

Research Article

Turk J Bot 34 (2010) 311-322 © TÜBİTAK doi:10.3906/bot-0906-64

Nitrogen and phosphorus resorption in a salt marsh in northern Turkey

Yasemin BEDİRKURUM ÇAKIR¹, Tuğba ÖZBUCAK^{2,*}, Hamdi Güray KUTBAY¹, Duygu KILIÇ³, Ali BİLGİN⁴, Rena HÜSEYİNOVA¹

 1 Ondokuz Mayıs University, Faculty of Science & Arts, Department of Biology, Samsun - TURKEY 2 University of Ordu, Faculty of Science & Arts, Department of Biology, 52750 Ordu - TURKEY 3 Bilsem High School, Amasya - TURKEY

⁴Rize University, Faculty of Science & Arts, Department of Biology, 53100 Rize - TURKEY

Received: 30.06.2009 Accepted: 18.04.2010

Abstract: This study was conducted to investigate foliar nitrogen and phosphorus dynamics, nitrogen and phosphorus resorption, and some leaf traits (i.e. SLA=specific leaf area) in some coastal salt marsh plants located in northern Turkey. The study area is located on the east bank of the Kızılırmak River in the central Black Sea Region. There were statistically significant differences among months (from June to October) in terms of SLA in all species except for *Artemisia santonicum*. However, no significant differences were found with respect to P in all species. The highest N and P resorption efficiencies were found in *Salicornia prostrata*, whereas the lowest resorption efficiencies were found in *Euphorbia paralias*. P resorption proficiency is biochemically complete in all species. N resorption proficiency is biochemically complete except for *E. paralias*. Additionally, Grime's CSR (C: competitive; S: stress-tolerant; R: ruderal) strategies were also investigated and it was found that all of the species exhibit competitive ruderal strategy.

Key words: Coastal salt marshes, Grime's strategies, resorption efficiency and proficiency

Kuzey Türkiye'deki bir tuzlu bataklıkta azot ve fosfor geri taşınımı

Özet: Bu çalışma Türkiye'nin Orta Karadeniz Bölgesinde bulunan Kızılırmağın doğu kısmında bulunan bazı kıyı tuzlu bataklık bitkilerinde yaprak azot ve fosfor dinamikleri, azot, fosfor ile yaprak özelliklerini (spesifik yaprak alanı SLA gibi) incelemek için gerçekleştirilmiştir. İstatistiksel olarak, *Artemisia santonicum* hariç bütün türlerde SLA yönünden aylar (Haziran-Ekim) arasında önemli farklılıklar bulunmuştur. Bununla beraber, bütün türlerde P içeriği yönünden önemli farklılıklar bulunmuştır. En düşük N ve P geri emilim yeterliliği *E. paralias*'de bulunurken, en yüksek N ve P geri emilim etkinliği *Salicornia prostrata*'da bulunmuştur. P yeterliliği bütün türlerde biyokimyasal olarak yeterlidir. N yeterliliği ise *Euphorbia paralias* hariç bütün türlerde biyokimyasal olarak yeterlidir. Ayrıca, Grime'in CSR stratejileri (C: rekabetçi; S: strese-dirençli; R: ruderal) de incelenmiş ve bütün türlerin Grime'in stratejilerinden rekabetçi ruderal stratejisi gösterdiği saptanmıştır.

Anahtar sözcükler: Kıyı bataklıklar, Grime'in stratejileri, geri taşınım etkinliği ve yeterliliği

^{*} E-mail: tsiozbucak@hotmail.com

Introduction

Salt-affected soils are commonly found in arid landscapes where evapotranspiration exceeds precipitation throughout most of the year and may occur naturally or due to human activities (Aschenbach, 2006). Coastal salt marshes comprise areas of land bordering the sea and lakes that are largely covered with vegetation and subject to periodic tidal inundation. These areas have certain qualities, related to the proximity to the sea or lake, that distinguish them from inland salt marshes (Asri & Ghorbanli 1997; Onaindia & Amezaga, 1999). Such ecosystems are generally dominated by halophytic species that have developed mechanisms to avoid and resist salt stress, such as salt glands, osmotic adjustment, increased growth, and selective ion uptake by roots (Kruger & Peinemann 1996; Allahverdiev et al. 1998; Kutbay & Demir, 2001).

Grime et al. (1997) suggested that mineral nutrients, especially nitrogen and phosphorus, should be considered as the major regulating factors of vegetation processes at scales ranging from individuals to ecosystems. Foliar resorption is an important mechanism of nutrient conservation, recycling 50% of maximum foliar N and P content in a wide range of perennial life-forms (Aerts, 1996; Kobe et al., 2005). The rate of nutrient resorption from senescing leaves may also vary with the availability of nutrients in a certain habitat. This implies that, in addition to leaf fall patterns, leaf chemistry can further amend the time-dependent controls on nutrient losses (Niinemets & Tamm, 2005).

