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Effect of thermo-coupled processes on the behaviour of a clay barrier submitted to heating and hydration

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ABSTRACT

The storage of high level radioactive waste is still an unresolved problem of the nuclear industry, being geological disposal the most favoured option and, naturally, the one requiring the strongest geo-mechanical input. Most conce tual designs for the deep geological disposal of nuclear waste envisage placing the canisters containing the waste horizontal drifts or vertical boreholes. The empty space surrounding the canisters is filled by an engineered barr often made up of compacted swelling clay. In the barrier and the near field, significant thermo-hydro-mechanical (THI phenomena take place that interact in a complex way. A good understanding of THM issues is, therefore, necessate to ensure a correct performance of engineered barriers and seals. The conditions of the bentonite in an engineer barrier for high-level radioactive waste disposal are being simulated in a *mock-up* heating test at almost scale, at the premises of CIEMAT in Madrid. The evolution of the main Thermo-Hydro-Mechanical (THM) variables of this to are analysed in this paper by using a fully coupled THM formulation and the corresponding finite element coor Special emphasis has been placed on the study of the effect of thermo-osmotic flow in the hydration of the clay barr at an advanced staged of the experiment.

Key words: expansive clays, heating test, nuclear waste disposal, thermo-hydro-mechanical coupled analysis.

INTRODUCTION

Conceptual designs for deep geological disposal of nuclear waste generally envisage placing the canisters containing the waste in horizontal drifts or vertical boreholes. The empty space surrounding the canisters is filled by an engineered barrier often made up of compacted bentonite in the form of blocks (Fig. 1). This clay-based isolation system has the multiple purposes of providing mechanical stability for the waste canister (by absorbing stresses and deformations); serving as a buffer around

it; sealing discontinuities in the emplacement bo and drifts; and delaying the water flow from t rock. The barrier behaviour is highly complex it involves coupled Thermo-Hydro-Mechanical phenomena that take place due to the simultaneouing (generated by the radioactive waste) and hy (due to the contribution of the surrounding rock barrier. A good understanding of THM issues is fore, necessary to ensure a correct performance of neered barriers and seals.

The work presented here started in the fram of FEBEX, which was a project for the study near field for a high level nuclear waste reposi-

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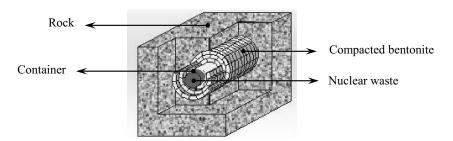


Fig. 1 – Scheme of the engineered barrier.

in situ test, under natural conditions and at full scale (Grimsel, Switzerland); a mock-up test, at almost full scale (CIEMAT, Madrid); and a series of laboratory tests to complement the information from the two large-scale tests. This paper focus mainly on the mock-up experiment, which avoids many of the uncertainties arising from the natural system. This is achieved mainly due to the good control of the initial and boundary conditions of the experiment. Two electrical heaters (simulating canisters containing heat-emitting waste) were placed in the centre of a steel cylinder 6 m long and with inner diameter of 1.62 m. The space between the heaters and the steel cylinder was filled with a 0.64 m-thick engineered barrier made up of compacted FEBEX bentonite (Fig. 2). The barrier is hydrated uniformly from all around the cylinder with an applied water pressure of about 0.5 MPa. Simultaneously, the barrier was heated maintaining a constant temperature of 100°C at the contact between heaters and bentonite. This very well instrumented test has provided the opportunity to study in detail the evolution of the main THM variables for a period of almost ten years.

The evolution of the main THM variables of the *mock-up* test has been analysed using the CODE_BRIGTH program (Olivella et al. 1996), which is a finite element code developed to handle coupled THM problems in porous media. An unexpected decay in the rate of hydration observed at advanced stages of the test has affected significantly the predictions performed using 'standard THM models'. The difference between model outputs and experimental results affect the reliability of the long-term predictions, especially the one related to the time required to reach the full saturation of the clay

the 'unexpected' behaviour of the clay barrier. A preliminary study has envisaged three main phenomena that are identified as: thermo-coupled effects, threshold gradient and evolution of the micro-fabric (Sánchez 2004). This paper focus on the analysis of the first phenomenon.

The paper is organized as follows: first the expansive clay used in the analysis is introduced; then, the general framework adopted to analyse the heating test is presented; afterwards, the main results of the numerical analyses are introduced; and finally, the paper closes with the main conclusion of this work.

MATERIALS

The clay barrier of the mock-up test has been built with the FEBEX bentonite, which was selected by ENRESA (the Spanish Agency for Radioactive Waste Management) as suitable material for the backfilling and sealing oh high level nuclear waste repositories. The FEBEX bentonite was extracted from the Cortijo de Archidona deposit (Almería, Spain). The processing of the material at the factory consisted in its disaggregation and gently grinding, drying at 60°C and sieving by 5 mm. For the large-scale tests of the FEBEX Project, the bentonite blocks were manufactured by uniaxial compaction of the granulated clay with its hygroscopic water content, at dry densities close to 1.7 g/cm³ (ENRESA 2000). The physico-chemical properties of the FEBEX bentonite, as well as its most relevant thermo-hydro-mechanical and geochemical characteristics, are summarised in the final reports of the project (ENRESA 2000, 2006) and are shown in detail in ENRESA (1998), Villar (2002), Lloret



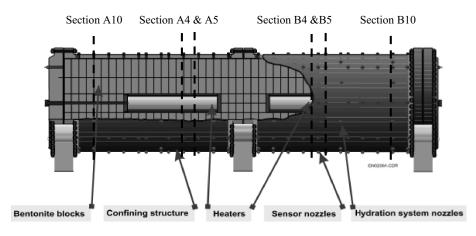


Fig. 2 – Layout of the large scale heating test.

