# Water transport in air-pores

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# **1. INTRODUCTION**

Air-pore structure of AAC has been well documented in terms of compressive strength [1] and thermal performance [2] while little is known regarding water transport characteristics, probably because air-pores are large enough not to be acting as a capillary [3]. This study emphasizes the characteristic role of air-pores in water transport based on the three experiments: water redistribution around unsaturated air-pores during freezing, threshold air-pore connectivity for water penetration and entrapped air dissolution during absorption. Low-temperature SEM observation of unsaturated AAC during freezing with a moisture content of 35 vol.% showed considerable ice segregation in air-pores implying unfrozen capillary water redistribution and stress relaxation. It was also shown that mean air-pore diameter of 0.2mm with a total air-pore volume of 45 vol.% exhibited abrupt water penetration implying a bond percolation in the connectivity of air-pores. The slow water uptake observed in the latter stage of water absorption test was attributed to the dissolution of the entrapped air bubbles in the water of air-pores.

# 2. UNFROZEN WATER TRANSPORT AND ICE SEGREGATION IN AIR-PORES 2.1 OBSERVATION OF AIR-PORE SURFACE DURING FREEZING

When unsaturated AAC is subjected to freezing, water held in large pores enough to freeze without freezing point depression freezes first and subsequent freezing extends gradually over smaller pores [4]. The unfrozen adsorbed or capillary water that exhibits freezing point depression due to surface forces of the solid shows a tendency to migrate towards ice as the chemical potential of ice decreases associated with a decrease in temperature, leading to a redistribution of water inside of a specimen without external water supply. The air-pores will provide unfrozen water with spaces for ice segregation.

A schematic image of AAC with a different degree of saturation *S* under freezing temperatures is shown in **Fig. 1**. It is show shown in **Fig. 1(b)** that moisture in a specimen saturates capillary pores but not extends over air-pores except for very fine sire pores. When specimen temperature decreases lower than the freezing temperature of free water, the capillary water begins to freeze and segregates in the air-pores. The direct observation of ice segregated in an air-pore was shown by Gruble [5] but the moisture content of the specimen during freezing was not specified and the conditions to obtain the observed surface was not reported. Corr



Fig. 1 Water distribution in air-pores before and after freezing

et al [6] and Monteiro et al. [7] conducted series of experiments with a cryo-SEM but the moisture condition was not controlled. In this study, the moisture content of specimen was controlled to be close to that shown in **Fig. 1(b)**. The specimen was then subjected to freezing and observed with a cryo-SEM.

# **2.2 TEST CONDITIONS**

AAC specimens with a bulk density of 486 kg/m<sup>3</sup> and 638 kg/m<sup>3</sup> were prepared using pre-foaming method in the laboratory while those of gas-foaming type with

Tuble 1 m pore sudeture and water transport properties					
Sample	Air pore	Apparent	Air pore	Standpipe	Immersion in
	volume	mean air pore	connectivity	adsorptivity	water 48 hours
	(vol.%)	diameter	index	(mm/24	(mass %)
		(mm)		hours)	
Α	43.2	0.142	0.20	249	67.5
В	44.6	0.153	0.14	11	48.5
С	45.8	0.155	0.28	250	53.1
D	46.1	0.162	0.21	219	69.5
Е	41.8	0.207	0.11	10	45.3
F	43.0	0.222	0.09	10	49.9
G	38.8	0.285	0.08	13	48.6

Table 1 Air pore structure and water transport properties

a bulk density of 492 kg/m<sup>3</sup> were a commercial product. The production conditions of the specimen are shown in **Table 1**. In this experiment, specimen with a bulk density of 486 kg/m<sup>3</sup> was used and cut out in a 7mm in length and 5 mm in diameter. They were first vacuum-saturated in water followed by a gradual drying to have a degree of saturation of 47.5%. The specimens were sealed and left in a freezing chamber of  $-15^{\circ}$ C for three days and then introduced to the specimen chamber of the field emission scanning electron microscope (Hitachi S-800) with a cryo-stage that was able to cool specimens to the liquid nitrogen temperature. Vacuumed at a pressure of 0.8torr, the specimen was cut in the specimen chamber to have a fresh cross-section for observation. The acceleration voltage was 5kV and no gold deposition was applied.

