Direct and legacy effects of plant-traits control litter decomposition in a deciduous oak forest in Mexico

Bruno Chávez-Vergara, Agustín Merino, Antonio González-Rodríguez, Ken Oyama, Felipe García-Oliva

Background. Litter decomposition is a key process in the functioning of forest ecosystems, because it strongly controls nutrient recycling and soil fertility maintenance. The interaction between the litter chemical composition and the metabolism of the soil microbial community has been described as the main factor of the decomposition process based on three hypotheses: substrate-matrix interaction (SMI), functional breadth (FB) and home-field advantage (HFA). The objective of the present study was to evaluate the effect of leaf litter quality (as a direct plant effect, SMI hypothesis), the metabolic capacity of the microbial community (as a legacy effect, FB hypothesis), and the coupling between the litter quality and microbial activity (HFA hypothesis) on the litter decomposition of two contiguous deciduous oak species at a local scale.

Methods. To accomplish this objective, we performed a litterbag experiment in the field for 270 days to evaluate mass loss, leaf litter quality and microbial activity in a complete factorial design for litter quality and species site.

Results. The litter of *Quercus deserticola* had higher rate of decomposition independently of the site, while the site of *Q. castanea* promoted a higher rate of decomposition independently of the litter quality, explained by the specialization of the soil microbial community in the use of recalcitrant organic compounds. The Home-Field Advantage index was reduced with the decomposition date (22% and 4% for 30 and 270 days, respectively).

Discussion. We observed that the importance of the coupling of litter quality and microbial activity depends on decomposition stage. At the early decomposition stage, the home-advantage hypothesis explained the mass loss of litter; however, in the advanced decomposition stage, the litter quality and the
metabolic capacity of the microbial community can be the key drivers.
Direct and legacy effects of plant-traits control litter decomposition in a deciduous oak forest in Mexico

Bruno Chávez-Vergara¹,², Agustín Merino³, Antonio González-Rodríguez³, Ken Oyama⁴, Felipe García-Oliva³*

¹ Departamento de Edafología, Instituto de Geología, Universidad Nacional Autónoma de México, 04510 Ciudad de México, (México).
² Department of Soil Science and Agricultural Chemistry, Escuela Politécnica Superior, Universidad de Santiago de Compostela, 27002 Lugo (Spain).
³ Instituto de Investigaciones en Ecosistemas y Sustentabilidad, Universidad Nacional Autónoma de México, 58090 Morelia (México).
⁴ Escuela Nacional de Estudios Superiores Campus Morelia, Universidad Nacional Autónoma de México, 58090 Morelia (México).
⁵ Laboratorio Nacional de Mineralogía y Geoquímica (LANGEM), 04510 Ciudad de México (México)

*Corresponding author. Felipe García-Oliva³. e-mail: fgarcia@cieco.unam.mx
Abstract

Background. Litter decomposition is a key process in the functioning of forest ecosystems, because it strongly controls nutrient recycling and soil fertility maintenance. The interaction between the litter chemical composition and the metabolism of the soil microbial community has been described as the main factor of the decomposition process based on three hypotheses: substrate-matrix interaction (SMI), functional breadth (FB) and home-field advantage (HFA).

The objective of the present study was to evaluate the effect of leaf litter quality (as a direct plant effect, SMI hypothesis), the metabolic capacity of the microbial community (as a legacy effect, FB hypothesis), and the coupling between the litter quality and microbial activity (HFA hypothesis) on the litter decomposition of two contiguous deciduous oak species at a local scale.

Methods. To accomplish this objective, we performed a litterbag experiment in the field for 270 days to evaluate mass loss, leaf litter quality and microbial activity in a complete factorial design for litter quality and species site.

Results. The litter of *Quercus deserticola* had higher rate of decomposition independently of the site, while the site of *Q. castanea* promoted a higher rate of decomposition independently of the litter quality, explained by the specialization of the soil microbial community in the use of recalcitrant organic compounds. The Home-Field Advantage index was reduced with the decomposition date (22% and 4% for 30 and 270 days, respectively).

Discussion. We observed that the importance of the coupling of litter quality and microbial activity depends on decomposition stage. At the early decomposition stage, the home-advantage hypothesis explained the mass loss of litter; however, in the advanced decomposition stage, the litter quality and the metabolic capacity of the microbial community can be the key drivers.
Key words: Litter decomposition, *Quercus*, Differential Scanning Calorimetry, 13C NMR, Enzymatic activity

1. Introduction

Litter decomposition is a key process in the functioning of forest ecosystems, because it strongly controls nutrient recycling and soil fertility maintenance (Austin et al., 2014). At the local scale, the decomposition rate is strongly affected by litter traits and microbial activity (Freschet et al., 2012). The litter traits that promote the decomposition are related with physical features as the rate of water uptake in litter (Makkonen et al., 2013). Additionally, some chemical characteristics of litter can promote its decomposition such as: a) a low C: N ratio (Ågren et al., 2013; Aponte et al., 2013; Bononomi et al., 2013; Osono et al., 2013), b) a high concentration of soluble organic forms (Berg, 2014), and c) a low proportion of lignin or phenolic compounds (Almendros et al., 2000; Prescott 2010; Ono et al., 2011), as well as changes in the proportions of lignin subunits (Chávez-Vergara et al., 2014; Talbot et al., 2012). The study of the effects of leaf litter traits on decomposition has been reported before by several authors (i.e., Grime and Anderson, 1986; Bass et al., 1989, Grime et al., 1996); and more recently, these traits as called “after life” traits, because they are products of the metabolism of living plant species, and they can regulate ecological process, as litter decomposition (Genung et al., 2013).

The decomposition rate of organic compounds is also associated with the composition of the soil microbial community and its metabolism (Austin et al., 2014; Freschet et al., 2012). For example, the presence of actinomycetes (Snajdr et al., 2011) and basidiomycetes species (Osono & Takeda 2002; Snajdr et al., 2011) favors the degradation of recalcitrant compounds (i.e. lignin,
polyphenols, aliphatics), because these microbial taxa are capable of producing exoenzymes which can cleave these organic molecules (Allison et al., 2014). Consequently, the inhibitory effect on litter decomposition of a high proportion of recalcitrant molecules can be reduced by the activity of specialized microbial species (Cleveland et al., 2004; Strickland et al., 2009; Snajdr et al. 2011; Chávez-Vergara et al., 2016). Therefore, the metabolic capacity of the microbial community can be considered as a “legacy” effect over litter decomposition (Wurst & Ohgushi, 2015). These authors defined legacy effect as “a specific case of long-term effects that persist after the biotic interaction that caused the effects ceases”. Recent studies have shown that the interaction between the chemical composition of the plant residues and the metabolic capacity of the microbial community is the most important factor in the regulation of the litter decomposition rate (Austin et al., 2004; Ayres et al., 2009; Fanin et al., 2016; García-Palacios et al., 2016; Hicks Pries et al., 2017). This interaction involves the functional traits of plant species (i.e. chemical characteristics of plant residues) and the activity of the microbial community of the forest floor (i.e. production of exoenzymes) (Ayres et al., 2009; Austin et al., 2014; Pearse et al., 2014; Fanin et al., 2016).

The coupling between litter chemical composition and metabolism of the microbial community of the forest floor has been described by the following hypotheses: A) home-field advantage (HFA), which states that the litter will be more easily decomposed by the microbial community in the same site where it was produced (Ayres et al., 2009, Austin et al., 2014); B) substrate-matrix interaction (SMI), in which exogenous litter can be decomposed at the same rate than endogenous litter if both have a similar chemical composition (Freschet et al., 2012) and, more recently, C) the functional breadth hypothesis (FB), according to which microbial communities that have been exposed to substrates with low chemical quality have developed
mechanisms for the use of substrates with different chemical quality; in other words, are functionally more diverse, and capable of using a wide-range of substrates (Fanin et al., 2016).

Therefore, the objective of the present study was to evaluate the effect of leaf litter quality (as a direct plant effect, SMI hypothesis), the metabolic capacity of the microbial community (as a legacy effect, FB hypothesis), and the coupling between leaf litter quality and microbial activity (HFA hypothesis), on the litter decomposition of two species of deciduous oaks, by using a controlled field experiment of litter decomposition.

In previous studies, we found that *Quercus deserticola* promoted higher nutrient availability than *Q. castanea*, because the former oak species produced leaf litter with higher chemical quality, therefore favoring microbial activity and litter chemical transformation (Chávez-Vergara et al., 2014; Chávez-Vergara et al., 2015). However, the microbial community in the litter of *Q. castanea* is dominated by microbial species specialized in the use of recalcitrant compounds, increasing the efficiency in the use of resources (Chávez-Vergara et al., 2016). Therefore, the main hypothesis of the present study is that the litter decomposition is regulated by the direct effect of the chemical composition of the plant residues, and the legacy effect on the specialization of the microbial community. Consequently, the site dominated by *Q. castanea* should have a higher potential of litter decomposition, but the *Q. deserticola* litterfall should be easier to decompose. Our study is the first report testing hypotheses on litter decomposition in species of the same genus, while most of the studies have been performed on taxonomically and functionally very distant plant species (i.e., Freschet et al., 2012; Pearse et al., 2014; Fanin et al., 2016).

