



HAL
open science

Linking Termite Feeding Preferences and Soil Physical Functioning in Southern-Indian Woodlands

Sougueh Cheik, Rashmi Ramesh Shanbhag, Ajay Harit, Nicolas Bottinelli,
Raman Sukumar, Pascal Jouquet

► **To cite this version:**

Sougueh Cheik, Rashmi Ramesh Shanbhag, Ajay Harit, Nicolas Bottinelli, Raman Sukumar, et al..
Linking Termite Feeding Preferences and Soil Physical Functioning in Southern-Indian Woodlands.
Insects, 2019, 10 (1), pp.4. 10.3390/insects10010004 . hal-02045875

HAL Id: hal-02045875

<https://hal.sorbonne-universite.fr/hal-02045875>


Submitted on 22 Feb 2019

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Article

Linking Termite Feeding Preferences and Soil Physical Functioning in Southern-Indian Woodlands

Sougueh Cheik^{1,2,*}, Rashmi Ramesh Shanbhag³, Ajay Harit⁴, Nicolas Bottinelli^{2,5} ,
Raman Sukumar⁶ and Pascal Jouquet^{1,2}

¹ Indo-French Cell for Water Science (IFCWS), Civil Engineering Department, Indian Institute of Science, Bangalore 560012, India; pascal.jouquet@ird.fr

² Institute of Ecology and Environmental Sciences (UMR 242 iEES-Paris), Sorbonne Universités, Centre IRD, 93143 Bondy, France; nicolas.bottinelli@ird.fr

³ Indian Plywood Industries Research and Training Institute, Bangalore 560022, India; rashmishanbhags@gmail.com

⁴ School of Environmental Sciences, Mahatma Gandhi University, Kottayam 686560, India; om.harit@gmail.com

⁵ Soils and Fertilizers Research Institute (SFRI), Dong Ngac, Tu Liem, Ha Noi, Vietnam

⁶ Centre for Ecological Sciences, Indian Institute of Science, Bangalore 560012, India; rsuku@iisc.ac.in

* Correspondence: sougueh.cheik@ird.fr; Tel.: +84-16-6284-0970

Received: 31 October 2018; Accepted: 27 December 2018; Published: 4 January 2019



Abstract: Termites are undoubtedly amongst the most important soil macroinvertebrate decomposers in semi-arid environments in India. However, in this specific type of environment, the influence of termite foraging activity on soil functioning remains unexplored. Therefore, this study examines the link between the quality of litter and the functional impact of termite feeding preferences on soil properties and soil hydraulic conductivity in a deciduous forest in southern India. Different organic resources (elephant dung: “ED”, elephant grass: “EG”, acacia leaves: “AL” and layers of cardboard: “CB”) were applied on repacked soil cores. ED appeared to be the most attractive resource to *Odontotermes obesus*, leading to a larger amount of soil sheeting (i.e., the soil used by termites for covering the litter they consume), more numerous and larger holes in the ground and a lower soil bulk density. As a consequence, ED increased the soil hydraulic conductivity (4-fold) compared with the control soil. Thus, this study highlights that the more *O. obesus* prefers a substrate, the more this species impacts soil dynamics and water infiltration in the soil. This study also shows that ED can be used as an efficient substrate for accelerating the infiltration of water in southern-Indian soils, mainly through the production of galleries that are open on the soil surface, offering new perspectives on termite management in this environment.

Keywords: *Odontotermes obesus*; sheeting; termite foraging activity; litter quality; organic resource consumption; soil water dynamic

1. Introduction

Soil biodiversity regulates a large number of ecological functions, such as the degradation of litter, the cycling of nutrients or the regulation of water dynamics in the soil [1,2]. Amongst soil organisms, termites are increasingly recognized as playing a role in the provisioning of key ecosystem services [3,4]. Termites are ubiquitously amongst the most abundant and active litter decomposers in tropical environments [5] and are also considered significant soil bioturbators or ecosystem engineers because of the biostructures they produce [6,7]. Indeed, a large body of literature describes the specific soil biological, chemical and physical properties of termite mound nests [8–17] and the link between

these properties and the ecological needs of termites (i.e., to main an homeostatic environment within their nest for the colony and the fungus to grow [18–20]).

However, less is known about the biostructures termites produce while they forage on the ground. Indeed, termites also build belowground galleries for foraging and translocate large quantities of soil to the surface (i.e., so-called sheetings) to cover the litter they consume (e.g., leaves, wood and herbivore dung) [7], and protect themselves from light, desiccation and predators [21–24]. A recent review revealed that sheetings have different soil properties than the surrounding top-soil, depending on the properties of the top-soil and the strategy developed by termites (i.e., favouring clay or organic matter for ensuring the stability of sheetings) [23]. Less is known, however, about the quantity of sheetings that termites produce on the ground and whether this quantity varies with the quality of the substrate termites consume [25,26]. The production of sheeting is associated with the construction of belowground galleries that are used by termites to bring litter into their nest. These galleries have a significant and positive impact on water infiltration in soil and therefore reduce the susceptibility of the soil to erosion [27–29], although the opposite can be observed when soil sheetings collapse and produce a structural crust on the ground, limiting water infiltration and increasing water runoff [30].

