

1 An idealised study for the long term evolution 2 of crescentic bars

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9 Abstract

10 An idealised study that identifies the mechanisms in the long term evolution of
11 crescentic bar system in nature is presented. Growth to finite amplitude (i.e., equi-
12 libration, sometimes referred to as saturation) and higher harmonic interaction are
13 identified as the leading nonlinear effects. These nonlinear effects are added to a
14 linear stability model and used to predict crescentic bar development along a beach
15 in Duck, North Carolina (USA). The equilibration prolongs the development of bed
16 patterns, thus allowing the long term evolution. Subsequently, higher harmonic in-
17 teraction enables the amplitude to be transferred from longer to shorter lengthscales,
18 which leads to the dominance of shorter lengthscales in latter post-storm stages, as
19 observed in the field. The conclusion is that these nonlinear effects should be in-
20 cluded in a model simulating the development of different bed patterns. This points
21 a way forward for long-term morphodynamical modelling in general. The compari-
22 son with observation indicates the importance of higher harmonic interaction in the
23 development of nearshore crescentic bar systems in nature.

24 *Key words:* Crescentic bed-patterns, linear stability analysis, field observations,
25 long term evolution, nearshore morphology, higher harmonic interaction

26 1 Introduction

27 Nearshore sea bed patterns are a common feature around the world and may
28 provide some protection to beach and coastal areas (*Hanley et al.*, 2014).

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29 As one of the most common nearshore sea bed patterns, crescentic bars are
30 observed worldwide, see e.g. *Van Enckevort et al. (2004)*. Because of their
31 prevalence, their possible role in coastal protection, and the need to gain more
32 understanding of nearshore coastal dynamics in general, it is important to
33 study the evolution of these bed patterns.

34 Increasingly, the genesis of such quasi-periodic patterns is thought to be due
35 to morphological instability (see *Ribas et al., 2015*). An often used method
36 for describing the development of crescentic bed-forms in idealised scenarios
37 is therefore linear stability analysis, see e.g. *Deigaard et al. (1999)*; *Falqués*
38 *et al. (2000)*; *Damgaard et al. (2002)*; *Calvete et al. (2005)*; *Van Leeuwen*
39 *et al. (2006)*; *Calvete et al. (2007)*. In this method, infinitesimally small per-
40 turbations are imposed on an equilibrium (basic) state. The interaction of flow
41 and sea bed may give rise to a so called fastest growing mode, a pattern with
42 largest growth rate, which will dominate the sea bed pattern after a period
43 of evolution. Linear stability analysis has proved to be useful in revealing the
44 initialization and short term evolution of crescentic bars.

45 Following this approach, *Tiessen et al. (2010)* predicted the development of
46 crescentic bed-patterns at Duck, North Carolina (USA), for a period of two
47 months, starting from an along-shore constant bed. The forcing used was the
48 measured wave and tidal data at the same field site. Although the predicted
49 crescentic pattern lengthscales were similar to those observed, they tended to
50 exhibit a much bigger fluctuation. Such significant discrepancy is believed to
51 be a combined result of missing nonlinear effects in the linear model and the
52 effect of pre-existing bed patterns in the natural environment. This is because
53 linear stability analysis is limited when pre-existing bed-forms are present,
54 since an alongshore constant initial bathymetry is assumed at each instant.
55 Another reason is that the exponentially growing bed form will violate the
56 small amplitude assumption after some time, and nonlinear effects will dom-
57 inate the evolution thenceforth. Therefore, a nonlinear analysis is necessary
58 for reliable long-term prediction of crescentic bars (*Dodd et al., 2003*).

59 Using fully numerical models, *Tiessen et al. (2011)* and *Smit et al. (2012)*
60 included nonlinear effects and investigated the impact of pre-existing bed-
61 patterns. *Smit et al. (2012)* showed that more pronounced pre-existing bed-
62 patterns are more likely to remain, and dominate subsequent development.
63 This suggests that, under certain circumstances, pre-existing modes are not
64 affected by the present forcing conditions and that once a certain threshold of
65 development is reached, only a reset-event, such as a storm, can remove pre-
66 existing bed-forms and the corresponding dominant crescentic bed-pattern
67 lengthscales.

68 On the other hand, *Tiessen et al. (2011)* showed that pre-existing modes can
69 modify the subsequent development of different crescentic bar lengthscales.

70 Pre-existing modes (patterns) of finite amplitude will persist if those same
71 modes show significant linear growth (i.e., initial growth from an infinitesi-
72 mally disturbed beach). On the other hand, pre-existing lengthscales that
73 show only limited growth or even decay when developing from an infinitesi-
74 mally disturbed beach, become overwhelmed by faster growing modes. How-
75 ever, the lengthscale of these pre-existing, slowly growing or decaying modes,
76 and that of the newly-arising crescentic bed-form are linked. This is because
77 the more rapid initial development of higher harmonics of the pre-existing
78 lengthscale can excite a linearly unstable mode at a smaller wavelength, prior
79 to decaying to insignificance. The findings of *Tiessen et al.* (2011) and *Smit*
80 *et al.* (2012) suggested a few important nonlinear effects in the long-term
81 evolution of crescentic bars.

82 The goal of this study is therefore to identify physical mechanisms for long-
83 term growth of crescentic bar systems by comparing with field observations.

84 To this end, we develop an idealised model that incorporates the processes
85 suggested by *Tiessen et al.* (2011) and *Smit et al.* (2012) into the linear sta-
86 bility analysis. The occurrence of pre-existing modes is also accounted for in
87 the model. This approach allows us to consider only those effects identified
88 earlier, and, moreover, is time efficient and so can be applied over substan-
89 tial durations. The model is used to predict the lengthscale of the crescentic
90 bed-forms for a period of two months in 1998 at Duck (NC, USA). The model
91 results are compared with field observation (*Van Enckevort et al.*, 2004) over
92 the same period.

93 The paper is organized as follows. In section 2 the model formulation is given,
94 as well as how linear stability theory is used in the amplitude evolution model.
95 In section 3 the amplitude evolution model is presented, and an example test
96 case used to illustrate its properties. Model results and a discussion are pre-
97 sented in section 4 and 5, respectively. Finally, a conclusion is given in sec-
98 tion 6.

99 **2 Model formulation: governing equations and linear stability anal-** 100 **ysis**

101 The model geometry describes an unbounded, straight open coast in the along-
102 shore direction. Quasi-steady flow conditions are assumed and the spatial co-
103 ordinate system, (x, y) , is aligned with cross- and long-shore directions. The
104 vertical direction is denoted by z , where $z = 0$ refers to mean sea level with
105 positive z points upwards.

106 The model-framework is composed of the phase-averaged shallow water equa-

107 tions, in combination with a description of the bathymetric evolution, the
 108 wave phase and the wave energy density (see *Calvete et al.* (2005) for a more
 109 extensive description of this model).

110 The equations of the model are:

111

$$\frac{\partial D}{\partial t} + \frac{\partial D u_j}{\partial x_j} = 0, \quad (1)$$

$$\frac{\partial u_i}{\partial t} + u_j \frac{\partial u_i}{\partial x_j} = -g \frac{\partial z_s}{\partial x_i} - \frac{1}{\rho D} \frac{\partial}{\partial x_j} \left(S'_{ij} - S''_{ij} \right) - \frac{\tau_{bi}}{\rho D}, \quad (2)$$

$$\frac{\partial E}{\partial t} + \frac{\partial}{\partial x_j} ((u_j + c_{g_j}) E) + S'_{ij} \frac{\partial u_j}{\partial x_i} = -\mathcal{D}, \quad (3)$$

$$\frac{\partial \Phi}{\partial t} + \sigma + u_j \frac{\partial \Phi}{\partial x_j} = 0 \quad (4)$$

$$\frac{\partial z_b}{\partial t} + \frac{1}{1-p} \frac{\partial q_j}{\partial x_j} = 0, \quad (5)$$

112 where $i, j = 1, 2$, with summation being on j ; $x_{1,2} = (x, y)$ and $u_{1,2} = (u, v)$,
 113 where u and v are the cross- and alongshore depth-averaged current respec-
 114 tively. t represents time. $z_s(x, y, t)$ is the mean sea level, $z_b(x, y, t)$ is the mean
 115 bed level and D is the total mean depth ($D = z_s - z_b$). $E(x, y, t)$ is the
 116 wave energy density, which can be expressed in terms of the wave height
 117 ($E = \frac{1}{8} \rho g H^2$). τ_{bi} represents the bed shear stress; here the expression of *Fed-*
 118 *dersen et al.* (2000) is used. g is the gravitational acceleration, Φ is the wave
 119 phase and σ is the intrinsic frequency. The sediment flux (q_i) is represented
 120 by the formula of Soulsby and Van Rijn (*Soulsby*, 1997). The bed porosity p
 121 is 0.4 and the seawater density (ρ) is 1024 kg m^{-3} . S'_{ij} is the radiation stress
 122 term and S''_{ij} represents the Reynolds stresses (*Calvete et al.*, 2005). \mathcal{D} is the
 123 wave energy dissipation due to wave breaking described according to *Church*
 124 *and Thornton* (1993).

