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1	Developing seagrass index for long term monitoring of Zostera japonica
2	seagrass bed: a case study in Yellow River Delta, China
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# Developing seagrass index for long term monitoring of *Zostera japonica* seagrass bed: a case study in Yellow River Delta, China

## 20 Abstract

Seagrass beds offer unique and vital ecological services as an important blue carbon ecosystem 21 22 in coastal wetlands. Zostera japonica is an intertidal seagrass species native to eastern Asia and is one of the most widely distributed seagrass species in China. However, little is known on the 23 long-term variations of Z. japonica extents. Automatic mapping method for Z. japonica 24 25 seagrass beds is in urgent need to fill this knowledge gap. In this study, we proposed a new SeaGrass Index (SGI) for automatic and rapid mapping of Z. japonica based on time-series 26 27 Landsat satellite imagery, aiming to alleviate the influence of tidal inundation and enhance the 28 separability from other coastal cover types. The SGI considers both spectral and phenological characteristics of Z. japonica, as well as the spatial location of Z. japonica. We took the Yellow 29 River Delta (YRD), China as our study area, where Z. japonica was first discovered and 30 reported in 2015. Based on SGI, Z. japonica extents during 1985-2018 were extracted using 31 32 multi-Otsu thresholding algorithm. Accuracy assessments based on field investigations and 33 high-resolution imagery showed that SGI has successfully separated seagrass beds from other 34 cover types, especially intertidal salt marshes, with overall accuracies >95%, producer's 35 accuracies >90% and user's accuracies >94%. Our study provides the first long-term maps of seagrass beds in YRD. The area of Z. japonica showed large variations during 1985-2018, 36 ranging from 149 ha in 2005-2006 to 1302.9 ha in 2011-2012. The spatial distribution of Z. 37

*japonica* varied with the morphological change of the estuary caused by river channel shifts.
Since 2011, *Z. japonica* seagrass beds have undergone area degradation due to the invasion of *S. alterniflora*. The area was only 332.3 ha in 2017-2018. Coastal erosion and extreme climate
events such as drought and typhoon might also explain degradation of seagrass beds in YRD.
We expect that the SGI will advance automatic and rapid mapping methods for intertidal
seagrass beds, and the *Z. japonica* maps will provide a baseline data for restoration and

45 Keywords: Seagrass mapping; Zostera japonica; Time-series analysis; Multi-Otsu algorithm;
46 Yellow River Delta

47

#### 48 1 Introduction

49 Seagrasses are marine plants found in shallow coastal and estuarine waters in many parts of the world from the equator to high latitude except Antarctica. Seagrass beds are one of the most 50 51 valuable marine and coastal ecosystems in the world (Programme, 2020). They form important 52 nursery habitats and support coastal food webs for thousands of marine species (Iacarella et al., 2018). In addition, they provide significant services including sediment stabilization, coastal 53 erosion reduction, water purification and nutrient cycling (Programme, 2020; Green et al., 54 55 2021). Meanwhile, seagrass beds are important blue carbon ecosystems. Although seagrass 56 beds cover less than 0.2% of the total ocean area, they store approximately 10% to 20% of the global marine carbon (Fourqurean et al., 2012; Campbell et al., 2022). They also bring valuable 57 socio-economic benefits by providing food and livelihoods for coastal communities. 58

Unfortunately, seagrass beds are one of the most threatened marine ecosystems. Researches 59 60 have shown that 29% of the known seagrass beds have disappeared since first recorded in 1879, and the global seagrass beds are being lost at a rate of 1.5% per year (Green et al., 2021). Global 61 62 climate change, including the elevated seawater temperature and the increasingly extreme 63 weather events such as hurricanes and ocean heat waves, has caused serious damage to seagrass 64 beds (Yue et al., 2021). Human activities such as beach aquaculture, coastal development and water pollution also pose threats to seagrass beds, resulting in habitat fragmentation and 65 degradation, presenting negative feedback to coastal ecosystem and climate change (Adams et 66 al., 2016; Salinas et al., 2020). 67

68 Zostera japonica (also called dwarf eelgrass or Japanese eelgrass) is a seagrass species native to eastern Asia from Vietnam to Russia. It is one of the most widely distributed seagrass 69 70 species in China's coastal areas and can be found primarily in intertidal zones. The Yellow River 71 Delta (YRD) is the youngest and broadest coastal wetland ecosystem in China (Li et al., 2019). In 2015, researchers discovered a large Z. japonica meadow in the intertidal zone of YRD 72 73 during field investigations, which was the first record of the seagrass distribution in YRD (Zhou et al., 2016). However, to date little is known about the spatial and temporal dynamics of Z. 74 japonica extents in YRD during the past decades. In recent years, YRD has experienced 75 76 dramatic morphological and landscape changes due to river channel shifts, coastline variations 77 and rapid expansion of invasive plant species Spartina Alterniflora (Li et al., 2019; Wang et al., 2021; Li et al., 2021). In this highly dynamic area, Z. japonica mapping and monitoring over 78 79 the past decades are vital for better understating their responses to climate change and human 80 activities, and for sustainable management and conservation of the coastal wetlands.

81 Compared to in-situ field investigations and acoustic-based techniques, satellite remote 82 sensing techniques provide unique opportunities for seagrass monitoring over large area in an 83 efficient and cost-effective manner (Hossain et al., 2015a; Programme, 2020; Veettil et al., 2020; 84 Li et al., 2021). As reviewed by Hossain et al. (2015) and Veettil et al., (2020), seagrass mapping 85 approaches should consider habitat conditions including water depth (exposed intertidal, submerged intertidal, shallow subtidal or deep subtidal) and water clarity (clear or turbid). For 86 87 submerged seagrasses in subtidal area (e.g., water depth > 2 m), water depth corrections are usually required to reduce the effects of water attenuation and retrieve bottom reflectance, based 88 on which seagrasses are identified (Kuhwald et al., 2021; Tu et al., 2021). For intertidal 89 90 seagrasses which are exposed during low tide, surface reflectance-based vegetation indices, such as Normalized Differential Vegetation Index (NDVI) have been adopted to delineate 91 92 seagrass extents (Valle et al., 2015; Zoffoli et al., 2020; Zoffoli et al., 2021). 93 At present, multispectral satellite data with spatial resolutions from hundreds of meters to

sub-meter including MODIS, Landsat series, Sentinel-2, SPOT series, QuickBird, WorldView-2/3 and PlanetScope have been used for seagrass mapping in different coastal areas (Hossain et al., 2015b; Veettil et al., 2020). For intertidal seagrasses such as *Zostera noltei* or *Zostera marina*, some researchers have investigated monitoring long-term variations of seagrass extents based on satellite imagery acquired on multiple years (Calleja et al., 2017; Zoffoli et al., 2020; Zoffoli et al., 2021). These studies generally selected single-date cloud-clear image acquired at low tide and identified seagrasses using supervised classification (Coffer et al., 2020; Kuhwald 101 et al., 2021; Carpenter et al., 2022; Lebrasse et al., 2022) or unsupervised classification (Barrell 102 and Grant, 2015; St-Pierre and Gagnon, 2020; Xu et al., 2021). For example, Calleja et al. (2017) 103 selected 10 cloudless Landsat images with lowest tidal level for each summer during 1984-104 2015 and developed rule-based thresholds to identify Zostera noltei in Bay of Santander, Spain. 105 Xu et al. (2021) selected low-tide Landsat images from 1974 to 2019 and used ISODATA clustering algorithm to map Zostera marina seagrasses in Caofeidian, Bohai Sea, China. 106 107 Fernandes et al. (2022) selected 21 cloud-free and low-turbidity Landsat images acquired 108 during summer from 1988 to 2018 and utilized Support Vector Machine to separate seagrasses 109 from sand in Adelaide, South Australia.