Since most of the research on the functional impact of termite foraging activity has been carried out in Africa and to a lesser extent in South-America and Australia, the aim of this research was to determine whether termite foraging activity varies with the quality of the litter in a southern-Indian forest. More precisely, this study also questioned the relationship between termite food preferences and the abundance of soil sheeting on the ground and water infiltration in the soil.

2. Materials and Methods

2.1. Study Site

This study was performed during the dry season from February to March 2016 in the forest of the Jubilee Garden of the Indian Institute of Science (13°01'17" N and 77°34'14" E) in Bangalore city, Karnataka state, India. This ecosystem has a tropical savannah climate with distinct wet and dry seasons, and the annual rainfall ranges from 900 to 1100 mm yr⁻¹ [22,31]. The soil is described as an Alfisol based on U.S. Soil Taxonomy (United States Department of Agriculture, USDA) or a Luvisol according to Food and Agriculture Organization (FAO) classification. The first soil layer (0–10 cm depth) is mainly sandy (≈60%) and contains approximately 10 to 20% clay, mainly kaolinite [32]. The soil pH is 5.7, and its C content reaches 2.2% on average [22]. The vegetation is a deciduous forest dominated by acacia trees, mainly *Acacia auriculiformis*.

In this study site, the litter-feeding termites are mainly *Odontotermes feae*, *O. obesus* and *O. feoides*, and their activity involves the production of soil sheetings on wood logs or fallen leaves on the ground [22,31]. *Odontotermes* spp. belong to the Macroterminae subfamily, also named fungus-growing termites. If *Odontotermes* spp. may appear as major crop pests in some environments, they also play an important role in more natural environments where they act as key decomposers and bioturbators.

2.2. Experimental Design

Repacked soil cores were prepared using plastic cylinders (PVC) (5 cm in height and 15 cm in diameter). The soil was sampled from the 0–10 cm top-soil layer, air dried and sieved through a 2 mm mesh before being compacted in the cylinders until reaching a density similar to that found in the field (i.e., 1.2 g cm⁻³). In total, each cylinder contained 500 g of soil (dry weight, DW). Cylinders were inserted into the soil to a depth of 3 cm at randomly chosen locations (mean average distance between samples: ≈2 m). The soil in the cylinders was covered by organic resources, namely leaves of elephant grass (*Pennisetum purpureum*: “EG”, 16 g DW) or acacia (*Acacia auriculiformis*: “AL”, 11 g DW), elephant dung pats (*Elephas maximus*: “ED”, 230 g DW) or 10 cm² layers of cardboard (“CB”, 14 g DW). These substrates were chosen because of their availability and attractiveness to termites ($n = 25$ with 4 treatments + 1 control without organic resource × 5 replicates).

2.3. Biodiversity and Food Consumption Rates

After two months, the soil macrofauna within or below the organic resources were sampled and preserved in 70% alcohol. Individuals were identified and their numbers were counted. Organic substrates were collected, air dried and weighed. The food consumption rate was calculated as the percentage of weight lost.

2.4. Termite Bioturbation

At the end of the experiment, soil sheetings that covered the organic resources were collected, dried at 40 °C for 2 days and weighed. The soil turnover activity was measured by dividing the dry weight of sheetings by the quantity of organic material removed [26]. The number and diameter of galleries open on the surface were also measured with a caliper. Soil hydraulic conductivity at saturation (K_{sat}) was measured with the Beerkan method [33–35]. A fixed volume of water (100 mL) was poured into the cylinder, and the time needed for the water to infiltrate was measured. The procedure was repeated between 7 to 10 times until a steady state of infiltration was reached. The soil cores ($\approx 530 \text{ cm}^3$) were then used to determine the soil bulk density, ρ (g cm^{-3}) and the initial volumetric water content, θ_i ($\text{m}^3 \text{ m}^{-3}$), was measured after sampling the surrounding top-soil environment (0–5 cm depth). The results were analysed with the original BEST algorithm [35].

Root biomass was measured in the soil cylinders. Samples were placed in beakers and soaked in water for at least 30 min, such that the soil aggregates could be easily broken down, after sieving through a 200 μm mesh. Roots were rinsed to remove soil particles and dried at 60 °C for 48 h.

2.5. Statistical Analyses

The normality of residuals was tested using the Shapiro–Wilk test. One-way analysis of variance (ANOVA) and least significant difference (LSD) tests were performed to assess differences between means. Partial least squares regression (PLSR) analysis was used to identify important variables linked to soil hydraulic conductivity. Correlations between variables were tested using the Pearson method. All statistical calculations were carried out using R version 3.5.1 (<https://www.r-project.org/>). Differences among treatments were declared significant at the < 0.05 probability level.

3. Results and Discussion

3.1. Organic Resource Consumption by Termites

At the end of the experiment, termites were only found in or beneath the elephant dung pads. A very small amount of earthworm casts was found beneath two cardboard pieces. Termite individuals were all identified as belonging to *O. obesus* (Termitidae, Macrotermitinae), thus confirming the importance of this termite species in southern-Indian ecosystems [17]. Termite occurrence was associated with a consumption of the organic material, as shown by the more important consumption rate of ED than the other organic resources (Figure 1, ANOVA test, $F_{4,20} = 5.30$, $p = 0.005$), while no differences occurred between the AL, EG and CB treatments ($p > 0.05$ between each pair).

