

1 Remote sensing of mangrove forest phenology and its environmental drivers

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12

13 **Abstract**

14 Mangrove forest phenology at the regional scale have been poorly investigated and its driving factors
15 remain unclear. Multi-temporal remote sensing represents a key tool to investigate vegetation
16 phenology, particularly in environments with limited accessibility and lack of *in situ* measurements.
17 This paper presents the first characterisation of mangrove forest phenology from the Yucatan
18 Peninsula, south east Mexico. We used 15-year time-series of four vegetation indices (EVI, NDVI,
19 gNDVI and NDWI) derived from MODIS surface reflectance to estimate phenological parameters
20 which were then compared with *in situ* climatic variables, salinity and litterfall. The Discrete Fourier
21 Transform (DFT) was used to smooth the raw data and four phenological parameters were estimated:
22 start of season (SOS), time of maximum greenness (Max Green), end of season (EOS) and length of
23 season (LOS). Litterfall showed a distinct seasonal pattern with higher rates during the end of the dry
24 season and during the wet season. Litterfall was positively correlated with temperature ($r=0.88$,
25 $p<0.01$) and salinity ($r=0.70$, $p<0.01$). The results revealed that although mangroves are evergreen
26 species the mangrove forest has clear greenness seasonality which is negatively correlated with
27 litterfall and generally lagged behind maximum rainfall. The dates of phenological metrics varied
28 depending on the choice of vegetation indices reflecting the sensitivity of each index to a particular
29 aspect of vegetation growth. NDWI, an index associated to canopy water content and soil moisture
30 had advanced dates of SOS, Max Green and EOS while gNDVI, an index primarily related to canopy
31 chlorophyll content had delayed dates. SOS ranged between day of the year (DOY) 144 (late dry
32 season) and DOY 220 (rainy season) while the EOS occurred between DOY 104 (mid-dry season) to
33 DOY 160 (early rainy season). The length of the growing season ranged between 228 and 264 days.
34 Sites receiving a greater amount of rainfall between January and March showed an advanced SOS and
35 Max Green. This phenological characterisation is useful to understand the mangrove forest dynamics
36 at the landscape scale and to monitor the status of mangrove. In addition the results will serve as a
37 baseline against which to compare future changes in mangrove phenology due to natural or
38 anthropogenic causes.

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40 **1. Introduction**

41 Mangroves are a taxonomically diverse assemblage of tree species which have common
42 morphological, biochemical, physiological and reproductive adaptations that allows them to colonise
43 and develop in saline, hypoxic environments (Alongi, 2016). These assemblages form intertidal
44 forests which are one of the most carbon rich ecosystems (Donato et al., 2011) because of their high
45 productivity (Twilley and Day, 1999), rapid sediment accretion (Bouillon et al., 2008) and low
46 respiration rates (Barr et al., 2010). Vegetation phenology, defined as the growing cycle of plants and
47 involving recurring biological events such as leaf unfolding and development, flowering, leaf
48 senescence and litterfall (Njoku, 2014), regulates the timing of plant photosynthetic activity and
49 influences directly the annual vegetation carbon uptake. Vegetation phenology has been a focus of
50 attention in recent years due to a strong and measurable link between biological events and climate
51 (Cleland et al., 2007; Richardson et al., 2013; Dannenberg et al., 2015).

52 Historically, vegetation phenology was based on field records of key biological events such as
53 budburst, flowering, seed set and leaf senescence (Fitter and Fitter, 2002). Recently, a network of
54 fine-resolution digital cameras installed in the field known as “phenocams”, emerged as a new method
55 to monitor vegetation phenology (Richardson et al., 2007). While this method reduces the subjectivity
56 of human observations, it is limited by its relatively small spatial extent across the globe (Mizunuma
57 et al., 2013; Klosterman et al., 2014). Alternatively, as the reflectance properties of vegetated land
58 varies seasonally in relation to vegetation phenology, the systematic, multi-temporal data collected by
59 optical satellite sensors offer a unique mechanism to monitor vegetation dynamics as this approach
60 allows monitoring of an entire ecosystem rather than individual trees (Reed et al., 2009; White et al.,
61 1997, 1999). This has led to the rise of a new field known as land surface phenology (LSP) (Hanes et
62 al., 2014).

63 In the last few decades, LSP, which uses time-series of satellite-derived vegetation indices, has
64 received considerable attention given its potential to characterise the interactions between vegetation
65 and climate at broad geographical scales. Pioneer work in the temperate latitudes of the American

66 continent showed that the start of vegetation greening was controlled by pre-season temperature
67 (White et al., 1997). Moulin et al. (1997) conducted one of the first attempts to map global vegetation
68 phenology using the Advanced Very High Resolution Radiometer (NOAA/AVHRR) at 1° x 1°
69 resolution. The study revealed patterns in the global vegetation phenology related to seasonal
70 variation in climate. For example, the start of vegetation greenness in temperate deciduous forests was
71 strongly influenced by temperature, whereas savannahs were more influenced by rainfall. The
72 capability to study global vegetation dynamics increases as more advanced sensors and new
73 algorithms become available. Zhang et al., (2006) mapped global vegetation phenology at a finer
74 spatial resolution (1 km x 1 km) achieving more realistic results using the Enhanced Vegetation Index
75 (EVI) derived from the Moderate Resolution Imaging Spectroradiometer (MODIS). In the last decade,
76 numerous studies have been carried out to analyse patterns in vegetation phenology at continental and
77 regional scales at a variety of latitudes including the boreal ecosystem (Xu et al., 2013; Jeganathan et
78 al., 2014), Europe (Stöckli and Vidale, 2004; Rodriguez-Galiano et al., 2015a), India (Dash et al.,
79 2010) and the Amazon Forest (Xiao et al., 2005b).

80 Despite increasing interest in the use of remote sensing to characterise vegetation phenology, most
81 studies of mangrove phenology rely on traditional field methods consisting on *in situ* collection of
82 different components of litterfall (e.g. leaves, branches, flowers and propagules) (Leach and Burgin,
83 1985). Studies of this nature were documented in mangrove forests across the globe (see Table 1), and
84 those studies have indicated that litterfall dynamics and reproductive phenology of mangroves is
85 driven by a complex interaction of ecological (species composition, competition, reproductive
86 strategy), climatic (air temperature, precipitation, evaporation, hours of sun, wind speed) and local
87 environmental factors (fresh water inputs, tides, flooding, soil salinity, soil nutrients) and natural
88 disturbances (e.g., hurricanes). Moreover, these studies revealed that although mangroves are
89 evergreen species that produce litterfall and replace old leaves continuously throughout the year they
90 generally present a peak of leaf fall, leaf emergence and reproductive structures in the wet season.
91 There are cases, however, where this pattern can be bimodal, with one leaf fall peak in the dry season
92 and one in the wet season.

93 Although the above studies provide a local perspective of the interaction between mangroves and
94 physical drivers, there are some limitations. For instance, mangroves are often distributed across
95 hundreds of kilometres of coastlines. Thus, spatially restricted studies do not support observation of
96 the phenology phases over the complete extent of the forest. In addition, a common characteristic of
97 those studies is the limited time span, ranging between 1 to 4 years. This relatively short period
98 prevents observing inter-annual variation and trends in phenological metrics and how they are driven
99 by any climatic factors. Spatially continuous and temporally rich information on mangrove phenology
100 would be useful to characterise mangrove forest dynamics at the landscape scale and understand their
101 contribution to biogeochemical cycles.

102 To date there has been no characterisation of mangrove forests phenology using remote sensing data.
103 In this paper, we estimate and map phenological metrics from time-series of medium resolution
104 satellite sensor imagery in a mangrove forest in the SE of Mexico and investigate their relationship
105 with environmental drivers. The objectives of this research were to (i) estimate phenological
106 parameters using a time-series of MODIS vegetation indices to explore the consistency among them,
107 (ii) map the spatial distribution of phenological metrics (start of growing season, time of maximum
108 greenness, end of growing season and end of season), and (iii) characterise the relationship between
109 phenology dynamics and environmental drivers.

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Table 1. Field studies addressing mangrove forest phenology.

Country	Reference
Australia	Coupland et al., 2005; Duke, 1990
Borneo	Sukardjo et al., 2013
Brazil	Mehlig, 2006; Fernandes, 1999
India	Upadhyay and Mishra, 2010; Wafar et al., 1997
Japan	Kamruzzaman et al., 2016; Sharma et al., 2014
Kenya	Slim et al., 1996
Malaysia	Hoque et al., 2015; Akmar and Juliana, 2012
Mexico	Agraz-Hernández et al., 2011; Utrera-López and Moreno-Casasola, 2008; Aké-Castillo et al., 2006; Arreola-Lizárraga et al., 2004; Day et al., 1987; Lopez-Portillo and Ezcurra, 1985
Panama	Cerón-Souza et al., 2014
South Africa	Rajkaran and Adams, 2007
United States of America	Castañeda-Moya et al., 2013

