



Durability based design of concrete

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Performance based service life design: durability of concrete

Contents:

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2. Getting a grip on service life
3. Example of the importance of the environment – tunnel example
4. Example of a mechanical degradation mechanism:
 1. Mechanical: freeze-thawing
 2. Chemical: carbonation
5. Conclusions



1. Introduction: what is durability

- › **Durability** is the resistance of concrete to degradation
- › **Degradation** is the decrease in properties of concrete under influence of loads, mostly from the outside
- › **Loads** include:
 - › - mechanical loads (e.g. self weight)
 - › - physical loads (e.g. T)
 - › - chemical loads (e.g. CO₂)





1. Introduction: why is durability important

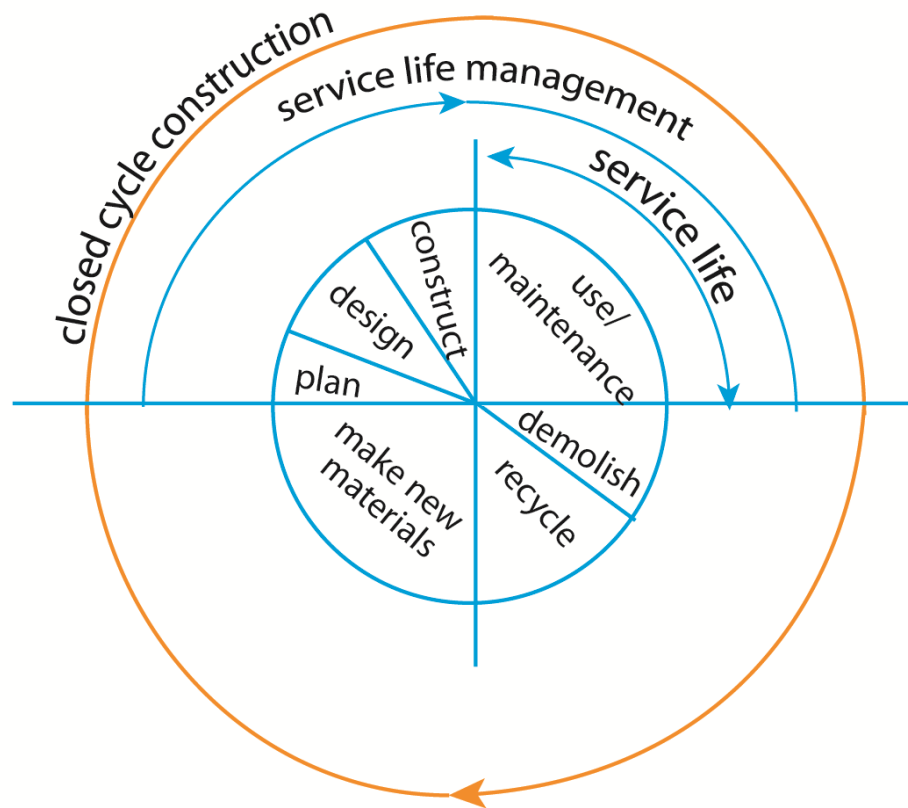
- **Degradation** is most often due to attack of the cement stone but also the aggregates and / or rebars.
- **Damage** include spalling & cracking, loss of cover and loss of rebar diameter upto bending / shear and collapse
- This may lead to extensive repair & costs or even early **end of service life**





1. Introduction

Service life is that stage of a life of a structure when it is in use

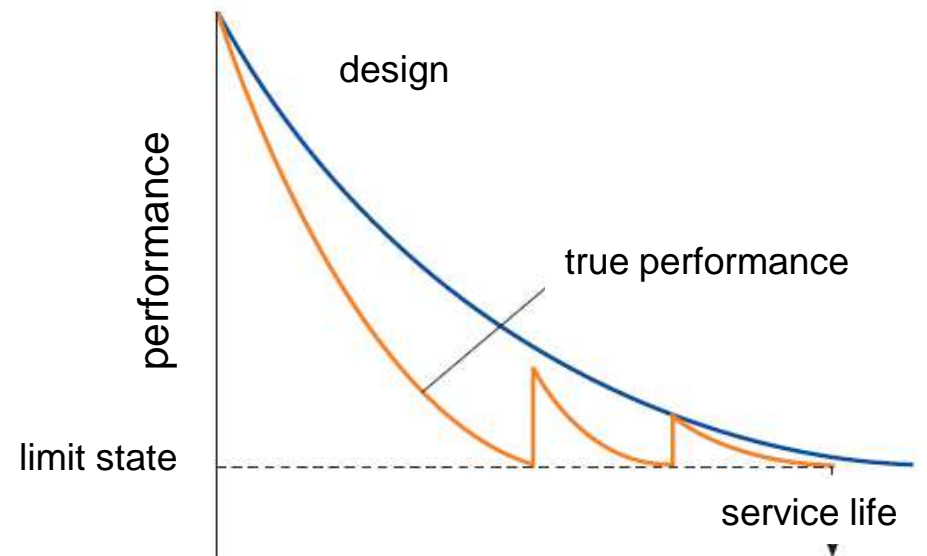


end of service life



1. Introduction

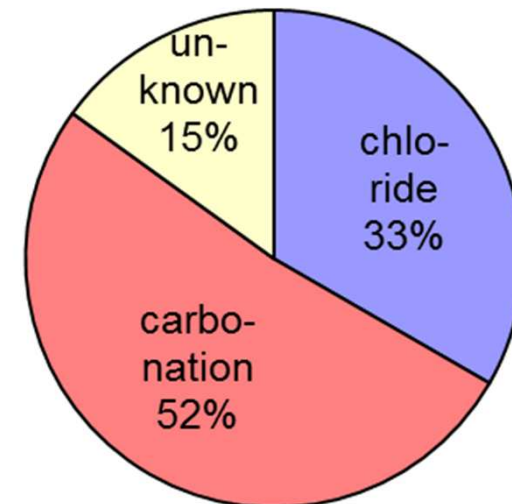
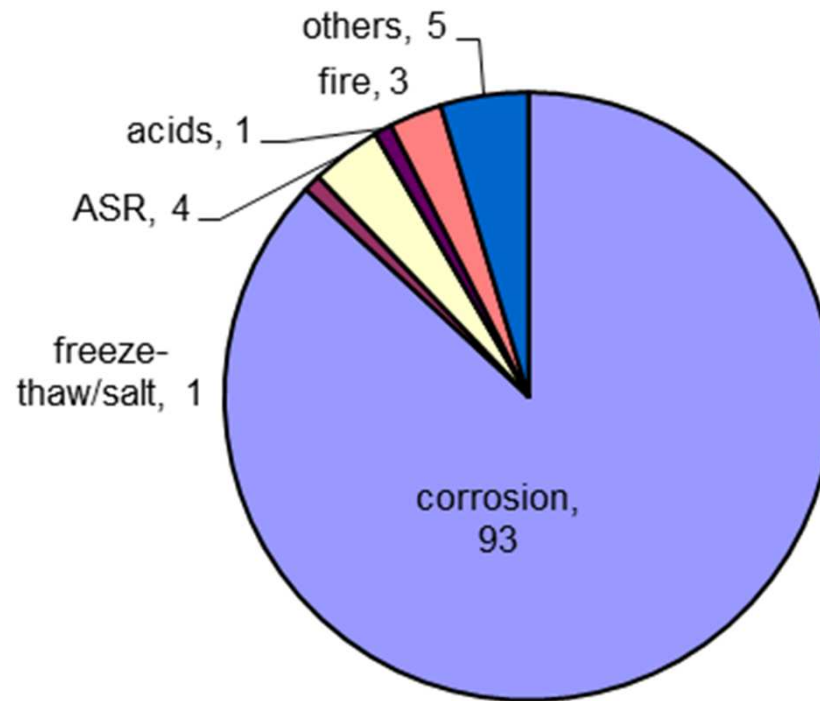
A **performance** is a **task** that is executed with a certain **quantified** result (limit state)





1. Introduction

- *The most often encountered durability issue is corrosion initiated by **chloride or carbonation***





1. Introduction

Why should you care
about the performance?

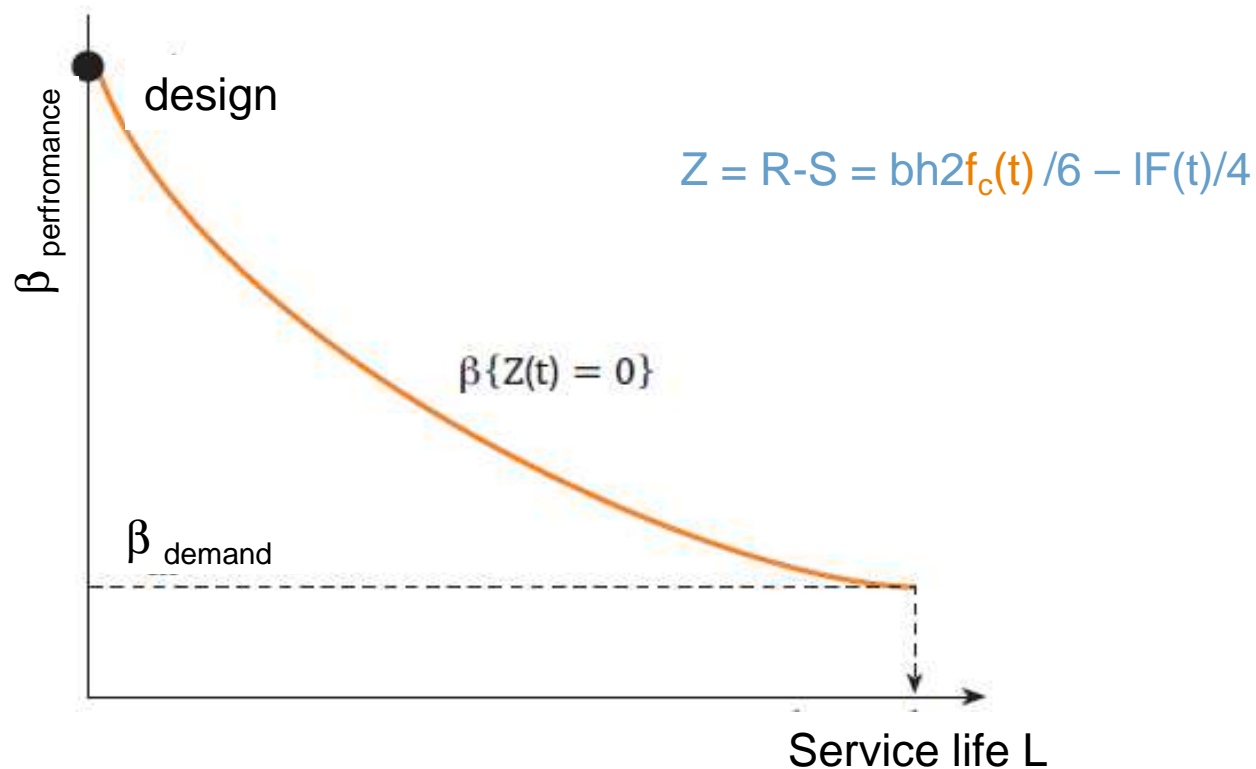




2. Performance in a service life design

› A performance consists of:

1. A behaviour model $Z(t)$ of which $Z(t) = 0$ the limit state describes
2. A pre-defined, agreed upon, reliability index β
3. A pre-defined, agreed upon service life L

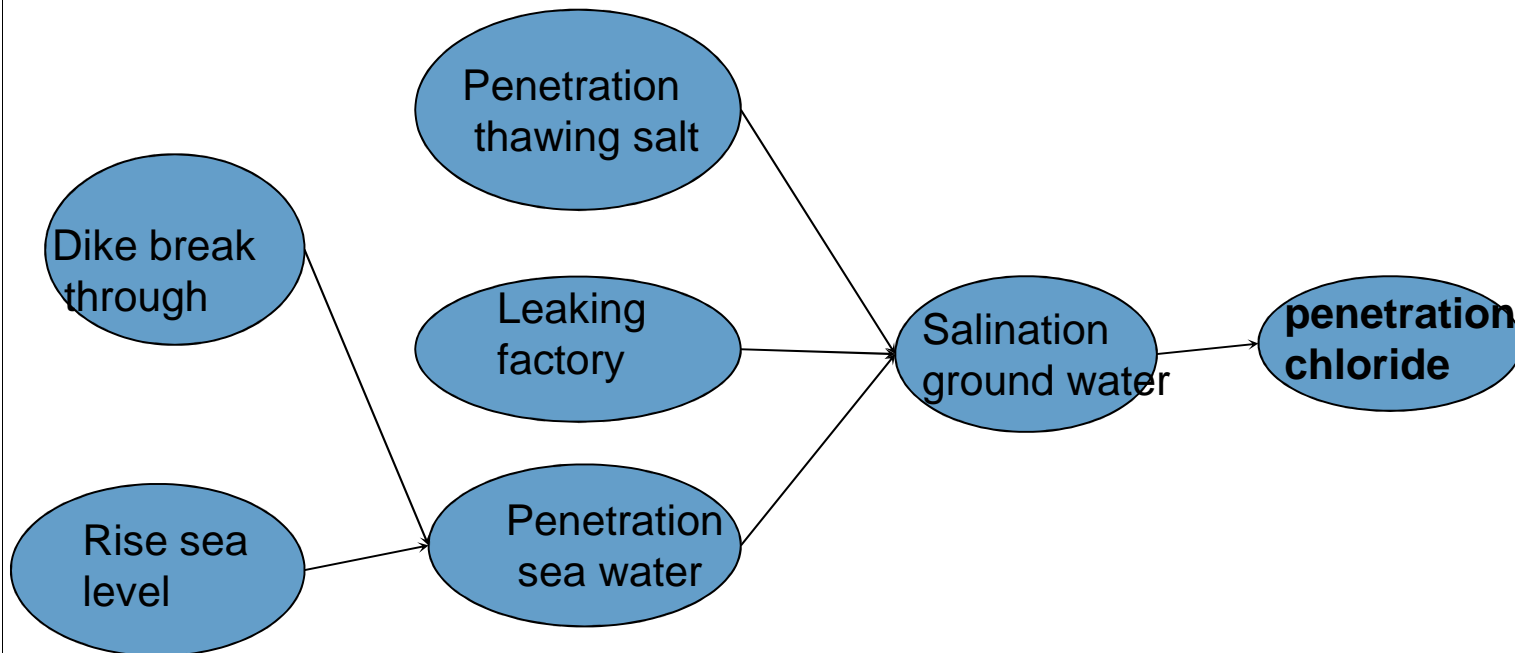




2. Performance in service life design

(1) Question 1: **which loads?**

- First: try to assess which of the loads are of importance, now, or **DURING SERVICE LIFE** (e.g. make an event tree)



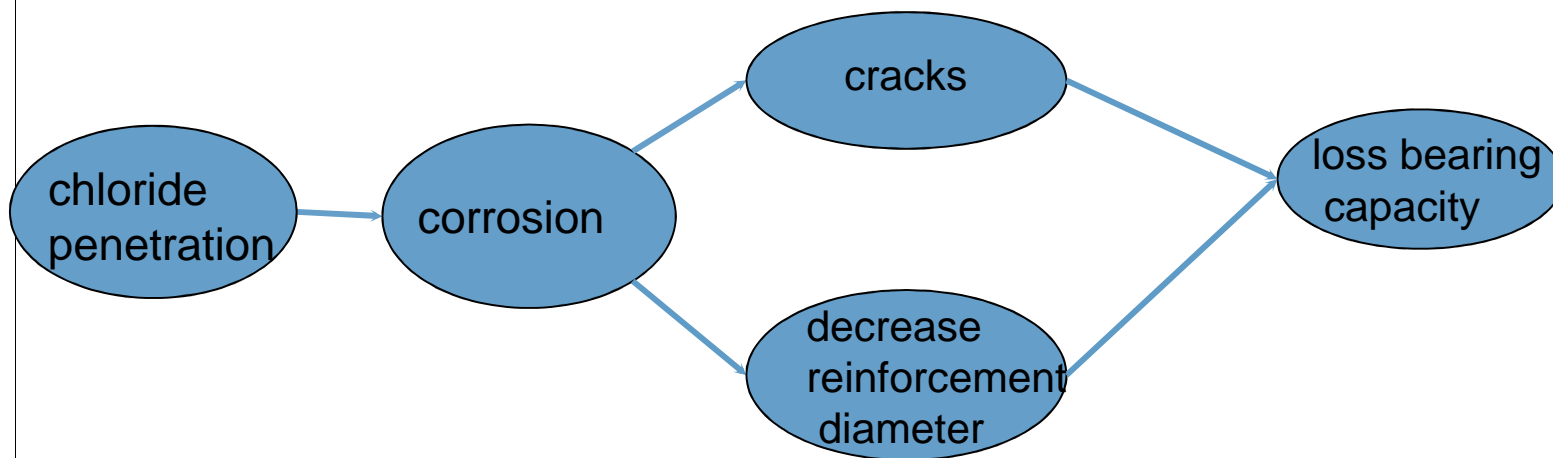
Reminder: structures fail more often due to forgotten treats rather than identified treats!



3. Performance in service life design

(1) Question 2: **does it matter?**

- Second: **check the effect of the load** (e.g. use a failure mode and effect analysis (FMEA))





3. Determining the performances to take into account

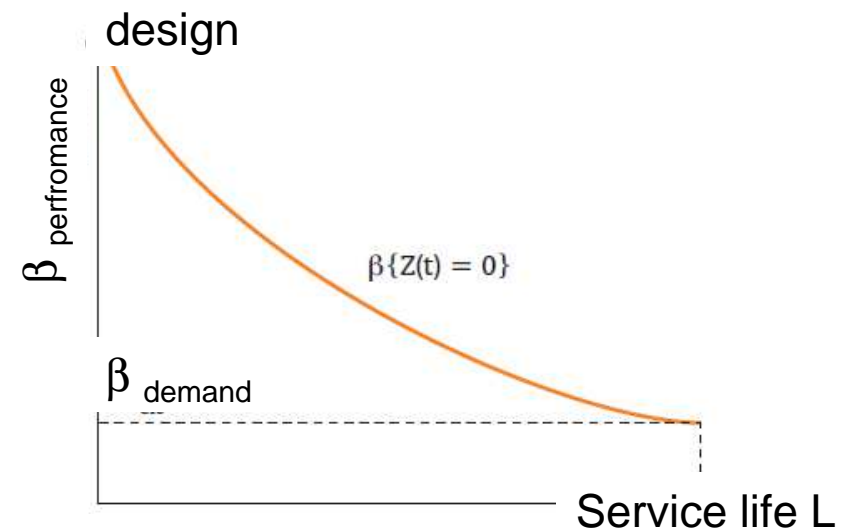
(1) Question 2: **does it matter?**

- Third, and last step (much later on in the process) will be to consider the consequence of the effects of failure (e.g. **make a risk analysis**):
 - loss of lives / health issues
 - Economical loss
 - Ecological loss etc.
- **Risk = probability of an event x the consequences**
- Most often, the risks are expressed as **costs**.



2. Performance in service life design

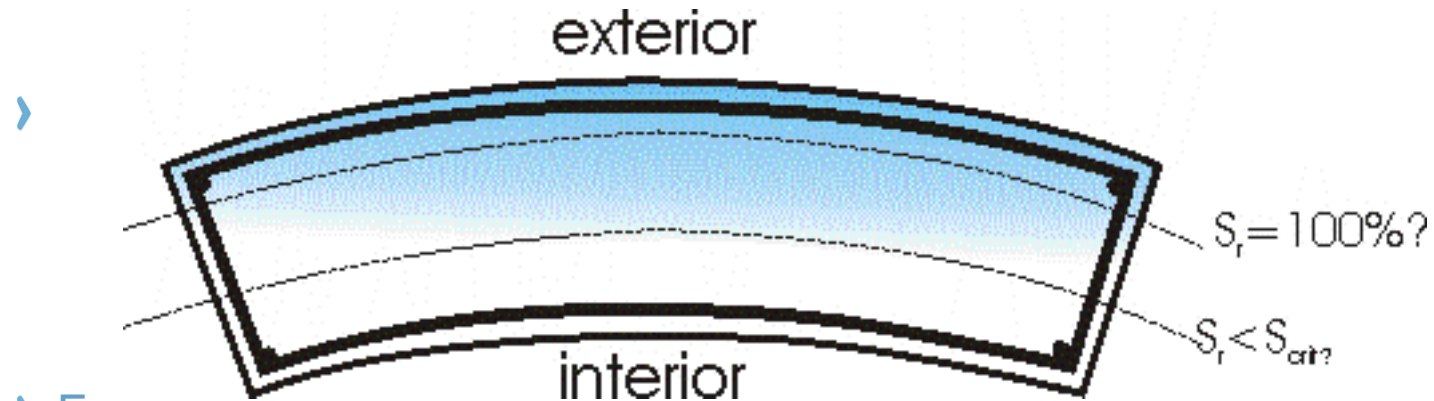
- › Recap:
 - › Performance consists of:
 1. A behaviour model $Z(t)$ of which $Z(t) = a$ the limit state describes
 2. A pre-defined, agreed upon, reliability index β
 3. A pre-defined, agreed upon service life L
 - › Its importance is determined from:
 1. Failure Mode
 2. And Effect
- Analysis (BOWTIE MODEL)





3. Example of the importance of the environment: moisture distribution in a tunnel

- › Problem statement: where is the moisture front?



