

# Conceptual modelling of the dynamics of soil macroporosity based on an *in situ* and direct observation of major soil structuring agents during early stage of pedogenesis

## Abstract

Many factors affect soil porosity but major of them are soil fauna, plant roots and the climate. Many studies attempted to investigate the effect of these factors on the evolution of soil porosity separately. The objective of this study is, based on an *in situ* experiment including three major

porosity factors, propose a conceptual model of soil macroporosity evolution. *In situ* observation and quantification of the evolution of macroporosity under the influence of each agent separately and the three agents combined allow to propose a model for each case. Results show that the evolution of microporosity due to plant roots linear and reach its maximum at the end of the plant cycle. Earthworms create and destruct macroporosity during the up and down movement for food searching. At long term, the consequence of earthworm action results to an increase of macroporosity. Wetting and drying cycle has the same effect as earthworm. At the beginning, when soil shrinks, it leads to a creation of macroporosity that could be disturbed by swelling during soil humectation. Some soil particles migrate in the shrink and reduce the surface of microporosity. When faced to many wetting drying cycle, the surface of macropores increase during time. Mathematical algorithms and computing are necessary to formalise this model and long-term experiment is needed to validate this model.

**Keywords:** Differential equations, Soil structure, Earthworms, Casts production, Swelling Shrinkage, Plant roots

## 1. Introduction

Soil, due to its importance in furnishing ecosystem services, attracted the attention of many soil scientists for many years. Many processes occurring in soil are defined and characterized but to better understand the fate of important factors, modelling approach has been developed. Since then, a various models concerning soil processes were developed and a historical point of view of this modelling is given by Vereecken *et al.* (2016). Among these processes soil structure, defined as the rearrangement of solid soil particles got a great interest. The consequence of this arrangement is the formation

10 of aggregates and voids. Many studies highlight the main factors that in-  
11 fluence soil structure and the five major of them are i) soil fauna, ii) plant  
12 roots, iii) environmental factors, iv) inorganic binding agents and v) soil mi-  
13 croorganisms (Six *etal.*, 2004). Research have been leaded to investigate  
14 the role of different agents on the evolution of structure. Plant roots influ-  
15 ence soil structure in different ways but the most studied is by introducing  
16 organic matter. This process is widely investigated and some models have  
17 been proposed. For example, Roth-C study the soil organic matter turnover  
18 in non-waterlogged soils (Coleman and Jinkinson, 1999) and how this pro-  
19 cess is involved in soil structure (aggregation formation). Based on Roth-C,  
20 Malamoud *etal.* (2009) proposed a new model which studies the dynamics  
21 of soil structure (aggregation and porosity) named Structure-C. This model  
22 written in Matlab consist of three submodels. An aggregation submodel,  
23 organic matter submodel and porosity submodel. The porosity submodel  
24 predicts the evolution of porosity during time under specific conditions of  
25 soil organic matter turnover.

26 Apart from its action in creating structure by introducing organic matter  
27 in soil, plant roots also embed soil particles and create aggregates between  
28 what pores are created. This aspect of research is not widely explored. Nev-  
29 ertheless, the process is detailed by Jangorzo *etal.* (2015). A more complex  
30 modelling platform has been developed to help understanding and simulate  
31 major processes governing soil evolution. This platform named Virtual Soil  
32 or Vsoil (Lafolie *etal.*, 2015) is designed to facilitate modelling chemical,  
33 physical and biological interactions occurring in soils and improve the simu-  
34 lation of anthropic activities and climate change impact on the soil ecosystem  
35 services. Different contributors may work together in order to develop mod-  
36 ules that can be incorporated in the platform. Since then, module of organic  
37 matter turnover, water dynamics are developed. An important module to de-  
38 velop will be that which concern the dynamic of soil structure (porosity and  
39 aggregation). Such a module exists, example Structure-C developed based  
40 on Roth-C. It describes the evolution of soil structure under the influence  
41 of soil organic matter dynamics and the agents that condition this evolution  
42 (Malamoud *etal.*, 2009). This model does not take into account the fact  
43 that soil porosity evolves according to the influence of other major processes  
44 like earthworm activities as well as the physical process occurring in soil like  
45 swelling and shrinkage. The role of earthworms in soil structure is also widely  
46 studied and an attempt to model this process has been undertaken.

47 Another phenomenon that influences soil porosity is the wetting-drying

48 cyclerelatedtopropertyofswellingandcrackingofthesoil. Manystud-  
 49 ieshavebeenundertakentostudytheprocessofsoilcracking(Konradand  
 50 Ayad,1997;Greco,2002;BraudeauandMohtar,2006;PengandHorn,2007;  
 51 Chertkov,2012b,a)anditsimpactonsoilporosity(Coppola<sup>etal.</sup>,2012,  
 52 2015).Inarecentwork,Stewart<sup>etal.</sup>(2016)showedthatmostofmod-  
 53 elsdevelopeddonotconsidertheprocessinitswhole.However,thetotal  
 54 porosityproducedduringsoilcrackinginaswellingsoilcouldbeapportioned  
 55 inthreeparts:aggregatesporosity,shrinkagecracksandverticalsubsidence  
 56 andthenproposedaunifiedmodel.Thereisanongoingdebateconcerning  
 57 theverticalsubsidenceaspartofsoilporosityinaswellingsoil.However,we  
 58 assumethattheverticalsubsidenceistakenintoaccountwhenconsidering  
 59 thedecreaseofaggregatesporosityduringsoilcompaction. Moreover,the  
 60 modelofStewartdidnotconsidertherepeatingwetting-dryingprocessora  
 61 successiveswelling-shrinkagewhichnaturallyhappenedinaVertisol.This  
 62 whyheassumedthatthemasstheirreversible(Stewart<sup>etal.</sup>,2016)is  
 63 But,inawetting-dryingcycle,massmovementisthemajordriverofcracks  
 64 dynamic(Jangorzo<sup>etal.</sup>,2015).Nevertheless,apartofthismodelcouldbe  
 65 integratedin anewmodeldevelopmentbyconsideringitasthestateofthe  
 66 porosityinthefirstswelling-shrinkageprocess.

