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Dendroecological study of *Pinus halepensis* and *Pinus pinea* in northeast coastal dunes in Tunisia according to distance from the shoreline and dieback intensity

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A B S T R A C T

Growth decline is a widespread phenomenon on sandy coasts throughout the Mediterranean region. In the northeastern pine coastal woodland of Tunisia, *Pinus pinea* and *Pinus halepensis* stands originally afforested to stabilize sand dunes are often damaged. Our objective was to estimate the effect of distance from the shoreline on ring- width variation of two pine species, *Pinus pinea* and *Pinus halepensis* located in the Remel and Menzel Belgacem coastal forest land. 15 to 30 trees of each species were selected based on three dieback levels (low, moderate and severe canopy damage). The results of dendroecological analysis show a significant reduction in the mean growth of trees growing near the shoreline for both species. Most stands show a positive correlation with precipitation between October of the previous year to April of the current year. The number of months, when precipitation is correlated positively with growth, is always lower for trees growing near the shoreline than for trees growing in distant stands. However, when the maximum temperature is correlated negatively with growth, the number of months is always higher for trees growing near the shoreline (2 months) than for trees those in distant stands (1 month). Moreover, there was an increased number of missing rings in the 1980s but only in those trees growing near the shoreline and their frequency was only statistically significant for *Pinus pinea* in the two forest sites. The severity of canopy damage due to being close to the shoreline negatively influenced the growth of both species and could threaten their survival in these areas.

1. Introduction

Forests, which cover about 30 percent of the world's land surface (FAO, 2006), are being rapidly and directly impacted in many areas by human expansion. Less evident are the pervasive effects of ongoing climatic changes on the condition and status of forests around the world. Increased greenhouse gas emissions are now widely acknowledged by the scientific community as a major cause of the recent rise in global mean temperature and changes in the world's hydrological cycle (IPCC, 2007). Understanding and predicting the consequences of these climatic changes on ecosystems is one of the greatest challenges for scientists, and forecasting the impacts on forests is particularly important (Boisvenue and Running, 2006; Bonan, 2008). Of these impacts on forests, tree dieback is a serious phenomenon on a local, regional and national level (Allen et al., 2010). This phenomenon alters the

external appearance of the tree due to organ mortality, reducing the quality and quantity of foliage, and may cause stunted growth or even death in some trees (Delatour, 1990). This dieback phenomenon is either caused by pathogens, parasites or conditions like acid rain and drought (Allen, 2009).

Examples of drought and heat-related forest stress and dieback are being documented by all forested continents, making it possible to begin to see global phenomena. Two of the nine tipping points for major climate changes forecast for the next century are directly related to forest dieback (IPCC, 2014). Examples of dieback have been particularly well documented in southern Europe (Breda et al., 2006 and Bigler et al., 2006) and in the temperate and boreal forests of western North America, where background mortality rates have increased rapidly in the past few decades (van Mantgem et al., 2009). The common causal factor in these examples is elevated temperatures and/or water

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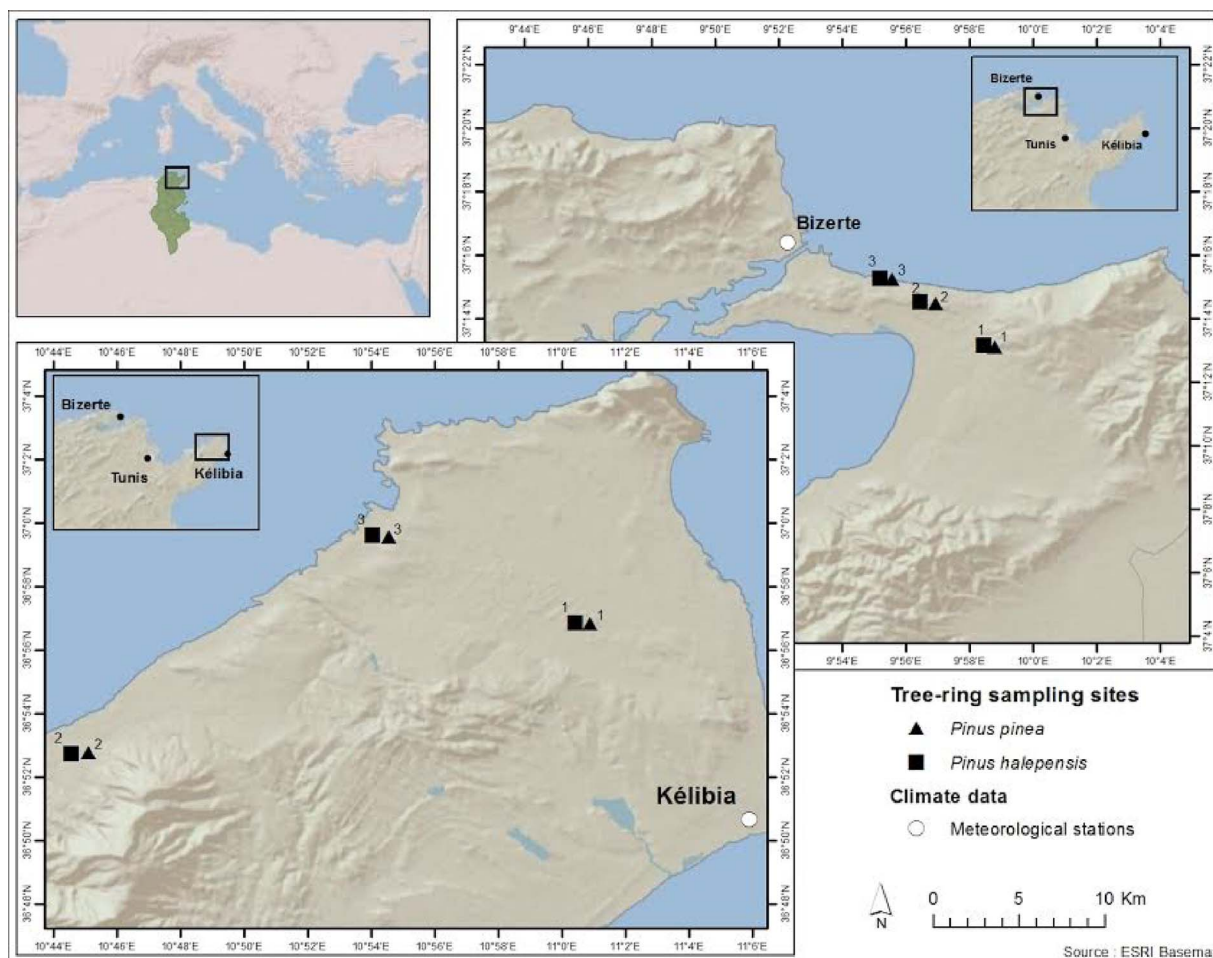


Fig. 1. Location of studied stands of *Pinus halepensis* and *Pinus pinea*, Northeast Tunisia.

stress. Recent extreme drought in North Africa (Touchan et al., 2008) has been linked to severe mortality of Atlas cedar (*Cedrus atlantica*) from Morocco to Algeria (El Abidine, 2003; Bentouati, 2008). Disturbing events have occurred in some forest areas in Tunisia over the last few years (Ben Jamaa et al., 2005, 2006a), raising a number of issues including the effects of drought and global warming (Ben Jamaa et al., 2006b; El Khorchani et al., 2013). Sudden and widespread mortality struck cork oak forests in Tunisia in the late 80s and early 90s after three years of consecutive drought, affecting more than 20% of the trees in different cork oak forests (Hasnaoui et al., 2005). The same dieback phenomenon was observed between 2000 and 2003 in Aleppo pine forests in a semiarid bioclimate (Aloui, 2010).

