



American Concrete Institute®
Advancing concrete knowledge

Recent Advances in ASR Test Methods and Understanding Mitigation Mechanisms, Part 2

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Karen Scrivener graduated from University of Cambridge in 1979 in Materials Science. She went on to do a PhD on “The Microstructural Development during the Hydration of Portland Cement” at Imperial College, London completed in 1984. She remained at Imperial College until 1995 as Royal Society Research Fellow and then lecturer, heading the Cement and Concrete Group in the Department of Materials. In 1995 she joined the Central Research Laboratories of Lafarge near Lyon in France as Head of research on Calcium Aluminate cements and expert of concrete durability in general. In March 2001 she was appointed as Professor and Head of the Laboratory of Building Materials, Department of Materials at the Swiss Federal Institute of Technology at Lausanne (EPFL, Ecole Polytechnique Fédérale de Lausanne), Switzerland. She created and is co-ordinating NANOCEM – the industrial-academic research network on cement and concrete which brings together 15 industrial and 24 academic partners. She is Editor-in-Chief of the leading academic journal in the field – Cement and Concrete Research.



Understanding the role of Supplementary Cementitious Materials in mitigating Alkali Silica Reaction

Théodore Chappex
Karen Scrivener



Alkali Silica Reaction



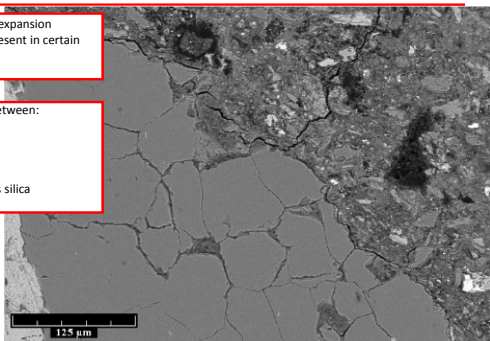
From J. Hildeter

Alkali Silica Reaction

Long-term expansion reaction present in certain concretes

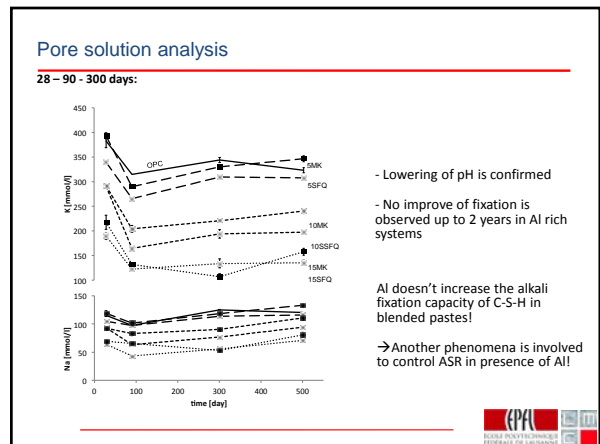
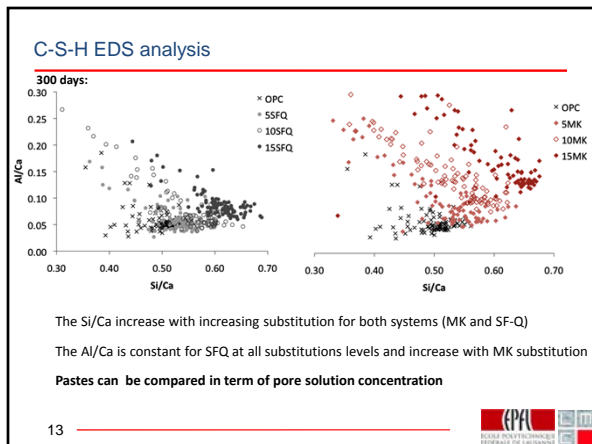
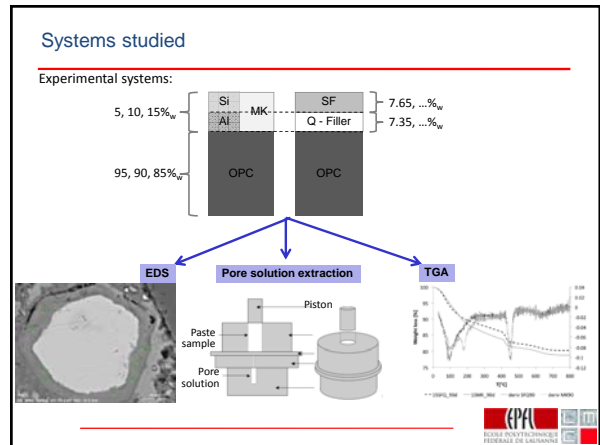
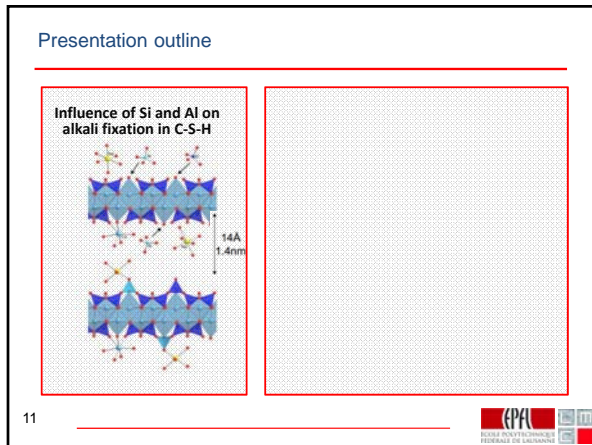
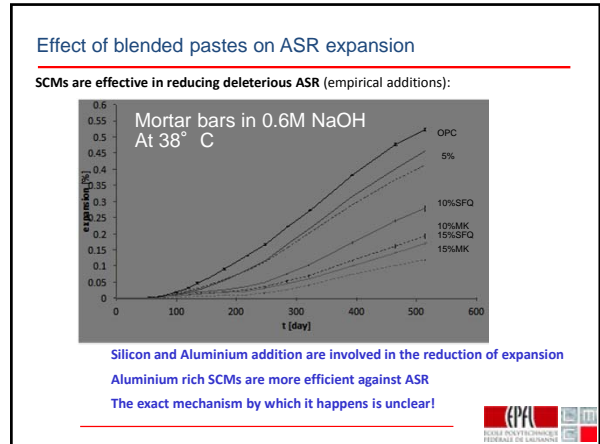
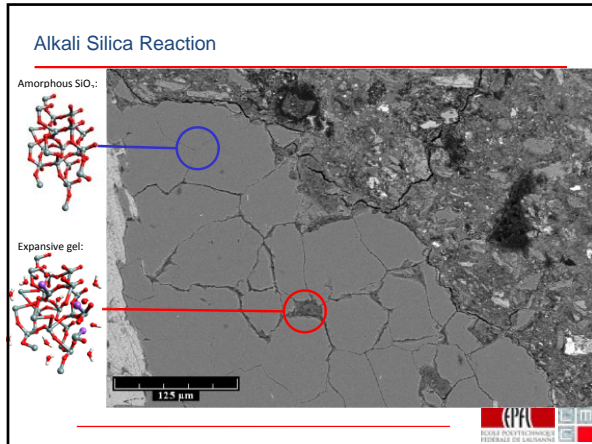
Reaction between:

- Alkali
- Water
- Amorphous silica



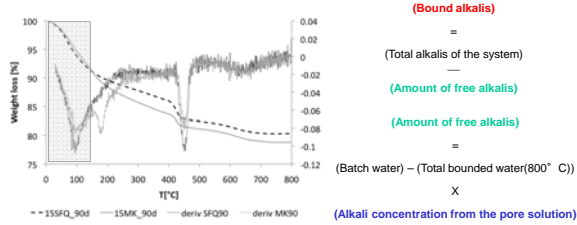
Alkali Silica Reaction



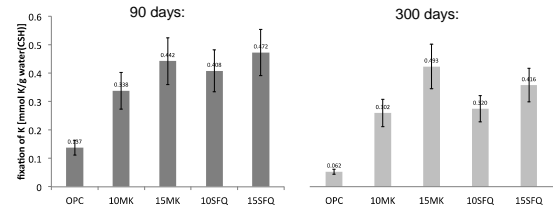


C-S-H fixation capacity estimation

Due to difficulties to quantify all the phases in presence in the blends (XRD/EDS/TGA), a comparative study was done, using the TGA water loss between 30 and 150° C, corresponding to the main C-S-H water loss.



