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Thermal and Hygric Expansion of High Performance Concrete

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The linear thermal expansion coefficient of two types of high performance concrete was measured in the temperature range from 20 °C to 1000 °C, and the linear hygric expansion coefficient was determined in the moisture range from dry material to saturation water content. Comparative methods were applied for measurements of both coefficients. The experimental results show that both the effect of temperature on the values of linear thermal expansion coefficients and the effect of moisture on the values of linear hygric expansion coefficients are very significant and cannot be neglected in practical applications.

Keywords: concrete, linear thermal expansion, linear hygric expansion, moisture, temperature.

1 Introduction

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Length changes of porous materials can be affected by changes of both temperature and moisture content [1]. Hygric expansion is often not taken into account in practical measurements and calculations, although a higher content of moisture, particularly in the liquid state, can lead to hygric stresses which are at least comparable with thermal stresses [2]. In this paper, we measure both the linear thermal expansion coefficient and the linear hygric expansion coefficient of two types of high performance concrete.

2 Thermal and hygric expansion parameters

The infinitesimal change of length due to change of temperature is defined by

$$l_T = l_0 \alpha_T \mathrm{d}T \;, \tag{1}$$

where l_0 is the length at the reference temperature T_0 , and α_T is the linear thermal expansion coefficient. In an analogous way, the linear hygric expansion coefficient α_u can be defined

$$dl_u = l_0 \alpha_u du , \qquad (2)$$

where

$$u = 100 \ \frac{m_m - m_d}{m_d} \tag{3}$$

is the moisture content in %, m_m is the mass of the moistened material, m_d the mass of the dried material.

Applying, in the first approximation, the superposition principle to the length changes due to temperature and moisture, we arrive at

$$dl = dl_{\rm T} + dl_u = l_0 (\alpha_T dT + \alpha_u du).$$
(4)

Defining the relative elongation as

 (\mathbf{n})

$$=\frac{\Delta l}{l} = \frac{1}{l_0} \int_{l_1}^{l_2} dl, \qquad (5)$$

we obtain

$$(u,T) = {}_{T}(u_{0},T) + {}_{u}(u,T_{0}) =$$
$$= \int_{T_{0}}^{T} \left(\frac{\partial}{\partial T}\right)_{u} dT + \int_{u_{0}}^{u} \left(\frac{\partial}{\partial u}\right)_{T} du = \int_{T_{0}}^{T} {}_{\alpha}{}_{T} dT + \int_{u_{0}}^{u} {}_{\alpha}{}_{u} du.$$
(6)

3 Experimental procedure

In measuring the high-temperature linear thermal expansion coefficient, we employed an experimental device developed in our laboratory (see [3]). The device consists of cylindrical, vertically placed electric furnace with two bar samples located in the furnace. The first sample is the measured material, the second sample is a reference material with the known dependence of the thermal expansion coefficient on temperature. The length changes of the samples are measured mechanically outside the furnace by thin ceramic rods which pass through the furnace cover and are fixed on the top side of the measured sample. These ceramic rods pass by an indefinite temperature field, so that their elongation cannot be determined mathematically and a comparative method of determining the elongation of the rod is used.

In measuring the linear hygric expansion coefficient, we measured the length changes by the Carl Zeiss optical contact comparator with $\pm 1 \mu m$ accuracy, and the mass changes were determined by the Sartorius electronic balance with $\pm 1 \text{ mg}$ accuracy. The basic set of points $l_i(m_i, T_i)$ necessary for determining the $\alpha(u)$ function was obtained in this way. The experiments were performed under isothermal conditions, with $T = (25.0 \pm 0.5) \,^{\circ}\text{C}$.

First, the dependence of the relative elongation on moisture content was determined. The samples were first dried in a hot-air dryer at T = 110 °C, then moistened in the environment with a known relative humidity of 40 %, moistened to full water saturation, and dried first in natural way in the laboratory and finally in the hot-air dryer at T = 110 °C. Both length and mass changes were simultaneously measured during the moistening and drying processes.

4 Materials and samples

The experimental work was done with two types of high performance concrete used in nuclear power plants: Penly concrete and Temelin concrete.

Penly concrete was used for a concrete containment building in a nuclear power plant in France (samples were obtained from M. Dugat, Bouygues Company, France). It had a dry density of 2290 kg/m³, and consisted of the following components: Cement CPA HP Le Havre (290 kgm⁻³), sand 0/5 size fraction (831 kgm⁻³), gravel sand 5/12.5 size fraction (287 kgm⁻³), gravel sand 12.5/25 size fraction (752 kgm⁻³),

calcareous filler PIKETTY (105 kgm⁻³), silica fume (30 kgm⁻³), water (131 kgm⁻³), retarder CHRYTARD 1.7, super-plasticizer Resine GT 10.62. The maximum water saturation was 4 %kg/kg.

The Temelin concrete used for the concrete containment building of the Temelin nuclear power plant in the Czech Republic had a dry density of 2200 kg/m³ and maximum water saturation 7 %kg/kg. The composition was as follows: Cement 42.5 R Mokrá (499 kgm⁻³), sand 0/4 size fraction (705 kgm⁻³), gravel sand 8/16 size fraction (460 kgm⁻³), gravel sand 16/22 size fraction (527 kgm⁻³), water (215 kgm⁻³), plasticizer 4.5 lm⁻³.

The measurements of both linear thermal expansion and linear hygric expansion were performed on 12 samples each. The dimensions of the samples were $40 \times 40 \times 120$ mm, and the centers of the 40×40 mm faces were provided with contact seats for use with the contact comparator.

5 Experimental results

5.1 Linear thermal expansion coefficient

Figs. 1a, b show the measured linear thermal expansion coefficient of both types of high performance concrete in the



Fig. 1a: Dependence of the linear thermal expansion coefficient of Temelin concrete on temperature

temperature range from 20 °C to 1000 °C. The experimental results for Temelin concrete show an abrupt change in the character of the $\alpha(T)$ function at approximately 500 °C. The course of the $\alpha(T)$ function for Penly concrete is even more dramatic, with several maxima and minima. This is



Fig. 1b: Dependence of the linear thermal expansion coefficient of Penly concrete on temperature

apparently a consequence of structural changes in the concrete due to the chemical processes taking place in the studied temperature range.

5.2 Linear hygric expansion coefficient

The measured results are summarized in Figs. 2a, b. Both types of high performance concrete exhibited a very similar behavior, and their linear hygric expansion coefficient decreased with the moisture content.



Fig. 2a: Dependence of the hygric expansion coefficient of Temelin concrete on temperature



Fig. 2b: Dependence of the hygric expansion coefficient of Penly concrete on temperature

6 Conclusions

The linear thermal expansion of two types of high performance concrete was determined in a wide temperature range, and the linear hygric expansion coefficient in wide moisture range. The changes both of linear thermal expansion coefficient with temperature and of linear hygric expansion coefficient with moisture were found to be very significant, so that they cannot be neglected in practical applications.

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