110 However, uncertainties exist when using single-date images for long-term mapping of 111 seagrasses. First, in coastal areas with frequent cloud coverage and tidal inundation, it is 112 difficult to find an adequate image acquired in the growing season that is neither contaminated 113 by cloud nor influenced by tide level. Second, when seagrasses and other coastal vegetation 114 (e.g., intertidal salt marshes such as S. alterniflora in YRD) are mixed, it is difficult to 115 discriminate them because they may have similar spectral features during growing seasons. 116 When it comes to the YRD, additional challenge exists due to the high turbidity of coastal water. 117 The Yellow River estuary is one of the most turbid estuaries in the world, with high 118 concentration of suspended particulate matter in the coastal water (Li et al., 2019). The 119 sediments may cover the seagrasses even during low tide and make it more difficult for 120 discrimination. Furthermore, supervised classifications that have been used for seagrass 121 mapping or coastal land cover classification in many studies are challenging for long-term timeseries seagrass mapping due to lack of field data for training sample collection for the retrospective analysis.

To address these issues, this study aims to: (1) present a SeaGrass Index (SGI) that is capable of discriminating *Z. japonica* from other coastal cover types and alleviating the influence of frequent cloud cover and tidal inundation, (2) develop an unsupervised classification approach for automatic extraction of *Z. japonica* in YRD based on SGI. Both SGI and the classification approach are to be developed on Google Earth Engine (GEE) platform to facilitate rapid mapping of *Z. japonica* extent, and (3) investigate long-term spatial and temporal variations of *Z. japonica* distribution during 1985-2018 in YRD.

131 2 Study area and data

#### 132 **2.1 Study area**

133 The YRD is located in Shandong Province, China, adjacent to the Bohai Sea in the north 134 and Laizhou Bay in the east. It belongs to temperate continental monsoon climate, with an average annual temperature of 11.5-12.9 °C and average annual precipitation of 592.2 mm. 135 136 Precipitation is mostly concentrated in July and August. The Yellow River estuary has weak tides, with an average tidal range of 0.73-1.77 m, and is dominated by irregular semi-diurnal 137 138 tide pattern (Yang et al., 2011). Our study area is located in the Yellow River Delta National 139 Natural Reserve with an area of 1530 km<sup>2</sup> (Fig. 1). Z. japonica meadows are distributed in the 140 lower intertidal area which are submerged at high tide and exposed to air for 4-8 hours at low 141 tide. The growing season is from June to August (Liu et al., 2019; Zhang et al., 2019). The 142 native salt marsh vegetations include S. alterniflora, Saueda salsa, Tamarix chinensis, and

143 Phragmites australis. S. alterniflora is an invasive species in YRD. It was first found in the 144 intertidal zones in 2008 and since then it has expanded rapidly (Wang et al., 2021). In recent 145 years, it was reported that S. alterniflora started to encroached ecological niches of Z. japonica 146 (Ren et al., 2019; Ma et al., 2020). S. salsa grows on both mid-high intertidal area and low 147 intertidal area near the coastline. T. chinensis and P. australis grow in high-tide area. The study 148 area is featured considerable sediment deposition because Yellow River carries an average 1.0 149 billion tons of sediment per year. The river channel in the delta has shifted many times in history. 150 During the past fifty years, it was artificially shifted from the Diaokou course to the 151 Qingshuigou course in 1976 (Fig. 1a), and then to Q8 in 1996 (Fig. 1b). In 2007, the downstream end of the Q8 course shifted naturally because of the change in riverine dynamics 152 153 (Fan et al., 2018; Li et al., 2019). As a results, the shape of the coastline in the estuary has 154 changed dramatically during the past a few decades. Based on the locations of river course 155 channels during the study period, we divided the study area into three zones, namely Zone A, 156 B and C (Fig. 1). Zone A covers the north bank of YRD after the river course shifted in 2007. 157 Zone C covers the south bank of YRD before the river course shifted in 1996. Zone B is located between Zone A and Zone C. 158



159

160 Fig. 1. The location of the study area. (a) - (c) are Landsat imagery in standard false color

- 161 acquired on August 27, 1993; August 5, 2003; August 10, 2016, respectively. (d) illustrates the
- spatial locations of field photos. Field photos in (1) (5) are Z. japonica, S. alterniflora, S. salsa,
- 163 tidal flat, and water, respectively.
- 164 **2.2 Datasets**
- 165 **2.2.1 Landsat imagery and pre-processing**

166 We collected all available Landsat 5 TM, Landsat 7 ETM+ and Landsat 8 OLI surface

167 reflectance (SR) images (Tire 1 Level 2 products) covering YRD (Path 121, Row 43) during

168	1985-2018 on the Google Earth Engine platform, with a total number of 801 images. The SR
169	products were generated by atmospheric correction using Landsat Ecosystem Disturbance
170	Adaptive Processing System (LEDAPS) and Landsat Surface Reflectance Code (LaSRC)
171	algorithms. For each image, the quality assurance (QA) band was used to mask out the bad-
172	quality pixels covered by cloud, cirrus, shadow, and snow. We grouped the time-series
173	observations during 34 years into 16 periods to ensure each period has enough valid
174	observations in each season (Fig. 2). In general, observations during every two-year period were
175	grouped except for 1993-1998. Because the river tail channel shifted in 1996, we grouped
176	observations before and after tail channel shift in to two periods, i.e., from 1993 to 1995 and
177	from 1996 to 1998, respectively. Fig. 2 shows the available Landsat images by sensors range
178	from 24 to 75, and the average number of valid observations ranges from 10 to 40 in the 16
179	periods (Fig. 2a). Most pixels have at least one observation during the summer (June to August)
180	and autumn (September to November), and the autumn witnessed slightly more valid
181	observations than the summer (Fig. 2 b-d).





Fig. 2 The number of Landsat images (a) by sensors (Landsat TM, Landsat ETM+, Landsat OLI), (b) by seasons (spring from March to May, summer from June to August, autumn from September to November, and winter from December to February), and percentage of pixels with various numbers of good-quality observations in (c) summer and (d) autumn on YRD in each period during 1985-2018.