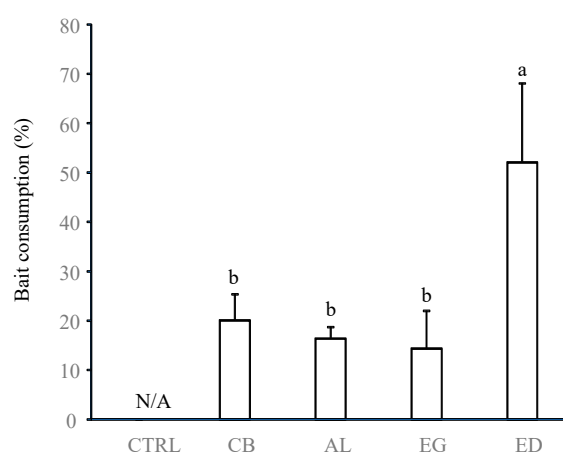


Figure 1. Rate of organic resource consumption by termites, in % of the initial organic resource weight loss (elephant dung (“ED”), acacia leaves (“AL”), elephant grass (“EG”) and cardboard (“CB”)), Error bars represent the standard error of the mean. Histograms with the same letter are not significantly different at $p \geq 0.05$ ($n = 5$). N/A for non-applicable in the case of the control.

The opportunistic consumption of mammalian dung by termites has been described in Africa for a long time [36,37]. Our study is, however, the first to stress the higher attractiveness of elephant dung to *O. obesus* in Asia compared to other organic resources. The rapid detection and consumption of ED by termites is in line with Cheik et al. [38], who found that elephant dung can be used to stimulate termite foraging activity in southern India. The low attractiveness of EG, CB, and AL to termites was more surprising since cellulose baits are commonly used to measure termite foraging activity in dry regions [39,40] and because termites respond the most to substrates with the highest C:N ratio [41]. We explain this low activity in EG, CB and AL treatments by the high attractiveness of ED in terms of nutrient content and/or because it provides a thermal shadow [42]. However, our experiment only lasted for 2 months, and it is likely that these substrates would have been more consumed if the experiment had lasted longer. Consequently, it can be concluded from this experiment that elephant dung is a sporadic and unpredictable but effective resource for termites and might be preferred over plant litter (EG and AL) or only cellulose (CB).

3.2. Relationship between Feeding Preferences and Soil Functioning

Figures 2–4 show that the preference of termites for ED was associated with a more important production of soil sheeting and more numerous and larger pores at the soil surface compared with the other substrates ($p < 0.05$ in all cases, Table 1). Conversely, no significant differences in sheeting production and diameter of the foraging holes were found between the AL, EG, CB and CTRL treatments ($p > 0.05$ in all cases), while the numbers of foraging holes in the CB and AL treatments were intermediate to those for the ED, EG and CTRL treatments.

Table 1. Results of the ANOVA test showing the influence of organic resource quality (control, elephant dung, elephant grass, acacia leaves and cardboard) on the food consumption rate (%), diameter (mm) and number of foraging holes, amount of sheetings (g), soil bulk density (g cm^{-3}), soil turnover activity ($\text{g soil g consumed}^{-1}$) and root biomass (mg g^{-1}) ($n = 5$ in all cases).

	$F_{4,20}$	p -Value
Consumption rate	5.29	0.004
Diameter of holes	7.78	<0.001
Number of holes	13.04	<0.001
Amount of sheetings	6.37	0.004
Soil bulk density	4.59	0.009
Turnover activity	1.63	0.207
Roots biomass	1.80	0.168

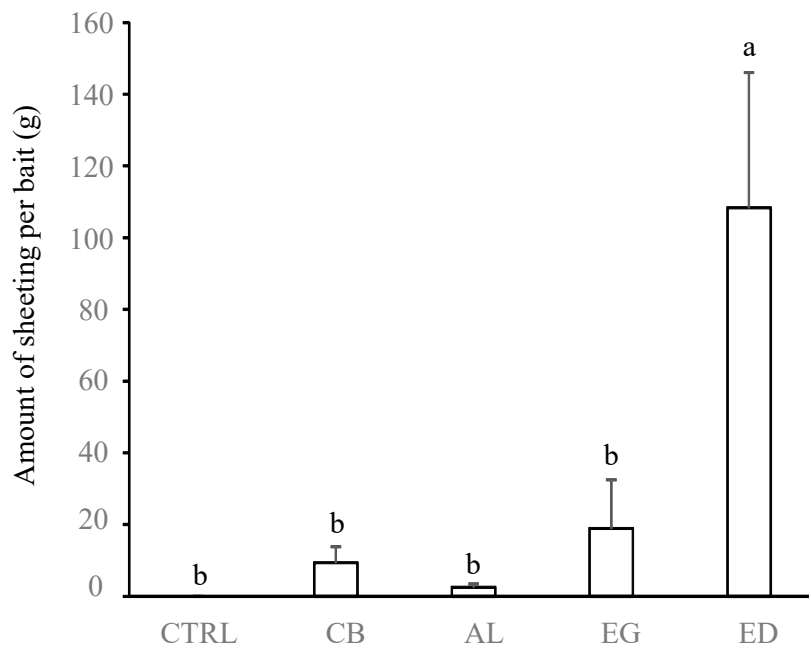


Figure 2. Amount of termite sheeting (g) for each treatment. Treatments are: control (“CTRL”), elephant dung (“ED”), acacia leaves (“AL”), elephant grass (“EG”) and cardboard (“CB”). Error bars represent the standard error of the mean. Histograms with the same letter are not significantly different at $p \geq 0.05$ (n = 5).

