

SEASONAL CHANGE OF ORIBATID MITE COMMUNITIES (ACARI, ORIBATIDA) IN THREE DIFFERENT TYPES OF MICROHABITATS IN AN OAK FOREST

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Abstract. Oribatid mites are one of the most abundant groups of the ground-dwelling mesofauna. They can be found in almost every terrestrial habitat all over the world and they are characterized by great species richness and great number of individuals. In spite of that not enough is known about their behaviour on community level and their spatial and temporal pattern in different habitats of the world. In our present study the seasonal behaviour of oribatid mite communities was analysed in three types of microhabitats in a temperate deciduous forest: in leaf litter, soil and moss. Samples were collected at a given site in a year and a half and the oribatid mite communities living there were studied on genus level along with the changes of meteorological factors characteristic of the area. The results show that corresponding to similar previous researches, the communities in our study do not have a seasonally changing, returning pattern either. Based on this, we can conclude that climatic differences and differences in other seasonally changing factors between the seasons do not have a significant role in the annual change of communities. Besides that we discovered that the communities of the three microhabitats are not completely the same. It is the oribatid mite community of the moss which differs mostly from communities in the leaf litter and in the soil. Our study calls attention among others to the fact that compositional changes of the oribatid mite communities living all over the world and their causes are unclear to date.

Keywords: *oribatid mites, communities, microhabitats, seasonality, temperate deciduous forest*

Introduction

In most habitats oribatid mites account for the biggest part of microarthropods (e.g. Schenker, 1986; Johnston and Crossley, 2002). They can be found in most terrestrial microhabitats: in soil, leaf litter, moss, underwood, foliage and in aquatic habitats as well (Behan-Pelletier, 1999). They can be found mostly in great species richness and abundance in their habitats (Behan-Pelletier, 1999). They play a significant role in decomposition processes because they fragment the organic matter and influence the biomass and species composition of fungi and bacteria (Wallwork, 1983; Seastedt, 1984; Yoshida and Hijii, 2005). As this group plays a significant role in soil processes, it is necessary to get to know its spatial pattern and the causes of pattern generation, which can be used later for indication (Behan-Pelletier, 1999).

Several researches have already been done to detect the pattern and composition of oribatid mite communities and their exact causes. However, it is hidden to date by which mechanisms the structure and functioning of the individual communities are affected.

Climatic factors belong to the most determinant ones. The above statement is reflected by the number of researches as well, because most of the studies on oribatid mite communities investigate how the communities react to the meteorological factors changing naturally or artificially, especially to temperature and the amount of precipitation (Gergócs and Hufnagel, 2009). First it is worth investigating the effect of the natural changes of the climatic factors, and the most appropriate way to do this is to study the seasonal changes of the communities. The mistake in most of these studies is that the research on seasonality is conducted over a maximum of a year (e.g. Schenker, 1984), so it cannot be determined whether the observed phenomena occur similarly each year. The most significant research on this topic was conducted by Irmeler (2006) in a beech forest in Germany. He investigated monthly changes of ground-dwelling oribatid mite communities over a seven-year period and found that there are no important changes among the communities.

Our research is similar to his one, however, it differs as well because we collected samples every two weeks and not once a month, and we investigated not one but three microhabitats over a one and a half year period in an oak forest in Hungary.

The other difference is that data on oribatid mites were recorded on genus level and not on species level. The decision on the genus level can be justified by several factors. The analysis of Caruso and Migliorini (2006) showed that there were not any important changes in the data when studying human disturbance on oribatid mites and changing from species level to genus level. Podani (1989) found a similar result in case of plants, namely, changing to genus level does not mean a significant difference when comparing the studied habitats. Osler and Beattie (1999), Hammer and Wallwork (1979) and Norton et al. (1993) concluded according to their studies that the many widespread genera and families indicate that the similarity between oribatid mite communities should be studied above species level. In their meta-analyses their assumption was confirmed that taxonomical levels above the species level are more suitable to compare habitats. According to this we also compared the data series recorded with time difference on genus level.

The aim of our study was to explore which seasonal changes occur in oribatid mite communities living in three types of microhabitats in an oak forest and what role the most important climatic factors that is the amount of precipitation and temperature have in these changes.

Materials and methods

Samples for our study were collected in 2005 and 2006 in a given quadrat of 100 m² in a dry oak forest near Törökbálint, Hungary (47°25'38" N, 18°54'16" E). In the sampling quadrat the most common tree species is turkey oak (*Quercus cerris*), common trees are field maple (*Acer campestre*), common ash (*Fraxinus excelsior*) and wild service tree (*Sorbus torminalis*). The most important herbaceous plants are broad-leaved Solomon's seal (*Polygonatum latifolium*) and garlic mustard (*Alliaria petiolata*). Samples were collected every two weeks from 15th March till 1st December 2005 and from 26th March till 30th July 2006. Every time samples were collected randomly from

three types of substrate within the quadrat: from 500 cm³ leaf litter, 300 cm³ foena (from under the leaf litter) and 0.5 dm² hypnum moss (*Hypnum cupressiforme*) living on tree trunks. So we obtained altogether 19 moss samples and 18 leaf litter and 18 foena samples in 2005 (the first time no leaf litter and foena samples were collected); in 2006 we collected 10 leaf litter, 10 moss and 9 foena samples (the tenth sample could not be analyzed for technical reasons). Components of the mesofauna were extracted with the help of a modified Berlese-Tullgren funnel (Coleman et al., 2004) and conserved in isopropyl alcohol, then they were sorted into larger groups, and finally oribatid mites were identified on genus level. For the identification the works of Balogh (1965), Balogh and Balogh (1992, 1972), Balogh and Mahunka (1980), Olsanowski (1996) and Hunt et al. (1998) were used.

