https://doi.org/10.15517/rev.biol.trop..v71i1.50333

Functional traits and performance of woody species on oil-affected soils of the Ecuadorian Amazon

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Received 04-III-2023. Corrected 10-IV-2023. Accepted 10-X-2023.

ABSTRACT

Introduction: Plant functional traits are widely used to predict community productivity. However, they are rarely used to predict the performance (in terms of growth diameter, growth height, survival, and integral response index) of woody species planted in degraded soils.

Objective: To evaluate the relationship between the functional traits and the performance of 25 woody species planted in disturbed soils affected by oil extraction activities in Ecuadorian Amazon.

Methods: Eighteen permanent sampling plots were established and five 6-month-old seedlings of each 25 species were randomly planted in each plot (125 individuals per plot), at a distance of 4×4 m. Eight quantitative functional traits (leaf size, specific leaf area, leaf nitrogen concentration, leaf phosphorus concentration, leaf minimum unit, leaf dry matter content, stem specific density and leaf tensile strength) were determined for each species.

Results: The woody species with high performance shows greater leaf size, specific leaf area and Stem Specific Density than those showing low performance. Leaf nitrogen concentration and stem specific density had a direct relationship with the integral response index. The leaf size, leaf phosphorus concentration, leaf dry matter content and leaf tensile strength showed a negative relationship with the integral response index.

Conclusions: Our study demonstrated that the performance of woody species o disturbed soils can be predicted satisfyingly by leaf and stem functional traits, presumably because these traits capture most of environmental and neighborhood conditions.

Key words: growing species; integral response index; disturbed soils; quantitative traits; multiple regression model.

RESUMEN

Rasgos funcionales y comportamiento de especies leñosas en suelos afectados por petróleo de la Amazonía ecuatoriana

Introducción: Los rasgos funcionales de las plantas han sido ampliamente utilizados para predecir la productividad (en términos de crecimiento en diámetro, crecimiento en altura, sobrevivencia e índice de respuesta integral) de las comunidades vegetales. Sin embargo, rara vez han sido utilizados para predecir el desempeño de las especies leñosas plantadas en suelos degradados.

Objetivo: Evaluar la relación entre el desempeño y los rasgos funcionales de 25 especies leñosas plantadas en suelos afectados por actividades de extracción de petróleo en la Amazonía ecuatoriana.

Métodos: Se establecieron 18 parcelas permanentes de muestreo y en cada parcela se sembraron aleatoriamente cinco plántulas de 6 meses de las 25 especies (125 individuos por parcela), a una distancia de 4×4 m. Se determinaron ocho rasgos funcionales (área foliar, área foliar específica, concentración de nitrógeno foliar, concentración de fósforo foliar, unidad mínima foliar, contenido foliar de materia seca, densidad específica del fuste y fuerza tensil foliar) de cada especie.

Resultados: Las especies leñosas con alto desempeño presentaron mayor área foliar, área foliar específica y densidad específica del fuste que las especies de bajo desempeño. La concentración de nitrógeno foliar y la densidad específica del fuste mostraron una relación directa. El área foliar, la concentración de fósforo foliar, el contenido de materia seca foliar y la fuerza tensil foliar presentaron una relación inversa con el Índice de Respuesta Integral. **Conclusión:** Se demostró que el desempeño de las especies leñosas plantadas en suelos alterados puede predecirse satisfactoriamente por rasgos funcionales de hoja y de tallo, debido posiblemente a que los rasgos influyen en el crecimiento y supervivencia de las especies, y reflejan la mayoría de las condiciones ambientales.

Palabras clave: crecimiento de especies; índice de respuesta integral; suelos perturbados; rasgos cuantitativos; modelo de regresión múltiple.

INTRODUCTION

Tropical forests cover 10 % of the terrestrial surface and account for 50 % of worldwide tree diversity (Mayaux et al., 2005). More than half of tropical forest areas are composed of tropical rainforests, which are characterized by a high diversity of tree species (Lewis et al., 2009). Tropical rainforests have a significant influence on global patterns of biodiversity, ecosystem ecology, productivity, and biogeochemical cycles (Malhi, 2010). According to Guevara-Andino et al. (2019) in the Ecuadorian Amazon rainforest there are 2296 tree species; however, approximately 4.2 million hectares of this ecosystem have been impacted by many anthropogenic activities related to oil extraction (opening of new roads, construction of platforms, mud and drill cutting cells, contaminated soil treatment units, and settlement of camps and heliports) (Rivera-Parra et al., 2020). These activities have resulted in high deforestation, acceleration of soil erosion, decrease in water infiltration, increase in superficial runoff (Bertzky et al., 2011), and

reduction of fauna species (Arroyo-Rodríguez et al., 2007; Pozo-Rivera et al., 2023).

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In response, the Ecuadorian government has implemented reforestation programs on sites affected by oil extraction processes since 2000 (Villacís, 2016). Reforestation activities began with the production of seedlings in plant nurseries, where various native and exotic tree species are produced (primarily timber, fruit, and ornamental plants), later, seedlings are transplanted into the affected soils when reach 30 cm in height. Studies to evaluate the most suitable species for reforestation sites (previously affected by oil extraction processes) have been conducted in plant nurseries (Villacís, Armas et al., 2016), as well as in open-field affected sites (Villacís, Casanoves et al., 2016).

Nevertheless, it is unknown whether the growth and survival rates of the sapling woody species on disturbed sites over a period of two years is a good indicator of their performance. In this sense, the long-term performance of woody species can be determined based on their functional traits (Poorter & Bongers, 2006), particularly leaf and stem traits (Martínez-Encino et al., 2013). The functional traits are fundamental to the understanding of plant adaptations and can be useful for selecting suitable species to restore degraded ecosystems (Wang et al., 2021). Additionally, functional traits encompass the morpho-phenological characteristics of plants that have indirect effects on their physical structure, affecting key biological processes such as growth, reproduction, and survival (Yadav et al., 2022). For example, Freschet et al. (2010) and Pérez-Harguindeguy et al. (2016) indicated that trees with high foliar concentrations of nitrogen and phosphorus, as well as large specific leaf areas, demonstrated rapid growth rates. Moreover, Sterck et al. (2006) found that trees with low stem specific density exhibited faster growth rates, whereas trees with high stem specific density had a higher survival rate.