One of the most vulnerable areas to climate change is the coast where about 30% of the entire coastline is affected by erosion (EEA, 2006). According to commonly accepted models of coastal dynamics (Bruun, 1988; Davidson-Arnott, 2005), an increase in coastal erosion should therefore be expected in the future together with an increase in vulnerability of the coastal ecosystem (IPCC, 2007). Tree dieback along the coast is due to a combination of factors unfavorable to tree growth such as drought, storms, and nutritional problems which lead to overall weakening of the tree and possible death (Garrec, 1994). Moreover, pollution from urbanization, industry, maritime traffic and agriculture converge in coastal waters. This ecosystem degradation directly affects vegetation under stress and demonstrates a rupture in forest management. Since the 80s, considerable damage has been observed in the maritime pine forests of northern Tunisia, in Bizerte (forests of M'hibes and Rimel) and Tabarka (Sidi Badr forests), following the attack of the insect *Tomicus piniperda* L. (Ben Jamaa et al., 2000). According to a dendroclimatological study of the impact of drought on the growth of

three pine species in Tunisia (maritime pine, stone pine and Aleppo pine), the number of dry years and observably narrower rings increased from 1978 to 2001 compared to the period between 1954 and 1977 (El Khorchani et al., 2006). Deterioration of coastal vegetation in the north of Tunisia is also related to the presence of several pollutants in the marine spray such as anionic surfactants and hydrocarbons (El Ayeub et al., 2004). The acceleration of climate change, extreme climatic events (storms, heat, frost, drought and excess water), anthropogenic changes in land use, and pollution are all factors which can contribute directly or indirectly, alone or in interaction, to the explosion of cases of ecosystem dysfunction. Sustainable and responsible management therefore requires new adaptation strategies. These may include assessing forest vulnerability to climate change, reviewing forest use expectations, identifying research and education needs, developing forest policies to facilitate and anticipate adaptation and maintaining the health and vitality of forest ecosystems (Spittlehouse, 2005). Knowing the extent of the territory affected by dieback will contribute greatly to the decision-making process concerning this crisis. For example, some species currently widely used for reforestation of coastal dunes in Tunisia may be restricted in the future and replaced by other species more resistant to drought, or their silvicultural management techniques may be revised in order to adapt to climate and territorial changes.

Two of the main species used to reforest Mediterranean dune ecosystems are *Pinus pinea* and *Pinus halepensis*. These pine species have to survive in hard environmental conditions with constant exposure to threatening marine spray and water stress conditions (Garrec, 1994). In southeast Spain, Raventós et al. (2001) showed that *Pinus pinea* and *Pinus halepensis* are highly sensitive to exposure both to sea spray and to surfactants in marine aerosols, as reported in other parts of the

Mediterranean coast of Spain (Diamentapolos et al., 2000). Reduced growth of *Pinus pinea* along the Tyrrhenian coast of Tuscany (Italy) is due to the crown's greater exposure to salty winds and surfactants and the increased salinity of the water table (Raddi et al., 2009).

Remel and Menel Belgacem forests, pine coastal woodlands of Tunisia, are good examples for studying the vulnerability of these two species under severe conditions of marine spray and drought. In these areas located along the northern coastal dunes, *Pinus pinea* and *Pinus halepensis* have clearly suffered canopy damage due to the proximity of the shoreline. In this study, a dendroecological approach was used to estimate the effect of distance from the shoreline on two pine species, *Pinus pinea* and *Pinus halepensis*, growing in two forest sites in northeast Tunisia.

2. Material and methods

2.1. Study area

The study was carried out in northeastern Tunisia (Fig. 1) on two coastal *Pinus pinea* and *Pinus halepensis* planted forests. The first site is located in Remel Forest (37°11'N, 10°04'E) in the region of Bizerte, and the second is in Menzel Belgacem Forest (36°57'N, 10°57'E) in Kelibia in the Cap Bon region. These forest sites have a sandy dune ecosystem and belong to the Mediterranean subhumid bioclimate. The mean annual rainfall in the Bizerte site is 656 mm, concentrated mainly in the winter. In Kelibia, the mean annual rainfall is 550 mm, mostly falling in autumn. Annual temperature varies from 7 °C (January) to 32 °C (August) in Bizerte and from 8 °C (January) to 31 °C (August) in Kelibia. Northwestern winds with a mean annual speed of 3.8 m s⁻¹ and 3 m s⁻¹ are most dominant in Bizerte and Kelibia, respectively.

The sand dunes of both sites have been afforested with species such as *Acacia sp.*, *Eucalyptus sp.*, *Pinus halepensis*, *Pinus pinea* and *Pinus pinaster* since the 1900s to fix the dunes along the coastal area.

2.2. Collection of data and measurements

2.2.1. Structural data

From 2010–2015, 12 plantation stands of *Pinus pinea* and *Pinus halepensis* with different environmental conditions were studied in the Remel and Menzel Belgacem forests. For each species in each forest site, three stands were sampled along a latitudinal transect according to distance from the shoreline. Characterization of the stands was analyzed by spiral plots according to the protocol used by the Department of Forest Health in France for determining tree dieback classes (Office National des Forêts, 2010). The center of the plot being fixed by the sampling plan, the first tree is the closest to the center of the plot and the following trees are chosen step by step in a spiral to reach the defined number (recommended: 20 trees within a maximum radius of 40 m). The method consists of estimating leaf deficit at the crown level of each sampled tree. Leaf deficit includes the absence of needles due to drop; twig drop, dried by wind or insects; reduced growth resulting in fewer branches or a smaller individual leaf area due to the phenomenon of microphylla or the impact of insects or foliar diseases. To assess leaf deficit, the appearance of the crown is described by assigning a dieback index. The dieback index in% is calculated by observing the foliar deficit in the crown. In order to limit the interference between phytosanitary problems and symptoms related to competition or lack of light, only dominant or codominant trees were sampled. For each plot, tree density (trees ha⁻¹), diameter at breast height (DBH, cm), height (m), height and crown diameter (m) for 20 trees were measured. Table 1 shows the characteristics of each stand based on the distance from the shoreline and dieback intensity.

2.2.2. Dendroecological data

For the dendroecological study, 15–30 trees were selected in each plot and 2–3 cores were extracted from each tree. Cores were collected

at breast height (1.30 m) with a Swedish increment borer 5 mm in diameter. All cores were placed on wooden supports and specimen details such as site name, sample number and collection date were recorded for identification purposes. These cores were polished using a sanding machine with progressively finer sandpaper (×100 × 180) to obtain a smooth surface.