- › E.g:

- › Reinforcement in saturated concrete: no corrosion
- › Reinforcement in dry concrete: no water, no corrosion

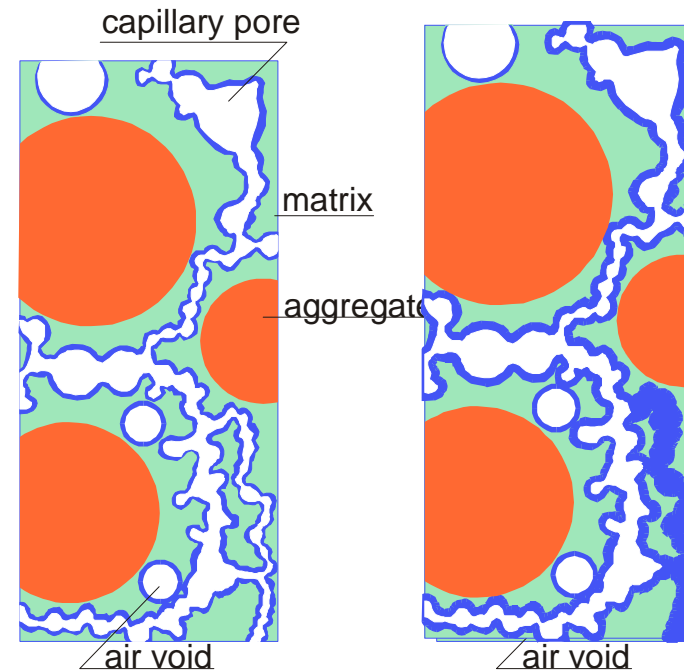




3. Example of the importance of the environment: moisture distribution in a tunnel

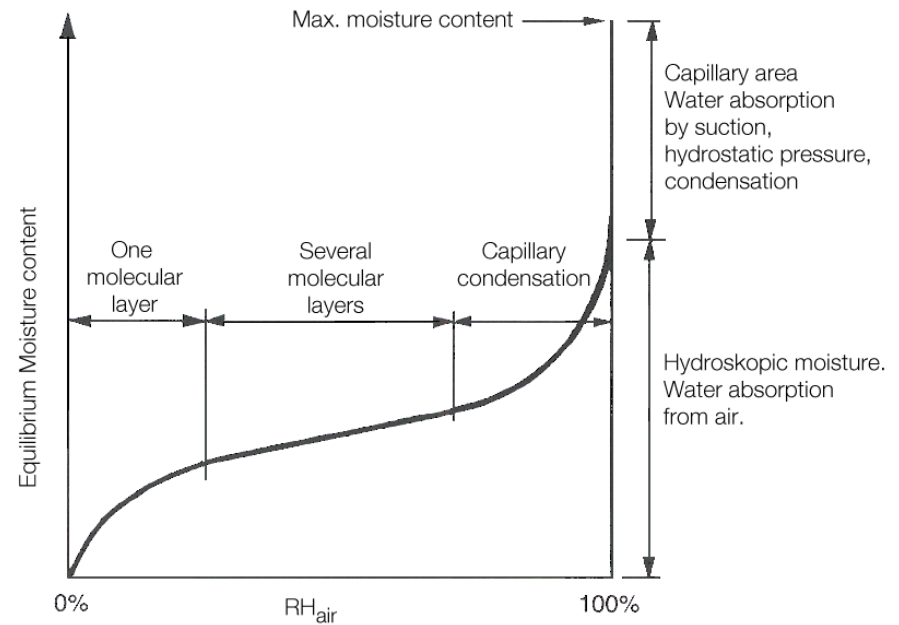
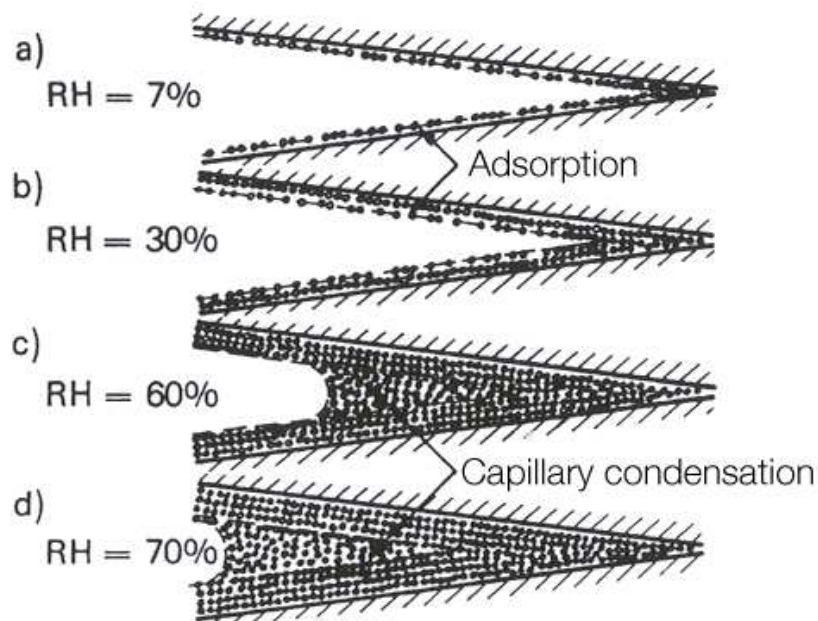
- › **Importance** of the structure and the moisture distribution
 - › Durability = transport + (chemical) reactions
 - › Transport rate / type depend on structure and related moisture distribution

- › Transport examples:
 - › Water / ions at $S > S_{crit}$
 - › gasses at $S < S_{crit}$





3. Example of the importance of the environment: moisture distribution in a tunnel

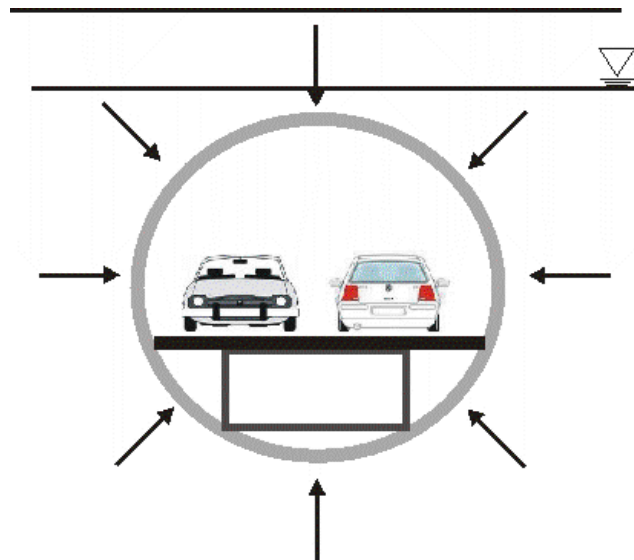




3. Example of the importance of the environment: moisture distribution in a tunnel

Exterior = below ground water table:

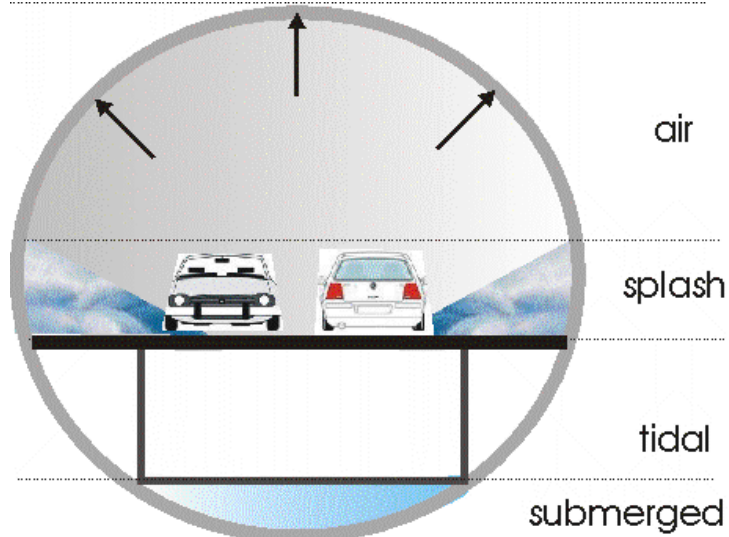
- › **pressure:** 0.1 – 0.5 bar
- › **temperature:** 12 °C (from -20 m)
- › **composition :** sweet (< XA1),
pH 6.0 – 7.1