Because the long-term growth assessment of woody species could be expensive, functional traits could be good indicators of forest species performance. Maire et al. (2013) found that functional traits were related to the growth of individual plants. In other studies, the high specific leaf area and low stem specific density of the trees have been found to be associated with their fast growth; furthermore, the low specific leaf area and high stem specific density of the trees were associated with high survival (Poorter et al., 2008; Rüger et al., 2012). Leaf functional traits have been used to explain species-specific growth rates and have contributed to a better understanding of tree growth strategies and the structure and dynamics of forest communities (Adler et al., 2014). Although the relationships between functional traits and species growth in tropical forest ecosystems have been often studied (Werden el at., 2018), no studies have been conducted in sites disturbed by oil extraction processes. The information about the performance of woody species in restored Amazonian tropical ecosystems and their relationships with functional traits will constitute a valuable contribution to scientific knowledge. In this study, the performance of twenty-five woody species planted on soils affected by oil extraction activities years was

evaluated. Using data for a 24 month growing period for twenty-five woody species, we correlated species performance (growth diameter, growth height, survival and integral response index) with seven functional traits related to leaf (Leaf Size, Leaf Minimum Unit, Specific Leaf Area, Leaf Dry Matter Content, Leaf Tensile Strength, Leaf Nitrogen Concentration and Leaf Phosphorus Concentration), and one stem functional traits (Stem Specific Density). The following two hypotheses were tested: (i) some plant functional traits differed between woody species according to their performance on soils affected for oil extraction activities, and (ii) the performance of all woody species can be predicted simultaneity by leaf and stem functional traits.

MATERIALS AND METHODS

Study area: The study was performed in the Sucumbíos (0°5' S & 76°53' W) province of Ecuador. The region has an average altitude of 328 m a.s.l., a mean annual rainfall of 3 000 mm, a mean annual temperature of 25 °C, a relative humidity of 85 % and a heliophany of 12 h day⁻¹. The area is classified as tropical rainforest (Peel et al., 2007). The soils of the Ecuadorian Amazon are acidic, have low levels of nutrients and high aluminum contents (Villacís, Armas et al., 2016).

Selection of species and disturbed sites: The 25 woody species selected were mostly produced in the Ecuadorian Amazon (Table 1). These species have been used in remediation programs of disturbed soils, mainly for their timber, forage and fruit uses (Villacís, Armas, et al., 2016; Villacís, Casanoves et al., 2016).

In sites affected by the oil extraction processes, the superficial layer of soil and all vegetation have been removed, and they contain residues of hydrocarbons and heavy metals. These sites present an average pH of 4.6, 0.5 % of organic matter, 5.3 mg kg⁻¹ of Al, 5 341.73 mg kg⁻¹ of total petroleum hydrocarbons, 5.3 mg kg⁻¹ of Cd, 33.2 mg kg⁻¹ of Ni and 23.76 mg kg⁻¹ of Pb. More details on the physical and chemical

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Taxonomic classification and uses of restoration woody species.

Species	Family	Use	Origin	Nitrogen fixer
Acnistus arborescens (L.) Schltdl.	Solanaceae	Timber/Medicinal	Native	No
Apeiba membranacea Spruce ex. Benth.	Malvaceae	Ornamental/Medicinal	Native	No
Averrhoa carambola L.	Oxalidaceae-	Timber/Fruit	Exotic	No
Cedrela odorata L.	Meliaceae	Timber	Native	No
Cedrelinga cateniformis (Ducke) Ducke.	Fabaceae	Timber	Native	Yes
Gmelina arborea Roxb. ex Sm.	Lamiaceae	Timber	Native	No
Guarea purusana C. DC.	Meliaceae	Timber	Native	No
Inga densiflora Benth.	Fabaceae	Timber/Fruit	Native	Yes
Leucaena leucocephala (Lam.) de Wit	Fabaceae	Forage	Exotic	Yes
Morinda citrifolia L.	Rubiaceae	Fruit	Exotic	No
Myrcia aff. fallax	Myrtaceae	Timber	Native	No
Myroxylon balsamum (L.) Harms	Fabaceae	Timber/Medicinal	Native	Yes
Nephelium lappaceum L.	Sapindaceae	Fruit	Native	No
Ochroma pyramidale (Cav. ex Lam.) Urb.	Malvaceae	Timber	Native	No
Ormosia macrocalyx Ducke	Fabaceae	Timber	Native	Yes
Piptadenia pteroclada Benth.	Fabaceae	Timber	Native	Yes
Platymiscium pinnatum (Jack.) Dougand	Fabaceae	Timber	Native	Yes
Pourouma cecropiifolia Mart.	Urticaceae	Fruit/Medicinal	Native	No
Schizolobium parahyba (Vell.) S.F. Blake	Fabaceae	Timber	Native	Yes
Stryphnodendron porcatum D.A.Neill & Occhioni f.	Fabaceae	Timber	Native	Yes
Syzygium jambos (L.) Alston	Myrtaceae	Fruit	Native	No
Syzygium malaccense (L.) Merr. & L.M.Perry	Myrtaceae	Fruit	Exotic	No
Tapirira guianensis Aubl.	Anacardiaceae	Timber	Native	No
Vitex cymosa Bertero ex Spreng.	Lamiaceae	Timber	Native	No
Zygia longifolia (Humb. & Bond. ex Willd.) Britton & Rose	Fabaceae	Timber	Native	Yes

characteristics of these sites can be found in Villacís, Casanoves et al. (2016). Eighteen permanent sampling plots were established and five 6-month-old seedlings of each 25 species were randomly planted in each plot (125 individuals per plot), at a distance of 4×4 m. The saplings that died due to post-transplant shock in the first month were replanted with new individuals of the same species. Every 4 months weed controls were carried out with a motor trimmer, eliminating all the vegetation within a 1 m radius from the stem of the seedlings.