The samples were analyzed based on oribatid mite communities with the help of multivariate computer aided methods using the software Syn-tax 2000 (Podani, 2001). The size difference between the samples from the three types of substrate was compensated by comparing the ratio and presence-absence data of the oribatid mite genera with the help of non-metric multidimensional scaling and classification method using several distances. Monte Carlo method was used to check these analyses. The means of the genus proportion of the substrate types were calculated, each mean was multiplied by a number created by a random generator twenty times, then these new values were divided by the sum of the random numbers. This way twenty data series were generated per substrate. These were compared using classification and ordination. Correspondence analysis and PCA analysis were used to identify the genera which are responsible for possible differentiations. The difference of the genera in specimen number and proportion was checked using Mann-Whitney test between the substrate types. The average diversity and the genus number were calculated in case of each substrate type.

In order to recognize seasonality patterns, each season was marked (in winter no sampling was conducted), and the changes in genus diversity and total abundance during a period of the study were displayed separately in case of the three substrate types. The relationship between the pattern changes of the communities and meteorological factors was analyzed with the help of data series from the meteorological station in Pestszentlőrinc (47°25'53" N, 19°10'57" E). This station can be found 21 air km eastwards from our sampling point. In the case of precipitation the total precipitation amount of the sampling day and that of the preceding 5 and 10 days, and the standard deviation of the precipitation amount of the preceding 5, 10, 15, 20, 25 and 30 days were considered. In the case of temperature minimum and maximum values of the given days were available in our database, so in our analyses the means of these two values were considered. As for temperature data, the mean temperature of the sampling day and that of the preceding 5 and 10 days, and the standard deviation of temperature of the preceding 5, 10, 15, 20, 25 and 30 days were involved in our analyses. A redundancy analysis was conducted for all substrates first and then separately. In order to make the relationships more exact, the correlation between certain genera and meteorological factors was studied based on Spearman's analysis.

Results

52 genera were found in the three types of substrate altogether, in leaf litter and foena 48-48 genera and in moss only 30 occurred. According to the ordination and

classification (*Fig. 1*) it can be stated that oribatid mite communities of the leaf litter and foerna do not differ from each other but they differ from moss samples. In these figures it can be seen as well that the samples collected in different times are completely mixed, the different markings of the three seasons are also mixed without a pattern in the figures, which confirms our assumption that no seasonal effects can be found in the case of oribatid mite genera.