67 Twomechanismsdrivetheswelling-shrinkagecycleinsoil. Mechanical  
 68 loadingmechanismandthesecondassociatedwiththechangeinsuction  
 69 knownashydraulicloadingmechanism(WangandWei,2014). Mecha-  
 70 nismofreversibleswelling-shrinkagestrainandtheMechanismofirreversible  
 71 swelling-shrinkagestrain(WangandWei,2014)showsthebehavioursofsoil  
 72 volumeduringswelling-shrinkagecycle.Thisvolumeisassumedtobeirre-  
 73 versible(Tessier,1984;Jangorzo<sup>etal.</sup>,2015)innaturalconditions.Morethe  
 74 dryingisimportantlessthesoiluptakewaterduringwettingwhichmeans  
 75 thatthesoilwillswellless(volumedecrease)(CroneyandColeman,1954).  
 76 Earthworm,consideredasecosystemengineer(Jones<sup>etal.</sup>,1994)havebeen  
 77 widelystudiedinthecontextoftheirroleinsoilstructure. Theefficiency  
 78 oftheseengineers(activity),particularlyearthwormsdependsonthreefac-  
 79 tors(Kretzschmar,1991).However,earthwormsaremoreactiveinarange  
 80 ofwaterpotential,whensoilislooseanddependingontheperiodofyear.  
 81 Thisactivityconditionedtheeffectofthisengineeronsoilstructureasthe  
 82 latterdependsontheintensityofburrowing(Jangorzo<sup>etal.</sup>,2015).Ac-  
 83 tivityofearthwormisalsoconditionedbysoiltemperatureandindividual  
 84 massoftheorganism(Kaneda<sup>etal.</sup>,2016).Basedonlaboratoryandfield  
 85 experiments,manymodelshavebeendevelopedtopredicttheroleofsome

86 earthworm species in soil aggregation (e.g. Kaneda *etal.*, 2016) or in soil  
 87 carbon sequestration (e.g. Komarov *etal.*, 2017). But these models neither  
 88 they did not predict the evolution of soil porosity nor they did not consider  
 89 the synergy that exist between plant roots and earthworm in soil structure  
 90 (Zangerlé *etal.*, 2011; Jangorzo *etal.*, 2015). One of the factor that most  
 91 incite earthworm burrowing is the food seeking. However, earthworm moves  
 92 into different layer of soil to find fresh organic matter that they decompose  
 93 and translocates sometimes deeper in the soil (Jangorzo, 2013). We could  
 94 then assume that more food is available, greater is the burrowing activity.  
 95 This assumption is modelled by Daniel (1991) when he studied the effect of  
 96 food consumption on aggregates formation. The degree of soil organic mat-  
 97 ter transformation by earthworm is conceptualized and modelled by Chertov  
 98 *etal.* (2017) and Komarov *etal.* (2017). If the soil aggregate formation is  
 99 known, it is not the case for the porosity existing in and between aggre-  
 100 gates. The aim of this work is to develop a mathematical integrated model  
 101 describing the porosity dynamics in model soils “constructed Technosols”.  
 102 This multi-agents model takes into account the effect of three ranked factors  
 103 -wetting-drying cycle, plant roots and soil fauna- (Jangorzo *etal.*; 2017, data  
 104 not published) in the evolution of soil structure.

## 105 **2. Model theory**

### 106 *2.1. General model of soil porosity dynamics*

107 We hypothesize that the changes of macroporosity  $P$  with time  $t$ ,  $\frac{dP}{dt}$ ,  
 108 depends on the variation of macroporosity due to i) soil moisture  $f(\Theta)$ ,  
 109 ii) the population dynamics of roots  $f(r)$ , and iii) the earthworm activity  $f(w)$  resp  
 110 ectively, as well as a porosity disappearance term  $D_p$ :

$$\frac{dP}{dt} = f(\Theta) + f(r) + f(w) - D_p. \quad (1)$$

### 111 *2.2. Changes in porosity due to variations in soil humidity (wetting and dry- 112 ing cycles)*

113 In clayey soils like Vertisols or in Technosols made of swelling  
 114 materials like paper or iron industry sludges, there are changes in soil  
 115 porosity with the variations of soil humidity (Peng and Horn, 2007; Huotet  
 116 *al.*, 2014). This shrinkage and swelling behaviour creates macroporosity in soil  
 when shrinks and decreases soil porosity during swelling processes.