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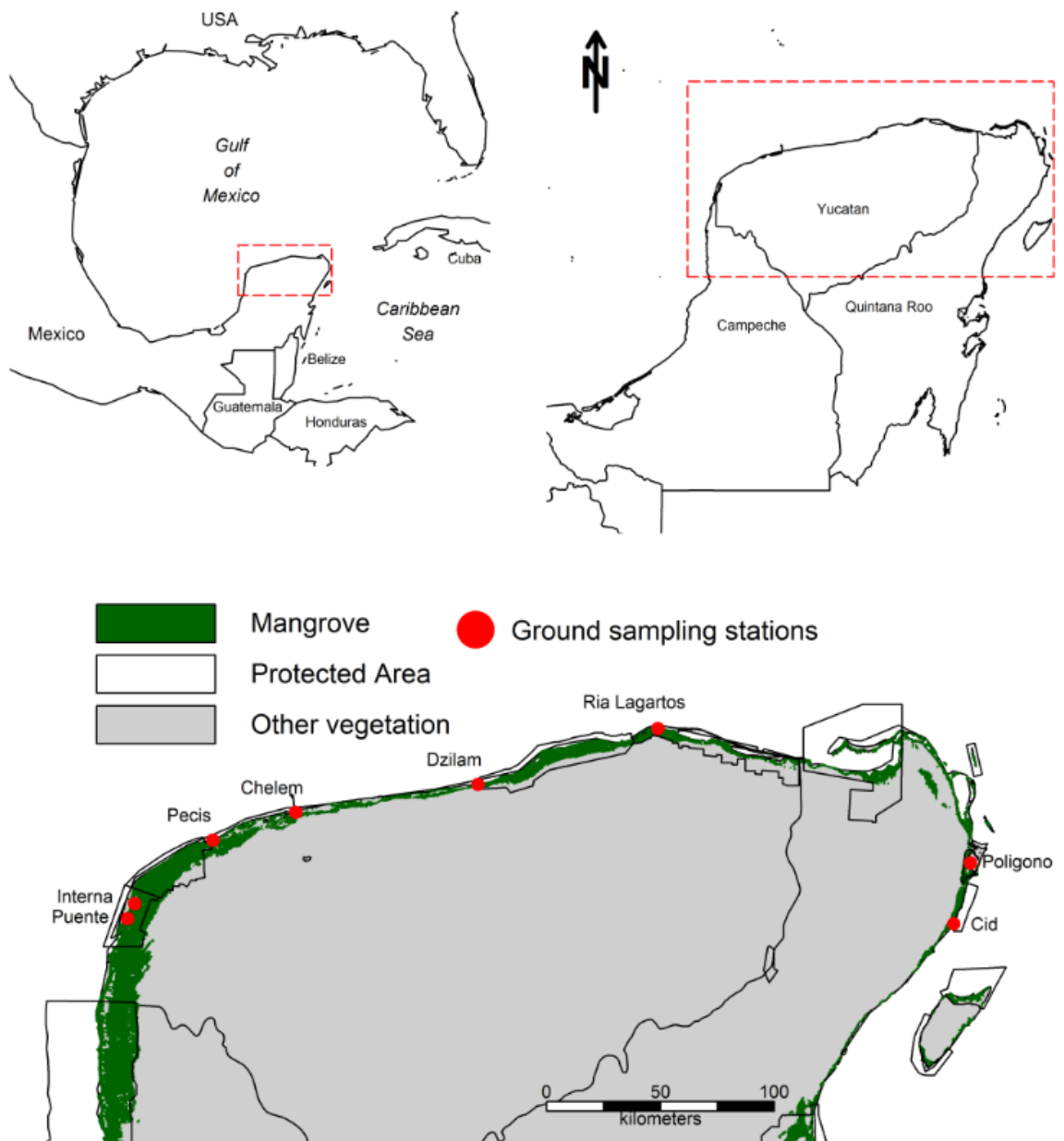
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Fig. 1. Study area showing eight ground sampling stations distributed along the Yucatan Peninsula coastline. The figure depicts the spatial distribution of the mangrove forest in green.

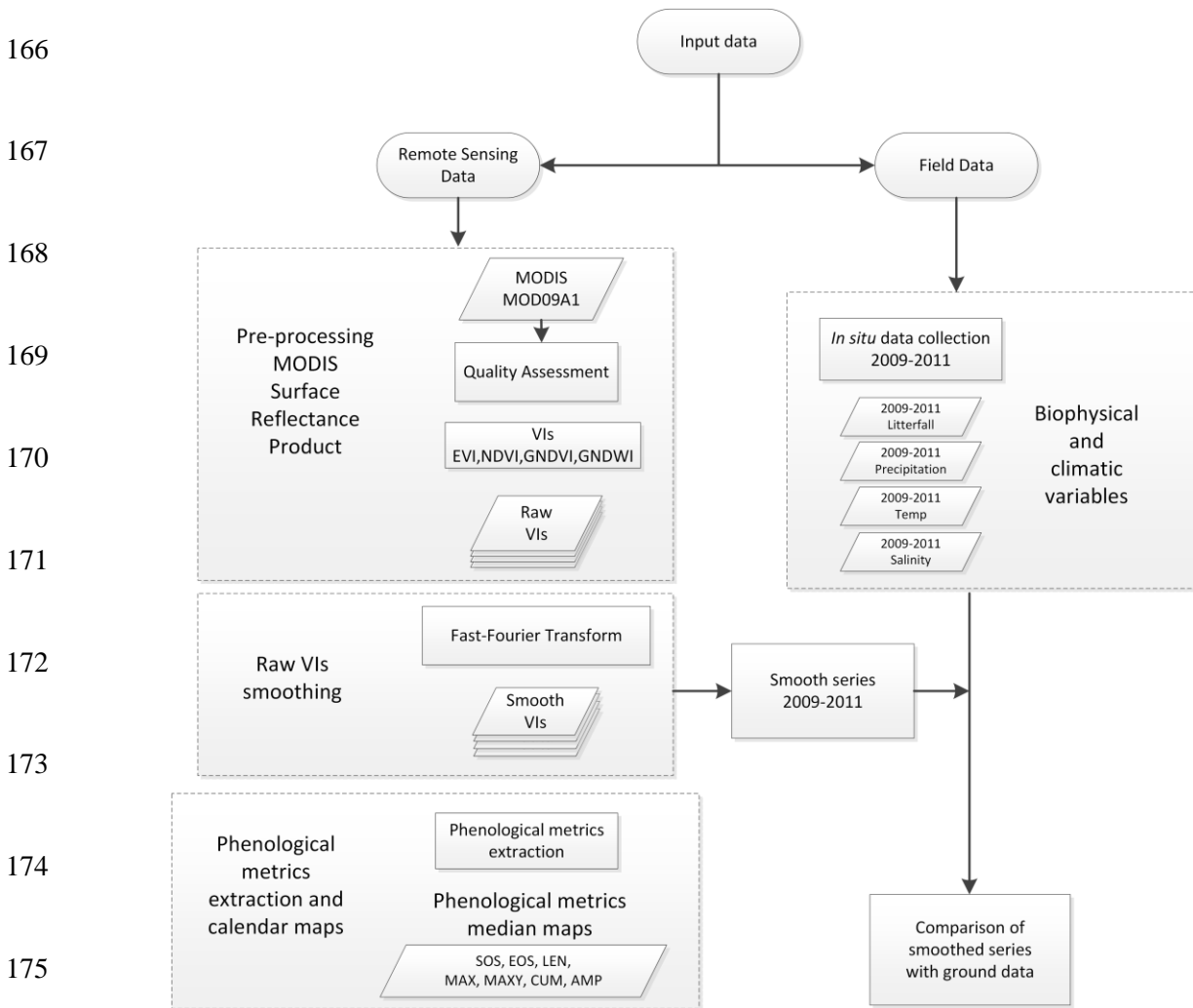
135 **2. Methods**

136 2.1. Study area

137 Yucatan Peninsula is located in SE Mexico (Fig. 1). To the west and north the Yucatan Peninsula is
138 bordered by the Gulf of Mexico and to the east it is bordered by the Caribbean Sea. The area
139 comprises the states of Campeche, Yucatan and Quintana Roo. Except for a narrow fringe of dry
140 climate in the north west (see Fig. S1-S3), the climate of the Yucatan Peninsula is predominantly hot
141 and humid with little precipitation all year and a distinct rainy season in summer (Roger Orellana et
142 al., 2009). The region experiences three seasons: a dry season from March to May, a rainy season
143 from June to October and a cold season from November to February (Herrera-Silveira et al., 1999).
144 The mean annual temperature ranges from 26.5 to 25.5 C and mean annual precipitation ranges from
145 600 mm to 1100 mm (Roger Orellana et al., 2009). Mangrove forest in the Yucatan Peninsula is found
146 on a flat karstic substrate that facilitates the infiltration of rainfall resulting in the absence of runoff
147 and the lack of important streams above the surface (Pope et al., 1997). The vertical and horizontal
148 range of the tides is variable across the study area as the tide depends on the morphology of a
149 particular location. For the Yucatan Peninsula the tidal range is estimated to be between 0.06 m to 1.5
150 m (Herrera-Silveira and Morales-Ojeda, 2010). The mangrove forest is separated from the sea by a
151 sand barrier and it extends in a fringe of varying widths parallel to the coast covering an area of
152 approximately 423,751 ha which represents 55% of Mexico's mangrove cover (CONABIO, 2009).
153 Four species of mangrove dominate the landscape: *Rhizophora mangle*, *Laguncularia racemosa*,
154 *Avicennia germinans* and *Conocarpus erectus*. The National Commission for Knowledge and Use of
155 Biodiversity (CONABIO) has a programme for mapping and monitoring mangrove forest based on
156 aerial photography and fine spatial resolution satellite sensor imagery which is updated every five
157 years.

161 2.2. Data processing

162 Four major steps were followed in this research as summarized in Fig. 2: (i) remote sensing data
 163 pre-processing and computation of vegetation indices, (ii) time-series smoothing and estimation of
 164 phenological metrics, (iii) *in situ* data collection and comparison of *in situ* data with mangrove
 165 phenology.



176 Fig. 2. Schematic diagram illustrating the methodology followed in this research.

177 Although there is a phenology product readily available it was not used in this research for several
 178 reasons. The MODIS Land Cover Dynamics (MCD12Q2) also known as the MODIS phenology
 179 product which primarily uses MODIS EVI, offers a global characterisation of vegetation phenology.
 180 However, according to the MODIS team the quality assurance layer of this product is not working as
 181 intended. In addition, the use of a standard product prevents the computation of other vegetation

182 indices which can provide complementary information about mangrove forest biophysical variables
183 such as canopy water thickness and chlorophyll content. Further, the use of a standard product limits
184 the user's control over the algorithms for smoothing the time-series and estimating phenology metrics.
185 An exploratory analysis revealed that for the study area, the MCD12Q2 product seems to capture the
186 start of greening season (SOS) at spurious peaks at the first half of the time-series (Fig. S12-S14).

187 2.2.1. Remote sensing data pre-processing and computation of vegetation indices

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189 To investigate the mangrove phenology this research employed a fifteen-year time-series (2000-2014)
190 of Moderate Resolution Imaging Spectroradiometer (MODIS) land surface reflectance product, 8-day
191 composite MOD09A1 at 500 m spatial resolution, tile h09v06 courtesy of the NASA EOSDIS Land
192 Processes Distributed Active Archive Center (LP DAAC; see <https://lpdaac.usgs.gov/>). The tiles were
193 reprojected using the MODIS Reprojection Tool. In addition, quality assessment (QA) was applied to
194 each pixel using the 32-bit QA band. Only those pixels that met the following criteria were included
195 in the analysis: MODLAND QA produced at ideal quality, the highest data quality for all reflectance
196 bands, atmospheric and adjacency correction performed.

