Name of indicator	2.15 Reed belt extent – the NDVI approach via high resolution satellite images
Type of Indicator	State indicator
Author(s)	Hanna Piepponen, Meri Koskelainen and Kirsi Kostamo
Description of the indicator	The indicator expresses the extent of coastal reed belts, using information from remote sensing and exposition-depth data. The reed belt extent in the archipelago of South-western Finland, in the MARMONI 3FIN study area, was examined from high resolution satellite images dating from September 2009 (RapidEye, 5m by 5m resolution) and July 2013 (WorldView, 2m by 2m resolution). For verifying the results, data from Kotka located in South-eastern Finland was used. The Kotka satellite images dated from July 2011 (RapidEye, 5m by 5m resolution) and September 2012 (WorldView, 2m by 2m resolution). The indicator is a combination of Normalized Difference Vegetation Index (NDVI) and the exposition-depth data (Isaeus & Rygg 2005, Tolvanen 2010) that shows the optimum growing area for reed vegetation.
	The indicator demonstrates the local extent of reed communities very well in the two test areas. NDVI calculated from satellite images shows the actual extent and location of reed communities in sheltered areas. A temporal aspect is possible to attain by comparing seasonal changes in reed vegetation, provided that suitable satellite images are available. However, both the temporal and spatial coverage of the indicator is still fairly restricted due to a lack of available high resolution satellite images, and for a better coverage of reed vegetation information in the Gulf of Finland a higher number of high resolution satellite images are needed. Images of a coarser resolution are not useful for local scale reed belt extent studies because they do not provide reliable spatial information for the calculation of the size and shape of the reed belt by the NDVI. Coarse resolution Landsat (30m by 30m resolution) satellite images were tested but the results are too rough for closer examination of changes in reed belt extent. The lack of high resolution satellite images also prevents comparing temporal changes in coastal areas in a long time scale, so the development of the index requires further studies with upcoming satellite images. Availability of usable high resolution satellite images is not certain because the weather conditions and scanning time limit the use of all available images from a study area. Summarizing the above, high resolution satellite images of sufficient spatial and temporal frequency are requisite for the indicator.
Relationship of the indicator to marine biodiversity	Common reed, <i>Pharagmites australis</i> (Cav.) Trin. Ex Steudel, is an erect perennial grass growing in lakes, trenches, shore and bog meadows in Northern Europe (Hämet-Ahti <i>et al.</i> 1998). It often forms dense and more or less monospecific patches or areas consuming all available growing space. Both the number and coverage of reed patches has increased in many areas both in disturbed and forested sites in Northern America (McCormick <i>et al.</i> 2010).
	Common reed is a rapidly expanding species which can reproduce both clonally and sexually. Even though it forms very dense patches, negative effects of shading on plant biodiversity have not been established (Güsewell & Edwards 1999). Furthermore, it has been discovered that the species richness does not decrease as a result of an increase in reed biomass (Grime 1973, 1979, Güsewell & Edwards 1999), but changes from aquatic macrophytes towards species occurring in the geolittoral (Munsterhjelm 1997). Furthermore, high reed biomasses may alter the proportion of species so that the plant community will develop so that light-demanding species from nutrient-poor sites are replaced by more shade-tolerant and more nutrient-demanding species. In light of these results, it is also likely that an increase in eutrophication may affect the plant community composition (Tilman 1982, 1987, Olff 1992, Eek & Zobel 1997).
	Since the light interception by common reed follows a clear seasonal pattern and is negligible until June, its impacts on species developing in spring or early summer are likely to be strongly reduced by phenological separation (Güsewell & Edwards 1999). Therefore species, growing until the end of the summer, are more likely to be influenced by <i>P. australis</i> than species that complete their annual growth in early summer or which are at least capable of doing so if light conditions decline. The relationship between species richness and reed biomass is further complicated by the different types of wetland communities (Güsewell & Edwards 1999). Thus the coexistence of other wetland species and succession of species composition can increase the biodiversity of the reed communities.