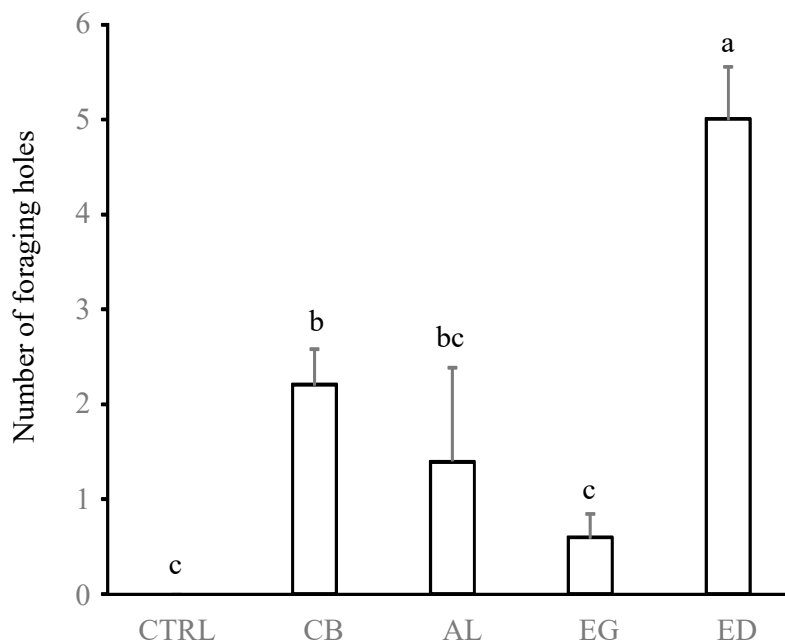


Figure 3. Number of foraging holes observed on the soil surface. Treatments are: control (“CTRL”), elephant dung (“ED”), acacia leaves (“AL”), elephant grass (“EG”) and cardboard (“CB”). Error bars represent the standard error of the mean. Histograms with the same letter are not significantly different at $p \geq 0.05$ (n = 5).

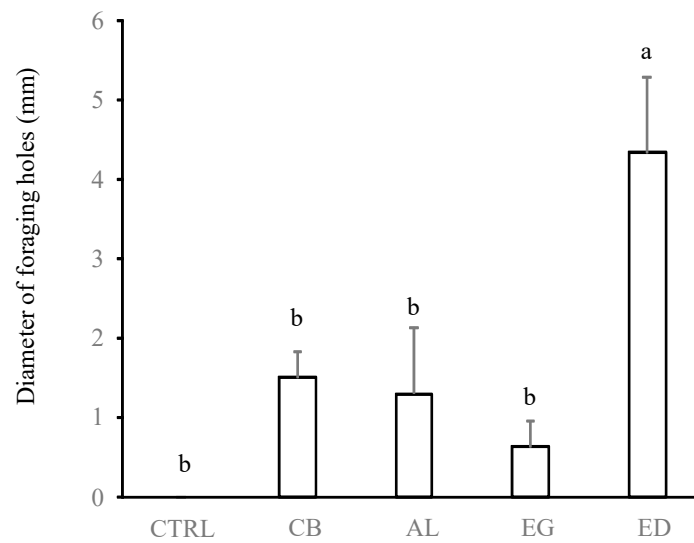


Figure 4. Average diameter of foraging holes (in mm). Treatments are: control (“CTRL”), elephant dung (“ED”), acacia leaves (“AL”), elephant grass (“EG”) and cardboard (“CB”). Error bars represent the standard error of the mean. Histograms with the same letter are not significantly different at $p \geq 0.05$ ($n = 5$).

These findings are clearly in line with those from studies carried out in Africa [25,43–45] that showed that fungus-growing termites bring soil from different soil layers to build the sheetings that they use to cover themselves and the food they consume. Although the specific chemical, physical and biological properties of termite sheetings have been previously described [23], demonstration of the relationship between termite feeding preferences and the production of sheetings above-ground or the construction of galleries below-ground remains limited to studies that have been carried out in Africa [25,46,47]. In our study, no significant difference in soil turnover activity was measured between treatments (Table 1), suggesting that the returns on investment (i.e., the amount of energy spent on building the sheeting per the amount of energy received from the organic resources) were similar despite the difference in food quality and preference. However, the production of sheeting was positively related to the consumption rate and the diameter of foraging holes (Figure 5).

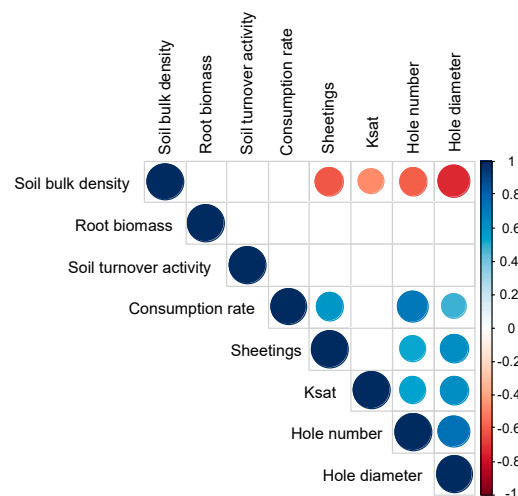


Figure 5. Matrix of correlations between variables. Only significant correlations ($p < 0.05$) are given. The scale colour indicates whether correlations between variables were positive (closer to 1, blue circles) or negative (closer to -1 , red squares).