C-S-H fixation capacity estimation: comparative

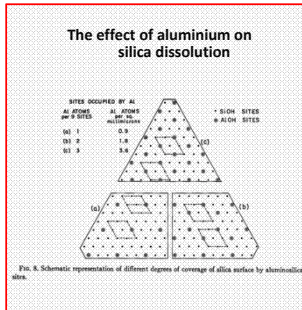
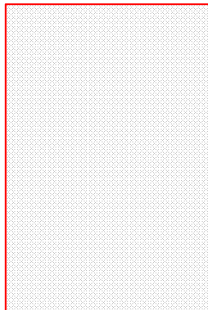


Comparatively, the fixation capacity of SFQ and MK are similar. Aluminium has no influence on the fixation capacity of alkalis

→ New approach: focus on the aggregates



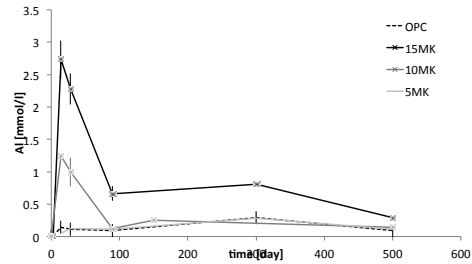
Presentation outline



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Pore solution composition of MK pastes



MK systems provide aluminium ions in the pore solution – A peak of aluminium appears during the first 90 days

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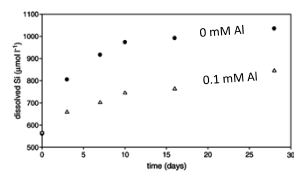
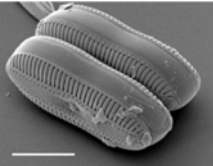
Silica dissolution in presence of aluminium ions

New approach: Focus on the aggregates

Marine chemistry and geology:

« Metal ions, particularly Al, reduce the rate of dissolution of amorphous silica » [Lewin, 1961]

Prepared frustules (amorphous silica) in sea water with and without Al: [Koning, 2007]



“only 1.35 Al atoms/nm² was adsorbed (less than one atom layer), and this was sufficient to minimize the solubility of silica”

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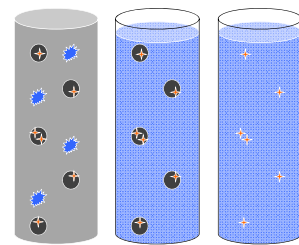


Multiscale approach

Is this mechanism present in concrete systems?

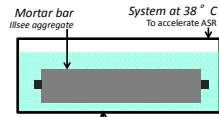
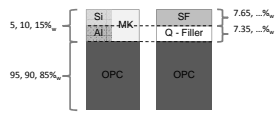
High alkalinity – Ca saturated

Multi scale approach: Concrete / Mortar, Reactive aggregates & pore solution, Amorphous silica & pore solution



Mortar expansion test

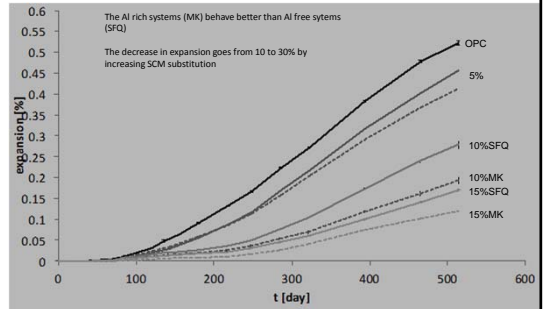
Concrete / Mortar



simulated pore solution
No leaching of alkalis, good reproducibility
0.6 M NaOH
Al(OH)₃ saturated



Mortar expansion test

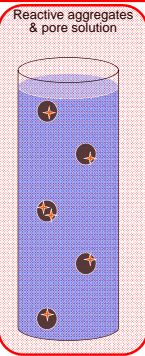
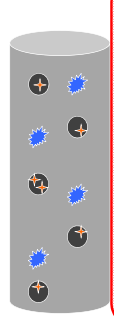


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The effect of Al on silica dissolution

Concrete / Mortar



Solutions:

- Al ions concentrations corresponding to blended pastes (0mM / 4mM / 10mM / 30mM / saturated)
- 38° C / 60° C
- Portlandite saturated
- 0.6 M NaOH

Aggregate:

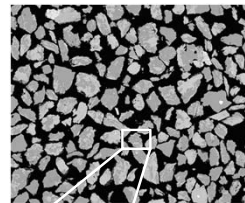
- Reactive Swiss Alps aggregates



Reactive aggregates test

Reacted fraction analysis (SEM-Image analysis):