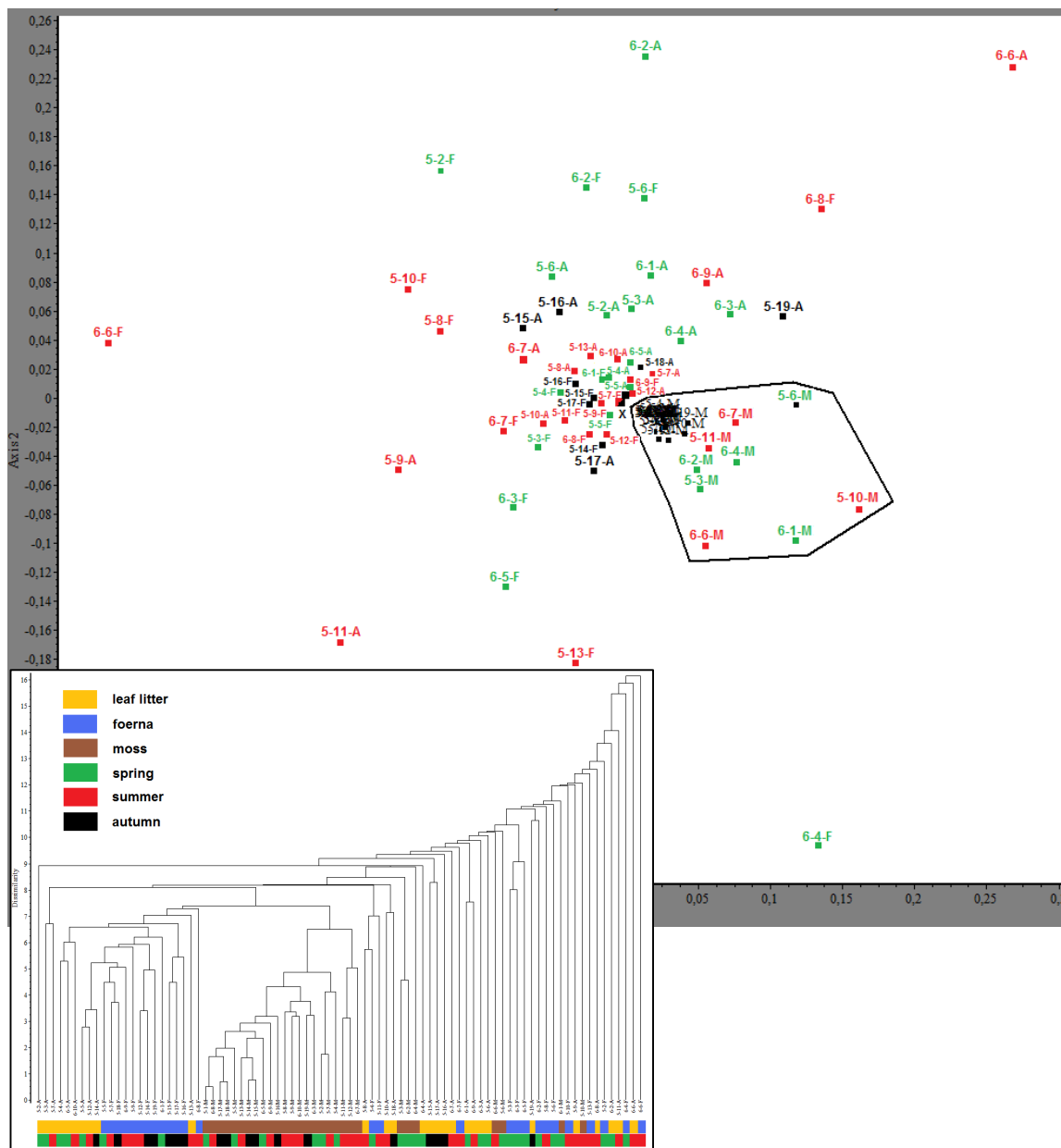


Figure 1. Large figure: Non-metric multidimensional scaling of the ratio of the genera and juvenile groups in the three substrates using Euclidean distance. The first number in the codes represents 2005 (5) or 2006 (6), the second one is the chronologic serial number of the samples and the third symbol in the code means the substrate. M=moss, F=foerna, A=leaf litter. Spring samples were marked with green, summer samples with red and autumn-winter samples with black colour. Moss samples are framed. X=5-14-A, 5-18-F, 5-19-F. **Small figure:** Classification of the same series using group-average method with Euclidean distance. Considered groupings were marked with a colour code based on substrate types and seasons

Monte Carlo experiments conducted in order to check the above analyses can be seen in *Fig. 2*. It shows that leaf litter and foerna substrates also differ from each other, however, not to the same degree as from the moss substrate.

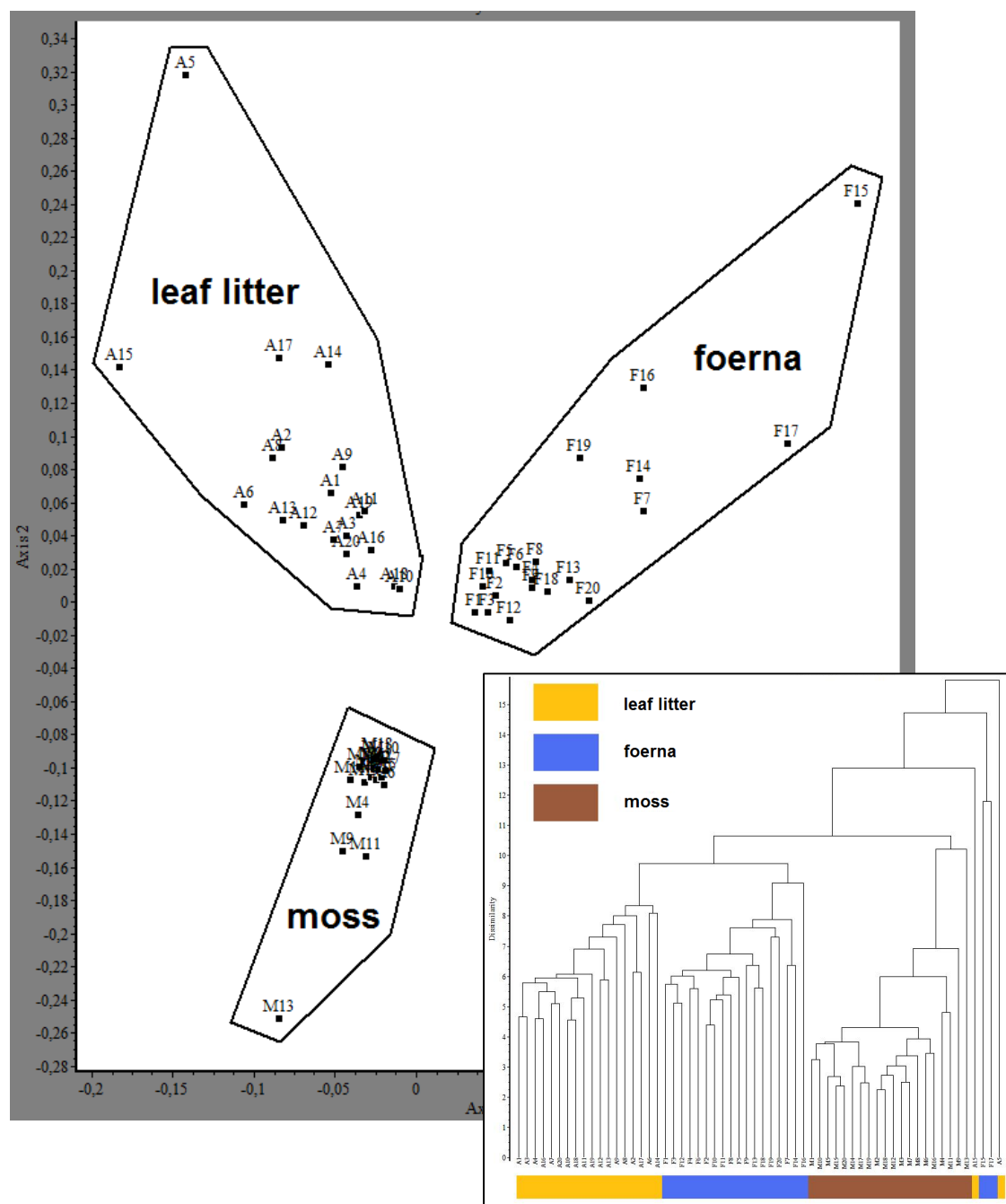
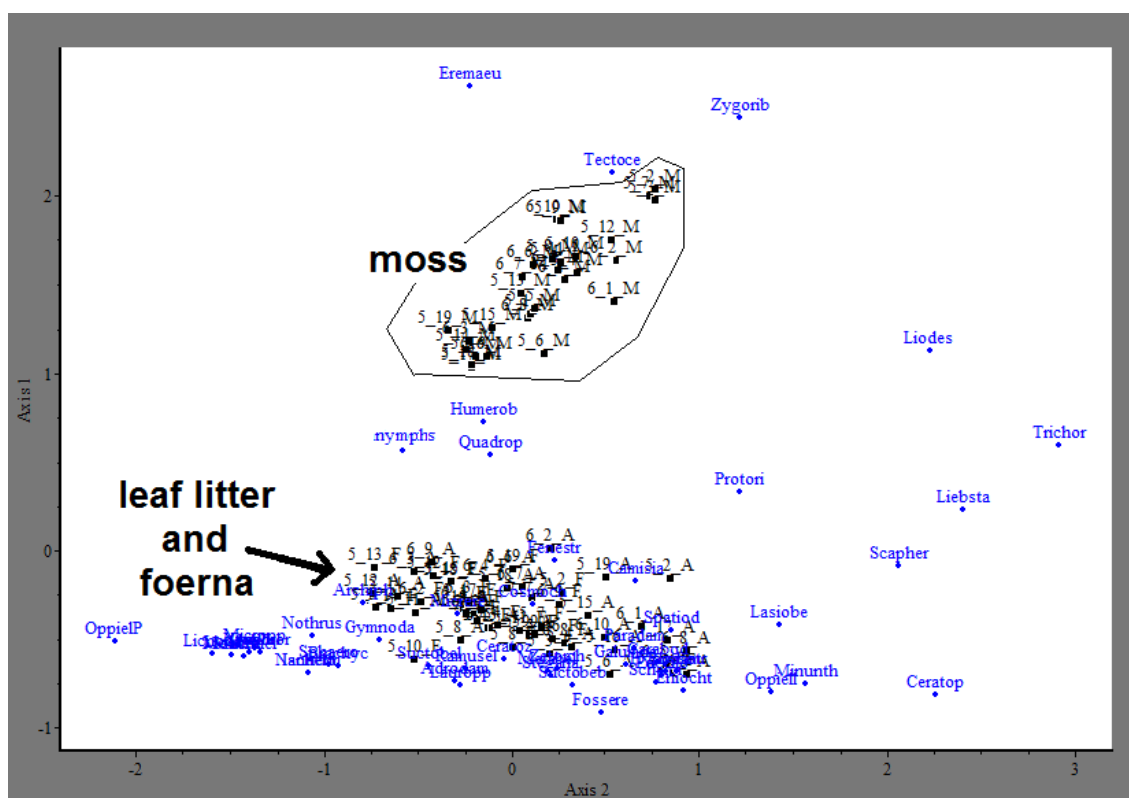