125

126 2.1 Linear stability analysis

In the usual way, our variables consist of an alongshore- and time invariant
 solution of (1)-(5), the basic state, denoted here with a zero subscript, and a
 small perturbation to that solution.

$$\begin{aligned} \{z_s, z_b, u_1, u_2, E, \Phi\} = & \{Z_{s0}(x), Z_{b0}(x), 0, V_0(x), E_0(x), \Phi_0(x, t)\} \\ & + \Psi(x) \exp(\omega t + iky). \end{aligned} \quad (6)$$

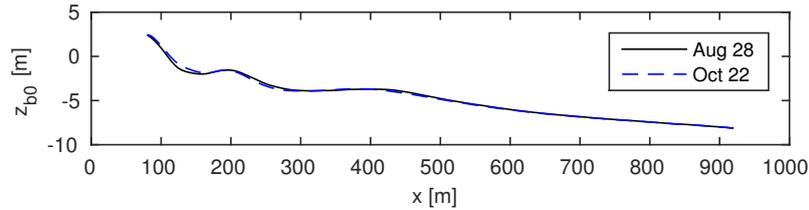


Fig. 1. Bed level profile resulting from alongshore averaging of the bathymetric surveys at the beginning and end of the two-month period.

127 The basic state corresponds to the wave conditions and water levels pertaining
 128 throughout the 2 months at Duck (see §2.2). It contains bed level Z_{b0} , mean
 129 water level Z_{s0} , alongshore current V_0 , wave density E_0 and phase Φ_0 . The
 130 second term on the right hand side of (6) is the perturbation. The disturbances
 131 considered are alongshore-periodic with arbitrary wavelength $\lambda = 2\pi/k$, and
 132 (complex) frequency $\omega = \omega_r + i\omega_i$. Thus the real part of the frequency w_r
 133 represents the growth rate of the periodic pattern, while the imaginary part
 134 $c_m = -w_i/k$ represents the corresponding migration rate. A pattern with
 135 positive w_r indicates a mode unbounded in time, i.e. an unstable mode. For a
 136 chosen k , the evolution of the perturbation is solved as an eigenvalue problem
 137 for eigenvalue ω and eigenfunction Ψ .

138 2.2 Basic state: field observation at Duck, 1998

139 The basic state is also referred to as the forcing. This forcing is the observed
 140 wave and tidal conditions recorded over a two month period in 1998, from
 141 August 20th (day 232) until October 22nd (day 294) (*Van Enckevort et al.*,
 142 2004). Wave data were recorded at three hour intervals. The same frequency
 143 was therefore used to obtain predictions from the model. Bathymetric evolu-
 144 tion was only recorded at the beginning and end of this 2-month period. So,
 145 the alongshore averaged bathymetric profile was determined every three hours
 146 by linear interpolation between the two alongshore-averaged profiles that were
 147 constructed from the full bathymetric surveys at the beginning and end of this
 148 period. In Fig. 1 we can see these two initial and final profiles.

149 Note that the tidal variation was included in the analysis by shifting the
 150 bathymetry vertically. The reproduced wave conditions and water depth are
 151 shown in Fig. 2. It can be seen that there are three times at which wave heights
 152 are increased for short durations (at about days 237, 263 and 272). We refer
 153 to these as storms 1, 2 and 3 respectively. Wave directions switch between
 154 northerly and southerly (with respect to the local coast), and so are likely
 155 to generate longshore currents in opposite directions at various times; some
 156 normally incident waves can also be seen. Periods are mostly confined within
 157 5 and 15s. A clear tidal signal can be seen in the depths.

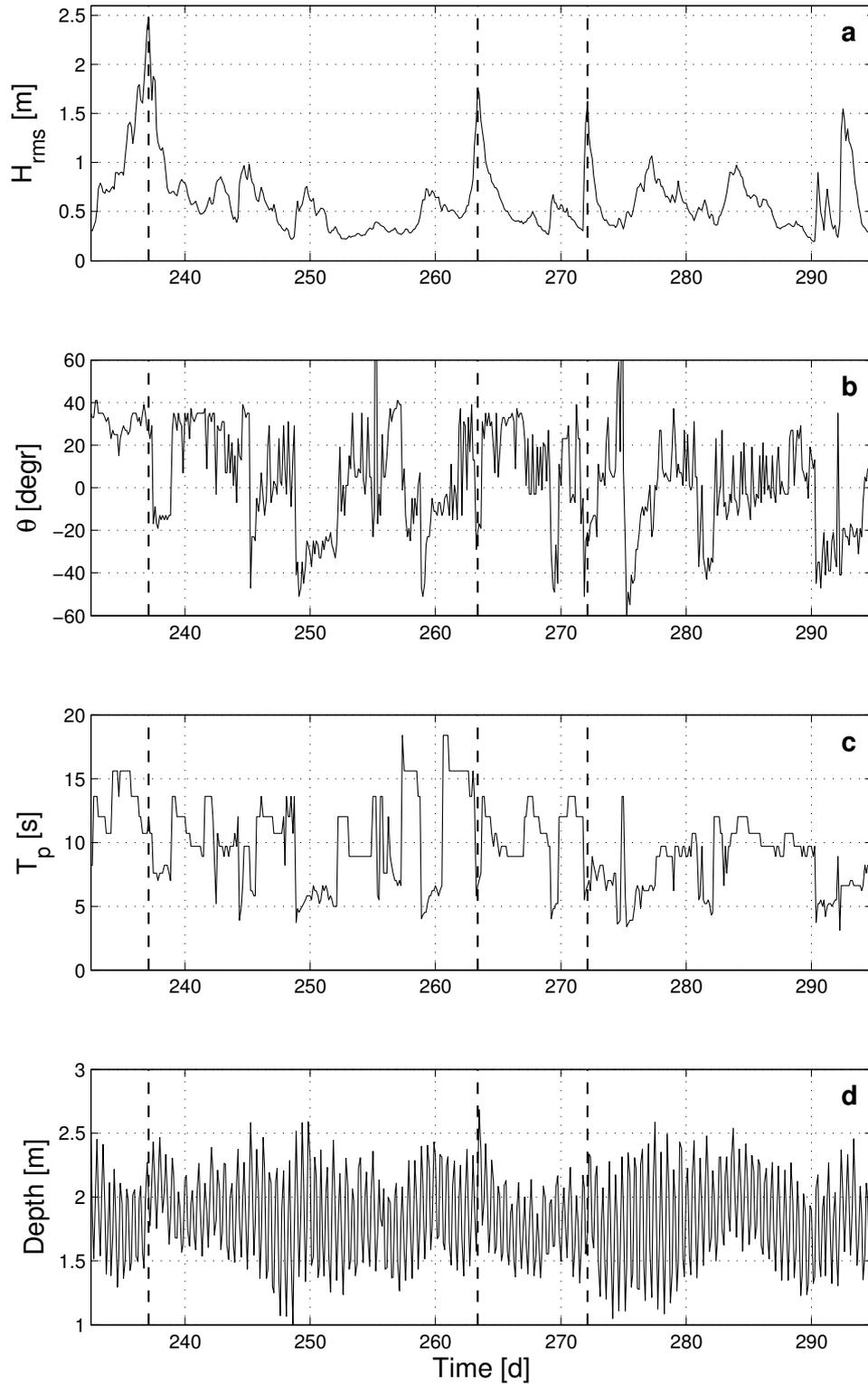


Fig. 2. Forcing conditions used in the linear stability model, as they were measured during observations period. (a) Wave height; (b) Wave angle relative to the coast (0° is perpendicular to the coast); (c) Wave peak period; (d) Water depth above the onshore bar. This water depth changes as a result of tides, surges and the bathymetric evolution. The vertical dashed lines represent storm events.