To test our hypothesis, we performed a factorial field experiment of decomposition bags during 30 and 270 days with litterfall of *Q. castanea* (low quality), *Q. deserticola* (high quality).
and a mix of both oak species litterfall (cumulative quality) in three sites based on the microbial activity specialization: fast degradation of recalcitrant compounds (under Q. castanea), slow degradation of recalcitrant compounds (under Q. deserticola) and an intermediate degradation, which represents a wider spectrum of resources utilization for the microbial community (under both Quercus species in interaction).

2. Materials and methods

2.1. Study site

This study was conducted within the Cuitzeo basin in El Remolino hill (19° 37' 01"N, 101° 20' 07"W; 11 km south of Morelia city, Michoacán, Mexico). The study site is an oak forest fragment (>12 ha) with low disturbance (about 80 years without wood extraction for charcoal production according to nearby inhabitants) and two dominant native oak species: Quercus castanea Née (section Lobatae) and Q. deserticola Trel. (section Quercus). The characteristics of this site and the species can be found in more detail in previous studies (Chavez-Vergara et al., 2014; Chávez-Vergara et al., 2015). Briefly, the predominant soil type is a chromic Luvisol developed over Quaternary basalts. The climate in the area is temperate subhumid, with annual mean temperature of 17.6 °C and annual mean precipitation of 805 mm concentrated in the summer months. In 2014, the annual rainfall was 850 mm and the average temperature was 16.6 °C. For the present study, three parallel plots of 30 X 150 m were established perpendicular to the main slope, where one of the studied species dominated in either of the two lateral plots; and both species were mixed in the central plot. Therefore, three species conditions were present: isolated Q. castanea (Qc), isolated Q. deserticola (Qd) and mixed Quercus species (Qx).
2.2. Litterfall collection

A circular trap of 0.5 m$^2$ was placed under each of five trees per species condition: isolated *Quercus castanea*, mixed species and isolated *Q. deserticola* (15 traps in total) to collect litterfall every month from December 2012 to May 2014. The fresh litter samples were weighed, and an aliquot was dried to constant weight at 70 °C for 72 h to determine water content, which was then used to calculate the dry mass of each sample. The fresh aliquot was stored at 4 °C in darkness prior to laboratory analysis (n = 5 for each species). The monthly dried subsamples from the two sampling years were mixed for the litterbag experiment.

2.3. Litterbags experiment

2.3.1. Field experiment

Brown color polyester mesh (1 mm) bags (10 X 10 cm) were used for the field decomposition experiment. The mesh size of 1 mm was chosen because it avoids losing small leaf litter debris but allows the activities of the aerobic microbial community and meso- and micro-fauna (Nguyen Tu et al., 2011), which play an important role in the initial fragmentation of litter (Gessner et al., 2010). These bags were filled with 11 g of oven-dried litterfall with the following arrangement: 30 bags with *Q. castanea* litterfall (QcL), 30 bags with *Q. deserticola* litterfall (QdL) and 30 bags with a mixture of both species litterfall (QxL) in the same proportion (5.5 g of *Q. castanea* and 5.5 g of *Q. deserticola* litterfall). In June 2014, two litterbags of each litterfall type (QcL, QdL and QxL) were randomly located above the litter and around the stem of each of the five-selected trees, distributed along the main slope, in each species condition plot (hereafter plots are referred as sites): *Q. castanea* site (QcS), *Q. deserticola* site (QdS) and the species mixture site (QxS). Therefore, the field design is a complete factorial 3 X 3 (site and litterfall condition).
conditions). One litterbag for each treatment per tree was harvested at 30 and 270 days after the bags were placed (5 bags for each treatment). The comparison of the two dates allows us to determine the decomposition effect on early and late decomposition stages, where the labile and recalcitrant molecules proportion changes over-time. The means (± standard deviation) of DBH for trees in each condition was Qc: 52.9 ± 11.7 cm, Qx: 48.9 ± 4.9 cm and Qd: 63.9 ± 11.4 cm.

In the collection dates, the content of each bag was carefully removed, fresh field weighed, and subsequently divided into two subsamples. The first one was stored in hermetic bags in the dark at 4 °C until laboratory analysis. The second subsample was dried to constant weight at 70 °C for 72 hours to calculate the water content. Then, the sample was milled in a ball mill at 350 RPM for 3 min and stored in sealed bags until chemical analysis.

2.3.2. Remaining mass and decomposition rate

Initial and remaining samples were combusted in a muffle furnace at 650 °C to determine inorganic particles to correct data on an ash-free basis. The litter decomposition rate was calculated using the simple exponential model: \( MR = M_i e^{-kt} \); where \( MR \) is the percentage of the remaining mass at 270 days, \( M_i \) is the initial mass percentage, \( k \) is the decomposition rate and \( t \) is the decomposition time in field conditions (Olsen, 1963).

2.4. Laboratory analysis

The nutrients and enzymatic activity analyses were done at Instituto de Investigaciones en Ecosistemas y Sustentabilidad, UNAM, Mexico, while the \(^{13}\text{C} \) Nuclear Magnetic Resonance and DSC analyzes were done at Universidad de Santiago de Compostela, Spain.

2.4.1. \(^{13}\text{C} \) Nuclear Magnetic Resonance (\(^{13}\text{C} \) NMR) spectroscopy
The $^{13}$C Nuclear Magnetic Resonance spectroscopy is a non-destructive analysis that improves the identification of the molecular composition from organic residues; it is useful tool for determination of the molecular composition of litterfall and decomposed litter. To characterize the chemical composition of litterfall and decomposed litter, the analysis of Cross Polarized Magic-Angle Spin $^{13}$C NMR in solid state was performed in samples previous to the field experiment (initial) and in samples at the end of the field experiment (remaining). The $^{13}$C NMR data were obtained at 298 K in a Varian Inova-750 17.6 T (operated at 750 MHz frequency proton), under the conditions described in Chávez-Vergara et al. (2014). The spectrogram obtained was processed with the program MestreNova V. 6 (Mestrelab Research Inc.).

For integration, the spectrogram was divided into four major regions representing different chemical environments of $^{13}$C nucleus according to position of relaxation signal in parts per million of chemical shift (ppm): C Alkyl (0-45 ppm), O-alkyl C (45-110 ppm), aromatic C (110-160 ppm) carbonyl and C (160-220 ppm). For more detailed analysis, spectra were divided according to Leifeld & Kögel-Knabner (2005) as: I) 10-45 ppm C alkyl: methyl groups, methylene groups on rings and aromatic chains. II) 45-110 ppm C O-alkyl: methoxy groups and C6 in some polysaccharides (45-60 ppm); C2-C5 hexoses C of some amino acids, aliphatic alcohols and fractions of lignin structure (60-90 ppm); Carbohydrate anomeric C, C2-C6 syringyl unit of lignin (90-110). III) 110-160 ppm aromatic C: and CC and CH carbon C2 guaiacil, C6 lignin (110-140 ppm, aryl C); COR aromatic or CNR (140-160 ppm, phenolic C) groups. IV) carboxyl 160-220 ppm C: carboxyl C, C carbonyl and C amide.

We also examined indexes associated with the decomposability of organic matter based on integrated specific regions: alkyl: O-alkyl ratio (A: OA), O-alkyl: aromatic ratio (OA: Ar), aromaticity (Ai), hydrophobicity (HB: HI) and characterization of lignin relations based on
subunits specific regions such as syringyl (S), guaiacyl (G) and p-hydroxyphenyl (H) as lignin relations S:G, S:H and G:H (Almendros et al., 2000; Spaccini et al., 2006; Talbot et al., 2012; Bonanomi et al., 2013; Chávez-Vergara et al., 2014).

2.4.2. Differential scanning calorimetry (DSC) and Thermogravimetry (TGA)

The Differential Scanning Calorimetry (DSC) and Thermogravimetry (TG) is a thermal analysis suitable for determination of organic matter stability (Angehrn-Bettinazzi et al., 1998). This method quantifies the energy release during different combustion temperatures of samples, like the energy required for biological oxidation of organic molecules (Rovira et al., 2008). Therefore, the thermograms can quantify the proportion of labile, recalcitrant and extra-recalcitrant compounds in the organic samples (Barros et al., 2007). The characterization of thermal properties of litterfall and decomposed litter was done by differential scanning calorimetry and thermogravimetric analysis (DSC-TGA, Mettler-Toledo International Inc.). The analysis was performed with 4 mg of powdered oven-dried sample placed in an aluminum pan in an atmosphere of dry air (flow rate, 50 ml min\(^{-1}\)) and the scan rate was 10 °C min\(^{-1}\). The temperature range used was 50 to 600 °C. An indium sample (melting point: 156.6 °C) was used to calibrate the calorimeter. All samples were analyzed in triplicate.