117 When a soil is set up and watered, the first process that affects its porosity  
 118 is water loss. The behaviour of water movement in soil has been largely stud-  
 119 ied, particularly the process of swelling-shrinkage (Smiles, 2000). According  
 120 to its texture, soil swells when it is wetted which causes an increase of vol-  
 121 ume, a decrease of bulk density, and an increase in macroporosity (Smiles,  
 122 2000). Conversely, when the soil dries, it shrinks and its volume decreases  
 123 which leads to an increase of soil bulk density (Peng and Horn, 2007), a de-  
 124 crease of aggregate porosity (microporosity), and an increase of shrinkage  
 125 cracks (macroporosity) (Stewart *et al.*, 2016). But, as the volume of cracks  
 126 is important, we assume that the total porosity of the soil increases as soon  
 127 as the water potential increases.

128 Stewart *et al.* (2016) have proposed and validated a unified model to  
 129 represent the variations of the different types of porosity with the changes in  
 130 soil humidity. They consider three types of porosity:

131 In absence of any other agents, here we assume that the effect of micro-  
 132 organisms is negligible, the main process that governs porosity formation is  
 133 water flow expressed by the swelling-shrinkage phenomenon. We used the  
 134 equation of Stewart *et al.* (2016) to model the variation of porosity due to  
 135 one wetting-drying cycle for cracks porosity:

$$\phi_{\text{crack}} = (\phi_{\text{pedon}} - \phi_{\text{min}}) \left( \frac{1 - U^q}{1 - \varepsilon U^q} \right), \quad (2)$$

136 with  $U = \Theta / \Theta_{\text{max}}$  and other parameters in Stewart *et al.* (2016).

137 Linking Stewart *et al.* (2016) equation with our framework:

$$f(\Theta) = \frac{\partial \phi_{\text{crack}}}{\partial t}. \quad (3)$$

138 We thus need to derivate  $\phi_{\text{crack}}$  with time  $t$ :

$$f(\Theta) = \frac{[-qU(t)^{q-1}U'(t)(1 + \varepsilon U(t)^q)] - [q\varepsilon U(t)^{q-1}U'(t)(1 - U(t)^q)]}{(1 - \varepsilon U(t)^q)^2} \quad (4)$$

139 and  $U'(t) = \partial \Theta / \partial t$

$$f(\Theta) = (1 - \varepsilon) \frac{[-qU(t)^{q-1}] - [2q\varepsilon U(t)^{2q-1}]}{(1 - \varepsilon U(t)^q)^2} \frac{\partial \Theta}{\partial t}, \quad (5)$$

140 where  $\varepsilon$  and  $q$  are fitting parameters.

141 **2.3. Changes in porosity due to the dynamics of roots population**

142 Rootshavethreedifferentimpactsonporosity:

- 143 •firstly,theyfillsomeavailableporositywhiletheyareappearingand  
144 growingandthenfreeupsomeporeswhenageinganddying(bynar-  
145 rowing)anddisappearing(bydegrading);thisaffectsmacroporosity;
- 146 •secondly,whentheyaregrowing,theycompacttheneighbouringsoil  
147 (rhizosphere)leadingtoadecreaseofwhatMalamoud<sup>etal.</sup>(2009)  
148 calledaggregatesporosityorthesoilmicroporosity. Theintensityof  
149 compactiondependsoneithercracksexistbeforetheappearanceof  
150 rootsornot;however,Jangorzo<sup>etal.</sup>(2015)showedthatrootsprefer-  
151 entiallyusedexistingporeslikecrackswhengrowing.
- 152 •third,whenrootsaredegraded,theyenterintothesoilorganicmatter  
153 turnoverwhichalsohasanotherimpactofsoilporosityasdemonstrated  
154 bymanyauthors(e.g.ColemanandJinkinson,1999;Malamoud<sup>etal.</sup>,  
155 2009).

156 Inthismodelweonlyconsidertheeffectofrootsonthedynamicsofmacro-  
157 porositybyinfilling.

158 Jangorzo(2013)andJangorzo<sup>etal.</sup>(2015)haveshownthattheimpact  
159 ofrootsonmacroporositydependsontheagecategoryoftheroots.When  
160 youngrootsappearandthenmature,theyaregrowingthusleadingtoa  
161 decreaseinporosity. Duringageinganddegradationafterdeath,theyare  
162 freeingssomespacewhich,byconsequenceleadtoanincreaseinporosity.  
163 Thuswhenrootsageandbecomeoldrootstheporosityincreases.Thus,if  
164 weconsiderthetotalsurfacefillbyrootswecandistinguishthreedifferent  
165 agecategories:

- 166 • surfacewithyoungroot $R_y(t)$ ,
- 167 • surfacewithmatureroot $R_m(t)$ ,
- 168 • surfacewitholdroot $R_o(t)$ ,

169 Therelationshipbetweenthosethreecategoriesoftherootpopulationcan  
170 bemodelledasshowninFigure1.Hereweconsiderthereisnodeathof  
171 youngandmatureroots.Thecomputationoftherootsurfacedynamicsis

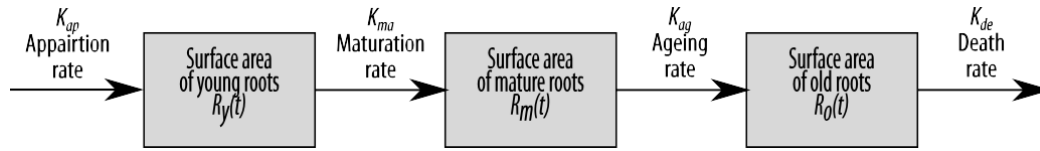


Figure 1: Conceptual model of the dynamic of roots surface.  
Possibility to consider only two categories for the root population.  $\kappa$  function of time?