# 187 **2.2.2 Field investigations and high spatial resolution imagery**

188 In August 2016, July 2017, and June 2018, we conducted several field investigations in 189 the seagrass bed habitats in the study area. During each survey, we recorded the locations of Z. 190 japonica patches using GPS RTK units. High spatial resolution imagery including Gaofen-1 191 (GF-1, 2 m), Gaofen-2 (GF-2, 1 m) and Sentinel-2 (10 m) images were used to examine the appearance of Z. japonica and other salt marsh species on the image. Visual analysis of these 192 images helped us locate Z. japonica on the images in other years when field surveys were not 193 carried out. In addition, we collected WordView-2 (0.5 m), ZY-3 (6 m), SPOT-6 (1.5 m), and 194 195 GF-1/2 high spatial resolution imagery during June-October from 2012 to 2018 (Table 1). Because of the scarcity of high-resolution data before 2012, we relied on Landsat satellite 196 197 images for visual interpretation. These images, together with the field investigation records

were used to create reference samples in order to analyze the phenological dynamics of typical
vegetation species in the study area (see Section 3.2) and to validate the seagrass mapping
results (Section 3.5).

201 Table 1. High spatial resolution imagery used for validation sample collection from 2012 to

202 2018.

Year	Date	Satellite sensors	Spatial resolution (m)
2018	June 5 to October 27	GF-1; GF-2	2; 1
2017	June 9 to November 4	GF-1; GF-2	2; 1
2016	June 22 to October 14	GF-1; GF-2	2; 1
2015	May 25 to October 27	GF-1; GF-2	2; 1
2014	June 6 to October 28	GF-1	2
2013	June 14 to October 17	GF-1; SPOT 6	2; 1.5
2012	May 27 to September 20	WorldView-2; ZY-3	0.5; 6

203

#### 204 **3** Methods

205 Fig. 3 illustrates the workflow of Z. japonica seagrass beds mapping. First, potential distribution area of Z. japonica in YRD was delineated based on time-series Landsat images 206 207 during each period (section 3.1). Second, SGI that considers both spectral and phenological 208 characteristics of Z. japonica was developed by analyzing time-series spectral indices (section 209 3.2). Third, separability analysis was conducted to evaluate the reliability of SGI (Section 3.3). 210 Fourth, SGI images were generated within the seagrass potential area for each period and an 211 unsupervised classification method was developed for seagrass identification (section 3.4); 212 Finally, the seagrass mapping results were validated and seagrass maps during 1985-2018 were 213 generated for YRD (Section 3.5).



# **3.1 Identify potential distribution area of seagrass beds**

According to our field surveys and previous reports (Zhou et al., 2016; Zhang et al., 2019), *Z. japonica* grows in the low intertidal zone of YRD within 500 m away from the coastline. In order to narrow down the image processing extent and reduce the influences of land cover types in the high tidal zone, we created 4 km buffer area around the coastline (2 km inside and outside the coastline) as the potential distribution area of *Z. japonica*.

We delineated coastline for each period based on the minimum composite image of Modified Normalized Difference Water Index (mNDWI) (hereafter  $mNDWI_{min}$ ). mNDWI (Eq. 1) is sensitive to open surface water bodies, and it can identify subtle differences between turbid and clear water (Xu, 2005). At present, mNDWI is one of the most widely used indices for coastline extraction.

$$mNDWI = \frac{\rho_{green} - \rho_{SWIR1}}{\rho_{green} + \rho_{SWIR1}} \tag{1}$$

where  $\rho_{green}$  is the reflectance of the green band (Band 2 for Landsat TM/ETM+ and Band 3 228 for Landsat OLI), and  $\rho_{SWIR1}$  is the reflectance of the short-wave infrared 1 (Band 5 for Landsat 229 230 TM/ETM+ and Band 6 for Landsat OLI). First, mNDWI was calculated from each Landsat image. Based on all mNDWI images during each period, we generated an mNDWI<sub>min</sub> image, 231 where the value at each pixel represents the lowest mNDWI during the period. The mNDWI<sub>min</sub> 232 233 image depicts the driest condition of every pixel. Then, Otsu algorithm was used to separate land from water on mNDWImin image and extract the coastline. The Otsu algorithm is a 234 nonparametric approach for image thresholding. The optimal threshold is determined by 235 traversing all potential thresholds using the image's histogram and selecting the one with the 236

minimal inter-class variation (Otsu, 1979). As *mNDWI<sub>min</sub>* generally represents the image at
the lowest tide, the delineated coastline and the 4 km buffer area can effectively remove the
features at mid to high tide area. As a results, the potential distribution area of *Z. japonica* was
identified for every periods. By field surveys and visual interpretation of the reference images,
land cover categories in the area consists of include *Z. japonica*, *S. alterniflora* (only after 2008), *S.salsa*, tidal flat and water.

# 243 **3.2 Construct seagrass index (SGI)**

# 244 **3.2.1** Analysis of time-series spectral indices

To enhance the separability between Z. japonica seagrass and other cover types (S. 245 alterniflora, S.salsa, tidal flat and water), time series analysis of three spectral indices including 246 247 NDVI, mNDWI and Tasseled Cap Brightness Index (TCBI) was conducted. NDVI has been 248 widely used to represent growing status of vegetation, with NDVI reaching peak during 249 growing season and declining during senescence season. mNDWI was selected as it is sensitive to water features mixed with vegetation (Singh et al., 2015). TCBI was derived from Tasseled 250 Cap Transformation, which incorporates six different bands of Landsat TM/ETM+/OLI 251 imagery (Kauth, 1976). The TCBI has proven to be suitable for soil moisture estimation and 252 253 inundation detection (Ludwig et al., 2019).

# Fig. 4 illustrates time-series NDVI, mNDWI and TCBI of pure *Z. japonica*, *S. alterniflora*, *S.salsa*, tidal flat and water pixels calculated based on valid observations during 2015-2016.

256 We merged all observations during 2015-2016 sorted by the day of year (DOY). Fig. 4a shows

257 that the NDVI of Z. japonica depicts water-like characteristics during late September-May

258 (NDVI<0) and strong vegetation features during June-August. It climbs from June to August, 259 with NDVI peak occurring in August, and then rapidly declines to negative values in the end of 260 September. We determined the DOY from 150 to 250 as the green period (Gr-P) and DOY from 261 260 to 330 as the senescence period (Se-P) of Z. japonica. The penology pattern of Z. japonica 262 observed from the NDVI time series is consistent with previous in-situ studies (Zhang et al., 2019; Zhang et al., 2021). During Gr-P, Z. japonica demonstrates similar NDVI as S. 263 alterniflora. During Se-P, Z. japonica demonstrates significantly lower NDVI than S. 264 265 alterniflora and S. salsa. Note that S.salsa shows low NDVI throughout the year because of its short plant height and low coverage, with peak value of around 0.3 in September. The mNDWI 266 267 and TCBI time series do not show much temporal variations throughout the year (Fig. 4b and 268 4c). Z. japonica has high mNDWI values. S. alterniflora and S. salsa has much lower mNDWI 269 (mostly negative) than other types, and Z. *japonica* pixel generally has the lowest TCBI values. 270 S. salsa had the highest TCBI. Due to the turbidity of the water in the YRD, the TCBI of the 271 water is higher than that of Z. japonica.





Fig. 4. The temporal profile of (a) NDVI (b) mNDWI (c) TCBI of *Z. japonica*, *S. alterniflora*, *S. salsa*, tidal flat and water pixel during 2015-2016. The maximum NDVI value was marked
as ×; and the corresponding mNDWI and TCBI values on the date when NDVI reaches the
maximum were also marked as ×.