The combustion heat release \((Q, \text{ J g}^{-1})\) was determined by integrating the DSC curves \((W \text{ g}^{-1})\) on the exothermic region (150-600 °C). Data recorded at temperatures <150 °C were discarded because they are associated with the loss of mass and energy release during moisture loss. The Q value was divided by the mass loss in each measurement \((Q', \text{ J mg}^{-1} \text{ MO}, \text{ Rovira et al.}, 2008)\).
Areas under the DSC curve were divided in three groups, representing different degrees of resistance to thermal oxidation (Dell'Abate et al., 2002; Fernandez et al., 2012): labile organic matter, comprising carbohydrates and other aliphatic compounds (200-375 °C); recalcitrant organic matter such as lignin and / or polyphenols (375-475 °C); and extra-recalcitrant organic matter, such as polycondensed aromatic forms (475-550 °C). The heat release by combustion in each region was designated as \( Q_1 \), \( Q_2 \) and \( Q_3 \), respectively. Also, the temperature at which the maximum heat flow was detected during the combustion of organic matter (\( T_1 \), \( T_2 \) and \( T_3 \)) and the temperature at which 50% of the total energy was released were recorded (\( T_{50 \text{DSC}} \)).

### 2.4.3. Nutrient analysis

All forms of C were determined on a total carbon analyzer (UIC model CM5012, Chicago, USA) by dry combustion and coulometric detection (Huffman, 1977), while forms of N were determined colorimetrically by the semi-Kjeldahl (Bremmer, 1996) method and the forms of P by molybdate colorimetric method after reduction with ascorbic acid (Riley & Murphy, 1962) in a Bran-Luebbe autoanalyzer (Autoanalyzer 3 Norderstedt, Germany) after acid digestion. The litterfall chemical analyses were performed from material collected from the respective traps (isolated *Q. castanea*, mixed species, and isolated *Q. deserticola*).

Soluble organic forms of C, N and P were extracted from 2 g of fresh material in deionized water, after stirring for 1 hour and filtered through a Whatman# 42 filter and on a vacuum system through a 0.45 µm nitrocellulose membrane. The dissolved organic carbon (DOC) was determined by combustion coulometric detection (Huffman, 1977). Dissolved organic nitrogen (DON) and dissolved organic phosphorus (POD) were determined after acid digestion. The DON was calculated as the difference between the acid digested nitrogen and
 soluble NH$_4^+$, as well as POD (acid digested P minus soluble inorganic P; Joergensen & Mueller, 1996).

Microbial biomass carbon (Cmic) and nitrogen (Nmic) were determined by direct extraction using chloroform fumigation (Brookes et al., 1984; Vance et al., 1987) from fresh samples. Two subsamples (2 g) were incubated at 30 °C for 24 h; one of the subsamples was maintained in chloroform atmosphere during incubation. Both samples were extracted in 0.5 M K$_2$SO$_4$ and percolated through # 42 Whatman filter paper and analyzed as DOC (as mentioned above) and removable N. The extractable N was quantified as total N after acid digestion (Brookes et al., 1984).

The microbial biomass P (Pmic) was determined by the fumigation-extraction method (Brookes et al. 1982) in fresh samples. A subsample of 0.5 g was fumigated for 24 h in a chloroform atmosphere and extracted with 30 mL of 0.5 M NaHCO$_3$ pH 8.5 for 30 min (van Meeteren et al., 2007). Extracts (fumigated and no-fumigated) were digested in a solution of sulfuric acid and persulfate ammonium according to Hedley sequential P fractionation (Tiessen & Moir, 1993) and quantified as orthophosphate, as described above. Cmic concentration, Nmic and Pmic were calculated from the difference between the fumigated and non-fumigated samples, then the concentration was corrected by applying the following factors: KeC 0.45 (Sparling et al., 1990), KeN 0.54 (Brookes et al., 1984; Jorgensen & Mueller, 1996) and KeP 0.40, respectively (Brookes et al. 1982).

2.4.4. Enzyme activity

As a measurement of microbial activity related with the use of organic molecules, the enzymatic activities of β-1,4-glucosidase (BG), cellobiohydrolase (CBH), β-N-acetyl-glucosaminidase
(NAG), polyphenol oxidase (POX) and dehydrogenase (DHG) were determined in fresh material of each treatment for all samples collected. The determination of hydrolases (BG, CBH and NAG) was performed according to Chávez-Vergara et al. (2014) by colorimetric measurement of p-nitrophenol (pNP) in a spectrophotometer (Evolution 201, Thermo Scientific Inc.) at 420 nm liberated from specific substrates during incubation (2 h in oscillatory shaking at 30 °C) and reported in g-mol pNP litter⁻¹ h⁻¹.

POX activity was determined through oxidation of 2, 2'-Azinobis[3-ethylbenzothiazoline-6-sulfonic acid]-diammonium salt (ATBS). One aliquot of the same extraction used for activity of hydrolases (Chávez-Vergara et al., 2014) was used, volume and time of preparation were the same than for hydrolases but in this case the result of the centrifugation was measured directly (without addition of NaOH and deionized water) on the same spectrophotometer described above by colorimetry at 460 nm and the result was reported as mol tyrosine g-litter⁻¹ h⁻¹, in this case the calibration curve is derived from tyrosine (tyr).

The dehydrogenase activity (DHG) was determined according to a modification of the method described by Alef (1995), which is based on the reduction of chloride of 2, 3, 5-triphenyltetrazolium (CTT) for the formation of triphenyltetrazolium formazan (TFF) in incubation at 30 °C for 24 h. To perform the assay 0.5 g of milled fresh material was weighed and placed in a conical tube with a capacity of 15 mL (the tube was covered with foil to keep out light) to which 1 mL of a solution CTT 1% in Tris buffer pH 7.6 was added. Blank samples were prepared only with the solution of CTT. Subsequently, all samples were placed and horizontally fixed in an incubator chamber; the incubation was for 24 h at 30 °C and 180 rpm. After incubation, 10 mL of acetone was added; the tube was stirred vigorously and allowed to react for 2 h in the dark at room temperature. Following, the supernatant was filtered through Whatman #
42 paper and measured at 546 nm by colorimetry. The measure of each sample was subtracted from the average value of blanks and adjusted by the equation described in Alef (1995). The results were expressed in g TPF g$^{-1}$ d$^{-1}$.

The efficiency of enzymatic activity according to the concentration of a nutrient immobilized in the microbial biomass was calculated as specific enzyme activity (SEA) according to the following equation (Chávez-Vergara et al., 2014):

$$\text{SEA} = \frac{A}{B_{mic}}$$

where SEA is expressed in mol of pNP or mol of tyr released per milligram of nutrient in the microbial biomass per hour (mol mg$^{-1}$ Bmic h$^{-1}$); A is the activity of any of the specific enzymes (BG, CBH, NAG, POX and DHG), and Bmic is the concentration of Cmic or Nmic in mg g$^{-1}$.

The association of enzymes with nutrients in the microbial biomass is as follows: BG, CBH, POX and DHG are associated with Cmic, and NAG with Nmic.

2.5. Statistical analysis

The initial nutrient concentration in the three litterfall conditions had five replicates, while the DSC and the $^{13}$C NMR parameters had only one composite sample for each Quercus species, as well as the DSC and $^{13}$C NMR parameters for the samples at 270 decomposition days. In contrast, the DSC parameter of the 30 decomposition days had a composite sample for each litter condition (litter origin and litter site, in total 9 samples). Therefore, the nutrient concentration of litterfall and DSC parameters according to litter quality and site effects at 30 days of decomposition were analyzed using one-way analyses of variance (ANOVA), while the remaining mass, total nutrient concentration and dissolved nutrients were analyzed with repeated measures ANOVA (RMANOVA), in which the between factor was litter species condition (QcL,
QxL and QdL) and the within factors where sampling date (litterfall, and litter after 30 and 270
days of field decomposition, n=5) and the interaction between litter condition and decomposition
time. The data of nutrients, enzymatic activities and specific enzymatic activities at 30 and 270
days of experiment were analyzed by a factorial ANOVA to test the effect of litter condition
(QcL, QdL and QxL) and site (QcS, QdS and QxS) in early and late decomposition. In this case,
all samples associated with each treatment in the factorial design (n = 15) were used. Afterwards,
statistically significant differences in one-way, RMANOVA and factorial ANOVA were
analyzed with the Tukey HSD post-hoc test.

To identify the relationship between the chemical quality of the litter and microbial
metabolism variables in the remaining mass at 30 days of decomposition, we performed
backward stepwise multiple regression analyses. As chemical quality variables, the C:N and C:P
ratios, dissolved forms of C, N and P were used, while as microbial activity variables the specific
enzymatic activities were used for the multiple regression model. We performed a correlation
model using remaining mass and thermal parameters from the DSC analysis as indicators of litter
changes during early decomposition process. All analyses were made in the statistical package
Statistica 7.0 (StatSoft, USA).