172 given by the following equations, with  $R_y(t)$ ,  $R_m(t)$  and  $R_o(t)$ , the surface  
173 areas occupied by young, mature, and old roots, respectively:

$$\frac{\partial R_y}{\partial t} = \kappa_{ap}(t) - \kappa_{ma}(t) \quad (6)$$

$$\frac{\partial R_m}{\partial t} = \kappa_{ma}(t) - \kappa_{ag}(t) \quad (7)$$

$$\frac{\partial R_o}{\partial t} = \kappa_{ag}(t) - \kappa_{de}(t), \quad (8)$$

174 with  $\kappa_{ap}(t)$  the apparition rate of young roots,  $\kappa_{ma}(t)$  the maturing rate of young  
175 roots,  $\kappa_{ag}(t)$  the ageing rate of mature roots, and  $\kappa_{de}(t)$  the death rate of old  
176 roots. No death of young and mature roots is considered. Thus  
177 the changes in the total surface occupied by roots is:

$$\frac{\partial R}{\partial t} = \kappa_{ap}(t) - \kappa_{de}(t) \quad (9)$$

178 The general equation for the changes in porosity due to the population dynamics of roots  
179 is function of  $\kappa_{ap}(t)$ ,  $\kappa_{ma}(t)$ ,  $\kappa_{ag}(t)$ , and  $\kappa_{de}(t)$ :

$$f(r) = -\kappa_{ap}(t) - \kappa_{ma}(t) + \kappa_{ag}(t) + \kappa_{de}(t). \quad (10)$$

180 This equation means that for each surface increase of young and  
181 mature roots there is a direct proportional loss of porosity. Reversibly, for  
182 each surface decrease of old roots by death, there is a direct proportional  
183 increase of porosity. When mature roots are ageing and become old, the consequenc  
184 eis the increase of porosity.  $dt_1$  is the time elapsed between the moment  
185 the first root appears and the moment it starts narrowing.  $dt_2$  is the time elapsed be  
186 tween  $dt_1$  and the lifetime of the plant or roots.  $dt_3$  is the time  
187 elapsed between  $dt_2$  and the moment all dead roots disappeared or are  
188 degraded. This time is soil moisture and biological activity dependent.

As young and

189 mature roots have the same effect on soil porosity, we include the young roots  
 190 and mature by extending the time. The previous equation then becomes: -

$$f(r) = -K_{ma}(dt_1) + K_{ag}(dt_2) + R_{de}(dt_3) \quad (11)$$

191 From equations 10, 6, 7, 8, ??, we can deduce:

$$f(r) = kR_o(t) - \kappa_{ap} \quad (t) - \frac{\partial R_m}{\partial t}. \quad (12)$$

192  $R_v(t)$ ,  $R_m(t)$  and  $R_o(t)$  and their time derivatives can be obtained from  
 193 the experimental data at each time step. If no experimental data are available  
 194 general forms for  $\kappa_{ap}(t)$ ,  $\kappa_{ma}(t)$ , and  $\kappa_{ag}(t)$  must be proposed.

#### 195 **2.4. Changes in porosity due to earthworm activity**

196 Jangorzo et al. (2013) showed that the intensity of earthworm  
 197 activity increases soil porosity.  
 198 This activity is a function of i) the number of earthworm, ii) the quantity of organic  
 199 matter and iii) the soil moisture. Many authors have described the soil organic  
 200 matter turnover and models were developed. Among these models we can  
 201 announce the RothC (Coleman and  
 202 Jinkinson, 1999), StructureC (Malamoud et al., 2009) and a module in Vsoil (Lafolie  
 203 et al., 2015). The general equation of porosity evolution according to  
 the intensity of activity is as follows:

$$204 \quad F(w) = k_{I_w} \times I_w \quad (13)$$

$$I_w = f(I_w^{\max} \times nb_w, \Theta, MO) \quad (14)$$

205  
 206 Where  $F(w)$  is the rate of porosity changes due to worm activity;  $I_w$  is the total intensity  
 207 of worm activity;  $I_w^{\max}$  is the maximal intensity of worm activity;  $k_{I_w}$  is the  
 208 conversion rate between the intensity of worm activity and the proportion  
 209 of pore surface;  $nb_w$  is the number of earthworm;  $\Theta$   
 is the soil moisture or water potential and  $MO$  is the soil organic matter.

##### 210 **2.4.1. Worm activity in relation with soil humidity**

211 In relation to earthworm development and activity, soil humidity is better  
 212 expressed as water potential due to the physiological characteristics of these  
 213 organisms (Kretzschmar, 1991; Kretzschmar and Bruchou, 1991; Holmstrup,  
 214 2001; Moreau-Valancogne et al., 2013). Some studies showed the influence

215 of water potential on earthworm development (Kretzschmar and Bruchou,  
 216 1991; Holmstrup, 2001; Eriksen-Hamel and Whalen, 2006) and activity of  
 217 earthworm (Kretzschmar, 1991; Hindell *etal.*, 1994; Daniel *etal.*, 1996). All  
 218 those articles assess the development or the activity of *Aporrectodea* spp.,  
 219 which are endogeic or anecic earthworms. Other factors can be tested in  
 220 parallel (compaction, temperature). The earthworms activity is assessed via  
 221 their cast production. Kretzschmar (1991); Hindell *etal.* (1994); Daniel *etal.*  
 222 (1996) show the same pattern for the relationship between cast production  
 223 and water potential: no or a negligible cast production for water potentials  
 224 more negative than a base potential  $\Psi_b$  and then a linear increase of cast  
 225 production with increasing water potential until  $\Psi=0$ . The estimated values  
 226 of  $\Psi_b$  vary with species but indicate a narrow moisture tolerance range (Ta-  
 227 ble 2.4.1).  $\Psi_b$  seem to vary with the ecological type: around -20 to -40 kPa  
 228 for anecic and around -10 kPa for endogeic.