Measured variables: In each plot sapling survival was measured by dividing the number of two-year-old living plants by the initial number of plants transplanted. The diameter and height at the time of transplantation and two years after transplantation were determined. The diameter of saplings was measured at 10 cm from the base of the plant, using a digital caliper and plant height was measured from the base to the apical meristem of the tallest stem, using a laser hypsometer. Subsequently, we estimated the integrated response index (IRI) which considers survival rates and growth variables and serves to determine the performance of saplings in an integrated way. The IRI was estimated according to the following equation:

IRI= survival percentage × RGR height × RGR diameter. Where: RGR height (height rate growth in cm / month) = [ln (final height) -ln (initial height)] / 24 months; RGR diameter (diameter rate growth in mm / month) = [ln (final diameter) -ln (initial diameter)] / 24 months (Elliott et al., 2003).

For the evaluation of functional traits, five adult individuals from the natural forest located along a longitudinal gradient in Sucumbios Province were selected (17 947 km²; Fig. 1).

Leaf and stem plant functional traits that are commonly used to characterize growth strategy of plants were evaluated. In each specimen, seven leaf functional traits [(leaf size (LS), leaf minimum unit (LMU), specific leaf size (SLA), leaf dry matter content (LDMC), leaf nitrogen concentration (LNC), leaf phosphorus concentration (LPC), leaf tensile strength (LTS], and one stem functional traits [(stem specific density (SSD)] were measured (Chave, 2006; Cornelissen et al., 2003).

The LS, LMU and SLA are indicators of the adaptation to the environment and to the strategies and the use of plant resources (Vendramini et al., 2002). The LDMC is related to the average density of leaf tissues (Cornelissen et al., 2003) and reflects an exchange between plant performance and rapid biomass production. To estimate the LS, SLA, and LDMC, five leaves of each individual were collected. Each leave using the Midebmp Image Analysis version 4.2 program was scanned and the LS in mm² was measured (Ordiales-Plaza, 2000). The SLA in mm² mg⁻¹ by dividing the leaf size by the leaf dry weight was estimated (Hunt, 1990). The LDMC in mg g⁻¹ by dividing the leaf dry weight by leaf fresh weight was calculated.

The LNC and the NPC foliar are vital elements of the photosynthesis functioning of the leaves (Bongers & Popma, 1990). A sample of fresh leaves without petioles or rachis of each sampled specimen to measure the LNC (Micro Kjheldal total combustion method) and the LPC (colorimeter method) was estimated (Cornelissen et al., 2003). The LTS is an indicator of the inverted carbon ratio in the structural protection of the photosynthetic tissues (Cornelissen et al., 2003). A rectangular section 1 cm wide by 4 cm long in the direction of the vein of



Fig. 1. Geographical location of 18 plots, and 125 sampled trees of 25 woody species in the Ecuadorian Amazon.



a fully sound expanded leaf was extracted; the section was subjected to traction in a Tearing Apparatus and the LTS in N mm⁻¹ (maximum force resisted by leaf section at the instant of its rupture) was obtained (Hendry & Grime, 1993). Finally, the SSD is directly related to the carbon accumulation and flows and inversely related to the stem growth rate, mortality rate and plant reproduction time (Swenson & Enquist, 2007). A cylindrical wood samples from each individual with a 6.2 mm auger drill was collected; the volume and dry weight of cylinder was measured, and the SSD in g cm⁻³ by dividing dry weight by cylinder volume was calculated. These data are available in the TRY database (Kattge et al., 2020).

Data analysis: In order to group woody species according to their performance a cluster analysis (Ward method and Euclidean distance) for integral response index (IRI) was performed to obtain two groups (high and low performance). Differences in IRI and quantitative traits between species of high and low performance were analyzed, by using general linear models; subsequently a less significant difference (LSD) post hoc test was performed (α = 0.05). Correlation analyses between functional traits and growth height, growth diameter and survival of species using Pearson correlations analysis were performed. To fit the performance of woody species in function of plant functional traits, a multiple regression models were built. Using leaf size, specific leaf area, leaf nitrogen concentration, leaf phosphorus concentration, leaf minimum unit, leaf dry matter content, stem specific density and leaf tensile strength, as predictor variables, we predicted the IRI. The multiple regression model was based on the assumption that residuals are normally distributed and their variance are homogeneous (Wright & Cannon, 2001). Normality and homogeneity of variances assumptions were evaluated with graphical analysis of residuals. We used partial residuals graphs to evaluate when was necessary or not the inclusion of polynomial terms for each trait. The selection and evaluation of the best regression model was based on the coefficient of determination (adjusted R²) and mean squared prediction error (MSPE), Akaike information criterion (AIC), Bayesian information criterion (BIC) and p-value. We use Variance Inflation Factor (VIF) values to evaluate multicollinearity. In order to compare the effect of measured traits on the IRI we use the relative variation explained for each trait. All analyzes were performed using InfoStat statistical software (Di Rienzo et al., 2020).

RESULTS

Performance of woody species: Based on the integral response index, cluster analysis exhibited two well-differentiated groups of plants (Fig. 2). The first group formed by Cedrelinga cateniformis, Myrcia aff. fallax, Piptadenia pteroclada, Platymiscium pinnatum, Tapirira guianensis, Vitex cymosa and Zygia longifolia species, showed high IRI mean value (high performance woody species). The second group formed for Acnistus arborescens, Apeiba membranaceae, Averrhoa carambola, Cedrela odorata, Gmelina arborea, Guarea purusana, Inga densiflora, Leucaena leucocephala, Myroxylon balsamum, Morinda citrifolia, Nephelium lappaceum, Ochroma pyramidale Ormosia macrocalyx Pourouma cecropiifolia, Schizolobium parahyba, Stryphnodendron porcatum, Syzygium jambos and Syzygium malaccensis species, displayed a low IRI mean value (low performance woody species). The growth diameter, growth height and integral response index was higher in high performance woody species than in low performance woody species. Nevertheless, the survival was similar for the high and low performance sapling woody species (Table 2).

Mean values for leaf size, specific leaf area and Stem Specific Density were higher in saplings of high performance woody species. The rest of plant functional traits did not differ between saplings woody species of high and low performance (Table 2).

