritima*) thus also has to be considered as an explanation for temporally varying wave attenuation.

The results of this study agree with previous measurements at Tillingham (Möller and Spencer, 2002) both in terms of significant wave height attenuation observed (around 30% over 10 m) and in terms of seasonal differences (for similar water depths, attenuation was higher in September than December). As measurements were made over three different vegetation transects, however, the results of this study provide some additional insight into the relative importance of wave/water

depth conditions and vegetation density/structure on wave attenuation.

Although it was hoped that incident wave conditions would be comparable between all three cross-shore transects, analysis of the results showed that Transect 3 consistently experienced significant wave heights 2–3 cm lower than those incident upon Transects 2 and 1. This thus allows no direct comparison between Transect 3 and the other two transects.

With respect to Transects 1 and 2, however, the results show that wave height attenuation, although highly variable, was consistently greater over Transect 1, in all water depth/ wave height conditions encountered. This difference in attenuation, however, only exceeded 1 standard deviation around the mean in the deeper water conditions (70 cm water depth in September and 50 cm water depth in December (Fig. 6)). Transect 1 contained the greatest proportion of Spartina (Fig. 4c) and was characterised by the highest vegetation density in September 2004 (Fig. 5), which may have led to greater friction and thus greater wave attenuation over this transect at that time of year. The average canopy height of Transect 1, however, was less than that of Transect 2 in September and December and its vegetation density was less than that of Transect 2 in December (although not significantly so), so that these vegetation parameters alone do not sufficiently explain the consistently higher wave attenuation. This suggests that vegetation density and height affect wave transformation differently. While the former is a measure of friction as experienced by the horizontal flow components of the wave motion, the latter affects the depth of the boundary layer affected by such friction and thus the vertical component of the wave motion. Furthermore, and possibly more importantly, the physical properties of the vegetation, such as its rigidity or flexibility, may also play a significant role in affecting wave transformations. Further studies investigating the relative differences between individual species with respect to such physical properties are needed to shed further light on this process. It must also be recognised that the method for determining canopy height used in this study differs from conventional measurements of vegetation height in that it constitutes an average of the highest parts of any vegetation component present in the digital image (rather than an average of the height of individual plants).

Transect 3 was characterised by a very different vegetation type (dominated by the annual, Salicornia (Fig. 4c)). It appears that this explains, at least partly, why this transect experienced less wave height attenuation than the other two transects and experienced seasonal differences in attenuation exceeding ± 1 standard deviation around the mean in intermediate (30 and 40 cm) water depths.

Variability in attenuation was high, however, and differences in vegetation characteristics between the three transects do not exert a statistically significant influence on wave attenuation. Other controls on wave attenuation must exist and the results of this study suggest that hydrodynamic controls, such as the wave height/water depth ratio, may override any differences in vegetation density/structure in this marsh edge environment. The significant positive relationship between the

 H_s/h ratios and wave height reduction (for conditions where $H_s/h < 0.55$) on Transects 1 and 2 in September and December provides evidence for this. Once H_s/h ratios fall below a critical threshold, further decreases in the H_s/h ratios are not expected to lead to further reduction in wave attenuation (i.e. water depths become too deep or waves too small for the surface waves to interact with the vegetation canopy on the bed). This critical point, however, had not been reached on Transects 1 and 2 (water depths remained relatively low (<70 cm)). At H_s/h ratios above 0.55, however, the relationship between H_s/h and wave attenuation also appears to break down. Although only few measurements were made in conditions of H_s/h ratios > 0.55, this may indicate that above H_s/h ratios of 0.55, the attenuation capacity of the vegetation canopy is reached and further attenuation cannot take place. It is also possible, however, that in conditions of high H_s/h ratios, attenuation is affected by dissipation due to wave breaking. This is an important observation in the context of the wider morphological evolution of the saltmarsh, as it suggests that this H_s/h ratio represents a threshold beyond which the vegetation acts as a significant energy dissipater, thus leading to energy dissipation by wave breaking, bed erosion and/or resuspension of surface sediments.

Fig. 7 also shows that, for values of $H_s/h > 0.55$, attenuation may vary greatly, e.g. when an H_s/h value of around 0.6 occurred, wave height attenuations of 11 and 34% (Transect 1) and 7 and 33% (Transect 2) were observed. This thus suggests that the H_s/h ratio controls the maximum attenuation that can be achieved, rather than allowing actual attenuation to be predicted exactly. Other factors, such as meteorological conditions (e.g. local wind direction and speed) will influence wave propagation and breaking processes and thus affect whether the potential maximum attenuation for a given H_s/h ratio is reached. Further research into the role of these factors is now needed.

On Transect 3, however, no relationship between H_s/h ratios and attenuation was present. Given that the vegetation on this transect was substantially shorter (6 cm compared to 15 and 23 cm on Transects 1 and 2, respectively), wave heights were 2-3 cm smaller, and water depths comparable to Transects 1 and 2, this suggests that (a) the canopy height on this transect was too low to allow the vegetation to occupy a large enough proportion of the water column, and/or (b) waves were too small to be affected by the vegetated layer. Furthermore, the different vegetation type (Salicornia instead of Spartina) provides greater flexibility for movement with the flow (i.e. reduced flow resistance). This, together with the significant decrease in vegetation density, may explain why, even in high H_s/h conditions of over 0.4 in December, attenuation remained low (<20%) on this transect, while it exceeded 30% on some occasions on the other two transects.