The Home-Field Advantage Index (HFAI) proposed by Ayres et al. (2009) was calculated as the
percentage of mass loss of each litterfall condition (QcL, QXL, QdL) at the site where it was
produced relative to the mass loss at all sites (QcS, QXS, QdS):

\[ A_{RMLa} = \frac{Aa}{Aa + Ba + Ca} \times 100 - 100 \]
where, $A_{RML_a}$ represents the relative mass loss of litter from condition A at site a, and Aa, Ba and Ca represent the mass loss of litter from conditions A, B and C decomposing at site a, respectively. The measures of relative mass loss for each condition and site combination were used to calculate HFAI.

$$\text{HFAI} = \left[ \frac{A_{RML_a} + B_{RML_b} + C_{RML_c}}{3} \right] \times 100$$

3. Results

3.1. Mass loss and residence time

Figure 1A shows that *Q. deserticola* had higher decomposition rate at all harvest dates in 270 days of field decomposition experiment, but not at 30 days (Fig. 1A), and the estimated residence time of litter for the three conditions was 1.2, 1.3 and 1.9 years for *Q. deserticola* (Qd), mixed species (Qx) and *Q. castanea* (Qc), respectively. We observed that at 30 days of decomposition, the slope of mass loss was higher in Qd and Qx than in Qc (Fig. 1A). For this reason, we analyzed the mass loss at 30 and 270 days with a factorial ANOVA to identify the litter and site effect over mass loss. We observed that the mass loss after 30 and 270 days showed differences for the two main factors analyzed in opposed ways: Qd litterfall (QdL) promotes higher mass loss (Fig. 1B), while in the Qd site (QdS) lower mass loss was observed (Fig. 1C). The Home-Field Advantage index was 22% and 4% for 30 and 270 days, respectively.

3.2. Chemical composition of litterfall and decomposed litter

3.2.1. $^{13}$C CP MASS NMR characterization of litterfall samples
Figure 2 shows the spectra and Table 1 the integration of regions of $^{13}$C CPMAS NMR of litterfall and decomposed litter at 270 days in both species. In the original litterfall, the most prominent compounds were O-Alkyl, Aryl C and Alkyl C. Particularly, we observed that the region between 160 and 220 ppm, assigned to carboxyl/amide and carbonyl C groups, is dominated by a peak at ca. 173 ppm (Fig. 2) with similar relative C distribution for both species (Table 1). In the aromatic and phenolic region (160–110 ppm), the litterfall produced by both species showed well defined peaks at ca. 153 and 145 ppm, which revealed the presence of C3, C5 syringyl lignin and tannins, respectively. An incipient peak at ca. 131 in Q. deserticola and well-defined in Q. castanea may be related to unsubstituted and C-substituted phenyl carbon of lignin monomers of syringyl units (De Marco et al., 2012) affecting the S:G ratio between species (Table 1). In the O-alkyl region (45-110 ppm), the most prominent signal was at ca. 73 ppm and it was particularly associated to the simultaneous resonance of C-2, C-3 and C-5 of pyranose rings in cellulose and hemi-cellulose; and a second prominent peak was at ca. 105 ppm, traditionally associated to crystalline cellulose, and it can be associated to non-protonated carbon arising from tannins (Almendros et al., 2000).

A shoulder at 56 ppm, attributable to N-methoxyl C compounds in lignin, was relatively well resolved in litter samples of both species. Meanwhile, the C-6 position in the pyranose ring of cellulose produced the peak or shoulder at ca. 64 ppm better resolved in Q. castanea. In a similar way, that shoulder at 84 ppm may correspond to C-4 in cellulose (Almendros et al., 2000). The alkyl region (46-0 ppm) showed two well-defined peaks at 30 and 21 ppm. The peak at ca. 30 ppm is related to polyethylene carbons in lipids and lipid polymers such as cutine or suberine and the peak at ca. 21 ppm is frequently attributed to acetate groups in hemicellulose and/or short chain aliphatic structures. The most noticeable changes after 270 days of...
decomposition occur in the reduction of peak intensity in ca. 145 ppm associated to tannins and the increment of peaks at ca. 56 ppm (N-methoxyl) and two peaks in the alkyl C region (Fig. 2). These changes were most intense in the *Q. deserticola* litter.

3.2.2. Thermal characteristics of litterfall samples by differential scanning calorimetry (DSC)

The litterfall thermograms of both species showed a bimodal shape (Figs. 3A and 3B). The first peaks were situated at 347°C and 352°C, and the second prominent peaks were at 449°C and 461°C for *Q. castanea* and *Q. deserticola*, respectively (Table 2). Additionally, the Qc litterfall had a second peak at 424°C in the Q2 region. However, the energy released from the recalcitrant region (Q2) was higher in the *Q. castanea* than in the *Q. deserticola* litterfall (Table 2). After 30 days of decomposition, the percentage of recalcitrant compounds (Q2) was higher in the Qc litter than in the Qd litter, while the Qd litter had the highest percentage of extra-recalcitrant compounds (Q3; Table 2; Fig. 3A and 3B). In the 270 days decomposed litter, the shape of the thermogram of *Q. castanea* remained as bimodal, while for *Q. deserticola* the third peak at the extra-recalcitrant region is well-defined and displaced to a higher temperature (503°C; Fig. 3). The decomposed litter at 270 days increased the energy released, compared to 30 days, in the Q1 region (Qc +4% and Qd +2%), the Q2 region showed a decrease in the litter of both species (Qc -7% and Qd -4%; Table 2) and the Q3 region increased for both species but more for *Q. castanea* (Qc +63% and Qd +15%; Table 2). The values of T50 were displaced to lower temperatures (Table 2) in both species at 270 days of decomposition in comparison to 30 days of decomposition.

In the one-way ANOVA of the thermal parameters at 30 days of decomposition, recalcitrant and extra-recalcitrant compounds were the highest and the lowest in QcL and QdL, respectively, without a significant effect of site (Table 3). Figure 4 shows the thermograms of
litter at 30 days of decomposition of each *Quercus* species condition within each site. The QdL showed the highest peak in the labile region at the three sites, while the QcL showed the highest peak in the recalcitrant region also in the three sites. However, only the QdL showed a small peak in the extra-recalcitrant region, mainly at the Qd site.

3.2.3. Nutrient concentration and stoichiometric ratios in litterfall and decomposed litter. The concentration of C was only affected by decomposition time (F= 3.5; p < 0.001) reducing its value at 270 days of decomposition (Table 4). In contrast, the interaction between litter quality and sampling date was significant for concentrations of N and P (F=5.2, p= 0.01 and F=5.5, p= 0.009, respectively). The QdL had higher N concentration in all decomposition dates in comparison with QcL and QxL (Table 4). However, the QxL and the QdL had higher P concentration than QcL at 30 and 270 days of decomposition, but these values were similar in the three species litterfall (Table 4). These results suggest that the QxL and QdL had microbial P immobilization after 30 days of decomposition. Therefore, the C:N and C:P ratios were affected by litter quality and sampling date (F= 25, p = 0.001 and F = 8.6, p= 0.004, respectively); the QcL had the highest values and the 270 days of decomposition had the lowest values for the three species conditions (Table 4). However, the N:P ratio was only affected by decomposition time (F=17, p<0.001), where the highest and the lowest values were in the litterfall and at 30 days of decomposition, respectively (Table 4).

In contrast, the dissolved organic carbon and nitrogen (DOC and DON, respectively) concentrations were affected by the interaction of litter quality and sampling date (F=18, p<0.001 and F= 21, p<0.001, respectively). The QdL had higher DOC concentration than the QcL in the two decomposition dates (Table 4), although the QxL had a higher DOC reduction at 270 days of decomposition than the QcL (64% and 30%, respectively). In contrast, the QcL had
lower DON than the QxL and QdL in the litterfall and the litter at 270 days of decomposition, but the three litters had no different DON values at 30 days of decomposition (Table 4).

Additionally, the QdL showed a reduction of DON concentration only at 270 days of decomposition, while the litter of the two other species conditions had increments or no changes in relation to their litterfall (QdL and QxL, respectively). The DOP concentration was affected by both litter quality and sampling date ($F=18$, $p=0.002$ and $F=26$, $p<0.00001$, respectively); the QdL and QcL had the highest and lowest DOP concentration, respectively, and the highest and the lowest DOP values were at 30 and 270 decomposition days, respectively (Table 4).

Similarly, the dissolved inorganic forms of N ($\text{NH}_4^+$ and $\text{NO}_3^-$) were also affected by the interaction between litter quality and sampling date ($F=66$, $p<0.001$ and $F=9$, $p=0.002$, respectively). Both dissolved $\text{NH}_4$ and $\text{NO}_3$ concentrations were highest in the Qd and lowest in the Qc in the litterfall, but this pattern changed at the 270 days of the experiment for $\text{NH}_4^+$ concentration ($\text{QcL}>\text{QdL}=\text{QxL}$; Table 4) and for $\text{NO}_3^-$ concentration ($\text{Qd}>\text{QxL}>\text{QcL}$; Table 4).

The dissolved inorganic P form (DiP) was affected by the main factors ($F=66$, $p<0.001$ and $F=26$, $p<0.001$ for litter quality and sampling date, respectively); the QdL and QcL had the highest and lowest DiP values, respectively, and the lowest DiP values were at 30 days of decomposition (Table 4).

