
This is the **accepted version** of the article:

Flores Moreno, Habacuc; Fazayeli, Farideh; Banerjee, Arindam; [et al.]. «Robustness of trait connections across environmental gradients and growth forms». *Global ecology and biogeography*, Vol. 28, issue 12 (Dec. 2019), p. 1806-1826. DOI 10.1111/geb.12996

This version is available at <https://ddd.uab.cat/record/218224>

under the terms of the  **COPYRIGHT** license

Appendix S1 List of studies including in the subset of TRY DB used in the current study.

- Adriaenssens S. (2012). Dry deposition and canopy exchange for temperate tree species under high nitrogen deposition. PhD thesis, Ghent University, Ghent, Belgium, 209p.
- Auger, S., Shipley, B. (2012). : Interspecific and intraspecific trait variation along short environmental gradients in an old-growth temperate forest. *Journal of Vegetation Science*. DOI: 1111/j.1654-1103.2012.01473.x
- B. Sandel, J. D. Corbin, and M. Krupa 2011. Using plant functional traits to guide restoration: A case study in California coastal grassland. *Ecosphere* 2:art23. <http://d>
- Bahn, M., G. Wohlfahrt, E. Haubner, I. Horak, W. Michaeler, K. Rottmar, U. Tappeiner, and A. Cernusca. 1999. Leaf photosynthesis, nitrogen contents and specific leaf area of 30 grassland species in differently managed mountain ecosystems in the Eastern Alps. Pages 247-255 in A. Cernusca, U. Tappeiner, and N. Bayfield, editors. *Land-use changes in European mountain ecosystems. ECOMONT- Concept and Results*. Blackwell Wissenschaft, Berlin.
- Bakker, C., J. Rodenburg, and P. Bodegom. 2005. Effects of Ca- and Fe-rich seepage on P availability and plant performance in calcareous dune soils. *Plant and Soil* 275:111-122.
- Bakker, C., P. M. Van Bodegom, H. J. M. Nelissen, W. H. O. Ernst, and R. Aerts. 2006. Plant responses to rising water tables and nutrient management in calcareous dune slacks. *Plant Ecology* 185:19-28.
- Baraloto, C., C. E. T. Paine, L. Poorter, J. Beauchene, D. Bonal, A.-M. Domenach, B. Herault, S. Patino, J.-C. Roggy, and J. Chave. 2010. Decoupled leaf and stem economics in rainforest trees. *Ecology Letters* 13:1338-1347
- Blonder, B., Buzzard, B., Sloat, L., Simova, I., Lipson, R., Boyle, B., Enquist, B. The shrinkage effect biases estimates of paleoclimate. *American Journal of Botany* (2012).
- Blonder, B., Vasseur, F., Violle, C., Shipley, B., Enquist, B., Vile, D. *Arabidopsis thaliana* rejects theories for the origin of the leaf economics spectrum. (in review, *New Phytologist*).
- Blonder, B., Violle, C., Enquist, B. Assessing the causes and scales of the leaf economics spectrum using venation networks in *Populus tremuloides*. (accepted, *Journal of Ecology*, 2013).
- Blonder, B., Violle, C., Patrick, L., Enquist, B. Leaf venation networks and the origin of the leaf economics spectrum. *Ecology Letters*, 2011.
- Bond-Lamberty, B., C. Wang, and S. T. Gower (2002), Above- and belowground biomass and sapwood area allometric equations for six boreal tree species of northern Manitoba, *Can. J. For. Res.*, 32(8), 1441-1450.
- Bond-Lamberty, B., C. Wang, and S. T. Gower (2002), Leaf area dynamics of a boreal black spruce fire chronosequence, *Tree Physiol.*, 22(14), 993-1001.
- Bond-Lamberty, B., C. Wang, and S.T. Gower (2004), Net primary production and net ecosystem production of a boreal black spruce fire chronosequence, *Global Change Biol.*, 10(4), 473-487.
- Brendan Choat, Steven Jansen, Tim J. Brodribb, Herve Cochard, Sylvain Delzon, Radika Bhaskar, Sandra J. Bucci, Taylor S. Feild, Sean M. Gleason, Uwe G. Hacke, Anna L. Jacobsen, Frederic Lens, Hafiz Maherali, Jordi Martinez-Vilalta, Stefan Mayr, Maurizio Mencuccini, Patrick J. Mitchell, Andrea Nardini, Jarmila Pittermann, R. Brandon Pratt, John S. Sperry, Mark Westoby, Ian J. Wright & Amy E. Zanne (2012) Global convergence in the vulnerability of forests to drought. *Nature* 491:752-755 doi:10.1038/nature11688
- Brown et al. (2011) *PLoS ONE* 6(9): e24107. doi:10.1371/journal.pone.0024107

Burrascano S., Del Vico E., Fagiani S., Giarrizzo E., Mei M., Mortelliti A., Sabatini F.M., Blasi C., submitted. Environmental filtering of wild boar rooting activity on understory composition and functional traits. *Plant Ecology*.

Butterfield, B.J. and J.M. Briggs. 2011. Regeneration niche differentiates functional strategies of desert woody plant species. *Oecologia*, 165:477-487.

Campetella, G; Botta-Duk\zelt, Z; Wellstein, C; Canullo, R; Gatto, S; Chelli, S; Mucina, L; Bartha, S (2011): Patterns of plant trait-environment relationships along a forest succession chronosequence. *Agriculture, Ecosystems & Environment*, 145(1), 38-48.
doi:10.1016/j.agee.2011.06.025

Carswell, F. E., Meir, P., Wandelli, E. V., Bonates, L. C. M., Kruijt, B., Barbosa, E. M., Nobre, A. D. & Jarvis, P. G. 2000 Photosynthetic capacity in a central Amazonian rain forest. *Tree physiology*. 20, 3, p. 179-186 8 p.

Cavender-Bares, J., A. Keen, and B. Miles. 2006. Phylogenetic structure of floridian plant communities depends on taxonomic and spatial scale. *Ecology* 87:S109-S122.

Cavender-Bares, J., L. Sack, and J. Savage. 2007. Atmospheric and soil drought reduce nocturnal conductance in live oaks. *Tree Physiology* 27:611-620.

Cerabolini B.E.L., Brusa G., Ceriani R.M., De Andreis R., Luzzaro A. & Pierce S. 2010. Can CSR classification be generally applied outside Britain? *Plant Ecology* 210: 253-261

Coomes, D. A., S. Heathcote, E. R. Godfrey, J. J. Shepherd, and L. Sack. 2008. Scaling of xylem vessels and veins within the leaves of oak species. *Biology Letters* 4:302-306.

Cornelissen, J. H. C. 1996. An experimental comparison of leaf decomposition rates in a wide range of temperate plant species and types. *Journal of Ecology* 84:573-582.

Cornelissen, J. H. C., B. Cerabolini, P. Castro-Diez, P. Villar-Salvador, G. Montserrat-Marti, J. P. Puyravaud, M. Maestro, M. J. A. Werger, and R. Aerts. 2003. Functional traits of woody plants: correspondence of species rankings between field adults and laboratory-grown seedlings? *Journal of Vegetation Science* 14:311-322.

Cornelissen, J. H. C., H. M. Quested, D. Gwynn-Jones, R. S. P. Van Logtestijn, M. A. H. De Beus, A. Kondratchuk, T. V. Callaghan, and R. Aerts. 2004. Leaf digestibility and litter decomposability are related in a wide range of subarctic plant species and types. *Functional Ecology* 18:779-786.

Cornelissen, J. H. C., N. Perez-Harguindeguy, S. Diaz, J. P. Grime, B. Marzano, M. Cabido, F. Vendramini, and B. Cerabolini. 1999. Leaf structure and defence control litter decomposition rate across species and life forms in regional floras on two continents. *New Phytologist* 143:191-200.

Cornelissen, J. H. C., P. C. Diez, and R. Hunt. 1996. Seedling growth, allocation and leaf attributes in a wide range of woody plant species and types. *Journal of Ecology* 84:755-765.

Cornwell, W. K., J. H. C. Cornelissen, K. Amatangelo, E. Dorrepaal, V. T. Eviner, O. Godoy, S. E. Hobbie, B. Hoorens, H. Kurokawa, N. Perez-Harguindeguy, H. M. Quested, L. S. Santiago, D. A. Wardle, I. J. Wright, R. Aerts, S. D. Allison, P. van Bodegom, V. Brovkin, A. Chatain, T. V. Callaghan, S. D\xedez, E. Garnier, D. E. Gurvich, E. Kazakou, J. A. Klein, J. Read, P. B. Reich, N. A. Soudzilovskaia, M. V. Vaieretti, and M. Westoby. 2008. Plant species traits are the predominant control on litter decomposition rates within biomes worldwide. *Ecology Letters* 11:1065-1071.

Cornwell, W. K., R. Bhaskar, L. Sack, S. Cordell, and C. K. Lunch. 2007. Adjustment of structure and function of Hawaiian *Metrosideros polymorpha* at high vs. low precipitation. *Functional Ecology* 21:1063-1071.

Craine JM, Nippert JB, Towne EG, Tucker S, Kembel SW, Skibbe A, McLauchlan KK (2011) Functional consequences of climate-change induced plant species loss in a tallgrass prairie. *Oecologia* 165: 1109-1117

Craine JM, Ocheltree TW, Nippert JB, Towne EG, Skibbe AM, Kembel SW, Fargione JE (in press) Global diversity of drought tolerance and grassland climate-change resilience. *Nature Climate Change*:

Craine JM, Towne EG, Ocheltree TW, Nippert JB (2012) Community traitscape of foliar nitrogen isotopes reveals N availability patterns in a tallgrass prairie. *Plant Soil* 356: 395-403

Craine, J. M., A. J. Elmore, M. P. M. Aida, M. Bustamante, T. E. Dawson, E. A. Hobbie, A. Kahmen, M. C. Mack, K. K. McLauchlan, A. Michelsen, G. B. Nardoto, L. H. Pardo, J. Penuelas, P. B. Reich, E. A. G. Schuur, W. D. Stock, P. H. Templer, R. A. Virginia, J. M. Welker, and I. J. Wright. 2009. Global patterns of foliar nitrogen isotopes and their relationships with climate, mycorrhizal fungi, foliar nutrient concentrations, and nitrogen availability. *New Phytologist* 183:980-992.

D\xedez, S., J. G. Hodgson, K. Thompson, M. Cabido, J. H. C. Cornelissen, A. Jalili, G. Montserrat-Mart\xed, J. P. Grime, F. Zarrinkamar, Y. Asri, S. R. Band, S. Basconcelo, P. Castro-D\xedez, G. Funes, B. Hamzehee, M. Khoshnevi, N. P\xedez-Harguindeguy, M. C. P\xedez-Rontom\xed, F. A. Shirvany, F. Vendramini, S. Yazdani, R. Abbas-Azimi, A. Bogaard, S. Boustani, M. Charles, M. Dehghan, L. de Torres-Espuny, V. Falczuk, J. Guerrero-Campo, A. Hynd, G. Jones, E. Kowsary, F. Kazemi-Saeed, M. Maestro-Mart\xednez, A. Romo-D\xedez, S. Shaw, B. Siavash, P. Villar-Salvador, and M. R. Zak. 2004. The plant traits that drive ecosystems: Evidence from three continents. *Journal of Vegetation Science* 15:295-304.

Demey A, J Staelens, L Baeten, P Boeckx, M Hermy, J Kattge, K Verheyen (2013) Nutrient input from hemiparasitic litter favors plant species with a fast-growth strategy. *Plant and Soil* DOI 10.1007/s11104-013-1658-4

Domingues TF, Meir P, Feldpausch TR, et al. (2010) Co-limitation of photosynthetic capacity by nitrogen and phosphorus in West Africa woodlands. *Plant, Cell & Environment* (33): 959-980.

Dunbar-Co, S., M. J. Sporck, and L. Sack. 2009. Leaf Trait Diversification and Design in Seven Rare Taxa of the Hawaiian *Plantago* Radiation. *International Journal of Plant Sciences* 170:61-75.

Fonseca, C. R., J. M. Overton, B. Collins, and M. Westoby. 2000. Shifts in trait-combinations along rainfall and phosphorus gradients. *Journal of Ecology* 88:964-977.

Frenette-Dussault, C., Shipley, B., L\xferger, J.F., Meziane, D. & Hingrat, Y. (2012). Functional structure of an arid steppe plant community reveals similarities with Grime's C-S-R theory. *Journal of Vegetation Science* 23: 208-222.

Freschet, G. T., J. H. C. Cornelissen, R. S. P. van Logtestijn, and R. Aerts. 2010. Evidence of the \x91plant economics spectrum\x92 in a subarctic flora. *Journal of Ecology* 98:362-373.

Fyllas, N. M., S. Patino, T. R. Baker, G. Bielefeld Nardoto, L. A. Martinelli, C. A. Quesada, R. Paiva, M. Schwarz, V. Horna, L. M. Mercado, A. Santos, L. Arroyo, E. M. Jimenez, F. J. Luizao, D. A. Neill, N. Silva, A. Prieto, A. Rudas, M. Silviera, I. C. G. Vieira, G. Lopez-Gonzalez, and J. Lloyd. 2009. Basin-wide variations in foliar properties of Amazonian forest: phylogeny, soils and climate *Biogeosciences* 6:2677-2708.

Garnier, E., S. Lavorel, P. Ansquer, H. Castro, P. Cruz, J. Dolezal, O. Eriksson, C. Fortunel, H. Freitas, C. Golodets, K. Grigulis, C. Jouany, E. Kazakou, J. Kigel, M. Kleyer, V. Lehsten, J. Lep\x9a, T. Meier, R. Pakeman, M. Papadimitriou, V. P. Papanastasis, H. Quested, F. Qu\x9tier, M. Robson, C. Roumet, G. Rusch, C. Skarpe, M. Sternberg, J.-P. Theau, A.

Thibault, D. Vile, and M. P. Zarovali. 2007. Assessing the effects of land-use change on plant traits, communities and ecosystem functioning in grasslands: A standardized methodology and lessons from an application to 11 European sites. *Annals of Botany* 99:967-985.

Givnish T.J., R.A. Montgomery and G. Goldstein. 2004. Adaptive radiation of photosynthetic physiology in the Hawaiian lobeliads: light regimes, static light responses, and whole-plant compensation points. *American Journal of Botany* 91: 228-246

Guerin G.R., Wen H. & Lowe A.J. (2012) Leaf morphology shift linked to climate change. *Biology Letters* 8, 882-886.

Gutiérrez AG, & Huth A (2012) Successional stages of primary temperate rainforests of Chiloe Island, Chile. *Perspectives in plant ecology, systematics and evolution*. 14: 243-256

Guy, A. L., J. M. Mischkolz, and E. G. Lamb. 2013. Limited effects of simulated acidic deposition on seedling survivorship and root morphology of endemic plant taxa of the Athabasca Sand Dunes in well watered greenhouse trials. *Botany* 91:176-181. <http://dx.doi.org/10.1139/cjb-2012-0162>

Han, W. X., J. Y. Fang, D. L. Guo, and Y. Zhang. 2005. Leaf nitrogen and phosphorus stoichiometry across 753 terrestrial plant species in China. *New Phytologist* 168:377-385.

Hao, G. Y., L. Sack, A. Y. Wang, K. F. Cao, and G. Goldstein. 2010. Differentiation of leaf water flux and drought tolerance traits in hemiepiphytic and non-hemiepiphytic *Ficus* tree species. *Functional Ecology* 24:731-740.

Hoof, J., L. Sack, D. T. Webb, and E. T. Nilsson. 2008. Contrasting structure and function of pubescent and glabrous varieties of Hawaiian *Metrosideros polymorpha* (Myrtaceae) at high elevation. *Biotropica* 40:113-118.

Jennifer S. Powers and Peter Tiffin 2012 Plant functional type classifications in tropical dry forests in Costa Rica: leaf habit versus taxonomic approaches. *Functional Ecology* 2010, 24, 927-936 doi: 10.1111/j.1365-2435.2010.01701.x

Kattge, J., W. Knorr, T. Raddatz, and C. Wirth. 2009. Quantifying photosynthetic capacity and its relationship to leaf nitrogen content for global-scale terrestrial biosphere models. *Global Change Biology* 15:976-991.

Kazakou, E., D. Vile, B. Shipley, C. Gallet, and E. Garnier. 2006. Co-variations in litter decomposition, leaf traits and plant growth in species from a Mediterranean old-field succession. *Functional Ecology* 20:21-30.

Kichenin et al. 2013. Contrasting effects of plant inter- and intraspecific variation on community-level trait measures along an environmental gradient. *Functional Ecology*, in press.

Kleyer, M., R. M. Bekker, I. C. Knevel, J. P. Bakker, K. Thompson, M. Sonnenschein, P. Poschlod, J. M. van Groenendael, L. Klimes, J. Klimesova, S. Klotz, G. M. Rusch, Hermy, M., D. Adriaens, G. Boedeltje, B. Bossuyt, A. Dannemann, P. Endels, L. Götzenberger, J. G. Hodgson, A.-K. Jackel, I. Kutsch, D. Kunzmann, W. A. Ozinga, C. Rötter, M. Stadler, J. Schlegelmilch, H. J. Steendam, O. Tackenberg, B. Wilmann, J. H. C. Cornelissen, O. Eriksson, E. Garnier, and B. Peco. 2008. The LEDA Traitbase: a database of life-history traits of the Northwest European flora. *Journal of Ecology* 96:1266-1274.

Kraft, N. J. B., R. Valencia, and D. Ackerly. 2008. Functional traits and niche-based tree community assembly in an Amazonian forest. *Science* 322:580-582.

Kurokawa, H. and T. Nakashizuka. 2008. Leaf herbivory and decomposability in a Malaysian tropical rain forest. *Ecology* 89:2645-2656.

Laughlin, D. C., J. J. Leppert, M. M. Moore, and C. H. Sieg. 2010. A multi-trait test of the leaf-height-seed plant strategy scheme with 133 species from a pine forest flora. *Functional Ecology* 24:493-501.

Laughlin, D.C., P.Z. Fulmer, D.W. Huffman, J. Crouse, and E. Laliberte. 2011. Climatic constraints on trait-based forest assembly. *Journal of Ecology* 99:1489-1499.