Reference	Species	Ecological type	$\Psi_b$ (kPa)
Kretzschmar (1991)	<i>Aporrectodea longa</i>	anecic	-
40 Hindell <i>etal.</i> (1994)	<i>Aporrectodea rosea</i>	endogeic	-
10 id.	<i>Aporrectodea caliginosa</i>	endogeic	-10
id.	<i>Aporrectodea caliginosa</i> <i>saf. trapezoides</i>	endogeic	-5
Daniel <i>etal.</i> (1996)	<i>Aporrectodea nocturna</i>	anecic	-20

Table 1: Value of the base water potential,  $\Psi_b$ , measured by different authors.

229 The relationship between the activity of *Lumbricus castaneus* (epi-anecic  
 230 worm, tested in Jangorzo (2013)) and the water potentials show the same  
 231 pattern as identified for anecic earthworms in the literature. So we hypothe-  
 232 sised that equations 13 and 14 can be inferred for *L. castaneus*. We assume  
 233 that the normalised earthworm activity is proportional to the normalised cast  
 234 production. Many equations were used to model the evolution of earthworms  
 235 activity as function of water potential but we are interested in that issuing  
 236 the cast production. For example Daniel *etal.* (1996) used an exponential  
 237 equation to predict the evolution of cast production by *Aporrectodea nocturna*.

$$C^*(P) = h e^{(Pi)} \quad \text{for } -0.06 \text{ MPa} \leq P \leq 0 \text{ MPa} \quad (15)$$

238 where  $C^*(P)$  is the transformed rate of cast production as a function of wa-  
 239 ter potential in MPa and  $h$  and  $i$  are constants. They assumed that cast

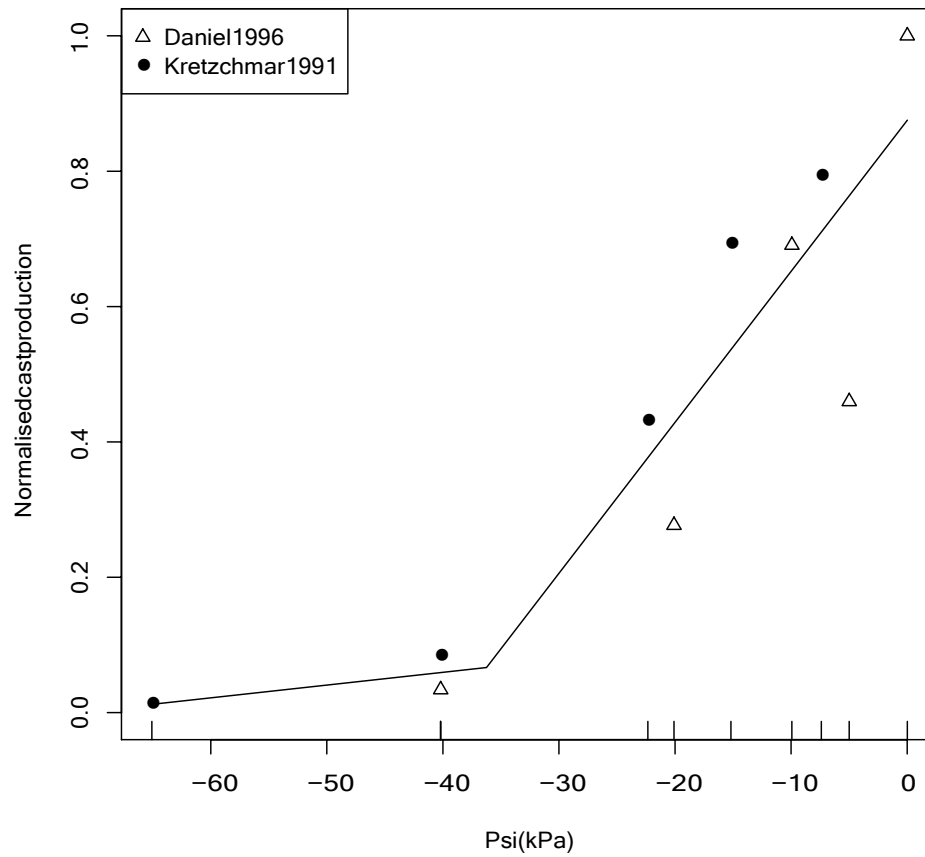


Figure 2: Model of the normalised cast production as a function of water potential for anecic earthworms. The data were extracted from Kretzschmar (1991) and Daniel *et al.* (1996). The fitted values are shown as points.