3.3. Microbial activity in decomposed litter.

The concentrations of microbial immobilized C, N and P ($\text{Cmic}$, $\text{Nmic}$ and $\text{Pmic}$, respectively) were highest in Qd, followed by Qx and lowest in Qc (Fig. 5) for litter decomposed on its original site. However, the C:N, C:P and N:P microbial ratios were not affected by any factor
analyzed (Fig. S1). The specific enzymatic activity of β-glucosidase (SEA BG), polyphenol oxidase (SEA POX) and dehydrogenase (SEA DHG) showed the highest values in Qc and Qx and the lowest values in Qd for litter decomposed on its original site (Fig. 6). At 30 days of decomposition, the microbial immobilization of C responded to the main factors (litter condition and site effects). The Qc site (QcS) had lower Cmic concentration than the other two sites (Fig. 5), while the Qd litter (QdL) had five-fold higher Cmic concentration than Qc (Fig. 5). However, the litter conditions promoted only differences for Nmic and Pmic. In both cases, the QdL had the highest concentrations, followed by the QxL, and the QcL had the lowest concentration values (Fig. 5).

SEA DHG was affected by both main factors (site and litter); the QcS and litter had the highest values (Fig. 6), SEA POX values were only affected by site, showing higher values in the QcS than in the QdS (Fig. 6). Meanwhile the value of SEA BG was only influenced by litter condition, with the QdL showing the lowest values (Fig. 6).

3.4. Relation of variables with remaining mass

The thermal parameters (heat released in the combustion from the DSC analysis) at 30 days of decomposition had relationships with remaining mass; the Q2 (375-475 °C) region was positively related \( r=0.77, p=0.025 \), while the Q3 region (475-550 °C) was negatively related \( r=-0.82, p=0.001 \). Therefore, these parameters can be used as indicators of the intensity of the decomposition process of litter (Table 5). Also, in the multiple regression model, the remaining mass at 30 days was positively explained by the C:N ratio and negatively explained by DON, DOP and dissolved NH\(_4^+\) \( R^2 = 0.78, p<0.001; \) Table 5). In addition, the remaining mass showed a positive relation to SEA DHG and a negative relation to SEA POX \( r^2 = 0.38, p=0.033; \) Table 5).
4. Discussion

Our results indicate that the factors which regulate litter decomposition are strongly affected by the decomposition date. At the early decomposition stage (30 days) when the labile molecules dominated, which regulate the decomposition rate (Berg, 2014), the coupling of litter quality and microbial activity (Home-Field Advantage hypothesis) is the main factor. However, at the advanced decomposition stage (270 days) when recalcitrant molecules dominated, the litter decomposition is regulated by the direct effect of the chemical composition of the plant residues (Substrate-Matrix Interaction hypothesis) and the legacy effect on the specialization of the microbial community in the use of organic compounds (Functional Breadth hypothesis). These conclusions are supported by the reduction of the Home-Field Advantage index with the decomposition date (22% and 4% for 30 and 270 days, respectively). Therefore, the hypotheses that have been raised to explain the process of decomposition of the litter are not mutually exclusive (Freschet et al., 2012; Fanin et al., 2016), which is only observable through cross-sowing experiments such as the one elaborated in the present study.

For example, the litter of *Q. deserticola*, when decomposed under its own canopy, showed a greater mass loss after 30 and 270 days as expected by the Home-Field Advantage hypothesis. These results are explained by a higher concentration of total nutrient and dissolved organic forms concentrations, and lower concentration of recalcitrant compounds than in the other two sites, conditions which promotes microbial activity (Almendros et al., 2000; Aponte et al., 2013, Bonanomi et al., 2013; Fanin et al., 2013; Freschet et al., 2013; Osono et al., 2013). However, when analyzing by means of a factorial design the mass loss on the analyzed dates (30 and 270 days of decomposition), we observed that this variable is explained by the type of litter
and by the site where it decomposes, since the material of *Q. deserticola* (QdL) loses more mass independently of the site where it decomposes, but it is in the sites of *Q. castanea* (QcS) and in the species mixture (QxS) where more mass is lost regardless of its origin. These results, suggest that the condition of the two other sites (QcS and QxS) decrease the importance of the Home-Field Advantage hypothesis, mainly at 270 decomposition days.

The mechanisms that explain the higher rate of mass loss in the litterfall of *Q. deserticola* are probably related to its chemical composition, since the higher concentrations of COD and N and P in soluble forms decrease the investment in energy of the microbial community for the production of exo-enzymes to obtain organic compounds of low molecular weight (Baldrian et al., 2010; Glanville et al., 2012; Allison et al., 2014).

In contrast, in places with natural incorporation of low quality material for decomposition, as the Qc site, the microbial community makes metabolic adjustments related to the chemical characteristics of the litter, which is also a key factor in the rate of decomposition. In this regard, we observed that when leaf litter of better chemical quality (QdL) is incorporated in the *Q. castanea* (QcS) site and in the mixed species site (QxS), it decomposes at a faster rate than in the site where it was produced, and also increases the immobilization of nutrients in the microbial biomass, mainly in the early decomposition stage (30 days). These results suggest that the microbial community of the litter under *Q. castanea* is more efficient in obtaining and using organic compounds, because of the continuous exposure to low quality litter (Van Meeteren et al., 2007; Baldrian et al., 2012; Allison et al., 2014; Chávez-Vergara et al., 2014; Chávez-Vergara et al., 2016), supporting the Functional Breadth hypothesis.

Although the litter of better chemical quality stimulated the immobilization of nutrients in the *Q. castanea* site, the microbial community of this site maintained a high SEA of POX and...
SEA of DG and did not modify the stoichiometric ratios in the microbial biomass. This indicates that the microbial community maintains carbon efficiency in a similar way than when it is exposed to the local litter, using more energy in the production of enzymes for the depolymerization of recalcitrant compounds than in the accumulation of biomass (Chávez-Vergara et al., 2014; Chávez-Vergara et al., 2016; Zederer et al., 2017). Therefore, we suggest that the greater availability of nutrients in the best quality litter (Q. deserticola) stimulates the growth of microbial populations, but nevertheless these populations maintain their ability to use recalcitrant compounds, in overall making the microbial community under Q. castanea more efficient in the decomposition of the litter, mainly at the advanced decomposition stage (270 days). This can be considered as a legacy effect on the microbial community of low quality compounds for decomposition (Fanin et al., 2016). In a previous work (Chávez-Vergara et al., 2016), we determined that the chemical composition of the litter influenced the composition of the fungal community. Under Q. castanea a greater proportion of basidiomycetes was observed, which have been reported to be specialized in the degradation of recalcitrant compounds (Osono & Takeda., 2002, Snajdr et al., 2011). Therefore, the composition of the microbial community reflects the physiological footprint of the plant (Wickings et al., 2012) and constitutes a legacy of the chemical traits of plant species (Wurst & Ohgushi, 2015; García-Palacios et al., 2016).

In general, we can suggest that the better litter quality regulates the accessibility of organic compounds for their use (Prescott 2010; Wickings et al., 2012; Freschet et al., 2013), while the microbial community, through its specialization, determines the efficiency in its use and therefore the speed of decomposition (Strickland et al., 2009; Snajdr et al., 2011; Cleveland et al., 2014). The above can explain the observed patterns in the thermal analysis of the decomposed material at 30 days. In the thermal analysis, we observed that it is at the Qc site that
the signal of the recalcitrant compounds is more intensely decreased, and it is at this site, as mentioned above, that there is a greater investment in the production of enzymes for the degradation of recalcitrant compounds. Likewise, it is at this site that a clear signal of reactive molecules derived from microbial metabolic activity is observed, which is inferred by the appearance of an exothermic peak (ca 500 °C) in the region of extra-recalcitrant compounds (Rovira et al., 2008). This suggests that it is at this site that a more intense transformation of microbial organic compounds occurs, more clearly detected with thermal analysis (DSC) than in the analysis of $^{13}$C NMR.

5. Conclusions

In this study, we observed that the importance of the coupling of litter quality and microbial activity depends on decomposition stage. At early decomposition stage, the Home-Field Advantage hypothesis explained the mass loss of litter; however, in the advanced decomposition stage, the litter quality and the metabolic capacity of the microbial community can be the key drivers, mainly under *Q. castanea* conditions (litter with low available nutrients). These results enhance our knowledge about the mechanisms that regulate the decomposition of the litter in oak deciduous forests.

Acknowledgements

The authors thank Rodrigo Velázquez and Ofelia Beltrán for field and laboratory assistance. We thank two Anonymous reviewers for comments on a draft of the manuscript. B. Chávez-Vergara
acknowledges the support from the Graduate Program in Biological Sciences of the National Autonomous University of México (UNAM).

Literature cited

Ågren GI, Hyvönen R, Berglund SL, Hobbie SE. 2013. Estimating the critical N:C from litter decomposition data and its relation to soil organic matter stoichiometry. Soil Biology and Biochemistry 67:312-318. DOI: 10.1016/j.soilbio.2013.09.010

Alef K. 1995. Dehydrogenase activity. In: Alef, K., Nannipieri, P. eds. Methods in Applied Soil Microbiology and Biochemistry. Academic Press, San Diego, California, 228–231.