Louault, F., V. D. Pillar, J. Aufrere, E. Garnier, and J. F. Soussana. 2005. Plant traits and functional types in response to reduced disturbance in a semi-natural grassland. *Journal of Vegetation Science* 16:151-160.

Markesteijn, L., L. Poorter, H. Paz, L. Sack, and F. Bongers. 2011. Ecological differentiation in xylem cavitation resistance is associated with stem and leaf structural traits. *Plant, Cell and Environment* 34:137-148.

Martin, R. E., G. P. Asner, and L. Sack. 2007. Genetic variation in leaf pigment, optical and photosynthetic function among diverse phenotypes of *Metrosideros polymorpha* grown in a common garden. *Oecologia* 151:387-400.

Medlyn, B. E., F.-W. Badeck, D. G. G. De Pury, C. V. M. Barton, M. Broadmeadow, R. Ceulemans, P. De Angelis, M. Forstreuter, M. E. Jach, S. Kellomäki, E. Laitat, M. Marek, S. Philippot, A. Rey, J. Strassmeyer, K. Laitinen, R. Liozon, B. Portier, P. Roberntz, K. Wang, and P. G. Jarvis. 1999. Effects of elevated CO₂ on photosynthesis in European forest species: a meta-analysis of model parameters. *Plant, Cell and Environment* 22:1475-1495.

Meir, P. & Levy, P. E. 2007 Photosynthetic parameters from two contrasting woody vegetation types in West Africa. *Plant Ecology*. 192, 2, p. 277-287 11 p.

Meir, P., Kruijt, B., Broadmeadow, M., Kull, O., Carswell, F. & Nobre, A. 2002 Acclimation of photosynthetic capacity to irradiance in tree canopies in relation to leaf nitrogen concentration and leaf mass per unit area. *Plant, Cell and Environment*. 25, 3, p. 343-357 15 p.

Messier, J., B. J. McGill, and M. J. Lechowicz. 2010. How do traits vary across ecological scales? A case for trait-based ecology. *Ecology Letters* 13:838-848.

Meziane, D. and B. Shipley. 1999. Interacting determinants of specific leaf area in 22 herbaceous species: effects of irradiance and nutrient availability. *Plant Cell and Environment* 22:447-459.

Milla & Reich 2011 *Annals of Botany* 107: 455-465, 2011.

Muller, S. C., G. E. Overbeck, J. Pfadenhauer, and V. D. Pillar. 2007. Plant functional types of woody species related to fire disturbance in forest-grassland ecotones. *Plant Ecology* 189:1-14.

Nakahashi, C. D., K. Frole, and L. Sack. 2005. Bacterial leaf nodule symbiosis in *Ardisia* (Myrsinaceae): Does it contribute to seedling growth capacity? *Plant Biology* 7:495-500.

Niinemets, U. 2001. Global-scale climatic controls of leaf dry mass per area, density, and thickness in trees and shrubs. *Ecology* 82:453-469.

Ogaya, R. and J. Penuelas. 2003. Comparative field study of *Quercus ilex* and *Phillyrea latifolia*: photosynthetic response to experimental drought conditions. *Environmental and Experimental Botany* 50:137-148.

Onoda, Y., M. Westoby, P. B. Adler, A. M. F. Choong, F. J. Clissold, J. H. C. Cornelissen, S. Diaz, N. J. Dominy, A. Elgart, L. Enrico, P. V. A. Fine, J. J. Howard, A. Jalili, K. Kitajima, H. Kurokawa, C. McArthur, P. W. Lucas, L. Markesteijn, N. Perez-Harguindeguy, L. Poorter, L. Richards, L. S. Santiago, Jr. E. Sosinski, S. Van Bael, D. I. Warton, I. J. Wright, S. J. Wright, and N. Yamashita. 2011. Global patterns of leaf mechanical properties. *Ecology Letters* 14:301-312.

Ordóñez, J. C., P. M. van Bodegom, J. P. M. Witte, R. P. Bartholomeus, J. R. van Hal, and R. Aerts. 2010. Plant Strategies in Relation to Resource Supply in Mesic to Wet Environments: Does Theory Mirror Nature? *American Naturalist* 175:225-239.

Pahl, A.T., Kollmann, J., Mayer, A. & Haider, S. (2013): No evidence for local adaptation in an invasive alien plant: field and greenhouse experiments tracing a colonization sequence. *Annals of Botany* 112 (9): 1921-1930. DOI: 10.1093/aob/mct246

Peco B., de Pablos I., Traba J. , & Levassor C. (2005) The effect of grazing abandonment on species composition and functional traits: the case of dehesa *Basic and Applied Ecology*, 6(2): 175-183

Penuelas, J., J. Sardans, J. Llusia, S. Owen, J. Carnicer, T. W. Giambelluca, E. L. Rezende, M. Waite, and Niinemets. 2010. Faster returns on leaf economics and different biogeochemical niche in invasive compared with native plant species. *Global Change Biology* 16:2171-2185.

PETER B. ADLER, DANIEL G. MILCHUNAS, WILLIAM K. LAUENROTH, OSVALDO E. SALA and INGRID C. BURKE (2004) Functional traits of graminoids in semi-arid steppes: a test of grazing histories. *Journal of Applied Ecology* 2004 41, 653-663

Pierce S., Brusa G., Sartori M. & Cerabolini B.E.L. 2012. Combined use of leaf size and economics traits allows direct comparison of hydrophyte and terrestrial herbaceous adaptive strategies. *Annals of Botany* 109(5): 1047-1053

Pierce S., Brusa G., Vagge I. & Cerabolini B.E.L. 2013. Allocating CSR plant functional types: the use of leaf economics and size traits to classify woody and herbaceous vascular plants. *Functional Ecology* (in press) DOI: 10.1111/1365-2435.12095.

Pierce S., Ceriani R.M., De Andreis R., Luzzaro A. & Cerabolini B. 2007. The leaf economics spectrum of Poaceae reflects variation in survival strategies. *Plant Biosystems* 141(3): 337-343.

Pierce S., Luzzaro A., Caccianiga M., Ceriani R.M. & Cerabolini B. 2007. Disturbance is the principal α -scale filter determining niche differentiation, coexistence and biodiversity in an alpine community. *Journal of Ecology* 95: 698-706.

Pillar, V. D. and E. E. Sosinski. 2003. An improved method for searching plant functional types by numerical analysis. *Journal of Vegetation Science* 14:323-332.

Prentice, I.C., Meng, T., Wang, H., Harrison, S.P., Ni, J., Wang, G., 2011. Evidence for a universal scaling relationship of leaf CO₂ drawdown along a moisture gradient. *New Phytologist* 190: 169-180

Preston, K. A., W. K. Cornwell, and J. L. DeNoyer. 2006. Wood density and vessel traits as distinct correlates of ecological strategy in 51 California coast range angiosperms. *New Phytologist* 170:807-818.

Price, C.A. and B.J. Enquist. Scaling of mass and morphology in Dicotyledonous leaves: an extension of the WBE model. (2007) *Ecology* 88(5): 1132-1141.

Pyankov, V. I., A. V. Kondratchuk, and B. Shipley. 1999. Leaf structure and specific leaf mass: the alpine desert plants of the Eastern Pamirs, Tadjikistan. *New Phytologist* 143:131-142.

Quero, J. L., R. Villar, T. Maranon, R. Zamora, D. Vega, and L. Sack. 2008. Relating leaf photosynthetic rate to whole-plant growth: drought and shade effects on seedlings of four *Quercus* species. *Functional Plant Biology* 35:725-737.

Quested, H. M., J. H. C. Cornelissen, M. C. Press, T. V. Callaghan, R. Aerts, F. Trosien, P. Riemann, D. Gwynn-Jones, A. Kondratchuk, and S. E. Jonasson. 2003. Decomposition of sub-arctic plants with differing nitrogen economies: A functional role for hemiparasites. *Ecology* 84:3209-3221.

- Reich, P. B., J. Oleksyn, and I. J. Wright. 2009. Leaf phosphorus influences the photosynthesis-nitrogen relation: a cross-biome analysis of 314 species. *Oecologia* 160:207-212.
- Reich, P. B., M. G. Tjoelker, K. S. Pregitzer, I. J. Wright, J. Oleksyn, and J. L. Machado. 2008. Scaling of respiration to nitrogen in leaves, stems and roots of higher land plants. *Ecology Letters* 11:793-801.
- Sack, L. 2004. Responses of temperate woody seedlings to shade and drought: do trade-offs limit potential niche differentiation? *Oikos* 107:110-127.
- Sack, L. and K. Frole. 2006. Leaf structural diversity is related to hydraulic capacity in tropical rain forest trees. *Ecology* 87:483-491.
- Sack, L., M. T. Tyree, and N. M. Holbrook. 2005. Leaf hydraulic architecture correlates with regeneration irradiance in tropical rainforest trees. *New Phytologist* 167:403-413.
- Sack, L., P. D. Cowan, N. Jaikumar, and N. M. Holbrook. 2003. The 'hydrology' of leaves: co-ordination of structure and function in temperate woody species. *Plant Cell and Environment* 26:1343-1356.
- Sack, L., P. J. Melcher, W. H. Liu, E. Middleton, and T. Pardee. 2006. How strong is intracanopy leaf plasticity in temperate deciduous trees? *American Journal of Botany* 93:829-839.
- Scherer-Lorenzen, M., Schulze, E.-D., Don, A., Schumacher, J. & Weller, E. (2007) Exploring the functional significance of forest diversity: A new long-term experiment with temperate tree species (BIOTREE). *Perspectives in Plant Ecology, Evolution and Systematics*, 9, 53-70.
- Schweingruber, F.H., Landolt, W.: The Xylem Database. Swiss Federal Research Institute WSL Updated (2005)
- Scoffoni, C., A. Pou, K. Aasamaa, and L. Sack. 2008. The rapid light response of leaf hydraulic conductance: new evidence from two experimental methods. *Plant Cell and Environment* 31:1803-1812.
- Shipley B., 2002. Trade-offs between net assimilation rate and specific leaf area in determining relative growth rate: relationship with daily irradiance, *Functional Ecology*(16) 682-689
- Shipley, B. 1995. Structured Interspecific Determinants of Specific Leaf-Area in 34 Species of Herbaceous Angiosperms. *Functional Ecology* 9:312-319.
- Shipley, B. and M. J. Lechowicz. 2000. The functional co-ordination of leaf morphology, nitrogen concentration, and gas exchange in 40 wetland species. *Ecoscience* 7:183-194.
- Shipley, B. and M. Parent. 1991. Germination Responses of 64 Wetland Species in Relation to Seed Size, Minimum Time to Reproduction and Seedling Relative Growth-Rate. *Functional Ecology* 5:111-118.
- Shipley, B. and T. T. Vu. 2002. Dry matter content as a measure of dry matter concentration in plants and their parts. *New Phytologist* 153:359-364.
- Soudzilovskaia, Nadejda A and Elumeeva, Tatiana G and Onipchenko, Vladimir G and Shidakov, Islam I and Salpagarova, Fatima S and Khubiev, Anzor B and Tekeev, Dzhamal K and Cornelissen, Johannes HC. 2013. Functional traits predict relationship between plant abundance dynamic and long-term climate warming. *PNAS*. 45:18180-18184
- Spasojevic, M. J. and K. N. Suding. 2012. Inferring community assembly mechanisms from functional diversity patterns: the importance of multiple assembly processes. *Journal of Ecology* 100:652-661.
- Swaine, E. K. 2007. Ecological and evolutionary drivers of plant community assembly in a Bornean rain forest. PhD Thesis, University of Aberdeen, Aberdeen.

Tomas F. Domingues, Luiz A. Martinelli, James R. Ehleringer (2007) Ecophysiological traits of plant functional groups in forest and pasture ecosystems from eastern Amazonia, Brazil. *Plant Ecol* (2007) 193:101-112 DOI 10.1007/s11258-006-9251-z

Tucker SS, Craine JM, Nippert JB (2011) Physiological drought tolerance and the structuring of tallgrass assemblages. *Ecosphere* 2(4): 48

van Bodegom, P. M., B. K. Sorrell, A. Oosthoek, C. Bakke, and R. Aerts. 2008. Separating the effects of partial submergence and soil oxygen demand on plant physiology. *Ecology* 89:193-204.

van Bodegom, P. M., M. de Kanter, C. Bakker, and R. Aerts. 2005. Radial oxygen loss, a plastic property of dune slack plant species. *Plant and Soil* 271:351-364.

Vanessa Minden, Michael Kleyer (2011): Testing the effect-response framework: key response and effect traits determining above-ground biomass of salt marshes. *Journal of Vegetation Science* 22: 387-401

Vanessa Minden, Sandra Andratschke, Janina Spalke, Hanna Timmermann, Michael Kleyer (2012): Plant trait-environment relationships in salt marshes: Deviations from predictions by ecological concepts. *Perspectives in Plant Ecology, Evolution and Systematics*, 14: 183-192

Vergutz, L., S. Manzoni, A. Porporato, R.F. Novais, and R.B. Jackson. 2012. A Global Database of Carbon and Nutrient Concentrations of Green and Senesced Leaves. Data set. Available on-line [<http://daac.ornl.gov>] from Oak Ridge National Laboratory Distributed Active Archive Center, Oak Ridge, Tennessee, U.S.A. <http://dx.doi.org/10.3334/ORNLDAAAC/1106>

Vile, D. 2005. Significations fonctionnelle et écologique des traits des espèces végétales: exemple dans une succession post-culturelle méditerranéenne et généralisations, PHD Thesis.

Von Holle, B. and D. Simberloff. 2004. Testing Fox's assembly rule: Does plant invasion depend on recipient community structure? *Oikos* 105:551-563.

Waite, M. and L. Sack. 2010. How does moss photosynthesis relate to leaf and canopy structure? Trait relationships for 10 Hawaiian species of contrasting light habitats. *New Phytologist* 185:156-172.

Wenxuan Han, Yahan Chen, Fang-Jie Zhao, Luying Tang, Rongfeng Jiang and Fusuo Zhang, 2011, Floral, climatic and soil pH controls on leaf ash content in China's terrestrial plants. *Global Ecology and Biogeography*, DOI: 10.1111/j.1466-8238.2011.00677.x

Williams, M., Y.E. Shimabokuro and E.B. Rastetter. 2012. LBA-ECO CD-09 Soil and Vegetation Characteristics, Tapajos National Forest, Brazil. Data set. Available on-line [<http://daac.ornl.gov>] from Oak Ridge National Laboratory Distributed Active Archive Center, Oak Ridge, Tennessee, U.S.A. <http://dx.doi.org/10.3334/ORNLDAAAC/1104>

WILSON K, D BALDOCCHI, P HANSON (2000) Spatial and seasonal variability of photosynthetic parameters and their relationship to leaf nitrogen in a deciduous forest. *Tree Physiology* 20, 565-578

Wirth, C. and J. W. Lichstein. 2009. The Imprint of Species Turnover on Old-Growth Forest Carbon Balances - Insights From a Trait-Based Model of Forest Dynamics. Pages 81-113 in C. Wirth, G. Gleixner, and M. Heimann, editors. *Old-Growth Forests: Function, Fate and Value*. Springer, New York, Berlin, Heidelberg.

Wright, I. J., D. D. Ackerly, F. Bongers, K. E. Harms, G. Ibarra-Manriquez, M. Martinez-Ramos, S. J. Mazer, H. C. Muller-Landau, H. Paz, N. C. A. Pitman, L. Poorter, M. R. Silman, C. F. Vriesendorp, C. O. Webb, M. Westoby, and S. J. Wright. 2007. Relationships among ecologically important dimensions of plant trait variation in seven Neotropical forests. *Annals of Botany* 99:1003-1015.

Wright, I. J., P. B. Reich, M. Westoby, D. D. Ackerly, Z. Baruch, F. Bongers, J. Cavender-Bares, T. Chapin, J. H. C. Cornelissen, M. Diemer, J. Flexas, E. Garnier, P. K. Groom, J. Gulias, K. Hikosaka, B. B. Lamont, T. Lee, W. Lee, C. Lusk, J. J. Midgley, M. L. Navas, U. Niinemets, J. Oleksyn, N. Osada, H. Poorter, P. Poot, L. Prior, V. I. Pyankov, C. Roumet, S. C. Thomas, M. G. Tjoelker, E. J. Veneklaas, and R. Villar. 2004. The worldwide leaf economics spectrum. *Nature* 428:821-827.

Wright, S. J., K. Kitajima, N. J. B. Kraft, P. B. Reich, I. J. Wright, D. E. Bunker, R. Condit, J. W. Dalling, S. J. Davies, S. D\xededaz, B. M. J. Engelbrecht, K. E. Harms, S. P. Hubbell, C. O. Marks, M. C. Ruiz-Jaen, C. M. Salvador, and A. E. Zanne. 2011 . Functional traits and the growth-mortality tradeoff in tropical trees. *Ecology* 91:3664-3674.

Yahan Chen , Wenxuan Han , Luying Tang , Zhiyao Tang and Jingyun Fang 2011 Leaf nitrogen and phosphorus concentrations of woody plants differ in responses to climate, soil and plant growth form. *Ecography* 34, doi: 10.1111/j.1600-0587.2011.06833.x

Yguel B., Bailey R., Tosh N.D., Vialatte A., Vasseur C., Vitrac X., Jean F. & Prinzing A. (2011). Phytophagy on phylogenetically isolated trees: why hosts should escape their relatives. *Ecol. Lett.*, 14, 1117-1124.

Appendix S2. Sample size for woody and non-woody species across climate regions, robustness test for the gap filling algorithm and examples of studies focused on multi-organ, multi-trait datasets.