Normally frost damage to AAC occurs after several freeze/thaw cycles. However major contribution of the freeze/thaw cycle is to increase moisture content to a critical degree of saturation  $S_{cr}$  [8] where only one freezing could cause frost damage. Hence just one freezing, employed in this experiment, is a sufficient condition to observe ice formation at the instant of frost damage.

# 2.3 RESULTS OF THE CRYO-SEM OBSERVATION OF ICE FORMATION

The degree of saturation of the specimen was 47.5 (38 vol.%) which was controlled to be close to the critical degree of saturation for freeze-thaw resistance as defined by RILEM recommendation. At this moisture condition, the capillary pores of 35 vol.% located in the matrix part between air-pores are fully saturated and some small air-pores may also be saturated [9]. The surface of air-pores frozen at this moisture condition is shown in **Fig. 2(a)** and **(b)**. At the center of **Fig. 2(a)**, a large air-pore shows small amount of ice while the other smaller air-pores on the right show considerable ice formation. **Figure 2(b)** is a



Fig. 2 Ice segregation in air-pores of unsaturated AAC

two-time magnified view of the upper-right part of Fig. 2(a).

The preliminary optical microscope observation showed that no water existed in such air-pores before freezing, hence the observed ice is supposed to come from the matrix part behind the air-pores. On the surface of the largest air-pore in **Fig.** 2(a), several parts show no ice formation, which is highlighted with a white dashed circle. It is obvious that this part has a hole behind which may be generated as a result of overlapping of air-pores. This channel plays an important role in the downward water absorption test as discussed later.

The ice observed on the surface of air-pores may come from the capillary pores between air-pores hence no ice can be formed on the channel that has no capillary pores behind. A straightforward evidence of the unfrozen water migration is shown in **Fig. 2(b)** as highlighted with a white circle, where the cross-section of the channel is clearly shown without any ice segregations.

#### **2.4 DISCUSSION**

Specimen with a moisture content of the critical degree of saturation for freeze/thaw resistance still contains ice-filled air-pores with a diameter less than 150 $\mu$ m as shown in **Fig. 2(b)** forming a hollow sphere of thick ice shell. Moreover the small air-pore shown in the right side of **Fig. 2(a)** is fully occupied with ice where unfrozen water migration from the matrix part is no longer possible resulting in a partial increase in hydraulic pressure. This corresponds with the fact that the specimen with a moisture content of  $S_{cr}$  showed expansion first among others with a lower moisture content exhibiting contraction [10].

As well known, unfrozen water in an unsaturated porous material shows redistribution during freezing and freezes into ice at large pores such as air voids. This is a major reason of the availability of entrained air for frost resistance of concrete. However, when its moisture content exceeds a certain limit, the air voids can no longer mitigate the hydraulic pressure, as directly shown in this experiment.

# **3. STANDPIPE ABSORPTIVITY AND AIR-PORE STRUCTURE 3.1 INTRODUCTION**

# 3.1 INTRODUCTION

AAC specimens with various air-pore structures and a fixed bulk density of approx. 500 kg/m<sup>3</sup> (total porosity of approx. 80%) were prepared using seven types of forming agent A to G. The dimension of the specimen was  $160 \times 160 \times 40$  mm. Because the bulk density was fixed, air-pore volumes of the specimens were all the same. However, due to the difference in the type of foaming agent, air-pore size distributions were different, which was supposed to have some impact on the standpipe absorptivity.

# **3.2 TEST CONDITIONS**

The standpipe absorptivity test was performed with a water-filled graduated cylinder standing on a specimen with a dimension of was  $160 \times 160 \times 40$  mm and the decease in the water level was recorded with time. A supplementary immersion test was also performed using specimens with a dimension of  $100 \times 100 \times 100$  mm, which was immersed in water 100 mm below the water level and the increase in the water content was recorded with time. Another specimens were cut out to have a cross-section for image analysis with an observation area of 8 x 6 mm. A sufficient contrast of air-pores was obtained simply by emitting visible light from the lateral direction. The video images were subjected to image analysis and the area fractions of air-pore corresponds to that of volume fraction according to the stereology principle while apparent mean air-pore diameters

(0.142 mm to 0.285mm) were not reliable because the cross-sections did not always show true diameter. Hence the connectivity between air-pores cannot be discussed in terms of the apparent mean air-pore diameters. Instead a new concept of air-pores connectivity was introduced. As shown in **Fig. 3**, number of holes originated from the overlapping of air-pores, white circles on the scanning electron micrograph, were counted per unit area and named as the air-pore connectivity index.