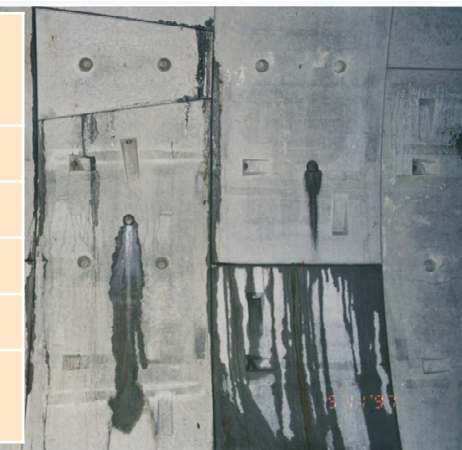


3. Example of the importance of the environment: moisture distribution in a tunnel

Interior – different environments



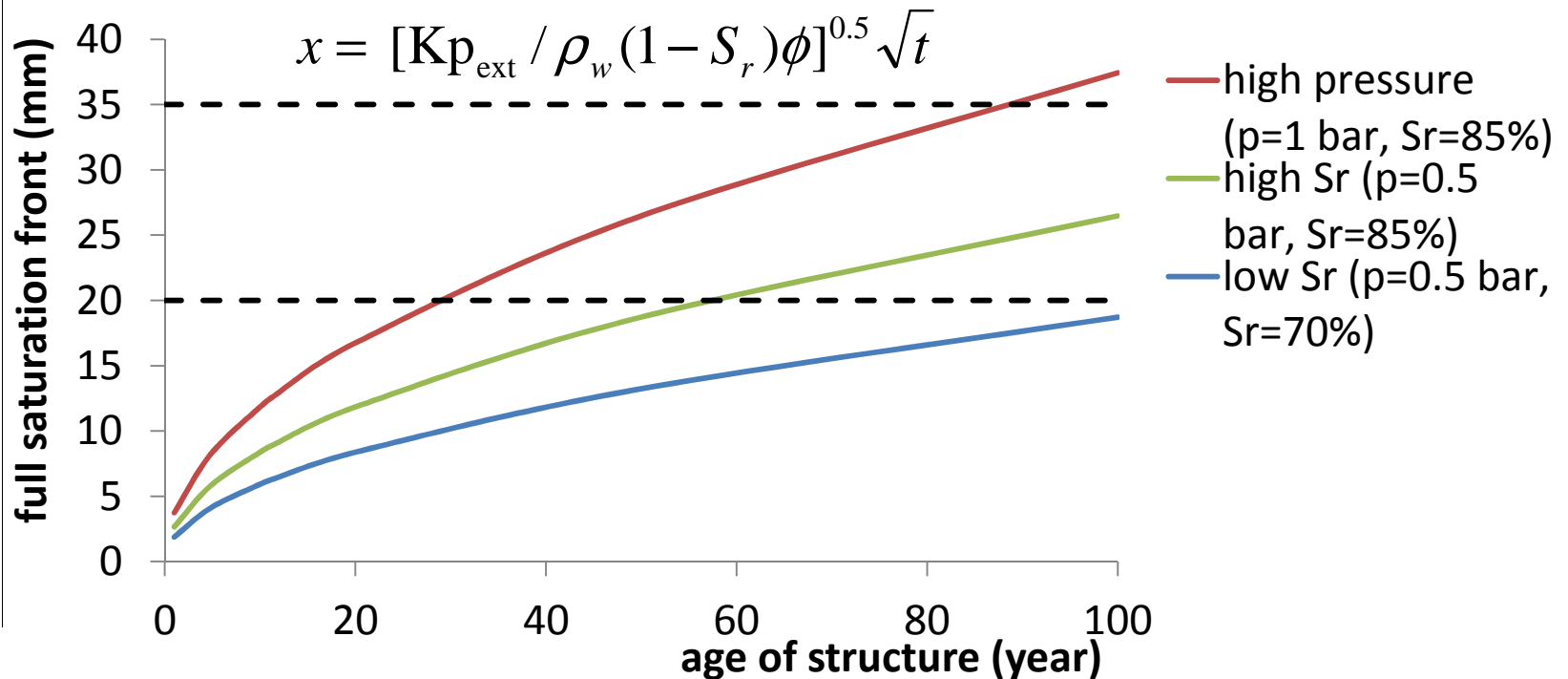
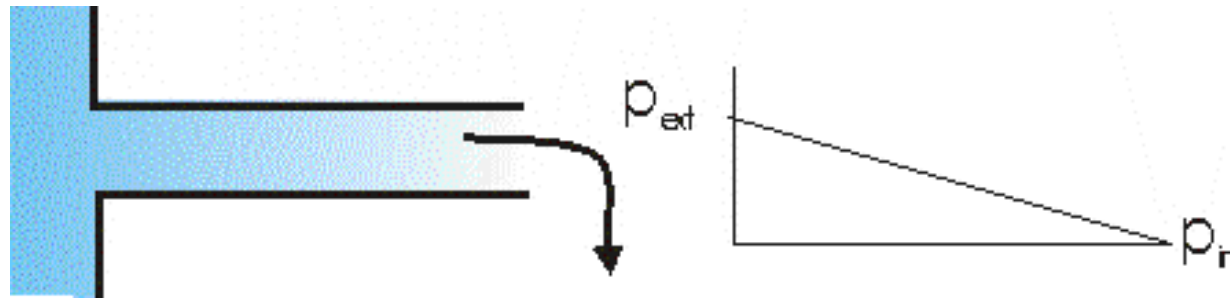
air zone	average	st.dev	minimum	maximum
RH _{int} (%)	70-65	10	20	92
RH _{out} (%)	80	10	24	100
T _{int} (°C)	12-14	6	-7 - -5	34-32
T _{out} (°C)	12	6	-8	32
CO ₂ (ppm)	400-580	20-50	400	650-1250





3. Example of the importance of the environment: moisture distribution in a tunnel

Further assessment by modelling: Darcy flow

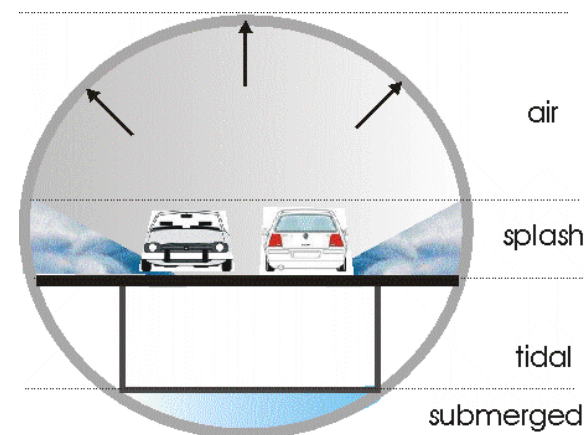
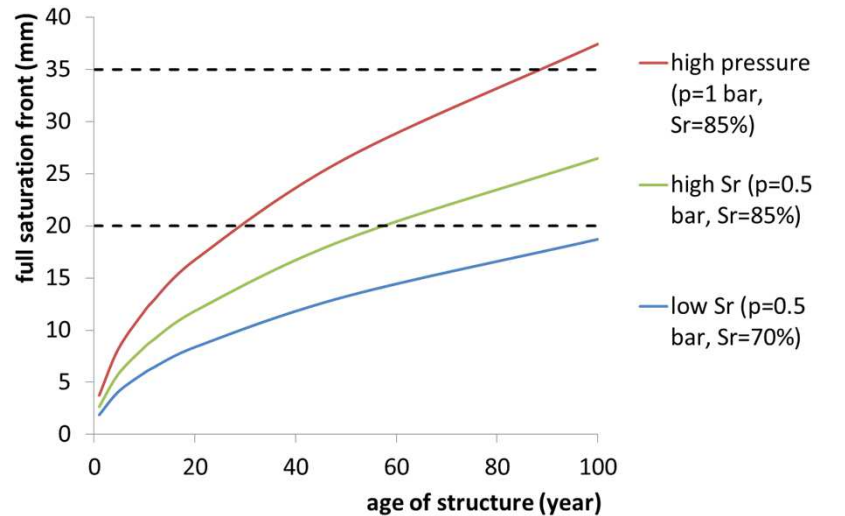




3. Example of the importance of the environment: moisture distribution in a tunnel

Conclusions:

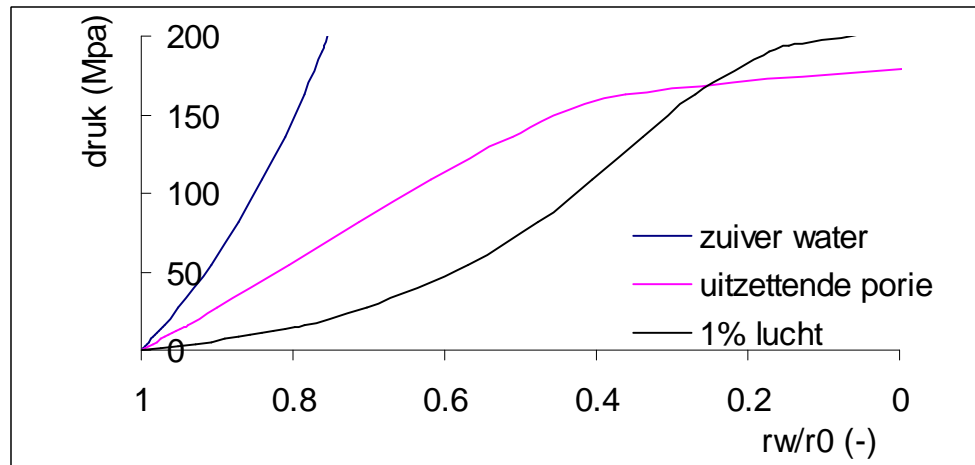
- Moisture conditions are in equilibrium with the internal RH
Water on the exterior functions as impervious
Moisture conditions further constant throughout, about 85%
- Corrosion of the reinforcement at the interior due to carbonation is a risk;
- Corrosion of the reinforcement at the exterior due to chloride penetration is much less likely



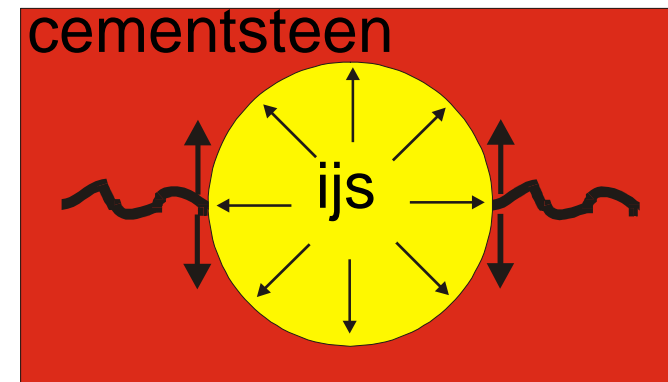


4.1 Degradation mechanisms: Example freeze-thawing

- › Mechanism 1: expansion of ice (compared to water approx. 10%)