Nitrogen and phosphorus are largely withdrawn from senescing leaves before abscission, and they are used for new growth or stored in vegetative tissue until the next growing season. Nutrient resorption efficiency (NRE) is defined as the proportion of a nutrient withdrawn from leaves before leaf abscission (van Heerwaarden et al., 2003; Sans-Pérez et al. 2009).

A new measure of resorption was introduced by Killingbeck (1996) as resorption proficiency. Killingbeck (1996) emphasized resorption proficiency (the content of a nutrient in senesced leaves) over resorption efficiency as proficiency is not subject to temporal variation in nutrient content in green leaves

and timing of sampling (Lajtha 1987; Zotz 2004; Kobe et al., 2005; Yasumuro et al. 2005; Yuan et al., 2005). Proficiency is simply the amount of a nutrient that remains in fully senesced leaves. From a biological perspective, an important advantage of measuring resorption as proficiency rather than efficiency is that proficiency is a more unequivocal measure of the degree to which selection has acted to minimize nutrient loss in ephemeral leaves (Killingbeck, 2004).

Nitrogen and salinity are 2 key factors that have been shown to be correlated with vegetation pattern and zonation in coastal salt marshes. Nitrogen is frequently the major nutrient limiting growth in salt marshes. Halophytes have been demonstrated to accumulate soluble nitrogenous compounds when subjected to stress, such as salinity (Skeffington & Jeffrey, 1988). Although nitrogen and phosphorus resorption have a great importance in salt marshes, the number of studies on nitrogen and phosphorus resorption in salt marshes are very scarce (Shaver & Melillo, 1984; Cartaxana & Catarino, 2002).

The present study addresses 3 main objectives: (a) to show foliar nitrogen and phosphorus dynamics in some coastal salt marsh plants, (b) to determine nitrogen and phosphorus resorption efficiency and proficiency in some coastal salt marsh plants, and (c) to classify salt marsh plants according to Grime's CSR strategies.

Materials and methods

The study area

This study is conducted on the east bank of the Kızılırmak River. It is located in the northern and north-eastern parts of Bafra District (41°42′17″ and 41°31′46″′N, and 36°05′40″ and 36°03′01″E) in the central Black Sea Region (northern Turkey).

The mean annual temperature and mean rainfall from June 2005 to October 2005 were 13.7 °C and 806.4 mm, respectively. The maximum temperature for the warmest month (July) was 33.8 °C and the minimum temperature for the coldest month (February) was –2.6 °C during the study period. The pluviometric quotient (Q) is 76.7, and the mean annual relative humidity is 79.81%. On the basis of these data and the classification scheme of Daget

(1977), the area has a semi-humid Mediterranean climate, whereas the precipitation regime is central Mediterranean (Winter, Autumn, Spring, and Summer) (Turkish Ministry of Agriculture, 2002). Taxonomic nomenclature in the present study follows that of Brummitt and Powell (2001).

The mean annual rainfall in the study area is more than is expected for an arid environment. However, it is substantially decreased during summer, and average monthly rainfall values are 39, 36, and 29.5 mm in June, July, and August, respectively. The mean temperatures in June, July, and August are 20, 23, and 23.3 °C, respectively. As a result of this excess evaporation that occurs during summer, the potential evapotranspiration rate is 19 mm during winter, whereas the potential evapotranspiration rate is 58 mm during summer. Additionally, the highest potential evapotranspiration rate is found in June (102 mm) and exceeds rainfall (Köksal, 1972; Engin et al., 1988). These high evaporation rates reduce the natural accumulation of surface water (Wadie, 2002). The Mediterranean climate is characterized by irregular and intense rainfall events and a harsh, dry summer period (Álvarez et al., 2000). Inundation also takes place in the study area and dissolved salts have been raised to the surface. As a result of this, saline soils are dominant in the study area (Avcı et al., 2001).

Two different plant groups occurr in the study area. One of them is halophytic species around salt pans. This group is represented by *Salicornia prostrata* Pall. (Chenopodiaceae) and *Spergularia marina* (L.) Gris. (Chenopodiaceae). The other group is salt pandune transitional species and this group is represented by *Artemisia santonicum* L. (Asteraceae), *Otanthus maritimus* (L.) Hoffmans. & Link (Asteraceae), and *Euphorbia paralias* L. (Euphorbiaceae).

Sampling and chemical analysis

This study was carried out during the growth period from June 2005 to October 2005 (June, July, August, September, and October) including 1 sampling date. On each sampling date 4 quadrats (0.25 m \times 0.25 m) were randomly selected and 7 leaf samples from each species were taken. Leaf area was calculated by a Placom digital planimeter (Cornelissen et al., 1997; Kutbay, 2001). For *S. prostrata*, due to cylindrical shape of leaves, surface

area was calculated as $\pi \times$ measured leaf area values (Cartaxana & Catarino, 2002). Leaf samples were dried at 70 °C until constant weight, and then ground and sieved. Samples were-digested in a mixture of nitric and perchloric acids prior to phosphorus (P) analysis. Nitrogen (N) was determined by the micro Kjeldahl method with a Kjeltec Auto 1030 Analyser (Tecator, Sweden) after digesting the samples in concentrated H_2SO_4 with a selenium catalyst._P was determined with stannous chloride method using a Jenway spectrophotometer.