The FEBEX bentonite has a content of montmorillonite higher than 90 percent. Besides, it contains variable quantities of quartz, plagioclase, K-feldspar, calcite and opal-CT (cristobalite-trydimite). The cation exchange capacity (CEC) varies from 96 to 102 meq/100g, and the major exchangeable cations are: Ca (35-42 meq/100g), Mg (31-32 meq/100g), Na (24-27 meq/100g) and K (2-3 meq/100g).

The liquid limit of the bentonite is 102 ± 4 percent, the plastic limit is 53 ± 3 percent, the specific gravity 2.70 ± 0.04 , and 67 ± 3 percent of particles are smaller than 2μ m. The hygroscopic water content in equilibrium with the laboratory atmosphere is 13.7 ± 1.3 percent. The value obtained for the external specific surface area using the BET technique is 32 ± 3 m²/g and the total specific surface area obtained using the hygroscopicity method is about 725 m²/g. The analysis of the mercury intrusion data reveals that the intra-aggregate pores (smaller than 0.006ν m) represents the 73-78 percent of total pore volume when the bentonite is compacted at a dry density of 1.7 g/cm³.

THEORETICAL FORMULATION AND COMPUTER CODE

An appropriate way to integrate the coupled THM phenomena and their mutual interactions that take place in expansive clays submitted to simultaneous heating and hydration is by means of a properly designed theoret-

thermo-hydraulic tests. A brief description of the lation is presented below.

The theoretical framework is composed of main parts: balance equations, constitutive equations, equilibrium restrictions. The framework is form using a multi-phase, multi-species approach. The scripts identify the phase ('s' for solid, 'l' for liq'g' for gas). The superscript indicates the spector mineral, 'w' for water and 'a' for air). The phase may contain water and dissolved air, and phase may be a mixture of dry air and water vaporation.

BALANCE EQUATIONS

Equations for mass balance were established for the compositional approach, which consists of bathe species rather than the phases. The mass balt solid present in the medium is written as:

$$\frac{\partial}{\partial t} (\rho_s (1 - \phi)) + \nabla \cdot (\rho_s (1 - \phi) \dot{\mathbf{u}}) = 0$$

where ρ_s is the mass of solid per unit volume of s is the porosity; \mathbf{j}_s is the flux of solid, t is time, \mathbf{v} divergence operator and $\dot{\mathbf{u}}$ is the solid velocity ve

Water is present in liquid and gas phases. T mass balance of water is expressed as:

$$\frac{\partial}{\partial t} \left(\theta_l^w S_l \phi + \theta_g^w S_g \phi \right) + \nabla \cdot \left(\mathbf{j}_l^w + \mathbf{j}_g^w \right) = f$$

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 $(\alpha = l, g)$; \mathbf{j}_l^w and \mathbf{j}_g^w denote the total mass fluxes of water in the liquid and gas phases (water vapour) with respect to a fixed reference system; and f^w is an external supply of water.

Dry air is considered as a single species and it is present in liquid and gas phases. The total mass balance of dry air is expressed as:

$$\frac{\partial}{\partial t} \left(\theta_l^a S_l \phi + \theta_g^a S_g \phi \right) + \nabla \cdot \left(\mathbf{j}_l^a + \mathbf{j}_g^a \right) = f^a \qquad (3)$$

where θ_l^a and θ_g^a are the masses of dry air per unit volume of liquid and gas, respectively; S_α is the volumetric fraction of pore volume, occupied by the alpha phase $(\alpha = l, g)$; \mathbf{j}_l^a and \mathbf{j}_g^a denote the total mass fluxes of dry air in the liquid and gas phases with respect to a fixed reference system; and f^a is an external supply of dry air.

Regarding the thermal problem, the equilibrium between the phases is assumed. Therefore, the temperature is the same for all the phases. Consequently, only one equation is needed for the energy balance. The total internal energy, per unit volume of porous media, is obtained adding the internal energy of each phase corresponding to each medium. Applying the balance equation to this quantity, the following equation is obtained:

$$\frac{\partial}{\partial t} \left(E_s \rho_s (1 - \phi) + E_l \rho_{lp} S_l \phi + E_g \rho_g S_g \phi \right) + \nabla \cdot \left(\mathbf{i}_c + \mathbf{j}_{Es} + \mathbf{j}_{El} + \mathbf{j}_{Eg} \right) = f^E$$
(4)

where E_s is the solid specific internal energy; E_l and E_g are specific internal energies corresponding to liquid and gas phases, respectively; ρ_s is the solid density; ρ_l and ρ_g are the liquid and gas phase densities; \mathbf{i}_c is the conductive heat flux; \mathbf{j}_{Es} is the advective energy flux of solid phase with respect to a fixed reference system; \mathbf{j}_{El} and \mathbf{j}_{Eg} are the advective energy flux of liquid and gas phases, respectively, with respect to a fixed reference system; f^E are the energies supply per unit volume of medium.