Cores were visually crossdated with each other by using regional pointer years (missing rings, false rings and narrow rings). The computer program COFECHA (Holmes 1983) was run to examine the statistical quality of the association between the aligned series. COFECHA standardizes raw ring width series and subsequently evaluates the statistical significance of the correlation of each whole series and series segment with a master series by using the Pearson coefficient (r-value) (Grissino-Mayer 2001). The segment length of 20 years lagged by 10 years was used for this study. Placement of missing rings was checked by direct comparison of the wood samples under a microscope and by COFECHA. The frequency of missing rings in each series was calculated for each calendar year. Ring widths were measured from bark to pith with a precision of 0.01 mm using the TSAP Win software and LINTAB® (Rinn, 2003) measuring table in the National Institute for Research in Rural Engineering, Water and Forests of Tunis and in the Mediterranean Institute of Biodiversity and Ecology of Marseille. Ring width series were standardized using the ARSTAN program (Cook and Peters, 1981; Holmes et al., 1986). The long-term trend was removed from each raw chronology by fitting a curve and calculating an index defined as measured ring width for each year divided by the curve-fit value. This made it possible to remove age trend, non-climatic factors and the effects of stand dynamics (Cook and Kairiukstis, 1990). Thus, we applied a double detrending with a negative exponential curve or linear regression equation in the first step and a cubic smoothing spline in the second step.

Finally, for each stand, a non-autoregressive chronology was computed by a technique known as “bi-weight robust estimate of the mean” designed to reduce the influence of isolated outlier values (Cook, 1985).

The ARSTAN program produces three chronologies: standard, residual and arstan. Residual chronology uses autoregressive modeling to remove any signal in the data that carries over from year to year and that may cause an unrealistic increase in the correlation with climate variables (Henderson and Grissino-Mayer, 2009). Because residual chronology showed the strongest and most interpretable relationship, we selected it to model the tree growth to climate relationship (Parker et al., 2014).

Dendrochronological variables calculated by ARSTAN and expressing the climate signal such as mean sensitivity, standard deviation and autocorrelation are good indicators for tree-ring to climate relationship studies (Fritts, 1976) (Table 2).

2.2.3. Climate data

To establish tree growth relationships to climate, reliable climate data close to the sampling sites are crucial. Available climate data from several meteorological stations around the sampling sites were scrutinized for their quality assessments especially in terms of homogeneity, duration and continuity of data. The monthly average of maximal and minimal temperature and monthly total precipitation data from 1950 to 2014 were collected from the two nearest meteorological stations: Bizerte (37°15'N, 09° 48'E) and Kelibia (36° 51'N, 11°05'E).

2.2.4. Tree-growth relationship to climate

A response function analysis (Fritts, 1976) and bootstrap technique (Guiot, 1991) were used to determine the relationship between tree rings and climate with 3PBase software (Guiot and Goeur, 1996). A response function analysis is a multiple regression technique after extracting the main components of monthly climatic data (predictors) to determine the significant climatic variable influencing ring width (predictants). With the bootstrap technique, 200 iterations were done to obtain 200 sets of regression coefficients, 200 multiple calibrations and

Table 1
Characteristics of each stand based on the distance from the shoreline and dieback intensity.

Region	Site name	Stand name	Distance from the shoreline (km)	Dieback index in% (dieback intensity)	Tree density (trees ha ⁻¹)	Age (year)
BIZERTE	RIMEL	BZPH1	3.154	12.66 (Low)	668	54
BIZERTE	RIMEL	BZPH2	1.155	26.27 (moderate)	684	52
BIZERTE	RIMEL	BZPH3	0.105	76.26 (very strong)	660	64
BIZERTE	RIMEL	BZPP1	3.187	13.25 (Low)	557	64
BIZERTE	RIMEL	BZPP2	1.263	26.31 (moderate)	716	90
BIZERTE	RIMEL	BZPP3	0.123	76.25 (very strong)	724	70
KELIBIA	MEZEL BELGACEM	KBPH1	7.464	14.65 (Low)	636	69
KELIBIA	MEZEL BELGACEM	KBPH2	1.725	24.93 (moderate)	652	52
KELIBIA	MEZEL BELGACEM	KBPH3	0.727	76.27 (very strong)	684	64
KELIBIA	MEZEL BELGACEM	KBPP1	7.619	9.53 (Low)	605	62
KELIBIA	MEZEL BELGACEM	KBPP2	1.635	27.67 (moderate)	668	50
KELIBIA	MEZEL BELGACEM	KBPP3	0.680	76.43 (very strong)	700	65

200 verification correlation coefficients. In tree-ring analysis, this method offers the advantage of avoiding errors caused by colinearity among variables and provides a more realistic estimate of tree response to climate. Whether for regression coefficients or for the overall significance of the response function, the different calculated ratios (mean multiple correlation coefficient or mean regression coefficient divided by standard deviation) give an approximate value of the t student coefficient (Gadbin-Henry, 1994).

Analyses were done between residual tree ring chronologies and monthly climatic variables (precipitation, maximal and minimal temperature). These climatic variables were organized according to the biological calendar for the Mediterranean region from October of the previous year to September of the current year (Serre, 1973).

2.3. Data analysis

Mean and standard deviation were calculated for each stand parameter (Mean Height, DBH, Height of the crown, Diameter of the crown and Mean ring width). For missing rings, frequency was calculated for each stand in ten-year increments from 1955 until 2014. An analysis of variance (ANOVA) under SAS software (9.1) was applied with distance from the shoreline as a source of variation to five stand parameters. To test the effect of distance from the shoreline and the effect of time period on missing rings, a two-way analysis of variance (ANOVA) was used. The main effects of distance from the shoreline and time periods were tested for statistical differences using the Newman-Keuls test.

3. Results

3.1. Crossdating

From the available samples, 627 ring width series from 12 stands were crossdated successfully. Statistics for each site are listed in Table 3. Using regional pointer years (missing rings, false rings and

Table 2
Summary statistics from computer program ARSTAN.

Station	Time Span	Mean sensitivity	Standard deviation	Skewness	Kurtosis	Autocorrelation 1 (RESID)	Autocorrelation 1 (STNDRD)
BZPH1	1960–2014	0.29	0.25	0.010	-0.172	-0.044	0.09
BZPH2	1962–2014	0.29	0.24	-0.248	0.412	-0.082	0.12
BZPH3	1950–2014	0.31	0.28	0.129	0.325	-0.009	0.10
BZPP1	1950–2014	0.40	0.36	0.372	-0.380	0.006	0.01
BZPP2	1924–2014	0.38	0.36	0.585	1.305	-0.31	0.27
BZPP3	1944–2014	0.45	0.40	-0.265	-0.06	0.04	0.24
KBPH1	1945–2014	0.41	0.36	1.023	1.415	-0.058	0.02
KBPH2	1962–2014	0.43	0.37	0.334	0.379	-0.035	0.23
KBPH3	1950–2014	0.59	0.48	0.570	0.850	-0.010	0.07
KBPP1	1950–2012	0.43	0.39	0.887	0.423	-0.019	0.08
KBPP2	1962–2012	0.45	0.37	0.036	-0.214	-0.029	0.17
KBPP3	1945–2010	0.47	0.44	-0.133	0.499	0.008	0.25

narrow ring) made crossdating much easier. Mean series inter-correlations were reasonable and gave high values for all sites (0.709–0.980). The number of 20-ring segments with possible problems was low. Comparing site composite skeleton plots and each series with the master series showed clear similarities in growth patterns. However, correlation coefficients for some segments were not statistically significant at 99% but no negative correlation was detected in any segment. The critical correlation coefficient at the 99% confidence level associated with the selected segment length (20 years) was 0.515. For each species at each site, localizing missing rings (Table 3) and false rings (Fig. 2) in all measurement series certified good crossdating.