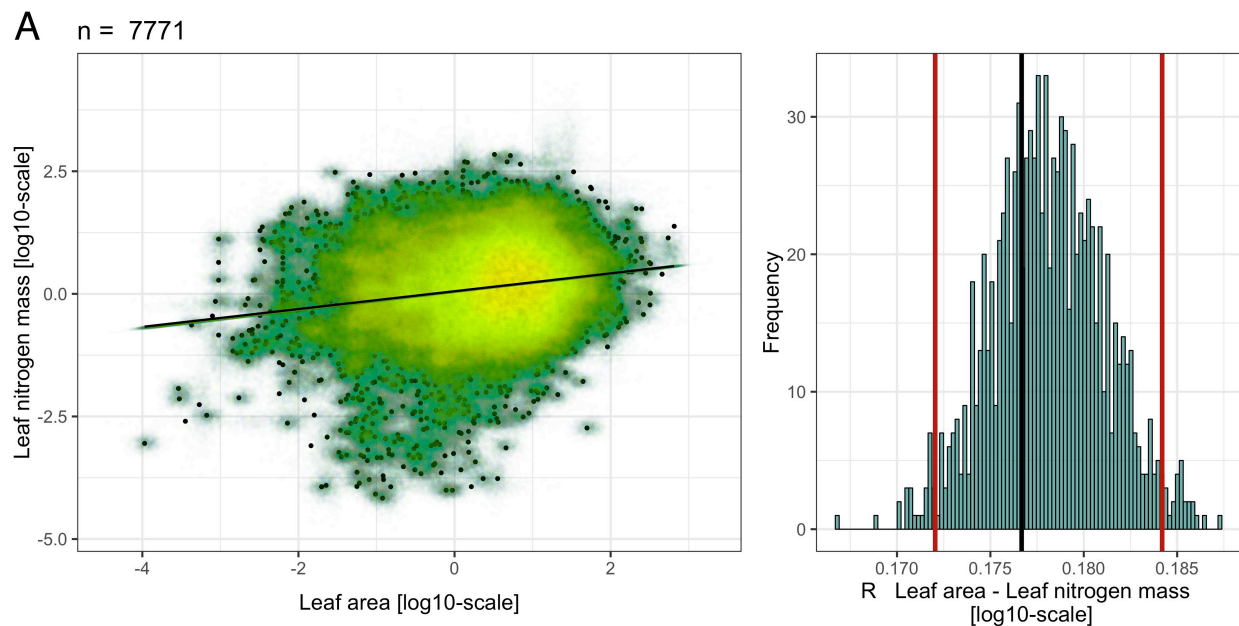
Table S2.1. Percentage of observation present by trait in the TRY subset used in this study. Traits in the table are order in ascendant order from the trait with the lowest coverage to the trait with the highest coverage in the dataset.

Trait	Percentage observation present
Leaf lifespan	0.6728772
Leaf phosphorus area	2.5561542
Leaf nitrogen area	5.7611659
Leaf phosphorus mass	6.8018047
Stem specific density	9.8029733
Leaf nitrogen mass	14.3595819
Seed mass	19.8256946
Leaf area	27.8021942
SLA	29.1242534
Plant height	35.6462607

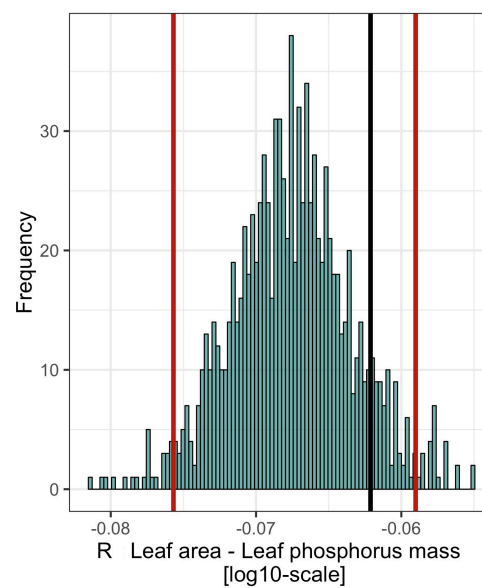
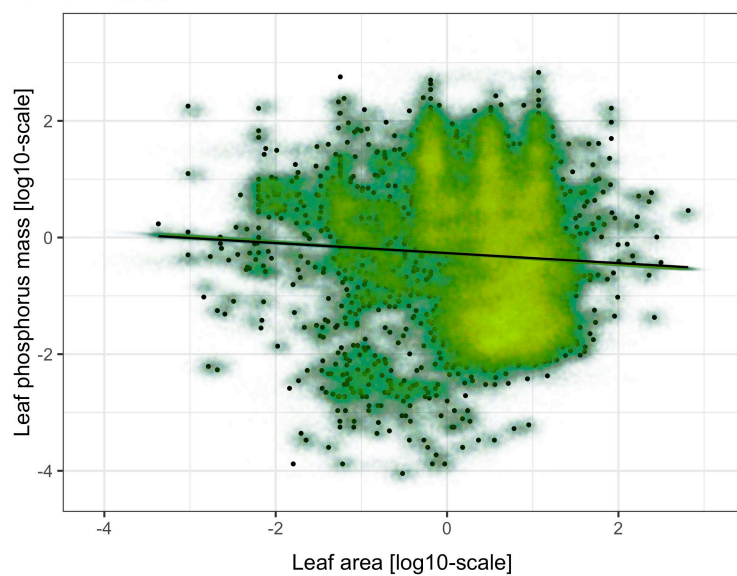
Table S2.3 Number of species by growth form across five climate regions.

Climate region	Woody	Non-woody
Tropical	4414	537
Temperate	2695	2352
Arid	1046	873
Cold	752	2050
Polar	146	419

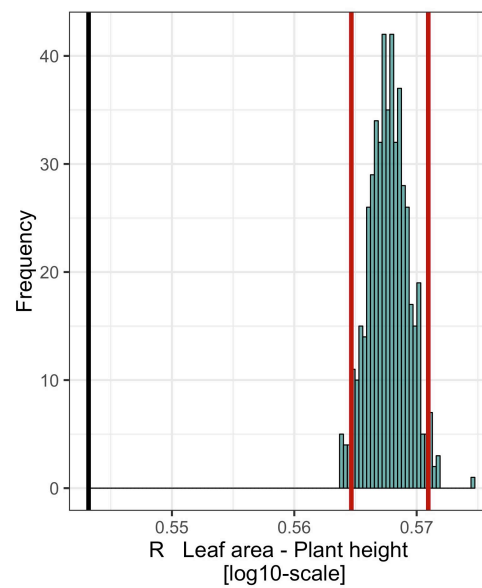
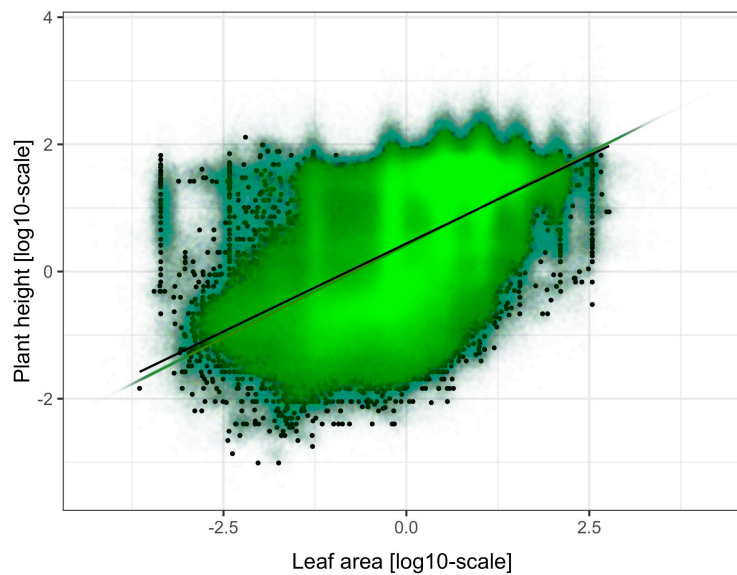
Figure S2.1 Comparison of trait-trait correlations using only gap-filled or only original trait values. Sample sizes (n) of the gap filled and original database are the same to ensure they are comparable. To test whether the gap-filling algorithm had an effect on trait-trait correlation we used a subset of 470769 observations from TRY for five traits (leaf area, SLA, leaf n, plant height and seed mass). We compare the trait-trait correlation for 1000 different gap-filled matrices with the trait-trait correlation of the original data. We did this comparison for trait-trait correlation with low and high sample size. We show that in most cases, the original trait-trait correlation lies inside the 95% CI of the 1000 gap-filled datasets. Sample size for each trait-trait correlation is given on the right top corner of each panel.



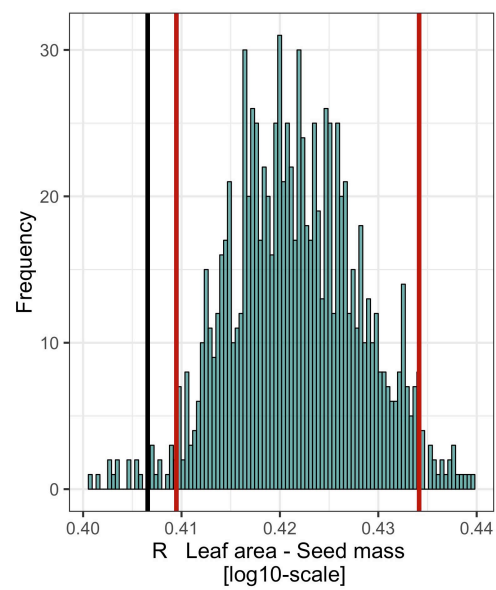
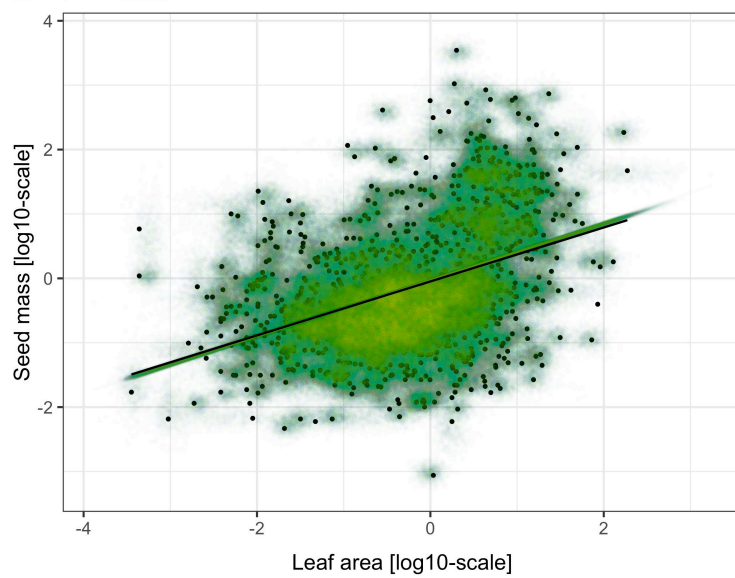
B n = 2739



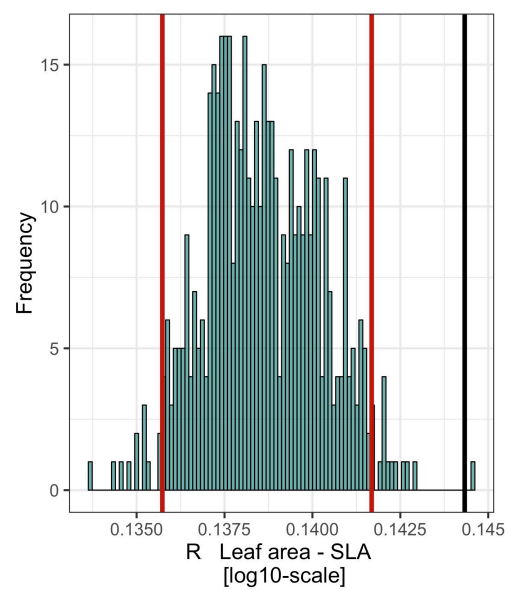
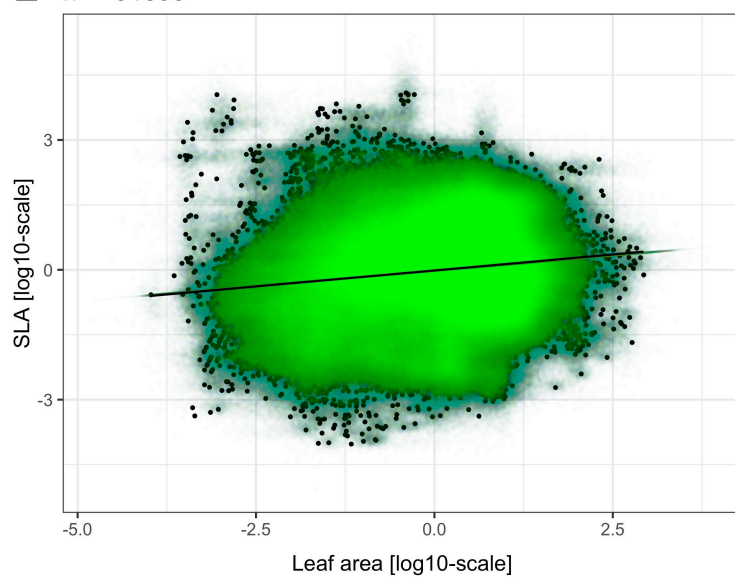
C n = 27562



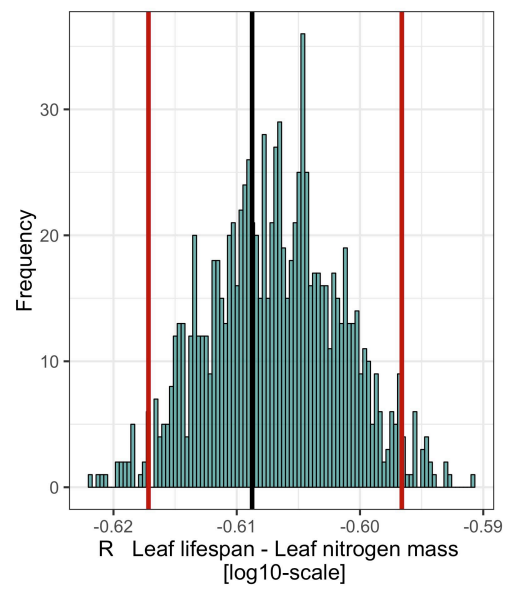
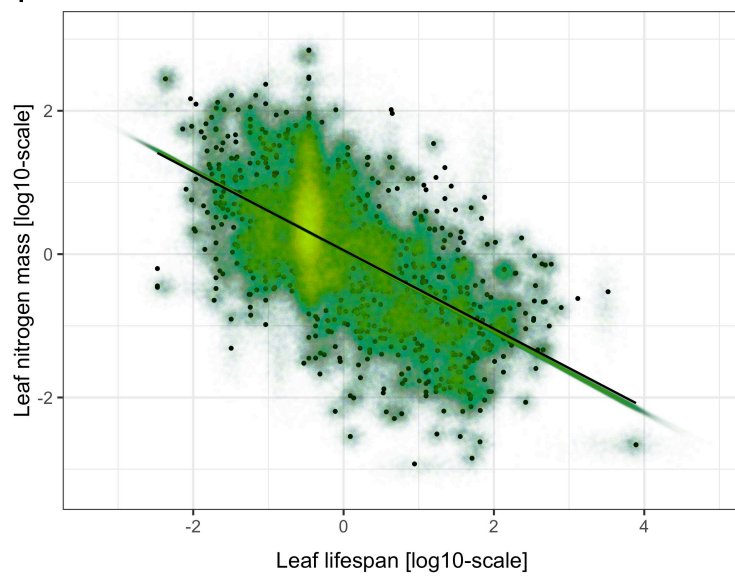
D $n = 1228$



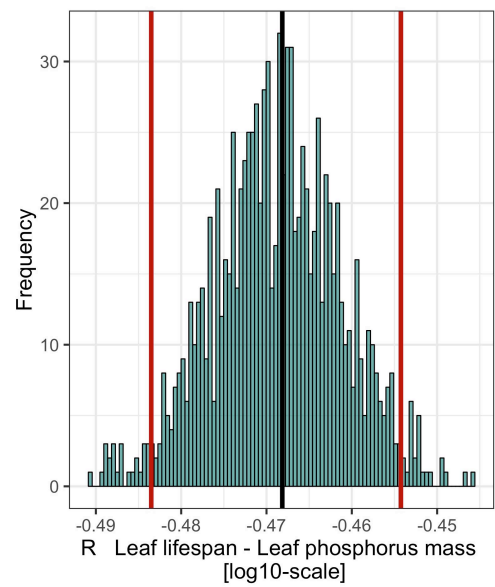
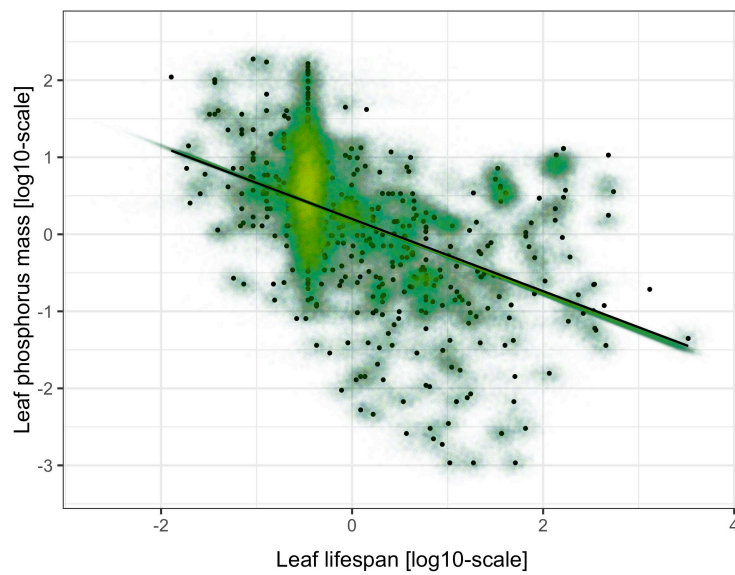
E $n = 51899$