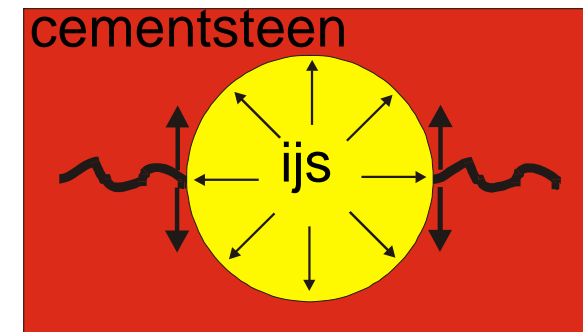
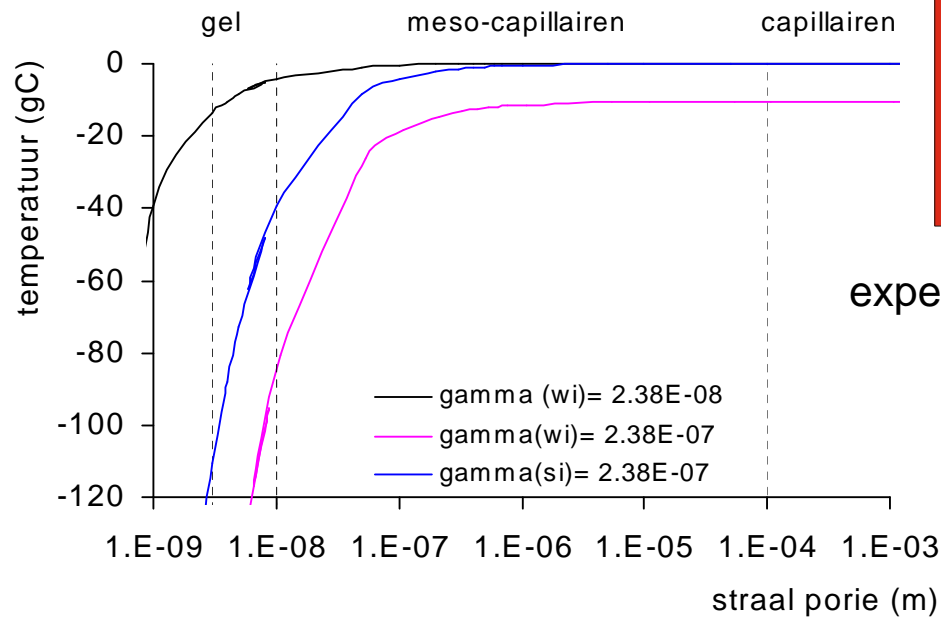
- › Conditions:
- › High enough S_r
- › Low enough temperature
- › Fast enough freezing





4.1 Degradation mechanisms: Freeze-thawing

- › Mechanism 1: expansion of ice (compared to water approx. 10%)
- › Mind: the freezing temperature increases with reduction in temperature and increase in salt concentration



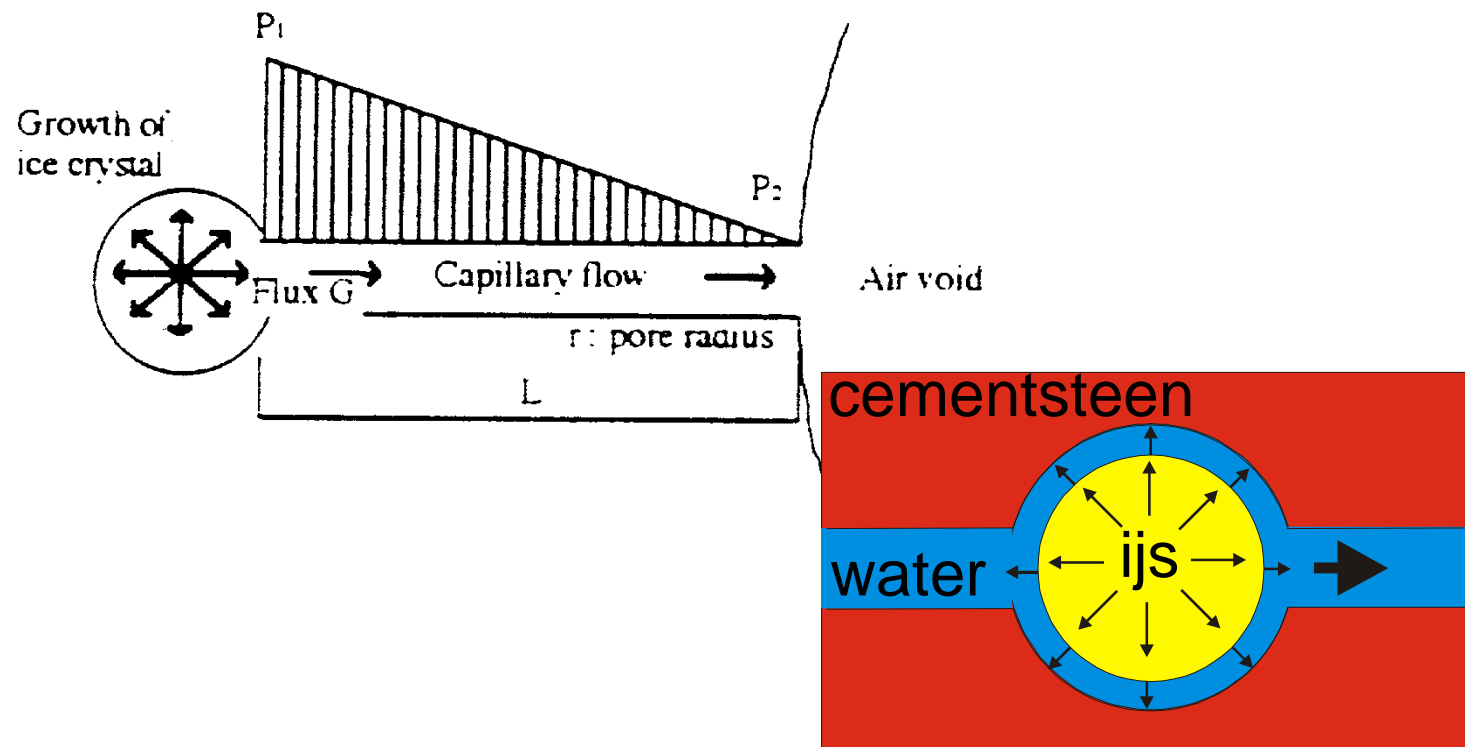
experiments

$r > 0.1 \text{ mm}$: 0 tot $-10 \text{ }^\circ\text{C}$
$r > 10 \text{ nm}$: -20 tot $-30 \text{ }^\circ\text{C}$
$3 < r < 10 \text{ nm}$: -35 tot $-45 \text{ }^\circ\text{C}$



4.1 Degradation mechanisms: Freeze-thawing

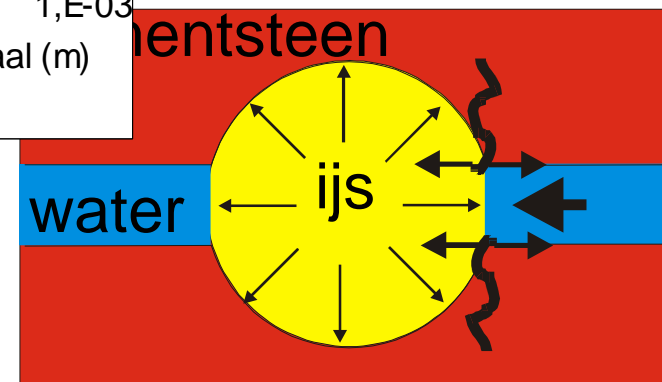
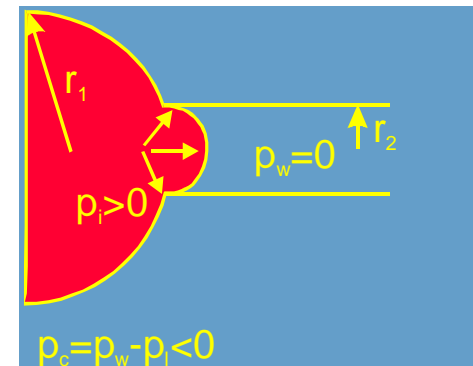
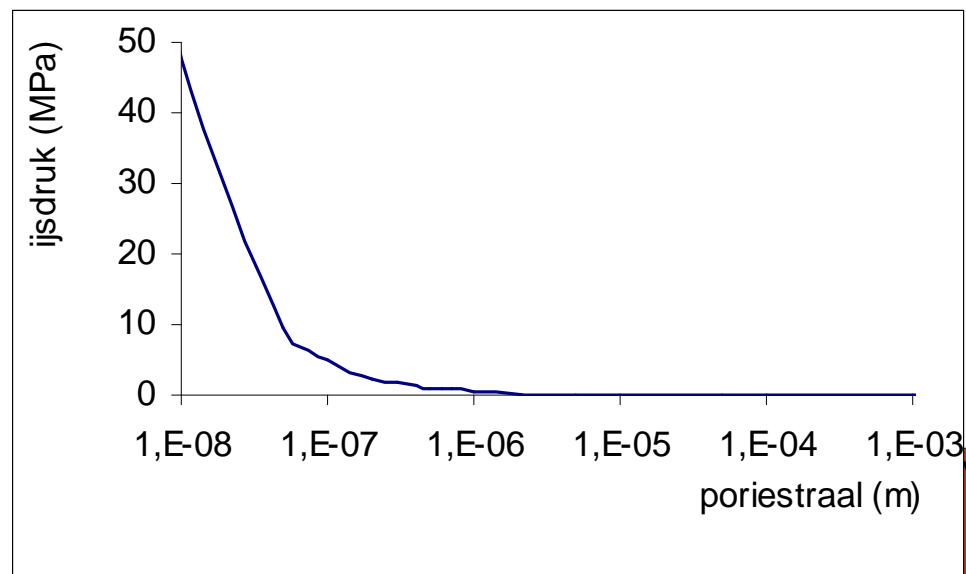
- › Mechanisms 2: hydraulic pressure (is why air entrainment works!)