Relevance of the indicator to different policy instruments	EU Marine Strategy Framework Directive: descriptor 1 Biodiversity, 1.5 Habitat extent, 1.5.1 Habitat area. HELCOM Baltic Sea Action Plan ecological objective for 'natural marine and coastal landscapes'. Habitats Directive: state of the coastal habitats and protected species.
Relevance to	1.5. Habitat extent

commission	1.5.1. Habitat area
decision criteria	
Method(s) for	The coverage of common reed was estimated by using information provided by satellite
values	remote sensing (RapidEye 5m by 5m resolution and WorldView-2 2m by 2m resolution). The reed vegetation presence was determined from the images by calculating the Normalized
	Difference Vegetation Index (NDVI), which was calculated from the band relations between
	red and infrared bands. NDVI areas were extracted to water areas by clipping the data by shoreline as we assumed that all the vegetation in water is reed vegetation. The threshold
	value of the NDVI was set in both summer and fall images to 0.2 to avoid errors caused by
	highly reflecting objects such as sailboats. In general, the reflection from vegetation was
	and shows the maximum vegetation cover. The indicator utilizes depth-exposition data to
	determine the potential growing area of reed belts. Used depth-exposition data covers the
	(sheltered and moderately sheltered areas, Munsterhjelm 1997, 2005); these conditions
	were considered to be the optimum growing area for reed vegetation. Over 95% of NDVI
	vegetation is coherent. Cloudy areas on each used data were removed for reducing errors
	and the pixel size of each data was resampled to the same size (4m by 4m). In addition to
	measurements during summer 2013 in Tammisaari.
	A comparison of summer and fall images revealed that it is possible to use both late
	summer and early fall satellite images for determining reed belt extent. Spring images
	(April, May) are not useful in Finnish coastal areas especially when growing season starts late due to elongated winter because the reed coverage reaches its maximum extent only in
	late June or July. Fall images (September) are usable but they underestimate the reed belt
	extent by about 10%. This was verified by calculations in Kotka, South-eastern Finland, from satellite images in July 2011 and in September 2012. The assumption was that reed
	vegetation has not changed during one year, and therefore, the reed extent in fall and
	summer images should be the same. As the reed extent in fall was smaller than in summer we derived a coefficient of 10% of underestimation for fall images. After adding 10% to reed
	vegetation data in fall 2009 in Tammisaari in the MARMONI 3FIN study area, we concluded
Documentation of	The main pressures affecting the extent of the reed belts are land use and eutrophication. In
relationship	the coastline of the Northern Baltic Sea, also land uplift influences the reed extent through a
and pressure	(Munsterhjelm 1997, 2005). Disturbance of upland habitats and eutrophication of estuaries
	have been shown to be positively correlated with the abundance of common reed (Bertness
	level of disturbance in developed watersheds can create open spaces for seedling
	emergence and rhizoid settlement and establishment and thus facilitate dispersal of this
	surrounding uplands (Burdick et al. 2001, Bertness et al. 2002), atmospheric enrichment of
	nitrogen and carbon dioxide, and altered climate, may also enhance invasions (Minchington
	from, e.g., CORINE-remote sensing data, but since this data exists currently only from 2000
	and 2006, more data is needed to confirm this relationship.
	The effects of eutrophication are more complex in macrophyte communities than in plankton
	or annual macroalgal communities, because macrophytes take up nutrients with roots from the bottom sediments and not directly from the water. The effects of eutrophication are
	therefore accumulative and should be studied more in relation to a temporal
	aspect, <i>e.g.</i> , the nutrient content of the water column versus the nutrient content in the sediment. When considering grasses, including common reed, the increase in reproductive
	output resulting from eutrophication may enhance the invaders ability to establish new,
	genetically distinct populations and enhance the spatial dominance in already invaded areas. Furthermore, increased input of atmospheric nitrogen and carbon dioxide levels can alter
	the competitive balance of marsh plants in favour of common reed (Jaworski <i>et al.</i> 1997).