Despite the fact that these results have to be interpreted with caution because variables reached important values only for the ED treatment but remained very low for the other treatments, the findings of our study suggest that the amount of sheeting can be used as an indicator of *O. obesus* feeding preference. These findings also suggest that the more *O. obesus* termites prefer a substrate, the more they circulate within galleries and, as a consequence, increase the diameters of the gallery openings. This hypothesis explains the significant reduction in soil bulk density in the ED treatment compared with the other treatments ($p < 0.05$ in all cases, Figure 6), as well as the negative relationship of soil bulk density with the diameter of foraging holes and the quantity of soil sheeting (Figure 5).

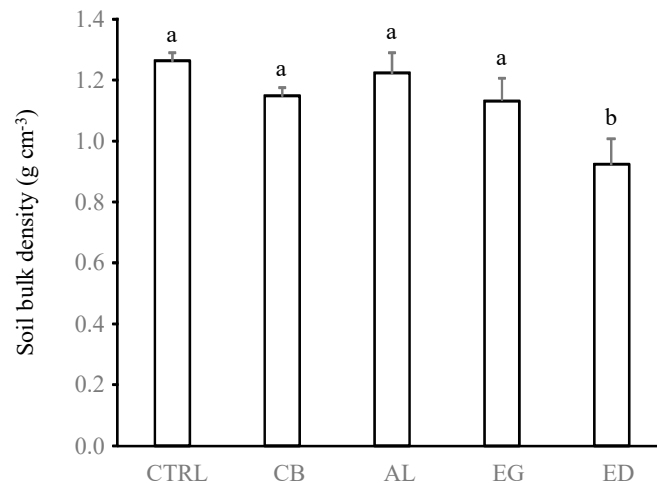


Figure 6. Soil bulk density (g cm^{-3}) below the treatments (control (“CTRL”), elephant dung (“ED”), acacia leaves (“AL”), elephant grass (“EG”) and cardboard (“CB”). Error bars represent the standard error of the mean. Histograms with the same letter are not significantly different at $p \geq 0.05$ ($n = 5$).

Odontotermes spp. mainly build superficial and horizontal galleries localized in the first centimetre of the soil [31,47]. Our study shows that the production of these foraging galleries was associated with a reduction in soil bulk density in the ED treatment as well as an increase in K_{sat} of 4 and 2.6-fold in the ED and CB treatments compared with the CTRL treatment, respectively (ANOVA test, $F_{4,10} = 8.36$, $p = 0.003$, Figure 7). Conversely, no significant differences were measured in the AL, EG and CTRL treatments ($p > 0.05$). These results are in line with those found in Cheik et al. [31] (3-fold) and Kaiser et al. [26] (1.5 to 9.28-fold), although termite foraging is usually increased by only 1.5- to 3-fold [27,47,48].

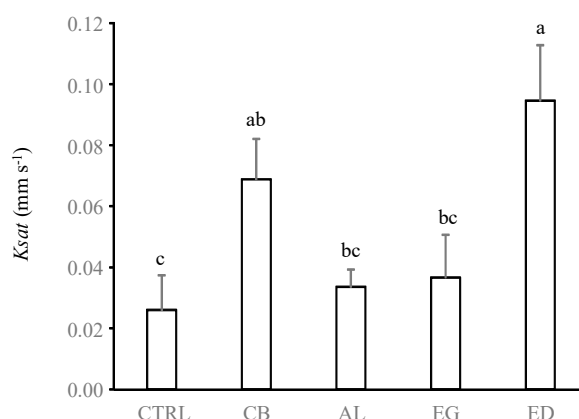


Figure 7. Soil hydraulic conductivity (K_{sat} , in mm s^{-1}) for the different treatments (control (“CTRL”), elephant dung (“ED”), acacia leaves (“AL”), elephant grass (“EG”) and cardboard (“CB”). Error bars represent the standard error of the mean. Histograms with the same letter are not significantly different at $p \geq 0.05$ ($n = 5$).

The diameter and number of holes were identified as the most important factors associated with K_{sat} (PLSR model, $Q^2 = 0.52$, Root Mean Square Error of Prediction (RMSEP) = 0.042, Table 2), which is in line with the general assumption that a higher soil macroporosity increases the soil hydraulic conductivity [49].

Consequently, the findings of this study confirmed the general assumption that the stimulation of termites can accelerate the infiltration of water in soil, mainly through the production galleries that are open on the soil surface [26–28,47,50].

Table 2. Variable important in the projection (VIP) scores from the most relevant variables used for the PLSR describing the evolution of soil hydraulic conductivity at saturation (K_{sat}).

Variables	VIP Scores
Amount of sheetings	6.97
Decomposition rate	0.99
Hole diameter	0.02
Hole number	0.01

4. Conclusions

Although a large number of studies have reported a link between termite feeding preferences and soil bioturbation in Africa, this relationship remains poorly understood in Asia where services provided by termites can differ, for instance due to different soil, climate and termite assemblage. This study is the first to stress that (i) the more termites prefer a substrate, the more they impact soil dynamic and water infiltration in the soil; (ii) elephant dung is an efficient amendment for stimulating termite activity in southern India; and (iii) its consumption by termites is associated with an increase in soil dynamic and water infiltration in the soil. In conclusion, we argue that an economic and agronomical study is now needed to determine whether this substrate can be used in cultivated agro-systems to attract termites and promote soil translocation, aeration and water infiltration.