Allison SD, Chacon SS, German DP. 2014. Substrate concentration constrains on microbial decomposition. Soil Biology and Biochemistry 79:43-49. DOI: 10.1016/j.soilbio.2014.08.021

Almendros G, Dorado J, González-Villa FJ, Blanco MJ, Lankes U. 2000. 13C NMR assessment of decomposition patterns during composting of forest and shrub biomass. Soil Biology and Biochemistry 32:793-804. DOI: 10.1016/S0038-0717(99)00202-3.

Angehrn-Bettinazzi C, Lüscher P, Hertz J. 1988. Thermogravimetry as a method for distinguishing various degrees of mineralization in macromorphologically defined humus horizons. Zeitschrift für Pflanzenernährung und Bodenkunde 151:177–183. DOI: 10.1002/jpln.19881510305
Aponte C, García LV, Marañón T. 2013. Tree species effects on nutrient cycling and soil biota: A feedback mechanism favoring species coexistence. *Forest Ecology and Management* 309:36-46. DOI: 10.1016/j.foreco.2013.05.035

Austin AT, Vivanco L, Gonzalez-Arzac A, Perez LI. 2014. There's no place like home? An exploration of the mechanisms behind plant litter-decomposer affinity in terrestrial ecosystems. *New Phytologist* 204:307-314 DOI: 10.1111/nph.12959

Austin AT, Yahdjian L, Stark JM, Belnap J, Porporato A, Norton U, Ravetta DA, Schaeffer SM. 2004. Water pulses and biogeochemical cycles in arid and semiarid ecosystems. *Oecologia* 141:221-235. DOI: 10.1007/s00442-004-1519-1

Ayres E, Steltzer H, Simmons BL, Simpson RT, Steinweg JM, Wallenstein MD, Mellor N, Parton WJ, Moore JC, Wall DH. 2009. Home-field advantage accelerates leaf litter decomposition in forests. *Soil Biology and Biochemistry* 41:606-610. DOI: 10.1016/j.soilbio.2008.12.022

Baas WJ. 1989. Secondary plant compounds, their ecological significance and consequences for the carbon budget: introduction to the carbon/nutrient-cycle theory. In: Lambers H, Cambridge ML, Konings H, Pons TL eds. *Causes and consequences of variation in growth rate and productivity of higher plants*. SPB Acad. Publ., The Hague, pp. 313-340

Baldrian P, Lopez-Mondejar R. 2014. Microbial genomics, transcriptomics and proteomics: new discoveries in decomposition research using complementary methods. *Applied Microbiology and Biotechnology* 98:1531-1537. DOI: 10.1007/s00253-013-5457-x
Baldrian P, Merhautová V, Cajthaml T, Petránková M, Šnajdr J. 2010. Small-scale distribution of extracellular enzymes, fungal, and bacterial biomass in *Quercus petraea* forest topsoil. *Biology and Fertility of Soils* 46:717-726. DOI: 10.1007/s00374-010-0478-4

Baldrian P, Kolarik M, Stursova M, Kopecky J, Valaskova V, Vetrovsky T, Zifcakova L, Snajdr J, Ridl J, Vlcek C, Voriskova J. 2012. Active and total microbial communities in forest soil are largely different and highly stratified during decomposition. *ISME Journal* 6:248-258. DOI: 10.1038/ismej.2011.95

Baldrian P, Lopez-Mondejar R. 2014. Microbial genomics, transcriptomics and proteomics: new discoveries in decomposition research using complementary methods. *Applied Microbiology and Biotechnology* 98:1531-1537. DOI: 10.1007/s00253-013-5457-x

Barros N, Salgado J, Feijóo S. 2007. Calorimetry and soil. *Thermochimica Acta* 458:11-17.

Berg B. 2014. Decomposition patterns for foliar litter – A theory for influencing factors. *Soil Biology and Biochemistry* 78:222-232. DOI: 10.1016/j.soilbio.2014.08.005.

Bonanomi G, Incerti G, Giannino F, Mingo A, Lanzotti V, Mazzoleni S. 2013. Litter quality assessed by solid state 13C NMR spectroscopy predicts decay rate better than C/N and Lignin/N ratios. *Soil Biology and Biochemistry* 56:40-48. DOI: 10.1016/j.soilbio.2012.03.003

Bremmer JM. 1996. Nitrogen-total. In: Spark DL, Page AL, Summer ME, Tabatabai MA, Helmke PA. eds. *Methods of Soil Analyses Part 3: Chemical Analyses*. Soil Science Society of America, Madison WI, pp. 1085–1121.

Brookes PC, Powlson DS, Jenkinson DS. 1982. Measurement of microbial biomass phosphorus in soil. *Soil Biology and Biochemistry* 14:319-329. DOI: 10.1016/0038-0717(82)90001-3
Brookes PC, Powlson DS, Jenkinson DS. 1984. Phosphorus in the soil microbial biomass. *Soil Biology and Biochemistry* 16:169-175. DOI: 10.1016/0038-0717(84)90108-1.

Chavez-Vergara B, Merino A, Vázquez-Marrufo G, García-Oliva F. 2014. Organic matter dynamics and microbial activity during decomposition of forest floor under two native neotropical oak species in a temperate deciduous forest in Mexico. *Geoderma* 235-236:133-145. DOI: 10.1016/j.geoderma.2014.07.005

Chávez-Vergara B, Rosales-Castillo A, Merino A, Vázquez-Marrufo G, Oyama K, García-Oliva F. 2016. Quercus species control nutrients dynamics by determining the composition and activity of the forest floor fungal community. *Soil Biology and Biochemistry* 98:186-195. DOI: 10.1016/j.soilbio.2016.04.015

Chávez-Vergara BM, González-Rodríguez A, Etchevers JD, Oyama K, García-Oliva F. 2015. Foliar nutrient resorption constrains soil nutrient transformations under two native oak species in a temperate deciduous forest in Mexico. *European Journal of Forest Research* 134:803-817. DOI: 10.1007/s10342-015-0891-1

Cleveland CC, Neff JC, Townsend AR, Hood E. 2004. Composition, Dynamics, and Fate of Leached Dissolved Organic Matter in Terrestrial Ecosystems: Results from a Decomposition Experiment. *Ecosystems* 7:275-285. DOI: 10.1007/s10021-003-0236-7

Cleveland CC, Reed SC, Keller AB, Nemergut DR, O'Neill SP, Ostertag R, Vitousek PM. 2014. Litter quality versus soil microbial community controls over decomposition: a quantitative analysis. *Oecologia* 174:283-294. DOI: 10.1007/s00442-013-2758-9

De Marco A, Spaccini R, Vittozzi P, Esposito F, Berg B, Virzo De Santo A. 2012. Decomposition of black locust and black pine leaf litter in two coeval forest stands on Mount
Vesuvius and dynamics of organic components assessed through proximate analysis and NMR spectroscopy. *Soil Biology and Biochemistry* 51:1-15. DOI: 10.1016/j.soilbio.2012.03.025

Dell’ Abate MT, Benedetti A, Trinchera A, Dazzi C. 2002. Humic substances along the profile of two Typic Haploxerert. *Geoderma* 107:281-296. DOI: 10.1016/S0016-7061(01)00153-7.

Fanin N, Fromin N, Bertrand I. 2016. Functional breadth and home-field advantage generate functional differences among soil microbial decomposers. *Ecology* 97:1023-1037. DOI: 10.1890/15-1263.1

Fanin N, Fromin N, Buatois B, Hattenschwiler S. 2013. An experimental test of the hypothesis of non-homeostatic consumer stoichiometry in a plant litter-microbe system. *Ecology Letters* 16:764-772. DOI: 10.1111/ele.12108

Fernández JM, Plaza C, Polo A, Plante AF. 2012. Use of thermal analysis techniques (TG-DSC) for the characterization of diverse organic municipal waste streams to predict biological stability prior to land application. *Waste Management* 32:158-164. DOI: 10.1016/j.wasman.2011.08.011

Freschet GT, Aerts R, Cornelissen JHC. 2012. Multiple mechanisms for trait effects on litter decomposition: moving beyond home-field advantage with a new hypothesis. *Journal of Ecology* 100:619-630. DOI: 10.1111/j.1365-2745.2011.01943.x

Freschet GT, Cornwell WK, Wardle DA, Elumeeva TG, Liu W, Jackson BG, Onipchenko VG, Soudzilovskaia NA, Tao J, Cornelissen JHC, Austin A. 2013. Linking litter decomposition of above- and below-ground organs to plant-soil feedbacks worldwide. *Journal of Ecology* 101:943-952. DOI: 10.1111/1365-2745.12092
Garcia-Palacios P, Shaw EA, Wall DH, Hattenschwiler S. 2016. Temporal dynamics of biotic and abiotic drivers of litter decomposition. *Ecology Letters* 19:554-563. DOI: 10.1111/ele.12590

Genung MA, Bailey JK, Schweitzer JA. 2013. The afterlife of interspecific indirect genetic effects: genotype interactions alter litter quality with consequences for decomposition and nutrient dynamics. *PLoS One* 8:e53718. DOI: 10.1371/journal.pone.0053718

Gessner MO, Swan CM, Dang CK, McKie BG, Bardgett RD, Wall DH, Hattenschwiler S. 2010. Diversity meets decomposition. *Trends in Ecology and Evolution* 25:372-380. DOI: 10.1016/j.tree.2010.01.010

Glanville H, Rousk J, Golyshin P, Jones DL. 2012. Mineralization of low molecular weight carbon substrates in soil solution under laboratory and field conditions. *Soil Biology and Biochemistry* 48:88-95. DOI: 10.1016/j.soilbio.2012.01.015

Hicks Pries CE, Bird JA, Castanha C, Hatton PJ, Torn MS. 2017. Long term decomposition: the influence of litter type and soil horizon on retention of plant carbon and nitrogen in soils. *Biogeochemistry* 134:5-16. DOI: 10.1007/s10533-017-0345-6

Huffman EWD. 1977. Performance of a new carbon dioxide coulometer. *Microchemical Journal* 22:567–573. DOI: 10.1016/0026-265X(77)90128-X

Joergensen RG, Mueller T. 1996. The fumigation-extraction method to estimate soil microbial biomass: Calibration of the KEN value. *Soil Biology and Biochemistry* 28:33-37. DOI: 10.1016/0038-0717(95)00101-8
Leifeld J, Kögel-Knabner I. 2005. Soil organic matter fractions as early indicators for carbon stock changes under different land-use. *Geoderma* 124:143–155. DOI: 10.1016/j.geoderma.2004.04.009.