	Nutrient enrichment results in taller stems, increase in floret and inflorescence production
	and overall biomass, but also an increase in asexual reproduction (Minchinton & Bertness
	Mozdzer & Zieman 2010, Kettenring <i>et al.</i> 2011). On one hand, it has been discovered that
	if the reed colony is under high nitrogen levels, it may invest heavily on rhizome production (Rickey & Anderson 2004). This enables dispersal to areas, where low pitrogen content
	prevents population establishment by seeds (Bart & Hartman 2002). On the other hand, in
	high nutrient regime, seedling emergence and establishment benefit from eutrophication

	and increase the probability that population establishment occurs via seeds (Saltonsall & Stevenson 2007).
	Surprisingly, according to some studies, it has also been discovered that under high nutrient levels the below-ground biomass does not increase, although this is expected in populations where dispersal occurs by fragmented pieces of rhizoids (Haslam 1965, Rickey & Anderson 2004). Instead, the above-ground biomass increases (Minchinton & Bertness 2003, Rickey & Anderson 2004). It has also been discovered that under a high nutrient regime, the species spreads to deeper water than in oligotrophic conditions (Haslam 1965, 1972).
	The increase in nitrogen levels has been linked to the successful dispersal of common reed in a large number of areas (Haslam 1965, Marks <i>et al.</i> 1994). However, it has also been discovered that an increase in nitrogen increases also the growth of native species, so the eradicative effects of nutrients might not be as strong as assumed earlier (Rickey & Anderson 2004). All in all however, present knowledge suggests that eutrophication favours reed belt extent.
Geographical relevance of indicator	4. Baltic Sea wide
How Reference Conditions (target values/thresholds) for the indicator were obtained?	The target is trend-based, expecting no increase of area covered by reed belts, as indicated by the NDVI, in order to achieve Good Environmental Status.
Method for determining GES	The trend-based target for reed belt extent, as indicated by the NDVI, is estimated for the years 2002-2013, using medium resolution images for 2002-2009 and high resolution images for 2009-2013.
References	Bart, D. & Hartman, J.M. 2002. Environmental constraints on early establishment of <i>Phragmites australis</i> in salt marshes. Wetlands 22:201-213.
	Bertness, M.D., Ewanchuk, P. & Sillman, B.R. 2002. Anthropogenic modification of New England salt marsh landscapes. Proceedings of the National Academy of Sciences of the United States of America 99:1395-1398.
	Burdick, D.M., Buchsbaum, R & Holt, E. 2001. Variation in soil salinity associated with expansion of <i>Phragmites autralis</i> in salt marshes. Environmental and Experimental Botany 46: 247-261.
	Burdick, D.M. & Konisky, R.A. 2003. Determinants of expansion for <i>Phragmites australis</i> , common reed, in natural and impacted coastal marshes. Estuaries 26: 407-416.
	Chambers, R.M., Havens, K.J., Killeen, S. & Berman, M. 2008. Common reed, <i>Phragmites australis</i> , occurrence and adjacent land use along estuarine shoreline in Chesapeake Bay. Wetlands 28:1097-1103.
	Eek, L. & Zobel, K. 1997. Effects of additional illumination and fertilization on seasonal changes in fine-scale grassland community structure. Journal of Vegetation Science 8:225-234.
	Grime, J.P. 1973. Control of species density in herbaceous vegetation. Journal of Environmental Management 1:151-167.
	Grime, J.P. 1979. Plant strategies and vegetation processes. Wiley & Sons, Chichester.
	Güswell, S. & Edwards, P. 1999. Shading by <i>Phragmites australis</i> : a threat for species-rich fen meadows. Applied Vegetation Science 2:61-70.
	Haslam, S.M. 1965. Ecological studies in the Breck fens. I. Vegetation in relation to habitats. Journal of Ecology 53:599-619.
	Haslam, S.M. 1972. Biological flora of the British Isles. Journal of Ecology 60:585-610.
	Hämet-Ahti, L., Suominen, J., Ulvinen, T. & Uotila, P. (eds) 1998: Retkeilykasvio, 4. uudistettu painos, 656 pp. Botanical museum of the Finnish Museum of Natural History. Helsinki.
	Isæus, M. & Rygg, B. 2005. Wave exposure calculations for the Finnish coast. Report.

Norwegian institute of water research.
