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Mechanics and Design of Concrete Structures:

**Degradation, corrosion and analysis of
the durability of concrete structures.**

REINFORCED CONCRETE DETERIORATION:

1) PHYSICAL PROCESSES

- Icing-deicing cycles
- Fire

2) CHEMICAL PROCESSES

- Sulphatic degradation
- Degradation due to seawater

3) MECHANICAL PROCESSES

- Abrasion
- Erosion
- Cavitation
- Impact

4) STRUCTURAL PROCESSES

- Overloads
- Settlements
- Fatigue

5) REBARS CORROSION

PHYSICAL PROCESSES

Icing-deicing cycles

➤ From the surface to the center of the section

Micro and macro cracks due to:

➤ Bigger volume of ice

➤ Higher water pressure in the inner pores

Fire

➤ <100 C (212 F) increase volume

➤ 100 C (212 F)-150 C (302 F) Decrease of volume due to thermal decomposition

➤ >500 C (932 F) fractures at the interfaces with rebars

CHEMICAL PROCESSES

Sulphatic degradation

- Occurs in environments rich in sulphates (Grounds, waters)
- Reaction between sulphates and hydrates components of concrete
- New products with bigger molecular volumes

Seawater degradation

- Usually between the high-low tide zone
- Mechanical processes (superficial erosion)
- Crystallization of salt into the pores of concrete

REBARS CORROSION:CAUSES

CONCRETE IS THE IDEAL ENVIRONMENT FOR REBARS. $\text{pH} > 13$

REBARS ARE PASSIVATED UNTILL:

- **CARBONATION**

Due to the presence of CO_2 in the surrounding environment

2) CHLORIDES ACTION

Due to the presence of chlorides in the environment or in the concrete itself

CORROSION DUE TO CARBONATION

➤ CHANGE IN THE CONCRETE pH

Consumption of OH⁻ ions due to
 $\text{CO}_2 + \text{Ca}(\text{OH})_2 = \text{CaCO}_3 + \text{H}_2\text{O}$

**Extensive destruction of the passivating
film**

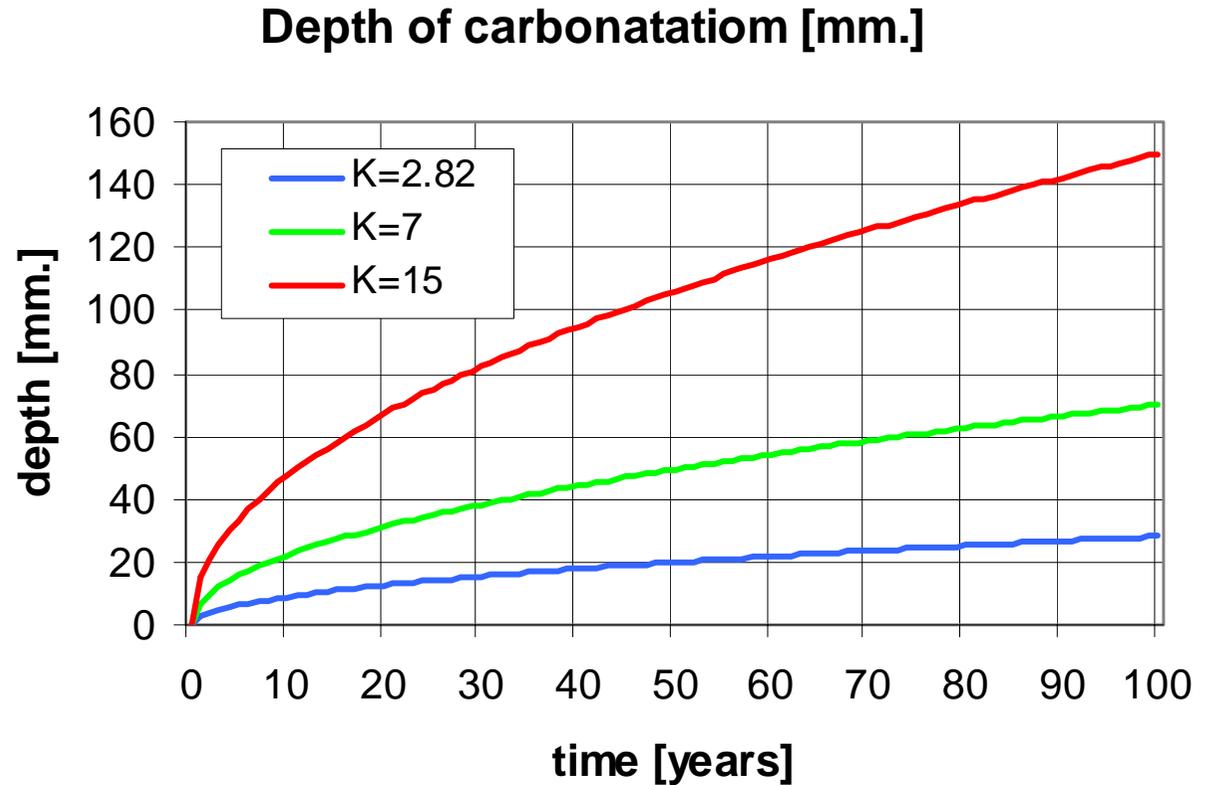
CORROSION DUE TO CARBONATION

Carbonation Depth:

Where K is the carbonation coefficient and it depends on:

- Concentration of CO₂
- External Environment
- Cement Used

$$s = K \cdot \sqrt{t}$$



CORROSION DUE TO CHLORIDES

- **High percentage of chlorides in either the external environment or in the concrete.**
- **The corrosion begins when the concentration of ions Cl^- reaches a critical value**
- **Corrosion localized to small zones of the rebar due to the circulation of electrons and ions.**

CORROSION DUE TO CHLORIDES

➤ DIFFUSION OF A CHEMICAL SPECIES
IN POROUS MEDIA:

$$\nabla^2 C = \frac{1}{D} \cdot \frac{\partial C}{\partial t}$$

➤ DIFFUSION OF CHLORIDES INTO
CONCRETE:

$$\nabla^2 C = \frac{1}{D_{eff}} \cdot \frac{\partial C}{\partial t}$$

D_{eff} is the effective diffusivity matrix
which is function of the concrete used .