Makkonen M, Berg MP, van Logtestijn RSP, van Hal JR, Aerts R. 2013. Do physical plant litter traits explain non-additivity in litter mixtures? A test of the improved microenvironmental conditions theory. *Oikos* 122:987-997. DOI: 10.1111/j.1600-0706.2012.20750.x

Murphy J, Riley JP. 1962. A modified single solution method for the determination of phosphate in natural waters. *Analytica Chimica Acta* 27:31e36. DOI: 10.1016/S0003-2670(00)88444-5

Nguyen Tu TT, Egasse C, Zeller B, Bardoux G, Biron P, Ponge JF, David B, Derenne S. 2011. Early degradation of plant alkanes in soils: A litterbag experiment using 13C-labelled leaves. *Soil Biology and Biochemistry* 43:2222-2228. DOI: doi:10.1016/j.soilbio.2011.07.009

Olsen JS. 1963. Energy storage and the balance of producers and decomposers in ecological systems. *Ecology* 44:322–331.

Ono K, Hiradate S, Morita S, Hirai K. 2011. Fate of organic carbon during decomposition of different litter types in Japan. *Biogeochemistry* 112:7-21. DOI: 10.1007/s10533-011-9682-z

Osono T, Azuma J-i, Hirose D. 2013. Plant species effect on the decomposition and chemical changes of leaf litter in grassland and pine and oak forest soils. *Plant and Soil* 376:411-421. DOI: 10.1007/s11104-013-1993-5

Osono T, Takeda H. 2002. Comparison of litter decomposing ability among diverse fungi in a cool temperate deciduous forest in Japan. *Mycologia* 94:421-427.
Pearse IS, Cobb RC, Karban R, Aerts R. 2014. The phenology-substrate-match hypothesis explains decomposition rates of evergreen and deciduous oak leaves. *Journal of Ecology* 102:28-35. DOI: 10.1111/1365-2745.12182

Prescott CE. 2010. Litter decomposition: what controls it and how can we alter it to sequester more carbon in forest soils? *Biogeochemistry* 101:133-149. DOI: 10.1007/s10533-010-9439-0

Rovira P, Kurz-Besson C, Coûteaux M-M, Ramón Vallejo V. 2008. Changes in litter properties during decomposition: A study by differential thermogravimetry and scanning calorimetry. *Soil Biology and Biochemistry* 40:172-185. DOI: 10.1016/j.soilbio.2007.07.021

Snajdr J, Cajthaml T, Valaskova V, Merhautova V, Petrankova M, Spetz P, Leppanen K, Baldrian P. 2011. Transformation of *Quercus petraea* litter: successive changes in litter chemistry are reflected in differential enzyme activity and changes in the microbial community composition. *FEMS Microbiology Ecology* 75:291-303. DOI: 10.1111/j.1574-6941.2010.00999.x

Spaccini R, Mbagwu JSC, Conte P, Piccolo A. 2006. Changes of humic substances characteristics from forested to cultivated soils in Ethiopia. *Geoderma* 132:9-19. DOI: 10.1016/j.geoderma.2005.04.015

Sparling GP, Feltham CW, Reynolds J, West AW. 1990. Estimation of soil microbial C by a fumigation–extraction method: use on soils of high organic matter content, and a reassessment of the KEC-factor. *Soil Biology and Biochemistry* 22, 301–307. DOI: 10.1016/0038-0717(90)90104-8.

StatSoft, Inc. 2014. STATISTICA (data analysis software system), version 12. www.statsoft.com.
Strickland MS, Osburn E, Lauber C, Fierer N, Bradford MA. 2009. Litter quality is in the eye of
the beholder: initial decomposition rates as a function of inoculum characteristics. *Functional
Ecology* 23:627-636. DOI: 10.1111/j.1365-2435.2008.01515.x

Talbot JM, Bruns TD, Taylor JW, Smith DP, Branco S, Glassman SI, Erlandson S, Vilgalys R,
Liao HL, Smith ME, Peay KG. 2014. Endemism and functional convergence across the North
American soil mycobiome. *Proceedings of the National Academy of Sciences* 111:6341-6346.
DOI: 10.1073/pnas.1402584111

Talbot JM, Yelle DJ, Nowick J, Treseder KK. 2012. Litter decay rates are determined by lignin
chemistry. *Biogeochemistry* 108:279-295. DOI: 10.1007/s10533-011-9599-6

Tiessen H, Moir JO. 1993. Characterization of available P by sequential extraction. In: Carter
MR, ed. Soil sampling and methods of analysis: Lewis Publishers, 75-85.

Van Meeteren MJM, Tietema A, Westerveld JW. 2007. Regulation of microbial carbon,
nitrogen, and phosphorus transformations by temperature and moisture during decomposition of
Calluna vulgaris litter. Biology and Fertility of Soils 44:103-112. DOI: 10.1007/s00374-007-
0184-z

Vance ED, Brookes PC, Jenkinson DS. 1987. An extraction method for measuring soil microbial
biomass C *Soil Biology and Biochemistry*. 19, 703–707. DOI: 10.1016/0038-0717(87)90052-6.

Wickings K, Grandy AS, Reed SC, Cleveland CC. 2012. The origin of litter chemical
complexity during decomposition. *Ecology Letters* 15:1180-1188. DOI: 10.1111/j.1461-
0248.2012.01837.x
Wurst S, Ohgushi T, Allen E. 2015. Do plant- and soil-mediated legacy effects impact future biotic interactions? *Functional Ecology* 29:1373-1382. DOI: 10.1111/1365-2435.12456

Zederer DP, Talkner U, Spohn M, Joergensen RG. 2017. Microbial biomass phosphorus and C/N/P stoichiometry in forest floor and A horizons as affected by tree species. *Soil Biology and Biochemistry* 111:166-175. DOI: 10.1016/j.soilbio.2017.04.009
Figure 1

Remaining mass in litterbags after 30 and 270 decomposition days

A) Remaining mass in litterbags after 30 and 270 days of decomposition in the field for each litter condition decomposed in the same site of production, B) Effect of litter quality over remaining mass at 30 days and 270 days, and C) Effect of site over remaining mass at 30 and 270 days. Different capital letters indicate differences among conditions at 30 days and lowercase letters indicate differences among conditions at 270 days.
Figure 2

$^{13}$C CPMAS NMR spectrograms

Litter decomposing on their site: A) *Quercus castanea* litterfall (solid line) and decomposed litter (dotted line) and B) *Quercus deserticola* litterfall (solid line) and decomposed litter (dotted line).
Figure 1. 

A. 

Quercus castanea 

- Litterfall 
- Decomposed at 270 days 

B. 

Quercus deserticola 

- Litterfall 
- Decomposed at 270 days 

Chemical shift (ppm)
Figure 3

DSC thermograms

Litter decomposing on their site: A) Quercus castanea and B) Quercus deserticola litterfall (solid line), decomposed at 30 days (broad dotted lines) and decomposed at 270 days (fine dotted lines).
A

Heat Flow (W g\(^{-1}\) sample)

Sample temperature (°C)

Manuscript to be reviewed

Quercus castanea

- Litterfall
- Decomposed at 30 days
- Decomposed at 270 days

B

Heat Flow (W g\(^{-1}\) sample)

Sample temperature (°C)

Quercus deserticola

- Litterfall
- Decomposed at 30 days
- Decomposed at 270 days
Figure 4

DSC thermograms of each litter condition decomposed

DSC thermograms of each litter condition decomposed on A) Quercus castanea site, B) mixed species site and C) Quercus deserticola site. Q. castanea litter (Qc, solid line), mixed species litter (Qx broad dotted lines) and Q. deserticola litter (Qd, fine dotted lines).
Figure 5

Microbial immobilization of C, N and P in litter decomposed

Microbial immobilization of C, N and P in litter decomposed on its original site (A-C), litter effect (D-F) and site effect (G-I). *Quercus castanea* (Qc), Mixed species (Qx) and *Quercus deserticola* (Qd). The suffixes -L and -S refer to litter and site, respectively. Different letters indicate statistical differences (p<0.05) according to the ANOVA model.
Figure 6

Specific enzymatic activities of dehydrogenase, β-glucosidase and polyphenol oxidase in litter decomposed on its original site (A-C), litter effect (D-F) and site effect (G-I). *Quercus castanea* (Qc), Mixed species (Qx) and *Quercus deserticola* (Qd). The suffixes -L and -S refer to litter and site, respectively. Different letters indicate statistical differences (p<0.05) according to the ANOVA model.
Table 1 (on next page)

Chemical characterization of litterfall and decomposed material in litterbags at 270 days by $^{13}$C CPMAS NMR.