159 As mentioned in §2.1, k is arbitrary. So, we calculate the growth rate of all
 160 realistic morphodynamic lengthscales: $0.001 < k < 0.1$ [rad m⁻¹], for in-
 161 crements $\Delta k = 0.001$ rad m⁻¹; corresponding λ values are approximately
 162 {6.3km, 3.1km, 2.1km, 1.6km, 1.3km . . . 65.4m, 64.8m, 64.1m, 63.5m, 62.8m}, for
 163 each set of forcing conditions (every three hours). It is assumed that the pre-
 164 dictions made for each set of forcing conditions are valid for the three hour
 165 period until a new set of conditions becomes available. We thus require an
 166 entire growth rate curve for this region of k space for each three-hour predic-
 167 tion. This allows us to identify a unique growth rate for each k , in order to
 168 determine the amplitude development of each lengthscale.

169 The identification of an entire growth rate curve corresponding to physical
 170 modes is complicated due to the presence of numerical solutions to the equa-
 171 tions. For each lengthscale, the number of possible solutions calculated equals
 172 the number (n) of computational cross-shore nodes, with most of these results
 173 only describing physically meaningless numerical (i.e. non-physical) solutions
 174 to the system. These numerical solutions generally display negative or near-
 175 zero growth rates and, therefore, obscure in particular the negative part of the
 176 physical growth rate curve.

177 For all modes we must be sure that we have correctly identified physical modes.
 178 These physical modes are identified by testing the convergence of eigenvalues
 179 and eigenfunctions as n increases. Runs were carried out with 300 ($n = 300$)
 180 and 450 nodes ($n = 450$). According to *Calvete et al.* (2005), 300 cross-shore
 181 nodes is sufficient to achieve convergence. Our tests lead to agreement with
 182 this condition.

183 This is done for all wavenumbers, resulting in multiple physical growth rate
 184 curves. An example of these curves is shown in Fig. 3. Among these physical
 185 growth rate curves, the one containing the highest growth rate for the region
 186 of k space being examined is chosen. This growth rate curve is considered to
 187 be the one that governs evolution of bed-forms for the 3 hours during which
 188 those forcing conditions pertain. Note, however (Fig. 3), that other physical
 189 curves do exist; we ignore these.

191 Every three hours, a separate prediction of the linear growth rate curve is
 192 created based on the new hydrodynamic forcing conditions and bathymetry.
 193 The variability of this growth rate curve over time is significant (see Fig. 4(a)).
 194 Calmer conditions (as occur from day 255 to 259, for instance) generally result

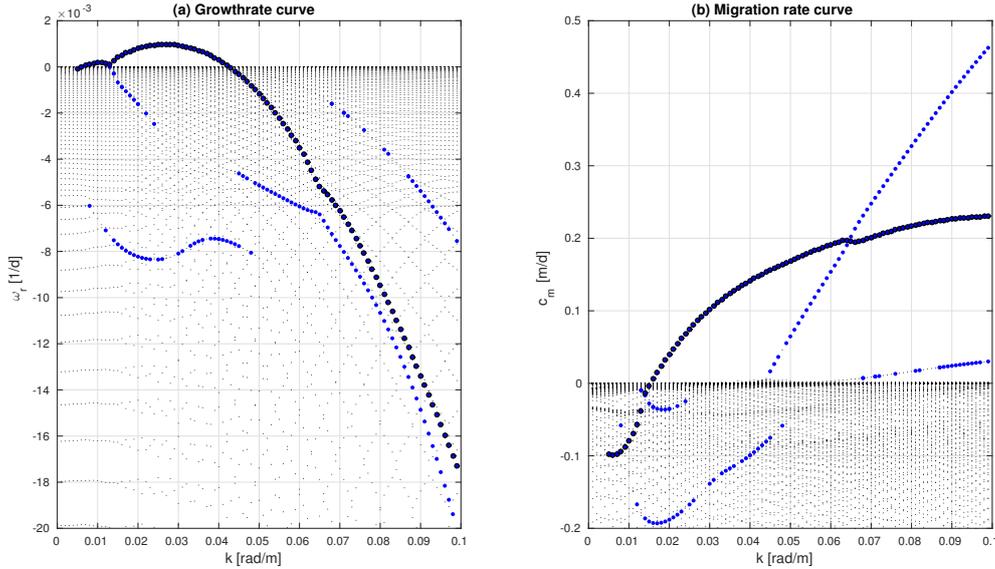


Fig. 3. (a) Growth rate (ω_r) curve; (b) Migration rate (c_m) curve. Shown are the distribution for all k -values of the solutions of the system of equations, with small black dots for all solutions from Morfo60, blue dots for all physical modes and black encircled blue dots for selected physical mode.

195 in very small growth rates, whereas bigger wave heights (as can be observed
 196 after day 237 in Fig. 2) result in both rapidly growing and decaying modes.
 197 The effect of the tidal variation can clearly be seen in the periodically varying
 198 growth rate.

199 The identification of the physical growth rates for each k -value has not been
 200 successful for all cases, as can be seen in Fig. 4(a, b). There are two situations
 201 when no physical growth rate could be obtained. Sometimes, the growth rate
 202 selected by the proposed method greatly deviates from neighbouring (in k
 203 space) growth rates. In these circumstances we deem that result non-physical,
 204 and to avoid seemingly unrealistic results, we set $\omega_r = 0$, see black dots in
 205 Fig. 4(b). Additionally, convergence is typically not achieved under more ex-
 206 treme storm conditions. When this occurred, it was assumed that all length-
 207 scales would show neither growth nor decay again ($\omega_r = 0$), see vertical black
 208 bars in Fig. 4(b). For most of the cases, however, a growth rate is available. As
 209 shown in Fig. 4(b), the percentage of lengthscales that lack a physical growth
 210 and migration rate over time is about 4%.

211 3 Model formulation: amplitude development

212 The bed-pattern lengthscale with the highest amplitude at any instant is
 213 deemed dominant and most likely to be observed in the field. *Tiessen et al.*
 214 (2010) took this lengthscale to be that corresponding to the *FGM* at different

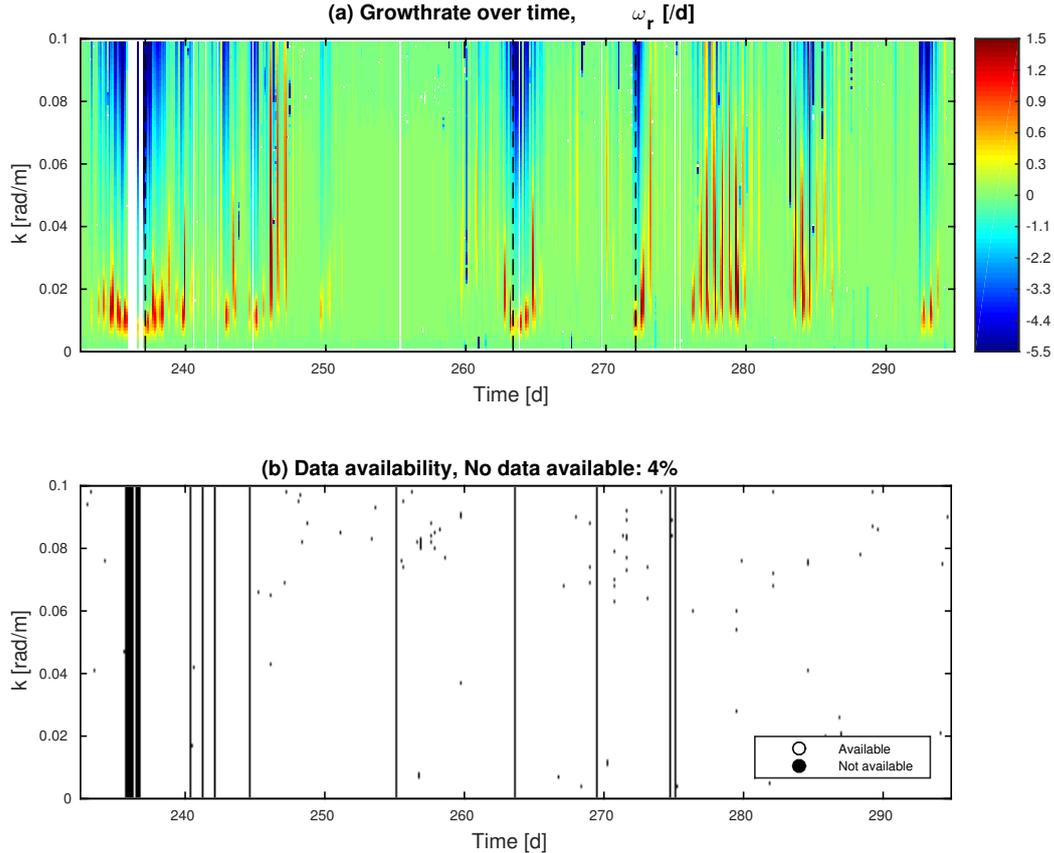


Fig. 4. (a) The growth rate curve at each time step as derived by selecting the physical growth rate curve as described in §§ 2.3 and 2.4. Blue indicates negative growth rate and red positive growth rate, and the black dashed line indicates the time of the peak of a storm. (b) Durations where no growth rate curve could be determined (in black).