Correlations between variables: Correlation analysis showed a negative relationship

Revista de Biología Tropical, ISSN: 2215-2075, Vol. 71: e50333, enero-diciembre 2023 (Publicado Oct. 30, 2023)



Fig. 2. Dendrogram of classification of 25 sapling woody species, obtained by hierarchical cluster analysis (Ward method and Euclidean distance) using integral response index (IRI).

Table 2

Performance variables and functional traits for high and low performance woody species on soils affected for oil extraction activities.

	Species per			
Variables	High	Low	F-Fisher	P-value
Performance				
Survival (%)	85.38 ± 4.52 a	85.18 ± 2.55 a	0.00	0.9687
Growth diameter (mm)	45.79 ± 3.08 a	19.50 ± 1.92 b	52.29	0.0001
Growth height (cm)	1.45 ± 0.12 a	$1.01\pm0.08~\mathrm{b}$	9.71	0.0049
Integral response index (IRI)	0.37 ± 0.08 a	$0.22 \pm 0.03 \text{ b}$	6,19	0.0205
Functional traits				
Leaf size (mm ²)	62 702.99 ± 14 146.56 a	26 696.30 ± 8 821.93 b	4.66	0.0415
Specific Leaf Area (mm ² mg ⁻¹)	59 564.60 ± 13 809.31 a	17 950.28 ± 8 611.62 b	6.54	0.0176
Leaf Nitrogen Concentration (%)	2.64 ± 0.41 a	2.79 ± 0.25 a	0.10	0.7593
Leaf Phosphorus Concentration (%)	0.20 ± 0.03 a	0.19 ± 0.02 a	0.08	0.7788
Leaf Minimum Unit (mm ²)	29 396.69 ± 12 782.49 a	13 104.35 ± 7 971.29 a	1.17	0.2907
Leaf Dry Matter Content (mg g ⁻¹)	0.38 ± 0.04 a	0.37 ± 0.02 a	0.17	0.6835
Stem Specific Density (g cm ⁻³)	0.56 ± 0.04 a	$0.45 \pm 0.03 \text{ b}$	4.52	0.0444
Leaf Tensile Strength (N mm ⁻¹)	4.63 ± 0.49 a	3.88 ± 0.30 a	1.72	0.2027

Values are means \pm error standard. Different letters in each row indicate significant differences (LSD, post hoc test, P < 0.05).

between Survival and Leaf Tensile Strength. The rest of correlation were non-significant (Table 3).

The best multiple regression model shows the higher adjusted R² and the lowest values of AIC and BIC. For LS, SLA, LDMC and LNC a second order polynomial was used (Table 4).

The estimated regression model predicted 90 % of the total IRI variability, and it is represented by the following equation:
$$\begin{split} IRI &= 0.066 - 1.19 \times LPC + 4.59 \ E^{-6} \times LMU + \\ 0.98 * SSD - 0.05 * LTS 5.53 \ E^{-6} \times LS + 2.68 \ E^{-11} \\ \times \ LS^2 + 1.31 \\ E^{-5} \times SLA - 6.22 \ E^{-11} \times SLA^2 - 3.12 \\ \times \ LDMC + 5.21 \times LDMC^2 + 0.23 \times LNC - 0.032 \\ \times \ LNC^2 \end{split}$$

The IRI showed a positive relationship with SLA, LMU, LNC and SSD and a negative relationship with LS, LPC, LDMC and LTS.

Table 3

Pearson correlation coefficients (r) between functional traits and average absolute growth in height (height growth), diameter (diameter growth) and survival after 24 months (survival, %) for 25 woody species planted on degraded soils in Amazonian Basin.

Functional traits	Growth diameter (mm year ⁻¹)		Growth height (cm year ⁻¹)		Survival (%)	
	r	P-value	r	P-value	R	P-value
Leaf traits						
Leaf Size (mm ²)	-0.05	0.8208	-0.22	0.2864	0.07	0.7289
Specific Leaf Area (mm ² mg ⁻¹)	-0.08	0.6907	-0.15	0.4790	-0.16	0.4378
Leaf Nitrogen Concentration (%)	0.30	0.1479	0.36	0.0798	-0.01	0.9610
Leaf Phosphorus Concentration FPC (%)	-0.01	0.9667	-0.09	0.6668	-0.09	0.6622
Leaf Minimum Unit (mm ²)	-0.10	0.6223	-0.28	0.1704	0.02	0.9109
Leaf Dry Matter Content (mg g ⁻¹)	0.18	0.3768	0.11	0.6081	0.18	0.4019
Leaf Tensile Strength (N mm ⁻¹)	-0.31	0.1305	-0.32	0.1131	-0.45	0.0236
Stem trait						
Stem Specific Density (g cm ⁻³)	-0.05	0.7973	0.07	0.7441	0.11	0.5915

Table 4

Regression coefficients and summary statistics for the best model to predict IRI from functional traits.

Coefficient	Estimation	Standard error	t-Student	P-value	CpMallows
Constant	0.0660	0.3077	0.2146	0.8337	
LPC %	-1.1901	0.5454	-2.1822	0.0497	15.7619
LMU (mm ²)	4.6E-06	1.6E-06	2.8510	0.0146	19.1281
SSD (g cm ⁻³)	0.9830	0.1751	5.6122	0.0001	42.4972
LTS (N mm ⁻¹)	-0.0479	0.0197	-2.4283	0.0318	16.8967
LS (mm ²)	-5.5E-06	2.1E-06	-2.6226	0.0223	17.8781
LS2 (mm ²)	2.7E-11	1.4E-11	1.9828	0.0708	14.9317
SLA (mm ² mg ⁻¹)	1.3E-05	2.6E-06	4.8415	0.0004	34.4398
SLA2 (mm ² mg ⁻¹)	-6.2E-11	1.8E-11	-3.3817	0.0055	22.4361
LDMC (mg g ⁻¹)	-3.1276	1.2474	-2.5073	0.0275	17.2863
LDMC ² (mg g ⁻¹)	5.2198	1.6910	3.0868	0.0094	20.5285
LNC %	0.2376	0.1011	2.3494	0.0368	16.5198
LNC ² %	-0.0323	0.0155	-2.0883	0.0588	15.3608

SLA = Specific Leaf Area; LDMC = Leaf Dry Matter Content; LTS = Leaf Tensile Strength; LS = Leaf Size; LMU = Leaf Minimum Unit; SSD = Stem Specific Density; LNC = Leaf Nitrogen Concentration; LPC = Leaf Phosphorus Concentration.