4.1 Degradation mechanisms: Freeze-thawing

- 3. Ice over pressure due to transport of vapour pressure and growth



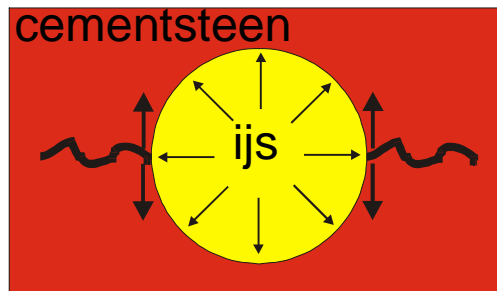


4.1 Degradation mechanisms: Freeze-thawing

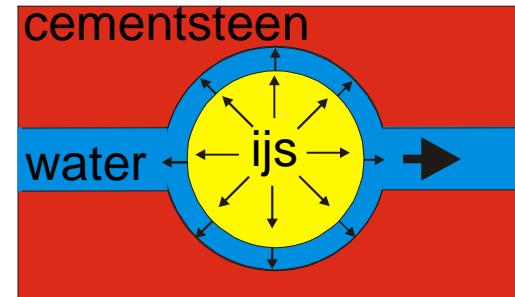
Summary mechanisms

› Fast freezing:

expansion of ice 10%;

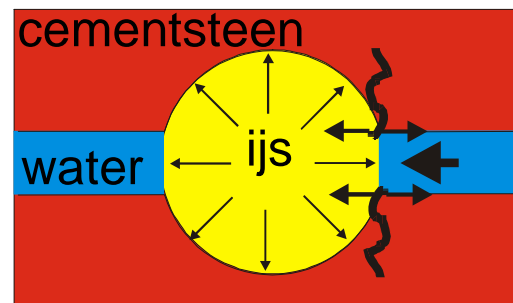


fast pressure build up



› Slow freezing (at modest T): transport of vapour to ice in larger pores

– ice over pressure

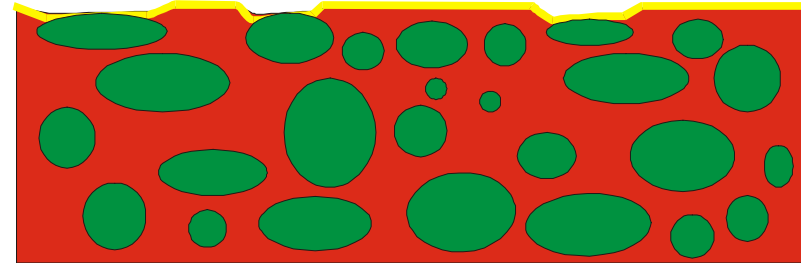




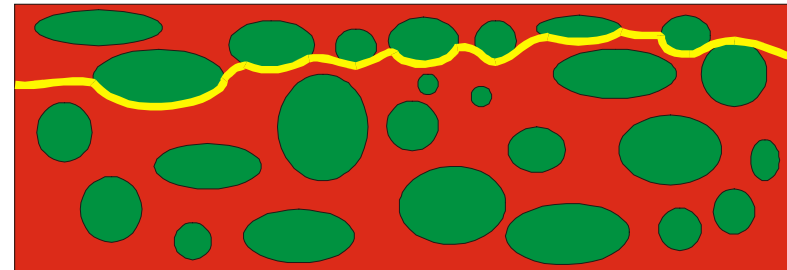
4.1 Degradation mechanisms: Freeze-thawing

Damage types:

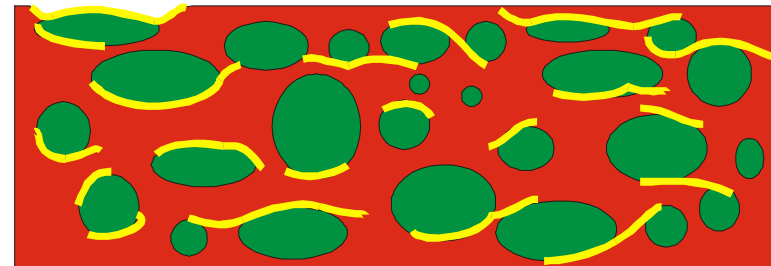
› Scaling



› Top layer delamination



› Total desintergration
(due to internal cracking)





4.1 Degradation mechanisms: Freeze-thawing

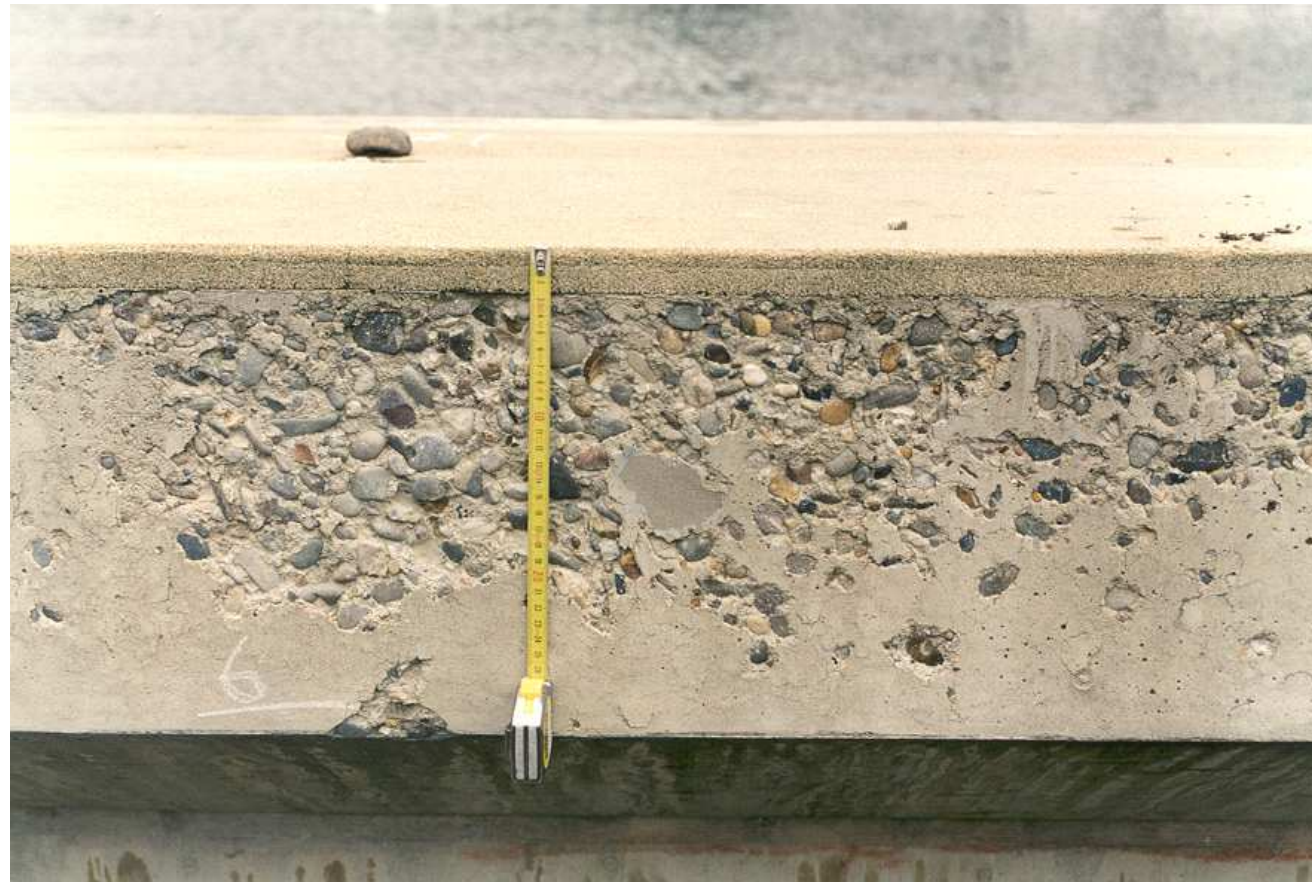
Conditions:

- › Sufficient low T
- › Sufficient fast (or slow) freezing
- › High enough degree of saturation
- › (depends of r_h , r and $c(\text{salt})$)





4.1 Degradation mechanisms: Freeze-thawing





4.1 Degradation mechanisms: Freeze-thawing





4.1 Degradation mechanisms: Freeze-thawing





4.2 Degradation mechanisms: chemical attack

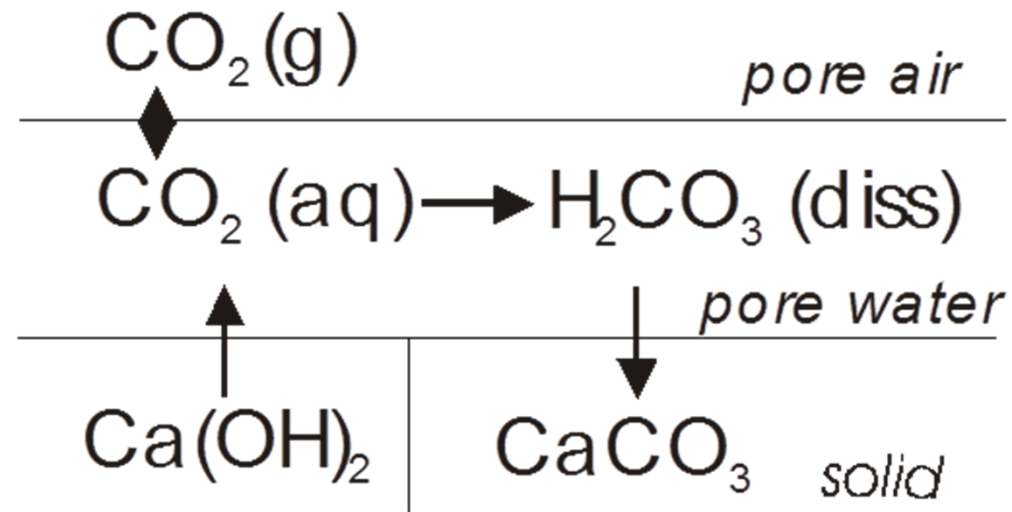
- › formation of new chemical products:
 - › volume increase / swelling
 - › lower binding capacity
 - › higher solubility

- › consequences
 - › cracking, scaling and spalling, debonding
 - › loss of stress bearing capacity
 - › reduction of service life