5.4. Implications for form—process interactions and marsh evolution

The results presented above provide evidence for a relationship between hydrodynamic controls (i.e. relative wave

heights) and different types (*Spartina* versus *Salicornia*), heights, and densities of vegetation canopies during individual tidal inundations. In the context of recent concerns regarding the future of saltmarsh systems under scenarios of relative sea-level rise and/or increased storminess (see e.g. French and Reed, 2001) and marsh restoration issues (Williams and Orr, 2002), it is instructive to consider the implications of these results for the longer-term form—process interactions at the marsh edge and thus marsh evolution.

While, over short, event-based time-scales, the topographic and vegetation characteristics discussed above can be considered invariant, this does not apply over longer (annual to decadal) time-scales, when the hydrodynamic processes themselves exert a feedback effect on marsh vegetation cover and morphology (see also Allen and Duffy, 1998a,b; Allen, 2000; Houwing, 2000). The H_s/h threshold of 0.55 identified above as the point when the *Spartina* canopy appears to reach its maximum attenuation capacity, for example, suggests that, above this value (i.e. in conditions of higher incident wave energy), more energy may be available for sediment resuspension/erosion. Over the longer term (from several spring-neap tidal cycles to several years), it is thus the frequency with which such thresholds are exceeded that will determine the morphological evolution of the marsh.

Evidence of previous studies does suggest that the evolution of the marshes of Dengie Peninsula has fluctuated between accretion and erosion phases. While the marshes are thought to have formed rapidly between 1870 and 1950, with subsequent progradation rates in the central sections of the Peninsula of 9.7 m a⁻¹ (Pye and French, 1993), subsequent marsh edge retreat of up to 270 m characterised the northern half of the Peninsula (i.e. the area around Tillingham) between 1953 and 1960 (Greensmith and Tucker, 1965; Harmsworth and Long, 1986). The placement of sunken barges to form wavebreakers to the immediate North of the Tillingham marshes in 1984 has had little long-term effect, however, and the characteristic mud-mound topography currently found at the marsh edge is thought to be typical of an eroding marsh (Greensmith and Tucker, 1975). In the central sections of the Peninsula, erosion has been extreme, with ca. 1000 m retreat between 1955 and 1985 (Pye and French, 1993), leading to the development of a ca. 1.5-2.0 m high marsh edge cliff. Once the marsh edge form has altered, the attenuation process changes (e.g. as reflection from the cliff face becomes an important additional process (see also Möller and Spencer, 2002 for an investigation of wave attenuation across the cliffed marsh edge)). Given the evidence for wave attenuation thresholds presented above, it is possible that changes in the frequency of occurrence of particular wave height/water depth conditions (e.g. events characterised by H_s/h ratios >0.55 at the marsh edge) over decadal time periods are indeed responsible for initiating such evolutionary trends. Further research is now necessary, however, to investigate this hypothesis in detail.

6. Conclusion

The method of side-on photographs of stands of vegetation as developed in prototype by Möller (2000) and, independently, by Zehm et al. (2003) for grasslands, has successfully been applied to marsh edge vegetation stands dominated by Spartina and Salicornia. A positive relationship was found to exist between %image obscuration and dry biomass of the photographed stands, although the nature of this relationship varied with vegetation type and canopy height. The technique provides a time-efficient method for estimating vegetation aboveground biomass and structure - a factor that is important when working in often harsh physical conditions. Furthermore, the technique allows a range of structural vegetation canopy parameters to be objectively quantified. Although this study focuses on vegetation canopy height and its variability only, other parameters such as the vertical and horizontal variation in vegetation density and the presence and size of gaps within the canopy can be quantified. It is intended that further analysis of the image data acquired as part of this study will focus on these aspects. Future studies should also focus on investigating further the relationship between obscuration ratio and biomass (as presented in Fig. 4) as well as the relationship between conventional vegetation height measurements and the canopy height measurements obtained from the digital images (as discussed above).

Through the use of the above vegetation photograph technique, it has been possible to acquire detailed information on vegetation canopy density and height on three 10-m long cross-shore wave recording transects. Wave attenuation over these transects was a function of a combination of hydrodynamic controls (relative wave heights (wave height/water depth ratios)) and vegetation (type, density, and height), although the former may override the latter. For the transects dominated by Spartina canopies, a relationship existed between H_s/h ratios and maximum observed wave height attenuation. For H_s/h ratios exceeding 0.55, no further increase in the maximum observed wave attenuation appeared to occur, indicating that the maximum 'attenuation potential' had been reached. For the transect characterised by a lower and more flexible Salicornia canopy as well as lower incident wave heights, no such relationship was present, suggesting that water depths and wave heights were too high and low, respectively, for wave heights to be affected by the low canopy at this location.

It is possible that, over longer time-scales (>annual), the existence of hydrodynamic thresholds such as the critical H_s/h value of 0.55 determined for the *Spartina* canopies in this study may control transitions from regimes of sediment deposition to erosion (e.g. H_s/h values < 0.55 allowing maximum attenuation to occur and thus encouraging sediment deposition/retention on the marsh surface, while conditions of H_s/h values > 0.55 may result in excess energy becoming available for sediment erosion/resuspension). While this study is based on a limited data set, the results are consistent with previous studies on wave attenuation and thus justify a call for further research to investigate this hypothesis in more detail.

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