197 Vegetation indices have been demonstrated to be a useful tool in assessing vegetation biophysical
198 variables. Therefore, in this research four vegetation indices were used to investigate mangrove
199 biophysical variables and to track their seasonality. The indices used were: Normalized Difference
200 Vegetation Index (NDVI), Enhanced Vegetation Index (EVI), Green Normalized Vegetation Index
201 (gNDVI) and Normalized Difference Water Index (NDWI). The NDVI is perhaps the most widely
202 employed index in vegetation phenology (Adole et al., 2016; Julien and Sobrino, 2009), which allows
203 comparison with previous studies. Although NDVI is used widely it tends to saturate at high biomass
204 or at high chlorophyll concentration, which is especially likely for mangroves, and it is affected by
205 soil background, atmospheric effects and aerosols. EVI has an improved sensitivity to high biomass
206 while it compensates for soil background and atmospheric effects using the canopy background
207 adjustment ($L=1$), the coefficients of aerosol resistance ($C_1=6$, $C_2=7.5$) and the gain factor ($G=2.5$)
208 (Huete et al., 2002). gNDVI uses the green band of MODIS and has been more directly related to leaf

209 chlorophyll concentration (Gitelson et al., 1996). NDWI is less sensitive than NDVI to atmospheric
210 scattering and is related to vegetation moisture content (Gao, 1996). The indices are computed as
211 follows:

$$212 \quad EVI = G * \frac{NIR-Red}{NIR+C1*Red-C2*Blue+L} \quad \text{Eq. (1)}$$

$$213 \quad NDVI = \frac{NIR-Red}{NIR+Red} \quad \text{Eq. (2)}$$

$$214 \quad gNDVI = \frac{NIR-Green}{NIR+Green} \quad \text{Eq. (3)}$$

$$215 \quad NDWI = \frac{NIR-SWIR}{NIR+SWIR} \quad \text{Eq. (4)}$$

216 For MOD09A1 data, band 1 corresponds to the reflectance in the red region of the spectrum (620-670
217 nm), band 2 represents the near infrared reflectance (871-876 nm), band 3 represents the blue region
218 of the spectrum (459-479 nm) and bands 4 and 5 correspond to the green (545-555 nm) and SWIR
219 (1230-1250) regions, respectively (see documentation at <https://lpdaac.usgs.gov>). In addition, non-
220 mangrove pixels were masked out using the mangrove distribution shapefile generated by the
221 National Commission for Knowledge and Use of Biodiversity (CONABIO, 2013). The dataset was
222 produced using SPOT-5 imagery from 2010 with a spatial resolution of 10 m. Further, only MODIS
223 pixels that had more than 60% mangrove cover were used in the analyses.

224 2.2.2. Time-series smoothing and phenological metrics estimation

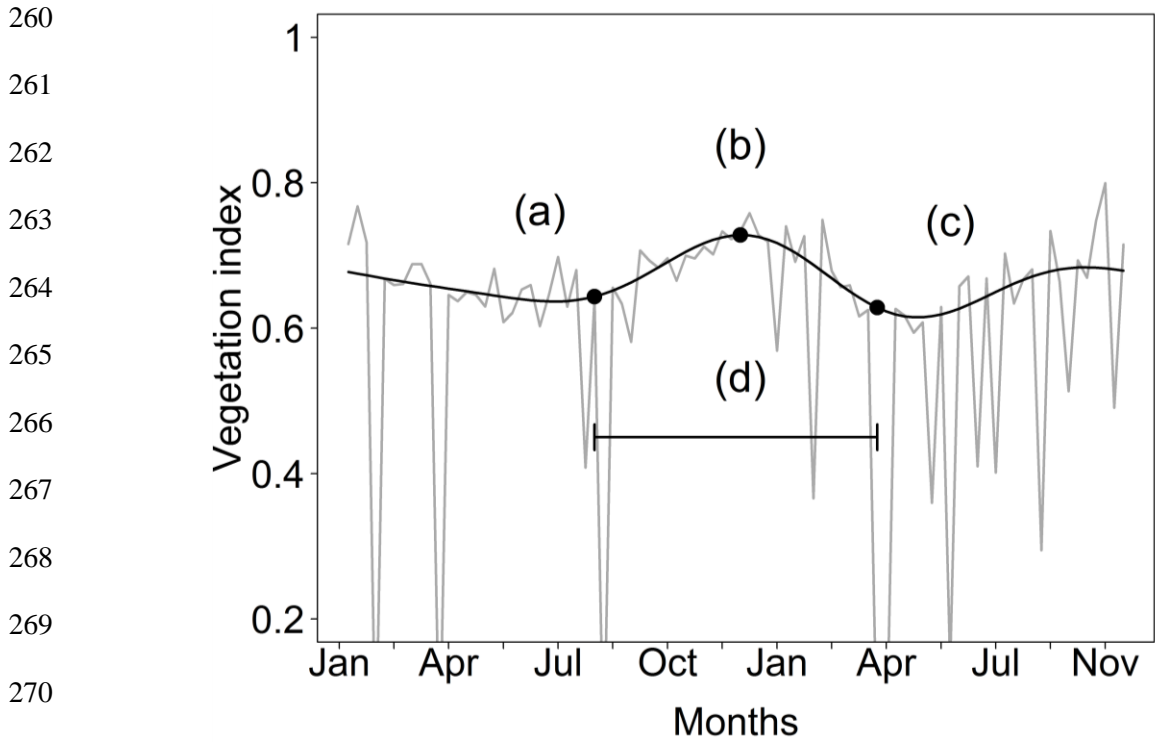
225 *Time-series smoothing*

226 Time-series of remote sensing data contain noise due to atmospheric conditions, aerosols and clouds
227 at the time of data acquisition. Therefore, it is necessary to remove this contamination by using curve
228 fitting methods which smooth the data and estimate the underlying signal prior to estimating
229 phenological metrics. Several smoothing methods are used commonly ranging from relatively simple
230 techniques such as the median filter (Reed et al., 1994) or moving averages to more complex
231 algorithms including the Savitzky-Golay, Asymmetric Gaussian, Double Logistic and Discrete Fourier
232 Transform (DFT) methods (Jonsson and Eklundh, 2002; Zhang et al., 2003; Atkinson et al., 2012).
233 The DFT algorithm was selected to decompose the mangrove signal because it has consistently
234 produced smaller root mean square error (RMSE) between the input and fitted data over a variety of

235 vegetation types including evergreen vegetation (Atkinson et al., 2012). The principle behind DFT is
236 that any complex vegetation growth cycle can be decomposed into a series of sinusoids of different
237 frequencies and a constant (Jakubauskas et al., 2001; Wagenseil and Samimi, 2006; Atkinson et al.,
238 2012). The sinusoids with their frequencies (harmonics) are then summed through a process known as
239 Inverse Fourier Transformation to form a complex smooth wave that resembles the original input
240 profile with the high frequency noise removed. The Inverse Fourier Transform has the advantage of
241 requiring minimum user input (it only needs the user to decide the number of harmonics to reconstruct
242 the profile). This technique has been used widely to smooth time-series of satellite sensor data
243 covering a wide range of environments (Roerink et al., 2000; Jakubauskas et al., 2001; Moody and
244 Johnson, 2001; Dash et al., 2010; Jeganathan et al., 2010a, 2010b; Atkinson et al., 2012; V.
245 Rodriguez-Galiano et al., 2015ab). Typically one to five harmonics are recommended to efficiently
246 reconstruct the natural annual phenological profile (Geerken, 2009). An exploratory analysis revealed
247 that the first four harmonics plus the mean satisfactorily reconstructed the phenological profile (Fig.
248 S11-S14); therefore, four harmonics plus the mean were used in in this research. Prior to
249 reconstructing the time-series with DFT, obvious dropouts from the raw input time-series are
250 removed. Then, the removed dropout is replaced by a moving average which considers valid
251 neighbouring values; in this way the new interpolated value preserves the trend of its temporal
252 neighbours. There is a limit for the number of missing values that are interpolated. If there is more
253 than a month of consecutive missing values the time-series is discarded and no phenological metrics
254 are estimated.

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257 Data processing, DFT based smoothing and phenological metrics estimation based on Dash et al.
258 (2010) and Atkinson et al. (2012) were developed and implemented in the R Programming Language
259 (R Core Team, 2015).



271 Fig. 3. Graphical illustration of phenological metrics estimation using DFT and inflection
272 point algorithms: (a) start of green up season (SOS), (b) time of maximum greenness (Max
273 Green), (c) end of growing season (EOS) and (d) length of growing season (LOS). Grey solid
274 line represents a raw data pixel profile and the black solid line represents the smoothed series using
275 four harmonics plus the mean. The pixel profile was extracted from the season 2010-2011 at the
276 coordinates 20.66601 N, -90.35937 W.

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278 The 15-year time-series was divided into 14 seasons. For each season four phenological metrics were
279 estimated: start of growing season (SOS), end of growing season (EOS), length of season (LOS), time
280 of maximum vegetation greenness (Max Green) (Fig. 3). Several methods are used to estimate
281 phenological metrics from smooth pixel profiles such as threshold based, trend derivative methods
282 and inflection point methods (Reed et al., 1994; Jönsson and Eklundh, 2002, 2004; Dash et al., 2010).
283 In this research, the inflection point method was adopted because it does not assume a pre-defined
284 value as transition date and it is easy to implement. The SOS was defined as the valley point at the
285 start of the increasing trend in the vegetation index values before the first half of the smooth curve,
286 while the EOS was defined as the valley point at the end of the decreasing trend in the vegetation
287 index values after the first half of the curve (Fig. 3). The process consists of extracting the dominant
288 peak (Max Green) in the curve and iterating backwards and forward to find the SOS and EOS (see
289 Dash et al., 2010 for more detail); LOS was defined as the difference between EOS and SOS. Finally,
290 the phenology characterisation maps were created by computing the median from the 14 seasons.