240 increased exponentially with the increase of water potential. Kaneda *etal.*  
 241 (2016) on the other hand used the modified Gompertz function expressed as  
 242 follows:

$$f(\Psi) = \zeta \exp^{-\left(\exp^{-(\Psi+\eta)/\theta}\right)} \quad \text{for} \quad -90.6 \text{ kPa} < P < -2 \text{ kPa} \quad (16)$$

243 where  $f(P)$  stands for the soil aggregate formation rate-modifying factor for  
 244 soil moisture as a function of water potential  $\Psi$ , in kilopascal,  $\zeta$ ,  $\eta$  and  $\theta$  are  
 245 constant experimental coefficients determining slope of the curve. They use  
 246 this function rather than Daniel's because their data showed a levelling-off  
 247 near the water potential 0 kPa. In contrary, experimental data obtained  
 248 by image analysis showed the increase of earthworm activity when water  
 249 potential increases (Jangorzo, 2013; Jangorzo *etal.*, 2015). We therefore  
 250 used the Daniel's exponential equation to predict the evolution of earthworm  
 251 activity. This relation could be formalized by the normalized cast production  
 252 equation:

$$I_w(\Psi) = h e^{(\Psi/\eta)} \quad (17)$$

253 Here we consider the casts production as a result of the intense activity of  
 254 earthworm and this is proportional to the porosity production. However,  
 255 when earthworms burrow, they dig, ingest soil particles mixed with organic  
 256 matter that they excrete like casts. So, the cast produced are proportional to  
 257 the void created in the soil. Say  $k_w$  the coefficient indicating the conversion  
 258 between  $I_w$  and porosity. We assume that  $0 < k_w < 1$  as it is a ratio between  
 259 the  $I_w$  and the  $I_w^{max}$ . It means that not all the casts produced are equivalent  
 260 to the porosity created.

#### 261 2.4.2. Worm activity in relation with number of earthworms

262 Most of experiments were undertaken with a single individual of earth-  
 263 worm. The effect of many individuals is not arithmetic but it has been shown  
 264 that more the number of earthworm is high, greater is the intensity and so  
 265 the porosity created (Jangorzo *etal.*, 2015). However, equation  
 17 could be rearranged as follows:

$$I_w(\Psi) = n b_w \times h e^{(\Psi/\eta)} \quad (18)$$

267 Where  $n b_w$  is the number of earthworms

268 **2.4.3. Worm activity in relation with organic matter content**

269 Organic matter is used by earthworms for feeding and they burrow in  
 270 order to find food. In this model, we assume that the organic matter content  
 271 is constant as the quantity introduced in the system does not vary. However,  
 272 the variation of earthworm activity as a function of organic matter is equal to  
 273 zero as well as that of soil temperature.

274 Combining the equations 13, 14 and 18 we can write the general equation  
 275 of porosity evolution as a function of earthworm activity as follows:

$$F(w) = k_i \times n_b w \times h e^{(\psi_i)} \quad (19)$$

276 **2.5. Porosity disappearance**

277 We suggest that a proportion of the total porosity disappeared at each  
 278 timestep: Two processes induce porosity disappearance according to our  
 279 assumption: i) water loading charged during wetting or watering and soil  
 280 fauna (earthworms) when burrowing.

$$D_P = f(\Theta, I_w) \quad (20)$$

281 **2.5.1. Porosity disappearance due to watering**

282 When water is added in a structured soil, according to the stability of  
 283 the soil, this water induces extinction of some aggregates. During water flow,  
 284 these particles are transported deep in the soil through porosity. At the limit of  
 285 water infiltration, these particles are deposited creating fillings. This process  
 286 leads to the decrease of porosity. However, as successive watering without  
 287 bioturbation leads undoubtedly to a disappearance of soil macropores. Say  
 288  $P_0$  the porosity of the system at  $T_0$ . After the first watering, the porosity  
 289 decreases into  $P_1$  at  $T_1$ . After  $n$  series of watering we have a porosity of  $P_n$ .  
 290 The total porosity that disappears is:

$$\Delta(P_\Theta) = \sum_{1}^n (P_{n+1} - P_n)$$

291 Where  $P$  is the porosity,  $n$  the number of watering series and the  $\Theta$  means that the de-  
 292 crease is due to watering.

293 **2.5.2. Porosity disappearance due to earthworm activity**

294 In presence of earthworms, we assume that the decrease of porosity due  
 295 to watering is null as during burrowing earthworms are willing to extract the

296 filings. The only action that decreases porosity is the deposit of casts in  
 297 burrows. This action depends on how many times an earthworm passes in  
 298 a burrow. Say  $P_0$  the porosity of the system at  $T_0$ . After the first passage  
 299 (burrowing), the porosity decreases into  $P_1$  at  $T_1$ . After  $n$  series of burrowing  
 300 we have a porosity of  $P_n$ . The total porosity that disappears is:

$$\Delta(P_w) = \sum_1^n (P_{n+1} - P_n)$$

301 Where  $P$  is the porosity,  $n$  the number of earthworm activity (burrowing)  
 302 and the  $w$  means that the decrease is due to burrowing.

303 The solution of equations 21 and 22 could be obtained using the least mean  
 304 square method as experienced by Daniel (1991); Daniel *etal.* (1996). The  
 305 principle of least mean square method is formalized by (Stocker *etal.*, 2002)  
 306 which stipulates that: Given a function  $y(x) = y(x; a)$  dependant on a pa-  
 307 rameter  $a$  and  $n$  couples of values  $(x_i; y_i)$ , the optimal parameter  $a$  is defined  
 308 by the Gauss function as follows:

$$\sum_i^n (y_i - y(x_i; a))^2 = Min_a$$

309 The idea behind this function is that it minimise the variable  $a$  and tends to  
 310 a residual. In the case of our study, it means that the disappearance of  
 311 porosity due to watering and earthworm activity is minimised. However the  
 312 porosity decreases at each time step and at each watering, but due to the  
 313 action of plants roots or burrowing, the decrease is compensate latter  
 314 particularly by earthworms. This is why, unless this temporary decrease, the  
 315 surface area of porosity increases with time.