276

Fig. 4 shows that the combination of NDVI, mNDWI and TCBI can help discriminate *Z*. *japonica* from others at the pixel level. The maximum NDVI (*NDVI<sub>max</sub>*) during each 2 or 3year period (marked as × in Fig. 4a) represents the greenest state of seagrass beds and thus reduces the tidal influence and possible cloud contamination in a single image. However, using

281 the NDVI<sub>max</sub> alone cannot discriminate Z. japonica from S. alterniflora as they have similar 282 NDVI<sub>max</sub> values (Fig. 4a). Note that Z. japonica has shorter green period than the intertidal salt marshes, and NDVI shows more rapid and higher magnitude of declination during senescence 283 284 period. Therefore, we consider to use  $NDVI_{max}$  to differentiate Z. japonica from non-285 vegetation types and use the difference between  $NDVI_{max}$  and the mean NDVI during 286 senescence period (SeNDVImean) to differentiate Z. japonica and other salt marshes. When 287 NDVI of S. alterniflora reaches the maximum, its mNDWI reaches the lowest (mNDWI = -288 0.4). This is because S. alterniflora has dense and tall plants, therefore, depicts terrestrial vegetation spectral characteristics during growing peak. 289

290 For each 2 or 3-year period, we generated the  $NDVI_{max}$  composite image using function 291 "imageCollection.qualityMosaic" on GEE platform. Each band of the NDVImax image 292 represents the band values for each pixel when NDVI reaches the maximum. Based on this 293 image, mNDWI ( $mNDWI_{vimax}$ ) and TCBI ( $TCBI_{vimax}$ ) images were derived (illustrated as × 294 in Fig. 4b and 4c at pixel level). We also generated SeNDVI<sub>mean</sub> image composite and derived  $NDVI_{max} - SeNDVI_{mean}$  layer. In order to examine the variation in  $NDVI_{max}$ ,  $mNDWI_{vimax}$ , 295 296  $TCBI_{vimax}$ , and  $NDVI_{max} - SeNDVI_{mean}$  indices of the land cover types, we generated 1000 297 reference sample points for each land cover type over the 16 periods (around 63 samples for 298 each period for each class) using field survey data and visual interpretation of reference images. 299 As shown in Fig. 5, NDVI<sub>max</sub> of Z. japonica seagrass beds demonstrates large variations, 300 ranging from 0.21 to 0.89. The histogram of  $NDVI_{max}$  of Z. japonica overlaps with that of S. 301 alterniflora in the right tail and that of S. salsa in the left tail (Fig. 5a). mNDWIvimax can 302 effectively distinguish Z. japonica from S. alterniflora because almost all Z. japonica sample pixels have positive mNDWIvimax and S. alterniflora sample pixels have negative 303 mNDWIvimax. However, mNDWIvimax cannot separate Z. japonica from S. salsa (Fig 304 305 5b). Note that S. salsa grows on both mid-high intertidal area and low intertidal area, for S. salsa on tidal flats near the water, mNDWIvimax is greater than 0 due to tidal influence. Fig. 5c 306 307 shows that the TCBI<sub>vimax</sub> of Z. japonica is considerably lower than that of S. salsa, while its histogram has obvious overlap with water and tidal flat. The  $NDVI_{max} - SeNDVI_{mean}$  of Z. 308 309 japonica is greater than other land cover types (Fig. 5d), while its histogram has small overlap 310 with S. alterniflora.



Fig. 5 Histograms and box plots of (a) NDVI<sub>max</sub>, (b) mNDWI<sub>vimax</sub>, (c) TCBI<sub>vimax</sub>, (d) NDVI<sub>max</sub>SeNDVI<sub>mean</sub> for *Z. japonica*, *S. alterniflora*, *S. salsa*, tidal flat and water.

314

# 315 **3.2.2** Formulation of SGI based on phenological-spectral characteristics

Based on the above analysis, we can utilize  $mNDWI_{vimax} > 0$  to remove *S. alterniflora* from the image. Compared to tidal flat and water, *Z. japonica* has much higher  $NDVI_{max}$ . Compared to *S. salsa*, it has higher  $NDVI_{max} - SeNDVI_{mean}$  and lower  $TCBI_{vimax}$ . To enhance the contrast, we formulated SGI as Eq. (4). We first assigned *NoData* value to the pixels with  $mNDWI_{vimax} \le 0$  as all *Z. japonica* pixels have positive  $mNDWI_{vimax}$  values. 321 For other pixels, we sum  $NDVI_{max}$  and  $NDVI_{max} - SeNDVI_{mean}$  in the numerator to 322 highlight the spectral and phenological characteristics of *Z. japonica* and use  $TCBI_{vimax}$  in the 323 denominator to differentiate with *S. salsa*.

$$SGI = \begin{cases} NoData & (mNDWI_{vimax} \le 0) \\ \frac{2NDVI_{max} - SeNDVI_{mean}}{TCBI_{vimax}} & (mNDWI_{vimax} > 0) \end{cases}$$
(4)

Fig. 6 shows that the histogram of SGI of *Z. japonica*, water, *S. salsa* and tidal flats sample pixels show three peaks. *Z. japonica* has obviously higher SGI than *S. salsa*, tidal flat and water. Water pixels have relatively lower SGI than other types, while *S. salsa* and tidal flats have similar SGIs. Overall, the SGI histogram of *Z. japonica* is visually more separable from others compared to any of  $NDVI_{max}$ ,  $NDVI_{max} - SeNDVI_{mean}$  or  $TCBI_{vimax}$  (Fig. 5).



329

330 Fig. 6 Histogram and box plot of SGI index for Z. japonica, S. salsa, tidal flat and water.

331

# 332 **3.3 Separability analysis**

333 Separability analysis is important to examine how well a target class can be discriminated
334 from others and can be used as indicators of the classification performances (Xu et al., 2021).
335 In this study, we utilized separability index (SI) presented by Somers and Asner (Somers et al.,

336 2010) to assess and compare the separability between *Z. japonica* and others in terms of 337  $NDVI_{max}$ ,  $mNDWI_{vimax}$ ,  $NDVI_{max} - SeNDVI_{mean}$ ,  $TCBI_{vimax}$  and SGI. SI incorporates 338 both intra-class and inter-class variabilities. The formula is as follows:

339 
$$SI_{zo} = \frac{\Delta inter_{zo}}{\Delta intra_{zo}} = \frac{|\overline{\mu_z} - \overline{\mu_o}|}{1.96 \times (\sigma_z + \sigma_o)}$$
(5)

where  $SI_{zo}$  denotes the separability index between *Z. japonica* and other categories.  $\overline{\mu_z}$  and  $\overline{\mu_o}$  denote the mean index values of *Z. japonica* and the other class, respectively.  $\sigma_z$  and  $\sigma_o$ denote the standard deviation of the index values.  $|\overline{\mu_z} - \overline{\mu_o}|$  denotes the inter-class variance between *Z. japonica* and the other class, and  $(\sigma_z + \sigma_o)$  denotes the intraclass variance. Higher SI values indicates greater disparities between *Z. japonica* and others and smaller within-class variances.