Fig. 3 Macoscopic channels between air-pores observed with SEM

# **3.3 RESULTS OF THE STANDPIPE ABSORPTIVITY TEST**

Results are shown in **Table 1** and the relationship between standpipe absorptivity and air-pore connectivity index is shown in **Fig. 4**. It is clearly shown that the standpipe absorptivity shows considerable increase when the air-pore connectivity index exceeds 20% implying the establishment of the first overall interconnection of air-pores. A similar tendency is also found in the relationship between the apparent mean air-pore diameter and the standpipe absorptivity as shown in **Fig. 5** where the abrupt increase in the standpipe absorptivity is shown at an apparent mean air-pore diameter of 0.15 mm. This is a transition phenomenon where a volume-filling parameter such as air-pore size distribution causes abrupt property jump and may be an experimental evidence of a percolation threshold.

#### **3.4 DISCUSSION**

In the standpipe absorptivity test, a pressure head is applied to the upper surface of a specimen and any defects such as air-pore connection channels, if any, drive





Fig. 4 Relationship between standpipe absorptivity and air-pore connectivity



water more quickly and massively than capillary suction. Hence the test can evaluate the macroscopic pore continuity such as air-pore connectivity. However in the water immersion test, the capillary suction and the macroscopic channel drive water simultaneously hence the experimental results provide less information on the relationship between pore structure and water transport characteristics. In the free water uptake test as described later, capillary suction dominates the entire flows lather than gravity-driven flow and the effect of air-pore structure would be manifested as the dissolution of entrapped air bubbles that are left behind by the capillary flows. It is shown in this experiment that an abrupt decrease in resistance to the gravity-driven flow may occur associated with an increase in air-pore connectivity when the apparent mean air-pore diameter lowers a limit value.

# 4. CAPILLARY WATER UPTAKE AND ENTRAPPED AIR BUBBLES

#### 4.1 FREE WATER UPTAKE TEST

When water is sucked form the bottom of an AAC specimen solely by capillary suction, the well-known two straight lines of the relationship between moisture content and square root of time can be observed. The first straight line has been attributed to the sorptivity [11] while the meaning of the other straight line observed at the latter stage of absorption still needs discussions. In this study, AAC specimens with different capillary and air-pore volumes were prepared and examine the later-stage absorption behavior with the free water uptake tests.

#### **4.2 TEST CONDITIONS**

Three types of AAC specimens with different bulk densities and foaming methods were prepared as shown in **Table 2**. The dimension of the specimen was 100mm in diameter and 40mm in thickness cut out from a cylinder of 100mm in diameter and 200mm in height and the surface was sealed with a silicone type

Foaming Bulk density Total Curing Raw materials method  $(kg/m^3)$ method porosity Gas Quartz powder, portland 492 0.810 Autoclave cement, Aluminium powder foaming curing Pre 180°C. 486 0.815 Quartz powder, portland foaming cement, Quick setting agent, 10 atm, Pre Surface active agent 8 hours 638 0.757 foaming

Table 2 AAC specimen

sealant except for the upper and bottom side to keep cutout surfaces. Specimens were placed in a glass desiccator to avoid drying and the shallow tray with a stainless steel mesh supporting the specimens makes it possible to keep water level constant as shown in **Fig. 6**. Because the top surface is the original cutout surface, permeability of gas is not interfered and the glass desiccator allows observation of the top surface of the specimen during water absorption. The specimens were taken out at specified intervals, wiped out the bottom surface with a PVA foam softly and then weighed quickly.

#### 4.3 RESULT OF THE FREE WATER UPTAKE TEST

The test result is shown in **Fig. 7** where absorption rate depends on both bulk density *d* and pore structure. In the case of specimen with a bulk density of d=486 kg/m<sup>3</sup>, entire capillary pore of 35 vl.% was filled quickly at the first 10 hours by the capillary suction. Because the moisture condition in this figure is expressed in terms of saturation, volume of water by the total pore volume, the measured values in the figure can be converted to volumetric moisture content by multiplying the total porosity. The broken two straight lines both show changes in moisture content with square-root of time and suggest two different diffusion mechanisms.