N and P resorption efficiency was calculated as $[(NG - NS) / NG] \times 100$, where NG and NS are either N or P contents in mature green-leaf (G) and senescent leaf (S), respectively. Resorption proficiency was the concentration of a nutrient in senesced leaves (Killingbeck, 1996).

Grime's CSR (C: competitive; S: stress-tolerant; R: ruderal) strategies were used to compare both sympatric species in terms of N and P usage strategies using a program developed in Microsoft Excel. Canopy height, dry matter content, flowering period, the onset of flowering, lateral spreading, dry leaf weight, and specific leaf area were used to find Grime's strategies for the studied species (Hodgson et al., 1999).

Statistical analysis

In order to determine if monthly changes in N (%), P (%), and SLA are statistically significant, we performed ANOVA tests on each species. The differences among the sampling periods regarding to the contents of N and P and the changes of SLA were tested by analysis of variance. Significant differences between stages and variables were determined by the Tukey's honest significant difference (HSD) test. All of the tests were performed at the significant level of α 0.001 with the SPSS Version 10.0.

Results and discussion

All of the species were characterized as CR (competitive ruderal strategy) strategy using the criteria presented in Table 1.

The studies on resorption efficiency of salt marsh species are rare and P resorption efficiency values have not been reported. N resorption efficiency values

Table 1. Allocation procedure for competitive, stress-tolerant, and ruderal (C-S-R) plants (based on Hodgson et al. 1999).

Variable			Definition	
Canopy Height	Six-point classification	1	1-49 mm	
		2	50-99 mm	
		3	100-299 mm	
		4	300-599 mm	
		5	600-999 mm	
		6	>999 mm	
Dry Matter Content	Mean of percent dry matter content in the largest, fully hydrated, full expanded leaves (%)			
Flowering Period	Normal duration of flowering period (months)			
Flowering Start	Six-point classification	1	First flowering in March or earlier	
		2	in April	
		3	in May	
		4	in June	
		5	in July	
		6	in August or later, or before leaves in spring	
Lateral Spread	Six-point classification	1	Plant short-lived	
		2	Loosely tufted ramets radiating about a single axis, no thickened root-stock (in graminoids)	
		2	Compactly tufted about a single axis, no thickened rootstock (in non-graminoids)	
		3	Compactly tufted ramets appressed to each other at base (in graminoids)	
		3	Compactly tufted about a single axis, thickened rootstock present (in non-graminoids)	
		4	Shortly creeping, < 40 mm between ramets	
		5	Creeping, 40-79 mm between ramets	
		6	Widely creeping, > 79 mm between ramets	
Leaf Dry Weight	Natural logarithm of mean dry weight in the largest, fully hydrated, fully expanded leaves (mg) plus 3			
Specific Leaf Area	Mean of area/dry weight quoitent in the largest, fully hydrated, fully expanded leaves (mm2/mg)			

were similar to other salt marsh species. However, P resorption efficiency values were rather low as compared to non-halophytes (Aerts 1996; Cartaxana & Catarino, 2002). The highest N and P resorption efficiencies were found in *S. prostrata*, whereas the lowest resorption efficiencies were found in *E. paralias*. N and P resorption efficiencies were higher in species found in salt pans compared to species found in the salt pan-dune transitional zone (Table 2).

Table 2. N and P resorption efficiency (%) in studied species.

Species	N resorption efficiency (%)	P resorption efficiency (%)
Salicornia prostrata	66.66	50.80
Spergularia marina	64.86	35.89
Artemisia santonicum	37.50	26.18
Otanthus maritimus	44.82	33.82
Euphorbia paralias	31.08	34.21

The highest N content was found in October in S. prostrata. N content was decreased from August to September in S. marina, whilst it was increased from August to October in A. santonicum. Monthly changes of N content in O. maritimus were somewhat similar to A. santonicum. N content was considerably higher in E. paralias as compared to the other species studied. Overall, the lowest N was found in June in all species except for S. marina in which the lowest N contents were found in October (Figure 1).

The highest P content was found in June in S. prostrata. P content of S. marina was remarkably decreased in October. The changes in terms of N and P contents from June to August were similar in A. santonicum. However, P content was slightly increased in October in A. santonicum. P content of E. paralias was decreased from June to August and then increased from August to October. The lowest P contents were found in October in S. marina and O. maritimus, while the lowest P contents were found in July in S. prostrata, in June in A. santonicum, and in August in E. paralias. P content of O. maritimus was constant from June to July, whilst increased in August and after August it had a decreasing tendency (Figure 2).