291 2.2.3. *In situ* data collection and comparison of *in situ* data with mangrove phenology

292 *In situ* data collection

293 This research used data produced by the regional programme for characterisation and monitoring of
294 mangrove ecosystems in the Gulf of Mexico and Mexican Caribbean: Yucatan Peninsula (Herrera-
295 Silveira et al., 2014). A field campaign was conducted from January 2009 to October 2011 in 16 sites
296 located along the coast of Yucatan Peninsula within MODIS tile h09v06 (Fig. 1, Table S1). The
297 objective was to characterise, and to establish a baseline for monitoring, the mangrove forest of the
298 region (Herrera-Silveira et al., 2014). The monitoring involved monthly *in situ* measurements of
299 interstitial salinity and litterfall. Rainfall and temperature data were obtained from automatic
300 meteorological stations. In this research, only those sites with meteorological data (Table S1) were
301 used in the analysis because one the main objectives was to examine the link between climatic
302 variables and mangrove forest phenology. The fieldwork was conducted by the Centre of Research
303 and Advanced Studies (CINVESTAV) campus Merida, Yucatan and it was funded by the National
304 Commission for the Knowledge and Use of Biodiversity (CONABIO). Two permanent sampling plots
305 of 10 m by 10 m were established at each site. Five litterfall traps were located at each sampling plot

306 at 1.3 m height from the ground. The vegetative material was collected monthly. It was then
307 dehydrated for 72 hours and litterfall components were separated. Dry litterfall was weighed and
308 represented in grams per square metre per month. Temperature and rainfall data were collected from
309 automatic meteorological stations located at the nearest town. The stations recorded the mean of the
310 meteorological variables every 10 minutes and were expressed as monthly means. Monthly
311 measurements of interstitial water salinity were conducted at each site using a YSI-30 multiprobe
312 sensor (YSI, Xylem Inc., Yellow Springs, Ohio, U.S.A.); the water was sampled at 30 cm depth and
313 expressed in grams per kilogram.

314 *Comparison of in situ data with mangrove phenology*

315 The *in situ* data described above were used for two purposes. First, litterfall was employed to validate
316 the mangrove phenology information derived from satellite sensor data. Second, physical variables
317 (rainfall, temperature and salinity) were used to identify the factors driving mangrove phenology.
318 Spearman's rank correlation analysis was conducted between VIs, litterfall, rainfall, temperature
319 using the monthly mean of all sites to explore the relationships between climatic variables and
320 mangrove greenness and between climatic variables and litterfall. In addition, Spearman's rank
321 correlation analysis was carried out to examine the relationship between cumulative rainfall and
322 phenological metrics SOS, Max Green across sites (eight sites) and years (2009 to 2011) to assess the
323 influence of climatic variables on the main phenological events in the mangrove forest. Spearman's
324 rank correlation was used because normality in the data and a linear relationship could not be assumed
325 in all cases and because building a predictive model was beyond the scope of the analysis. Data
326 analysis was carried out in R (R Core Team, 2015).

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328 **3. Results**

329 3.1. Relationship between litterfall, physical variables and vegetation indices temporal profiles

330 Fig. 4 shows the seasonal variation profiles of monthly litterfall, soil interstitial salinity, temperature,
331 rainfall and vegetation indices at one representative ground sampling station located in the north west
332 of Yucatan Peninsula from January 2009 to October 2011. At this site, during the *in situ* data
333 collection period air temperature increased from March to August with a concomitant increase in

334 salinity. From January to May rainfall remained below 50 mm. Moreover, an increment in rainfall
335 appears to be followed by a decrease in salinity. This seasonal pattern seems to be consistent across
336 all sampling sites (Fig. S4-S10).

337 3.1.1. Litterfall and physical variables

338 Litterfall production is continuous throughout the year in the mangrove forest of Yucatan Peninsula.
339 Minimum values of litterfall were recorded between December and February during the cold season.
340 Two peaks in litterfall were observed, one between April and June and the second between August
341 and October. The first peak in litterfall corresponds to the end of dry season and beginning of rainy
342 season while the second peak corresponds to the late rainy season (Fig. 4a). Both peaks are composed
343 of leaves, stems and reproductive structures, but reproductive structures are more common in the
344 second litterfall peak. A large positive correlation ($r=0.88, p<0.01$) was observed between litterfall
345 and temperature, a moderate significant positive correlation ($r=0.70, p<0.01$) was observed between
346 litterfall and salinity while a weak and not significant correlation ($r=0.25, p>0.1$) was observed
347 between litterfall and rainfall.

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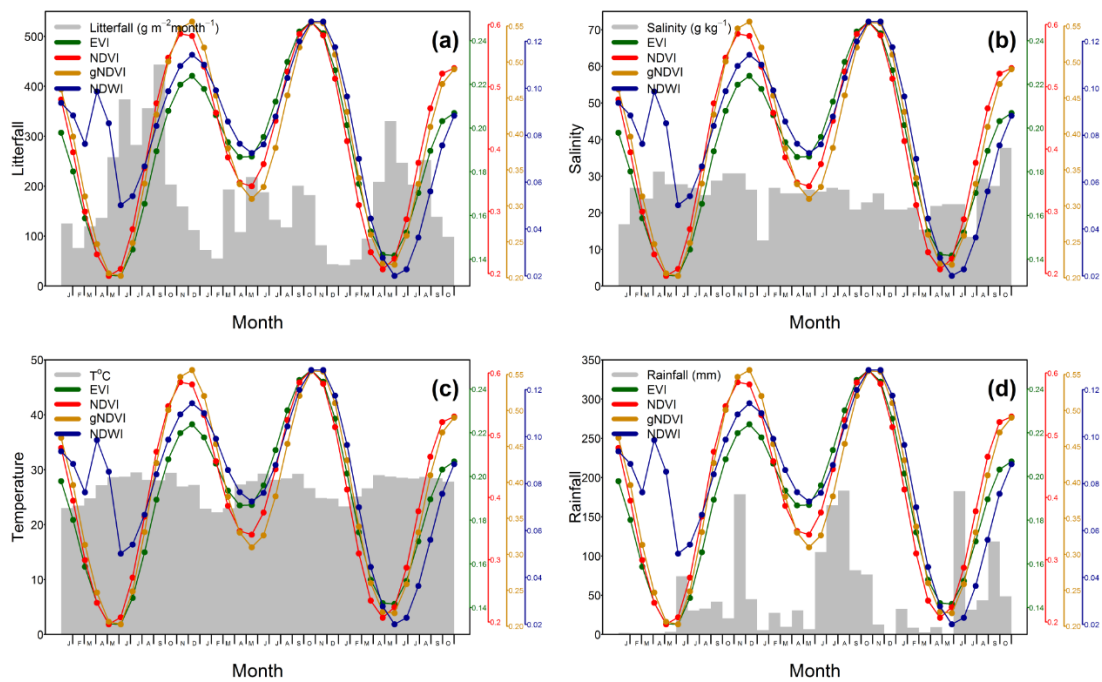


Fig. 4. Seasonal variation in vegetation indices, litterfall (a), salinity (b), temperature (c) and rainfall (d) at the ground station Interna located in the north west of the Yucatan Peninsula between January 2009 and October 2011.

3.1.2. Litterfall and vegetation indices profiles

In Fig. 4, the vegetation indices profiles were averaged monthly to match the temporal resolution of the litterfall data. The temporal pattern of vegetation indices shows that the seasonality of mangrove forest growth is unimodal in nature, it presents a continuous decrease from January to March 2009, reaching a minimum between April and June; then vegetation indices increase steadily from July to September. Mangrove greenness had broadly a negative correlation with litterfall; periods of minimum greenness were coincident with maximum litterfall, and significant negative correlations were observed for gNDVI ($p < 0.05$) (Fig. 5). Further, mangrove greenness appears to lag behind the second peak of litterfall by two to three months. The vegetation indices seasonal pattern described is consistent across all sampling sites (Fig. S4-S10).