Figure 2. Large figure: Non-metric multidimensional scaling of the samples created with Monte Carlo method from the ratios of the three substrates: M1-20; A1-20; F1-20 (standardised Euclidean distance). **Small figure:** Classification of the same series using group-average method with standardised Euclidean distance. Separation of the three substrate types was marked with a colour code

Separation of moss samples is mainly caused by their smaller genus diversity and smaller genus richness and by the fact that the following genera: *Tectocephus*, *Eremaeus*, *Zygoribatula* and the juvenile group, to which larvae and nymphs belong, are present in a larger number here (Fig. 3). In moss samples 7, in leaf litter 19 and in foerna samples 22 genera were present on average during a period of one and a half years. Genus diversity of the samples from all three substrates differ from each other significantly ($p < 0.01$). Moss samples have the lowest diversity and foerna samples have the highest one (Table 1).

	Foerna	Leaf litter	Moss
Shannon mean ($\pm$ standard deviation)	2.20 ($\pm$ 0.21)	1.871 ($\pm$ 0.34)	0.98($\pm$ 0.26)
Simpson mean ($\pm$ standard deviation)	0.82 ($\pm$ 0.07)	0.73 ($\pm$ 0.11)	0.52 ($\pm$ 0.10)
Average number of genera	21.7 ($\pm$ 4.5)	19.3 ($\pm$ 4.3)	6.6 ($\pm$ 2.7)



The proportions of the genera *Tectocephus* and *Zygoribatula* and the juveniles in the three substrates can be seen in *Table 2*. The difference of these values in the three substrates was significant in all cases ($p < 0.01$). The genus *Eremaeus* occurred only in

moss samples except for only one foerna sample, with 10.9 specimens on average, 15 times out of 29, but with rather varying abundance and proportion in time. Besides the above, two other genera near the moss group: *Humerobates* and *Quadroppia* can be seen in Fig. 3. *Humerobates* is a rare component, and in the case of *Quadroppia* no difference was found between the proportions compared to the other two substrates.

Table 2. Proportions of *Tectocephus*, *Zygoribatula* and the juveniles in the three substrates. The values in moss samples are highlighted because these values are significantly different from the values of the two other substrates.

	Leaf litter	Foerna	Moss
juvenile group / total number of specimens	0.188±0.106	0.201±0.104	0.397±0.206
juvenile group / adults	0.249±0.174	0.280±0.169	0.919±0.775
proportion of <i>Tectocephus</i>	0.020±0.037	0.017±0.020	0.261±0.170
proportion of <i>Zygoribatula</i>	0.017±0.023	0.005±0.004	0.230±0.187

The reason of the difference between foerna and leaf litter can be seen in the PCA analyses (Fig. 4). Table 3 represents in which genera the numbers of specimens and the ratios significantly differ between the two substrates. In all cases it can be shown that the genera displayed in Table 3 are present with larger abundance and relative abundance in foerna samples.