215 times. Here we identify amplitude development for all lengthscales and derive
 216 the dominance of one lengthscale based on competition between these ampli-
 217 tudes, each of which is influenced by, but not solely dependent on, the linear
 218 growth rate.

219 A systematic approach to doing this is a weakly nonlinear perturbation expan-
 220 sion (see e.g. *Schielen et al.*, 1993). This approach results in a rapidly increas-
 221 ing number of different harmonics of k . Motivated by *Tiessen et al.* (2011)
 222 we limit our investigation to linear growth, self-limitation of that growth (i.e.,
 223 equilibration, or saturation), and the generation of the first harmonic. This
 224 approach is in keeping with that of *Knaapen and Hulscher* (2001), who used
 225 data-assimilation techniques to derive coefficients of an amplitude evolution
 226 equation that would result from a weakly nonlinear analysis. We thus hy-
 227 pothesise that the two most important nonlinear effects in the long-term de-
 228 velopment of crescentic bars are: i) equilibration of growing modes for all k
 229 values; and ii) generation of higher harmonics by growing modes, which there-
 230 fore allow energy to be transferred to smaller wavelengths. This generation is

	$-3k$	$-2k$	$-k$	0	k	$2k$	$3k$
$O(\epsilon^0)$				✓			
$O(\epsilon^1)$			✓		✓		
$O(\epsilon^2)$		✓		x		✓	
$O(\epsilon^3)$	x		✓		✓		x

Table 1

Schematic depiction of the harmonics included in the amplitude evolution model; a ✓(x) indicates inclusion (exclusion). ϵ represents the (small) amplitude of the bed pattern.

231 depicted schematically in Table. 1. The $O(\epsilon^0)$ term is our basic state, which
 232 remains unchanged. We consider the linearly growing (fundamental) mode (at
 233 $O(\epsilon^1)$), and the first harmonic ($O(\epsilon^2)$) that it generates by self-interaction. As
 234 noted, we exclude alterations to the mean bed (basic state). Being a mean
 235 component this will not affect lengthscale evolution. However, interaction of
 236 the mean term with the fundamental mode (that of the linear instability) will
 237 give rise to an equilibration (saturation) term at $O(\epsilon^3)$; this is included. Sec-
 238 ond and higher harmonics are excluded. Note also that we assume this model
 239 to pertain for all k values.

We choose the generic amplitude equation that can result from a weakly nonlinear analysis, which embodies the energy transfers described above (see *Drazin and Reid*, 1981). This is:

$$\frac{dA_k}{dt} = \omega_{rk}(t_n) A_k - l_k(t_n) A_k^3 + m_{k/2}(t_n) A_{k/2}^2. \quad (7)$$

Note that $A_k(t)$ is here our bed-form (mode) amplitude hereafter, where the k subscript refers to the lengthscale to which this amplitude pertains (also for ω_{rk}). The other coefficients in (7) are:

$$l_k(t_n) = |\omega_{rk}(t_n)|, \quad m_{k/2} = \alpha(1 - A_k^{10}),$$

240 where α is a constant. The first term on the right represents the linear growth
 241 (or decay). The amplitude ($A_k(t)$) is therefore an initially exponentially grow-
 242 ing (or decaying) quantity, assuming a small enough initial amplitude, with
 243 growth rate $\omega_{rk}(t_n)$. $A_k(t = 0) = A_{min} = 0.1$ is the same for all lengthscales;
 244 this is also the minimum amplitude. During storm events, all pre-existing bed-
 245 forms are expected to be erased. This is simulated by resetting the amplitudes
 246 of all lengthscales to A_{min} . The maximum amplitude $A_{max} = 1$; as amplitudes
 247 approach this value it is assumed that nonlinear effects will become domi-
 248 nant, and so further linear development is assumed to cease as this limit as
 249 approached. The values of A_{min} and A_{max} do not convey any intrinsic mean-
 250 ing themselves, except that choosing $A_{max} = 1$ is consistent with the weakly
 251 nonlinear nature of the expansion (i.e. all powers of $A_k < 1$) and can be done

252 without loss of generality. The value of A_{min} therefore is arbitrary, except
 253 that a ten-fold growth seems to represent roughly the duration it takes for a
 254 crescentic bathymetry to reach a new stable situation after a storm.

255 This assumption regarding A_{max} motivates the choice for $l_k = |\omega_{rk}(t_n)|$, the
 256 coefficient of the second term on the right. This ensures the desired long-term
 257 behaviour. This $O(\epsilon^3)$ term represents the equilibration, and the amplitude
 258 equation including just the first two terms on the right is the Stuart-Landau
 259 equation (*Drazin and Reid*, 1981). The final term in (7) allows energy transfer
 260 to A_k from lengthscales twice those of the lengthscale $\lambda = \frac{2\pi}{k}$. The energy
 261 transfer factor, $\alpha = 0.3$, is chosen based on the rate of energy transfer observed
 262 by *Tiessen et al.* (2011). In § 5.4 we examine the sensitivity of the simulations
 263 to changes in α . The dependence of $m_{k/2}$ on A_k is included here to ensure that
 264 all modes can only achieve the same maximum amplitude, so that this term, if
 265 operational, accelerates growth only, and becomes inoperational as $|A_k| \rightarrow 1$.
 266 This dependence is the only part of (7) that would not result from a weakly
 267 nonlinear analysis.

268 3.1 Numerical experiment on synthetic data

269 Before applying (7) to the data-set for Duck, we first illustrate the effect of
 270 the various terms on the right of (7) by means of an idealised but repre-
 271 sentative example. This example consists of two different forcing conditions
 272 consecutively applied for 12.5 days each. In Fig. 5 (a) and (b) we show the
 273 (time-invariant) growth rate curves corresponding to these two sets of forcing
 274 conditions. In Fig. 5 (c), (d) and (e) this results in the development of differ-
 275 ent crescentic bed-patterns with regards to lengthscale λ (or k) and amplitude
 276 (A_k), for three scenarios: Fig 5 (c) linear evolution (first term on the right of
 277 (7) only); Fig 5 (d) equilibration (first two terms on the right of (7) only); and
 278 Fig 5 (e) full model, i.e., linear evolution, equilibration and higher harmonic
 279 generation (all terms on the right of (7)).

280 In the early stages of linear evolution (Fig. 5(c)) there is rapid development of
 281 the lengthscale $\lambda_1 = 700$ m. This is the lengthscale of the *FGM* for the first
 282 forcing condition (denoted here *FGM*₁, green line, see caption). After the first
 283 forcing conditions (Fig. 5(a)) have been applied for 12.5 days, the second set
 284 of forcing conditions (Fig. 5(b)) results in a decay of *FGM*₁, which remains
 285 dominant until the *FGM* of the new conditions (*FGM*₂, blue line) surpasses
 286 it. During day 23, A_{k_2} exceeds A_{max} , so further development is terminated.
 287 Note also the growth of lengthscale $\lambda'_1 = 785$ m (k'_1) in the first 12.5 days: see
 288 Fig. 5 (a) and (c). This corresponds to that of the mode *FGM*'₁ with growth
 289 rate almost as large as that of *FGM*₁. This mode grows and decays much like
 290 *FGM*₁.