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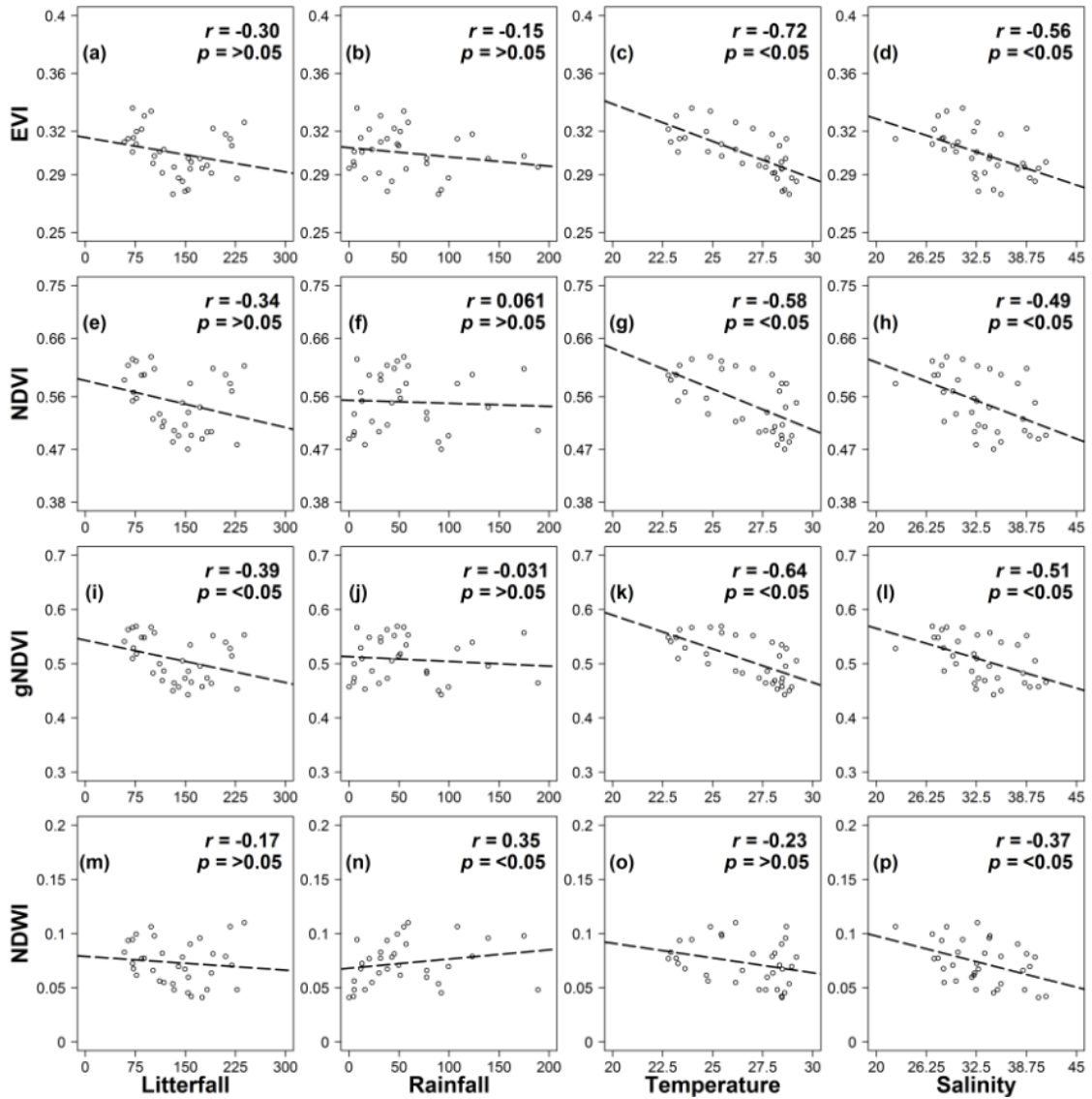
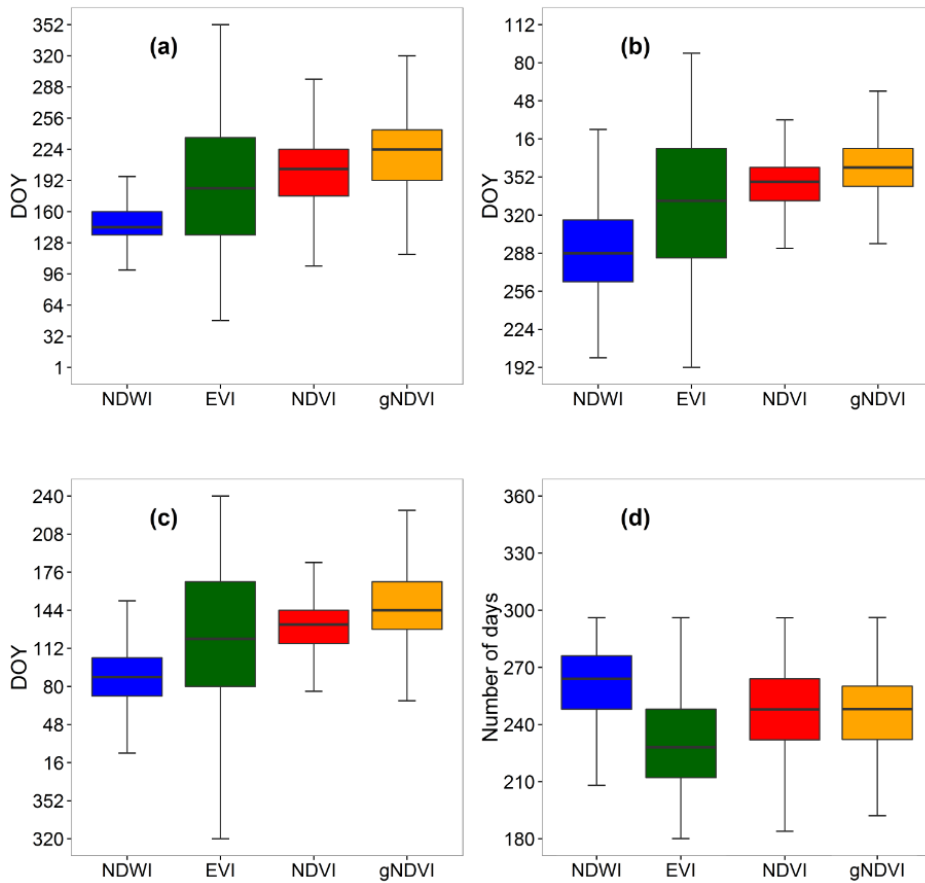


Fig. 5. Scatterplots describing the relationship between monthly vegetation indices values and monthly values of four physical *in situ* variables: litterfall, rainfall, temperature and salinity from January 2009 to October 2011 ($n=34$). Dominant correlations are observed for EVI, gNDVI and NDVI.

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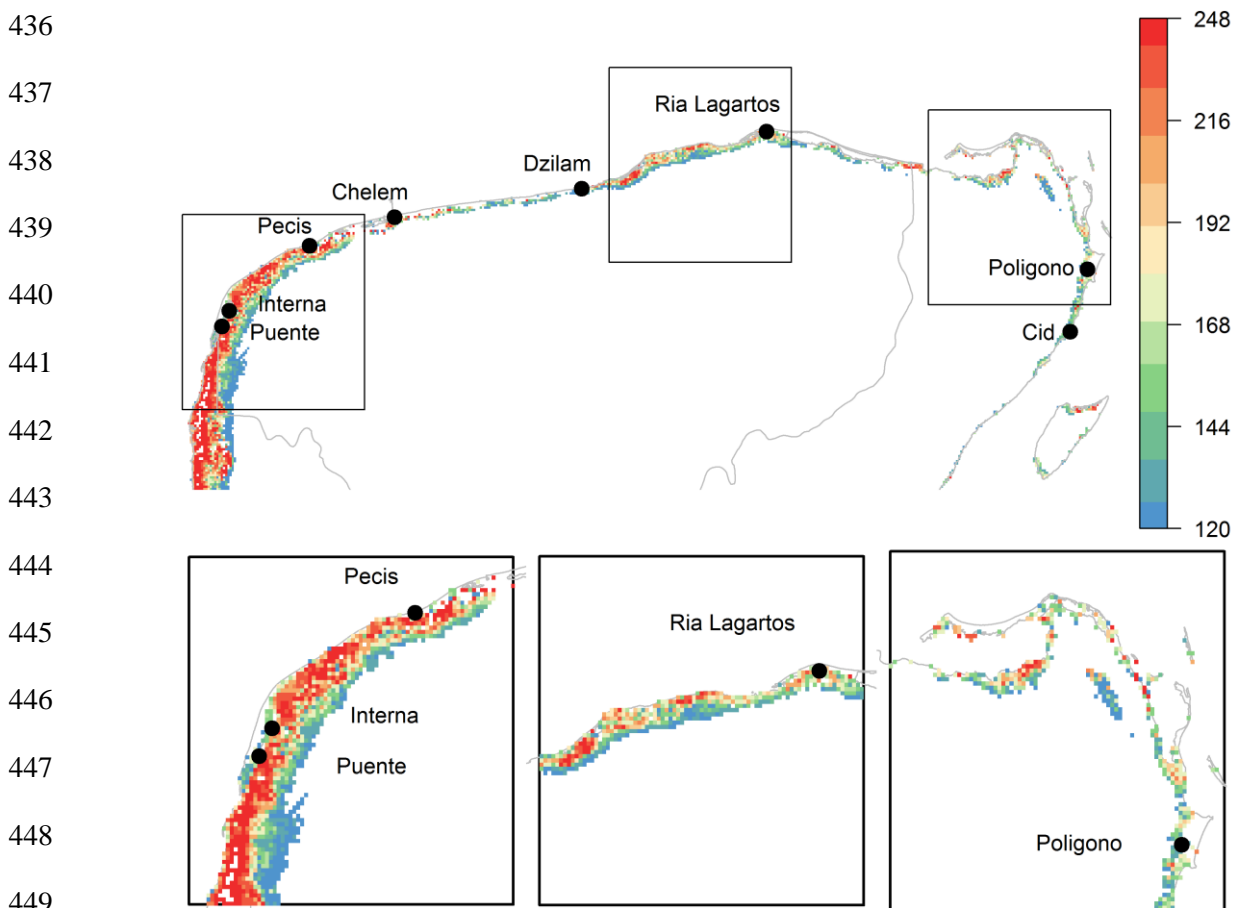


415 Fig. 6. Distribution of phenological metrics for the mangrove forest of Yucatan Peninsula. The green,
 416 red, orange and blue boxes denote EVI, NDVI, gNDVI and NDWI, respectively. The boxplots are
 417 ordered from earlier to later dates (or minimum to maximum, depending on the phenological metric).
 418 The letters on the plots denote the phenological metrics as follows: (a) SOS, (b) Max Green, (c) EOS
 419 and (d) LOS. The middle line of the box represents the median, the lower and upper boundaries of the
 420 box are located at the first and third quartile, respectively, and the bars indicate maximum and
 421 minimum values.

422 In general, the temporal profiles of the four vegetation indices were negatively correlated with the
 423 physical variables (Fig. 5). Temperature and salinity were significant for NDVI, EVI and gNDVI
 424 ($p < 0.05$). Maximum NDVI, EVI and gNDVI seem to lag behind maximum rainfall by two to three
 425 months (Fig. 4; Fig. S4-S10). NDWI and rainfall have a weak, positive, but not significant correlation.