The abrupt absorption rate observed immediately after the specimen-water



Fig. 6 Apparatus of the Fig.7 Water absorption rate of various types of AAC free water uptake test



Fig. 8 Schematic view of pore structure, water and air in unsaturated AAC

contact is well known as the sorptivity. At the end of this process, i.e. 10 hours later in this experiment, water reached to the top surface and the specimen exhibited a wet color while the moisture content was not as high as 80 vol.% that is the total pore volume of the specimen. Hence it is concluded that considerable air still remain in the specimen.

#### **4.4 DISCUSSIONS**

A stage of moderate increase in water content observed after the rapid absorption has been attributed to the diffusion of water to a finer gel pores [12] or to the escape of air through macroscopic channels [13]. It is however more realistic to take it a gradual dissolution of entrapped air bubbles and associated volume change can be compensated with incoming water [9, 15].

Configurations of water and air at different degree of saturations are schematically shown in **Fig. 8**. When water is in contact with a dried specimen, it

travels quickly through the capillary pores distributed in the matrix between air-pores, and the air inside of the air-pores is likely to be left as bubbles in water. The cryo-SEM observation of a cutout surface of an unsaturated frozen AAC also demonstrated a bubble-like spherical cross section within ice in an air-pore as shown in **Fig. 9**. This ice-air configuration agrees well with the predicted one shown in Fig. 8(2).

Diameter of the entrapped air bubble tends to be larger when the host air-pore is large. The dissolution rate of air in the entrapped air bubble becomes slow when the diameter of air bubble



Fig. 9 Direct observation of an entrapped air bubble in ice (see Fig. 8(2))

becomes larger resulting in slower water uptake. For comparison, AAC specimens with different mean air-pore diameters were tested in this experiment while their bulk densities are very close, 486 and 492 kg/m<sup>3</sup>, showing the total air-pore volume is the same. At the first stage of quick absorption, the sorptivity is almost the same while at the second stage, the water absorption slope is gentler in specimen with a larger mean air-pore diameter where air dissolution rate is slow as shown in Fig. 7.

#### **5. CONCLUSIONS**

The total porosity of AAC with a bulk density of approx.  $500 \text{ kg/m}^3$  extends over 80%. The pore volume comprises air-pore of 45% and capillary pore of 35%. Characteristics of this study is to execute various water transport experiments under well-characterized pore structure and well-controlled moisture contents.

A reverse from contraction to expansion may occur in unsaturated porous hydrophilic bodies at a critical moisture content. AAC specimens were prepared to have the critical moisture content and observed with the cryo-SEM. Some evidences were confirmed that the ice segregated on the surface of air-pores could be a result of redistribution of unfrozen capillary water coming from the matrix part between air-pores.

When water is applied downward, the gravity-driven bulk flow may exceed the capillary flow provided that a sufficient amount of macroscopic channels develop between air-pores leading to an early-stage saturation. The standpipe absorptivity test was performed for AAC specimens with an air-pore volume of approx. 45 vol.% while having different mean air-pore diameters. The permeability exhibited an abrupt jump when the mean air-pore diameter was smaller than 0.15mm that corresponded to the air-pore connectivity of 20%. The critical air-pore connectivity that provides a number of air-pore channels from the top to the bottom of the specimen could cause such a transition phenomenon.

In the free water uptake test using a dried AAC specimen, the capillary suction precedes other water transports and saturates the capillary pore of 35 vol.% instantly followed by a slow absorption process, i.e. escaping or dissolution of air bubbles entrapped in air-pores, that controls the subsequent absorption rate until the full saturation.

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KEY WORDS: Autoclaved aerated concrete, Cryo-SEM, Ice formation, Air pore, Entrapped air, Sorptivity, Standpipe absorptivity.

**Abstract**: Water transport experiments were performed on AAC with well-characterized pore structure and well-controlled moisture contents. Specimens with the critical moisture content for freeze/thaw resistance were observed with the cryo-SEM. Evidences were confirmed that the ice segregated on the surface of air-pores as a result of redistribution of unfrozen water coming from the matrix part between air-pores. The standpipe absorptivity test for specimens with an air-pore volume of approx. 45 vol.% while having different mean air-pore diameters showed an abrupt jump of permeability when the mean air-pore diameter was smaller than 0.15mm or the air-pore connectivity was 20%. In the free water uptake test using a dried AAC specimen, the capillary suction precedes other water transports and saturates the capillary pore of 35 vol. % instantly followed by a slow absorption process, i.e. escaping or dissolution of air bubbles entrapped in air-pores that controls the subsequent absorption rate