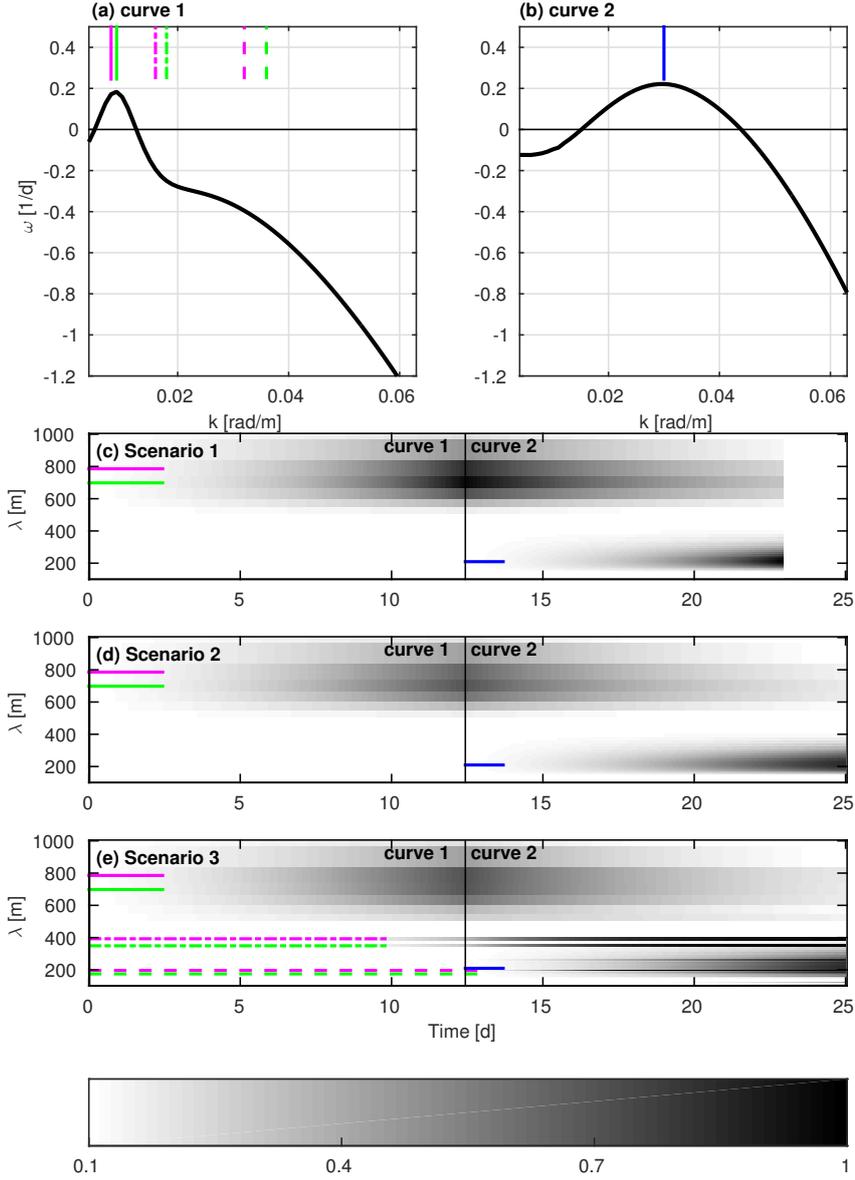


Fig. 5. Example of the three different cases: (a,b) Two different growth rate curves applied consecutively for 12.5 days; (c) linear evolution only; (d) equilibrated solution; (e) full model. Light (dark) shading indicates low (high) amplitude. Coloured lines indicate the position in k space (a,b) or λ space (c-e) of modes that exhibit significant growth in one or more cases. Solid lines: modes that only grow linearly. Green: FGM_1 (FGM corresponding to growth rate curve from the first forcing conditions, at $k = k_1 = 0.009$ rad m^{-1}); Magenta: FGM'_1 (mode adjacent to FGM_1 , for which ω_r is only slightly smaller than that for FGM_1 under first forcing conditions, $k = k'_1 = k_1 - \Delta k = 0.008$ rad m^{-1}); Blue: FGM_2 (FGM corresponding to the growth rate curve from second forcing conditions, at $k = k_2 = 0.03$ rad m^{-1}). Dash-dotted lines: Green: higher harmonic of FGM_1 ($2k_1$); Magenta: higher harmonic of FGM'_1 ($2k'_1$). Dashed lines: further higher harmonics ($4k_1, 4k'_1$) of FGM_1 and FGM'_1 . The lengths of the lines is for illustrative purpose only.

291 For the equilibration case (Fig. 5(d)) bathymetric evolution is self limiting.
292 As the amplitudes increase, again, centred around k_1 for the first 12.5 days,
293 the rate of increase decreases, especially toward the end of this period. The
294 subsequent transition from the first to the second forcing conditions (growth
295 centred on k_1 to growth centred on k_2) leads to similar behaviour. However,
296 now the amplitude development levels off when the amplitude approaches 1.

297 For the full model (Fig. 5(e)) we see qualitatively different behaviour. A small
298 but significant amount of energy is fed into $2k_1$ and $2k'_1$ during the first 12.5
299 days, by higher harmonic generation. Under the second set of forcing condi-
300 tions these wavelengths correspond to linearly growing modes, and so these
301 continue to evolve during the latter 12.5 days. Additionally, $4k_1$ and $4k'_1$ are
302 similarly excited, and these modes lie close to k_2 , so that even though they ini-
303 tially possess only limited amplitudes they ultimately grow rapidly. The result
304 is a broader range of lengthscales (modes) containing significant amplitudes.

305 4 Results

306 4.1 *The evolution of crescentic bars*

307 The model predictions representing the two months of field observations at
308 Duck (NC) for the three cases are shown in Fig. 6, where the amplitude de-
309 velopment for all examined lengthscales is shown over time. We show the
310 equivalent three cases to illustrate the effects of the inclusion of these physical
311 mechanisms on predictions. For the predictions made solely by linear growth
312 rates (Fig. 6(c)), the amplitude development is terminated when the fastest
313 growing lengthscale reaches A_{max} (about day 246, after storm 1). Predictions
314 only resume immediately after a storm (shown as dashed lines), which is as-
315 sumed to erase all existing bed-patterns. This eradication of pre-existing bed-
316 forms during a storm is also applied for the other cases. During the subsequent
317 bed evolution, the development of crescentic bars starts again from A_{min} .

318 The rate of development after the first and third storms is similar, which can
319 be seen in the emergence of significant amplitudes at similar post-storm times.
320 This development is larger than that after the second storm. The growth rate
321 curve (Fig. 4(a)) shows why this difference happens. The only large growth
322 rates after the second storm occur immediately after it, as the wave height
323 is subsiding from its peak. In contrast, both the first and third post-storm
324 periods exhibit significant durations when growth rates are significant (see
325 the regions with 'red' growth rates in Fig. 4(a)). These durations roughly
326 correspond to times when $H_{rms} > 0.5m$ (see Fig. 2(a)). Furthermore, the time
327 interval between second and third storms is shorter than that between first