#### 346 3.4 Identify seagrass bed extent

347 We adopted multi-Otsu thresholding algorithm (Liao et al., 2001), an unsupervised 348 clustering approach, to extract Z. japonica automatically based on SGI image without training 349 process. The conventional Otsu algorithm automatically select one optimal threshold and 350 partitions the dataset into two classes based on the histogram of the dataset. It has been 351 successfully used for image binarization and target detection in remote sensing. Unlikely, multi-Otsu thresholding algorithm calculates multiple thresholds with the number of categories 352 353 defined by user (>2). From Fig. 6, we suppose that two thresholds can be used to separate Z. japonica, water and tidal flat/S. salsa based on SGI image. In this case, multi-Otsu algorithm 354 355 aims to find two thresholds which maximize the inter-class variance of SGI.

356 
$$\sigma_t^2(t_1, t_2) = \sum_{i=1}^3 P_i (\mu_i - \mu_T)^2$$
(6)

357 
$$(T_1, T_2) = \arg \max_{0 < T_1 < T_2} \{ \sigma^2(t_1, t_2) \}$$
(7)

358 
$$P_i = \frac{\sum pixels in class i}{Total pixels}$$
(8)

where  $\sigma_t^2(t_1, t_2)$  denotes the inter-class variance at threshold  $(t_1, t_2)$ . The algorithm goes 359 through all possible thresholds to find the optimal thresholds  $(T_1, T_2)$  which maximize 360  $\sigma_t^2(t_1, t_2)$ .  $P_i$  represents the probability of a pixel belonging to class *i*,  $\mu_i$  represents the mean 361 362 SGI value of class *i*, and  $\mu_T$  represents the mean value of the whole SGI histogram. We used 363 the multi-Otsu algorithm to segment SGI images into three classes. According to Fig. 5, the 364 class with SGI values greater than  $T_2$  is identified as Z. japonica seagrass bed. Finally, a total of 16 seagrass maps were generated by re-classifying the results in each period into two 365 categories: Z. japonica and non- Z. japonica. 366

#### 367 **3.5 Accuracy assessment**

368 Considering the availability of high spatial resolution imagery, we validated all Z. japonica 369 seagrass maps during 2011-2018, and selected two Z. japonica seagrass maps before 2000 370 (1993-1995) and after 2000 (2007-2008) for validation. First, we employed stratified random 371 sampling strategy to select validation samples. From the seagrass maps during each period, 150 and 300 pixels were randomly selected for Z. japonica and non-Z. japonica, respectively. The 372 sample pixels were visually interpreted based on the field investigations during 2016-2018 373 374 combined with high spatial resolution imagery acquired between 2012 to 2018 (Table 1). Before 375 2010, the sample pixels were interpretated based on the time-series Landsat images. These 376 validation pixels should have consistent class (Z. japonica or non- Z. japonica) during the two or three-year period. In rare cases that the class of a sample pixel has changed from Z. japonica 377

to non- *Z. japonica* or vice versa, the sample pixel was excluded and a pixel with unchanged
class around it was added.

380 4 Results

381 4.1 Separability analysis and SGI maps

Fig. 7 shows the SI values of each index including mNDWIvimax, NDVImax, TCBIvimax, 382  $NDVI_{max} - SeNDVI_{mean}$  and SGI for seagrasses between different classes based on the 1000 383 reference samples. It is obvious that  $mNDWI_{vimax}$  has much greater SI value (1.68) between 384 385 Z. japonica and S. alterniflora than other indices, indicating that S. alterniflora can be well separated from Z. japonica by mNDWI<sub>vimax</sub>. SGI shows much greater SI values than other 386 indices in separating Z. japonica from S. salsa, tidal flat and water (SI value = 1.42, 1.41 and 387 388 2.17 respectively). We also calculated the SI value of the 5 indices between Z. japonica and 389 non-Z. japonica. Similarly, the SI value of SGI between Z. japonica and non-Z. japonica 390 (SI=1.41) was higher than any other indices (0.81 for NDVI<sub>max</sub>; 0.42 for TCBI<sub>vimax</sub> and 0.68 391 for  $NDVI_{max} - SeNDVI_{mean}$ ;). The highest SI value of SGI between Z. japonica and non-Z. 392 japonica indicates that SGI generates good inter-class separability and low intra-class variability. 393



394

Fig.7. SI values for each index between different classes. Class acronyms are: ZJ- *Z. japonica*,
SA- *S. alterniflora*, SS- *S. salsa*, TF- Tidal flat, WT- water, Non ZJ- non ZJ

397 Fig. 8 presents some exemplar SGI maps within the potential distribution area (left column), as well as corresponding histograms (center column) and the extracted Z. japonica 398 399 map (right column). The SGI maps show substantially higher SGI values at the lower intertidal 400 area around the north and east coastline of YRD. Two peaks in the low range of SGI (<3) and 401 one flatter peak in the high range of the SGI  $(3 \sim 10)$  are observed in the histograms, which is 402 quite similar as Fig. 6. The three-class Otsu algorithm identified two thresholds, 0.73 and 3.21 403 for 2013-2014 (Fig. 8a); -0.09 and 4.07 for 2003-2004 (Fig. 8b); 0.46 and 3.16 for 1993-1995 404 (Fig. 8c), and the pixels with SGI greater than 3.21, 4.07, 3.16 were classified as Z. japonica 405 Fig. 9 illustrates that the SGI thresholds for Z. japonica seagrass beds, respectively. 406 identification are relatively stable, varying from 2.22 to 4.11 with an average threshold of 3.22. 407 The SGI thresholds for the images before 2001 (2.22~3.16) were slightly lower than those after 408 2001 (3.18~4.11). Examinations of the images showed that the seagrass beds demonstrated 409 lower  $NDVI_{max}$  in those years than other years, which might be explained by lower biomass or

410 coverage.

411



412

Fig. 8. SGI maps in the potential distribution area (left column), the corresponding SGI histogram and the thresholds identified by multi-Otsu algorithm (center column), and the classification results of *Z. japonica* (right column) in (a) 2013-2014, (b)2003-2004, (c)1993-1995.



418 Fig. 9. SGI thresholds for *Z. japonica* extraction in each period during 1985-2018. The area
419 with SGI greater than the threshold is identified as *Z. japonica*.