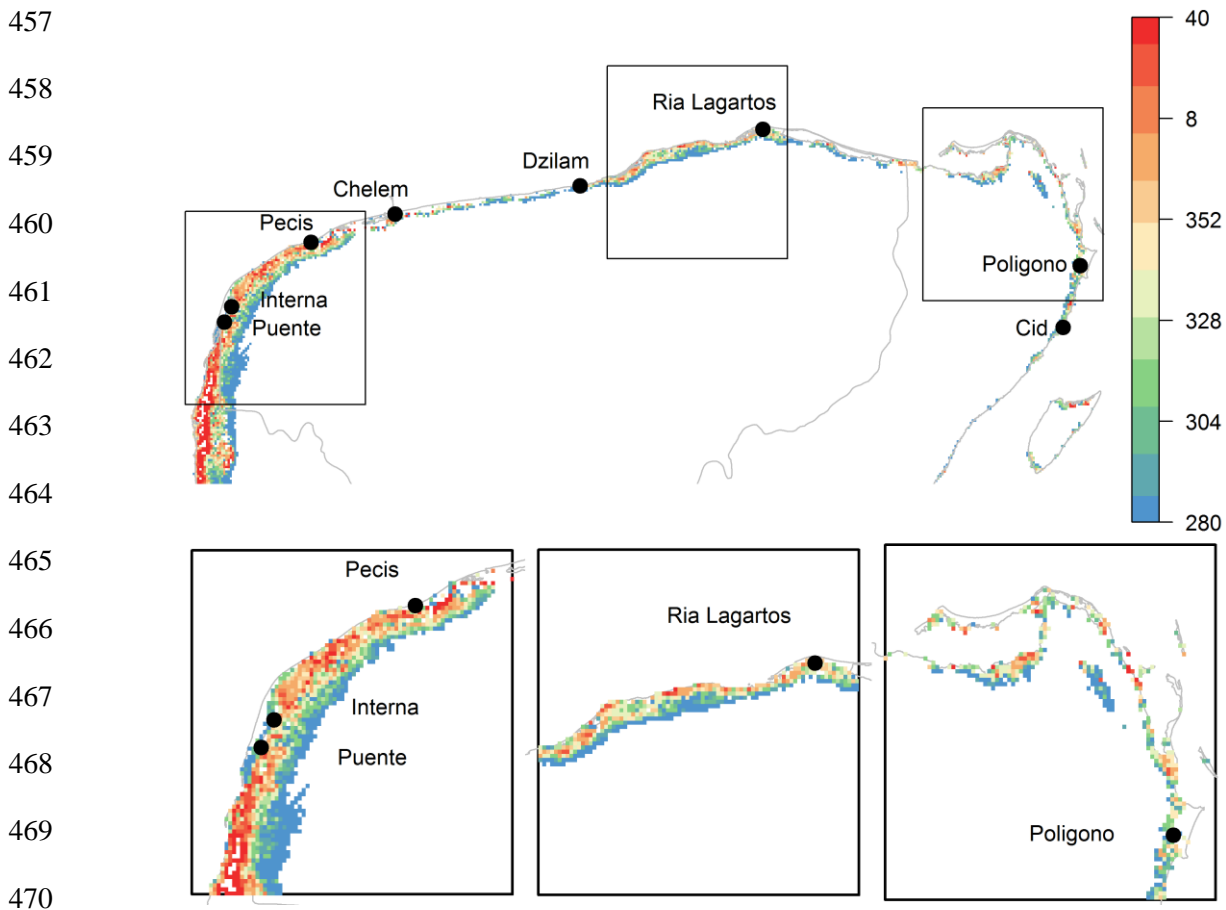
426 3.2. Regional mangrove phenology

427 Fig. 6 illustrates the distribution of the median of the phenological variables derived from the calendar
428 maps. To examine the spatial distribution of the phenological variables, calendar maps of SOS, Max
429 Green, EOS and LOS were created by computing the median from the 14 seasons. For brevity, only
430 the EVI phenological maps are presented in the main paper (Fig. 7-10) and the maps for NDVI,
431 gNDVI and NDWI are presented as supplementary figures (Fig. S15-S26).
432 The dates of phenological variables depend upon the choice of VI. Overall, the timing of SOS, Green
433 Max and EOS is reached first by NDWI, next by EVI, then by NDVI and finally by gNDVI (Fig. 6a-
434 c). SOS occurred at DOY 144 (third week of May), DOY 184 (July), DOY 200 (mid-July) and DOY
435 220 (August) for NDWI, EVI, NDVI and gNDVI, respectively.

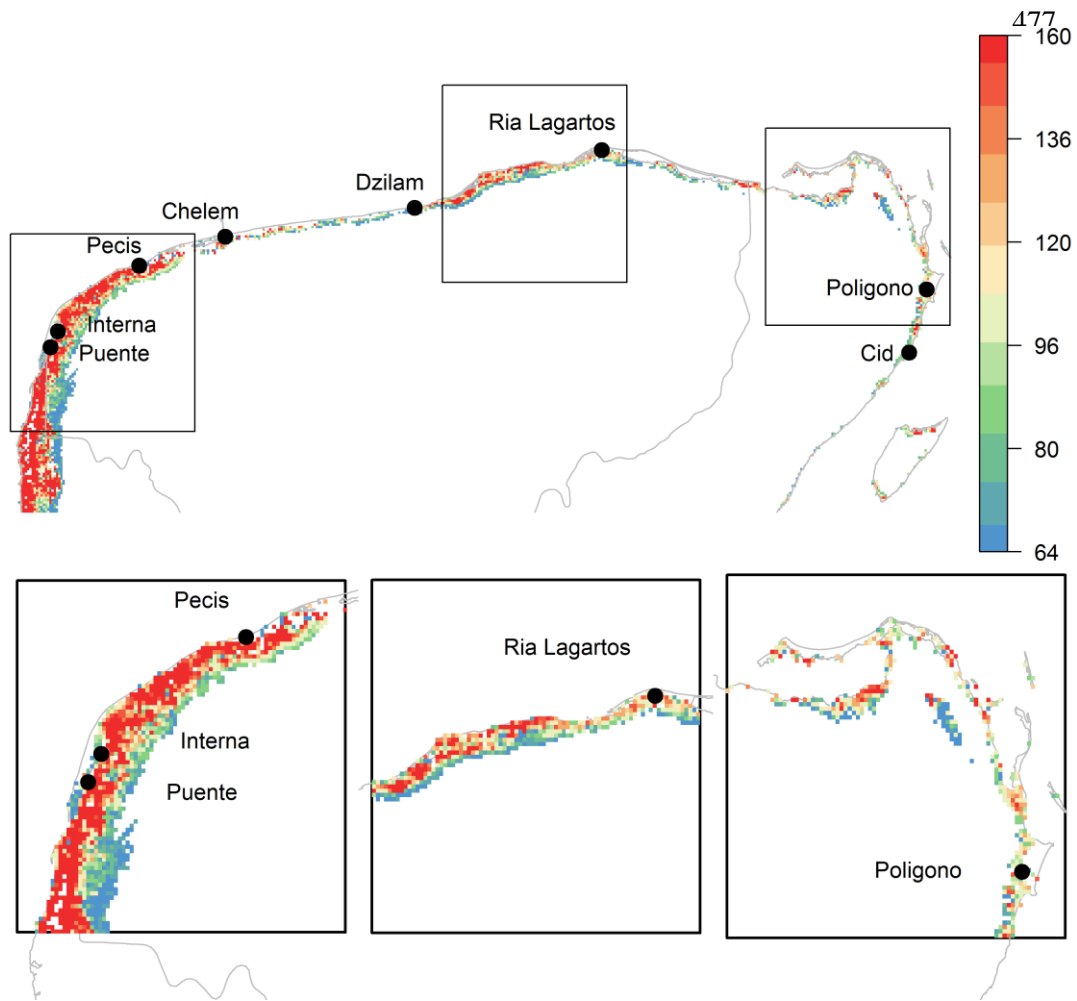


450 Fig. 7. Spatial distribution of EVI SOS in the mangrove forest of Yucatan Peninsula. The
451 map shows the integrated median for the period 2000-2014. The top figure is a map of the
452 entire study area, while at the bottom, from left to right, the figure shows a zoom of the west,

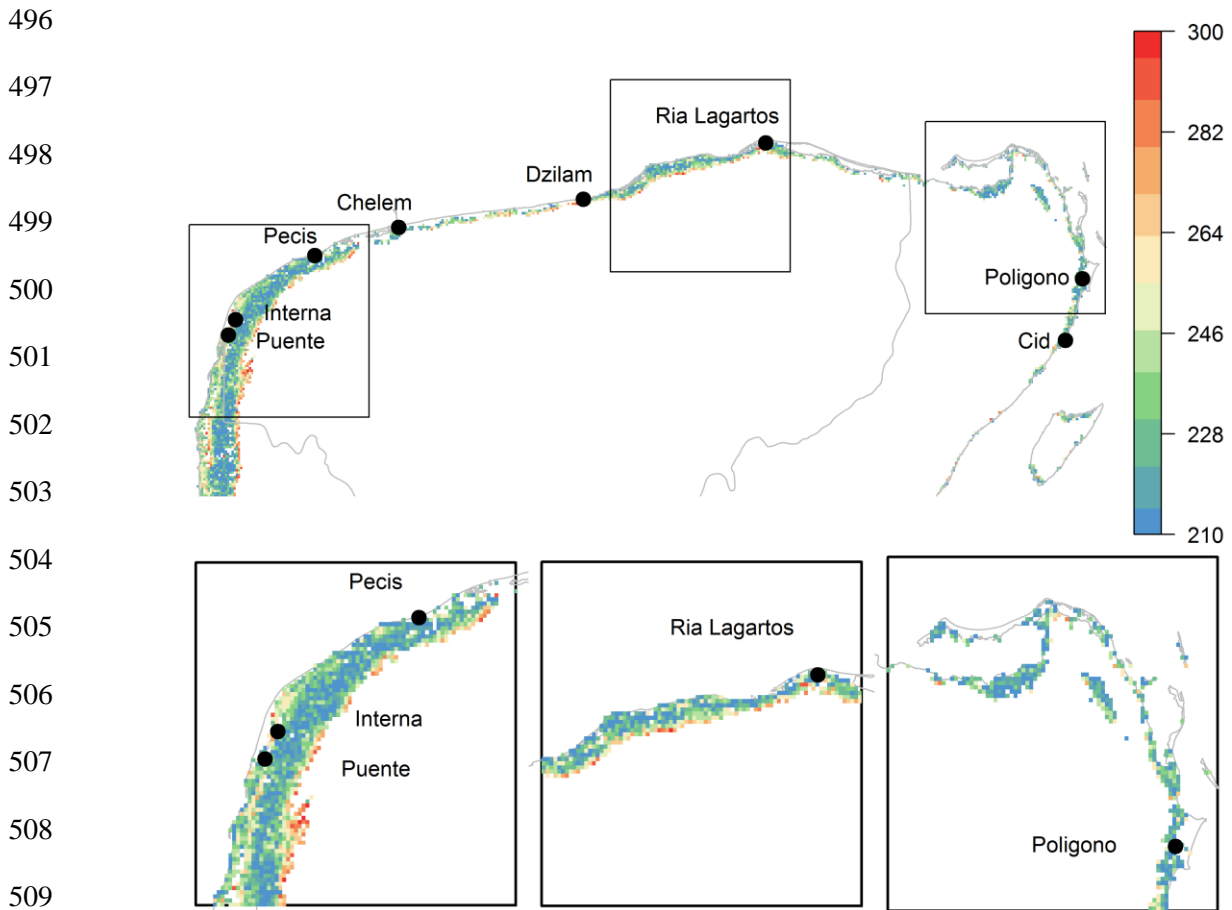
453 north and north-east of the Yucatan Peninsula. The continuous colour ramp on the left
 454 indicates the DOY of SOS, with earlier SOS in blue and later SOS in yellow and red. Non-
 455 mangrove pixels were masked out using the land cover map of the Yucatan Peninsula
 456 (CONABIO, 2013)



471 Fig. 8. Spatial distribution of time of EVI maximum greenness (Max Green) in the mangrove
 472 forest of Yucatan Peninsula. The map shows the integrated median for the period 2000-2014.
 473 The top figure is a map of the entire study area, while at the bottom, from left to right, the
 474 figure shows a zoom of the west, north and north-east of the Yucatan Peninsula. The
 475 continuous colour ramp on the left indicates the DOY of Max Green, with earlier Max Green
 476 in blue and later Max Green in yellow and red.