3.2. Stand characteristics

The results of the analysis of variance (ANOVA) revealed a significant difference in the structural measurement between stands of each species in each forest site. Along the transect, the closer the distance to the shoreline, the greater the decrease in diameter at breast height, tree height, height and crown diameter (Table 4). When comparing the three stands in each transect for each species, trees growing near the shoreline were more sensitive and smaller in radial growth on average. Stands with the highest percentage of dieback index always showed the lowest growth and structural values regardless of stand age. BZPP2 is the oldest stand (90 years) for *Pinus pinea* transect in Bizerte; however, the lowest growth and structural values were recorded in BZPP3 (70 years). KBPH1 is the oldest stand (69 years) for *Pinus halepensis* transect in Kelibia; however, the lowest growth and structural values were recorded in KBPH3 (64 years).

3.3. Missing and false rings

Table 3 shows that missing rings were observed only in stands near the shoreline for both species in both sites. The number of missing rings was higher for *Pinus pinea* than *Pinus halepensis*. The number of series

Table 3
COFECHA results and ring width statistics per site.

	KBPH1	KBPH22	KBPH3	KBPP1	KBPP2	KBPP3	BZPH1	BZPH2	BZPH3	BZPP1	BZPP2	BZPP3
No. of dated series	30	49	30	29	58	84	30	30	74	30	93	90
Time span	65	54	50	60	52	60	55	53	65	65	65	65
Mean series inter-correlation	0.888	0.922	0.959	0.915	0.908	0.864	0.980	0.709	0.802	0.964	0.819	0.845
Segments with possible problems	0	0	0	0	0	3	0	5	7	0	3	5
No. Of missing rings	0	0	36	0	0	291	0	0	17	0	0	240
No. of series with missing ring												
1967			5			15						17
1968						3						
1981			1			15						10
1988			8			67		2				70
1989						41		1				42
1990						10						
1995			4			9		6				17
1998						7						
1999			2			37						1
2001						2						
2002			13			78		8				59
2005												6
2006			3									
2007												4
2008						7						14

Segments with possible problems is the number of 20-ring segments where the correlation with the master series is less than 0.515.

with missing rings could indicate the years when there were more affected trees with missing rings. The results of crossdating showed that in the years 1988–1989 and 2002, there were more affected trees with missing rings.

Fig. 2 shows the distribution of false rings by species and site. At each site, there was a similarity in the distribution of false rings. However, this distribution was often lacking in some stands.

Table 5 shows the statistics on missing rings for the period

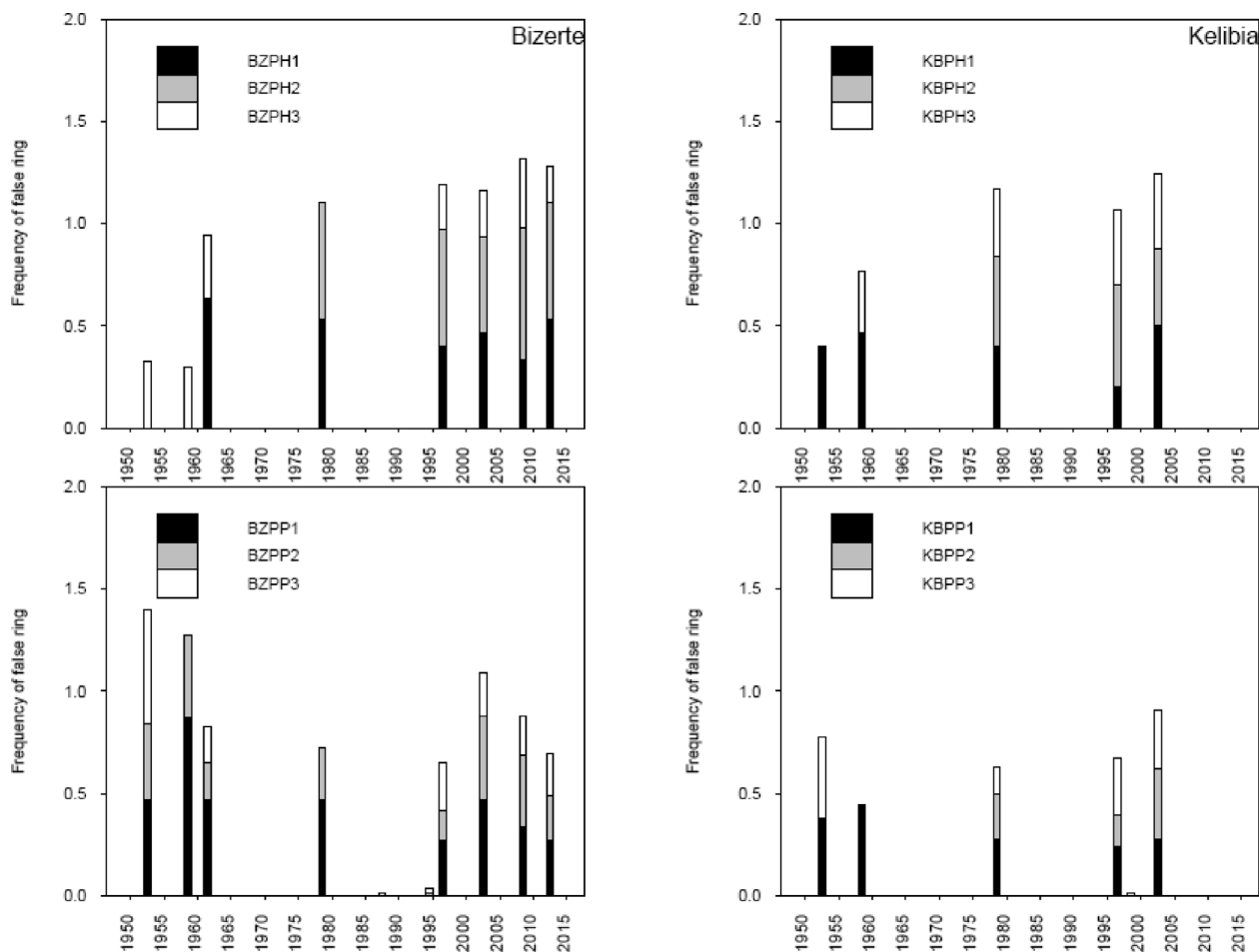


Fig. 2. shows the distribution of false rings by species and site. At each site, there was a similarity in the distribution of false rings. However, this distribution was often lacking in some stands.

Table 4
Analysis of variance (ANOVA) with distance from the shoreline as source of variation to 5 stand parameters.