### 420 **4.2** Accuracy assessment of seagrass bed maps

417

Table 2 lists the Z. japonica seagrass mapping accuracies in 1993-1995, 2007-2008, 2011-421 422 2012, 2013-2014, 2015-2016, and 2017-2018. The overall accuracies of the seagrass bed maps are higher than 95%, and the kappa coefficients are above 89%. The user's accuracies of Z. 423 424 japonica seagrass bed are generally higher than 94.67%, indicating very small commission errors. The producer's accuracies of seagrass beds are higher than 95.33% except for 2015-425 426 2016. In 2015-2016, the producer's accuracy is 90.06%, suggesting omission errors of around 427 10%. Examinations of Landsat images in 2015-2016 found that frequent cloud cover in both years may have resulted in insufficient acquisition of effective image pixels at low tide during 428 429 the green period. Overall, the validation results show that the seagrass maps generated from 430 SGI and multi-Otsu thresholding algorithm are accurate for further analysis of spatiotemporal dynamics of Z. japonica distribution. 431

432 Table 2. Confusion matrix of Z. japonica seagrass bed (ZJ) maps accuracy assessment. PA=

Veer	Class	Ground truth samples (pixels)		$\mathbf{D}\mathbf{A}$ (0/)		OA(0/)	Vanna
Year		ZJ	Non-ZJ	PA (%)	UA (%)	OA (%)	карра
1003 1005	ZJ	148	2	95.48	98.67	98.00	05.54
1993-1995	Non-ZJ	7	293	99.32	97.67		95.54
2007 2008	ZJ	142	8	98.61	94.67	07 79	04.05
2007-2008	Non-ZJ	2	298	97.39	99.33	97.78	94.93
2011 2012	ZJ	143	7	95.33	95.33	96.89	93.0
2011-2012	Non-ZJ	7	293	97.67	97.67		
2012 2014	ZJ	147	3	95.45	98.0	07.76	05.02
2013-2014	Non-ZJ	7	293	98.99	97.67	97.70	95.05
2015 2016	ZJ	145	5	90.06	96.67	05.22	80.60
2013-2010	Non-ZJ	16	284	98.27	94.67	95.55	89.09
2017 2018	ZJ	144	6	99.31	96.0	09.44	06 47
2017-2018	Non-ZJ	1	299	98.03	99.67	98.44	90.4/

433 producer's accuracy; UA = user's accuracy; OA = overall accuracy.

434

# 435 **4.3 Spatial-temporal dynamics of** *Z. japonica* seagrass bed

Fig. 10 shows that the area of Z. japonica fluctuated greatly, ranging from 149 ha in 2005-436 437 2006 to 1302.9 ha in 2011-2012. From 1985 to 1995, the area of Z. japonica generally showed 438 an increasing trend from 762.2 ha in 1985-1986 to 1060.9 has in 1993-1995, except that only 439 374.9 ha was detected in 1989-1990. From 1996 to 2010, the area of Z. japonica varied from 440 149.0 ha to 924.6 ha. From 2011 to 2018, the area presented obvious decreasing trend, from 441 1302.9 ha to 332.3 ha in 2017-2018. During the 34 years, no Z. japonica was detected in Zone C. Z. japonica was distributed in zone A throughout the 34 years but showed an overall 442 443 decreasing trend. Before 1999, Zone A had an average area of 904.4 ha Z. japonica except for 1989-1990, when only 374.3 ha of Z. japonica was detected. After 1999, the average Z. japonica 444 area was 652.2 ha. Z. japonica first appeared in Zone B in 1991. The area varied from 10 ha in 445 2005-2006 to 822 ha in 2011-2012. From 2011 to 2018, the area of Z. japonica decreased to 446

#### 447 166.1 ha. In 2001-2004 and 2009-2018, the area in Zone B exceeded that in Zone A.







#### 450 **4.3.1** Spatial distribution of seagrass bed before and after river course diversion

The river course of the Yellow River shifted twice in YRD during 1985-2018. The first 451 452 shift occured in 1996 when the Q8 channel was artifically constructed and the river course 453 migrated from the southeastward to the eastward. The second one occurred in 2007, with the end of the channel natually shifted northward. Fig.11 presents the Z. japonica seagrass maps 454 455 during 1985-2018, superimposed on the flase color Landsat imagery acquired in growing season. The Landsat imagery illustrates that the morphology of the shorelines changed 456 significantly after each river channel shift, and the spatial distribution of Z. japonica changed 457 accordingly. In 1985, only one large patch of Z. japonica was detected in the east of the artificial 458 coastline of Gudong Oil Field. Since 1985, this patch shrank substantially (Fig. 11). From 1985 459

to 1996, the shoreline around the river mouth extended southeastward for aournd 4.5 km.
Correspondingly, new patches of *Z. japonica* grew in the northern intertidal area from 1991 to

462 1995 (marked as 1) in Zone B in Fig. 11 d-e).







Fig. 11. Spatial distribution of Z. japonica seagrass bed in YRD.

464 In 1996, a large part of the patch in Zone B disappeared when the tidal flat formed with the sediment deposition during the river course diversion. Only a small patch remained in 1996 465 466 and then expanded from 1999 to 2004. After 1996, the river mouth extended eastwards. 467 Correspondingly, a big patch of Z. japonica was observed in the newly formed intertidal area of the north shoreline from 2001 (marked as 2) in Fig. 11h). In 2002, Yellow River Conservancy 468 469 Commission (YRCC) initiated water sediment regulation scheme (WSRS), aiming to sour the elevated riverbed in the lower Yellow River and maintaining the storage capacity in the 470 reservoirs in the downstream of Yellow River. From 2002 to 2018 except for 2016 and 2017, 471 WSRS was implemented every summer by creating man-made flood peak and expelling 472 473 accumulated sediment from the large reservoirs (e.g., Xiaolangdi Reservoir). During WSRS, significant increase in water and sediment delivery was observed and thus the sediment 474 deposition in the estuary resulted in rapid land gain in YRD (Li et al., 2019). Fig. 11 i-j 475

illustrates substantial expansion of tidal flats in YRD during the early stage of WSRS (2003-476 477 2006). This might cause Z. japonica seeds to be buried by the rapid deposition of sediments, 478 resulting in shrinkage of Z. japonica segrass beds patches in 2005-2006 (Fig. 11j). In 2007, the 479 flow channel in the river mouth diverted naturally to the northward, which again changed the 480 mophology of the shoreline in the river mouth (Fig. 11k). The rapid deposition of sediments 481 caused the damage of Z. japonica bed in the north bank of the new river mouth (Fig. 11k), while 482 new patches started to grow in 2011 (Fig. 11m). From 2007 to 2018, the northeastern shoreline 483 continued extending to the sea, while the eastern shoreline retreated toward the land. The Z. *japonica* seagrass beds in the southeastern bank remained relatively stable (marked as ③ in Fig. 484 485 10k-p).

#### 486 **4.3.2** Changes of seagrass beds after *S. alterniflora* invasion

S. alterniflora was first discovered in the YRD's intertidal zone in 2008. Since then, the 487 488 invasion of S. alterniflora has experienced three stages: slow expansion during incubation 489 period before 2011, rapid expansion during outbreak period from 2011 to 2017, and slower 490 expansion after 2017 (Ren et al., 2019; Wang et al., 2021). By 2019, the area of S. alterniflora reached 4672.38 ha (Wang et al., 2021). S. alterniflora expanded both landward and seaward. 491 492 As illustrated in Fig. 12, before 2011, only small patches of S. alterniflora were found in the 493 north bank of YRD. After 2011, S. alterniflora expanded rapidly to the seaward in both north 494 bank and south bank of YRD, and started to enchroach the habitat of Z. japonica since 2013. 495 Correspondingly, the seagrass bed patches shrank substantially from 2013 to 2018, especially 496 in the south bank of YRD. Fig. 10 showed that the area of Z. japonica decreased from 1302.9

ha in 2011 to 332.3 ha in 2018. This is consistent with previous report by Wang et al. (2021) that *S. alterniflora* has enchroached *Z. japonica* seagrass by around 902.32 ha. On the Landsat image acquired during growing season in 2019 and 2020, no obvious patches of *Z. japonica* seagrass beds were deteced (Fig. 12f). Our field investigations also showed that no large patches of *Z. japonica* were found in 2019 and 2020. In summer 2021, the investigators could hardly find intact *Z. japonica* patches which is larger than 1 m<sup>2</sup>.