DISCUSSION

According to our knowledge, this is the first report that approach the functional traits and the performance of woody species planted on soils affected for oil extraction activities. The woody species of high performance showed greater leaf size, specific leaf area and Stem Specific Density than low performance woody species. The performance of woody species using IRI was explained using leaf and stem functional traits as regressor variables.

Differences in functional traits between woody species according to their performance: The High performance woody species (Cedrelinga cateniformis, Myrcia aff. fallax, Piptadenia pteroclada, Platymiscium pinnatum, Tapirira guianensis, Vitex cymosa and Zygia longifolia) showed a high Leaf Size, that enables them to maximize light capture (Poorter at al., 2006). Moreover, the nutrient cycling processes of these species are more efficient due to their high photosynthetic capacity per unit of leaf area (Denslow, 1996; Le Roux et al., 2001; Mooney et al., 1981; Reich et al., 1998). This character trait is favorable at the beginning of restoration processes because the disturbed sites are totally exposed to the sun and do not have any type of vegetation. Similar results were reported by Poorter and Evans (1998), who found that fast-growing tree species presented high leaf Area. Additionally, high performance woody species tend to have a high Leaf Dry Matter Content. This allows them capture and use available resources in an efficient way (Tecco et al., 2013), due to the greater stiffness of its leaves (Niinemets, 2001) because of the lignification of the thick and rigid cell walls (Niinemets & Kull, 1998). Similar results were reported by Martínez-Garza et al., (2013), who found that tree plants with higher Leaf Dry Matter Content had higher survival. The high Specific Stem Density of high performance woody species reflects an elevated growth rate (Santiago et al., 2004), which is essential to compete with aggressive colonizing vegetation. This allows the saplings

increase their performance under the adverse conditions of disturbed soils (Villacís, Casanoves et al., 2016). However, other studies have reported that trees with higher Specific Stem Density have presented higher survival, but lower diameter growth (Fortunel et al., 2016; Poorter et al., 2008). Finally the low Leaf Tensile Strength of high performance woody species favors the rapid growth and decomposition of its biomass (Obando-Vargas, 2003).

On the other hand, the low performance woody species showed a low Leaf Size and Specific Leaf Area, that indicate that the species of this group must invest more amounts of energy into the structural protection of their leaves (Wright et al., 2004). In addition, species with low Leaf Size and Specific Leaf Size have low assimilation and respiration rates, and high survival (Lusk, 2002). Aditionally, the low Leaf Dry Matter Content reflects a slow process of nutrient conservation in low performance woody species (Messier et al., 2010). Pérez-Harguindeguy et al. (2016), reported that the species with low growth rates presented low levels of Leaf Dry Matter Content. The low performance woody species showed less Specific Stem Density than other species. Similar results were found in several studies concerning rainforest saplings (Muller-Landau, 2004) and trees (King et al., 2005). Finally, due to their high Leaf Tensile Strength, low performance woody species have a greater proportion of vascular tissues, fibers or sclerenchyma, and high tissue density (Tecco et al., 2010; Wright & Cannon, 2001), which was reflected in lower growth rates and lower performance compared to high performance woody species.

Relationship between functional traits and performance of woody species: Considering that stem specific density is directly related with the content, accumulation, and fluctuations of carbon (Wiemann & Williamson, 2002) and inversely related with stem growth rate, mortality rate, and breeding time (Swenson & Enquist, 2007), woody species with lower stem specific density will have greater possibilities of growing under the adverse conditions of degraded sites, due to their capacity to produce biomass quickly (Díaz et al., 2004) and survive the adverse conditions of the disturbed soils.

In our study, the leaf and stem functional traits were good predictors of the performance of the woody species evaluated with IRI. Similar studies carried out with plants grown under controlled conditions and in the open field, have also reported mathematical models that have explained the performance of tree species based on their functional traits (Grime & Hunt, 1975; Rüger et al., 2012). However, Paine et al. (2015) point out that functional traits are not good predictors of tree plant growth at a global scale. These differences could possibly be explained because the woody species performance is not only affected by its functional traits, but also by the adaptation of the traits to its environment. In this sense, the conditions of the disturbed soils may have influenced the performance of the trees in addition to their functional traits, for which more in-depth studies are necessary to validate this hypothesis. The estimated regression model predicted 90 % of the total IRI variability, indicating that there are other traits or factors that should be considered to explain greater variability. Within the best multiple regression model estimated to predict the performance of the trees of the woody species planted on disturbed soils, eight quantitative functional traits (seven of the leaf and one of the stems) explained the performance of the woody species.

The functional traits SLA, LMU, LNC and SSD, showed a directly proportional relationship with the IRI of the woody species. Among these traits, SSD explained the highest relative percentage of the variance. Although the literature does not report studies on relationships between the IRI and functional traits (growth in diameter + growth in height + survival), studies carried out by Iida et al. (2014) and Gibert et al. (2016) reported that the stem specific density represents the best trait to predict the woody species growth and survival. On the other hand, LS, LPC, LDMC and LTS showed an inverse relationship with the IRI of the woody species. LDMC explained the highest relative percentage of the total variability. Negative relationships between the LDMC and the relative growth rates of the species have been reported in studies by Pérez-Harguindeguy et al. (2016). The negative rate of change between IRI and LPC differs from the results found by Chaturvedi et al., (2011) and Cernusak et al., (2010), who reported positive relationships between the growth rate and the LPC.

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The multiple regression model estimated by the combination of leaf and stem functional traits, explained better the woody species performance. This finding provides empirical evidence that the woody species performance is the integrated result of several leaf and stem functional traits (Iida et al. 2014). Therefore, in our research the woody species performance was explained better by several foliar and stem functional traits, than for a single functional trait (Li et al., 2017).