Finally, the balance of momentum for the porous medium reduces to the equilibrium equation in total stresses:

$$\nabla \cdot \boldsymbol{\sigma} + \mathbf{b} = 0 \tag{5}$$

that are negligible in comparison with the stress terms. Through an adequate constitutive model, the equilibrium equation is transformed into a form expressed in terms of the solid velocities, fluid pressures and temperatures. The assumption of small strain rate is also made.

CONSTITUTIVE EQUATIONS

The constitutive equations establish the link between the independent variables (or unknowns) and the dependent ones. A more detailed explanation of the mathematical formulation in general and constitutive equations in particular is given in Gens et al. (1998). The constitutive laws and the corresponding parameters presented in the section correspond to the case use in the numerical analysis coded as 'OBC' (Operational Base Case). Mode details in section 4.2.

Hydraulic problem

The advective flows of the phases (*i.e.*, liquid or gas), with respect to the solid phase, are assumed to follow Darcy's law:

$$\mathbf{q}_{\alpha} = -\mathbf{K}_{\alpha} (\nabla P_{\alpha} - \boldsymbol{\rho}_{\alpha} \mathbf{g}); \quad \alpha = l, g$$
 (6)

where P_l and P_g are liquid and gas pressure, respectively; ρ_l is the liquid density; ρ_g is the gas density; \mathbf{g} is the gravity vector; and \mathbf{K}_{α} is the permeability tensor of the alpha phase $(\alpha = l, g)$, which is given by:

$$\mathbf{K}_{\alpha} = \mathbf{k} \frac{k_{r\alpha}}{\mu_{\alpha}}; \quad \alpha = l, g \tag{7}$$

The intrinsic permeability tensor (\mathbf{k}) depends on the pore structure of the porous medium. $k_{r\alpha}$ is the value of relative permeability that controls the variation of permeability in the unsaturated regime, and μ_{α} denotes the dynamic viscosity. α may stand for either l or g, depending on whether liquid or gas flow is considered.

The saturated permeability to deionised water, \mathbf{k} (7), of samples of untreated FEBEX bentonite compacted at different dry densities, can be related with porosity through a modified Kozeny's law according to:

$$\mathbf{k} = k_0 \frac{\phi^3}{(1 - \phi_0)^2} \mathbf{I}$$
 (8)



adopted together with experimental data obtained for FEBEX bentonite permeated at room temperature with deionised water. A k_0 =1.9 e^{-21} m² for a reference porosity (ϕ_0) of 0.40 has been adopted. It was checked that the permeability of the FEBEX bentonite to granitic water (salinity 0.02%) is analogous to that obtained for deionised water (Villar 2002).

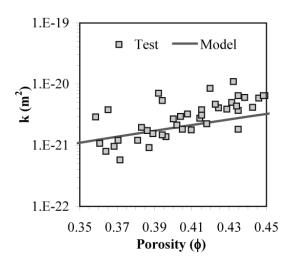


Fig. 3 – Variation of saturated permeability with porosity. Experimental data and adopted model for the intrinsic permeability law.

The relative permeability of liquid (k_{rl}) phases is made dependent on S_e (effective degree of saturation) according to:

$$k_{rl} = S_e^n \tag{9a}$$

$$S_e = \frac{S_l - S_{lr}}{S_{ls} - S_{lr}}$$
 (9b)

where S_l is degree of saturation; S_{lr} and S_{ls} are residual and maximum degree of saturation, respectively; and n is a material parameter. Based on back-analyses (performed with CODE_BRIGHT) of isothermal infiltration tests, a cubic law (*i.e.* n = 3) has been adopted for this law (Lloret et al. 2002, Pintado et al. 2002).

The retention curve of the bentonite was determined in samples compacted to different dry densities (Lloret et al. 2004). The volume of the samples remained constant during the determinations, since they were confined modified van Genuchten law has been adopted to the dependence of the degree of saturation on su

$$S_e = \left[1 + \left(\frac{s}{P_0} \right)^{\frac{1}{1 - \lambda_0}} \right]^{-\lambda_0} f_d;$$

where:
$$f_d = \left(1 - \frac{s}{P_d}\right)^{\lambda_d}$$

where $s(s=P_g-P_l)$ is the suction; P_0 is a parelated to the capillary pressure (the air entry value) λ_0 is a parameter that controls the shape of the cure Genuchten 1978). The function f_d is included to more suitable values at high suctions; P_d is a parelated with the suction at 0 degree of saturation; is a model parameter (when it is null, the original is recovered). Figure 4 presents the retention cure tained with these expressions for different values density together with the experimental data. The aparameters are: $P_0 = 28$ MPa, $\lambda_0 = 0.18$, $P_d =$

$$P_0 = P_{T_0} \frac{\sigma_T}{\sigma_{T_0}}$$

where the surface tension (in N/m) has been o fitting values of surface tension with the follow pression (Olivella 1995):

$$\sigma_T = 0.03059 \exp\left(\frac{252.93}{273.15 + T^0}\right)$$

By varying P_0 in accordance with this expression pendence of the retention curve with temperature troduced. Therefore, suction will decrease with a ature for a given degree of saturation (Olivella and 2000). Experimental results have shown that, it cases, the influence of temperature on retention of small (Lloret et al. 2004).