## 316 2.6. Submodels integration

317 In our approach, we assumed that different factors acted successively in  
 318 soil structuring. Beginning by the soil moisture which is the limiting factor  
 319 to biological activity. Second plants find ideal conditions to develop and then  
 320 soil fauna like earthworms. To maintain an equilibrium, a proportion of soil  
 321 structure disappears due to the same structuring agents, primarily water.

322 Rearranging we obtained:

$$\frac{dP}{dt} = [(I_w^{max} \times nb_w)^{-zv} \times \left\{ \begin{array}{l} 0, 0224\psi + 0, 8764, \text{ for } \psi < \Psi_b \\ 0, 0019\psi + 0, 1379, \text{ for } \psi > \Psi_b \end{array} \right\}] \quad (24)$$

$$+ [(3 + \eta_{ar}) - \frac{(D_{byr} + D_{bar} + D_{bor})}{D_s}] \quad (25)$$

$$+ [\int_1^n (\frac{\varepsilon + 1}{\varepsilon + \Theta^{-q}})] - D_P \quad (26)$$

### 323 3. Model implementation

324 An experiment was set up in a climate chamber where the effect of  
 325 wetting-drying cycle, plant roots and earthworm activity on soil structure  
 326 has been studied. This experiment was run during 14 months; images were  
 327 recorded using Soilinsight (INRA, 2015), a device of *insitu* monitoring  
 328 of soil structure developed by Jangorzo (2013) and soil structure parameters  
 329 were quantified by image analysis (Jangorzo *etal.*, 2013, 2014, 2015). Three  
 330 replicates of each factor were realized. The experimental design was fully de-  
 331 scribed in Jangorzo (2013).

#### 332 3.1. Initial conditions

333 At the beginning of the experiment  $t=0$ , the surface areas of porosity  
 334 and aggregates were quantified but the surface area of roots was null as well  
 335 as the earthworm activity.

$$P(t=0) = P_0 \quad (27)$$

$$r(t=0) = 0 \quad (28)$$

$$l_w(t=0) = 0 \quad (29)$$

#### 336 3.2. Swelling-Shrinkage parameters

337 The mesocosm used in this experiment were filled with a constructed  
 338 Technosol 500  $\mu$ m sieved. The soil was made by a mixture of Treated Industrial  
 339 Soil (TIS) and Paper mill-Sludge (PS). At the top of the cosm a thin  
 340 layer of compost was layed out. Cosms were first moistened by capillarity  
 341 uptake until saturation and gravimetric water content at field capacity was  
 342 measured by weighing. Then four wetting-drying cycles were applied. After  
 343 saturation, the system was let for drying until 20%. Then it is wetted un-  
 344 til saturation and the excess water was collected as leachate. Regularly the

345 cosmswereweightedtodeterminethewatercontent.Basedontheseinfor-  
 346 mationtherelationbetweenmatricpotentialandgravimetricwatercontent  
 347 wasestablishusinghydrus. Imageswereanalysedandcracksparameters  
 348 werequantified.

### 349 3.3. Plant roots parameters

350 Inothercosms,fourseedsof*Lupinus albus*weresownandrootdevel-  
 351 opmentismonitoredasdescribedbyJangorzo<sup>etal.</sup>(2015). Fourdays  
 352 afterthebeginningoftheexperiment,thefirstrootsappeared.Say $D_1$ this  
 353 date. Rootscontinuegrowingduringtimeandthesurfaceiscalculatedby  
 354 imageanalysis. Bycomparingdifferentimagesgenerated,weidentifythe  
 355 timewhenrootsstopgrowing.Say $D_2$ thisday.Thenthe $dt_1$ isdeducedas  
 356 follows: $dt_1 = D_2 - D_1$ . Thevariationofrootssurfaceaccordingtotime  
 357  $\frac{\partial R_m}{\partial t}$ iscalculated.Theexperimentcontinuedrunningandat amomentplant  
 358 rootsweredeadunlessthesystemwaswatered.Say $D_3$ thisdate.Thetime  
 359 elapsedduringrootsnarrowing $dt_2 = D_3 - D_2$ isdetermined.Knowingthat  $\frac{\partial R_o}{\partial t}$   
 360 thetotalsurfaceofdeadrootsisquantified,thenthevariationrate  $\frac{\partial R_o}{\partial t}$  is  
 361 calculated.Therateofdeadrootsdegradationisdeterminedassoonasroots  
 362 disappeared.Thiscoefficientcanalsobedeterminedusingtheequationof  
 363 LafolieusedinVsoil.

### 364 3.4. Earthworms activity parameters

365 Incosmscontainingplantroots,wereintroducedsixindividuals( $nb_w$ )of  
 366 *Lumbricus castaneus*.Thesoilmoisturewasmaintainedinarangebetween  
 367 60%fieldcapacityand80%fieldcapacity.Usinghydrus,thismoisturecon-  
 368 tentisequivalenttoacorrespondingsuction $\Psi$ .Asinthepreviousystems,  
 369 imagesweregeneratedeachtwohourswhatallowsustomonitortheearth-  
 370 wormactivity.Thedifferentoperationsrecordedarethefollowing:  
 371 actionsmadebyaworm: i)creatingaburrow,ii)fillingaburrow,iii)en-  
 372 largingburrow;iv)unchangedburrow{(Jangorzo<sup>etal.</sup>,2015).Thenthe  
 373 intensityofearthwormsactivity/ $w$ wasdeterminedknowingthenumberof  
 374 individualsandthetimeelapsed.Moreover,imagewereanalysedtoquantify  
 375 theevolutionofsoilporosity.