Implications for forest restauration: Our results are specific to predict the performance of young woody species planted in soils affected by oil extraction processes in Amazon Basin. However, our findings should not be correlated with the performance of adult woody species growing in other environments. Additionally, the performance of saplings not only depends on the local environment, but also of the functional traits of the species and their interaction (Uriarte et al., 2016). The environmental variation was not considered in our models and therefore contributes to the non-explained variability. On the other hand, the search of additional functional traits that provide predictions on the performance of woody species must also be specific for each phonological stage. In this sense, others functional traits such as physiological or allometric, which are well correlated with common performance must be identified. We must need to consider the possibility that the relationship between any functional trait and performance of woody species depends on the values of other unidentified functional traits (Marks & Lechowicz, 2006).

Ethical statement: the authors declare that they all agree with this publication and made significant contributions; that there is no conflict of interest of any kind; and that we followed all pertinent ethical and legal procedures and requirements. All financial sources are fully and clearly stated in the acknowledgments section. A signed document has been filed in the journal archives.

REFERENCES

- Adler, P. B., Salguero-Gómez, R., Compagnoni, A., Hsu, J. S., Ray-Mukherjee, J., Mbeau-Ache, C., & Franco, M. (2014). Functional traits explain variation in plant life history strategies. Proceedings of the National Academy of Sciences of the United States of America, 111(2), 740–745.
- Arroyo-Rodríguez, V., Mandujano, S., Benítez-Malvido, J., & Cuende-Fanton, C. (2007). The influence of large tree density on howler monkey (*Alouatta palliata mexicana*) presence in very small rain forest fragments. *Biotropica*, 39, 760–766.
- Bertzky, M., Ravilious, C., Araujo, A. L, Kapos, V., Carrión, D., Chíu, M., & Dickson, B. (2011). Carbono, biodiversidad y servicios ecosistémicos: Explorando los beneficios múltiples. Ecuador. UNEP-WCMC.
- Bongers, F., & Popma, J. (1990). Leaf Characteristics of the Tropical Rain Forest Flora of Los Tuxtlas, Mexico. *Botanical Gazette*, 151(3), 354–365.
- Cernusak, L. A., Winter, K., & Turner, B. L. (2010). Leaf nitrogen to phosphorus ratios of tropical trees: Experimental assessment of physiological and environmental controls. *New Phytologist*, 185, 770–779.
- Chaturvedi, R. K., Raghubanshi, A. S, & Singh, J. S. (2011). Leaf attributes and tree growth in a tropical dry forest. *Journal of Vegetation Science*, 22, 917–931.
- Chave, J. (2006). Measuring wood density for tropical forest trees a field manual. PAN-AMAZONIA. https:// rainfor.org/wp-content/uploads/sites/129/2022/06/ wood_density_english1.pdf
- Cornelissen, J. H. C., Lavorel, S., Garnier, E., Díaz, S., Buchmann, N., Gurvich, D. E., Reich, P. B., ter Steege, H., Morgan, H. D., van der Heijden, M. G. A., Pausas, J. G., & Poorter, H. (2003). A handbook of protocols for standardised and easy measurement of plant functional traits worldwide. *Australian Journal of Botany*, 51(4), 335–380.
- Denslow, J. S. (1996). Functional Group Diversity and Responses to Disturbance. In G. H. Orians, R. Dirzo, & J. H. Cushman (Eds.), *Biodiversity and Ecosystem*

Processes in Tropical Forests. Ecological Studies (p. 122). Springer.

- Di Rienzo, J. A., Casanoves, F., Balzarini, M. G., Gonzalez, L., Tablada, M., & Robledo, C. W. (2020). *InfoStat* (Versión 2020, Software). Centro de Transferencia Infostat, Argentina. https://www.infostat.com.ar/
- Díaz, S., Hodgson, J. G., Thompson, K., Cabido, M., Cornelissen, J. H., Jalili, A., Montserrat-Marti, G., Grime, J. P., Zarrinkamar, F., Asri, Y., Band, S. R., Basconcelo, S., Castro-Diez, P., Funes, G., Hamzehee, B., Khoshnevi, M., Pérez-Harguindeguy, N., Perez-Rontome, M. C., Shirvany F. A., ... Zak, M. R. (2004). The plant traits that drive ecosystems: evidence from three continents. *Journal of Vegetation Science*, 15(3), 295–304.
- Elliott, S., Navakitbumrung, P., Kuarak, C., Zangkum, S., Anusarnsunthorn, V., & Blakesley, D. (2003). Selecting framework tree species for restoring seasonally dry tropical forests in northern Thailand based on field performance. *Forest Ecology and Management*, 184, 177–191.
- Fortunel, C., Valencia, R., Wright, S. J., Garwood, N. C., & Kraft, N. J. B. (2016). Functional trait differences influence neighbourhood interactions in a hyperdiverse Amazonian forest. *Ecology Letters*, 19, 1062–1070.
- Freschet, G. T., Cornelissen, J. H. C., van Logtestijn, R. S. P., & Aerts, R. (2010). Substantial nutrient resorption from leaves, stems and roots in a subarctic flora: What is the link with other resource economics traits? *New Phytologist*, 186, 879–889.
- Gibert, A., Gray, E. F., Westoby, M., Wright, I. J., & Falster, D. S. (2016). On the link between functional traits and growth rate: meta-analysis shows effects change with plant size, as predicted. *Journal of Ecology*, 104, 1488–1503.
- Grime, J. P., & Hunt, R. (1975). Relative Growth-Rate: Its Range and Adaptive Significance in a Local Flora. *The Journal of Ecology*, 63, 393.
- Guevara-Andino, J. E., Pitman, N. C., Ulloa, C., Romoleroux, K., Fernández-Fernández, D., Ceron, C., Palacios, W., Neill, D. A., Oleas, N., Altamirano, P., Rivas-Torres, G., & ter Steege, H. (2019). Trees of Amazonian Ecuador: a taxonomically verified species list with data on abundance and distribution. *Ecology*, 100(12), e02894.
- Hendry, G. A., & Grime, J. P. (1993). Methods in comparative plant ecology: a laboratory manual. Springer Science & Business Media.
- Hunt, R. (1990). Basic Growth Analysis. Plant Growth Analysis for Beginners. Springer Science & Business Media.
- Iida, Y., Kohyama, T. S., Swenson, N. G., Su, S. H., Chen, C. T., Chiang, J. M., & Sun, I. F. (2014). Linking

functional traits and demographic rates in a subtropical tree community: The importance of size dependency. *Journal of Ecology*, *102*, 641–650.