SLA was decreased from June to October in *S. prostrata*. The highest SLA was found in July in *S. marina*. SLA was decreased in September and increased in October in *A. santonicum*. The changes of SLA in *O. maritimus* were similar to *A. santonicum* like N content. SLA in *E. paralias* was increased from June to October. Except for *E. paralias*, SLA was decreased during September in all species (Figure 3). For example SLA was also decreased over the growth period in *S. prostrata*. SLA may be decreased due to the contraction of cell for loss of water (Wang & Lin, 1999).

The content of elements in the leaves of halophytes under saline conditions changes during leaf development and senescence (Wang & Lin, 1999). N contents (%) were increased during autumn in all species except for *S. marina*. In other words, "accretion" was observed during autumn (September and October) because vegetative and generative plant parts have been competed with each other during autumn (Duchesne et al., 2001). Accretion was only found in *E. paralias* with respect to P (%) contents.

N dynamics and N resorption efficiency values was similar to those of the other salt marsh species. Cartaxana and Catarino (2002) reported 31%–76% resorption efficiency for salt marsh species in Portugal. In the present study, resorption efficiency for all species varied between 31%–67%. No data were available on P resorption efficiency. However, P resorption efficiency of the species studied was similar to those of evergreen species (Mayor & Rodá, 1992).

N and P contents in the present study were rather low. In most salt marshes, nitrogen is very limited. This is in accordance with competition among saline ions (Ca⁺², Mg⁺², K⁺, Na⁺, Cl⁻, SO₄²⁻, HCO₃⁻) and N and P (Donovan et al., 1997; Kutbay & Demir, 2001). Increase in salinity may decrease a plant's ability to absorb essential nutrients (Yeo, 1983; Marschner, 1995; Aschenbach, 2006).

Killingbeck (1996) stated that resorption is highly proficient in plants that reduce nitrogen and phosphorus concentrations below 0.7% and 0.05%, respectively, during their senescent stages. According to this threshold values, P resorption proficiency is highly proficient in all species. N resorption proficiency is also highly proficient except for *E*.

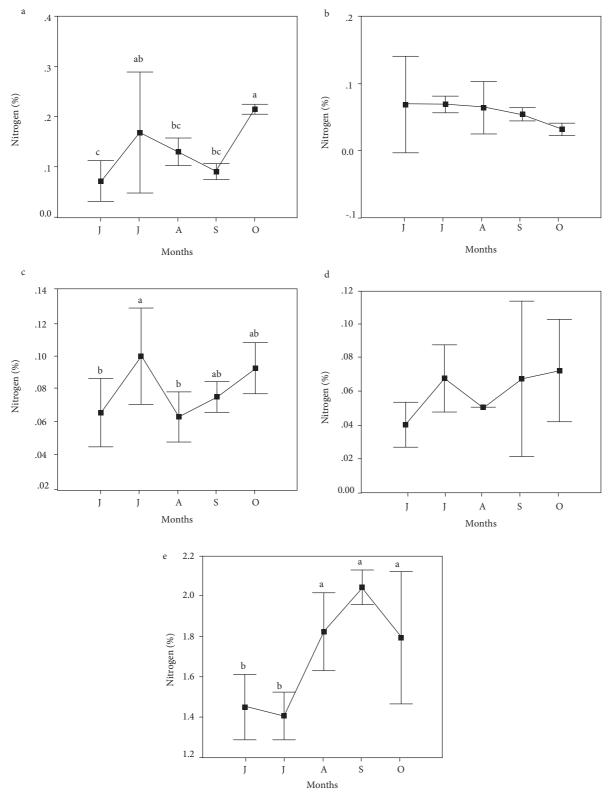


Figure 1. Seasonal changes in N (%) contents of the leaves of (a) *S. prostrata*, (b) *S. marina*, (c) *A. santonicum*, (d) *O. maritimus*, and (e) *E. paralias* (Different letters indicate significant differences between the groups according to Tukey's HSD test) (rejection level 0.05). Error bars depict standart errors of means (n = 7).