The molecular diffusion of vapour water phase is governed by Fick's law, through:

$$\mathbf{i}_{g}^{w} = -\mathbf{D}_{g}^{w} \nabla \omega_{g}^{w} = -(\phi \rho_{g} S_{g} \tau D_{m}^{w} \mathbf{I} + \rho_{g} \mathbf{D}_{g}^{\prime}) \nabla \alpha$$

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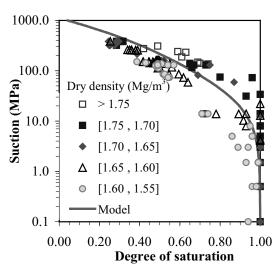


Fig. 4 – Retention curve adopted in the analyses, together with the experimental data for FEBEX bentonite (symbols).

dispersion tensor. \mathbf{D}_{m}^{w} is the molecular diffusion coefficient of vapour in gas. A tortuosity factor of 0.8 has been adopted based on numerical back analysis of heat flow tests at constant overall water content (Lloret et al. 2002, Pintado et al. 2002).

Thermal problem

The Fourier's law has been adopted for the conductive flux of heat. Thermal conductivity depends on the hydration state of the clay and is expressed by a variant of the geometric mean:

$$\mathbf{i}_c = -\lambda \nabla T \tag{14}$$

where

$$\lambda = \lambda_{sat}^{s_e} \lambda_{dry}^{(1-s_e)} \tag{15}$$

and

$$\lambda_{dry} = \lambda_{solid}^{(1-\phi)} \lambda_{gas}^{\phi}; \quad \lambda_{sat} = \lambda_{solid}^{(1-\phi)} \lambda_{liq}^{\phi}$$
 (16)

where λ is the thermal conductivity. The thermal conductivity $\lambda[W/m \cdot K]$ (14) of the compacted bentonite at laboratory temperature is related to the degree of saturation (S_e) (15). Figure 5 presents the experimental values obtained for the FEBEX bentonite together with

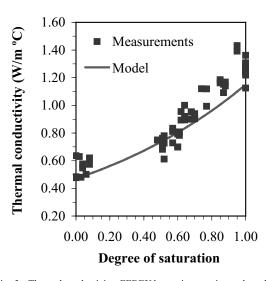


Fig. 5 – Thermal conductivity: FEBEX bentonite experimental results (symbols) and model fitting.

Mechanical problem

The mechanical constitutive law adopted is the Barcelona Basic Model (BBM). This model extends the concept of critical state to the unsaturated conditions, including the dependence of yield surface on suction. Two stress variables are considered: the net stresses ($\sigma-P_gm$) and capillary suction. Net stress is the excess of total stress over gas pressure. If full saturation is achieved, net mean stress becomes effective stress. The main equations are introduced as follows, more details are provided in the Appendix. For simplicity, net stress will also be denoted by σ . It is assumed that the generalised yield surface depends not only on stresses and history variables, but on temperature and suction, as follows:

$$F = \frac{3J^2}{g_y^2} - L_y^2(p + P_s)(P_o - p) = 0$$
 (17)

where M is the slope of the critical state; p_o is the apparent unsaturated isotropic pre-consolidation pressure; g is a function of Lode's angle; and p_s considers the dependence of shear strength on suction and temperature (see Appendix). The hardening law is expressed as a rate relation between volumetric plastic strain and the



where e is void ratio; κ is the elastic compression index for changes in net mean stress; and $\lambda_{(0)}$ is the compression index for changes in net mean stress for virgin states of the soil in saturated conditions. More details about BBM can be found in the elsewhere (Alonso et al. 1990, Gens 1995).

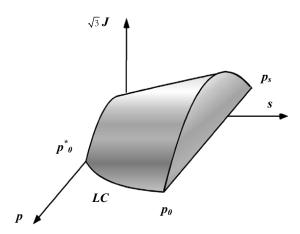


Fig. 6 – A three-dimensional representation of the *BBM* yield surface.

Due to the high compaction the bentonite blocks have been subjected to, the description of the behaviour of the material inside the yield surface is particularly important. The variation of stress-stiffness with suction and, especially, the variation of swelling potential with stress and suction have been considered (Sánchez and Gens 2006). The resulting elastic model is the following:

$$\dot{\varepsilon}_v^e = \frac{\kappa_i}{(1+e)} \frac{\dot{p}}{p} + \frac{\kappa_s}{(1+e)} \frac{\dot{s}}{(s+0.1)} + (\alpha_0 + \alpha_2 \Delta T) \dot{T}$$
(19)

$$\dot{\varepsilon}_s^e = \frac{\dot{J}}{G} \tag{20}$$

where

$$\kappa_i = \kappa_i (1 + \alpha_s s) \tag{21}$$

$$\kappa_s = \kappa_{s0} (1 + \alpha_{sp} \ln p / p_{ref})$$
 (22)

is the young module. Figure 7 presents the

$$G = \frac{E}{2(1+v)} \tag{23}$$

initial drying (in this case close to the 200 MP then, a wetting. In the same figure, the results swelling pressure tests coded as SP1 and SP2 are Test SP1 and SP2 are described in detail in Llor (2003). The model results can be considered s tory, the stress path is quite well reproduced, as the predicted value of the swelling pressure. The model parameter are summarised in Table I. More can be found in Sánchez and Gens (2006).