4.2 Degradation mechanisms: chemical attack

- › *Chemical equilibrium (carbonation example):*
 - › *Transport of CO₂ due to concentration difference*
 - › *Dissolution of CO₂ in pore water and formation carbonic acid and dissociated (H₂CO₃, HCO₃⁻, CO₃²⁻)*
 - › *Precipitation CaCO₃ - dissolution Ca(OH)₂*

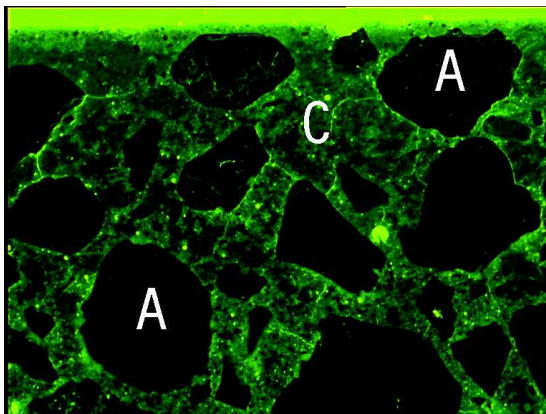




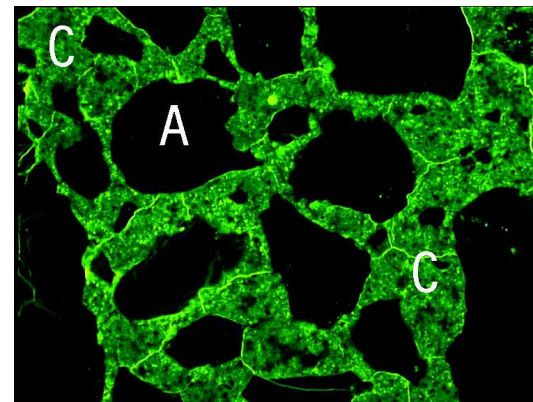
4.2 Carbonation

- Carbonation lead to **a change in the cement paste** (e.g. $\text{Ca}(\text{OH})_2$ into CaCO_3)
- Carbonation leads to **a change in the pore structure** (e.g. more dense for OPC, more porous for BFSC)
- Carbonation leads to a **change in the pH** of the cement stone below $\text{pH} < 9$; this breaks the passivation of the reinforcement when the carbonation front reaches it, leading to **corrosion**

non-carbonated BFSC



carbonated BFSC

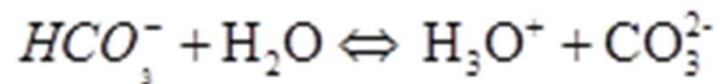
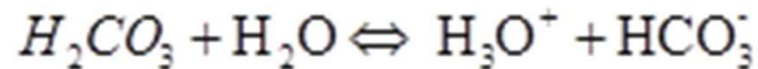




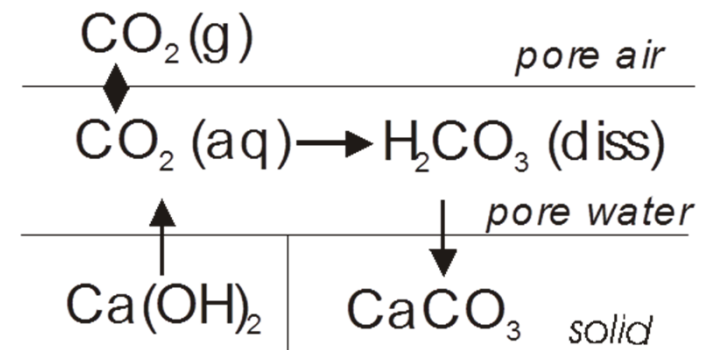
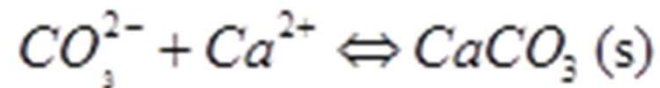
4.2 Carbonation