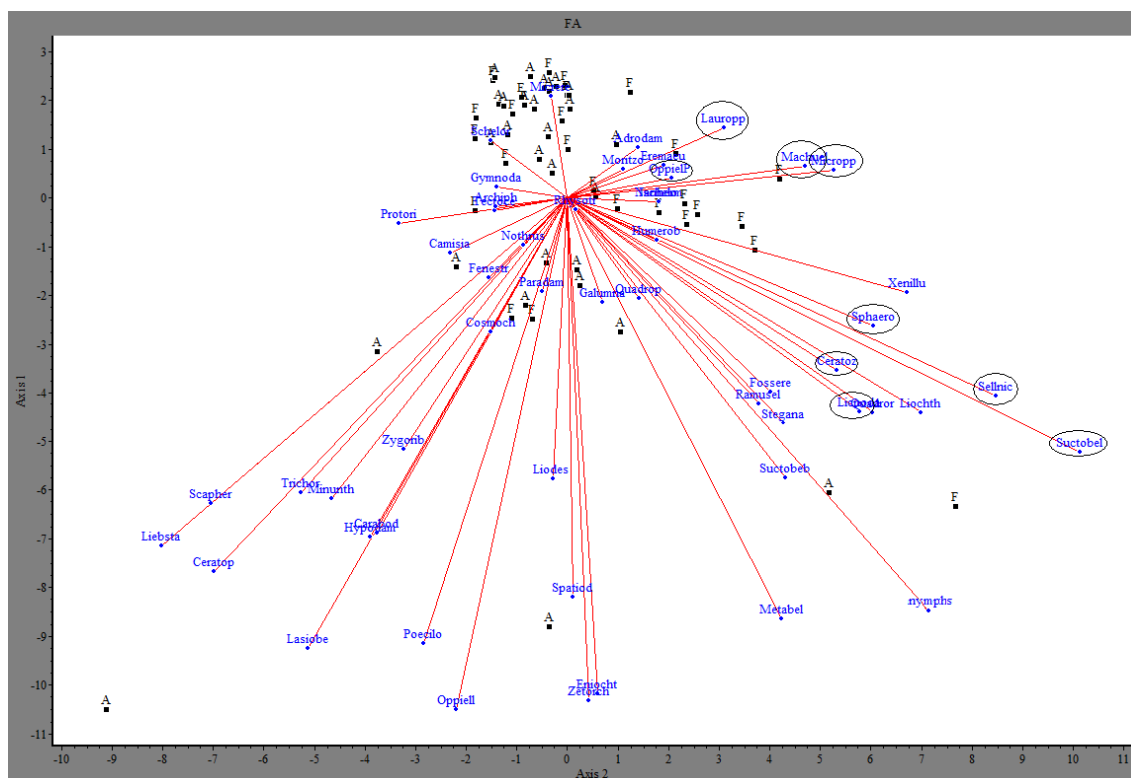


Figure 4. PCA analysis with the ratios of the juvenile group (“nymphs” in the figure) and the genera of the leaf litter and foerna samples (standardised). A=leaf litter, F=foerna; the first 7 letters of the genera are displayed and the genera, the abundance and proportion of which significantly differ from each other in the two types of substrate, are circled

Table 3. Abundance and proportion of the genera causing the difference between leaf litter and foerna samples

genera	with the numbers of specimens			with proportions		
	leaf litter	foerna	p(same)	leaf litter	foerna	p(same)
<i>Sellnickochthonius</i> sp.	0.8	2.3	0.028	0.0014	0.0043	0.025
<i>Ceratozetes</i> sp.	1.9	3.0	0.055	0.0048	0.0092	0.049
<i>Lauroppia</i> sp.	0.3	8.1	9.76E-08	0.0012	0.0250	3.06E-07
<i>Licnodamaeus</i> sp.	60.1	68.7	0.04	0.1323	0.1577	0.0604
<i>Machuella</i> sp.	0.1	2.4	1.05E-05	0.0002	0.0058	9.19E-06
<i>Microppia</i> sp.	0.0	2.0	2.32E-05	0.0002	0.0048	0.000034
<i>Oppiella</i> (<i>Perspiciuoppia</i>) sp.	0.0	1.2	0.017	0.0001	0.0030	0.018
<i>Sphaerorchthonius</i> sp.	7.2	11.4	0.002	0.0200	0.0274	0.0006
<i>Suctobelbella</i> sp.	11.8	29.5	0.0002	0.0298	0.0750	4.95E-05

Seasonal changes could not be detected in the communities of the three substrates. These results can be seen on *Fig. 1* and 5-6. The second number in the codes of the samples in *Fig. 1* (the large one) means the progress of time during the year. It can be observed well that the numbers do not form groups according to the date of sampling. This proves that the progress of time that is seasonality does not generate patterns in oribatid mite communities. In *Fig. 5* change in the Shannon diversity of the genera in the case of the three substrates and in *Fig. 6* change in abundance of the oribatid mites can be observed, however, a seasonal pattern recurring the following year cannot be detected.