Fig. 12. Spatial distribution of *Z. japonica* and *S. alterniflora* in YRD from 2009 to 2020. Note
that no *Z. japonica* was detected in 2019-2020.

# 505 **5 Discussion**

# 506 5.1 Advantages of SGI in seagrass mapping

In this study, we constructed SGI by taking into account both spectral, phenological and spatial characteristics of *Z. japonica* seagrass beds. Using SGI, we generated the first maps of *Z. japonica* seagrass beds from 1985 to 2018 in YRD. The resultant seagrass bed maps demonstrate good accuracies, with producer's accuracies higher than 90.06% and user's accuracies higher than 94.67%. The successful mapping can be explained by three advantages of SGI.

513 First, SGI was constructed by temporal composite approach on a pixel-by-pixel basis, 514 which effectively alleviates the influence of frequent cloud cover and tidal inundations in 515 coastal areas. Especially for seagrass beds that are submerged at high tide and exposed at low 516 tide, the maximum NDVI composite image (i.e.,  $NDVI_{max}$ ) highlights the greenest growth

517 condition of each Z. japonica pixel and eliminates the possible influence of tidal inundation on 518 a single image. In recent years, the pixel-based approach has been increasingly utilized for 519 coastal wetlands mapping and proved to achieve better accuracy than those from a single or 520 multiple cloudless images because it allows all pixels with valid observations to be utilized 521 regardless of the cloud coverage at a scene (Zhang et al., 2020; Jia et al., 2021; Hu et al., 2021; 522 Sun et al., 2021; Xu et al., 2021). For example, Jia et al., (2021) proposed to use maximum 523 mNDWI (mNDWI-MSIC) and maximum NDVI image composites (NDVI-MSIC) to extract 524 tidal flat, with the former extracting coastline at high tidal level and the latter extracting coastline at low tidal level. Note that in our study, we used the minimum mNDWI composite 525 526 to obtain the coastline at low tidal level instead of NDVI-MSIC. This is because the coastal 527 water in YRD is highly turbid. For the water area in YRD, the NDVI-MSIC composite generates 528 the most turbid pixels throughout a year, which are difficult to be distinguished from tidal flats 529 as they have similar NDVI values. In contrast, mNDWI was less influenced by water turbidity 530 (Xu, 2005). The minimum mNDWI of tidal flat pixels represents the status when they are fully 531 exposed at low tide level, and thus help extract more accurate coastlines at low tide level. Second, SGI considers the spectral and phenological characteristics, and spatial location 532 of Z. japonica seagrass beds as well as salt marsh vegetations such as S. alterniflora and S. 533

534 salsa. With S. alterniflora invading seaward, its niche overlaps with that of Z. japonica (Fig.

535 11). It is difficult to distinguish S. alterniflora and Z. japonica solely based on  $NDVI_{max}$  (Fig.

536 5a). However, S. alterniflora is distributed in higher intertidal area and has much taller and

537 dense plants compared to Z. japonica, thus its mNDWI at low tide level is negative and similar

as terrestrial vegetation. As *Z. japonica* has short plants and is frequently submerged at high
tide level, it has low brightness value than *S. salsa* and tidal flats. In addition, *Z. japonica* has
much shorter growing length than salt marsh vegetation. Compared to any of the vegetation
indices, the integration of spectral and phenological indices in SGI exaggerate the difference
between *Z. japonica* and others.

Third, SGI allows automatic and rapid mapping of Z. japonica using unsupervised 543 classification algorithm. As Z. japonica and other types showed distinct SGI values, the 544 545 unsupervised extraction using multi-class Otsu algorithm achieved good accuracy. Most previous studies utilized supervised classification algorithms for seagrass mapping (Coffer et 546 al., 2020; Kuhwald et al., 2021; Fernandes et al., 2022; Lebrasse et al., 2022). Sample collection 547 548 is the most time-consuming and labor-intensive task in supervised classifications. It is 549 especially difficult for the coastal wetlands as many areas are hard to reach for field surveyors. 550 In addition, the long-term distribution of seagrass beds is usually poorly documented, 551 hampering collection of sufficient training datasets for retrospective mapping. Compared to the 552 supervised classification method, the unsupervised Otsu threshold segmentation algorithm does not require the collection of training samples and the classification process is automated without 553 554 human intervention, which is suitable for the long time series and large-scale dynamic 555 monitoring of seagrass beds in YRD. Therefore, SGI combined the multi-Otsu algorithm can 556 be easily applied to other time-series satellite imagery such as Sentinel-2 or the Harmonized 557 Landsat-8 Sentinel-2 surface reflectance for seagrass extraction. In 2021, Shandong Province 558 started to implement seagrass beds restoration in the south of YRD. Our proposed method has

559 potential to provide technical support and basic data for regular seagrass restoration monitoring.

560 5.2 Uncertainties and limitations of our seagrass mapping method

561 Although our method achieved high accuracy in long-term seagrass mapping YRD, 562 uncertainties and limitations still exist. First, successful application of this method relied on 563 temporally dense observations during specified phenological periods. Due to limited Landsat 564 data before 1999 and scan-off failure in Landsat 7 ETM+ after 2003, our method generated 565 seagrass bed maps every two years instead of performing annual mapping. Although using 566 Landsat imagery during two-year period effectively increases the number of valid observations at pixel level and can better represent vegetation phenology, it may not capture the rapid annual 567 568 change in Z. japonica seagrass beds. For example, research has shown that S. alterniflora can 569 push over 10 m into Z. japonica region during several months (Yue et al., 2021). Because S. alterniflora has generally higher NDVI than Z. japonica, using NDVI<sub>max</sub> during two-year 570 571 period probably identified S. alterniflora instead of Z. japonica for those pixels where Z. japonica was replaced by S. alterniflora. In this case, the seagrass bed maps represent the 572 573 minimum area of Z. *japonica* during the two-year period. Nevertheless, the problem can be well 574 resolved when dense observations are available during a year. Second, our method is suitable 575 for intertidal seagrass extraction where the seagrass beds are exposed at low tide. For submerged 576 seagrass beds in subtidal area, it may not work as water column correction is needed to retrieve 577 correct bottom reflectance. Third, the 30 m-resolution Landsat images inevitably produce mixed-pixel problem. Small patches of Z. japonica seagrass beds with low biomass may not be 578 579 identified. In July 2019, we found some small patches of Z. japonica in our field investigation 580 in Zone B. Unfortunately, our method did not detect any of Z. japonica in this year.