- Kattge, J., Bönisch, G., Díaz, S., Lavorel, S., Prentice, I. C., Leadley, P., Tautenhahn, S., Werner, G. D. A., Aakala, T., Abedi, M., Acosta, A. T. R., Adamidis, G. C., Adamson, K., Aiba, M., Albert, C. H., Alcántara, J. M., Alcázar, C., Aleixo, I., Ali, H., ... Wirth, C. (2020). TRY plant trait database–enhanced coverage and open access. *Global Change Biology*, 26(1), 119–188.
- King, D. A., Davies, S. J., Nur Supardi, M. N., & Tan, S. (2005). Tree growth is related to light interception and wood density in two mixed dipterocarp forests of Malaysia. *Functional Ecology*, 19, 445–453.
- Le Roux, X., Walcroft, A. S., Daudet, F. A., Sinoquet, H., Chaves, M. M., Rodrigues, A., & Osorio, L. (2001). Photosynthetic light acclimation in peach leaves: Importance of changes in mass:area ratio, nitrogen concentration, and leaf nitrogen partitioning. *Tree Physiology*, 21, 377–386.
- Lewis, S. L., Lloyd, J., Sitch, S., Mitchard, E. T. A., & Laurance, W. F. (2009). Changing ecology of tropical forests: Evidence and drivers. *Annual Review of Ecology, Evolution, and Systematics*, 40, 529–549.
- Li, Y., Kröber, W., Bruelheide, H., Härdtle, H., & Von Oheimb, G. (2017). Crown and leaf traits as predictors of subtropical tree sapling growth rates. *Journal of Plant Ecology*, 10(1), 136–145.
- Lusk, C. H. (2002). Leaf area accumulation helps juvenile evergreen trees tolerate shade in a temperate rainforest. Oecologia, 132, 188–196.
- Maire, V., Gross, N., Hill, D., Martin, R., Wirth, C., Wright, I. J., & Soussana, J. F. (2013). Disentangling Coordination among Functional Traits Using an Individual-Centred Model: Impact on Plant Performance at Intra- and Inter-Specific Levels. PLOS ONE, 8, 1–16.
- Malhi, Y. (2010). The carbon balance of tropical forest regions, 1990-2005. Current Opinion in Environmental Sustainability, 2, 237–244.
- Marks, C. O., & Lechowicz, M. J. (2006). A holistic tree seedling model for the investigation of functional trait diversity. *Ecological Modelling*, 193(3), 141–181.
- Martínez-Encino, C., Villanueva-López, G., & Casanova-Lugo, F. (2013). Densidad y composición de árboles dispersos en potreros en la Sierra de Tabasco, México. Agrociencia, 47, 483–496.
- Martínez-Garza, C., Bongers, F., & Poorter, L. (2013). Are functional traits good predictors of species performance in restoration plantings in tropical abandoned pastures? *Forest Ecology and Management*, 303, 35–45.
- Mayaux, P., Holmgren, P., Achard, F., Eva, H., Stibig, H. J, & Branthomme, A. (2005). Tropical forest cover change

in the 1990s and options for future monitoring. *Philosophical Transactions of the Royal Society B: Biological Sciences*, 360, 373–384.

(i) (ii)

- Messier, J., McGill, B. J., & Lechowicz, M. J. (2010). How do traits vary across ecological scales? A case for traitbased ecology. *Ecology Letters*, 13, 838–848.
- Mooney, H. A., Field, C., Gulmon, S. L., & Bazzaz, F. A. (1981). Photosynthetic capacity in relation to leaf position in desert versus old-field annuals. *Oecologia*, 50, 109–112.
- Muller-Landau, H. C. (2004). Interspecific and inter-site variation in wood specific gravity of tropical trees. *Biotropica*, 36, 20–32.
- Niinemets, Ü. (2001). Global-scale climatic controls of leaf dry mass per area, density, and thickness in trees and shrubs. *Ecology*, 82, 453–469.
- Niinemets, Ü., & Kull, O. (1998). Stoichiometry of foliar carbon constituents varies along light gradients in temperate woody canopies: implications for foliage morphological plasticity. *Tree Physiology*, 18(7), 467–479.
- Obando-Vargas, F. V. (2003). Árboles del trópico húmedo. Importancia socioeconómica. Editorial Tecnológica de Costa Rica.
- Ordiales-Plaza, R. (2000). MideBMP (Versión 4.2, Software). Estación experimental de zonas áridas, España.
- Paine, C. T., Amissah, L., Auge, H., Baraloto, C., Baruffol, M., Bourland, N., Bruelheide, H., Dainou, K., de Gouvenain, R. C., Doucet, J. L., Doust, S., Fine, P. V. A., Fortunel, C., Haase, J., Holl, K. D., Jactel, H., Li, X., Kitajima, K., Koricheva, J., ... Hector, A. (2015). Globally, functional traits are weak predictors of juvenile tree growth, and we do not know why. *Journal of Ecology*, *103*(4), 978–989.
- Peel, M. C., Finlayson, B. L., & McMahon, T. A. (2007). Updated world map of the Köppen-Geiger climate classification. *Hydrology and Earth System Sciences*, 11, 1633–1644.
- Pérez-Harguindeguy, N., Diaz, S., Garnier, E., Lavorel, S., Poorter, H., Jaureguiberry, P., Bret-Harte, M. S., Cornwell, W. K., Craine, J. M., Gurvich, D. E., Urcelay, C., Veneklaas, E. J., Reich, P. B., Poorter, L., Wright, I. J., Ray, P., Enrico, L., Pausas, J. G., de Vos, A. C., ... Cornelissen, J. H. C. (2016). Corrigendum to: New handbook for standardised measurement of plant functional traits worldwide. *Australian Journal of Botany*, 64(8), 715–716.
- Poorter, L., & Bongers, F. (2006). Leaf traits are good predictors of plant performance across 53 rain forest species. *Ecology*, 87, 1733–1743.