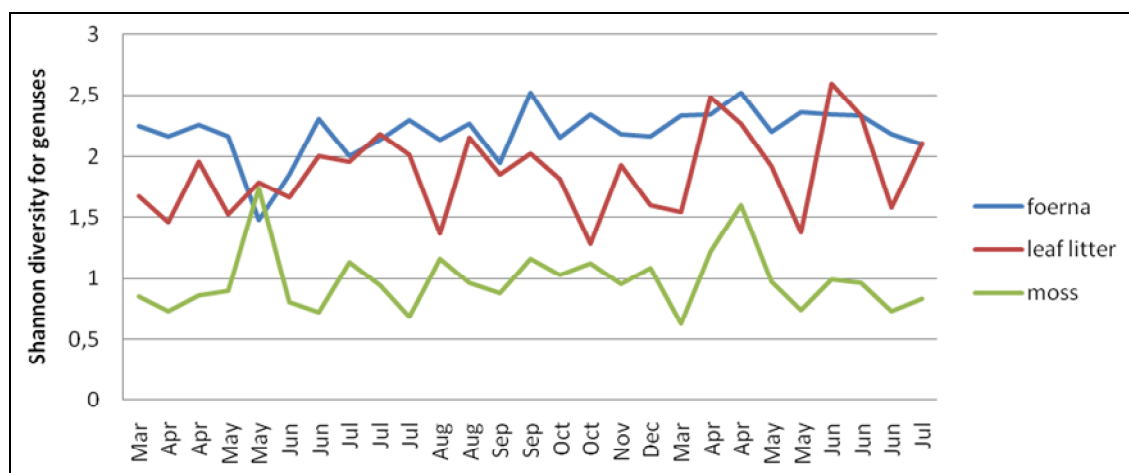


Figure 5. Change in the Shannon diversity of the genera in the case of the three substrates (2005-2006)

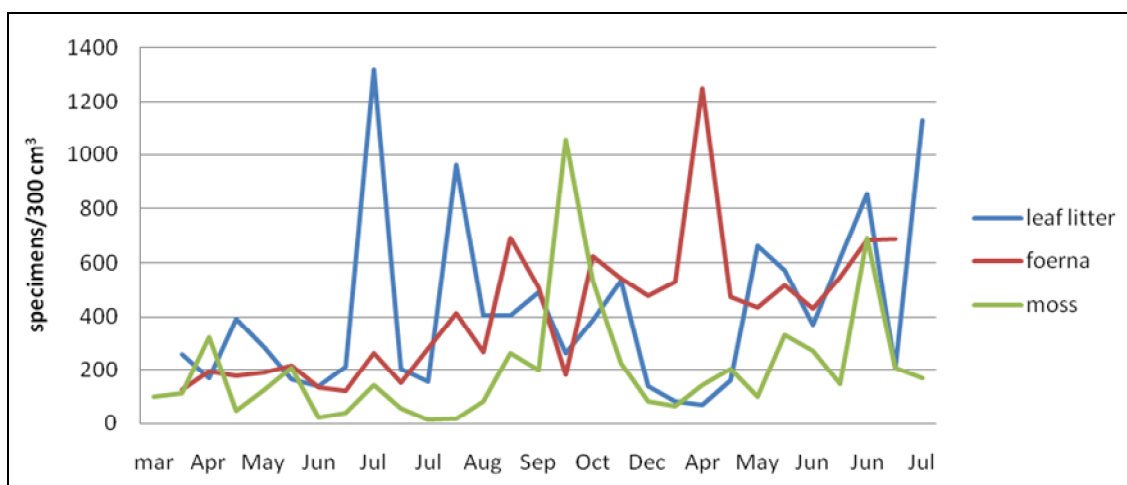


Figure 6. Change in abundance of the oribatid mites (2005-2006)