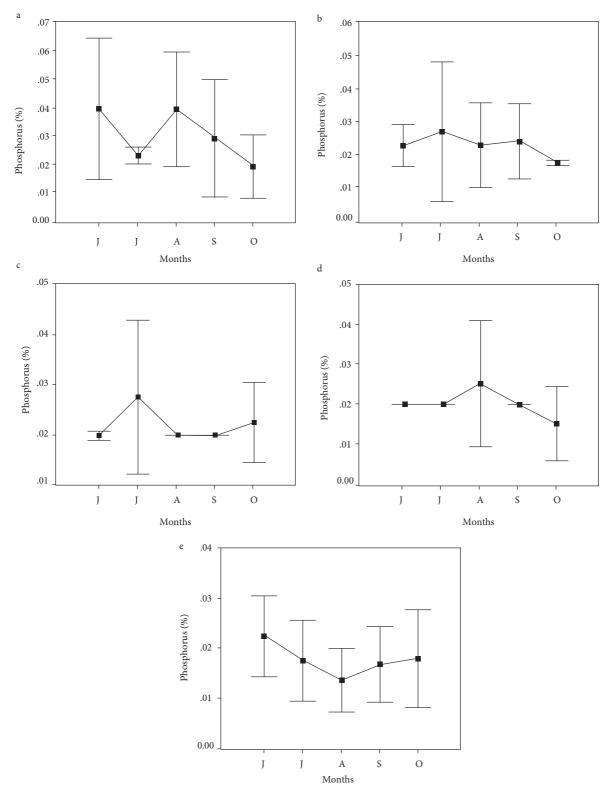


Figure 2. Seasonal changes in P (%) contents of the leaves of (a) *S. prostrata*, (b) *S. marina*, (c) *A. santonicum*, (d) *O. maritimus*, and (e) *E. paralias* (Different letters indicate significant differences between the groups according to Tukey's HSD test) (rejection level 0.05). Error bars depict standart errors of means (n = 7).

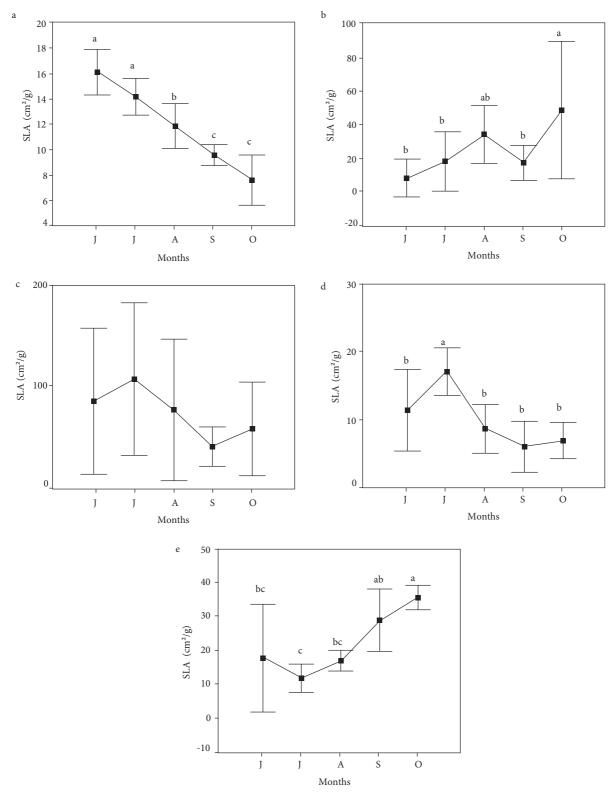


Figure 3. Seasonal variations in SLA of the leaves harvested from (a) *S. prostrata*, (b) *S. marina*, (c) *A. santonicum*, (d) *O. maritimus*, and (e) *E. paralias* (Different letters indicate significant differences between the groups according to Tukey's HSD test) (rejection level 0.05). Error bars depict standart errors of means (n = 7).

paralias. This species is a salt pan-dune transitional species and N content was higher than the other species. In addition to this, the lowest N resorption efficiency was found in *E. paralias*.

Ungar et al. (1979) and Munir & Aftab (2009) reported that Salicornia species were highly competitive in highly saline soils, because they achieved an optimal growth at salinity levels that would be limiting to other species. Yeo (1983) stated plants that are able to cope with harsh conditions benefit from high availability of water, light, and nutrients, compensate for the "extra cost for tolerance", and produce a large amount of biomass. Salicornia L. and Spergularia (Pers.) J. & C. Presl species were also classified as high phenotypic plasticity species (Wilkoń-Michalska, 1985). The studied species were characterized as Grime's CR (competitive ruderal strategy). CR species usually occurred in seasonally inundated and arid areas. Competitive species, i.e. species with dominant competitive ability (C s.l. strategy: C+CR+CS), which presented high vegetative development, ecological plasticity, and occasionally some allelopathic potential (Vidal et al., 2000). In the study area inundation occurred periodically and dryness was observed due to excess evaporation during summer (Kutbay & Demir, 2001). Environmental conditions in these areas were harmful only once during the life cycle of a certain species. However, they never cause completely removing of the species (Grime, 2002). Plant zonation in coastal salt marshes depends upon competitive processes with poorer competitors being displaced to areas of greater disturbance or stress, which provide refuges from superior competitors in addition to physicochemical factors (Reynolds et al., 2001).

Disturbances as a result of wrack deposition and sedimentation are also significant in salt marshes (Levine et al., 1998; Pennings & Richards, 1998). Because substrate salinity in coastal marshes varies through the growing season in relation to weather conditions and tidal activity (Adam, 1990), there is a high probability of interactions between the temporal pattern of substrate salinity and plant phenological development. Such interactions are likely to affect plant population abundance and distribution in coastal marshes, particularly for annuals that must

necessarily go through germination and establishment phases every year (Reynolds et al., 2001). SLA was increased from August to October in *E. paralias*. In addition, SLA was also increased in *S. marina*, *A. santonicum*, and *O. maritimus* from September to October. Vegetative growth period of these species may be occurred several times during the year and new leaves emerge through autumn.