Jaworski, N.A., Howarth, R.W. & Hetling, L.J. 1997. Atmospheric deposition of nitrogen oxids onto the landscape contributes to coastal eutrophication in the northeast United States. Environmental Science and Technology 31: 1995-2004.
Kettenring, K.M., McCormick, M.K., Baron, H.M. & Whigham, D.F. 2011. Mechanisms of <i>Phragmites</i> <i>australis</i> invasion: feedbacks among genetic diversity, nutrients, and sexual reproduction. Journal of Applied Ecology 48: 1305-1313.
King, R.S., Deluca, W.V., Whigham, D.F. & Marra, P.P. 2007. Threshold effects of coastal urbanization on <i>Phragmites australis</i> (Common Reed) abundance and foliar nitrogen in Chesapeake Bay. Estuaries and Coasts 30:1-13.
Luther, H. 1951a. Verbreitung und Ökologie der Höheren Wasserpflanzen im Brackwasser der Ekenäs-Gegend in Südfinnland. I. Allgemeiner Teil. – Acta Bot. Fennica 49: 1-231.
Luther, H. 1951a. Verbreitung und Ökologie der Höheren Wasserpflanzen im Brackwasser der Ekenäs-Gegend in Südfinnland. II. Spezieller Teil. – Acta Bot. Fennica 50: 1-370.
Marks, M., Lapin, B. & Randall, J. 1994. <i>Phragmites australis (P. communis</i>): Threats, management, and monitoring. Natural Areas Journal 14: 285-294.
McCormick, M.K., Kettenring, K.M., Baron, H.M. & Whigham, D.F. 2010. Extent and reproductive mechanism of <i>Phragmites australis</i> spread in brackish wetlands on Chesapeake Bay, Maryland (USA). Wetlands 30:67-74.
Minchinton, T.E. 2002. Disturbance by wrack facilitates spread of <i>Phragmites australis</i> in a coastal marsh. Journal of Experimental Marine Biology and Ecology 281:89-107.
Minchinton, T.E. Bertness, M.D. 2003. Disturbance-mediated competition and the spread of <i>Phragmites australis</i> in a coastal marsh. Ecological Applications 13:1400-1416.
Mozdzer, T.J. & Zieman, J.C. 2010. Ecophysiological differences between genetic lineages facilitate the invasion of non-native <i>Phragmites australis</i> in North American Atlantic coast wetlands. Journal of Ecology 98:451-458.
Munsterhjelm, R. 1997. The aquatic macrophyte vegetation of flads and gloes, S coast of Finland. Acta Bot. Fennica 157: 1-68.
Munsterhjelm, R. 2005. Natural succession and human-induced changes in the soft-bottom macrovegetation of shallow brackish bays on the Southern coast of Finland. – W. & A. de Nottbeck Foundation Sci. Rep. 26: 1-53.
Olff, H. 1992. Effects of light and nutrient availability on dry matter and N allocation on six successional species: Testing for resource ratio effects. Oecologia (berl.) 89:412-421.
Rickey, M.A. & Anderson, R.C. 2004. Effects of nitrogen addition on the invasive grass <i>Phragmites australis</i> and a native competitor <i>Spartina pectinata</i> . Journal of Applied Ecology 41:888.
Saltonsall, K., Peterson, P.M. & Soreng, R.J. 2004. Recognition of <i>Phragmites australis subsp. americanus (Poeaceae: Arundinoideae</i>) in North America: evidence from morphological and genetic analyses. SIDA 21:683-692.
Saltonsall, K. & Stevenson, J.C. 2007. The effect of nutrients on seedling growth of native and introduced <i>Phragmites australis</i> . Aquatic Botany 86:331-336.
Silliman, B.R. & Bertness, M.D. 2004. Shoreline development drives invasion of <i>Phragmites australis</i> and the loss of plant diversity of New England salt marshes. Conservation Biology 18:1424-1434.
Tilman, D. 1982. Resource competition and community structure. Princeton University Press, Princeton, NJ.
Tilman, T. 1987. On the meaning of competition and the mechanisms of competitive superiority. Functional Ecology 1:304-315.