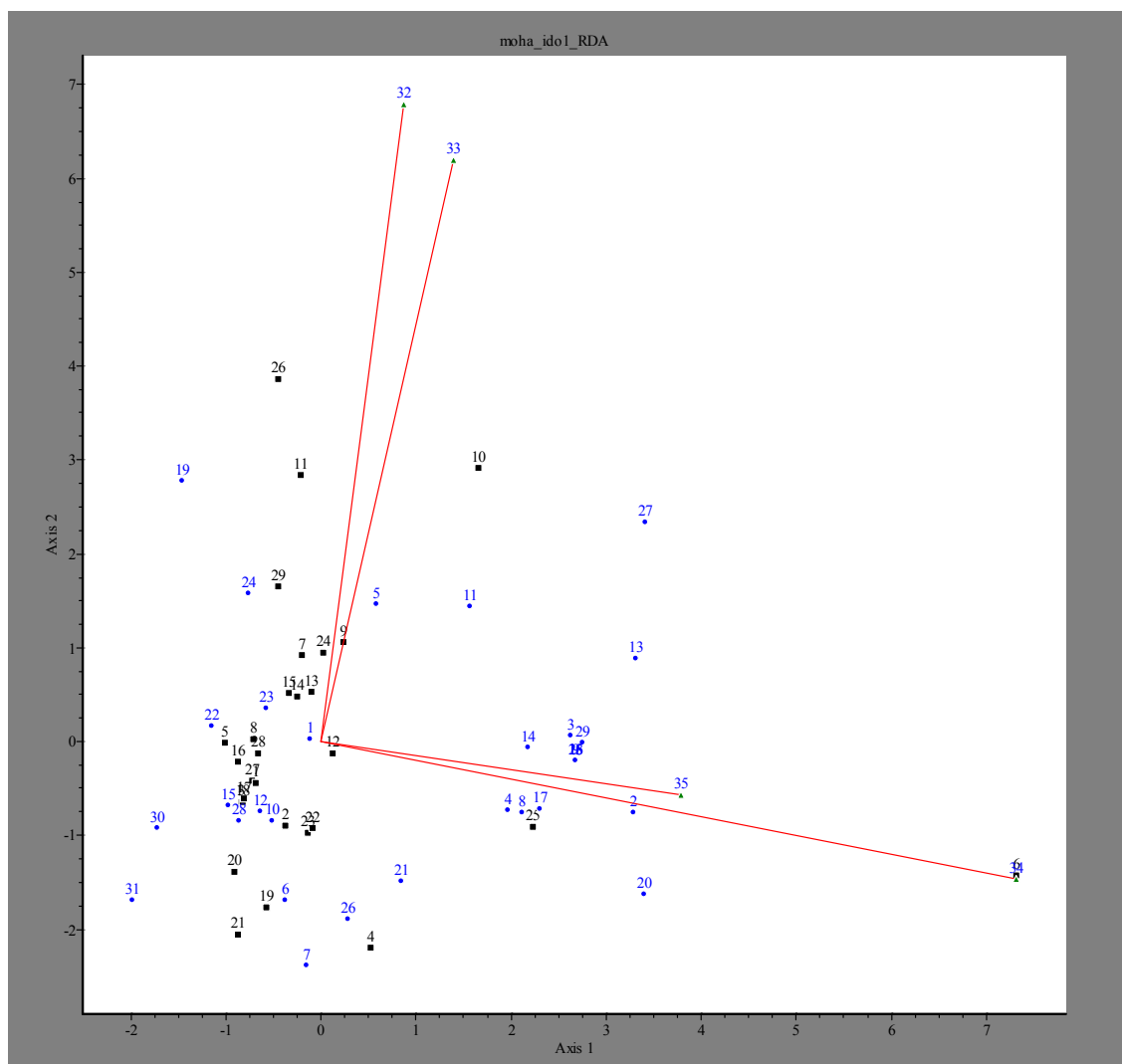


Figure 7. Redundancy analysis of moss samples collected during one and a half year regarding the precipitation amount and mean temperatures. (32: mean temperature (of 5 days); 33: mean temperature (of 10 days); 34: total amount of precipitation (in 5 days); 35: total amount of precipitation (in 10 days). The first axis is for 46% and the second one is for 31%.)

We have not found any relationships between the annual changes in precipitation amount and temperature and the composition and structure of oribatid mite communities in leaf litter and foerna, however, in the case of moss we have. The result of the redundancy analysis is displayed in Fig. 7. The location of the samples is affected by mean temperature to a larger degree than by the precipitation amount. This can be caused by the genus *Tectocephus*, which is one of the most frequent genera in moss samples (43% of the adult specimens on average), and its correlation is $r=0.38$ regarding the five-day mean temperature (Spearman's correlation; $p=0.043$); and regarding the ten-day mean temperature its correlation is $r=0.45$ (Spearman's correlation; $p=0.014$) (Fig. 8).