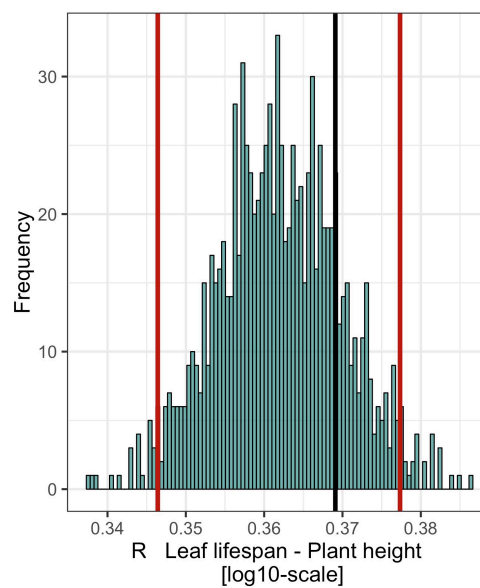
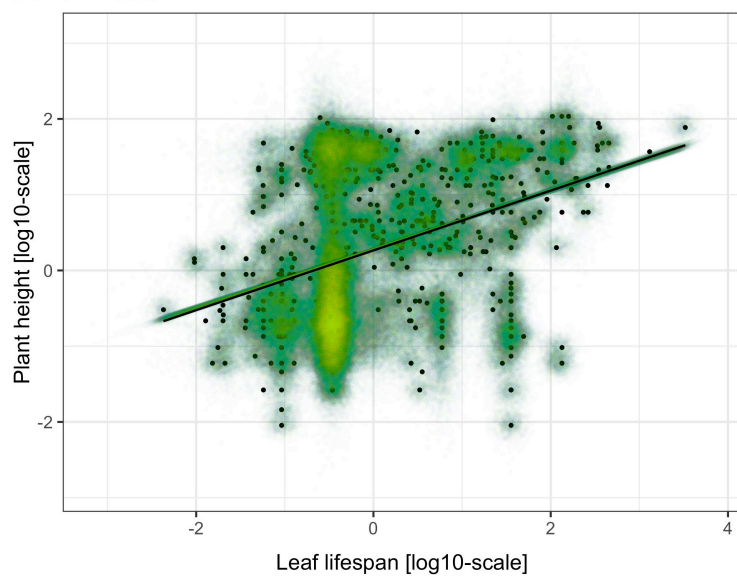
F n = 1303



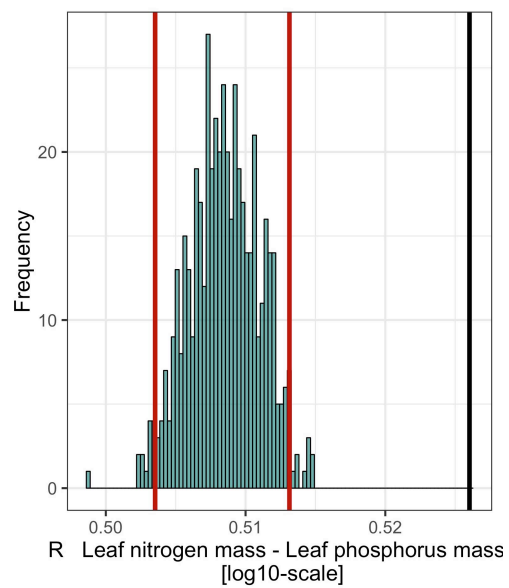
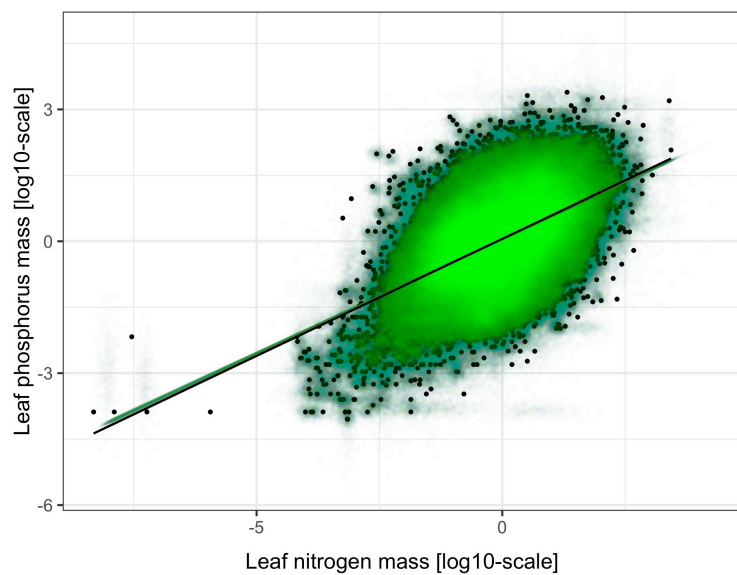
G n = 664



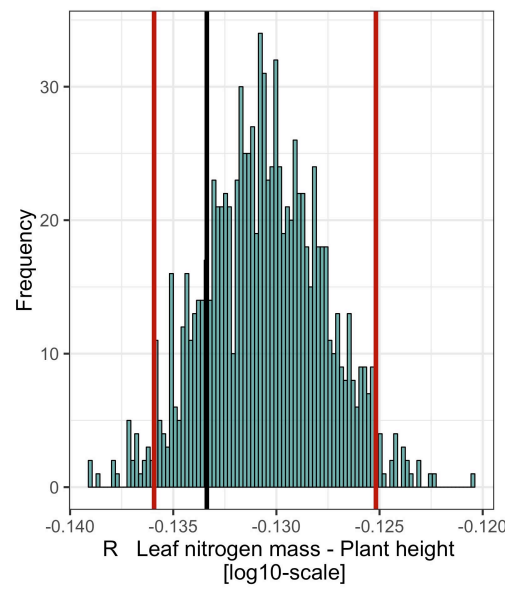
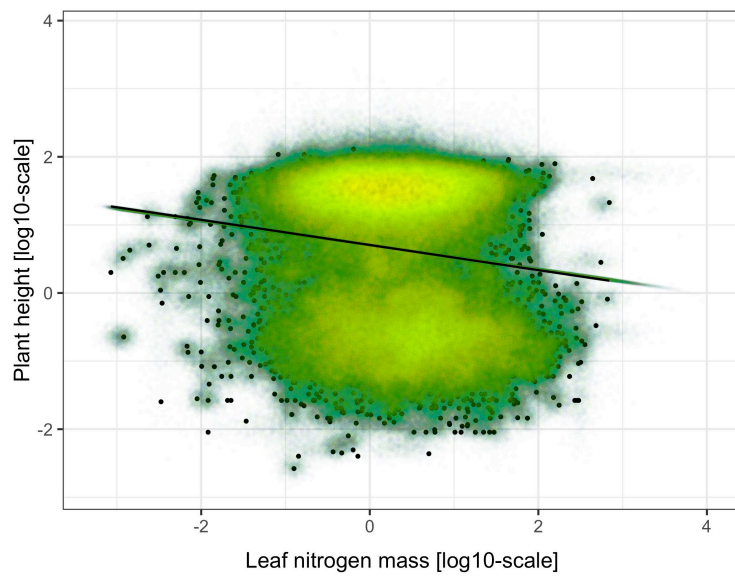
H n = 906



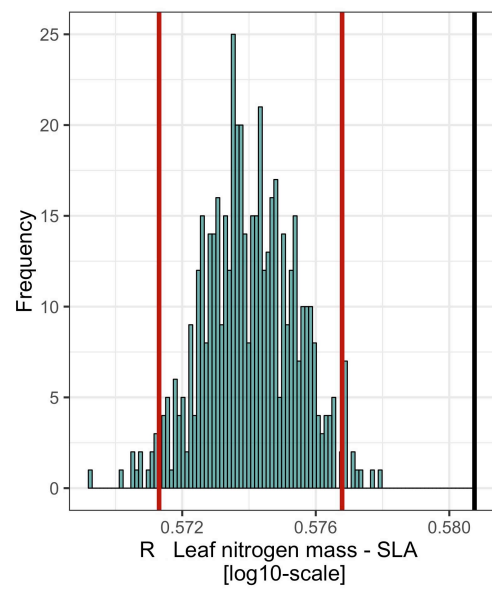
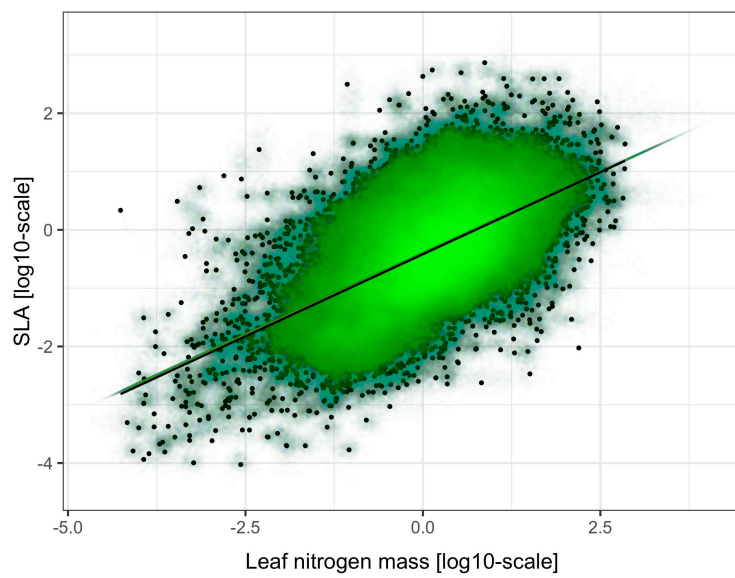
I n = 18301



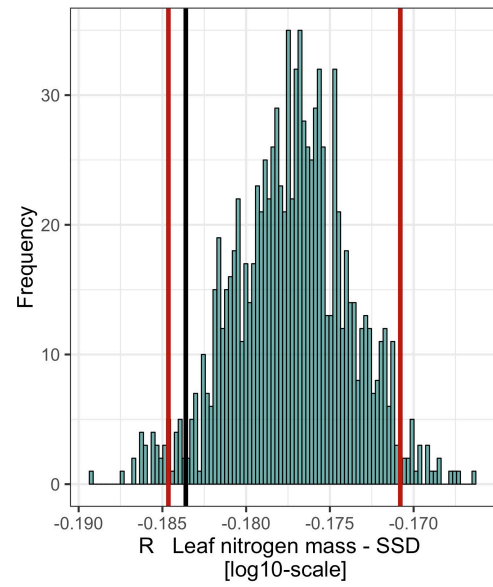
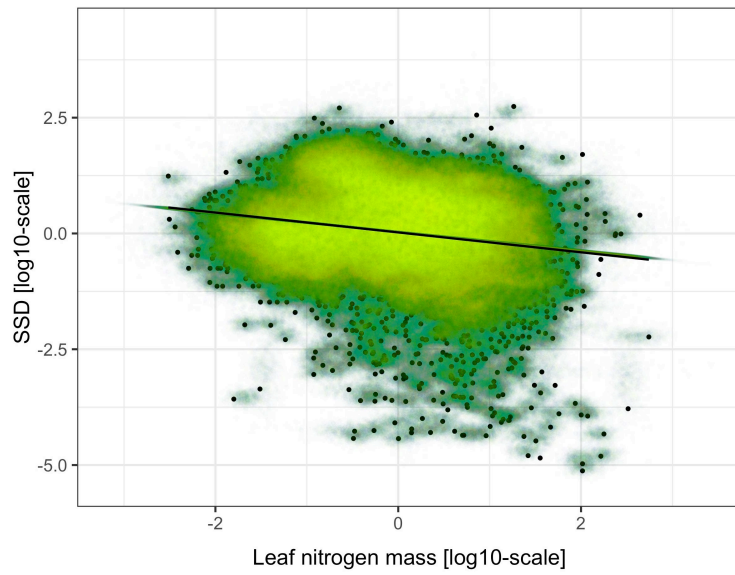
J n = 6397



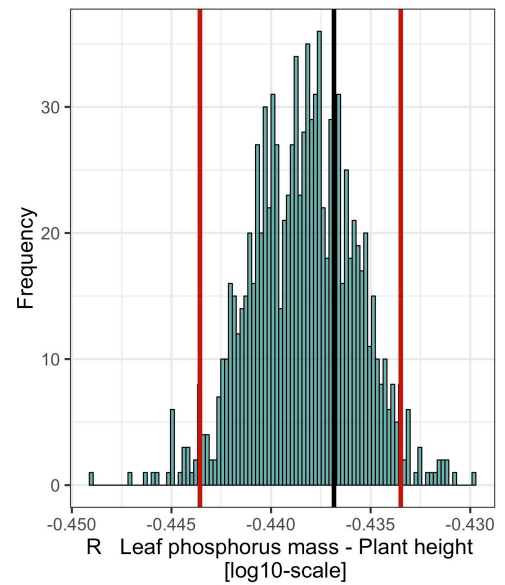
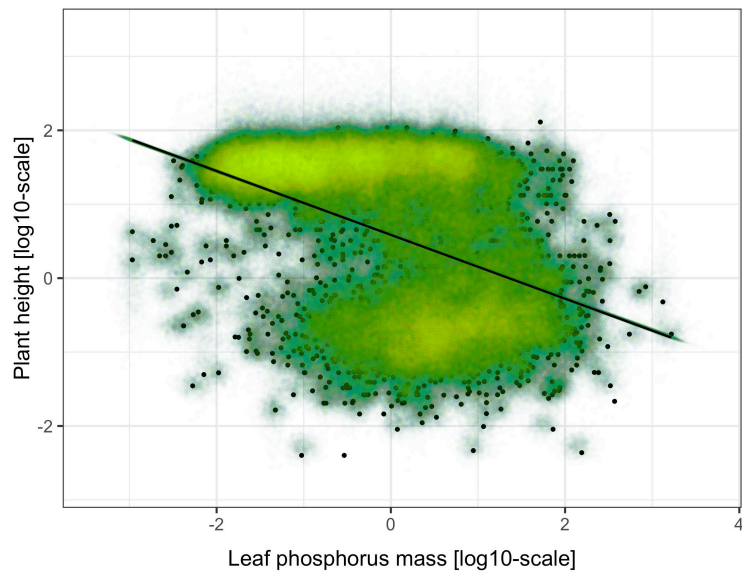
K n = 16236



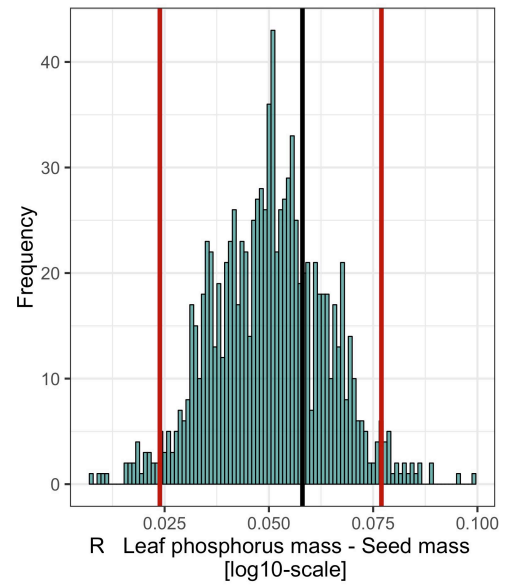
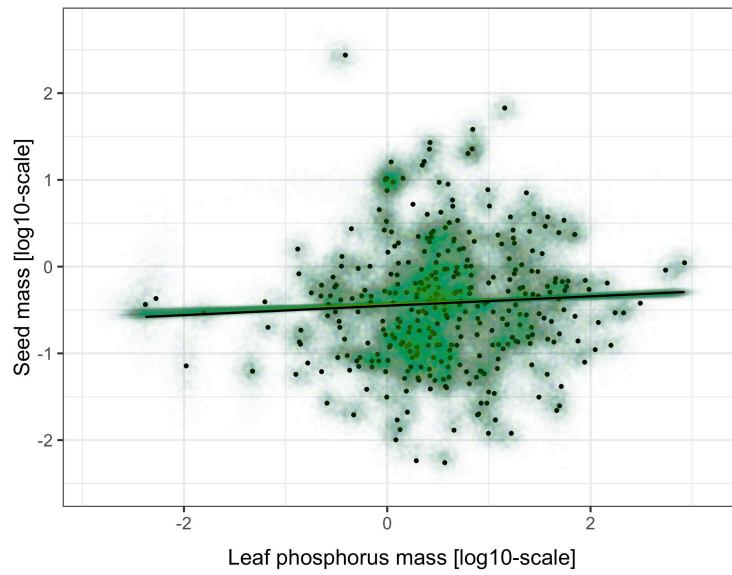
L $n = 5904$



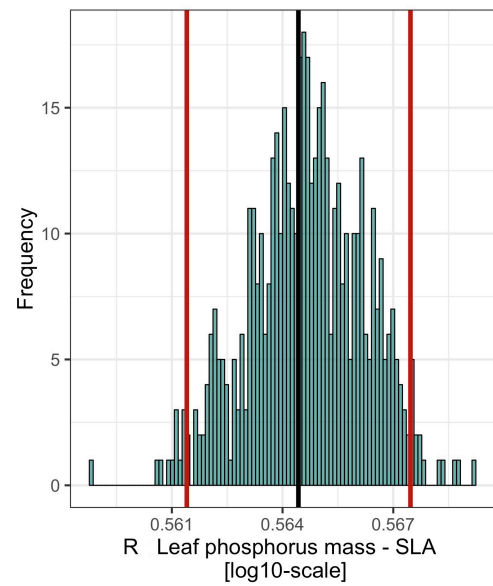
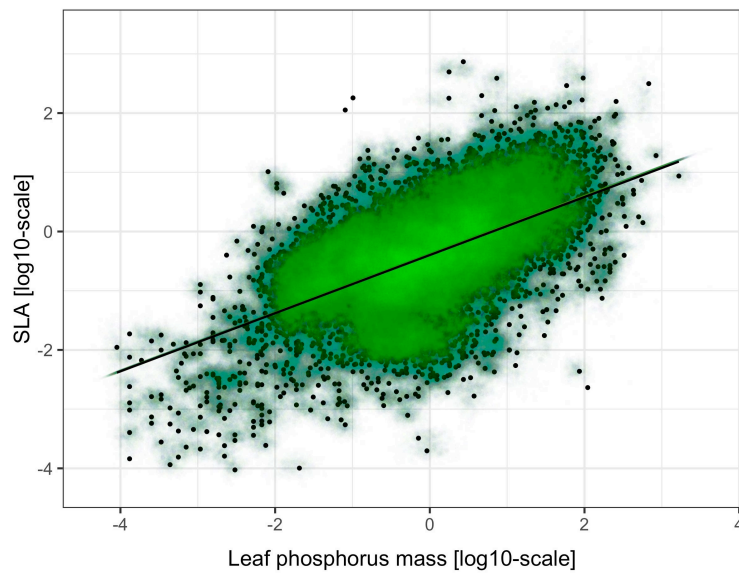
M $n = 3324$