Author Contributions: S.C., P.J., A.H. and R.R.S. designed and performed the experimental setup and implementation of the research; R.S. provided access to the study site; analyses were carried out by S.C., N.B. and P.J. The article was written by S.C. and P.J. in consultation with all authors.

Funding: This research was funded by the Indo-French Cell for Water Science (LMI IFCWS/CEFIRSE, IRD) and the French National Program EC2CO-Biohefect “MACROFLUX”.

Acknowledgments: We would like to thank S. Subramanian, Nirmala. S. and J. Riotte for their supports and for allowing us to have access to facilities from Indian Institute of Science (IISc). This project was supported by the French Institute for research for development (IRD) through “Allocations de recherche pour une thèse au Sud” (ARTS) grant.

Conflicts of Interest: The authors declare no conflict of interest.

References

1. Lavelle, P.; Decaëns, T.; Aubert, M.; Barot, S.; Blouin, M.; Bureau, F.; Margerie, P.; Mora, P.; Rossi, J.-P. Soil invertebrates and ecosystem services. *Eur. J. Soil Biol.* **2006**, *42*, S3–S15. [[CrossRef](#)]
2. Wall, D.H.; Nielsen, U.N.; Six, J. Soil biodiversity and human health. *Nature* **2015**. [[CrossRef](#)] [[PubMed](#)]
3. Holt, J.A.; Lepage, M. Termites and Soil Properties. In *Termites: Evolution, Sociality, Symbioses, Ecology*; Abe, T., Bignell, D.E., Higashi, M., Eds.; Springer: Dordrecht, The Netherlands, 2000; pp. 389–407, ISBN 978-90-481-5476-0.
4. Jouquet, P.; Traoré, S.; Choosai, C.; Hartmann, C.; Bignell, D. Influence of termites on ecosystem functioning. Ecosystem services provided by termites. *Eur. J. Soil Biol.* **2011**, *47*, 215–222. [[CrossRef](#)]
5. Bignell, D.E.; Eggleton, P. Termites in Ecosystems. In *Termites: Evolution, Sociality, Symbioses, Ecology*; Abe, T., Bignell, D.E., Higashi, M., Eds.; Springer: Dordrecht, The Netherlands, 2000; pp. 363–387, ISBN 978-90-481-5476-0.
6. Bottinelli, N.; Jouquet, P.; Capowiez, Y.; Podwojewski, P.; Grimaldi, M.; Peng, X. Why is the influence of soil macrofauna on soil structure only considered by soil ecologists? *Soil Tillage Res.* **2015**, *146*, 118–124. [[CrossRef](#)]
7. Jouquet, P.; Guilleux, N.; Caner, L.; Chintakunta, S.; Ameline, M.; Shanbhag, R.R. Influence of soil pedological properties on termite mound stability. *Geoderma* **2016**, *262*, 45–51. [[CrossRef](#)]
8. Jouquet, P.; Tessier, D.; Lepage, M. The soil structural stability of termite nests: Role of clays in *Macrotermes bellicosus* (Isoptera, Macrotermitinae) mound soils. *Eur. J. Soil Biol.* **2004**, *40*, 23–29. [[CrossRef](#)]
9. Jouquet, P.; Bottinelli, N.; Lata, J.-C.; Mora, P.; Caquineau, S. Role of the fungus-growing termite *Pseudacanthotermes spiniger* (Isoptera, Macrotermitinae) in the dynamic of clay and soil organic matter content. An experimental analysis. *Geoderma* **2007**, *139*, 127–133. [[CrossRef](#)]
10. Jouquet, P.; Guilleux, N.; Shanbhag, R.R.; Subramanian, S. Influence of soil type on the properties of termite mound nests in Southern India. *Appl. Soil Ecol.* **2015**, *96*, 282–287. [[CrossRef](#)]
11. Abe, S.S.; Kotegawa, T.; Onishi, T.; Watanabe, Y.; Wakatsuki, T. Soil particle accumulation in termite (*Macrotermes bellicosus*) mounds and the implications for soil particle dynamics in a tropical savanna Ultisol. *Ecol. Res.* **2012**, *27*, 219–227. [[CrossRef](#)]
12. Tilahun, A.; Kebede, F.; Yamoah, C.; Erens, H.; Mujinya, B.B.; Verdoodt, A.; Van Ranst, E. Quantifying the masses of *Macrotermes subhyalinus* mounds and evaluating their use as a soil amendment. *Agric. Ecosyst. Environ.* **2012**, *157*, 54–59. [[CrossRef](#)]
13. Horiuchi, Y.; Ohno, T.; Hoshino, M.; Shin, K.-C.; Murakami, H.; Tsunematsu, M.; Watanabe, Y. Geochemical prospecting for rare earth elements using termite mound materials. *Miner. Depos.* **2014**, *49*, 1013–1023. [[CrossRef](#)]
14. Menichetti, L.; Landi, L.; Nannipieri, P.; Katterer, T.; Kirchmann, H.; Renella, G. Chemical Properties and Biochemical Activity of Colonized and Abandoned Litter-Feeding Termite (*Macrotermes* spp.) Mounds in Chromic Cambisol Area on the Borana Plateau, Ethiopia. *Pedosphere* **2014**, *24*, 399–407. [[CrossRef](#)]
15. Seymour, C.L.; Milewski, A.V.; Mills, A.J.; Joseph, G.S.; Cumming, G.S.; Cumming, D.H.M.; Mahlangu, Z. Do the large termite mounds of *Macrotermes* concentrate micronutrients in addition to macronutrients in nutrient-poor African savannas? *Soil Biol. Biochem.* **2014**, *68*, 95–105. [[CrossRef](#)]
16. Erens, H.; Mujinya, B.B.; Mees, F.; Baert, G.; Boeckx, P.; Malaisse, F.; Van Ranst, E. The origin and implications of variations in soil-related properties within *Macrotermes falciger* mounds. *Geoderma* **2015**, *249–250*, 40–50. [[CrossRef](#)]
17. Shanbhag, R.R.; Kabbaj, M.; Sundararaj, R.; Jouquet, P. Rainfall and soil properties influence termite mound abundance and height: A case study with *Odontotermes obesus* (Macrotermitinae) mounds in the Indian Western Ghats forests. *Appl. Soil Ecol.* **2017**, *111*, 33–38. [[CrossRef](#)]
18. Lüscher, M. Air-Conditioned Termite Nests. *Sci. Am.* **1961**, *205*, 138–145. [[CrossRef](#)]