Plant zonation in salt marshes is affected by physical stress and nutrient limitation. Salinity and sodium toxicity act as ecological filters that reduce species richness in salt marshes (Flowers et al., 1977; Yeo, 1983; García et al., 1993, El Demerdash, 1996 & Abd El-Ghani, 2000). In addition, inundation causes stressful conditions in salt marshes, especially near sea level (Curcó et al., 2002). The notable aspects of seasonal fluctuations in terms of salinity due to excess evaporation occur during summer in salt marshes (Apaydın et al. 2009; Ababou et al., 2010). Notable changes especially in July in the studied plant characters may probably be due to increasing salinity during summer as a result of excess evaporation. Owing to the strong environmental gradients present in salt marsh habitats, marked intra- and inter-specific variation is probably the rule for all species of salt marsh plants (Richards et al., 2005). Jiang et al. (2009) stated that salt marsh plants subject to seasonal variations with respect to specific leaf area owing to the strong environmental gradients especially during summer.

It has been proposed that plant species competition might change the spatial distribution pattern of vegetation. Possibly, the competition played a role for the distribution pattern of the natural vegetation in the studied salt marsh so that the vegetation types have adapted to the local environmental conditions (Ji et al., 2009). The efficiency of the resorption process seems to be related to the plant species (Cartaxana & Catarino, 2002). Salt marshes contain steep environmental gradients, for instance conditions are fairly mild near the terrestrial border of the marsh but become severe in salt pans and these traits are effective on microhabitat preferences and zonation of salt marshes (Richards et al. 2005; Apaydın et al. 2009). The highest N and P resorption efficiency was found in 2 species belonging to Chenopodiaceae, namely S. prostrata

and *S. marina*, while the lowest N and P resorption efficiency was found in *E. paralias*. In other words, the species that are better adapted to saline conditions have also higher resorption efficiencies. *S. prostrata* and *S. marina*, which occur around salt-pans, can use nitrogen more effectively as compared to salt pandune transitional species, such as *O. maritimus* and *E. paralias*. Although all species inhabit areas of similar flooding conditions, some species seem to have a higher competitive capability than others. The differences among species in terms of resorption efficiency are probably due to the different adaptation capacities of species against harsh and selective saline

conditions (Sánchez et al, 1998; Kutbay & Demir, 2001; Cartaxana & Catarino, 2002).

High N resorption efficiency values of *S. prostrata* and *S.marina* indicated that nutrient resorption may play an important role on nitrogen dynamics in coastal salt marshes. It is possible to generalize the obtained results to other similar coastal salt marshes for reclamation and rehabilitation purposes (Álvarez et al., 2001; Jafari et al., 2003). When the temporal and spatial factors are established more clearly, the factors affecting nutrient resorption in salt marshes will be explained in more detail.

References

- Ababou A, Chouieb M, Khader M, Mederbal K & Saidi D (2010). Using vegetation units as salinity predictors in the Lower Cheliff Algeria. *Turk J Bot* 34:73-82.
- Abd El-Ghani MM (2000). Vegetation composition of Egyptian inland salt marshes. *Bot Bull Acad Sin* 41: 305–314.
- Adam P (1990). Saltmarsh Ecology. Cambridge: Cambridge University Press.
- Allahverdiev SR, Mavituna M, Ganieva R & Nafisi S (1998). Effects of salt stress and synthetic hormone polystimuline K on photosynthetic activity of *Trianea bogotensis* Karst. *Turk J Bot J Bot* 22: 19-23.
- Aschenbach TA (2006). Variation in growth rates under saline conditions of *Pascopyrum smithii* (Western Wheatgrass) and *Distichlis spicata* (Inland Saltgras) from different source populations in Kansas and Nebraska: Implications for the restoration of salt-affected plant communities. *Rest. Ecology* 14: 21–27
- Aerts R (1996). Nutrient resorption from senescing leaves of perennials. Are there general patterns? *J Ecol* 84: 597–608.
- Álvarez-Rogel J, AF Alcaraz & Silla RO (2000). Soil salinity and moisture gradients and plant zonation in Mediterranean salt marshes of Southeast Spain. *Wetlands* 20: 357–372.
- Álvarez-Rogel J, Silla RO & Alcaraz AF (2001). Edaphic characterization and soil ionic composition influencing plant zonation in a semiarid Mediterranean salt marsh. *Geoderma* 99: 81–98.
- Apaydın Z, Kutbay HG, Özbucak T, Yalçın E & Bilgin A. (2009). Relationship between vegetation zonation and edaphic factors in a salt-marsh community (Black Sea Coast). *Pol J Ecol* 57: 99-112.
- Asri Y & Ghorbanli M (1997). The halophilous vegetation of the Orumieh lake salt marshes, N. W. Iran. *Plant Ecol* 132: 155–170.