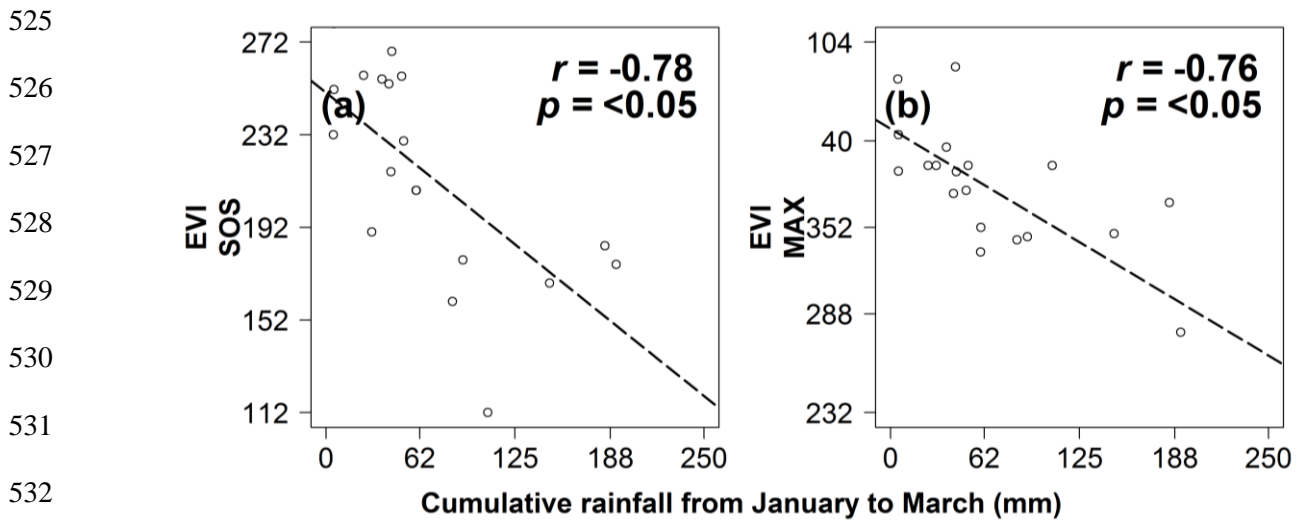
486 Fig. 9. Spatial distribution of time of EVI EOS in the mangrove forest of Yucatan Peninsula.
 487 The map shows the integrated median for the period 2000-2014. The top figure is a map of
 488 the entire study area, while at the bottom, from left to right, the figure shows a zoom of the
 489 west, north and north-east of the Yucatan Peninsula. The continuous colour ramp on the left
 490 indicates the DOY of EOS, with earlier EOS in blue and later EOS in yellow and red.
 491
 492 Spatially, EVI SOS of the mangrove forest of Yucatan Peninsula is reached first in the east of the
 493 peninsula and later in the north and the west (Fig. 7). A notable feature particularly in the north and
 494 west of the peninsula is the earlier SOS dates at the edge of the mangrove forest and later dates at the
 495 interior.



510 Fig. 10. Spatial distribution of time of EVI LOS in the mangrove forest of Yucatan Peninsula. The
 511 map shows the integrated median for the period 2000-2014. The top figure is a map of the entire study
 512 area, while at the bottom, from left to right, the figure shows a zoom of the west, north and north-east
 513 of the Yucatan Peninsula. The continuous colour ramp on the left indicates number of days, shorter
 514 LOS in blue and longer LOS in yellow and red.

515
 516 Max Green was estimated at DOY 288 (mid-October) for NDWI, at DOY 332 (end of November) for
 517 EVI, at DOY 348 (mid-December) for NDVI and at DOY 360 (end of December-beginning of
 518 January) for gNDVI (Fig. 6b). EOS occurred at DOY (104) for NDWI, at DOY (136) for EVI, at
 519 DOY (148) for NDVI and at DOY (160) for gNDVI (Fig. 6c). The spatial distribution of NDVI Max
 520 Green and EOS is generally similar to that of the SOS. That is, delayed Max Green and EOS dates are
 521 coincident with delayed dates of SOS (Fig. 8-9; Fig. S15-S23). The mangrove LOS duration ranged
 522 between 228 to 264 days; shorter for EVI and longer for NDWI (Fig. 6d). The NDVI LOS was

523 homogeneous across the study area; longer EVI LOS durations were observed in the north or north
524 west of the peninsula (Fig. 10).



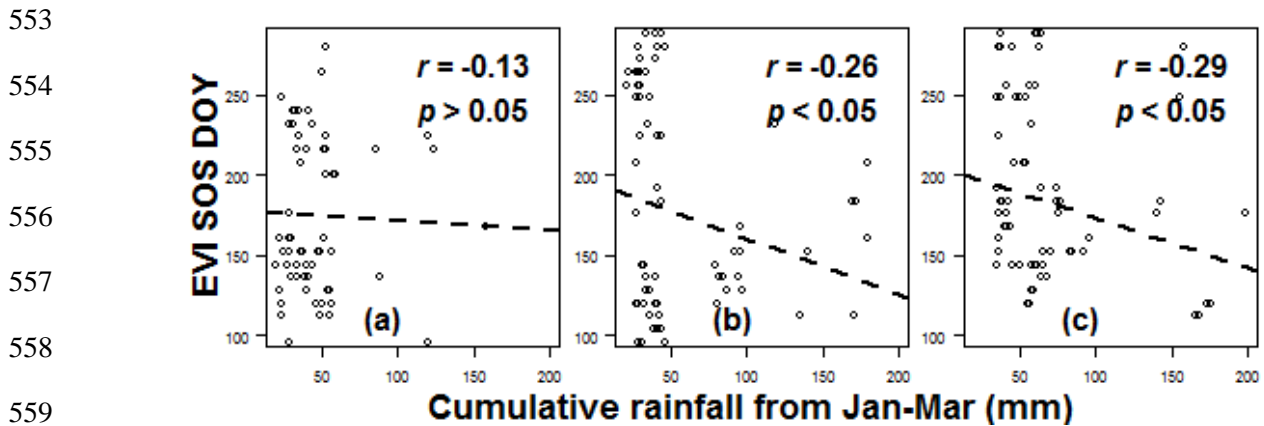
533 Fig. 11. Relationship between cumulative rainfall and mangrove forest phenological
534 variables. The scatterplots describe the relationship between EVI SOS and MAX with the
535 accumulated rainfall (mm) between January and March. The scatterplots suggest that an
536 increase in rainfall during the first three months of the year leads to an earlier maximum
537 greenness.

538

539 3.3. Relationship between satellite-derived phenology and rainfall

540 To investigate the relationships between the mangrove forest phenological variables and
541 physical drivers, correlation analysis was conducted between the timing of phenological
542 variables and the cumulative temperature, rainfall and salinity. Correlations were weak and
543 not significant for temperature and salinity (results not shown); however, cumulative rainfall
544 from January to March produced a large negative correlation ($r > -0.75$, $p < 0.05$) with the
545 timing of EVI SOS and Max Green (Fig. 11) and this pattern was also shown by the other
546 three vegetation indices (Fig. S27). Further analysis was carried out to examine if the effect
547 of rainfall on the phenological variables observed using *in situ* data was consistent using
548 rainfall gridded data. Fig. 12 shows the relationship between EVI SOS and cumulative

549 (January to March) rainfall using CHIRPS gridded data for 2009 to 2011 at 0.05 ° spatial
550 resolution. The figure confirms the negative correlation between the cumulative precipitation
551 and EVI SOS across the Yucatan Peninsula. Overall, this pattern was consistent across
552 seasons (Fig. S28).



560 Fig. 12. Panels a, b and c represent the Spearman correlation between cumulative rainfall
561 from January to March and the timing of EVI start of greening season (SOS) for 2009, 2010
562 and 2010, respectively. Rainfall corresponds to Climate Hazards Group InfraRed
563 Precipitation with Station data (CHIRPS) gridded data at 0.05 °. Correlations were calculated
564 using pixels with mangrove cover greater than 30% for each CHIRPS rainfall pixel.

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575 **4. Discussion**

576 In this paper, 15 years of 8-day MODIS composites were used to characterise the phenology of the
577 mangrove forest of Yucatan Peninsula, SE Mexico. The present study used four vegetation indices,
578 EVI, NDVI, gNDVI and NDWI computed from the MODIS surface reflectance product to derive the
579 SOS, Max Green, EOS and LOS of the mangrove forest. The vegetation indices temporal profiles
580 suggest that although mangrove is an evergreen forest it exhibits seasonal variation in greenness
581 detectable using moderate spatial resolution (500 m) remotely sensed data. *In situ* data on litterfall,
582 temperature, precipitation and salinity were used as possible explanatory drivers of the mangrove
583 forest phenology. To our knowledge, this is the first attempt to provide a regional scale
584 characterisation of the main phenological events of a mangrove forest and assess its relationship with
585 physical drivers and litterfall.