19. Korb, J.; Linsenmair, K.E. Thermoregulation of termite mounds: What role does ambient temperature and metabolism of the colony play? *Insectes Sociaux* **2000**, *47*, 357–363. [[CrossRef](#)]
20. Ndlovu, M.; Pérez-Rodríguez, A. Temperature fluctuations inside savanna termite mounds: Do size and plant shade matter? *J. Therm. Biol.* **2018**, *74*, 23–28. [[CrossRef](#)]
21. Mackay, W.P.; Blizzard, J.H.; Miller, J.J.; Whitford, W.G. Analysis of Above-ground Gallery Construction by the Subterranean Termite *Gnathamitermes tubiformans* (Isoptera: Termitidae). *Environ. Entomol.* **1985**, *14*, 470–474. [[CrossRef](#)]
22. Jouquet, P.; Guilleux, N.; Chintakunta, S.; Mendez, M.; Subramanian, S.; Shanbhag, R.R. The influence of termites on soil sheeting properties varies depending on the materials on which they feed. *Eur. J. Soil Biol.* **2015**, *69*, 74–78. [[CrossRef](#)]
23. Harit, A.; Moger, H.; Duprey, J.-L.; Gajalakshmi, S.; Abbasi, S.A.; Subramanian, S.; Jouquet, P. Termites can have greater influence on soil properties through the construction of soil sheetings than the production of above-ground mounds. *Insectes Sociaux* **2017**, *64*, 247–253. [[CrossRef](#)]
24. Fernandes, G.W.; Murcia, S.L.; Santos, J.C.; DeSouza, O.; Constantino, R.; Hafig, I. Termite Foraging on Plants of a Brazilian Savanna: The Effects of Tree Height. *Sociobiology* **2018**, *65*, 48. [[CrossRef](#)]
25. Rouland, C.; Lepage, M.; Chotte, J.L.; Diouf, M.; Ndiaye, D.; Ndiaye, S.; Seugé, C.; Brauman, A. Experimental manipulation of termites (Isoptera, Macrotermitinae) foraging patterns in a Sahelo-Sudanese savanna: Effect of litter quality. *Insectes Sociaux* **2003**, *50*, 309–316. [[CrossRef](#)]
26. Kaiser, D.; Lepage, M.; Konaté, S.; Linsenmair, K.E. Ecosystem services of termites (Blattoidea: Termitoidea) in the traditional soil restoration and cropping system Zaï in northern Burkina Faso (West Africa). *Agric. Ecosyst. Environ.* **2017**, *236*, 198–211. [[CrossRef](#)]
27. Mando, A.; Stroosnijder, L.; Brussaard, L. Effects of termites on infiltration into crusted soil. *Geoderma* **1996**, *74*, 107–113. [[CrossRef](#)]
28. Mando, A.; Brussaard, L.; Stroosnijder, L. Termite- and Mulch-Mediated Rehabilitation of Vegetation on Crusted Soil in West Africa. *Restor. Ecol.* **1999**, *7*, 33–41. [[CrossRef](#)]
29. Léonard, J.; Perrier, E.; Rajot, J.L. Biological macropores effect on runoff and infiltration: A combined experimental and modelling approach. *Agric. Ecosyst. Environ.* **2004**, *104*, 277–285. [[CrossRef](#)]
30. Jouquet, P.; Janeau, J.-L.; Pisano, A.; Sy, H.T.; Orange, D.; Minh, L.T.N.; Valentin, C. Influence of earthworms and termites on runoff and erosion in a tropical steep slope fallow in Vietnam: A rainfall simulation experiment. *Appl. Soil Ecol.* **2012**, *61*, 161–168. [[CrossRef](#)]
31. Cheik, S.; Bottinelli, N.; Sukumar, R.; Jouquet, P. Fungus-growing termite foraging activity increases water infiltration but only slightly and temporally impacts soil physical properties in southern Indian woodlands. *Eur. J. Soil Biol.* **2018**, *89*, 20–24. [[CrossRef](#)]
32. Kandasami, R.K.; Borges, R.M.; Murthy, T.G. Effect of biocementation on the strength and stability of termite mounds. *Environ. Geotech.* **2016**, *3*, 99–113. [[CrossRef](#)]
33. Haverkamp, R.; Ross, P.J.; Smettem, K.R.J.; Parlange, J.Y. Three-dimensional analysis of infiltration from the disc infiltrometer: 2. Physically based infiltration equation. *Water Resour. Res.* **1994**, *30*, 2931–2935. [[CrossRef](#)]
34. Braud, I.; De Condappa, D.; Soria, J.M.; Haverkamp, R.; Angulo-Jaramillo, R.; Galle, S.; Vauclin, M. Use of scaled forms of the infiltration equation for the estimation of unsaturated soil hydraulic properties (the Beerkan method). *Eur. J. Soil Sci.* **2005**, *56*, 361–374. [[CrossRef](#)]
35. Lassabatère, L.; Angulo-Jaramillo, R.; Soria Ugalde, J.M.; Cuenca, R.; Braud, I.; Haverkamp, R. Beerkan Estimation of Soil Transfer Parameters through Infiltration Experiments—BEST. *Soil Sci. Soc. Am. J.* **2006**, *70*, 521. [[CrossRef](#)]
36. Ferrar, P.; Watson, J.A.L. Termites (isoptera) associated with dung in Australia. *Aust. J. Entomol.* **1970**, *9*, 100–102. [[CrossRef](#)]
37. Freymann, B.P.; Buitenwerf, R.; Desouza, O.; Olf, H. The importance of termites (Isoptera) for the recycling of herbivore dung in tropical ecosystems: A review. *Eur. J. Entomol.* **2008**, *105*, 165–173. [[CrossRef](#)]
38. Cheik, S.; Bottinelli, N.; Soudan, B.; Harit, A.; Chaudhary, E.; Sukumar, R.; Jouquet, P. Effects of termite foraging activity on topsoil physical properties and water infiltration in Vertisol. *Appl. Soil Ecol.* **2018**. [[CrossRef](#)]
39. Davies, A.B.; Eggleton, P.; van Rensburg, B.J.; Parr, C.L. The pyrodiversity-biodiversity hypothesis: A test with savanna termite assemblages: *Savanna termites* and fire. *J. Appl. Ecol.* **2012**, *49*, 422–430. [[CrossRef](#)]