- Avcı K, Kutbay HG & Gençoğlu S (2001). Bafra Ovasında Halofitik (Tuzcul) Vejetasyonun Tesbiti In: T.C. Başbakanlık Köy Hizmetleri Genel Müdürlüğü (ed.) *Toprak ve Su Kaynakları Araştırma Yıllığı*, pp. 346-356, Ankara: APK Dairesi Başkanlığı.
- Brummitt RK & Powell CE (2001). *Authors of Plant Names*. Kew: Royal Botanic Gardens.
- Cartaxana P & Catarino F (2002). Nitrogen resorption from senescing leaves of three salt marsh plant species. *Plant Ecol* 159: 95–102.
- Cornelissen JHC, Werger MJA, Castro-Diez P, van Rheen JWA & Rowland AP (1997). Foliar nutrients in relation to growth, allocation, and leaf traits in seedlings of a wide range of woody plant species. *Oecologia* 111: 460–469.
- Curcó A, Ibàñez CJ, Day WWW & Pratt N (2002). Net primary production and decomposition of salt marshes of the Ebre Delta (Catalonia, Spain). *Estuaries* 25: 309–324.
- Daget PH (1977). Lé bioclimat méditerranéen: Analyse des formes climatiques par le système d'Emberger. *Vegetatio* 34: 87–103.
- Donovan LA, Richards JH & Schaber EJ (1997). Nutrient relations of the halophytic shrub, *Sarcobatus vermiculatus*, along a soil salinity gradient. *Plant Soil* 190: 105–117.
- Duchesne L, Ouimet R, Camiré C & Houle D (2001). Seasonal nutrient transfers by foliar resorption, leaching, and litter fall in a northern hardwood forest at Lake Clair Watershed, Quebec, Canada. *Can J For Res* 31: 333-344.
- El-Demerdash MA (1996). The vegetation of the Farasãn Islands, Red Sea, Saudi Arabia. *J Veg Sci* 7: 81–88.
- Engin A, Korkmaz H & Şahin N (1988). Preliminary observations of floristic investigation of Bafra-Altınkaya Dam Lake (in Turkish). In: Bilaloğlu R, Çelik N, Erdem Ü, Dere Ş, Yanıkoğlu A, Gey H, Erdem U (ed.) *IX. National Biological Congress Booklet Section of General and Systematic Botany* pp. 353-361. Sivas: Özemek Matbaa.

- Flowers TJ, Troke PF & Yeo AR (1977). The mechanism of salt tolerance in halophytes. *Ann Rev Physiol* 28: 89–121
- García LV, Maraňón T, Moreno A & Clemente L. (1993). Aboveground biomass and species richness in a Mediterranean salt marsh. J Veg Sci 4: 417–424.
- Grime JP, Thompson K, Hunt R, Hodgson JG, Cornelissan JHC & Rorison IH. (1997). Integrated screening validates primary axes of specializations in plants. *Oikos* 79: 251–281.
- Grime JP (2002). Plant Strategies, Vegetation Processes and Ecosystem Properties, Second Edition. Chichester: Wiley & Sons Ltd.
- Hodgson JG, Wilson PSR, Hunt JP & Thompson R (1999). Allocating C-R-S plant functional types; a soft approach to a hard problem. *Oikos* 85: 282–294.
- Jafari M, Chahouki MAZ, Tavili A & Azarnivand H (2003). Soilvegetation relationships in Hoz-e-Soltan region of Qom province, Iran. Pak J Nut 2: 329–334.
- Ji Y, Zhou G & New T (2009). Abiotic factors influencing the distribution of vegetation in coastal estuary of the Liaohe Delta, Northeast China. *Estuaries Coasts* 32: 937-942.
- Jiang Li-Fen, Luo Yi-Qi, Chen Jia-Kuan & Li B (2009). Ecophysiological characteristics of invasive Spartina alterniflora and native species in salt marshes of Yangtze River estuary, China. Estuar Coast Shelf Sci 81: 74-82.
- Killingbeck KT (1996). Nutrients in senesced leaves: keys to the search for potential resorption and resorption proficiency. *Ecology* 77: 1716–1727.
- Killingbeck KT (2004). Nutrient resorption In: Noodén LD (ed.) Plant Cell Death Processes, pp. 215–223. USA: Elsevier Science.
- Kobe RK, Lepczyk CA & Iyer M. (2005). Resorption efficiency decreases with increasing green leaf nutrients in a global data set. *Ecology* 86: 2780–2792.
- Köksal A. (1972). Geographical Analysis of Bafra Plain. Ankara: Faculty of Letter Publications.
- Krüger HR, Peinemann N (1996). Coastal plain halophytes and their relation to soil ionic composition. *Vegetatio* 122: 143–150.
- Kutbay HG (2001). Nutrient content in leaves from different strata of a swamp forest from Northern Turkey. *Pol J Ecol* 49: 221–230.
- Kutbay HG & Demir M (2001). The changes in ionic contents of salt marsh species and the importance of edaphic physicochemical factors. Arab Gulf J Sci Res 19: 35–43.
- Lajtha K (1987). Nutrient resorption efficiency and the response to phosphorus fertilization in the desert shrub *Larrea tridentata* (DC) Cov. *Biogeochemistry* 4: 265–276.
- Levine J.. Brewer MJS & Bertness MD (1998). Nutrients, competition and plant zonation in a New England salt marsh. *J Ecol* 86: 285–292.
- Marscher H (1995). *Mineral nutrition of higher plants Second edition*. Cambridge: Academic Press.
- Mayor X & Roda F (1992). Is primary production in holm oak forests nutrient limited? *Vegetatio* 99: 209–217.