## 376 4. Conclusion and perspectives

377 Basedonaninnovativeexperimentalset-upofsoilobservationandquan-  
 378 tification,weproposedaconceptualmodelofsoilporositydynamicsunder

379 the influence of three aggregation factors. For this, three agents were exper-  
 380 imented: wetting-drying cycle for environmental factor, *Lupinus albus* for  
 381 plant roots and *Lumbricus castaneus* for soil fauna. To validate this model,  
 382 it is important to set up a new experiment and resolve the different equations  
 383 by computing them.

384 **Appendix A. Variable definitions**

385 All the variables used in the model are summarised in Table A.1.

Var.	Definition	Units	Source
$a_{ry}$	Apparition rate of young roots	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2} \times \text{h}^{-1}$	
$A(t)$	Surface area of the soil aggregates in proportion to the total surface of the picture	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2}$	Obtained by image analysis on experimental data (Jangorzo, 2013)
$d_{rd}$	Degradation rate of dead roots	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2} \times \text{h}^{-1}$	
$D_p$	Destruction rate of the porosity	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2} \times \text{h}^{-1}$	Literature review on soil compaction and collapse, experimental framework
$\vartheta$	Rate of porosity changes due to soil humidity (wetting and drying cycles)	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2} \times \text{h}^{-1}$	
$f(w)$	Rate of porosity changes due to worm activity	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2} \times \text{h}^{-1}$	
$f(r)$	Rate of porosity changes due to root dynamics	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2} \times \text{h}^{-1}$	
$I_w$	Total intensity of worm activity	Number of worm actions per hour	Computed by image analysis from experimen-

			<b>taldata</b>
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**Table A.1—continued from previous page**

<b>Var.</b>	<b>Definition</b>	<b>Units</b>	<b>Source</b>
$I_w^{\max}$	Maximal intensity of worm activity	Number of worm actions per hour and per individual	Computed by image analysis from experimental data Jangorzo(2013)
$k$	Factor of porosity destruction (collapsed due to humidity)	Dimensionless	Calibration or literature review
$k_{I_w}$	Conversion rate between the intensity of worm activity and pore surface proportion	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2}$ per worm action	
$k_{r_d}$	Conversion rate between the surface of degraded roots and the pore surface	Dimensionless	Calibration or computation from experimental data Jangorzo(2013)
$k_{r_y}$	Conversion rate between the surface area of new young roots and the pore surface area (proportion of porosity occupied by the surface area of new young roots)	Dimensionless	Calibration or computation from experimental data Jangorzo(2013)
$k_{r_{y \rightarrow o}}$	Conversion rate between the surface area of new old roots and the pore surface area	Dimensionless	Calibration or computation from experimental data Jangorzo(2013)
$k_{r_{o \rightarrow d}}$	Conversion rate between the surface area of new dead roots and the pore surface area	Dimensionless	Calibration or computation from experimental data Jangorzo(2013)

**Table A.1—continued from previous page**

<b>Var.</b>	<b>Definition</b>	<b>Units</b>	<b>Source</b>
$k_{\theta}$	Conversion rate between humidity variations and pores surface proportion	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2}$ per soil humidity unit	Literature review on shrinking, experimental framework
$m_{r_d}$	Death rate/mortality of old roots	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2} \times \text{h}^{-1}$	
MO	Soil organic matter content	$\text{g} \cdot \text{g}^{-1}$	Literature review
$nb_w$	Number of earthworms	Individuals	
$P(t)$	Surface area of the porosity ( $>50 \mu\text{m}$ ) in proportion to the total surface of the picture	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2}$	Obtained by image analysis on experimental data (Jangorzo, 2013)
$P_0$	Surface area of the porosity ( $>50 \mu\text{m}$ ) in proportion to the total surface of the picture at the initial time $t=0$	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2}$	Obtained by image analysis on experimental data (Jangorzo, 2013)
$R(t)$	Total surface area occupied by roots in proportion to the total surface of the picture	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2}$	Obtained by image analysis on experimental data (Jangorzo, 2013)
$R_o(t)$	Surface area with old roots in proportion to the total surface of the picture	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2}$	Obtained by image analysis on experimental data (Jangorzo, 2013)
$R_m(t)$	Surface area with mature roots in proportion to the total surface of the picture	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2}$	Obtained by image analysis on experimental data (Jangorzo, 2013)

**Table A.1—continued from previous page**

<b>Var.</b>	<b>Definition</b>	<b>Units</b>	<b>Source</b>
$R_y(t)$	Surface area with young roots in proportion to the total surface of the picture	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2}$	Obtained by image analysis on experimental data (Jangorzo, 2013)
$t$	Time	Hour (h)	Experimental time step is 2h for total length of the experiment of 14 months
$v_{ry}$	Ageing rate of young roots (conversion in old roots)	$\text{mm}^2 \times 10^{-2}$ $\text{mm}^{-2} \times \text{h}^{-1}$	
$K_{d_r}$	Coefficient of root degradation (proportion of root degradation in relation to the population of dead roots)	$\text{h}^{-1}$	Calibration or literature review
$\Theta$	Soil humidity or matric potential ( $\Psi$ )	Units in relation with the chosen variables	Literature review

Table A.1: Description of the used variables. Var. : Variables.

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