HB: hydrophobic compounds; HI: hydrophilic compounds; S: syringyl; G: guaiacyl; H: p-hidroxypheny.
| Chemical shift (ppm) | Q. castanea | Q. deserticola |
|---------------------|-------------|----------------|
|                     | Litterfall  | 270 days       | Litterfall | 270 days |
| **Principal regions** |             |                |            |           |
| Alkyl C (0-45 ppm)  | 11          | 11             | 14         | 18        |
| O-Alky C (45-110 ppm) | 66         | 64             | 63         | 62        |
| Aryl C (110-160 ppm)| 18          | 17             | 17         | 15        |
| Carboxyl C (160-220 ppm) | 6          | 7              | 6          | 6         |
| **Ratios and indexes** |           |                |            |           |
| Alkyl C:O-Alkyl C   | 0.17        | 0.17           | 0.22       | 0.29      |
| O-Alky C:Aromatic C | 3.66        | 3.74           | eb3.71     | 4.13      |
| Hidrophobicity (HB:HI) | 0.40       | 0.39           | 0.45       | 0.48      |
| Aromaticity         | 0.23        | 0.22           | 0.22       | 0.18      |
| S:G                 | 1.35        | 1.75           | 1.00       | 1.50      |
| S:H                 | 2.30        | 1.75           | 1.50       | 1.50      |
| G:H                 | 1.18        | 1.00           | 1.45       | 1.00      |
Table 2 (on next page)

Thermal characteristics of litterfall and decomposed material in litterbags at 30 and 270 days analyzed by DSC-TG.

$Q'$: Energy released per gram of organic matter; $T_{50}$: Temperature in which the 50% of energy release occurs; Percent of energy released in: labile region ($Q_1$), recalcitrant region ($Q_2$) and extra-recalcitrant region ($Q_3$). Temperature in which the peak occurs in: the labile region ($T_1$), recalcitrant region ($T_2$) and extra-recalcitrant region ($T_3$).
|                   | Q. castanea |         |         | Q. deserticola |         |         |
|-------------------|------------|---------|---------|---------------|---------|---------|
|                   | Litterfall | 30 days | 270 days| Litterfall    | 30 days | 270 days|
| Q’ (J gMOS⁻¹)     | 11122      | 11103   | 11848   | 9085          | 9361    | 11060   |
| T₅₀ (°C)          | 396        | 379     | 375     | 400           | 377     | 375     |
| Q₁ (200-375 °C) (%)| 40.3       | 47.7    | 49.8    | 39.0          | 48.9    | 49.9    |
| Q₂ (375-475 °C) (%)| 57.0       | 49.7    | 46.1    | 55.9          | 45.5    | 43.6    |
| Q₃ (475-550 °C) (%)| 2.7        | 2.5     | 4.2     | 5.2           | 5.6     | 6.5     |
| T₁ (°C)           | 347        | 344     | 343     | 352           | 342     | 350     |
| T₂ (°C)           | 424/449    | 410     | 425     | 461           | 414     | 432     |
| T₃ (°C)           | -          | -       | -       | -             | 490     | 502     |
Table 3 (on next page)

Means of thermal parameters obtained through DSC at 30 days of decomposition.

Different capital letters indicate significant differences (P < 0.05) according to a one-way ANOVA model. Bold letters are statistically significant. Qc: *Quercus castanea* litter; Qx: Mixed species litter; Qd: *Quercus deserticola* litter. The suffixes -L and -S refer to litter and site, respectively. Q*: Energy released per gram of organic matter; T_{50}: Temperature in which the 50% of energy release occurs; Percent of energy released in: the labile region (Q₁), recalcitrant region (Q₂) and extra-recalcitrant region (Q₃). F: value of the test statistic; p: significance level.
| Litter quality | Site |
|----------------|------|
|                | QcL  | QxL  | QdL  | F(p) |
| Q' (W g⁻¹)     | 10815| 10804| 11083| 0.53(0.60) |
| T₅₀ (°C)       | 376  | 375  | 377  | 0.60(0.58) |
| Q₁ (%)         | 49.2 | 49.7 | 48.8 | 0.80(0.49) |
| Q₂ (%)         | 48.6ᴬ | 46.8ᴬ | 45.6ᴬ | 17.8(<0.01) |
| Q₃ (%)         | 2.2ᴮ | 3.4ᴮ | 5.6ᴬ | 20.4(<0.01) |

|                | QcS  | QxS  | QdS  | F(p) |
|----------------|------|------|------|------|
| Q' (W g⁻¹)     | 10716| 11154| 10832| 1.35(0.32) |
| T₅₀ (°C)       | 378  | 375  | 375  | 3.99(0.07) |
| Q₁ (%)         | 48.4 | 49.7 | 49.6 | 3.36(0.10) |
| Q₂ (%)         | 47.2 | 46.8 | 46.9 | 0.04(0.95) |
| Q₃ (%)         | 4.3  | 3.5  | 3.4  | 0.25(0.78) |
Table 4

Means of the concentration of total and dissolved C, N and P in litterfall and decomposed litter at 30 and 270 days for *Q.castanea*, mixture of species and *Q.deserticola* at their site of origin.

Different uppercase letters indicate significant differences (p < 0.05) between litter types within the same date. Different lowercase letters indicate significant differences (p < 0.05) between dates within the same litter type according to a repeated measures ANOVA (RMANOVA).
|                | Q. castanea | Mixed species | Q. deserticola |
|----------------|-------------|---------------|----------------|
| Litterfall     |             |               |                |
| 30 days        |             |               |                |
| 270 days       |             |               |                |
| C              | 489 \text{a} | 459 \text{ab} | 404 \text{b}  |
| N              | 7.3 \text{Ba} | 7.5 \text{Ba} | 7.5 \text{Ca} |
| P              | 0.21 \text{Aa} | 0.31 \text{Ba} | 0.23 \text{Ba} |
| C:N            | 68 \text{Aa}  | 62 \text{Aab} | 55 \text{Ab}  |
| C:P            | 2316 \text{Aa} | 1527 \text{Ab} | 1806 \text{Ab} |
| N:P            | 34 \text{a}   | 25 \text{c}   | 34 \text{b}   |

Dissolved organic nutrients (\(\mu g \text{ g}^{-1}\))

|                | Q. castanea | Mixed species | Q. deserticola |
|----------------|-------------|---------------|----------------|
| DOC            | 2933 \text{Ca} | 1194 \text{Cc} | 2051 \text{Cb} |
| DON            | 76 \text{Cb}   | 153 \text{Aa}  | 121 \text{Ba}  |
| DOP            | 26 \text{Bb}   | 32 \text{Ba}   | 10 \text{Bc}   |

Dissolved inorganic nutrients (\(\mu g \text{ g}^{-1}\))

|                | Q. castanea | Mixed species | Q. deserticola |
|----------------|-------------|---------------|----------------|
| \text{NH}_4^+  | 13 \text{Cb} | 18 \text{c}   | 53 \text{Aa}  |
| \text{NO}_3^-  | 2.9 \text{Ba}  | 3.0 \text{Aa}  | 4.6 \text{Ca}  |
| \text{HPO}_4^- | 22 \text{Ca}  | 8 \text{b}    | 21 \text{Ca}  |
Table 5 (on next page)

Multiple regression models at 30 days of decomposition between litter remnant mass and litter chemical quality and microbial metabolism variables.

Bold letters refer to significant variables according to a multiple regression model.
| Factors               | Included variables | Significant variables (β) | Multiple R² (p) |
|----------------------|--------------------|---------------------------|-----------------|
| Chemical quality     | C:N ratio          | 0.29                      |                 |
|                      | C:P ratio          | NS                        |                 |
|                      | DOC                | NS                        |                 |
|                      | DON                | -0.56                     | 0.78 (<0.001)   |
|                      | DOP                | -0.23                     |                 |
|                      | NH₄⁺               | -0.24                     |                 |
|                      | NO₃⁻               | NS                        |                 |
|                      | PO₄⁻               | NS                        |                 |
| Microbial metabolism | SEA BG             | NS                        |                 |
|                      | SEA CBH            | NS                        |                 |
|                      | SEA POX            | -0.45                     | 0.38 (0.033)    |
|                      | SEA NAG            | NS                        |                 |
|                      | SEA PHO            | NS                        |                 |
|                      | SEA DHG            | 0.41                      |                 |