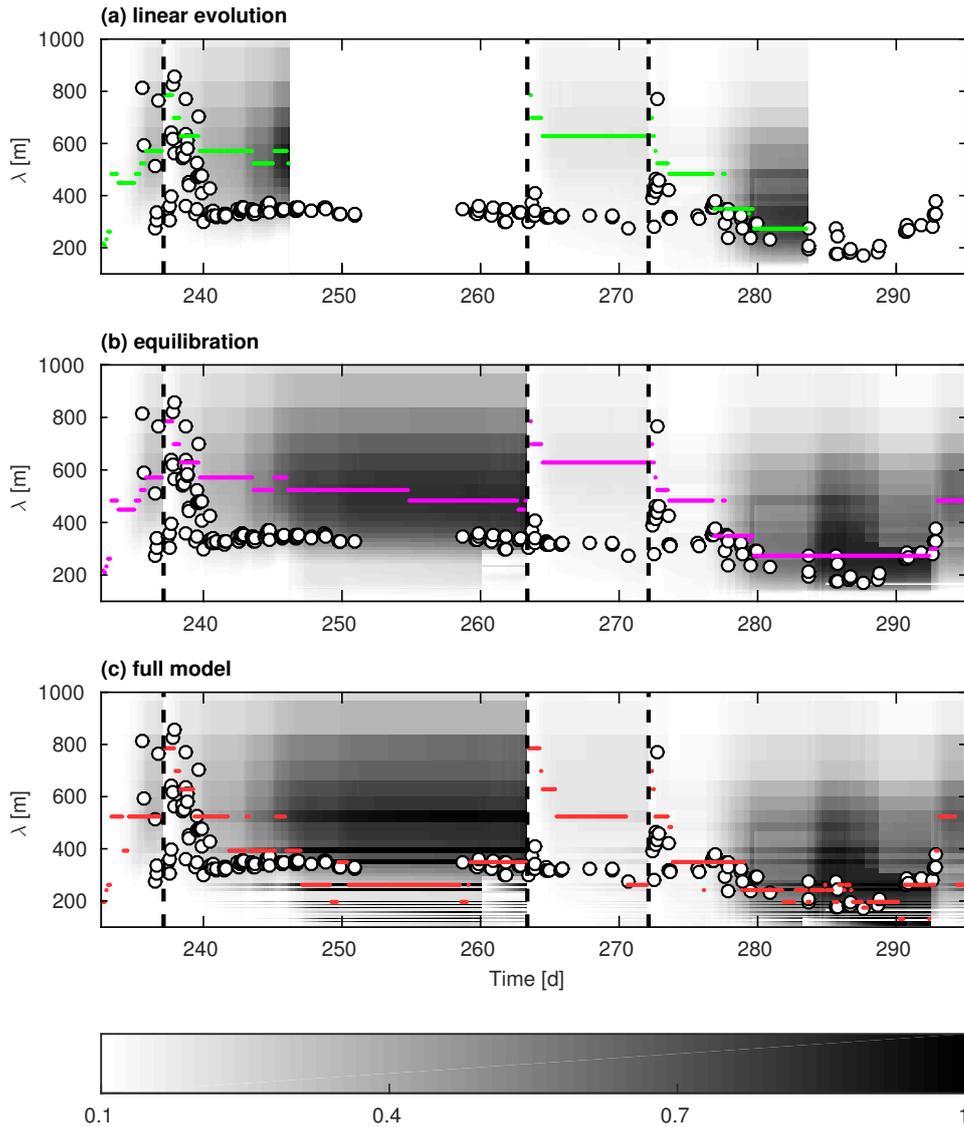


Fig. 6. Amplitude development for the three cases compared to the observed lengthscales (large white circles) (*Van Enckevoort et al., 2004*), where coloured dots denote the predicted dominant lengthscales. (a) linear evolution; (b) equilibration; (c) full model.

328 and second storms, thus allowing less time for development of these bed-forms.

329 For the equilibration case, development rates are reduced by the equilibration
 330 term during the latter post-storm stages. As a result, more gradual growth is
 331 seen latterly, but qualitatively behaviour is the same, except that the whole
 332 time period can now be accommodated.

333 In the case of higher harmonic interaction (full model), the simulation shows
 334 a significant amplitude transfer occurring from longer lengthscales to shorter
 335 lengthscales. This gives rise to a wider range of developing modes than is the
 336 case when only the linear evolution or equilibration are considered.

337 A comparison of the predicted and observed lengthscale evolution is also shown
338 in Fig. 6. The predicted dominant lengthscale (that of the biggest amplitude
339 at each time $t = t_n$) is shown as a coloured dot, and the observed lengthscales
340 are shown as larger white dots. In between storms, amplitude development
341 based on linear evolution and equilibration generally over-predict the domi-
342 nant lengthscale. Higher harmonic interaction (full model) results in a more
343 rapid development of shorter lengthscales which is more in line with field
344 observations (Fig. 6(c)). But after storm 1 the observed stabilisation of the
345 bed-form lengthscales is not reproduced in the full model. Fluctuations in pre-
346 dicted lengthscale are apparent, which, as can be seen from the amplitudes in
347 Fig. 6(c), are due to relatively small amplitude differences between a number
348 of co-existent modes.

349 4.2 Amplitude evolution

350 Due to the lack of observational data of the vertical amplitude of crescen-
351 tic bars, a straight comparison of the amplitude of the predicted dominant
352 lengthscales with field observation is not possible. However, in *Van Enckevort*
353 *et al.* (2004), the horizontal amplitude (A_y) of the crescentic bar at Duck is
354 recorded. This amplitude was calculated as half the average cross-shore dis-
355 tance between the bay and the two horns (Figure 7). We hypothesize that the
356 vertical amplitudes of crescentic bars is proportional to A_y . In figure 8, the
357 predicted amplitude of the dominant lengthscale (black curve) is compared
358 with the observed A_y (blue curve). The full model evolution of amplitude in
359 general fits well with observation, except after the second storm. Amplitude
360 growth and equilibration after storm 1 is consistent with that observed. Af-
361 ter storm 3 the model produces more rapid growth to a higher amplitude
362 than that observed, but, nonetheless, qualitatively similar behaviour. Again,
363 the effect of the higher harmonic interactions may be observed by compar-
364 ing figure 8 (b) and (c). The differences are small, but remember that the
365 simulated amplitudes are those of the dominant lengthscale, and these are in
366 general overpredicted by the equilibration model. The main difference is the
367 very limited amplitude development after storm 2 and that observed, which is
368 substantial. This, as also noted by *Tiessen et al.* (2010), points to the persis-
369 tence of bed-forms through the second storm. This will be further discussed
370 in §5.2.

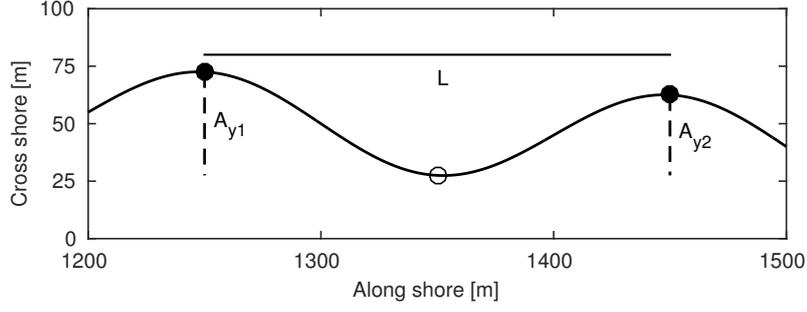


Fig. 7. A single crescent from the crescentic bar with length L and horizontal amplitude $A_y = 0.5 \times (A_{y1} + A_{y2})/2$. The horn of the crescent is labelled with filled circles, whereas the bay is labelled with open circle.