40. Muvengwi, J.; Davies, A.B.; Parrini, F.; Witkowski, E.T.F. Contrasting termite diversity and assemblages on granitic and basaltic African savanna landscapes. *Insectes Sociaux* **2018**, *65*, 25–35. [[CrossRef](#)]
41. Ouédraogo, E.; Mando, A.; Brussaard, L. Soil macrofaunal-mediated organic resource disappearance in semi-arid West Africa. *Appl. Soil Ecol.* **2004**, *27*, 259–267. [[CrossRef](#)]
42. Filser, J.; Faber, J.H.; Tiunov, A.V.; Brussaard, L.; Frouz, J.; De Deyn, G.; Uvarov, A.V.; Berg, M.P.; Lavelle, P.; Loreau, M.; et al. Soil fauna: Key to new carbon models. *Soil* **2016**, *2*, 565–582. [[CrossRef](#)]
43. Bagine, R.K.N. Soil translocation by termites of the genus *Odontotermes* (Holmgren) (Isoptera: Macrotermitinae) in an arid area of Northern Kenya. *Oecologia* **1984**, *64*, 263–266. [[CrossRef](#)] [[PubMed](#)]
44. Diouf, M.; Miambi, E.; Mora, P.; Delgarde, S.; Rouland, C. The impact of termite sheetings age on their fungal communities. *Eur. J. Soil Biol.* **2006**, *42*, S85–S91. [[CrossRef](#)]
45. Mora, P.; Seuge, C.; Rossi, J.P.; Rouland, C. Abundance of biogenic structures of earthworms and termites in a mango orchard. *Eur. J. Soil Biol.* **2006**, *42*, S250–S253. [[CrossRef](#)]
46. Lepage, M.G. Étude de la prédation de *Megaponera foetens* (F.) sur les populations récoltantes de Macrotermitinae dans un écosystème semi-aride (Kajiado-Kenya). *Insectes Sociaux* **1981**, *28*, 247–262. [[CrossRef](#)]
47. Léonard, J.; Rajot, J.L. Influence of termites on runoff and infiltration: Quantification and analysis. *Geoderma* **2001**, *104*, 17–40. [[CrossRef](#)]
48. Elkins, N.Z.; Sabol, G.V.; Ward, T.J.; Whitford, W.G. The influence of subterranean termites on the hydrological characteristics of a Chihuahuan desert ecosystem. *Oecologia* **1986**, *68*, 521–528. [[CrossRef](#)]
49. Beven, K.; Germann, P. Macropores and water flow in soils. *Water Resour. Res.* **1982**, *18*, 1311–1325. [[CrossRef](#)]
50. Mando, A.; Miedema, R. Termite-induced change in soil structure after mulching degraded (crusted) soil in the Sahel. *Appl. Soil Ecol.* **1997**, *6*, 241–249. [[CrossRef](#)]



© 2019 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (<http://creativecommons.org/licenses/by/4.0/>).