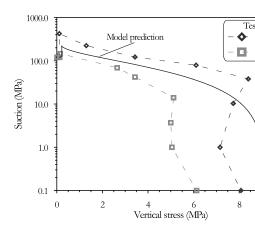


Fig. 7 – Computed stress path for swelling pressure tests BBM. SP1 and SP2 experimental results are provided for cor

TABLE I

Model parameters related to the mechanical pr
lem (defining the Barcelona Basic Model).

κ	0.04	κ_{S}	0.25
ν	0.4	α_{is}	-0.003
α_{sp}	-0.147	α_{ss}	0.00
α_o	$1.5 \times 10^{-4} (^{\circ}\text{C}^{-1})$	α_2	0.00
λ_o	0.15	p_{c}	0.10 MPa
p_o^*	14 MPa	α	0.395
r	0.75	β	0.05
M	1.5	T_{O}	20°C
k	0.1	ρ	0.2

EQUILIBRIUM RESTRICTIONS

Other types of relationships that relate depende

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in phases (liquid, gas). This assumption is sufficiently adequate because these chemical processes are fast compared to the transport processes that take place in porous media and, for this reason; they are not rate-controlled. In this problem, the concentration of water vapour in the gas phase is controlled by the psychometric law, and the solubility of dry gas in water is given by Henry's law (Olivella et al. 1994).

COMPUTER CODE

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The mathematical formulation presented above has been implemented in the finite element computer program CODE-BRIGHT (Olivella et al. 1996). CODE-BRIGHT is a tool designed to analyse numerically coupled THM problems in geological media. One unknown "state variable" is associated with each of the balance equations presented in Section 3.1 (Equations 1 to 5). The unknowns are obtained by solving the system of PDE's (Partial Differential Equations) numerically in a coupled way. From state variables, dependent variables are calculated using the constitutive equations or the equilibrium restrictions. The numerical approach can be viewed as divided into two parts: spatial and temporal discretization. Galerkin finite element method is used for the spatial discretization, while finite differences are used for the temporal discretization. The discretization in time is linear, and an implicit scheme is used. Finally, since the problem presented here is non-linear, the Newton-Raphson method was adopted as an iterative scheme. More details can be found in Olivella (1995) and CODE BRIGHT User's Manual (2008).

RESULTS

INTRODUCTION

A 2-D axis-symmetric model has been adopted for the numerical analysis of the *mock-up* test. The study has been focused on two characteristic cross-sections: 'hot' and 'cold' cross-section. The first one corresponds to a section in the heater zone (*i.e.*, Section A4 and B4, Fig. 2). The second one is located away from the heater (Section A10 and B10, Fig. 2). In this way, two different patterns of behaviour can be examined. A complete

The adopted initial and boundary conditions are introduced as follows.

INITIAL CONDITIONS

The initial global degree of the saturation of the *mock-up* test just before the switch on of the heater was 71.50%. The same initial degree of the saturation has been assumed in the model. An initially uniform temperature of 20°C is assumed in the entire domain. This is consistent with CIEMAT Report (2002) data. As for the mechanical problem, a hydrostatic value of 0.11 MPa has been adopted for the initial stress, approximately equal to the weight of the bentonite in the mid diameter of the buffer.

Boundary conditions

A water pressure of 0.55 MPa is applied in the external boundary in accordance with CIEMAT Report (2002) data. For the mechanical problem, a stress free outer boundary has been prescribed. Finally, for the thermal boundary conditions at a radius equal to 0.15 m (radial coordinate of the heater elements), the applied conditions are as follows: i) 0-6 days: constant power (250 W/heater); ii) 6 days – t_{100} : constant power (500 W/heater); iii) t_{100} : $T = 100^{\circ}$ C. Where t_{100} is the time at which the temperature reaches 100° C at some point in the bentonite (15.6 days for the analysis presented herein). On the external boundary, the following radiation condition has been applied:

$$j_e = \gamma_e(T^0 - T) \tag{24}$$

where j_e is the heat flow, T^0 is the prescribed temperature ($T^0 = 20^{\circ}$ C) and γ_e is the radiation coefficient. A coefficient $\gamma_e = 5$ has been adopted. This value has been adjusted in order to ensure the prescribed condition (T^0) on the outer boundary.

BASE CASE ANALYSIS

The heating and hydration of the test, known as the 'operational stage', started in February 1997 and is ongoing nowadays. The parameters of the model coded as 'OBC' (Operational Base Case) were determined at the



b) present the evolution of the global variables of the problem, which are: a) the heater power and b) the water intake. Figures 9 a) and b) show the time evolution of temperature at different points of the barrier, whereas Figures 9 c) and d) present the evolution of the relative humidity at different radii of the barrier for the two selected sections. Finally, Figures 10 a) and b) show the time evolution of stresses in selected positions of the barrier (P_T : tangential stress, P_Z : longitudinal stress and P_R : radial stress). In the following sections, the main results (experimental and numerical) of the *mock-up* test are analysed considering the different problems (thermal, hydraulic and mechanical) in a separately way.

Thermal problem

The barrier is heated maintaining a constant maximum temperature of 100°C at the contact between heaters and bentonite. It can be stated that the model yields good results regarding the thermal problem. This is reflected either in the evolutions of a global variable (*i.e.*, power emitted by the heaters on Fig. 8 a), or in the measures of the local ones (such as the evolution of temperatures in different points of the test Figs. 9 a) and b). It can be noted that there are some small differences in the temperature field, especially in zones close to the heater. They may be an effect of the lower thermal conductivity in the discontinuity that may exist between heater and bentonite.