Chronology name	Common Intervals	DBH (cm)	H (m)	Height of the crown (m)	Diameter of the crown (m)	Mean radial growth (1/100 mm)
BZPH1	1962–2014	33.22 ± 6.03 a	19.96 ± 1.41 a	5.81 ± 1.06 a	8.07 ± 0.60 a	379.03 ± 242.90 a
BZPH2	1962–2014	32.30 ± 4.42 a	17.52 ± 0.89 b	4.65 ± 0.52 b	7.18 ± 0.99 b	321.12 ± 222.96 b
BZPH3	1962–2014	25.41 ± 6.50 b	12.86 ± 1.03 c	1.90 ± 0.31 c	3.02 ± 0.75 c	72.57 ± 37.95 c
BZPP1	1950–2014	33.45 ± 5.67 a	20.17 ± 0.89 a	5 ± 0.96 a	8.01 ± 0.32 a	321.12 ± 317 a
BZPP2	1950–2014	32.05 ± 5.29a	17.83 ± 0.82 b	4.43 ± 0.46 b	7.44 ± 0.95 b	122.68 ± 70.71 b
BZPP3	1950–2014	25.47 ± 4.24b	13.44 ± 0.62 c	1.82 ± 0.20 c	2.94 ± 0.40 c	89.57 ± 107.85 c
KBPH1	1961–2014	34.71 ± 4.34 a	20.13 ± 1.07 a	4.83 ± 0.66 a	7.44 ± 0.63 a	160.37 ± 107.06 a
KBPH2	1961–2014	26.03 ± 4.24 b	15.05 ± 1.18 b	3.61 ± 0.59 b	6.52 ± 0.74 b	167.87 ± 98.34 a
KBPH3	1961–2014	21.38 ± 5.05 c	10.08 ± 1.42 c	2.25 ± 0.32 c	3.22 ± 0.65 c	104.95 ± 82.07 b
KBPP1	1961–2010	36.07 ± 4.11 a	20.66 ± 1.53 a	5.45 ± 0.49 a	8.74 ± 1.15 a	109.38 ± 78.28 a
KBPP2	1961–2010	25.95 ± 4.16 b	14.89 ± 1.14 b	3.60 ± 0.56 b	5.87 ± 0.87 b	217.12 ± 148.92 b
KBPP3	1961–2010	21.84 ± 3.82 c	11.30 ± 1.06 c	2.28 ± 0.33 c	4.45 ± 0.78 c	49.37 ± 33.05 c

Table 5
Mean number of missing rings in *Pinus pinea* and *Pinus halepensis* per decade for various types of canopy damage.

Site	1955–1964	1965–1974	1975–1984	1985–1994	1995–2004	2005–2014
BZPH1	0ab	0ab	0ab	0ab	0ab	0ab
BZPH2	0ab	0ab	0ab	0ab	0ab	0ab
BZPH3	0b	0b	0b	0.40b	1.89b	0b
BZPP1	0b	0b	0b	0b	0b	0b
BZPP2	0b	0b	0b	0b	0b	0b
BZPP3	0b	1.88b	1.11b	12.44a	8a	2b
KBPH1	0ab	0ab	0ab	0ab	0ab	0ab
KBPH2	0ab	0ab	0ab	0ab	0ab	0ab
KBPH3	0ab	0ab	0.33ab	2.66ab	5.66ab	1ab
KBPP1	0ab	0ab	0ab	0ab	0ab	0ab
KBPP2	0ab	0ab	0ab	0ab	0ab	0ab
KBPP3	0ab	2.10ab	1.97ab	15.39a	16.71a	0.92ab

In the same line the followed figures by the same letter indicate the absence of significant differences to the threshold of 5%.

1955–2014 using a variance analysis. For both species, no missing rings were observed for the first period (1955–1964), but the appearance of missing rings was observed from the period 1965–1974 until 2005–2014 but only for trees growing near the shoreline. In contrast, no missing rings were recorded in trees growing far from the shoreline. Their frequency was only statistically significant for *Pinus pinea* in the two forest sites for the periods 1985–1994 and 1995–2004.

3.4. Climatic influence on tree-growth

The tree-growth to climate relationship of the two pine species in both studied forest sites was illustrated using a response function analysis (Figs. 3 and 4). Only response functions for the combination of maximal temperature and precipitation were significant. The calculated ratios (mean multiple correlation coefficient divided by standard deviation) show that all response functions were significant. The climate-growth relationship showed two types of correlation: positive correlation with precipitation and negative correlation with maximal temperature. The monthly variables significantly correlated with growth varied between stands depending on the species and the distance from the shoreline. Most stands showed positive correlation with precipitation between October of the previous year and April of the current year. Concerning the negative correlation with maximal temperature, difference occurred along the transect, between stands for both species in both forest sites. In the Kelibia forest site, both species growing in stands far from the shoreline showed negative correlation only with the March maximal temperature, while in stands near the shoreline both species showed negative correlations with both March and April maximal temperatures. In the Bizerte forest site, negative correlation with maximal temperature was observed in May for both species growing in stands far from the shoreline, in February and March for the *Pinus halepensis* stand and in March and May for the *Pinus pinea* stand near the shoreline.

Fig. 5 shows a mixed graph of missing rings, residual site chronologies, temperature and precipitation based on the results of response function. With this graph, the reason for the increased number of missing rings in the coastal sites can be understood clearly. Dry conditions from October to April and high temperatures from March to April in Kelibia and from February to May in Bizerte seem to be the main climatic elements causing missing rings in both species. In Bizerte, the rainfall deficit was 55% and 50% for 1988 and 2002, respectively, and the increase in maximum temperature was 1.3 °C and 1.4 °C, respectively. In Kelibia, this deficit was 39% and 45% and the increase in maximum temperature was 0.67 °C and 1.67 °C, respectively. Comparisons of site chronologies for each species and site showed clear similarities in growth patterns. Missing rings occurred where rings were usually narrow. This is explained by the fact that, for a year, when a tree shows a missing ring, another tree shows a narrow ring. Missing rings are also included when computing the master series.

4. Discussion

Pines are important components of coastline landscapes in North Africa. Pine forests can be found along a coastal strip of this region. Of the Mediterranean pines which occur at low altitude close to the coast, *Pinus halepensis*, *P. pinaster* and *P. pinea* are the most frequent. *P. pinea*'s natural range is difficult to define because it has been planted widely for so long (Barbero et al., 1998). Pine forests play a crucial role in this region in preventing water erosion by decreasing the effect of erosive forces and keeping the soil in place (Mazzoleni et al., 2004). The extensive root system and large crown make the species resistant to strong winds, which is why *P. pinea* has been used to consolidate sand dunes and protect soil from erosion in exposed coastal areas. Moreover, the multi-layered root system makes it possible for mature trees to obtain water from different levels of the soil and from various sources of water: precipitation, soil water and the water table (Frattegianni et al., 1994).