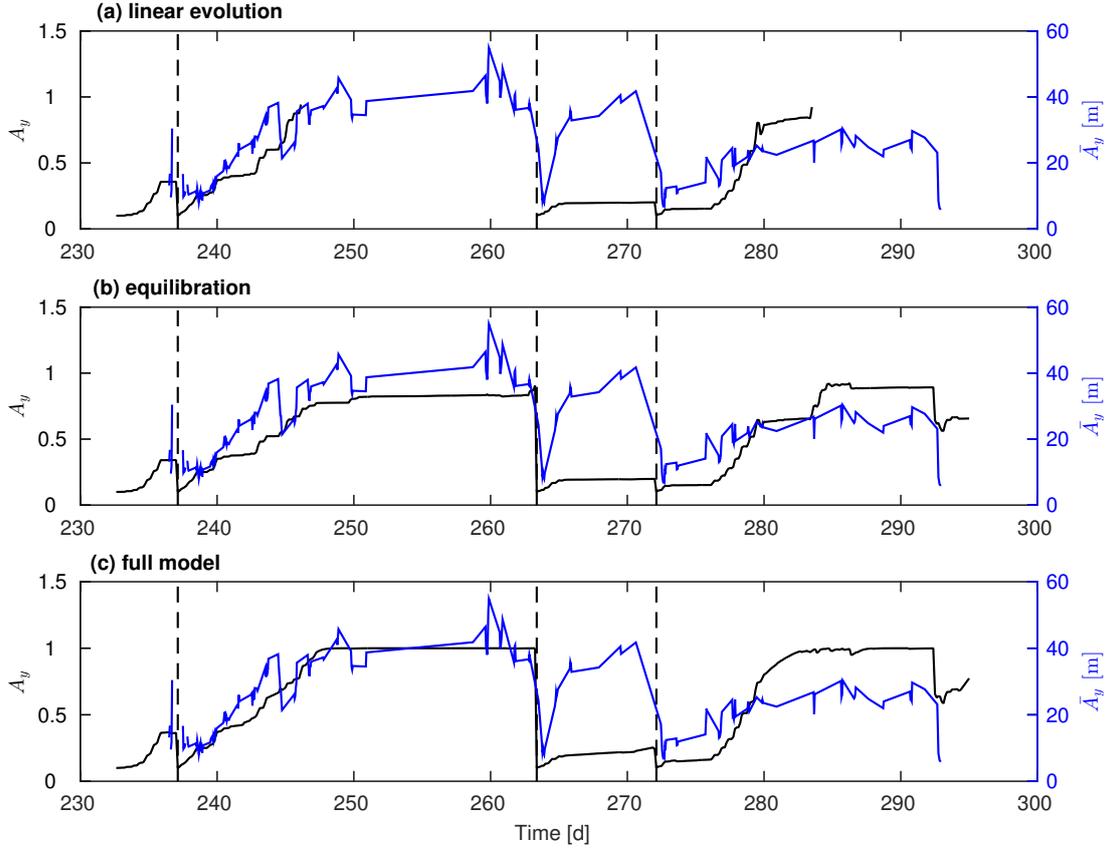


Fig. 8. Comparison between the observed and predicted dominant amplitudes, for three simulated cases: (a) linear evolution; (b) equilibration; (c) full model. The dark curve describes the amplitude of dominant lengthscale, whereas the blue curve refers to the observed longshore averaged horizontal amplitude (\bar{A}_y).

371 5 Discussion

372 5.1 Importance of nonlinear effects

373 The most striking effect is the higher harmonic interaction. A quantitative
 374 comparison between the observed and predicted lengthscales (Table 2) shows

	Absolute error [m]	Relative error [-]
Linear evolution	190	0.54
Equilibration	168	0.49
Full model	108	0.31

Table 2

The error between prediction and observation of the different scenarios. Note that the comparison is taken at the moments when observation could be made, and both the absolute and relative error are averaged values.

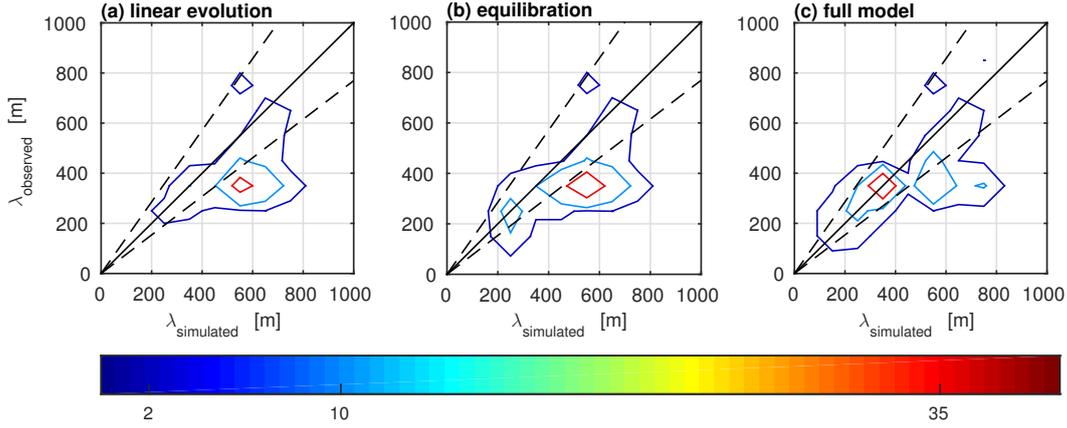


Fig. 9. Comparison between the observed and predicted dominant lengthscales, for three simulated : (a) linear evolution; (b) equilibration; (c) full model. The area between the dashed lines corresponds to relative error < 0.3 . The contour line denotes the density of data points, with red referring to high density and blue to low density.

375 that the inclusion of higher harmonic interaction reduced the absolute error
376 from 168 m (equilibration) to 108 m (full model), and relative error from
377 0.49 to 0.31. The improvement in correspondence with the inclusion of higher
378 harmonic interaction is also apparent in Fig. 9 where the predicted dominant
379 lengthscale is compared to the observed lengthscales at the moments when
380 observations could be made. The incorporation of the equilibration term is
381 necessary. Without the self limitation effect inherent in the equilibration term,
382 the bed development ceases when the linearly growing amplitude reaches the
383 maximum amplitude.

384 5.2 The persistence of bed pattern after storms

385 In the model we have assumed that all pre-existing bed-forms have been eradi-
386 cated after each storm, and the development of all lengthscales starts from the
387 same A_{min} . This assumption is based on the notion that each storm is pow-
388 erful enough and of long enough duration for an alongshore constant sandbar

389 to be formed. However, our model findings for the crescentic bed-pattern de-
 390 velopment after the second storm (similar to those presented in *Tiessen et al.*
 391 (2010)) are distinctly different to the field observations. As previously postu-
 392 lated in *Tiessen et al.* (2010), this might be due to the persistence of crescentic
 393 bed-forms throughout a comparatively less powerful storm. Moreover, apart
 394 from one observation at $\sim 700m$ (see Fig. 6) the observed lengthscales right
 395 after the third storm stay in a narrow band close to the dominant wavelength
 396 after the second storm. This is distinctly different from the fluctuation of
 397 lengthscales observed after the first storm, and consistent with the aforemen-
 398 tioned persistence of bedforms through the second storm.

To investigate this effect, we introduce a so-called persistence ratio (μ) of
 pre-existing bed patterns after storm,

$$\mu = \frac{A_{k,t_s^+} - A_{min}}{A_{k,t_s^-} - A_{min}},$$

399 where t_s^- (t_s^+) refers to the time immediately before (after) the storm. The
 400 value of μ therefore ranges from 0 to 1, where $\mu = 0$ (1) means that all
 401 pre-existing bed-forms have been eradicated (preserved). Previously (Fig. 6)
 402 $\mu = 0$ for all storms. Here we relate the value of μ to storm strength which is
 403 represented by the maximum wave height of each storm. From this perspective,
 404 storm 2 and 3 are of similar strength, whereas storm 1 is more powerful, see
 405 Fig. 2. We thus assume $\mu = 0$ after the first, and investigate the effect of
 406 varying the (same) value of μ after second and third storms for the full model
 407 (7). In Fig. 10 (black dashed line) we see the effect of this variation in μ .
 408 By allowing more bed amplitude to be preserved we observe a reduction in
 409 relative error of lengthscale as μ increases from 0 (its value in Fig. 6), and
 410 thereafter a modest increase. In fact, there is a max. error for $\mu = 0$. Further
 411 research is required to clarify the mechanism lying beneath μ . The sensitivity
 412 of model behaviour on μ is further discussed in §5.4.

413 5.3 Energy transferred to higher harmonics

414 The energy transferred from λ to $\frac{\lambda}{2}$ is characterised by a factor α (see § 3). As
 415 mentioned in § 3, the value of α in this study was chosen based on the rate of
 416 energy transfer observed by *Tiessen et al.* (2011). A high value of α indicates a
 417 rapid transfer of energy to $\frac{\lambda}{2}$ and hence probably leads to an earlier post-storm
 418 dominance of short wavelength. It is apparent (see Fig. 10 for $\mu = 0$) that the
 419 value used in Fig. 6 (following *Tiessen et al.*, 2011) gives something close to
 420 the minimum relative error for the full model.

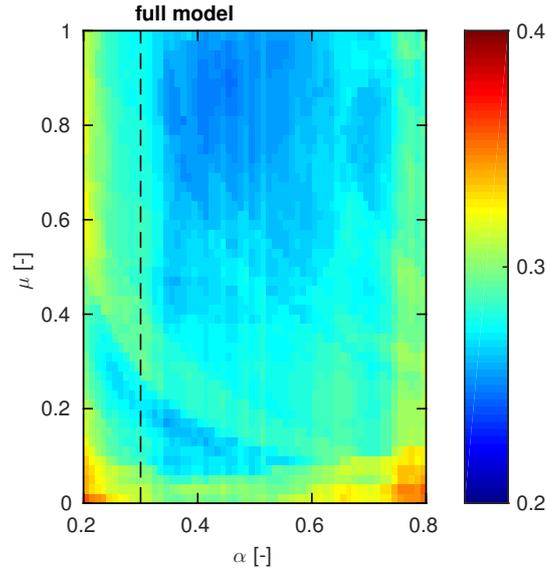


Fig. 10. sensitivity of full model behaviour on persistence ratio μ of pre-existing bed patterns and energy transfer factor α . The vertical black dashed line refers to the choice of $\alpha = 0.3$ in section 4. Colours indicate the relative error of the predicted dominant lengthscales and observed lengthscales, with blue for low relative error and red for high relative error.