Hydraulic problem

The barrier is hydrated uniformly from the periphery, maintaining a practically constant water pressure of about 0.5 MPa. Figure 8 b) shows the time evolution of the water entry in the test and the rate of the water intake as well. Good model predictions can be observed at the beginning of the experiment, but overestimation of the water entry is computed at advances stages. Figures 9 a) and b) show the evolution of the relative humidity. It can be seen that until the day 900 of the test, approximately, the model offers a good reproduction of the hydraulic variables. Up to that time, the overall behaviour

slow hydration. The behaviour of zones close to a of 0.37 m shows an initial wetting, due to the castion of the water vapour coming from the inner and, then, a drying.

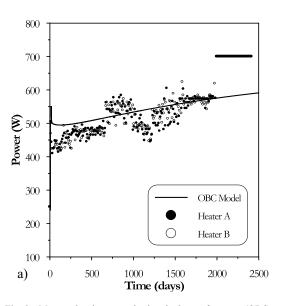
From about the day 900, some differences served between the hydraulic behaviour of the tomodel predictions. In terms of the water entry, the results move away progressively, in relation to perimental data. This fact can be clearly detecterms of the rate of the water intake (Fig. 8 b)), in the test data values (from the day 900 of the todergo an important reduction in comparison walues of the 'OBC' model. A recovery of the imental values can be seen when a non-planne took place (the overheating episode, close to to 1381; for more details, see Sánchez et al. 2005). It to the tendency observed previous to the overheating recovered, that is, a clear slowing down of the rate water intake compared to the model predictions.

Regarding the evolution of the relative hum the heater regions Sections A4 and B4, Figure significant reduction in the rate of the relative ity increase can be observed. The effects of th heating can be clearly noted in the 'hot cross-se especially just after this episode. It seems that the sient period induced by the overheating was prafinished after the day 1700. On the other hand, 'cold cross-section' is examined, the hydration prare less perceptible (Sections A10 and B10, Figure Note that some small differences between most test results can also be detected in these section the day 2000 (approximately).

Mechanical problem

Similar trends to the ones observed in the hyproblem appear in the mechanical problem. The up to the day 900 approximately, in sections in heater, there is a good agreement between predictional and observations. From that moment on, meatresses respond to the modifications of the hypattern, and a tendency to keep constant values of can be observed. This is ascribed to the strong

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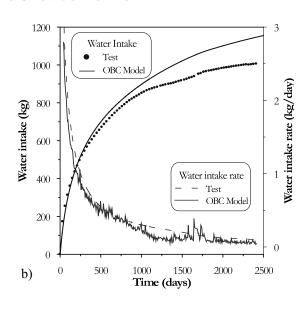


Fig. 8 – Measured and computed values in the mock-up test (OBC model). a) Evolution of heater. b) Evolution of water entry in the mock-up test.

sections. In sections far from the heaters, the values and tendency registered previous to this event have been recovered practically immediately (Section A10 and B10), whereas in sections involving heaters, a more marked influence on the evolution of the stress field can be observed. In these sections, the tendency to keep a practically constant stress level (observed previous to this episode) has been recovered after the overheating, but the stresses are now lower compared with the ones measured before this event (Sections A6 and B6). Arguments based in double structure concepts have been presented in Sánchez et al. (2005) to explain this behaviour.

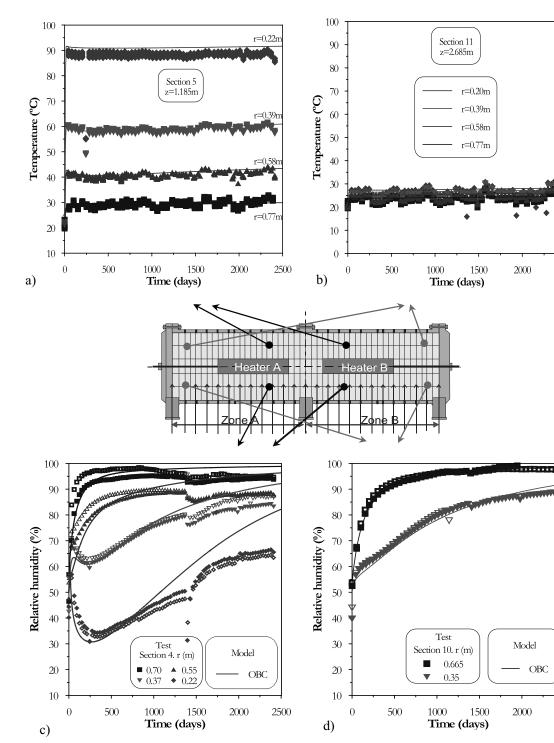
THERMO-COUPLED EFFECTS

The understanding and explanation of the apparent decay in the rate of the barrier hydration are crucial aspects concerning the performance of reliable long-term predictions. Firstly, it was explored whether minor modifications of the constitutive laws or their parameters were possible to explain and reproduce more closely the global evolution of the test. A wide sensitivity study was carried out to that end, and it was unable to obtain a set of constitutive laws and materials parameters (with physical).

to the barrier has also been investigated. It was explored whether there were any artefact in the hydration system, or a geotextile blockage had occurred that could affect the normal hydration of the barrier. It was confirmed experimentally that there was no obstruction in the hydration system or geotextile, and that the water intake was nearly uniform over the entire hydration front. Discarding those problems, research was focused on the identification of other processes that could be responsible for the apparent slowing down of barrier hydration. Three main physical phenomena, not included in the original THM formulation presented in Section 3, have been considered in the analysis of the mock-up test (Sánchez and Gens 2006). These phenomena are as follows: a) the presence of thermo-osmosis coupled processes; b) the existence of a threshold gradient in Darcy's law; and c) the change of intrinsic permeability due to the modifications of microfabric that occur during hydration of the compacted clay. This paper focused on the analysis of the first one. a more detailed analysis of the mock-up test is presented in Sánchez and Gens (2006).