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- Poorter, L., Bongers, L., & Bongers, F. (2006). Architecture of 54 moist-forest tree species: Traits, trade-offs, and functional groups. *Ecology*, 87, 1289–1301.
- Poorter, H., & Evans, J. R. (1998). Photosynthetic nitrogenuse efficiency of species that differ inherently in specific leaf area. Oecologia, 116, 26–37.
- Poorter, L., Wright, S. J., Paz, H., Ackerly, D. D., Condit, R., Ibarra-Manríquez, G., Harms, K. E., Licona, J. C., Martínez-Ramos, M., Mazer, S. J., Muller-Landau, H. C., Peña-Claros, M., Webb, C. O., & Wright, I. J. (2008). Are functional traits good predictors of demographic rates? Evidence from five neotropical forests. *Ecology*, 89(7), 1908–1920.
- Pozo-Rivera, W. E., Quiloango-Chimarro, C., Paredes, X., Landívar, M., Chiriboga, C., Hidalgo, D., García, K., & Villacís, J. (2023). Response of dung beetle diversity to remediation of soil ecosystems in the Ecuadorian Amazon. *PeerJ*, 11, e14975.
- Reich, P. B., Walters, M. B., Tjoelker, M. G., Vanderklein, D. W., & Buschena, C. (1998). Photosynthesis and respiration rates depend on, leaf and root morphology in nine boreal tree species differing in relative growth rate. *Functional Ecology*, 12, 395–405.
- Rivera-Parra, J. L., Vizcarra, C., Mora, K., Mayorga, H., & Dueñas, J. C. (2020). Spatial distribution of oil spills in the north eastern Ecuadorian Amazon: A comprehensive review of possible threats. *Biological Conservation*, 252, 108820.
- Rüger, N., Wirth, C., Wright, S. J., & Condit, R. (2012). Functional traits explain light and size response of growth rates in tropical tree species. *Ecology*, 93(12), 2626–2636.
- Santiago, L. S., Goldstein, G., Meinzer, F. C., Fisher, J. B., Machado, K., Woodruff, D., & Jones, T. (2004). Leaf photosynthetic traits scale with hydraulic conductivity and wood density in Panamanian forest canopy trees. *Oecologia*, 140, 543–550.
- Sterck, F. J., Poorter, L., & Schieving, F. (2006). Leaf traits determine the growth-survival trade-off across rain forest tree species. *The American Naturalist*, 167(5), 758–765.
- Swenson, N. G., & Enquist, B. J. (2007). Ecological and evolutionary determinants of a key plant functional trait: wood density and its community-wide variation across latitude and elevation. *American Journal of Botany*, 94(3), 451–459.
- Tecco, P. A., Díaz, S., Cabido, M., & Urcelay, C. (2010). Functional traits of alien plants across contrasting climatic and land-use regimes: do aliens join the locals or try harder than them? *Journal of Ecology*, 98(1), 17–27.
- Tecco, P. A., Urcelay, C., Díaz, S., Cabido, M., & Pérez-Harguindeguy, N. (2013). Contrasting functional trait

syndromes underlay woody alien success in the same ecosystem. *Austral Ecology*, *38*, 443–451.

- Uriarte, M., Lasky, J. R., Boukili, V. K., & Chazdon, R. L. (2016). A trait-mediated, neighbourhood approach to quantify climate impacts on successional dynamics of tropical rainforests. *Functional Ecology*, 30(2), 157–167.
- Vendramini, F., Díaz, S., Gurvich, D. E., Wilson, P. J., Thompson, K., & Hodgson, J. G. (2002). Leaf traits as indicators of resource-use strategy in floras with succulent species. *New Phytologist*, 154(1), 147–157.
- Villacís, J. (2016). Evaluación de las técnicas de remediación vegetal utilizadas en plataformas petroleras mediante estudios del desempeño de especies y análisis de diversidad funcional (Doctoral dissertation). Universidad Nacional de Córdoba, Argentina.
- Villacís, J., Armas, C., Hang, S., & Casanoves F. (2016). Selection of Adequate Species for Degraded Areas by Oil-Exploitation Industry as a Key Factor for Recovery Forest in the Ecuadorian Amazon. Land Degradation and Development, 27, 1771–1780.
- Villacís, J., Casanoves, F., Hang, S., Keesstra, S., & Armas, C. (2016). Selection of forest species for the rehabilitation of disturbed soils in oil fields in the Ecuadorian Amazon. Science of the Total Environment, 566, 761–770.
- Wiemann, M. C., & Williamson, G. B. (2002). Geographic variation in wood specific gravity: Effects of latitude, temperature, and precipitation. *Wood and Fiber Science*, 34, 96–107.
- Wang, C., Jian, S., Ren, H., Yan, J., & Liu, N. (2021). A Web-Based Software Platform for Restoration-Oriented Species Selection Based on Plant Functional Traits. *Frontiers in Ecology and Evolution*, 9, 570454.
- Werden, L. K., Alvarado, J. P., Zarges, S., Calderón, E., Schilling, E., Gutierrez, M., & Powers, J. (2018). Using soil amendments and plant functional traits to select native tropical dry forest species for the restoration of degraded Vertisols. *Journal of Applied Ecology*, 55, 1019–1028.
- Wright, I. J., & Cannon, K. (2001). Relationships between leaf lifespan and structural defences in a low-nutrient, sclerophyll flora. *Functional Ecology*, 15, 351–359.
- Wright, I. J., Groom, P. K., Lamont, B. B., Poot, P., Prior, L. D., Reich, P. B., Schulze, E. D., Veneklaas, E. J., & Westoby, M. (2004). Leaf trait relationships in Australian plant species. *Functional Plant Biology*, 31, 551–558.
- Yadav, M. R., Choudhary, M., Singh, J., Lal, M. K., Jha, P. K., Udawat, P., Gupta, N. K., Rajput, V. D., Garg, N. K., Maheshwari, C., Hasan, M., Gupta, S., Jatwa, T. K., Kumar, R., Yadav, A. K., & Prasad, P. V. (2022). Impacts, tolerance, adaptation, and mitigation of heat stress on wheat under changing climates. *International Journal of Molecular Sciences*, 23(5), 2838.