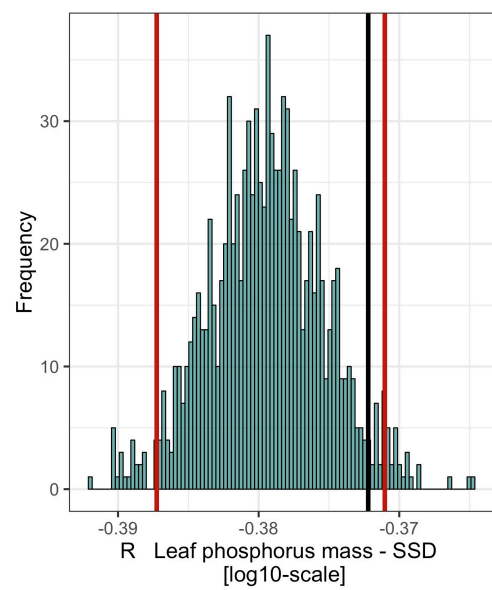
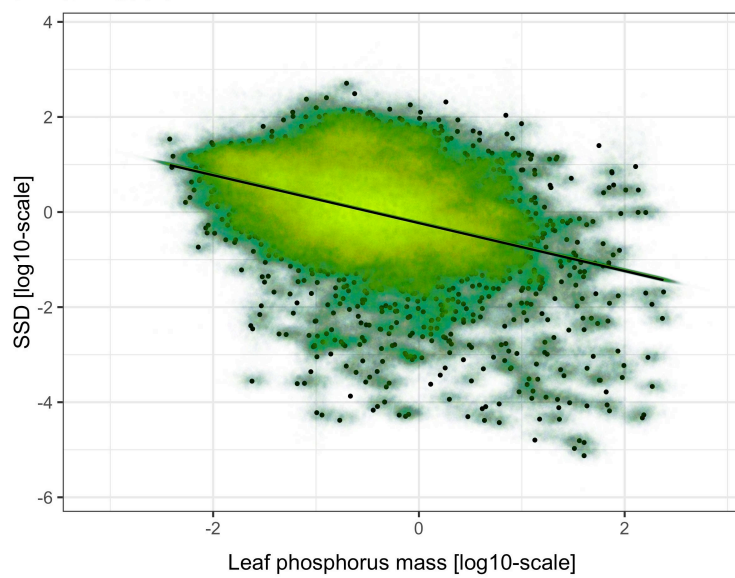
N n = 363



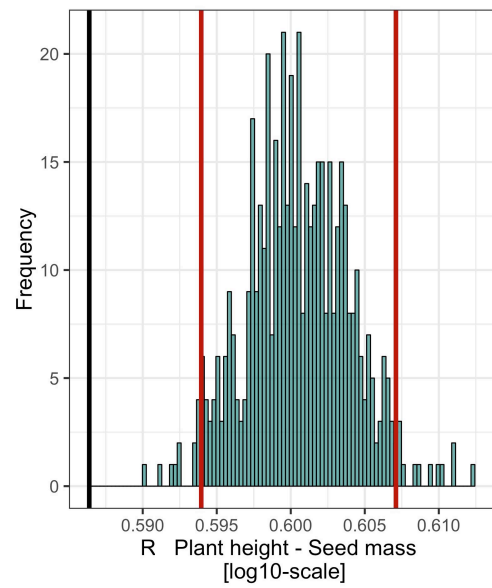
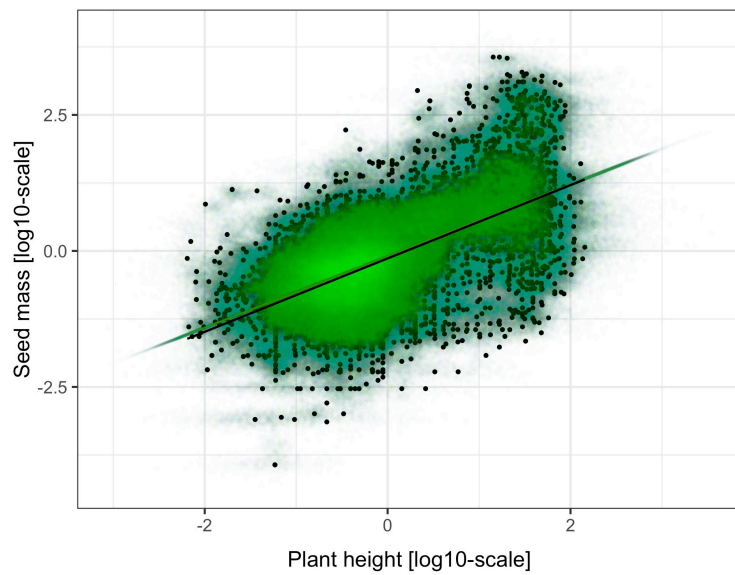
O n = 7036



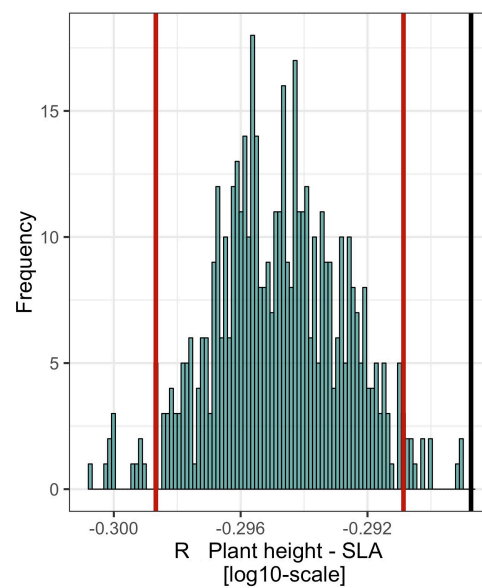
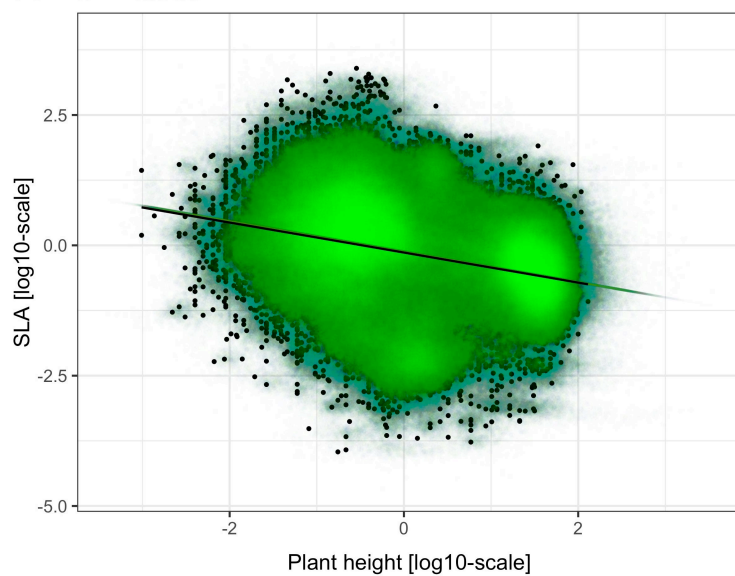
P n = 3774



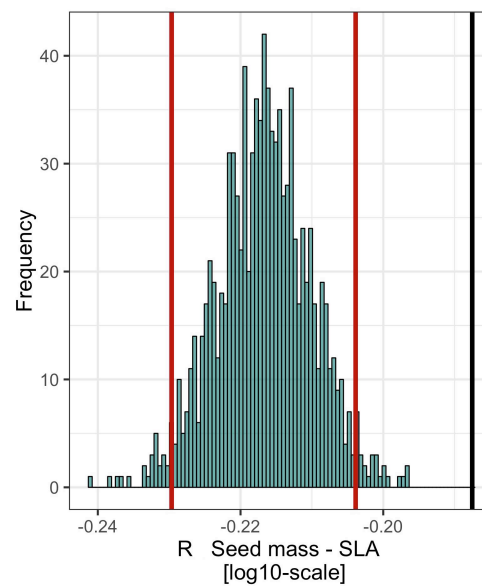
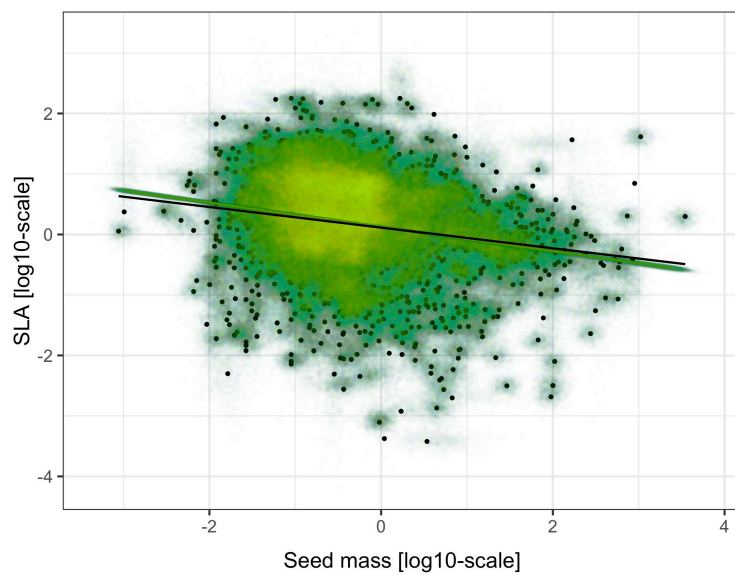
Q n = 5667



R n = 19606



S n = 1971



T n = 314

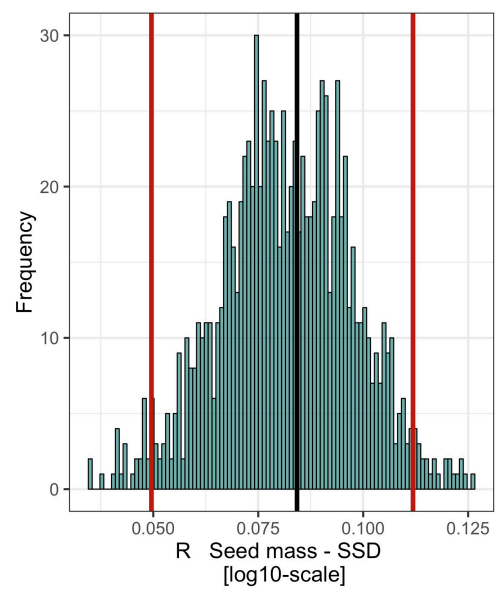
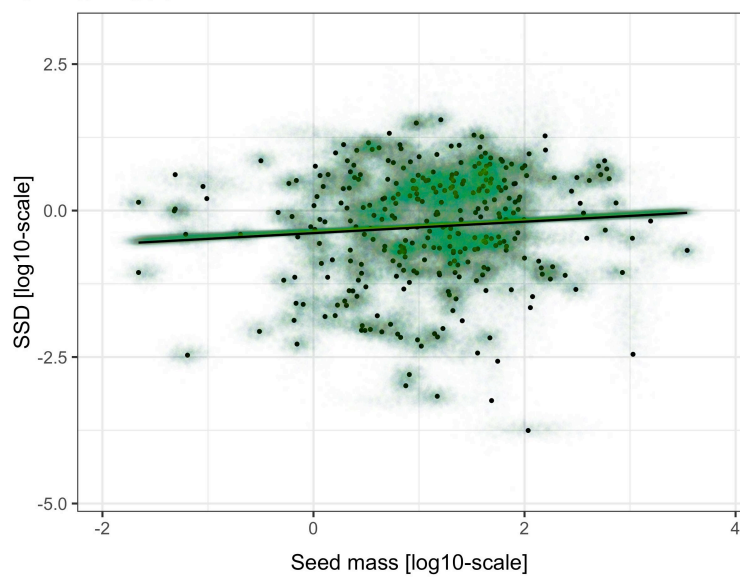


Figure S2.2 Correlation between Standard deviation and Root mean square error in dataset

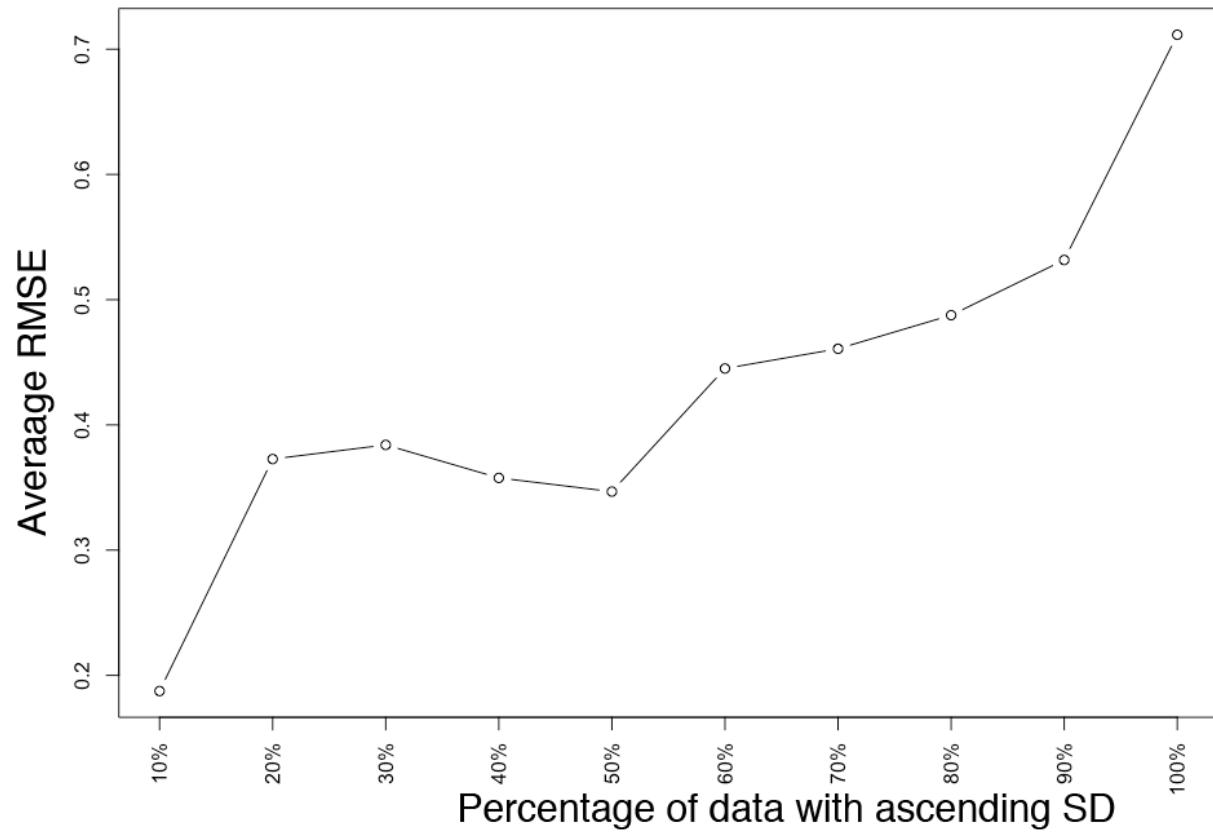


Table S2.3 Examples of studies focused on multi-organ, multi-trait datasets.

Study	Growing conditions of the plants	Spatial scale	Environmental gradients	Are environmental gradients explicitly considered?	Plant groups included	Plant groups taken into consideration?	Number of species	Life stage	Organs
Ackerly 2004	field	local	Water availability	no	woody evergreen, woody deciduous	yes	20	adults	leaf 23 stem 11 seed 2
Baraloto et al. 2010	field	regional	Precipitation, Soil nutrients	no	trees	no	668	adults	leaf 12 stem 5
Cheng et al. 2015	field	local	Temperature, Precipitation, Soil nutrients	yes	xerophytes, intermediate xerophytes, mesophytes mesophytes, mesophytes shallow vs deep-rooted	yes	55	adults	leaf 1 roots 4
de la Riva et al. 2015	field	local	Humidity (soil water)	yes	woody		38	adults	leaf 7 stem 1 roots 5 seed 1
Diaz et al. 2004	field	regional	Precipitation, Temperature, Altitude	no	eudicotyledons, monocotyledons; Asteraceae; Fabaceae; Poaceae	yes	640	adults	leaf 5 stem 4 seed 2 other 2
Diaz et al. 2016	field	global	Precipitation, Temperature, Altitude	no	woody, non-woody; angiosperms, gymnosperms, pteridophyte	yes	46085	adults	
Fortunel et al. 2012	field	regional	Precipitation, Soil nutrients	yes	trees	no	758	adults	leaf 11 stem 2 root 1

Freschet et al. 2010	field	local	Soil nutrients (C &N), Temperature (soil litter), Humidity (soil litter), growing season	yes	woody evergreen, woody deciduous, fern allies, club mosses, monocots, terrestrial forbs, aquatic forbs	yes	40	adults	leaf 8 stem 7 roots 7
Ishida 2008	field	no	trees, creeping trees, ruderal trees, climber, C3 shrubs and forbs, CAM, palm	yes	angiosperm trees, conifer trees, palm trees, fern trees	yes	32	adults	leaf 14 stem 1
Jager et al. 2015	field	local	Terrain slope, Soil nutrients	yes	yes	yes	30	> 10 dbh	leaf 8 stem 4 seed 1
Kramer et al. 2016	glasshouse	local, regional	Soil nutrients	yes	conifer, eudicot, palm, magnoliid, tree fern; AM(nodules)*, AM*, Non*, Dual AM/EM*, EM*, Ericoid*	yes	66	seedlings	leaf 2 stem 2 root 5
Li and Bao 2015	common garden	woody deciduous, woody xerophytic	no	trees, shrubs, lianas	no	no	23	1 year old	leaf 3 stem 1 roots 4
Wright 2007	field	regional	Precipitation	no	trees, shrubs, lianas	no	122	adults	leaf 2 stem 2 seed 2

When several plant group classifications were used, a semicolon divides them. The numbers next to the name of the organs included in the study in the ‘Organs’ column refers to the number of traits per organ. * Represent types of mychorrhizal association: arbuscular mychorrhizal (AM), no mychorrhizal association (Non), ectomycorrhizal (EM), Dual (association with both arbuscular and ectomycorrhizal).