421 5.4 Model sensitivity to μ and α

422 The full sensitivity of the full model behaviour to μ and α is shown in Fig. 10,
 423 with $0.2 \leq \alpha \leq 0.8$ and $0 \leq \mu \leq 1$ (note that we still assume that $\mu = 0$ for the
 424 first, larger storm). The relative error of the predicted dominant lengthscales
 425 and observed lengthscales is smaller for non-zero μ . This suggests that part
 426 of pre-existing bed pattern persists after second and third storms, and, by
 427 implication, that the second and third storm are not strong enough to erase
 428 all the existing bed forms. There is a region of broadly minimum error for
 429 about $0.2 \leq \mu \leq 1$ and $0.3 \leq \alpha \leq 0.6$. The conclusion appears to be that a
 430 higher μ after storm 2 and 3 leads to slightly better correspondence between
 431 prediction and observation.

432 The minimum error is actually achieved (Fig. 10) for $\alpha = 0.41$ and $\mu =$
 433 0.78 , resulting in a relative error of 0.24 (as compared to 0.31 for $\mu = 0$,
 434 $\alpha = 0.3$, see Table 2). Using these values we re-run the model for the full
 435 duration, and results are shown in Fig. 11. Additionally, we see results of the
 436 predicted dominant amplitude plotted against that observed. The predicted
 437 dominant amplitude now shows better correspondence with observation after
 438 the second storm, but poorer correspondence after the third storm. This and
 439 Fig. 9 suggest that these two storms correspond to different μ values.

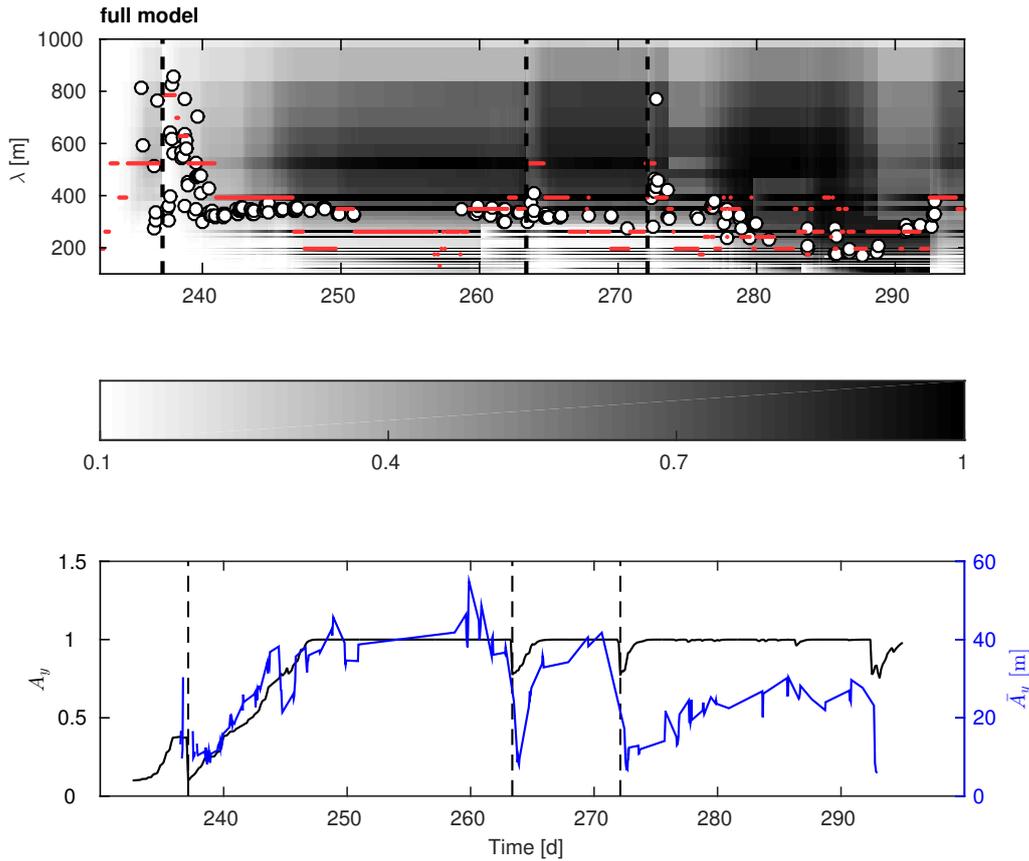


Fig. 11. Amplitude (top) and dominant amplitude (bottom) development for the full model with $\alpha = 0.41$ and $\mu = 0.78$.

440 6 Conclusions

441 In this study, we hypothesize that the dominant mechanisms for evolution of
 442 crescentic bar systems in nature are linear growth allied to equilibration (self-
 443 limitation) and higher harmonic generation by self-interaction. These mecha-
 444 nisms have been implemented into a model that would result from a weakly
 445 nonlinear perturbation analysis, but in which the coefficients of the nonlin-
 446 ear terms (in particular, that governing higher harmonic interactions) are set
 447 based on observations. This model is then used to investigate the bathymetric
 448 evolution of a crescentic-barred beach at Duck (NC). The model was used to
 449 reproduce a 2-month period, over which field observations were analysed by
 450 *Van Enckevort et al. (2004)*. Results show that nonlinear effects of equilibra-
 451 tion and higher harmonic interaction lead to significantly improved reproduc-
 452 tion of long-term evolution of a crescentic bar system in terms of observed
 453 lengthscales.

454 In between storms when crescentic bars develop, their initial development
 455 corresponds well with the results from a basic linear stability analysis. The

456 addition of a self-limitation term (*Drazin and Reid, 1981*) extends the predic-
457 tive range of the linear stability model to the entire post-storm period. The
458 inclusion of the term describing generation of higher-harmonics (as suggested
459 by *Tiessen et al., 2011*) leads to a significant improvement in prediction of
460 observed lengthscales. With these extra effects, an approach based on linear
461 stability analysis can describe the observed change from immediately post-
462 storm large lengthscales to the subsequent shorter lengthscales, related to
463 calmer conditions in between storm events, and the subsequent stabilisation
464 of the bed.

465 Note that the present approach is a significantly larger undertaking than that
466 of just determining a single fastest growing mode (*FGM*), i.e. corresponding to
467 a single k at one time, as done by *Tiessen et al. (2010)*. Here we must determine
468 a whole, unique growth rate curve at each time. Nonetheless, the present ap-
469 proach is still significantly less demanding in terms of computational time than
470 the simulations typically required to describe the development of the whole sea
471 bed over this area (this is generally done using a fully nonlinear model, and ei-
472 ther 2DH or 3D). An additional advantage of the currently proposed method is
473 the significantly reduced need for beach-specific parametrisation, because de-
474 tailed, spatially-variable planform-bathymetric data is not required. Similarly,
475 only relatively idealised and schematised conditions regarding wave climate
476 and tidal elevation are needed for a linear stability approach.

477 Whilst these findings represent an improvement on a linear stability model
478 (*Tiessen et al., 2010*), several effects are not yet included or fully understood.
479 For instance, the occurrence of a storm-related eradication of the crescentic
480 bed-forms needs to be further investigated. The current research suggests that
481 certain storms might not be strong enough to cause a complete wipe-out.
482 Additionally, the energy transferred in the higher harmonic interaction is not
483 yet quantified. More work is needed on developing a systematic approach to
484 deriving the amplitude equations. Note also that in our approach we consider
485 discrete wavelengths as opposed to the continuum of wavelengths that are
486 described by a Ginsburg-Landau equation (*Schielen et al., 1993*). Finally, note
487 that for some forcing conditions there is likely to be more than one physically
488 relevant growth rate curve (see Fig. 3).

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