- Munir N & Aftab F (2009). The role of polyethylene glycol (PEG) pretreatment in improving sugarcane's salt (NaCl) tolerance. *Turk J Bot* 34:73-82.
- Niinemets U& Tamm U (2005). Species differences in timing of leaf fall and foliage chemistry modify nutrient resorption efficiency in deciduous temperate forest stands *Tree Physiol* 25: 1001-1014.
- Onaindia M & Amezaga I (1999). Natural regeneration in salt marshes of northern Spain. *Ann Bot Fenn* 36: 59–66.
- Pennings SC & Richards CL (1998). Effects of wrack burial in saltstressed habitats: *Batis maritima* in a southwest Atlantic salt marsh. *Ecography* 21: 630–638.
- Reynolds CE, Houle G & Marquis C (2001). Light and salinity affect growth of the salt marsh plant *Aster laurentianus*. *New Phytol* 149: 441–448.
- Richards CL, Pennings SC & Donovan LA (2005). Habitat range and phenotypic variation in salt marsh plants. *Plant Ecol* 176: 263-273.
- Sánchez JM, Otero XL & Izco J (1998). Relationship between vegetation and environmental characteristics in a salt-marsh system on the coast of Northwest Spain. *Plant Ecol* 136: 1–8.
- Sans-Pérez V, Castro-Díez P & Millard P (2009). Effects of drought and shade on nitrogen cycling in the leaves and canopy of Mediterranean *Quercus* seedlings. *Plant Soil* 316: 45-56.
- Shaver GR, Melilo JM (1984). Nutrient budgets of marsh plants: efficiency concepts and relation to availability. *Ecology* 65: 1491-1510.
- Skeffington MJS & Jeffrey DW (1988). Response of *Armeria maritima* (Mill.) Willd. and *Plantago maritima* L. from an Irish salt marsh to nitrogen and salinity. *New Phytol* 110: 399-408.
- Turkish Ministry of Agriculture (2002). Meteorological Bulletin, Mean and Extreme Temperature and Precipitation Values. Ankara: State Meteorological Service.
- Ungar IA, Benner DK & McGraw DC (1979). The distribution and the growth of *Salicornia europaea* on an inland salt pan. *Ecology* 60: 329–336.
- Van Heerwaarden LM, Toet S, & Aerts R (2003). Current measures of nutrient resorption efficiency lead to a substantial underestimation of real resorption efficiency: facts and solutions. *Oikos* 101: 664–669.
- Vidal E, Médail F, Tatoni T & Bonnet V (2000). Seabirds drive plant species turnover on small Mediterranean islands at the expense of native taxa. *Oecologia* 122: 427–434.
- Wadie HA (2002). Floristic composition and vegetation of Wadi Talha, Aseer Mountains, South West Saudi Arabia. J Biol Sci 2: 285-288.
- Wang W & Lin P (1999). Transfer of salt and nutrients in *Bruguiera* gymnorrhiza leaves during development and senescence. Mang Salt Marsh 3: 1–7.
- Wilkoń-Michalska J (1985). Structure and dynamics of the inland populations of *Salicornia patula*. *Vegetatio* 61: 145–154.

- Yasumura Y, Onoda Y, Hikosaka K & Hirose T (2005). Nitrogen resorption from leaves under different growth irradiance in three deciduous woody species. *Plant Ecol* 178:29-37.
- Yeo AR (1983). Salinity resistance: Physiologies and prices. *Physiol Plant* 58: 214–222.
- Yuan ZY, Li LH, Han XG, Huang JH, Jiang GM & Wan SQ (2005). Soil characteristics and nutrient resorption in *Salix krylovii* native to northern China. *Plant Soil* 273: 257–268.
- Zotz G, (2004). The resorption of phosphorus is greater than that of nitrogen is senescing leaves of vascular epiphytes of lowland Panama. *J Trop Ecol* 20: 693-696.