› *Chemical equilibrium – in detail:*

› *Dissociation is acidification process*

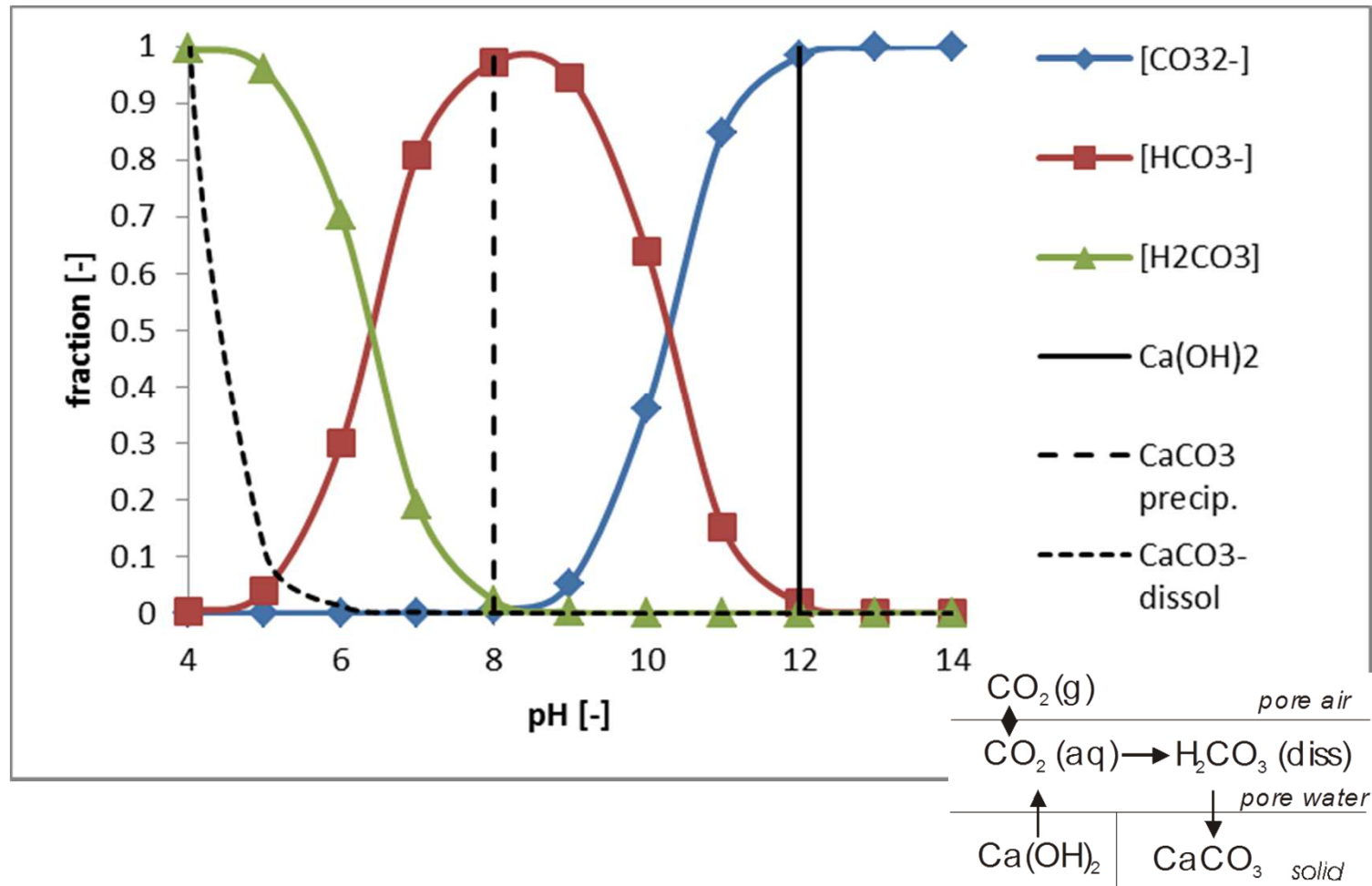


› *Precipitation of Ca^{2+} goes to completion:*





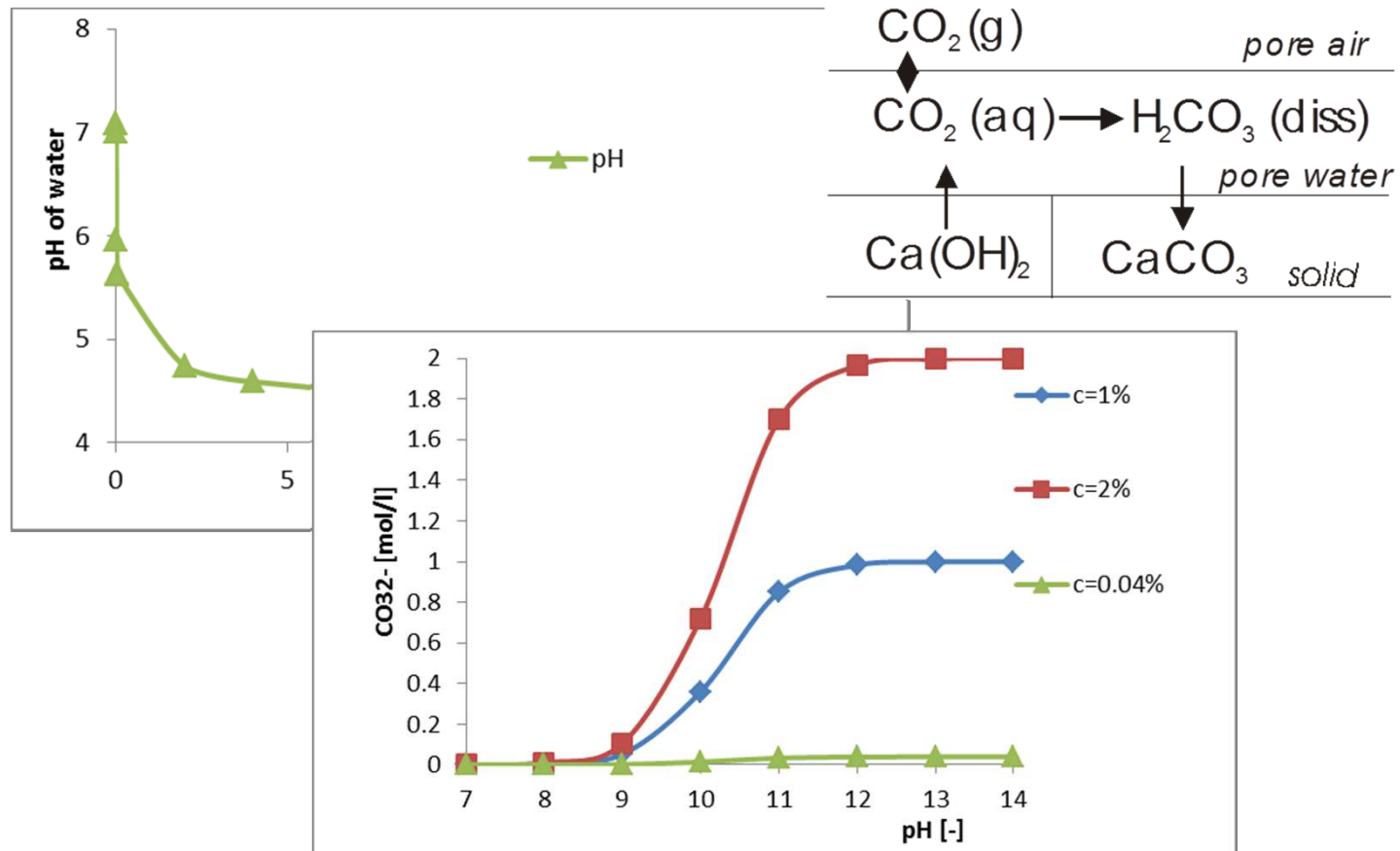
4.2 Carbonation





4.2 Carbonation

- *Equilibrium pH depending on CO₂ concentration*





4.2 Carbonation

- *Stability & pH*

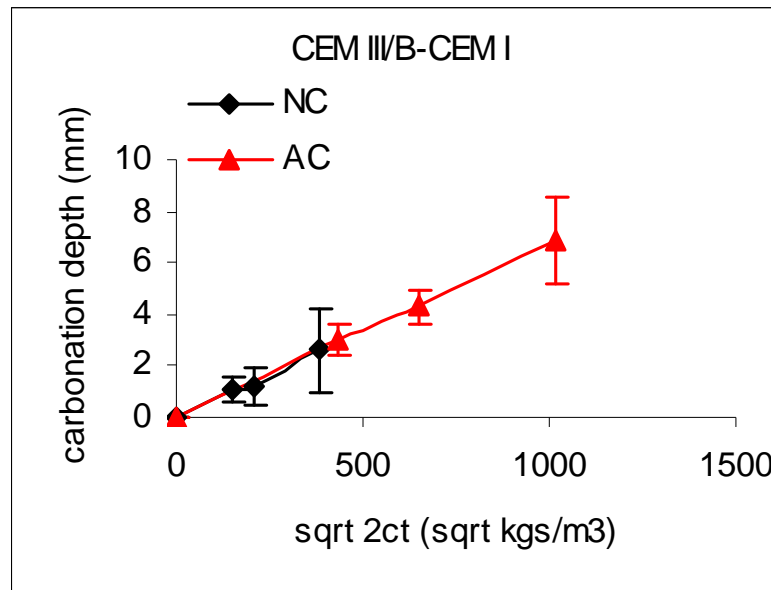
Stage	pH	stable phases
1 (non-carbonated)	>12.6	Ca(OH) ₂ , CSH (Ca/Si > 1.8 or at high common ion effect), AFt, AFm
2	11.6 – 12.6	CSH (Ca/Si< 1.8), AFt, AFm
3	10.5 – 11.6	CSH (Ca/Si< 1.05), AFt, Al(OH) ₃
4	10.0 – 10.5	CSH (Ca/Si< 0.85), Fe(OH) ₃ , Al(OH) ₃
5 (fully carbonated)	< 10	SiO ₂ with some CaO, Fe(OH) ₃ , Al(OH) ₃

Data from literature, mostly from Lagerblad, 2005



4.2 Carbonation

- *Carbonation is a very slow process*
- *Testing taking often a year to obtain reliable results*
- *In a design-by-testing a year is too long*
- *Acceleration of the test is required*



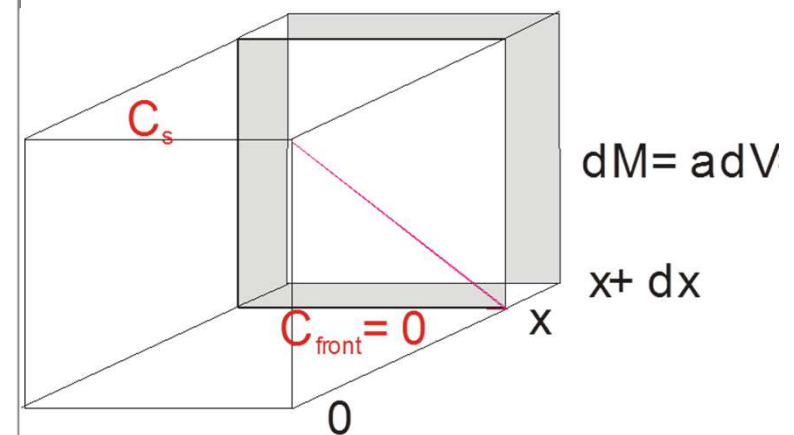
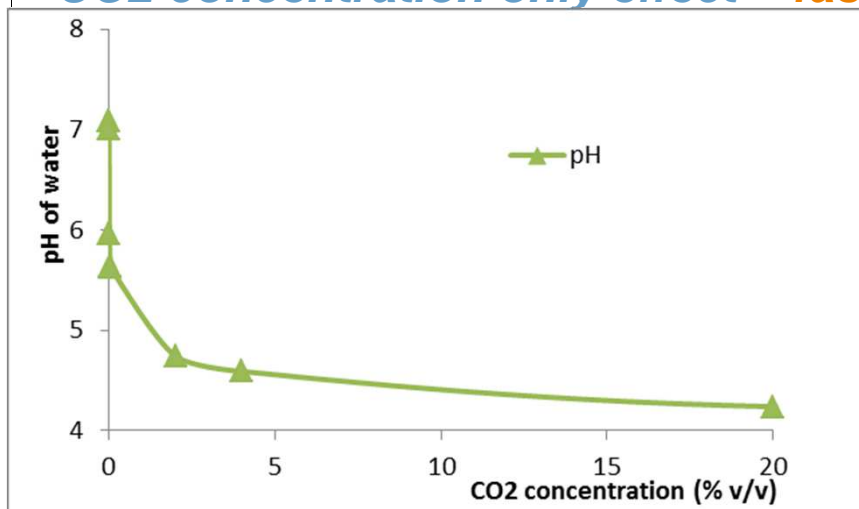
$$x_c = \sqrt{\frac{D}{a}} \sqrt{2ct} = \sqrt{1/R_{carb}} \sqrt{2ct}$$

E.g.
Acceleration concentr.: $c = 2\%$
Natural concentration: $c = 0.04\%$
Acceleration = $2/0.04 = 50 \times$



4.2 Carbonation

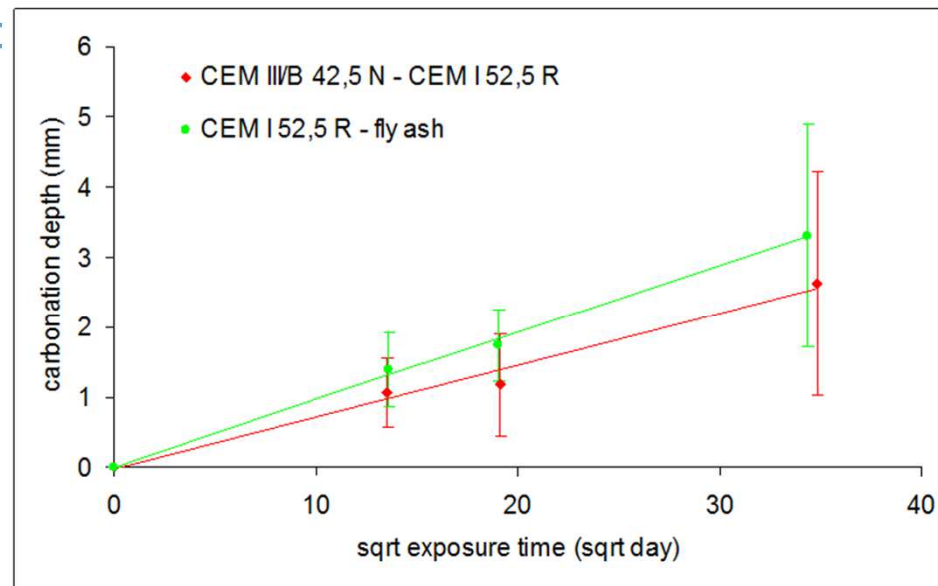
- Carbonation starts with **diffusion of CO₂-molecules**
- At the carbonation front, it **dissolves in the pore water**
- It is instantly **consumed by Ca²⁺ in the pore fluid**
- Buffer capacity of cement phases **releases new Ca²⁺**
 - First Ca(OH)₂ & high Ca CSH, next the other phases at succ. Lower pHs
- Concentration CO₂ at the front remains 0 until no more buffer
- CO₂ concentration only effect = **faster delivery CO₂ molecules**





4.2 Carbonation

- › Carbonation of each cement phase occurs at **phase stability pH**
- › **No cement phase is stable** below $\text{pH} = 7$ ($c(\text{CO}_2)$ approx. 0)
- › All cement phases react in a neutral way so can **go to completion**





4.2 Carbonation: FMEA

- › Carbonation of each cement phase occurs at **phase stability pH**
- › **No cement phase is stable** below $\text{pH} = 7$ ($c(\text{CO}_2)$ approx. 0)
- › All cement phases react in a neutral way so can **go to completion**
- › **All calcium is consumed**: buffer capacity = Ca content of cement





4.2 Degradation mechanisms: carbonation





Performance based service life design: durability of concrete

Conclusions:

1. Understanding all successive steps in the degradation mechanism is of importance; also from the viewpoint of prevention
2. Manipulating the structure of the concrete / cement paste is also a good way to either prevent or slow down degradation
3. There is however not one type of degradation: freeze-thawing, corrosion due to carbonation or chloride, ASR, sulfate resistance and so on may be of importance – a good assessment of the environment thus is important