Appendix S3. Trait network analyses across plant groups species and climate regions excluding leaf lifespan (LLS)

Table S3.1. Trait centrality (i.e. Degree) for all plants, woody, and non-woody species and 95% confidence interval (CI). We ran this analysis for each network of all plants, woody and non-woody species for 1000 bootstrapped trait matrices across 7 traits, excluding LLS.

Life form	Trait	2.5%	50%	97.5%
All	Plht	1.00	1.00	1.00
	SLA	1.00	1.00	1.00
	Sm	0.83	1.00	1.00
	SSD	0.83	1.00	1.00
	Leaf N	0.67	0.83	1.00
	Leaf P	0.67	0.83	0.83
	Leaf area	0.67	0.67	0.83
Non-woody	Leaf area	0.50	0.83	0.83
	Leaf N	0.67	0.83	0.83
	Leaf P	0.67	0.67	0.83
	Plht	0.67	0.67	0.83
	SLA	0.50	0.67	0.68
	Sm	0.67	0.67	0.83
	SSD	0.00	0.17	0.33
Woody	Sm	0.83	1.00	1.00
	Leaf area	0.83	0.83	0.83
	Leaf P	0.67	0.83	0.83
	SLA	0.83	0.83	0.83
	Leaf N	0.50	0.50	0.67
	Plht	0.50	0.50	0.50
	SSD	0.50	0.50	0.67

Table S3.2 Multiple-comparison adjusted 95% confidence interval (CI) from the trait centrality (i.e. degree) analyses for seven traits. We ran the trait centrality analyses for the trait of all plant species, woody and non-woody species across 1000 trait matrices. Differences in centrality between trait pairs significant when the 95% CI for the differences between trait pairs does not overlap with zero.

Trait-pairs	All plants		Woody		Non-woody	
	2.5%	97.5%	2.5%	97.5%	2.5%	97.5%
Leaf area-SLA	-0.17	0.00	-0.17	0.00	-0.17	0.33
Leaf area-Leaf N	0.17	0.33	0.17	0.33	-0.33	0.17
Leaf area-Leaf P	0.00	0.17	0.00	0.17	-0.33	0.17
Leaf area-Plht	0.33	0.33	0.33	0.33	-0.33	0.17
Leaf area-SSD	0.00	0.33	0.00	0.33	0.17	0.67
Leaf area-Sm	-0.17	0.00	-0.17	0.00	-0.33	0.17
SLA-Leaf N	0.17	0.33	0.17	0.33	-0.50	0.17
SLA-Leaf P	0.00	0.17	0.00	0.17	-0.33	0.17
SLA-Plht	0.33	0.50	0.33	0.50	-0.17	0.00
SLA-SSD	0.17	0.33	0.17	0.33	0.17	0.83
SLA-Sm	-0.17	0.00	-0.17	0.00	-0.33	0.17
Leaf N-Leaf P	-0.33	0.00	-0.33	0.00	-0.17	0.17
Leaf N-Plht	0.00	0.17	0.00	0.17	-0.17	0.33
Leaf N-SSD	0.00	0.00	0.00	0.00	0.33	0.83
Leaf N-Sm	-0.50	-0.17	-0.50	-0.17	-0.17	0.17
Leaf P-Plht	0.17	0.33	0.17	0.33	-0.17	0.33
Leaf P-SSD	0.00	0.33	0.00	0.33	0.33	0.83
Leaf P-Sm	-0.17	-0.17	-0.17	-0.17	-0.17	0.17
Plht-SSD	-0.17	0.00	-0.17	0.00	0.17	0.83
Plht-Sm	-0.50	-0.33	-0.50	-0.33	-0.17	0.17
SSD-Sm	-0.50	-0.17	-0.50	-0.17	-0.83	-0.33

Table S3.3 Edge density for woody and non-woody species and 95% confidence interval (CI). We ran this analysis for each network of woody and non-woody species across climate regions for 1000 bootstrapped trait matrices.

Life-form	Climate	2.5%	50%	97.5%
Woody	Arid	0.52	0.57	0.67
	Cold	0.43	0.52	0.57
	Polar	0.29	0.38	0.48
	Temperate	0.52	0.57	0.62
	Tropical	0.62	0.67	0.71
Non-woody	Arid	0.38	0.48	0.57
	Cold	0.52	0.62	0.67
	Polar	0.33	0.43	0.52
	Temperate	0.52	0.62	0.71
	Tropical	0.29	0.38	0.52

Table S3.4 Multiple-comparison adjusted 95% confidence interval (CI) from the edge density analyses among climate regions for woody and non-woody species. We ran the edge density comparison for woody and non-woody species across five climate regions for 1000 trait matrices. Differences in edge density between trait pairs significant when the 95% CI for the differences between trait pairs does not overlap with zero.

Region-pairs	Woody		Non-woody	
	2.5%	97.5%	2.5%	97.5%
Arid-Cold	-0.05	0.19	-0.29	0.00
Arid-Polar	0.05	0.38	-0.19	0.19
Arid-Temperate	-0.14	0.14	-0.33	0.05
Arid-Tropical	-0.19	0.05	-0.14	0.24
Cold-Polar	0.00	0.29	0.00	0.33
Cold-Temperate	-0.19	0.05	-0.19	0.14
Cold-Tropical	-0.24	-0.05	0.00	0.38
Polar-Temperate	-0.38	-0.05	-0.33	0.05
Polar-Tropical	-0.43	-0.14	-0.19	0.24
Temperate-Tropical	-0.19	0.05	0.00	0.38

Table S3.5 Modularity for woody and non-woody species and 95% confidence interval (CI). We ran this analysis for each network of woody and non-woody species across climate regions for 1000 bootstrapped trait matrices.

Life-form	Climate	2.5%	50%	97.5%
Woody	Arid	0.06	0.15	0.22
	Cold	0.15	0.21	0.28
	Polar	0.17	0.27	0.37
	Temperate	0.10	0.15	0.21
	Tropical	0.02	0.06	0.10
Non-woody	Arid	0.10	0.20	0.31
	Cold	0.02	0.05	0.13
	Polar	0.13	0.20	0.32
	Temperate	0.02	0.05	0.13
	Tropical	0.09	0.22	0.32

Table S3.6 Multiple-comparison adjusted 95% confidence interval (CI) from the modularity analyses among climate regions for woody and non-woody species. We ran the modularity comparison for woody and non-woody species across five climate regions for 1000 trait matrices. Differences in modularity between regions is significant when the 95% CI for the differences between regions pairs does not overlap with zero.

Region pairs	Woody		Non-woody	
	2.5%	97.5%	2.5%	97.5%
Arid-Cold	-0.21	0.11	-0.02	0.33
Arid-Polar	-0.31	0.08	-0.24	0.22
Arid-Temperate	-0.15	0.16	0.00	0.34
Arid-Tropical	-0.04	0.20	-0.22	0.23
Cold-Polar	-0.23	0.13	-0.33	0.05
Cold-Temperate	-0.06	0.18	-0.11	0.16
Cold-Tropical	0.00	0.26	-0.30	0.06
Polar-Temperate	-0.06	0.30	0.00	0.34
Polar-Tropical	0.03	0.35	-0.19	0.21
Temperate-Tropical	-0.04	0.19	-0.32	0.00

Table S3.7. Trait centrality (i.e. Degree) for woody species across five climate regions and 95% confidence interval (CI). We ran this analysis over 1000 bootstrapped trait matrices.

Climate	Trait	2.50%	50%	97.50%
Tropical	Sm	0.83	1.00	1.00
	SLA	0.83	0.83	0.83
	Leaf area	0.67	0.67	0.67
	Leaf P	0.50	0.67	0.67
	Leaf N	0.50	0.50	0.67
	Plht	0.50	0.50	0.67
	SSD	0.50	0.50	0.67
Temperate	Leaf area	0.67	0.67	0.67
	SLA	0.67	0.67	0.83
	Sm	0.50	0.67	0.67
	Leaf N	0.33	0.50	0.50
	Leaf P	0.50	0.50	0.67
	Plht	0.50	0.50	0.67
	SSD	0.50	0.50	0.67
Arid	Leaf area	0.67	0.67	0.83
	SLA	0.67	0.67	1.00
	SSD	0.50	0.67	0.83
	Leaf P	0.50	0.50	0.67
	Plht	0.33	0.50	0.67
	Sm	0.50	0.50	0.67
	Leaf N	0.33	0.33	0.50
Cold	Plht	0.33	0.67	0.83
	SLA	0.50	0.67	0.67
	Leaf area	0.33	0.50	0.50
	Leaf P	0.50	0.50	0.67
	Sm	0.50	0.50	0.50
	SSD	0.33	0.50	0.50
	Leaf N	0.33	0.33	0.33
Polar	Leaf area	0.17	0.50	0.67
	Leaf P	0.17	0.50	0.67
	SLA	0.33	0.50	0.67
	Sm	0.33	0.50	0.50
	Leaf N	0.33	0.33	0.50
	Plht	0.17	0.33	0.50
	SSD	0.17	0.33	0.34

Table S3.8. Trait centrality (i.e. Degree) for non-woody species across five climate regions and 95% confidence interval (CI). We ran this analysis over 1000 bootstrapped trait matrices.

Climate	Trait	2.50%	50%	97.50%
Tropical	Leaf area	0.33	0.50	0.67
	Leaf N	0.50	0.50	0.67
	Sm	0.33	0.50	0.83
	Leaf P	0.17	0.33	0.67
	Plht	0.17	0.33	0.50
	SLA	0.17	0.33	0.50
	SSD	0.00	0.17	0.50
Temperate	Leaf N	0.67	0.83	1.00
	Leaf area	0.50	0.67	0.83
	Leaf P	0.33	0.67	0.83
	Plht	0.50	0.67	0.83
	Sm	0.67	0.67	0.83
	SLA	0.50	0.50	0.50
	SSD	0.00	0.17	0.33
Arid	Leaf area	0.50	0.50	0.67
	Leaf N	0.33	0.50	0.67
	SLA	0.33	0.50	0.67
	Sm	0.33	0.50	0.83
	Leaf P	0.33	0.33	0.67
	Plht	0.17	0.33	0.50
	SSD	0.17	0.17	0.33
Cold	Leaf area	0.67	0.83	0.83
	Leaf N	0.50	0.67	0.83
	SLA	0.50	0.67	0.83
	Sm	0.50	0.67	0.67
	Leaf P	0.50	0.50	0.67
	Plht	0.17	0.50	0.67
	SSD	0.17	0.33	0.50
Polar	Sm	0.50	0.67	0.83
	Leaf area	0.33	0.50	0.50
	Leaf N	0.50	0.50	0.67
	Leaf P	0.33	0.50	0.67
	SLA	0.33	0.50	0.50
	Plht	0.33	0.33	0.50
	SSD	0.00	0.17	0.33

Table S3.9 Multiple-comparison adjusted 95% confidence interval (CI) from the trait centrality (i.e. degree) analyses for seven traits of woody species across five climate regions.

We ran the trait centrality analyses for the trait of woody species across five climate regions across 1000 trait matrices. Differences in centrality between trait pairs significant when the 95% CI for the differences between trait pairs does not overlap with zero.

Trait-pairs	Tropical		Temperate		Arid		Cold		Polar	
	2.5%	97.5%	2.5%	97.5%	2.5%	97.5%	2.5%	97.5%	2.5%	97.5%
Leaf area-SLA	-0.33	-0.17	-0.17	0.17	-0.33	0.33	-0.33	0.17	-0.50	0.33
Leaf area-Leaf N	0.00	0.17	0.17	0.33	0.17	0.50	0.00	0.33	-0.17	0.33
Leaf area-Leaf P	-0.17	0.17	0.00	0.33	0.00	0.17	-0.33	0.17	-0.50	0.33
Leaf area-Plht	0.00	0.17	0.00	0.33	0.00	0.50	-0.33	0.17	-0.17	0.50
Leaf area-SSD	-0.17	0.17	-0.17	0.33	-0.17	0.33	-0.17	0.17	-0.17	0.50
Leaf area-Sm	-0.33	-0.17	0.00	0.33	-0.17	0.33	-0.17	0.17	-0.33	0.33
SLA-Leaf N	0.17	0.50	0.17	0.50	0.00	0.67	0.00	0.50	-0.17	0.33
SLA-Leaf P	0.00	0.50	0.00	0.33	-0.17	0.50	-0.17	0.33	-0.33	0.33
SLA-Plht	0.17	0.50	0.00	0.33	0.00	0.50	-0.17	0.17	-0.17	0.50
SLA-SSD	0.17	0.33	0.00	0.33	-0.17	0.33	0.00	0.33	-0.17	0.50
SLA-Sm	-0.17	0.17	0.00	0.33	-0.17	0.33	0.00	0.33	-0.17	0.33
Leaf N-Leaf P	-0.33	0.17	-0.33	0.17	-0.50	0.00	-0.33	0.00	-0.33	0.33
Leaf N-Plht	-0.17	0.17	-0.33	0.17	-0.33	0.17	-0.50	0.00	-0.17	0.33
Leaf N-SSD	-0.17	0.00	-0.33	0.00	-0.50	0.00	-0.33	0.00	-0.17	0.33
Leaf N-Sm	-0.50	-0.17	-0.33	0.00	-0.33	-0.17	-0.33	0.00	-0.33	0.17
Leaf P-Plht	0.00	0.17	-0.17	0.17	-0.17	0.33	-0.33	0.17	-0.33	0.50
Leaf P-SSD	-0.33	0.33	-0.33	0.17	-0.33	0.17	-0.17	0.33	-0.17	0.50
Leaf P-Sm	-0.33	-0.17	-0.17	0.17	-0.33	0.33	-0.17	0.17	-0.33	0.50
Plht-SSD	-0.33	0.17	-0.17	0.17	-0.33	0.17	-0.17	0.50	-0.33	0.50
Plht-Sm	-0.50	-0.17	-0.33	0.17	-0.50	0.17	-0.17	0.33	-0.50	0.17
SSD-Sm	-0.50	0.00	-0.17	0.33	-0.17	0.33	-0.17	0.17	-0.50	0.17

Table S3.10. Multiple-comparison adjusted 95% confidence interval (CI) from the trait centrality (i.e. degree) analyses for seven traits of non-woody species across five climate regions. We ran the trait centrality analyses for the trait of non-woody species across five climate regions across 1000 trait matrices. Differences in centrality between trait pairs significant when the 95% CI for the differences between trait pairs does not overlap with zero.

Trait-pairs	Tropical		Temperate		Arid		Cold		Polar	
	2.5%	99.9%	2.5%	97.5%	2.5%	97.5%	2.5%	97.5%	2.5%	97.5%
Leaf area-SLA	-0.17	0.50	-0.17	0.33	-0.17	0.33	0.00	0.17	-0.17	0.33
Leaf area-Leaf N	-0.17	0.17	-0.50	0.00	-0.33	0.33	-0.17	0.33	-0.33	0.17
Leaf area-Leaf P	-0.33	0.50	-0.33	0.33	-0.17	0.33	0.00	0.33	-0.17	0.17
Leaf area-Plht	-0.17	0.50	-0.33	0.33	0.00	0.50	0.00	0.67	-0.17	0.33
Leaf area-SSD	0.00	0.50	0.17	0.67	0.17	0.67	0.17	0.67	0.00	0.67
Leaf area-Sm	-0.50	0.50	-0.33	0.17	-0.33	0.50	0.00	0.33	-0.50	0.17
SLA-Leaf N	-0.50	0.17	-0.50	0.00	-0.33	0.50	-0.33	0.33	-0.33	0.17
SLA-Leaf P	-0.50	0.33	-0.33	0.17	-0.17	0.50	-0.17	0.33	-0.33	0.17
SLA-Plht	-0.17	0.33	-0.33	0.00	-0.17	0.67	-0.17	0.50	-0.17	0.33
SLA-SSD	-0.33	0.50	0.17	0.50	0.17	0.50	0.17	0.50	0.00	0.50
SLA-Sm	-0.50	0.17	-0.33	-0.17	-0.33	0.50	-0.17	0.33	-0.50	0.00
Leaf N-Leaf P	-0.33	0.50	-0.17	0.67	-0.17	0.50	-0.17	0.33	-0.17	0.33
Leaf N-Plht	0.00	0.50	-0.17	0.50	-0.17	0.50	0.00	0.50	0.00	0.33
Leaf N-SSD	0.00	0.67	0.33	0.83	0.00	0.50	0.17	0.50	0.00	0.67
Leaf N-Sm	-0.33	0.33	-0.17	0.33	-0.33	0.33	-0.17	0.33	-0.33	0.33
Leaf P-Plht	-0.17	0.50	-0.33	0.33	-0.17	0.33	0.00	0.33	0.00	0.50
Leaf P-SSD	-0.17	0.67	0.00	0.83	0.00	0.50	0.00	0.50	0.17	0.50
Leaf P-Sm	-0.50	0.33	-0.33	0.00	-0.50	0.33	-0.33	0.17	-0.50	0.17
Plht-SSD	-0.33	0.50	0.17	0.67	-0.17	0.50	-0.17	0.50	-0.17	0.50
Plht-Sm	-0.67	0.33	-0.33	0.17	-0.50	0.17	-0.33	0.00	-0.50	0.00
SSD-Sm	-0.67	0.00	-0.83	-0.33	-0.67	0.17	-0.50	0.00	-0.67	-0.17

Appendix S4 Conditional dependency and its determination through the calculation of precision matrix and its confidence interval.

Conditional Independency

Two random variables X and Y are independent if the occurrence of one variable does not affect the probability of the other or formally if their joint distribution equals the product of their probabilities:

$$p(X, Y) = p(X)p(Y)$$

Similarly, we say two random variables X and Y are conditionally independent given a third variable Z if the knowledge of one variable does not provide any information on the likelihood of the other after observing variable Z , or formally if their conditional joint distribution equals the product of their conditional probabilities:

$$p(X, Y | Z) = p(X|Z)p(Y|Z)$$

and the conditional mutual information of X and Y given Z , $I(X, Y|Z)$, is zero (i.e. the “amount of information” obtained about X through Y after observing Z is zero):

$$\begin{aligned} I(X, Y | Z) &= \int_{z \in Z} p(Z = z) \iint_{x \in X, y \in Y} p(X = x, Y = y) \log \frac{p(X = x, Y = y | Z = z)}{p(X = x | Z = z)p(Y = y | Z = z)} \\ &= \int_{z \in Z} p(Z = z) \iint_{x \in X, y \in Y} p(X = x, Y = y) \log \frac{p(X = x | Z = z)p(Y = y | Z = z)}{p(X = x | Z = z)p(Y = y | Z = z)} \\ &= 0 \end{aligned}$$

Intuitively, if X and Y are conditionally independent given Z , then any relationship that may exist between X and Y can be explained by Z . In other words, X and Y may appear to be correlated if Z is not considered, but if Z is controlled by holding it constant then any relationship between X and Y would disappear.