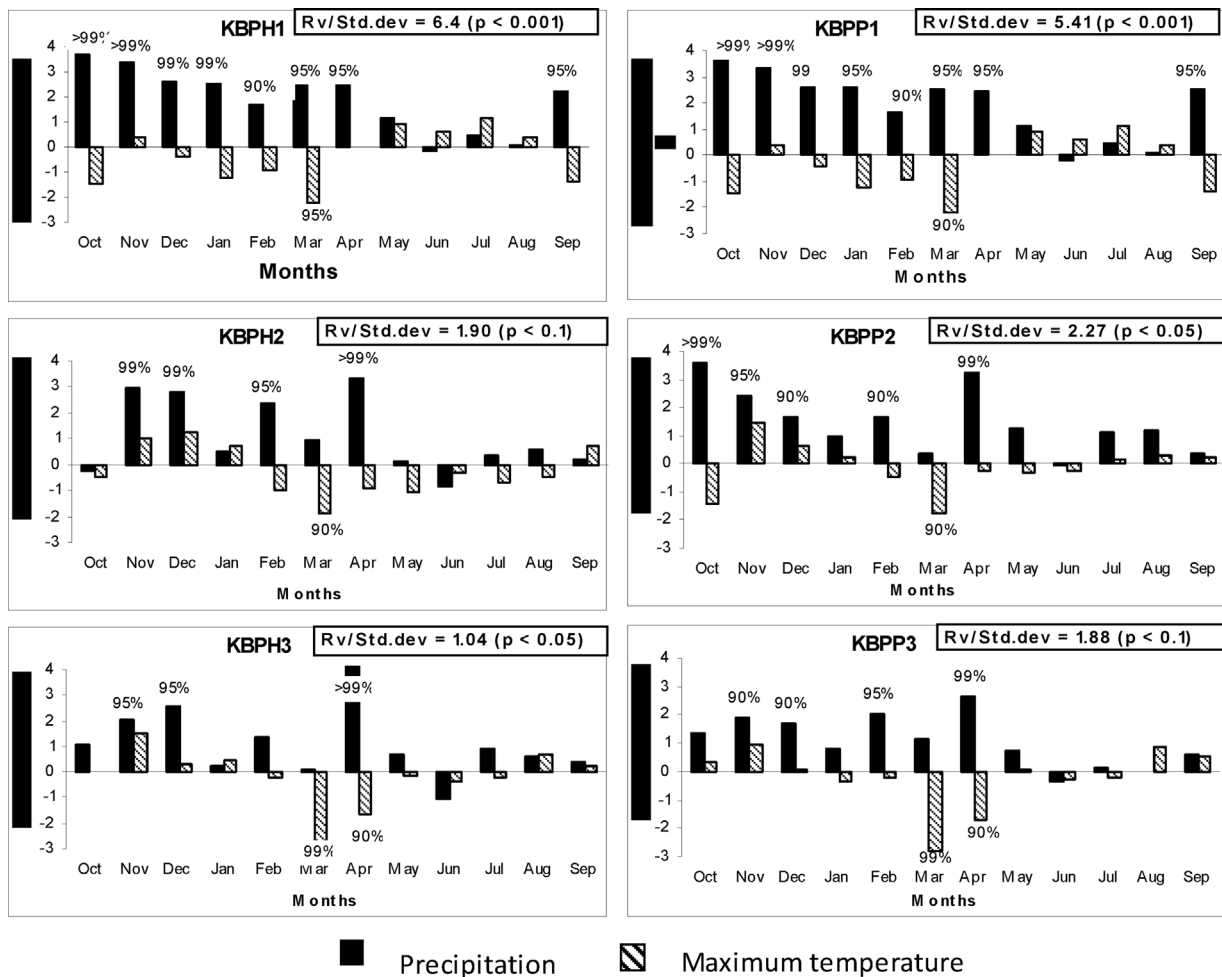


Fig. 3. Response functions for *Pinus halepensis* and *Pinus pinea* in Kelibia with monthly average precipitation and maximal temperature.

In this study, the differences in *Pinus halepensis* and *Pinus pinea* stand structure and growth reflect the effect of the distance from shoreline. Stands with the highest percentage of dieback index always showed the lowest growth and structural values regardless of stand age. Tree age should also be taken into consideration as one of the reasons for differences in growth between stands but its effect is masked by that of dieback intensity and remains minor. However, the effect of age could be important during the first years of life, when even-aged stands of *Pinus spp.* have high radial growth rates. These high rates are only induced by competition from shrubs and understory vegetation because of the light-demanding behavior typical of the pioneer species. However, competition in planted pinewoods is not as strong as in naturally regenerating pines. (Mazza and Manetti, 2013).

The distance from the shoreline influences environmental variables such as mean temperature, water availability, growth period extension, land use, erosion and exposition to sea spray. This characteristic was observed for these species on the Mediterranean coast in Italy (Gellini et al., 1986; Bussotti et al., 1995; Nicolotti et al., 2005; Raddi et al., 2009) and in Spain (Diamantopoulos et al., 2000; Raventos et al., 2001). According to Raddi et al. (2009), growth decline in *Pinus pinea* on the eroded coastal dunes in Italy may be attributed to greater exposure of the crowns to salty winds and surfactants and increased salinity of the water table. On the other hand, *Pinus pinea* is highly sensitive to exposure both to sea spray and to surfactants in marine aerosols (Raventos et al., 2001; Rettori et al., 2005). In the Mezél Belgacem forest, El Ayebe et al. (2004) noted increased Na^+ concentration in leaf tissue when trees were closer to the shoreline (Na^+ increased from 0.54 to 0.62% in *Acacia* and from 0.39 to 2.3% in *Eucalyptus*). A

synergistic effect due to seawater contamination with anionic surfactants was recognized to be the cause of growth decline on the north coast of Tunisia (El Ayebe et al., 2004). Sea-spray containing the surfactants and hydrocarbons had a significant impact on leaves with particular deterioration of the cuticle. Indeed, deleterious effects of Na^+ and Cl^- ions and the pollutants present in the spray on leaf tissue occurred when these substances crossed the cuticular barrier (Bussotti et al., 1997). This phenomenon could explain the reduction of photosynthetic activity in decaying trees and thereafter the growth reduction in trees growing near the shoreline.

The results of this study show that the frequency of missing rings was surprisingly high in *Pinus pinea* compared to *Pinus halepensis*. However, in a semiarid Mediterranean site in Spain, *Pinus halepensis* was shown to be more sensitive to drought than *Pinus pinea* (Novak et al., 2011). These results confirm that, with increasingly severe environmental conditions, production of wood may be partly or totally absent. This might be the case for the trees in our study, where the frequency of missing rings has increased since the 1980s and there have been more and more missing rings in those stands nearest the shoreline. Several factors may be the cause of this increase in missing ring frequency. It may be associated with an increase in water stress, which could be of climatic origin (dry years) or edaphic origin (sandy soils of the littoral dunes). However, dry conditions from October to April and high temperature in late winter and spring were found to be the main climatic factors causing missing rings in both species. In a study on the impact of drought on the growth of three tree species in Tunisia (*Pinus halepensis* Mill., *Pinus pinea* L., and *Pinus pinaster* Sol.), El Khorchani et al. (2007) found a correlation between increased drought and the