The hydraulic gradient is the main force influencing the movement of water in the soils. It is, however, not the only one. Figure 11 presents all the different





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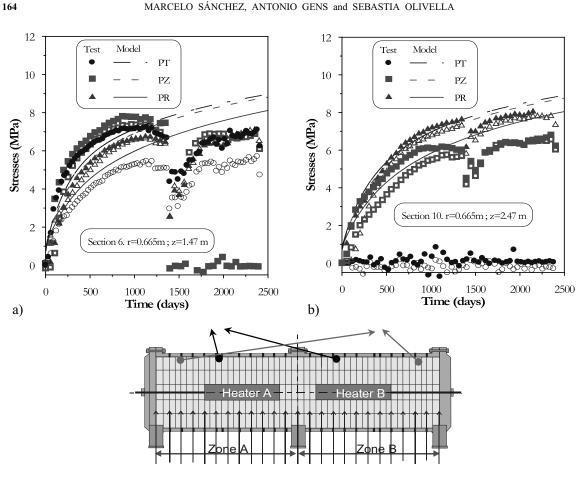


Fig. 10 - Evolution of stresses in the mock-up test. a) Sections A6-B6, and b) A10-B10. Observed versus computed values (OBC model).

is used for the diagonal terms associated to the direct flow phenomena, and the name 'effect' is reserved for the non-diagonal ones, called 'coupled processes' (i.e., Bear 1972, de Marsily 1986, Mitchell 1993). The 'phenomenological coefficient' that links each flow with the corresponding driving force must be measured experimentally (Djeran 1993, Mitchell 1993, Soler 1999).

Generally, the non-diagonal coefficients are relatively small and negligible compared to the diagonal terms, and the coupled process can be ignored. However, there are certain problems in which, due to their particular conditions, the coupled process may play a more influential role. The thermal conditions imposed on the mock-up test correspond to a practically constant thermal gradient during the test and, in consequence, a

tial high suction, but diminishes as the hydration of the barrier progresses. The liquid flows associated to these two gradients have opposite directions. Generally, the advective flow due to the pressure gradients (Darcy's law) is the dominant flow. However, at advanced stages of the test (when the hydraulic gradient becomes smaller), it is possible that the flow of liquid due to the coupled phenomena (thermo-osmotic flow) could have a noticeable effect on the behaviour of the test in the hot sections, causing a tendency to slow down the hydration in the hot zones close to the heaters.

The magnitude of this effect will depend on the relation between the gradients of the two mechanisms and on the relationship between the phenomenological coefficients associated to either flow. On the other hand,



	Gradients		
Flow	Hydraulic Head	Chemical Concentration	Temperature
Fluid	Darcy's Law (Hydraulic Conduction)	Chemical Osmosis	Thermo Osmosis
Solutes	Ultra Filtration	Fick's Law (Diffusion)	Soret Effect (Thermal Diffusion)
Heat	Thermo Filtration (Isothermal Heat Transfer)	Dufour Effect	Fourier's Law (Thermal Conduction)

Fig. 11 – Direct and coupled flow processes.

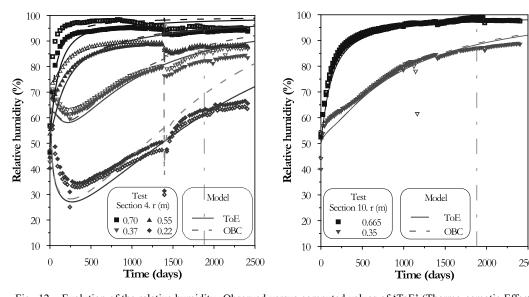


Fig. 12 – Evolution of the relative humidity. Observed versus computed values of 'ToE' (Thermo-osmotic Effect and 'OBC' (Operational Base Case) models. a) Sections A4-B4. b) Sections A10-B10.

A series of analyses including the thermo-osmosis effect were carried out to explore the possibility that this coupled process may help the explanation of the *mock-up* test observations. It is necessary to mention the lack of experimental data for FEBEX bentonite concerning the phenomenological coefficient associated to the thermo-osmotic flow. The thermo-osmotic constant adopted in the analysis is 2.73×10^{-13} m²/K/s and falls in the range of possible values found in the literature for low permeability materials (Soler 1999, Djeran 1993).

Figure 12 shows, with solid lines, the results obtained considering the Thermo-osmotic Effects (ToE) for

close to the heater. As expected, the responses tions A10 and B10 do not change under this lesis. Finally, Figure 13 presents the evolution water intake. It has been shown that, in generate the inclusion of the thermo-osmotic flows in the assimproves the performance of the model. Howeversults presented should be considered only as a cise performed with the aim of investigating the thetical response of the barrier under these assumuntil more experimental evidence on the phenomenance available.