➤ WITH THE SOLUTION:

$$C_x(t) = C_s \left(1 - \operatorname{erf} \frac{x}{2 \cdot \sqrt{D_{eff} \cdot t}} \right)$$

DESIGN LIFE OF A STRUCTURE

➤ CORROSION DUE TO CARBONATION

$$t = t_i + t_p = \left(\frac{d}{K} \right)^2 + \frac{D^{cr}}{V^{corr}}$$

Where:

d=Cover thickness

K=carbonation coefficient

D_{cr}=Critical depth of corrosion in the rebars

V_{corr}=velocity of corrosion

$$K = \frac{d}{\sqrt{t - \frac{D^{cr}}{V^{corr}}}}$$

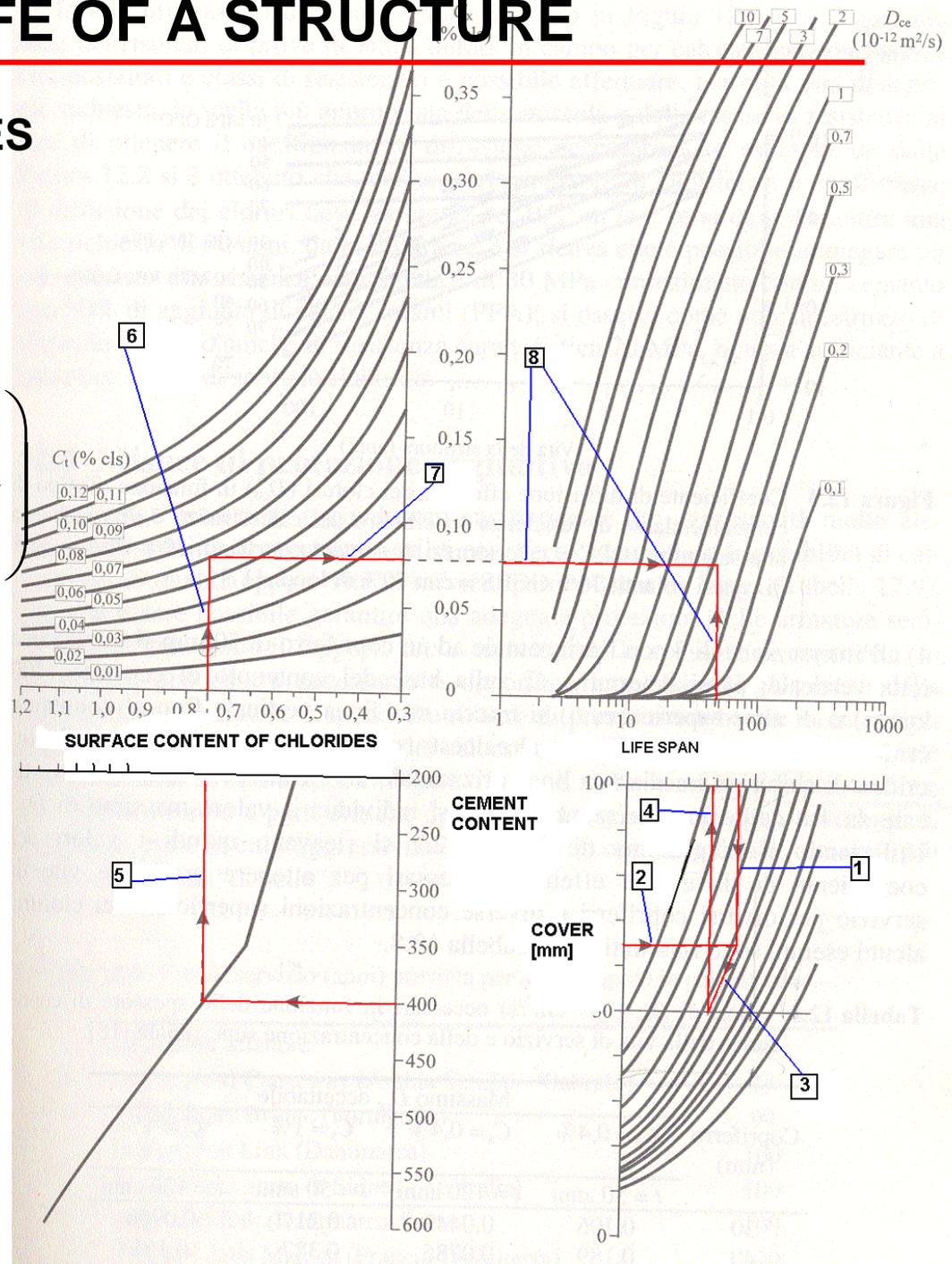
DESIGN LIFE OF A STRUCTURE

➤ CORROSION DUE TO CHLORIDES

In this case the design life is only function of t_i

$$C_x(t) = C_s \left(1 - \operatorname{erf} \frac{x}{2 \cdot \sqrt{D_{eff} \cdot t}} \right)$$

1. It is possible to find the effective diffusivity that will provide the design life as expected
2. Then the effective diffusivity is related to the concrete that must be used



QUESTIONS

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SELECTED REFERENCES

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