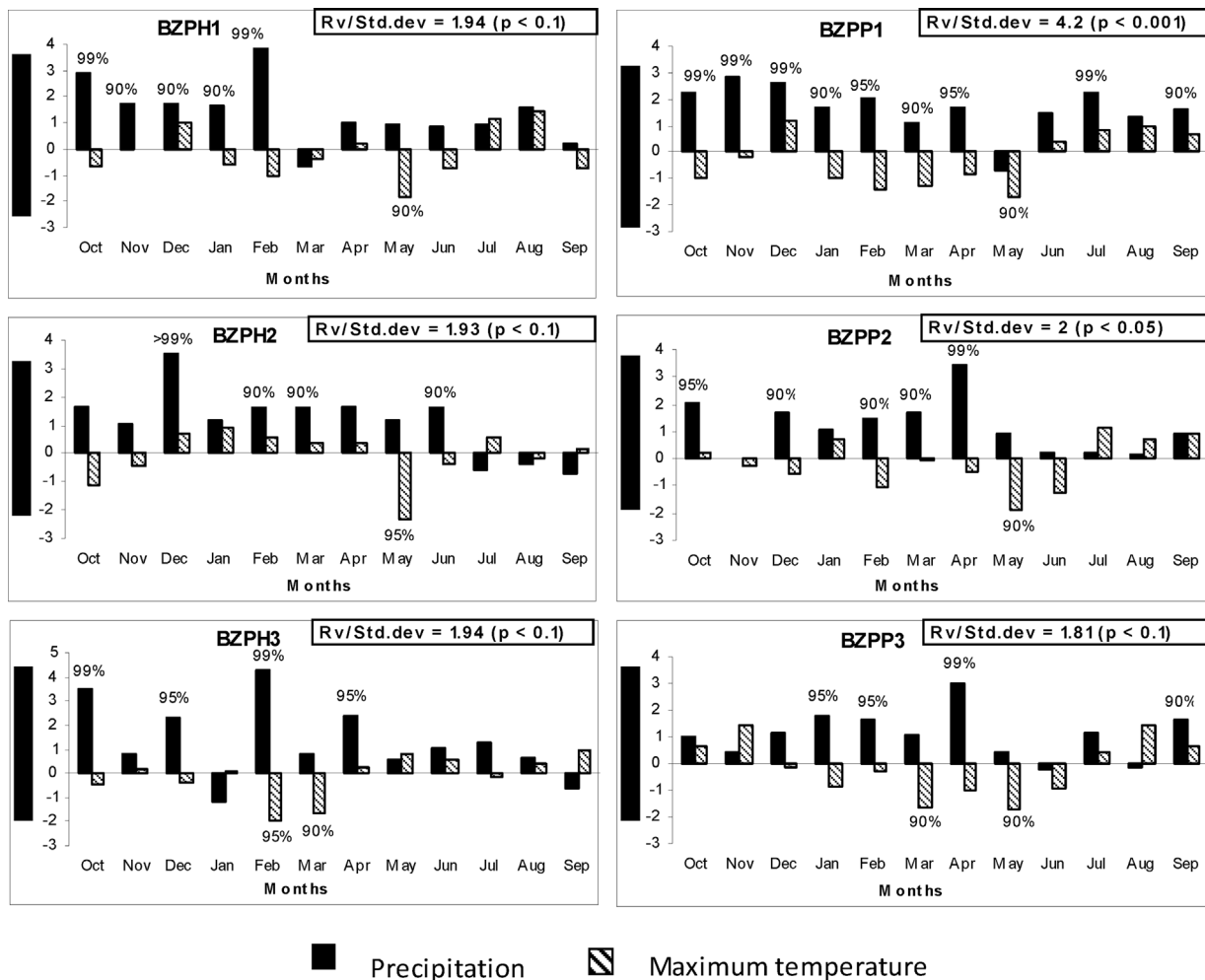


Fig. 4. Response functions for *Pinus halepensis* and *Pinus pinea* in Bizerte with monthly average precipitation and maximal temperature.

appearance of narrow rings in the period from 1978 to 2001 compared to the 1954–1977 period. [de Luis et al. \(2009, 2010\)](#) showed that the first potential cause of an observed increase of missing rings could be related to increased stress conditions due to rising temperatures and decreased precipitation recorded in Spanish Mediterranean areas since the mid-1970s. Other dendrochronological studies on *Pinus pinea* and *Pinus halepensis* from semiarid coastal sand dunes in southeastern Spain have showed a high frequency of missing rings in recent years due to dryness and other disturbance factors such as marine spray ([Raventos et al., 2001; Novak et al., 2011](#)).

In addition to climate, other stressors might contribute to the frequency of missing rings. The coastal vegetation in north Tunisia is harmfully affected by chemical components contained in marine spray ([El Ayeb et al., 2004](#)). As reported in other parts of the Mediterranean coast in Spain ([Raventos et al., 2001](#)) and in Italy ([Bussotti et al., 1995](#)), the appearance of missing rings might accompany alteration or death of needles, decreased photosynthetic production and reduced availability of assimilates for wood production ([Raddi et al., 2009](#)).

Trees on coastal sites also often suffer from underground problems (e.g. root damage). This problem has become more common on sandy coasts throughout Mediterranean regions due to the filtering factor which limits the water retention capacity of the sandy soils of the coastal dunes where the availability of water could decrease abruptly during dry years. Accordingly, underground problems are associated with disturbed water and mineral uptake, leading to reduced wood production ([Mc Laughlin et al., 2002](#)).

A missing ring in a particular year should not endanger tree survival ([Cherubini et al., 2002; Sass-Klassen et al., 2008](#)). However, long-term

disturbances of cambial activity often lead to physiological problems associated with disturbed water transport and reduced storage capacity of reserve assimilates ([Bigler et al., 2004; Wunder et al., 2008](#)).

The relationship between tree-ring index and precipitation from October to April shows that water availability is the main factor influencing tree growth of both pine species in Tunisia. These results are consistent with several previous dendroecological studies in Mediterranean regions ([Raventos et al., 2001; El Khorchani, 2006; Campelo et al., 2007; Raddi et al., 2009; Mazza and Manetti, 2013](#)) and showed that a large part of a tree ring is formed during the late winter and spring. These findings reflect the behavior of evergreen conifers that grow in regions with mild winters and are characterized by a photosynthetic activity in winter that produces carbohydrates for early wood formation in the following year ([Kozłowski et al., 1991](#)). The number of months that precipitation is positively correlated with growth is always less for trees growing near the shoreline than for those growing in distant stands. Thus, low amounts of precipitation over the past months in Tunisia, which is the only available freshwater, has reduced soil water content above the water table. The degree to which trees regulate water loss during drought may explain patterns of carbohydrate production and subsequent susceptibility to drought or biotic attack ([McDowell et al., 2008](#)). Stomatal responses to drought vary from drought avoidance, in which stomata close at a threshold water potential to minimize further transpiration, to drought tolerance, in which stomatal closure is less severe and transpiration continues at relatively high rates ([McDowell et al., 2008](#)).