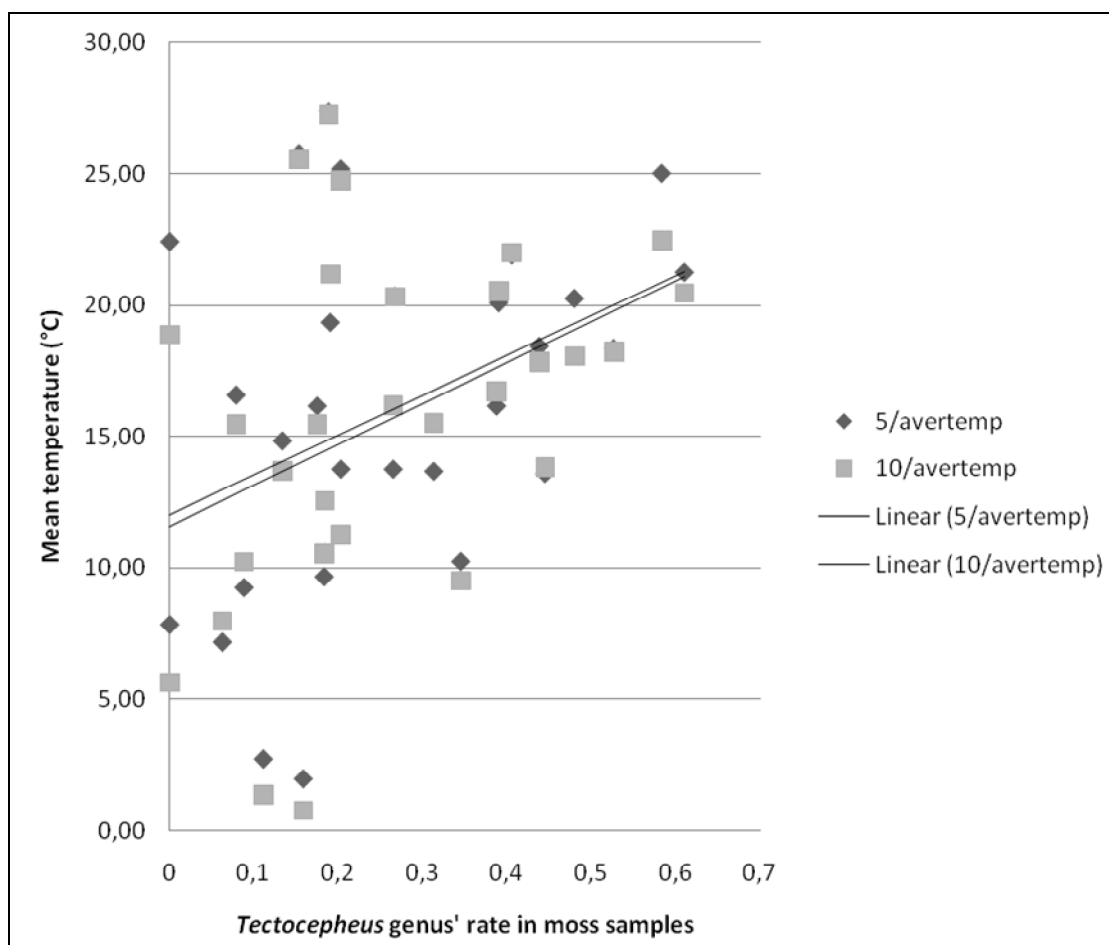


Figure 8. Correlation between the genus *Tectocephus*' rate and the five-day and ten-day mean temperature

The most frequent groups within the moss are the genera *Tectocephus* and *Zygoribatula* and the juvenile group. Table 4 shows that the ratio of the two most frequent genera is inversely proportional to the juvenile group that is the proportion of the juvenile group is only larger if the proportion of the two genera is smaller. The proportion of juveniles and adults changes during the one and a half year (Fig. 9), however, this pattern does not recur the following year, and it is not in connection either

approximately in the same abundance further away from the oak trunks, than in the soil around it. For our study, data were collected in an oak forest. This may be the reason for the fact that we have not found any seasonal changes or changes in the genus composition, either. However, since Irmeler (2006) conducted his studies in a beech forest and could not observe any seasonality either, and Schenker (1984) collected samples for only one year, it cannot be stated that the lack of seasonality is characteristic of oak forests.

If changes in communities do not occur seasonally, this can be brought into connection with micro- and macroclimatic effects. According to the results of Stamou and Sgardelis (1989) it could be concluded that the density of oribatid mites is largely influenced by temperature, although several later studies showed that temperature does not have the power to shape communities (Haimi et al., 2005). Irmeler (2006) found that the structure of the community was in connection with the annual mean temperature only, and only some species showed significant correlation with some climatic factors. Webb et al. (1998) showed in the case of oribatid mite species living in polar areas that these species do not depend on seasonal changes, life cycle of the studied oribatid mite species is mainly influenced by temperature fluctuation. In our study proportional change correlating with temperature could be observed in the case of the genus *Tectocephus* only and only in moss, however, no such connection can be found in the study of Irmeler (2006), for example. Based on our results – just like based on those of Irmeler (2006) – it can be concluded that the structure of oribatid mite communities is not affected by climatic factors in leaf litter and foena substrates. In moss samples the connection with temperature was due to the genus *Tectocephus*.

Seasonality can also be observed in the decomposition of plant material. Quantity and quality of the decomposing plant material change seasonally in the leaf litter and in the soil so it can be assumed that oribatid mite communities may change correspondingly during the year. However, the exact role of oribatid mites in the decomposition of the leaf litter is not completely clear till this day (Lindo and Winchester, 2007). The most important role of oribatid mites in the decomposition is the spreading of microbiota as they feed mainly on fungi and bacteria, and they are not in direct connection with the leaf litter input, accordingly (Maraun et al., 2001). This corresponds with the phenomenon observed by us, that the quantity of leaf litter may not have influenced the compositional changes of the communities.

Our result that leaf litter and foena substrates differ from moss was not interpreted by other literature yet, however, there are observations regarding other types of substrates. A common result is for example that the oribatid mite community living in the foliage of the trees differs significantly from the one living in the soil under the trees (e.g. Yoshida and Hijii, 2005; Karasawa and Hijii, 2008). Karasawa and Hijii (2004) showed that the substrate of oribatid mite communities living in the soil, in the foliage, on the bark of the tree trunks and on the remnants of algae accumulating on soil significantly differ from each other in seaside forests. In our study the community living in hypnum moss was simpler than the one living in the soil in the forest. Communities of moss and lichen are always relatively simple (Gjelstrup, 1979; Gjelstrup and Søchtig, 1979; Smrz, 1992; Smrz and Kocourková, 1999; Smrz, 2006). The three frequent genera found in moss turned up in the observations of others as well. *Zygoribatula exilis* is assumed to be a species living in moss (Gjelstrup, 1979), however, the *Zygoribatula* species found by us could be found on the forest ground as well. The genus *Tectocephus* occurs everywhere from drier and more disturbed habitats to intact

forests, accordingly it can also be found in moss in great quantities, especially because climatic fluctuation is larger in moss, what only some species can tolerate (Gjelstrup, 1979). A common epiphyte is *Eremaeus oblongus* (Srnz and Kocourková, 1999), in our study the genus *Eremaeus* could be found only in moss (except for only one foerna sample).

Therefore our research showed that oribatid mite communities living in soil, leaf litter and hypnum moss, in Hungarian oak forests – similarly to those living in German beech forests – did not show seasonal changes. This result is important on the one hand because according to this, we are not bound to a season regarding sampling. However, besides that it would be important to detect the cause of the still occurring changes and patterns exactly. Furthermore it is unclear as well whether non-woody biocoenoses in the temperate zone or oribatid mite communities living in other climatic zones show seasonal changes.

Communities living in the soil differ from the oribatid mite community of moss living on tree trunks more significantly than from the community of the leaf litter. Oribatid mites in moss, especially the genus *Tectocepheus*, may be influenced by climatic factors to a large degree. In future it would be necessary to study oribatid mite communities of various microhabitats in order to detect exactly by which factors and to what degree their composition, changes and patterns are affected.

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