Probabilistic Graphical Models

A graph $G = (V, E)$ is a data structure consisting of a set of nodes $V = \{X_1, X_2, \dots, X_p\}$ and a set of edges E that connect pairs of nodes. A pair of nodes X_i and X_j can be connected by either a directed edge $X_i \rightarrow X_j$ or an undirected edge $X_i - X_j$. If all edges of the graph are directed edges, we say that the graph is directed. If all edges of graph are undirected edges, we

say that the graph is undirected.

Graphs are an intuitive way of representing and visualizing the relationships between random variables, where nodes correspond to random variables, and edge represents dependency or relationship between variables. For example, correlation graphs (also called correlation networks) represent the pairwise correlation between variables. In a correlation graph, a pair of nodes X_i and X_j are connected if the pairwise correlation of the data associated with X_i and X_j is beyond a threshold.

Probabilistic graphical models (PGMs) use a graph-based representation as the basis for encoding a complex distribution over a high-dimensional space. In this representation, the edges correspond to direct probabilistic interactions between nodes. The graph is a compact representation of a set of conditional independencies that hold in the distribution. In particular, if there is no edge between two nodes X_i and X_j , then the random variable X_i is conditionally independent of the random variable X_j given all other variables. Two approaches for graphically representing the distributions are commonly used: (i) *Bayesian networks* that uses a directed graph (where the edges have a source and a target) and (ii) *Markov networks* which uses an undirected graph representation.

Precision Matrix

One special example of PGMs is Gaussian graphical models which gives an efficient and compact representation for Gaussian multivariate data. In particular, let $\mathbf{X} = \{X_1, X_2, \dots, X_p\}$ be a set of random variables with a multivariate Gaussian design ensemble (i.e., $\mathbf{X} \sim \mathcal{N}(\mathbf{0}, \Sigma)$; for simplicity and without loss of generality, here we assume that the mean of $\mathbf{X}$ is zero). The precision matrix (the inverse of covariance matrix; $\Omega = \Sigma^{-1}$) reveals the statistical conditional independence structure among features i.e., if $\Omega_{ij} = 0$, then feature X_i is conditionally independent of X_j given all other variables and there is no edge in the undirected graph structure associated with $\mathbf{X}$ (Figure 1).

To illustrate the relationship between the precision matrix and the graph structure, we consider the probabilistic graphical models as a skeleton for compactly representing a high-

dimensional distribution P . We “break” the distribution into smaller factors, each over a much smaller space of possibilities. Then, we can define the overall joint distribution as a product of these factors. The graph structure defines the factorization of a distribution P associated with it — the set of factors and the variables that they encompass.

In the Gaussian graphical models, the set of factors can be represented in terms of elements of the precision matrix Ω . Mainly, the density function of $\mathbf{X}$, is written as

$$\begin{aligned}
 f(\mathbf{X}) &= \frac{1}{\sqrt{(2\pi)^p |\Sigma|}} \exp \left\{ -\frac{1}{2} \mathbf{X}^T \Sigma^{-1} \mathbf{X} \right\} \\
 &= \frac{1}{\sqrt{(2\pi)^p |\Sigma|}} \exp \left\{ -\frac{1}{2} \mathbf{X}^T \Omega \mathbf{X} \right\} \\
 &= \frac{1}{\sqrt{(2\pi)^p |\Sigma|}} \exp \left\{ -\frac{1}{2} \sum_{i,j=1}^p X_i X_j \Omega_{ij} \right\} \\
 &= \frac{1}{\sqrt{(2\pi)^p |\Sigma|}} \prod_{i,j=1}^p \exp \left\{ -\frac{1}{2} X_i X_j \Omega_{ij} \right\} \\
 &= \frac{1}{\sqrt{(2\pi)^p |\Sigma|}} \prod_{i=1}^p \psi_i(X_i) \prod_{(i,j) \in E} \psi_{ij}(X_i, X_j)
 \end{aligned}$$

where $|\Sigma|$ is determinant of Σ , the second equality uses the fact that $\Omega = \Sigma^{-1}$, and the third equality is the expansion of matrix multiplication, $\psi_i(X_i) = \exp \left\{ -\frac{1}{2} X_i^2 \Omega_{ii} \right\}$ can be viewed as the node potential and $\psi_{ij}(X_i, X_j) = \exp \left\{ -\frac{1}{2} X_i X_j \Omega_{ij} \right\}$ can be viewed as the edge potential. As shown in the above equation, the density function of $\mathbf{X}$ is factorized to a set factors written as $\exp \left\{ -\frac{1}{2} X_i X_j \Omega_{ij} \right\}$. If $\Omega_{ij} = 0$, then the factor $\exp \left\{ -\frac{1}{2} X_i X_j \Omega_{ij} \right\} = 1$ with no effect in the density of $\mathbf{X}$. In other words, if the edge potential $f_{ij}(X_i, X_j) = 0$, and there is no edge between nodes X_i and X_j in the graph structure, then we say that the feature X_i is conditionally independent of X_j given all other variables.

Interestingly, the elements of a precision matrix are related to the partial correlation. In particular, the partial correlation coefficient between variables X_i and X_j measures their conditional correlation after removing the effect of the variables $X_{-ij} = X - \{X_i, X_j\}$. The partial correlation values can be calculated by normalizing the off-diagonal entries of the precision matrix (Lauritzen 1996) as

$$r_{ij} = \frac{\text{cov}(X_i, X_j | X_{-ij})}{\sqrt{\text{var}(X_i | X_{-ij}) \text{var}(X_j | X_{-ij})}} = - \frac{\Omega_{ij}}{\sqrt{\Omega_{ii} \Omega_{jj}}}$$

Why are we not using sample covariance matrix and instead using Precision matrix?

Consider an independent and identically distributed random sample $\{\mathbf{X}_1, \mathbf{X}_2, \dots, \mathbf{X}_n\}$ from the distribution of $\mathbf{X}$. In the classical situation where number samples $n \rightarrow \infty$, the sample covariance matrix $\hat{\Sigma}_n = \sum_{i=1}^n \mathbf{X}_i \mathbf{X}_i^T$ is a consistent and well behaved estimator of $\Sigma = E[\mathbf{X}_i \mathbf{X}_i^T]$ and $\hat{\Omega}_n = \hat{\Sigma}_n^{-1}$ is a natural and good estimator of Ω . However, if the number of samples is not sufficient, then $\hat{\Sigma}_n$ is no longer a consistent estimate of Σ and in some cases (e.g., $p > n$), $\hat{\Sigma}_n$ is not invertible, thus $\hat{\Omega}_n$ is not well-defined. Unlike the covariance matrix, the precision matrix is usually sparse (it contains many zero values). This characterization yields to several efficient and consistent algorithms for estimating Ω . Thus, in this paper, we focus on estimating Ω directly.

Difference between Correlation and Dependency Graphs

Correlation graphs capture the similarity between variables. Two variables are similar if their pairwise correlation exceeds a threshold. However, the similarity between two variables may come from a third source (e.g., another variable). Thus, the correlation graph captures indirect similarities. The covariance matrix essentially indicates the correlation between variables; if an element in the covariance matrix is zero, that means that two variables are not correlated or marginally independent without observing other variables. However, this kind of independency is rarely expected to happen in real world applications, because it reveals that two random variables have no active relationship even without observing other nodes.

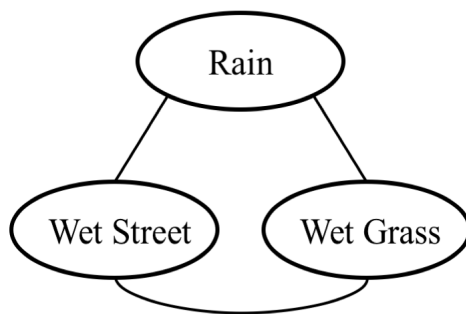
Precision matrices are related to the partial correlations, as they can calculate the correlation between two variables after removing the effect of other variables. Thus, the precision matrix provides information about the direct similarity between two variables. In particular, two variables that are connected in correlation graphs, may or may not be connected in a precision graph and vice versa. Below, we provide some examples to illustrate the difference between correlation graph and precision graph.

a. Example 1

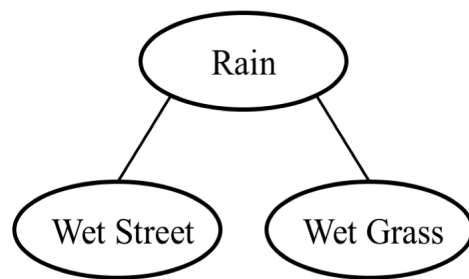
Consider three binary variables:

- Rain? (yes/no) i.e., set to yes if it was rainy today.
- Wet Grass? (yes/no) i.e., set to yes if it the grass is wet.
- Wet Street? (yes/no) i.e., set to yes if it the street is wet.

When it is raining there is a high chance that both the grass and the street get wet, yielding a high correlation between Rain-Wet Street, and Rain-Wet Grass. At the same time, if we calculate the correlation between Wet Grass -Wet Street, they will likely be highly correlated, however, neither Wet grass, nor Wet street is causing the occurrence of the other variable (Figure 1 a). Rain is the main reason that Wet Street and Wet Grass are correlated, and if we remove the effect of Rain from both variables, they will no longer be correlated. As a result, Wet Street and Wet Grass are not connected in the dependency graph i.e., they are conditionally independent given Rain (Figure 1 b).



a. Correlation Graph



b. Dependency Graph

Figure 1. Rain Example. Wet Street and Wet Grass are connected in the correlation graph (highly correlated) but are independent given Rain in the dependency graph.

b. Example 2

Consider three variables:

- Lung Cancer? (yes/no) i.e., set to yes if the person has lung cancer.
- Age i.e., indicates the age of a person.
- Smoking? (yes/no) i.e., set to yes if the person is smoking.

Whether a person gets lung cancer depends on several factors such as age and whether the person is smoking or not. As a result, in the correlation graph there is an edge between Lung Cancer-Age and Lung Cancer-Smoking. However, for the population considered in this example, Age and Smoking are not correlated i.e., the age of a person does not significantly impact whether he/she is smoking (Figure 2a). On the other hand, if a person with lung cancer diagnosis is young, this raises the chances that the person is smoking, as lung cancer patients often have at least one of the two major risk factors, increased age or are smokers. As a result, there is an edge between Age-Smoking in the dependency graph (Figure 2b).

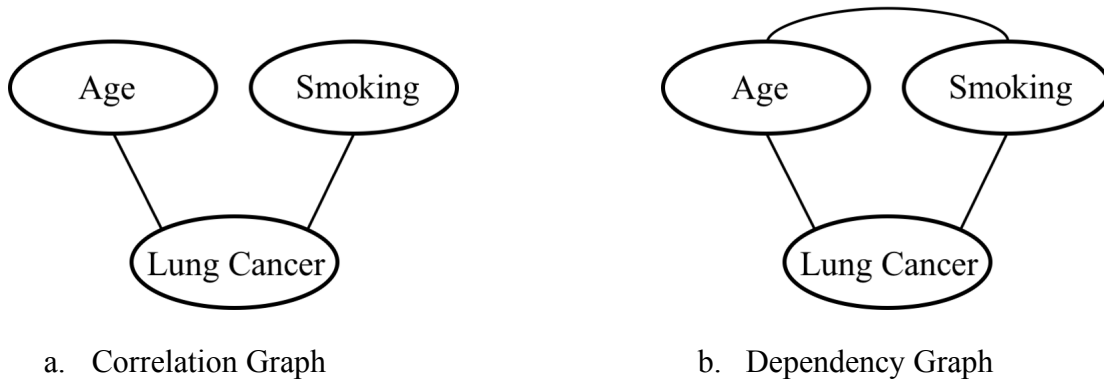


Figure 2. Lung Cancer Example. Age and Smoking are not connected in the correlation graph (not correlated) but are dependent given Lung Cancer in the dependency graph.

Analysis

We estimated the precision matrix for each plant group (see methods section) separately. To estimate the precision matrix the glasso algorithm assumes a multi-variate Gaussian

distribution and estimates the precision matrix by minimizing the negative log-likelihood among all measurements (in this case traits).

The log-likelihood of a Gaussian multivariate distribution is:

$$\begin{aligned}\log f(x) &= \log \left(\frac{1}{\sqrt{(2\pi)^p |\Sigma|}} \exp \left\{ -\frac{1}{2} \mathbf{X}^T \Omega \mathbf{X} \right\} \right) \\ &= -\log \left(\sqrt{(2\pi)^p} \right) - \frac{1}{2} \log |\Sigma| - \frac{1}{2} \text{Tr}(\mathbf{X}^T \Omega \mathbf{X}) \\ &= -\log \left(\sqrt{(2\pi)^p} \right) + \frac{1}{2} \log |\Omega| - \frac{1}{2} \text{Tr}(\hat{\Sigma} \Omega)\end{aligned}$$

where the last equality is from the determinant property: $|\Sigma| = \frac{1}{|\Sigma^{-1}|} = \frac{1}{|\Omega|}$. Thus, ignoring the constants, glasso finds the precision matrix Ω , by solving the following optimization problem:

$$\arg \min_{\Omega} -\log |\Omega| + \text{Tr}(\hat{\Sigma} \Omega) + \lambda \|\Omega\|_1$$

where $\|\Omega\|_1$ is the L_1 norm of Ω .

Selecting penalty parameter λ

Since the precision matrix is usually sparse (i.e. most of the precision matrix elements are zero), glasso adds a penalty term (L_1 norm) to the objective function which encourages to have a sparse precision matrix. The glasso algorithm includes a penalty parameter λ , which controls the sparsity level of the precision matrix. If $\lambda = 0$, then all of the elements of the precision matrix are non-zero. In contrast, with a very large value of λ , all the elements become zero. Following Jankova and van de Geer (2015), we accounted for differences in sample size across precision matrices and their effect on λ as follows: $\lambda = 2 \sqrt{(\log p / n)}$, where n refers to the sample size and p denotes the number of variables, in this case traits (Appendix S4).

Significance of trait-trait connections in the trait network

Given a data sample matrix $X \in \mathbb{R}^{n \times p}$ and estimated precision matrix $\Omega \in \mathbb{R}^{p \times p}$, the test statistic is defined as $T = \Omega + \lambda \Omega Z \Omega$ where $Z \in \mathbb{R}^{p \times p}$ is the sub-gradient of norm $\|\Omega\|_1$, and its variance $\sigma_{ij}^2 = \Omega_{ii} \Omega_{jj} + \Omega_{ij}^2$. Then under this scenario, thresholding T_{ij} at level $\Phi^{-1}(1 - \frac{\alpha}{p(p-1)}) \frac{\hat{\sigma}_{ij}}{\sqrt{n}}$ for all i, j will remove all zero entries (i.e., non-significant trait-trait interactions) with probability $1 - \alpha$ asymptotically (Jankova & van de Geer, 2015).

Specifically, considering the data for a particular group represented in a matrix $X \in \Re^{n \times p}$ which contains the trait information for p traits. First, we fix λ at a particular value and estimate the precision matrix $\Omega \in \Re^{p \times p}$ by running glasso. Next, we estimate the $(1 - \alpha)\%$ asymptotic confidence interval of all the obtained edges as

$$I_{ij} = \left[T_{ij} - \Phi^{-1} \left(1 - \frac{\alpha}{p(p-1)} \right) \frac{\hat{\sigma}_{ij}}{\sqrt{n}}, T_{ij} + \Phi^{-1} \left(1 - \frac{\alpha}{p(p-1)} \right) \frac{\hat{\sigma}_{ij}}{\sqrt{n}} \right]$$

References

- Friedman J, Hastie T, Tibshirani R. 2008.** Sparse inverse covariance estimation with the graphical lasso. *Biostatistics* **9**(3): 432-441.
- Jankova J, van de Geer S. 2015.** Confidence intervals for high-dimensional inverse covariance estimation. *Electronic Journal of Statistics* **9**(1): 1205-1229.
- Lauritzen SL. 1996.** *Graphical models*: Clarendon Press.

Appendix S5. Analysis of trait network across plant growth forms and climate regions.

Table S5.1. Trait centrality (i.e. Degree) for all plants, woody, and non-woody species and 95% confidence interval (CI). We ran this analysis for each network of all plants, woody and non-woody species for 1000 bootstrapped trait matrices.

Life form	Trait	2.50%	50%	97.50%
All	LLS	1	1	1
	SSD	1	1	1
	SLA	0.86	1	1
Non-woody	Seed mass	0.86	0.86	0.86
	Plant height	0.71	0.86	0.86
	Leaf area	0.71	0.71	0.86
	Leaf p	0.71	0.71	0.86
	Leaf n	0.71	0.71	0.71
	Leaf n	0.71	0.71	0.86
	Leaf area	0.57	0.71	0.86
	Leaf p	0.57	0.71	0.86
	LLS	0.57	0.71	0.86
	Plant height	0.57	0.57	0.86
	Seed mass	0.57	0.57	0.86
	SLA	0.43	0.57	0.57
	SSD	0.29	0.29	0.43
	LLS	0.86	0.86	1
	Leaf area	0.86	0.86	0.86
Woody	Seed mass	0.71	0.86	0.86
	Leaf n	0.71	0.71	0.71
	Leaf p	0.71	0.71	0.71
	SLA	0.57	0.71	0.71
	Plant height	0.57	0.57	0.57
	SSD	0.43	0.43	0.57

Table S5.2. Multiple-comparison adjusted 95% confidence interval (CI) from the trait centrality (i.e. degree) analyses for eight traits. We ran the trait centrality analyses for the trait of all plant species, woody and non-woody species across 1000 trait matrices. Differences in centrality between trait pairs significant when the 95% CI for the differences between trait pairs does not overlap with zero.