The negative relationships with maximal temperatures show that the number of months that maximal temperature is correlated

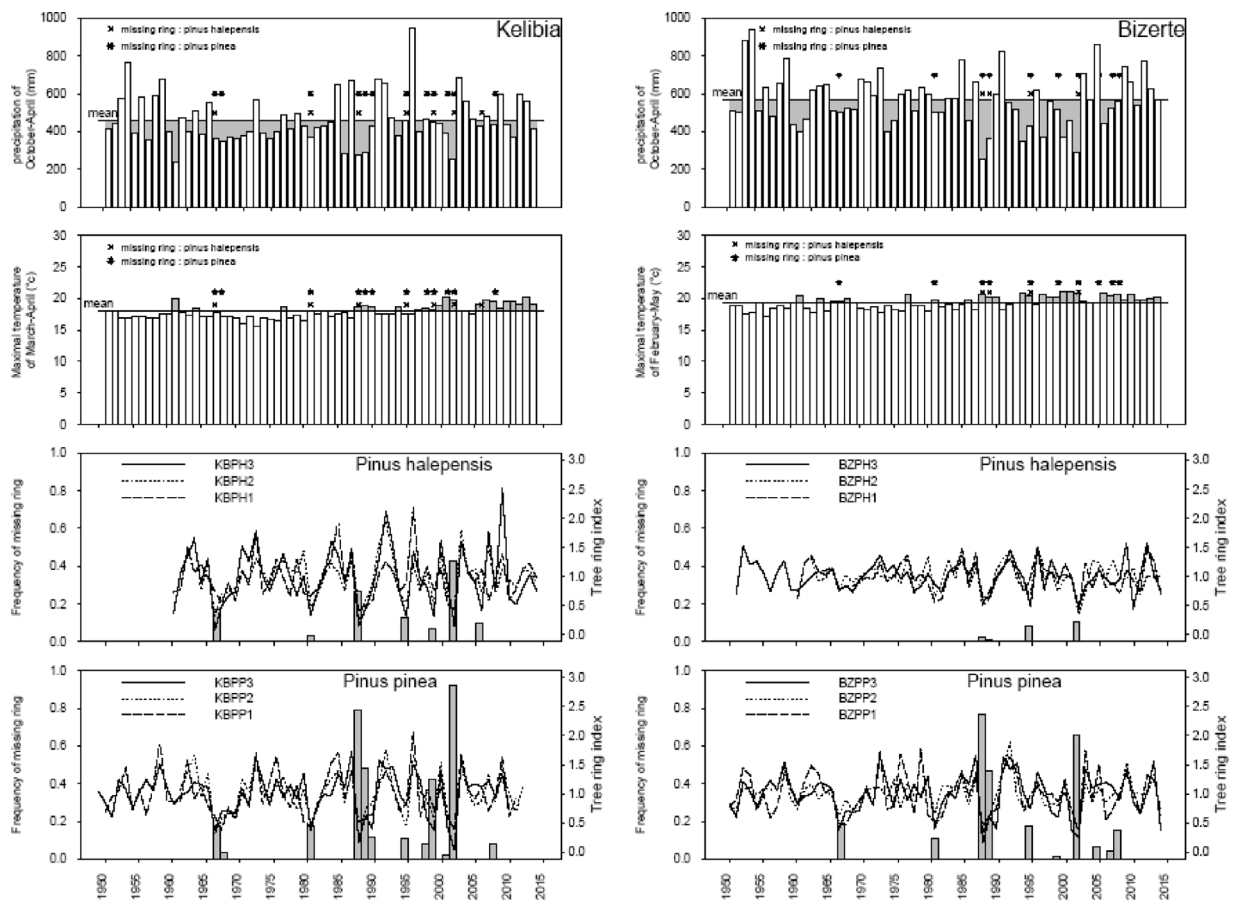


Fig. 5. Mixed graph for missing rings, temperature and precipitation based on the result of response function.

negatively with growth is always higher for trees growing near the shoreline (2 months) than for trees growing in distant stands (1 month). This observation could be interpreted by the possibility of intense water stress for trees growing near the shoreline. However, precipitation remains the primary explanatory variable of tree growth decline. High temperatures might also affect water shortage because of increased evapotranspiration and soil water evaporation (Campelo et al., 2006). Furthermore, recent research indicates that warmer temperatures alone can increase forest water stress, independent of the amount of precipitation (Barber et al., 2000). Moreover, on coastal sand dunes where water availability could suddenly decrease in dry years, the negative effect of maximal temperature on radial growth in the spring could accentuate the phenomenon of water stress (Aussenac, 1976; El Khorchani et al., 2007). Water availability still remains a factor more limiting than temperature. Thus, it might be the case that *Pinus pinea*, like other Mediterranean species (e.g. *Pinus halepensis*), responds by becoming increasingly dependent on moisture from deeper soil horizons that have not been exhausted in previous years (Sarris et al., 2007). In a study on the process based model shedding light on climate sensitivity of Mediterranean tree-ring width, conducted on *Pinus halepensis* from semi-arid areas in Tunisia, Touchan et al. (2012) showed that the average duration of the growing season is 191 days, and that on average soil moisture limits tree-ring growth for 128 days and temperature for 63 days in that 191-day window. Due to water deficiency and subsequent stomata closure, dryness causes a decrease in photosynthesis and consequently fewer carbohydrates are available for radial growth (Sitte et al., 1999).

The combination of response function, dendrochronological measurements and structural characteristics of *Pinus halepensis* and *Pinus pinea* stands has shown that distance from the shoreline influences environmental conditions. The closer the trees are to the shoreline, the

more they are put under stress conditions. Among these stress conditions, drought can operate as a trigger (“inciting factor”) that may ultimately lead to mortality or dieback in trees that are already under stress (by “predisposing factors” such as old age, poor site conditions and air pollution) and succumb to subsequent stem and root damage by biotic agents (“contributing factors” such as wood-boring insects and fungal pathogens) (Manion and Lachance, 1992).

Dieback of pine forests involves multiple interacting factors, ranging from particular sequences of climate stress and stand life histories to insect pests and diseases (Manion, 1981; Miao et al., 2009). Of the many factors of climate stress, drought is a concern affecting forests in coastal zones. This type of damage causes long-term degradation that may predispose trees to attack by pathogens and pests. Forests may become ecologically unstable, reducing their efficiency to protect coastal dunes and increasing forest fire risks due to the abundance of dry canopies.

5. Conclusion

The northeastern pine coastal woodland of Tunisia has great ecological and economic importance in protecting the soil from erosion and controlling water shortage. The major threats to the coastal landscapes of Tunisia, are common along the Mediterranean coastal zone, and related to uncontrolled development, urbanization, increasing national and international tourism and unplanned or over-exploitation of natural resources. Depletion and degradation of resources, including freshwater, forests and coastal areas, is a growing concern, as is pollution within the region itself (Vogiatzacos and Cassar, 2007). Our study indicates that *Pinus pinea* and *Pinus halepensis* on subhumid Mediterranean coastlines are growing under stress conditions when trees are close to the shoreline. These stress conditions are manifested

by a negative effect on radial growth and structural characteristics of the trees, namely the development of the crown and the needles which can lead to conditions of dieback. Several climate models indicate that drought frequency and intensity are expected to increase in the Mediterranean basin (Giorgi and Lionello, 2008; Lelieveld et al., 2012; Ozturk et al., 2015). In sensitive areas such as the coastal dunes of Rimel and Menzel Belgacem, maximum temperature is significantly negative in late winter and spring, high precipitation from autumn of the previous year to mid-April of the current year has a significantly positive influence on tree-ring width. Long-term low precipitation and high temperature during these months in Tunisia influence the availability of water resources and cause drought stress. The higher water stress induced by an increase in air temperature and a decrease in rainfall for successive years could seriously reduce growth accompanied by an increased number of missing rings of the Mediterranean pine species, despite their drought-tolerant behavior. Such conditions trigger physiological disturbances in trees and predispose them to be attacked by pathogens and pests and thus accelerate their decline. The *Pinus pinaster* decline observed in recent years on northern coastal dunes of Tunisia could be the consequence of this imbalance caused by such factors (drought, sea spray...). Our results suggest that stands growing in stressed conditions need different management approaches in order to achieve sustainability.

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