# 581 **5.3 Driving factors for** *Z. japonica* seagrass beds variations in YRD

582 Previous studies have reported that many factors can cause the degradation and loss of 583 seagrass beds, including coastal erosion, extreme climate events, and human activities such as 584 coastal development and water pollutions (Kim et al., 2015; Kendrick et al., 2019; Oprandi et 585 al., 2020). The YRD is probably one of the most active deltas in the world's estuaries because 586 of large interannual and intra-annual variations in sediment transport and recent invasions of S. 587 alterniflora. In YRD, the large variations of the spatial extents of Z. japonica during 1985-2018 (Fig. 9 and Fig. 10) indicates that the driving factors can be very complex. To date, no previous 588 589 study has investigated the driving factors for the long-term spatial variations of Z. japonica 590 meadows in YRD. Our results demonstrated two convincing factors for the loss of Z. japonica 591 seagrass beds: (1) rapid and large amount of sediment deposition in the estuary due to river 592 channel diversion and early implementation of WSRS, and (2) S. alterniflora invasion in the recent decade. 593

After each river channel diversion (1996 and 2007) and during the early stage of WSRS (2003-2006), the morphology of estuarine land changed substantially owing to rapid sediment deposition in the river mouth. Several patches were buried by the newly formed land, and the sediment burial hampers the germination of seeds (Cabaço and Santos, 2007), which could cause rapid loss of the patches (e.g., the patches in Zone B in 1996-1998 and Zone A in 2005-2006, Fig. 11f and Fig. 11j). Meanwhile, new seagrass patches started to grow in the newly formed intertidal area (e.g., the patches in Zone A in 2001-2002, Fig. 11h). Note that after 2007, the continuous implementation of WSRS seemed not affect seagrass beds, regardless of the highly turbid water around coastal area (Li et al., 2019). From 2007 to 2012 (before explosion of *S. alterniflora*), we did not observe rapid declination of *Z. japonica* area. This is consistent with recent in-situ field investigations and laboratory experiment by Hou et al. (2020) and Zhang et al. (2021), which concluded that *Z. japonica* in YRD shows good short-term resistance to high turbidity during WSRS.

S. alterniflora invasion is the primary reason for the gradual Z. japonica's degradation in 607 608 the recent decade. S. alterniflora has great reproduction capacity with both sexual and asexual reproduction. Our results showed that from 2013 to 2018, the area of Z. japonica has been 609 610 encroached by S. alterniflora with 868.4 ha. Ma et al., (2020) reported that S. alterniflora, 611 regardless of the plant densities, have significant inhibition effects on the symbiotic Z. japonica. 612 The stem density, height and total biomass of Z. japonica decreased dramatically once the 613 invading patches of S. alterniflora arise. Yue et al., (2021) explained three steps of S. 614 alterniflora invading Z. japonica: first, seeds of S. alterniflora float into the Z. japonica and the 615 sparse patches of the invader grow; subsequently, sediment accumulation increased with the growing density of clonal ramets, and the taller S. alterniflora patches blocked the sunlight 616 617 needed for Z. japonica, gradually inhibiting the growth of Z. japonica; finally, the patches of S. 618 alterniflora connected and replace the Z. japonica community.

619 Other possible factors that affect the growth of *Z. japonica* seagrass beds in YRD include 620 soil erosion in local area and extreme climate events including drought and typhoon (timeline

621 illustrated in Fig. 9). The loss in the Z. japonica patch near Gudong Oil field from 1985 may be

622 explained by severe coastal erosion in this area. Ji et al., (2018) reported that Gudong nearshore 623 experienced severe erosion (-0.1 m/yr) due to reduction of sediment supply and strong wave 624 currents. In 2000-2002, the rapid loss of Z. japonica may be explained by severe drought events 625 in 2000, typhoon Toraji in summer 2001 and typhoon Fengshen in summer 2002. The severe 626 drought in Shandong Province in 2000 resulted in hyper-salinity and water column stratification 627 in coastal YRD (Xi et al., 2001; Hall et al., 2016), which can lead to mortality of Z. japonica. Typhoon can be another factor. Recent investigations found that the super typhoon Lekima in 628 629 2019 resulted in over 100-fold loss of the area of Z. japonica in YRD due to strong winds, heavy rainfall and sudden soil erosion (Yue et al., 2021). 630

#### 631 **5.4 Implications for seagrass management and restoration**

632 This study provides the first long-term Z. japonica seagrass bed maps in YRD, filling the 633 knowledge gaps on the seagrass bed extents in coastal China. We hope our results can 634 significantly facilitate in-depth understanding on the mechanisms and driving factors for Z. 635 japonica variations in YRD, as well as the understanding on the ecosystem services they 636 provide. It is unfortunate that the typhoon Lekima led to severe loss of Z. *japonica* meadows in YRD in 2019 (Yue et al., 2021). Restoration efforts are required urgently for seagrass meadow 637 638 recovery. Our long-term Z. japonica seagrass maps can serve as an intrinsic basis for the 639 development of seagrass restoration measures. In fact, in recent years numerous restoration 640 efforts have been implemented in the coastal provinces in China, including Hebei, Shandong, Guangxi and Hainan provinces (Liu et al., 2016; Yu et al., 2019; Xiao et al., 2020). The long-641 642 term seagrass mapping methods developed in this study have great potential to be applied for timely monitoring and evaluation of the effectiveness of the restoration efforts at regional scale.
As seagrass bed is one of the most important blue carbon ecosystems, our study provides
baseline efforts for seagrass carbon storage estimation and long-term monitoring, which is
critical to maintain coastal sustainability.

647

648 6 Conclusion

This paper proposed a new seagrass index, namely SGI, for automatic mapping of Z. 649 japonica, an intertidal seagrass species widely distributed in China. SGI alleviates the influence 650 651 of tidal inundation and enhances the spectral and phenological separability between seagrass 652 beds and other cover types by incorporating temporal composites of NDVI, mNDWI and TCBI 653 based on time-series remote sensing imagery. Using SGI, we then extracted Z. japonica extents 654 in YRD based on all available Landsat 5/7/8 images during 1985-2018 with SGI thresholds automatically determined by multi-Otsu algorithm. The results showed that SGI has 655 656 successfully discriminated the Z. japonica seagrass beds and non-Z. japonica types such as salt 657 marshes, tidal flat and water. The SGI thresholds were relatively stable, ranging from 2.22 to 658 4.11. The overall accuracies were greater than 95%, producer's accuracies and user's accuracies 659 of Z. japonica were greater than 90% and 94%, respectively, which were validated through field 660 inventory data, high resolution satellite imagery and Landsat imagery. From 1985 to 2018, the area and spatial distribution of Z. japonica showed large variations (from 149 ha in 2005-2006 661 to 1302.9 ha in 2011-2012). River channel shifts in YRD altered the spatial distribution of Z. 662 japonica, and the expansion of invasive salt marsh vegetation S. alterniflora caused gradual 663

degradation of *Z. japonica* in recent years. Coastal erosion and extreme climate events such as drought and typhoon are other possible factors explaining *Z.* japonica area decline. In sum, this paper provides the first long-term seagrass bed maps in YRD. We expect that the SGI will advance automatic and rapid mapping methods for intertidal seagrass beds, and the *Z. japonica* maps will provide a baseline data for restoration and management of seagrasses at regional scale.

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