Trait-pairs	All		Woody		Non-woody	
	2.5%	97.5%	2.5%	97.5%	2.5%	97.5%
Leaf area-SLA	-0.29	0	0	0.29	-0.14	0.29
Leaf area-Leaf n	0	0.14	0	0.29	-0.29	0.14
Leaf area-LLS	-0.29	-0.14	-0.14	0	-0.29	0.29
Leaf area-Leaf p	-0.14	0	0.14	0.14	-0.29	0.29
Leaf area-Plant height	-0.14	0.14	0.14	0.43	-0.29	0.29
Leaf area-SSD	-0.29	-0.14	0.14	0.43	0.29	0.43
Leaf area-Seed mass	-0.29	0	0	0.14	-0.29	0.29
SLA-Leaf n	0.14	0.29	-0.29	0.14	-0.43	0
SLA-LLS	-0.14	0	-0.43	0	-0.43	0.14
SLA-Leaf p	0	0.29	-0.14	0.14	-0.43	0.14
SLA-Plant height	0.14	0.14	0	0.29	-0.29	0.14
SLA-SSD	-0.14	0	-0.14	0.43	0	0.43
SLA-Seed mass	0	0.14	-0.14	0	-0.29	0.14
Leaf n-LLS	-0.29	-0.29	-0.29	0	-0.14	0.29
Leaf n-Leaf p	-0.29	0	-0.14	0.14	-0.14	0.29
Leaf n-Plant height	-0.14	0	0	0.29	0	0.14
Leaf n-SSD	-0.29	-0.29	0	0.29	0.29	0.57
Leaf n-Seed mass	-0.29	-0.14	-0.29	0.14	-0.14	0.29
LLS-Leaf p	0	0.29	0.14	0.29	-0.14	0.29
LLS-Plant height	0.14	0.29	0.14	0.57	-0.14	0.29
LLS-SSD	0	0	0.14	0.43	0.14	0.71
LLS-Seed mass	0	0.14	0	0.29	-0.14	0.29
Leaf p-Plant height	-0.14	0.14	0	0.29	-0.14	0.29
Leaf p-SSD	-0.29	0	0	0.29	0.14	0.57
Leaf p-Seed mass	-0.14	0	-0.14	0	-0.14	0.14
Plant height-SSD	-0.29	-0.14	-0.14	0.29	0.14	0.57
Plant height-Seed mass	-0.14	0	-0.43	0	-0.29	0.29
SSD-Seed mass	0	0.14	-0.43	0	-0.57	-0.14

Table S5.3. Edge density for woody and non-woody species and 95% confidence interval (CI). We ran this analysis for each network of woody and non-woody species across climate regions for 1000 bootstrapped trait matrices.

Life form	Climate	2.50%	50%	97.50%
Woody	Tropical	0.61	0.64	0.68
	Temperate	0.61	0.68	0.71
	Arid	0.61	0.64	0.68
	Cold	0.53	0.57	0.64
	Polar	0.29	0.39	0.46
Non-woody	Tropical	0.29	0.36	0.46
	Temperate	0.5	0.57	0.64
	Arid	0.39	0.46	0.57
	Cold	0.57	0.61	0.68
	Polar	0.36	0.43	0.5

Table S5.4. Multiple-comparison adjusted 95% confidence interval (CI) from the edge density analyses among climate regions for woody and non-woody species. We ran the edge density comparison for woody and non-woody species across five climate regions for 1000 trait matrices. Differences in edge density between trait pairs significant when the 95% CI for the differences between trait pairs does not overlap with zero.

Regions-pairs	Woody		Non-woody	
	2.5%	97.5%	2.5%	97.5%
Arid-Cold	-0.04	0.18	-0.29	0.04
Arid-Polar	0.14	0.39	-0.11	0.21
Arid-Temperate	-0.11	0.07	-0.29	0.07
Arid-Tropical	-0.07	0.11	-0.07	0.29
Cold-Polar	0.04	0.36	0.07	0.32
Cold-Temperate	-0.21	0.04	-0.11	0.18
Cold-Tropical	-0.18	0.07	0.07	0.39
Polar-Temperate	-0.43	-0.14	-0.29	0.04
Polar-Tropical	-0.39	-0.11	-0.14	0.21
Temperate-Tropical	-0.07	0.14	0.04	0.39

Table S5.5. Modularity for woody and non-woody species and 95% confidence interval (CI). We ran this analysis for each network of woody and non-woody species across climate regions for 1000 bootstrapped trait matrices.

Life form	Climate	2.5%	50%	97.5%
Woody	Tropical	0.05	0.08	0.11
	Temperate	0.05	0.08	0.11
	Arid	0.08	0.11	0.15
	Cold	0.10	0.14	0.19
	Polar	0.16	0.25	0.39
Non-woody	Tropical	0.12	0.22	0.34
	Temperate	0.07	0.12	0.19
	Arid	0.09	0.19	0.28
	Cold	0.04	0.09	0.13
	Polar	0.13	0.21	0.34

Table S5.6. Multiple-comparison adjusted 95% confidence interval (CI) from the modularity analyses among climate regions for woody and non-woody species. We ran the modularity comparison for woody and non-woody species across five climate regions for 1000 trait matrices. Differences in modularity between regions is significant when the 95% CI for the differences between regions pairs does not overlap with zero.

Regions-pairs	Woody		Non-woody	
	2.5%	97.5%	2.5%	97.5%
Arid-Cold	-0.14	0.06	-0.04	0.25
Arid-Polar	-0.32	0.01	-0.25	0.17
Arid-Temperate	-0.07	0.11	-0.08	0.25
Arid-Tropical	-0.05	0.09	-0.21	0.16
Cold-Polar	-0.29	0.03	-0.33	0.03
Cold-Temperate	-0.02	0.17	-0.14	0.07
Cold-Tropical	-0.02	0.15	-0.3	0.02
Polar-Temperate	0.03	0.34	-0.06	0.32
Polar-Tropical	0.04	0.32	-0.2	0.22
Temperate-Tropical	-0.08	0.07	-0.28	0.07

Table S5.7. Trait centrality (i.e. Degree) for woody species across five climate regions and 95% confidence interval (CI). We ran this analysis over 1000 bootstrapped trait matrices.

Climate	Trait	2.5%	50%	97.5%
Tropical	Sm	0.86	0.86	0.86
	Leaf area	0.57	0.71	0.71
	LLS	0.57	0.71	0.86
	SLA	0.71	0.71	0.86
	Leaf N	0.57	0.57	0.71
	Leaf P	0.57	0.57	0.57
	Plht	0.57	0.57	0.71
	SSD	0.43	0.43	0.57
Temperate	LLS	0.71	0.86	1.00
	Leaf area	0.71	0.71	0.86
	Leaf P	0.43	0.71	0.86
	SLA	0.57	0.71	0.71
	Sm	0.71	0.71	0.71
	Leaf N	0.57	0.57	0.71
	Plht	0.43	0.57	0.71
	SSD	0.43	0.57	0.71
Arid	LLS	0.71	0.86	0.86
	Leaf area	0.71	0.71	0.86
	SLA	0.57	0.71	0.71
	Sm	0.57	0.71	0.71
	Leaf N	0.43	0.57	0.57
	Leaf P	0.43	0.57	0.71
	SSD	0.43	0.57	0.71
	Plht	0.43	0.43	0.57
Cold	LLS	0.57	0.71	1.00
	Plht	0.57	0.71	0.86
	Leaf area	0.57	0.57	0.57
	Leaf P	0.43	0.57	0.71
	SLA	0.57	0.57	0.71
	Sm	0.43	0.57	0.57
	Leaf N	0.43	0.43	0.57
	SSD	0.29	0.43	0.57
Polar	Leaf area	0.29	0.43	0.57
	Leaf N	0.29	0.43	0.43
	Leaf P	0.14	0.43	0.57
	LLS	0.29	0.43	0.57
	SLA	0.43	0.43	0.57
	Sm	0.29	0.43	0.43
	Plht	0.14	0.29	0.43
	SSD	0.14	0.29	0.29

Table S5.8. Trait centrality (i.e. Degree) for non-woody species across five climate regions and 95% confidence interval (CI). We ran this analysis over 1000 bootstrapped trait matrices.

Climate	Trait	2.5%	50%	97.5%
Tropical	Leaf area	0.29	0.43	0.57
	Leaf N	0.43	0.43	0.71
	LLS	0.14	0.43	0.57
	Sm	0.29	0.43	0.71
	Leaf P	0.14	0.29	0.71
	Plht	0.29	0.29	0.57
	SLA	0.14	0.29	0.57
	SSD	0.00	0.14	0.43
Temperate	Leaf N	0.57	0.71	0.86
	LLS	0.57	0.71	0.86
	Sm	0.57	0.71	0.86
	Leaf area	0.43	0.57	0.57
	Leaf P	0.43	0.57	0.86
	Plht	0.43	0.57	0.71
	SLA	0.43	0.43	0.43
	SSD	0.14	0.29	0.43
Arid	Leaf area	0.43	0.57	0.71
	Leaf N	0.43	0.57	0.86
	LLS	0.43	0.57	0.71
	SLA	0.29	0.57	0.71
	Sm	0.29	0.43	0.71
	Leaf P	0.29	0.29	0.57
	Plht	0.14	0.29	0.57
	SSD	0.29	0.29	0.43
Cold	Leaf area	0.71	0.86	0.86
	Leaf N	0.71	0.71	0.86
	LLS	0.57	0.71	0.86
	Sm	0.57	0.71	0.71
	Leaf P	0.43	0.57	0.71
	SLA	0.43	0.57	0.71
	Plht	0.29	0.43	0.57
	SSD	0.29	0.43	0.57
Polar	Leaf N	0.57	0.57	0.71
	Sm	0.43	0.57	0.71
	Leaf area	0.29	0.43	0.43
	Leaf P	0.29	0.43	0.57
	LLS	0.29	0.43	0.57
	Plht	0.29	0.43	0.57
	SLA	0.29	0.43	0.57
	SSD	0.00	0.14	0.43

Table S5.9. Multiple-comparison adjusted 95% confidence interval (CI) from the trait centrality (i.e. degree) analyses for eight traits of woody species across five climate regions. We ran the trait centrality analyses for the trait of woody species across five climate regions across 1000 trait matrices. Differences in centrality between trait pairs significant when the 95% CI for the differences between trait pairs does not overlap with zero.

Trait-pairs	Tropical		Temperate		Arid		Cold		Polar	
	2.5%	97.5%	2.5%	97.5%	2.5%	97.5%	2.5%	97.5%	2.5%	97.5%
Leaf area-SLA	-0.29	0.14	-0.14	0.29	-0.14	0.29	-0.14	0.14	-0.43	0.43
Leaf area-Leaf N	-0.14	0.14	0	0.29	0	0.43	0	0.14	-0.29	0.43
Leaf area-LLS	-0.14	0.14	-0.29	0.14	-0.29	0.14	-0.43	0	-0.29	0.29
Leaf area-Leaf P	-0.14	0.14	0	0.29	0	0.29	-0.14	0.29	-0.29	0.57
Leaf area-Plht	-0.14	0.29	0	0.43	0.14	0.43	-0.3	0.14	-0.14	0.43
Leaf area-SSD	-0.14	0.29	0	0.43	-0.01	0.43	0	0.29	-0.14	0.57
Leaf area-Sm	-0.43	-0.14	0	0.14	-0.14	0.29	-0.14	0.14	-0.29	0.43
SLA-Leaf N	0	0.43	-0.14	0.29	-0.01	0.29	0	0.29	-0.14	0.43
SLA-LLS	-0.14	0.29	-0.43	0	-0.29	0	-0.43	0.14	-0.29	0.43
SLA-Leaf P	0	0.43	-0.29	0.29	-0.14	0.29	-0.16	0.43	-0.29	0.43
SLA-Plht	0	0.43	-0.14	0.29	0	0.29	-0.29	0.14	-0.14	0.57
SLA-SSD	0	0.43	0	0.29	0	0.29	0	0.43	0	0.57
SLA-Sm	-0.29	0.14	-0.14	0.14	-0.14	0.14	-0.14	0.29	-0.29	0.43
Leaf N-LLS	-0.29	0.14	-0.43	0	-0.43	-0.14	-0.57	0	-0.29	0.29
Leaf N-Leaf P	-0.14	0.14	-0.29	0.29	-0.29	0.14	-0.29	0.14	-0.43	0.29
Leaf N-Plht	-0.14	0.14	-0.14	0.29	-0.14	0.14	-0.43	0	-0.14	0.43
Leaf N-SSD	0	0.29	-0.14	0.29	-0.29	0.29	-0.14	0.14	-0.14	0.43
Leaf N-Sm	-0.43	-0.14	-0.14	0.14	-0.14	0	-0.29	0.14	-0.29	0.29
LLS-Leaf P	-0.14	0.29	0	0.43	0.13	0.43	0	0.57	-0.29	0.43
LLS-Plht	0.14	0.57	-0.29	0.43	-0.29	0.43	0	0.57	-0.14	0.29
LLS-SSD	0	0.43	0.14	0.57	-0.14	0.57	0.14	0.43	-0.14	0.43
LLS-Sm	0	0.3	0	0.43	-0.29	0.43	0	0.29	-0.43	0
Leaf P-Plht	0	0.29	-0.43	0.01	-0.29	0.43	-0.14	0.29	-0.14	0.14
Leaf P-SSD	-0.16	0.29	-0.14	0.3	-0.14	0.43	-0.29	0.43	-0.14	0.29
Leaf P-Sm	-0.29	0.14	-0.29	0.29	-0.29	0.43	-0.29	0.14	-0.29	-0.14
Plht-SSD	-0.16	0.14	0	0.57	-0.14	0.43	-0.29	0.29	-0.14	0.29
Plht-Sm	-0.29	0	-0.14	0.43	-0.43	0.29	-0.29	0.14	-0.43	-0.14
SSD-Sm	-0.29	0.14	-0.29	0.14	-0.43	0.14	-0.29	0.14	-0.57	-0.14

Table S5.10. Multiple-comparison adjusted 95% confidence interval (CI) from the trait centrality (i.e. degree) analyses for eight traits of non-woody species across five climate regions. We ran the trait centrality analyses for the trait of non-woody species across five climate regions across 1000 trait matrices. Differences in centrality between trait pairs significant when the 95% CI for the differences between trait pairs does not overlap with zero.

Trait-pairs	Tropical		Temperate		Arid		Cold		Polar	
	2.5%	97.5%	2.5%	97.5%	2.5%	97.5%	2.5%	97.5%	2.5%	97.5%
Leaf area-SLA	-0.29	0.57	-0.14	0.29	-0.14	0.29	0	0.29	-0.29	0.29
Leaf area-Leaf N	-0.29	0.14	-0.43	0.14	-0.43	0.29	-0.14	0.29	-0.43	0
Leaf area-LLS	-0.29	0.43	-0.43	0.14	-0.29	0.29	-0.29	0.43	-0.29	0.29
Leaf area-Leaf P	-0.43	0.57	-0.29	0.29	-0.14	0.43	0	0.43	-0.29	0.14
Leaf area-Plht	-0.29	0.43	-0.29	0.29	0	0.57	0.14	0.71	-0.29	0.29
Leaf area-SSD	-0.14	0.57	0	0.57	0	0.57	0.14	0.57	-0.14	0.43
Leaf area-Sm	-0.43	0.43	-0.43	0.14	-0.29	0.57	-0.14	0.43	-0.43	0.14
SLA-Leaf N	-0.57	0.29	-0.57	0	-0.43	0.29	-0.43	0.14	-0.43	0.14
SLA-LLS	-0.29	0.43	-0.43	0	-0.43	0.29	-0.43	0.14	-0.29	0.29
SLA-Leaf P	-0.43	0.43	-0.43	0.14	-0.29	0.57	-0.29	0.29	-0.43	0.29
SLA-Plht	-0.29	0.29	-0.43	0	-0.14	0.57	-0.14	0.43	-0.29	0.29
SLA-SSD	-0.29	0.71	-0.14	0.57	-0.14	0.43	0	0.43	-0.14	0.57
SLA-Sm	-0.43	0.29	-0.43	0	-0.29	0.43	-0.29	0.14	-0.43	0.14
Leaf N-LLS	-0.14	0.43	-0.29	0.29	-0.29	0.43	-0.29	0.29	0	0.57
Leaf N-Leaf P	-0.43	0.57	-0.29	0.57	-0.14	0.57	0	0.43	0	0.43
Leaf N-Plht	-0.14	0.43	-0.14	0.43	-0.14	0.57	0.14	0.57	0	0.43
Leaf N-SSD	-0.14	0.57	0.14	0.71	0.14	0.43	0.14	0.57	0	0.71
Leaf N-Sm	-0.29	0.43	-0.29	0.29	-0.14	0.57	-0.14	0.29	-0.14	0.43
LLS-Leaf P	-0.43	0.29	-0.29	0.43	-0.14	0.57	-0.14	0.29	-0.43	0.29
LLS-Plht	-0.29	0.29	-0.29	0.43	-0.14	0.57	0	0.57	-0.29	0.14
LLS-SSD	-0.29	0.43	0.14	0.71	-0.14	0.57	0	0.57	-0.14	0.43
LLS-Sm	-0.57	0.29	-0.29	0.29	-0.29	0.57	-0.14	0.29	-0.43	0.14
Leaf P-Plht	-0.29	0.43	-0.29	0.29	-0.29	0.43	-0.14	0.43	-0.29	0.43
Leaf P-SSD	-0.29	0.57	-0.14	0.86	-0.14	0.43	-0.14	0.29	-0.14	0.43
Leaf P-Sm	-0.43	0.43	-0.43	0.14	-0.43	0.43	-0.29	0.14	-0.43	0.43
Plht-SSD	-0.14	0.57	0	0.57	-0.29	0.29	-0.29	0.29	-0.14	0.57
Plht-Sm	-0.43	0.29	-0.43	0.29	-0.43	0.29	-0.43	0	-0.43	0.14
SSD-Sm	-0.57	0.14	-0.86	-0.14	-0.43	0.29	-0.43	0	-0.57	0