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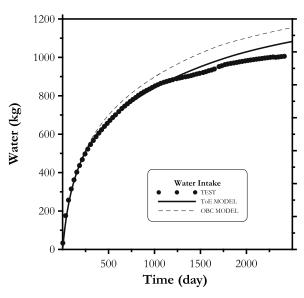


Fig. 13 – Evolution of the water intake. Observed versus computed values of 'ToE' (Thermo-osmotic Effect) and 'OBC' (Operational Base Case) models.

carried out under controlled thermo-hydra-mechanical conditions at CIEMAT laboratory (Madrid, Spain). This research has been performed in the context of the European projects FEBEX and NF-PRO. The main components of THM formulation adopted for the numerical analysis of this experiment have also been introduced. The comprehensive laboratory works carried out in the context of the FEBEX project have been used to identify the main parameters of the THM constitutive laws required for the modelling. An initial model, coded as OBC, was developed at the beginning of the heating and hydration of the *mock-up* test. Good model predictions have been obtained for, approximately, the first three years of the experiment. However, the results obtained are not totally satisfactory because the modelling over-predicts the hydration rate of the barrier at advanced stages of the hydration. A tendency to keep a constant and very low rate of the water intake and nearly constant values of the main variables (relative humidity and stresses) at different radii of the barrier can be observed in all the sections of the test, with a more marked trend in the cross-sections that involve heater. Changes in the

the mock-up test could be ascribed to other THM and chemical processes or phenomena, which were not contemplated in the original OBC model. To extend the capability of the original THM formulation, three main phenomena have been considered: a) the presence of thermo-osmosis coupled processes; b) the existence of a threshold gradient in Darcy's law; and c) the change of intrinsic permeability due to the modifications of microfabric that occur during hydration of the compacted clay. Each one of these phenomena could explain, to some extent, with physical basis, the unforeseen response of the clay barrier (Sánchez and Gens 2006). This paper has been focused on the effect of thermo-osmotic flows in the behaviour of the clay barrier. Improvements in the model outputs have been obtained under this hypothesis. Finally, it is important to mention that each of these developments (i.e., a; b; and c) does not exclude the others, and it is possible that an explanation for the whole behaviour of the clay barrier requires the combinations of some of them. In contrast, it must be pointed out that one problem associated with these new developments is the scarcity of experimental data available for calibrating



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RESUMO

O armazenamento de rejeitos altamente radioativos é ainda um problema em aberto na área de engenharia nuclear sendo os sítios geológicos ainda a opção mais favorável e naturalmente aquela que demanda maior conhecimento na área de geomecânica. A maioria dos projetos conceituais de armazenamento do lixo nuclear objetiva a alocação de cilindros que contêm os rejeitos em poços verticais ou horizontais. O espaço vazio que circunda os cilindros é preenchido por uma barreira de engenharia na maioria dos casos composta por uma argila expansiva. Na barreira e na vizinhança fenômenos significativos de acoplamento termo-hidro-mecânico(THM) tomam lugar e interagem entre si de maneira bastante complexa. Um bom entendimento dos tópicos em acoplamento THM é portanto necessário para assegurar a boa performance das barreiras de engenharia. As condições da bentonita em uma barreira de engenharia para o armazenamento do lixo nuclear foram simuladas em um teste de calor experimental nas premissas do CIEMAT em Madrid. A evolução das variáveis que governam o acoplamento termo-hidro-mecânico neste teste são analisadas neste artigo utilizando uma formulação totalmente acoplada juntamente com o código computacional de elementos finitos. Ênfase especial é dada ao estudo do escoamento termo-osmótico na hidratação da barreira argilosa no estágio avançado do experimento.

Palavras-chave: argilas expansivas, teste de calor, armazenamento do lixo nuclear, análise do acoplamento termo-hidromecânico.

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APPENDIX

The BBM yield surface (F_{LC}) is given by (17) and the plastic potential (G) is expressed as

$$G = \alpha 3J^{2} - \left[\frac{g(\theta)}{g(-30^{\circ})}\right]^{2} M^{2}(p+p_{s})(p_{0}-p) = 0$$
(A1)

where α is determined according to Alonso et al. (1990). The dependence of the tensile strength on suction and temperature is given by

$$p_s = ks \ e^{-\rho \Delta T} \tag{A2}$$

where k and ρ are model parameters. The dependence of p_0 on suction is given by

$$p_0 = p_c \left(\frac{p_{0T}^*}{p_c}\right)^{\frac{\lambda_{(0)} - \kappa}{\lambda_{(s)} - \kappa}} \tag{A3a}$$

$$p_{0T}^* = p_0^* + 2(\alpha_1 \Delta T + \alpha_3 \Delta T | \Delta T|)$$
 (A3b)

where p_c is a reference stress, α_1 and α_3 are models parameters. $\alpha_{(s)}$ is the compressibility parameter for changes in net mean stress for virgin states of the soil. This parameter depends on suction according to

$$\lambda_{(s)} = \lambda_{(0)} \left[r + (1 - r) \exp(-\zeta s) \right] \tag{A4}$$

where r is a parameter that defines the minimum soil compressibility (at infinite suction), and ζ is a parameter that controls the rate of decrease of soil compressibility with suction.