586 Mangrove litterfall is commonly used as an indicator of forest productivity. Attempts to understand
587 the drivers of mangrove forest litterfall dynamics have tried to establish correlations with physical
588 variables such as wind speed, temperature, rainfall, evaporation and hours of sunshine, but given the
589 disparity in results it is difficult to generalize conclusions. In our study, temperature and salinity were
590 linearly correlated with litterfall. The dynamics of litterfall in the Yucatan Peninsula was in agreement
591 with reports from the Gulf of Mexico (Gill and Tomlinson, 1971; Day et al., 1987; Day Jr. et al.,
592 1996; Utrera-López and Moreno-Casasola, 2008) and previous records in the study area (Zaldivar-
593 Jimenez et al., 2004). These studies have reported continuous litterfall throughout the year with leaf
594 litter the main component, low litterfall values during the cold season, a peak in litter at the end of the
595 dry season and most of the litterfall occurring in the rainy season with a higher proportion of
596 reproductive structures. The pattern described responds to environmental adaptations. On the one
597 hand, during periods of water stress characterised by increases in evapotranspiration and soil salinity
598 due to higher temperatures or decreases in soil moisture due to lack of precipitation and low tides,
599 mangrove trees shed leaves, potentially with a decrease in leaf cover (Medina, 1999). Leaf shedding
600 in mangroves has been widely cited as an adaptation to saline environments (Hasegawa et al., 2000),

601 more specifically as a mechanism to eliminate salt from the tissue (Tomlinson, 1995; Zheng et al.,
602 1999). Further, the physiological stress caused by high pool water salinity has been related to
603 reductions in photosynthetic radiation-use-efficiency (Barr et al., 2013; Song et al., 2011). On the
604 other hand, during the wet season litterfall contains a higher proportion of flowers, fruits and
605 seedlings. Fresh water input from rainfall leads to higher water levels and milder salinity, and these
606 conditions favour the dispersal of seedlings (Lopez-Portillo and Ezcurra, 1985).

607 In other tropical evergreen and deciduous forests canopy seasonality is regulated by rainfall
608 seasonality (Chave et al., 2010; Zhang et al., 2014) but the response of vegetation is not immediate.
609 Rainfall increases soil water and atmospheric humidity reducing water stress in the plants. Similarly,
610 in this research, mangroves did not respond immediately to rainfall, the mangrove greenness was
611 weak and not significantly correlated with rainfall (Fig. 5), and mangrove greenness generally lagged
612 behind rainfall (Fig. 4).

613 Regarding the phenological variables, the results revealed differences between VIs. One obvious
614 contrast was the earlier dates of SOS, EOS and Max Green for NDWI compared to the rest of the
615 vegetation indices (see Fig. 6). This asynchrony in phenological metrics may be explained by the
616 sensitivity of the indices to different features of vegetation growth that may not necessarily occur
617 simultaneously. The advanced SOS for NDWI observed in this study could be due to an increase in
618 soil moisture before the development of new leaves as NDWI is not only sensitive to vegetation water
619 content (Gao, 1996; Chen et al., 2005) but is also sensitive to soil moisture (Gao, 1996; Xiao et al.,
620 2005a). Increases in NDWI proportional to soil moisture were observed in paddy fields before leaf on
621 and proportional to leaf water content during leaf development (Xiao et al., 2002; De Alwis et al.,
622 2007; Tornos et al., 2015). In addition, delayed phenological variables for NDWI were observed in
623 the northwest of the Yucatan Peninsula, an area characterised by an arid and semi-arid climate (Fig.
624 S1-S3). The ability of NDWI to vary proportionally according to soil moisture and canopy water
625 stress makes it suitable to monitor mangrove forest water stress.

626 It is evident that EVI presents greater variability in the phenological metrics (Fig. 6). This variability
627 may respond to the attributes of this vegetation index. EVI is a vegetation index primarily sensitive to
628 canopy structural properties such as LAI, canopy cover and leaf structure (Gao et al., 2000).

629 Mangrove forests are heterogeneous landscapes composed of several species and individuals of
630 different age which may not respond uniformly to the environmental drivers. Therefore, the
631 mangrove's canopy structural properties vary spatially within a mangrove stand, and this structural
632 diversity is reflected in a greater variation in EVI phenological metrics. Another contrast was the
633 delayed SOS, Max Green and EOS dates for gNDVI. This delay could respond to mangrove leaf
634 development processes. Previous work suggests that gNDVI is more sensitive to mangrove leaf
635 chlorophyll concentration (Pastor-Guzman et al., 2015), a photosynthetic pigment that increases with
636 leaf age during the leaf development stage (Wang and Lin, 1999). The time to fully develop leaf
637 anatomy and photosynthetic pigments depends on the species and the time of the year that leaves are
638 born. Soto (1988), reported that new leaves of *A. germinans* reached their maximum size at between 4
639 to 6 months, whereas leaves from the genus *Rhizophora* reached their maximum mass, area and
640 chlorophyll concentration in 3 to 4 months (Wang and Lin, 1999; Mehlig, 2001; Sharma et al., 2014).
641 Therefore, it could be suggested that the delay in gNDVI phenological variables could be due to its
642 sensitivity to canopy chlorophyll content.

643 With respect to the drivers of mangrove forest phenology, our results revealed that sites receiving a
644 greater amount of rainfall between January and March experience earlier SOS and Max Green (Fig.
645 11 and Fig. S27-S28). The geology of the Yucatan Peninsula prevents the formation of rivers due to
646 the karstic permeable soil. Thus, fresh water contributions from runoff and rivers are practically non-
647 existent (Perry et al., 1995). The mangrove forest in the region receives fresh water inputs mainly
648 from rainfall and ground water discharges via springs when the aquifer recharge is greatest. This fresh
649 water is characterized by high inorganic nitrogen content, silicates and low particulate matter
650 (Herrera-Silveira, 1994; Herrera-Silveira et al., 1999). Given that milder salinity and greater
651 availability of nutrients leads to more favourable conditions for mangrove growth, it is reasonable to
652 speculate that cumulative rainfall plays an important role in mangrove phenology.

653 The results obtained in this research serve as a reference for future studies addressing long-term
654 phenological changes in the mangrove forest of the Yucatan Peninsula. Nevertheless, there are some
655 limitations that need to be taken into account. For example, the study focused on mangrove as a single
656 cover type; however, in reality the mangrove forest is not a homogeneous landscape but a complex

657 mosaic, a fragmented landscape that in many cases includes individuals of different ages, heights,
658 open areas, and gaps with water ponds. Although the four mangrove species are dominant, in some
659 areas there may be other communities embedded in the mangrove matrix such as grassland (Batllori-
660 Sampedro and Febles-Patrón, 2007). Therefore, at 500 m spatial resolution the phenological pattern
661 could be masked. Additional analyses showed that the mangrove forest phenology is clearly
662 distinguished from that of terrestrial vegetation but it resembles that of surrounding flooded land
663 cover types (Fig. S36-S37). In order to control for this source of uncertainty in this research we used
664 only pixels that had more than 60% mangrove cover. Further examination revealed that the difference
665 in the timing of SOS is marginal with a mangrove cover between 60% and 80% per MODIS pixel. At
666 this coverage, it is assumed that the mangrove forest dominates the reflectance signal, supporting the
667 use of this threshold. However, as the percentage of mangrove increases above 80% the SOS median
668 shifts towards later DOY for about one month, suggesting pure mangrove stands may have delayed
669 response to the environmental drivers (Fig. S29-S35). Another potential source of uncertainty can be
670 the mismatch in the temporal and spatial resolution between the *in situ* data and the MODIS data. *In*
671 *situ* data were collected monthly while MODIS data were 8-day composites; further, the sampling
672 plots are not be evenly distributed within the MODIS pixel footprint. These sources of error could be
673 minimized by adopting a sampling method designed to validate remote sensing observations
674 (Elmendorf et al., 2016). Further, it should also be noted that other factors exist that may affect the
675 seasonal dynamics of the vegetation indices on the mangrove forest such as clouds and aerosols. To
676 reduce this source of potential errors, the QA MODIS layer was used to include only pixels with the
677 highest quality.

678 **5. Conclusion**

679 This paper based on the Yucatan Peninsula is the first phenological characterisation of a mangrove
680 forest using remote sensing data. The study used 15-year time-series of four vegetation indices
681 computed from MODIS surface reflectance and the phenology was compared with climatic variables,
682 salinity and litterfall. The DFT algorithm was used to smooth the time-series and four phenological
683 variables were estimated: SOS, EOS, Max Green and LOS. The results revealed clear seasonality in
684 mangrove forest greenness. Periods of lower greenness were typically associated with the dry season
685 while periods of maximum greenness were associated with the months following maximum rainfall.
686 The timing of the phenological variables differs depending on the vegetation index employed. In
687 general, NDWI showed advanced phenological parameters whereas gNDVI showed delayed dates.
688 SOS ranged between DOY 144 (late dry season) and DOY 220 (rainy season) while the EOS occurred
689 between DOY 104 (mid-dry season) to DOY 160 (early rainy season). The length of the growing
690 season ranged between 228 and 264 days. Interestingly, sites receiving a greater amount of rainfall
691 between January and March have an advanced SOS and Max Green. These results show the potential
692 of MODIS to monitor mangrove phenology at 500 m spatial resolution. MODIS constitutes a cost-
693 effective tool to monitor temporal variation in mangroves as data are freely available at fine temporal
694 resolution. The phenological calendar maps obtained are up-to-date and represent a reference for
695 future research and the length of the time-series processed gives robustness to the phenological
696 parameters. It is acknowledged, however, that the MODIS phenology product could be used in future
697 studies if the tools or expertise required for the analysis are not available, or if the time available for
698 fitting is limited. The results have implications for understanding mangrove forest dynamics at the
699 landscape scale, and provide the potential to monitor biophysical variables such as water stress and
700 canopy chlorophyll and their link to climatic variables at the global scale; information that could be
701 used as input to bio-geochemical models. Finally, this study highlights the need to continue the long-
702 term *in situ* data collection network in the mangrove forest.

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706

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713

714 **7. References**

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