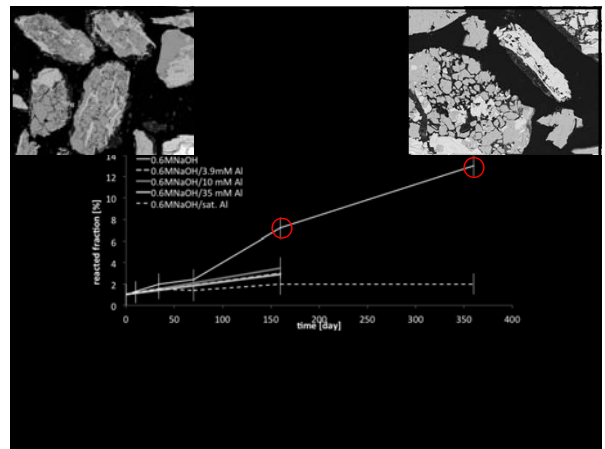
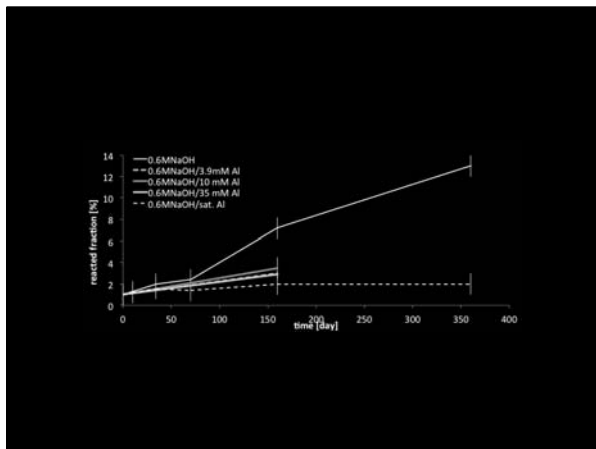
[Ben Haha & al.]

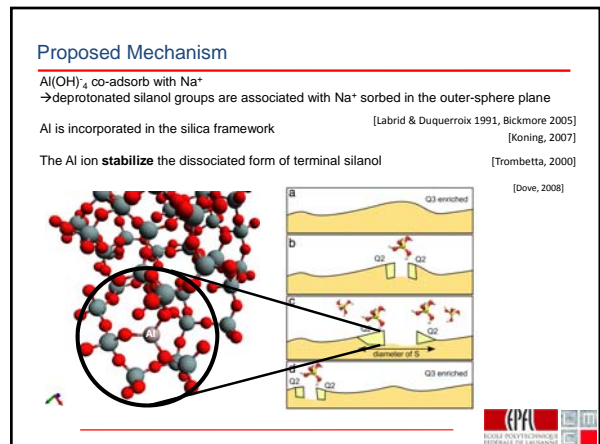
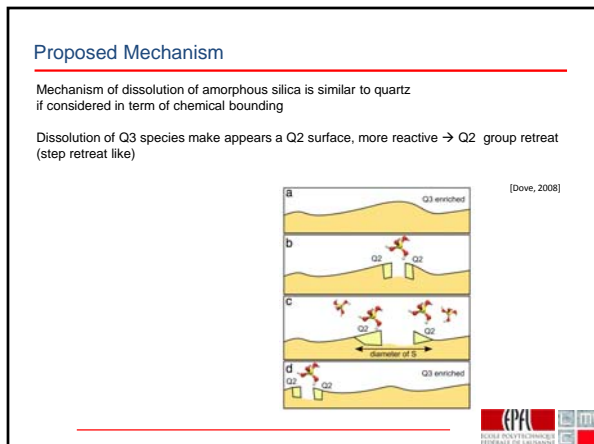
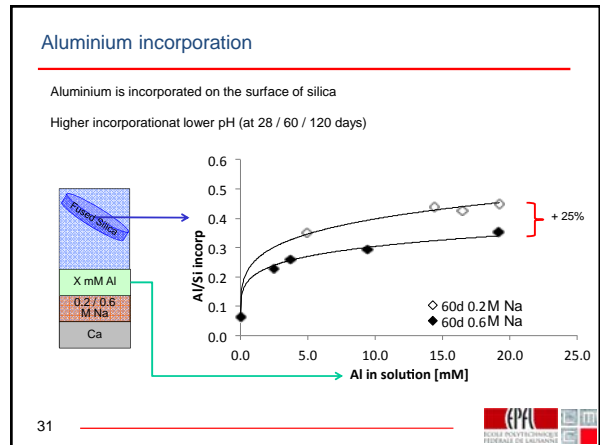
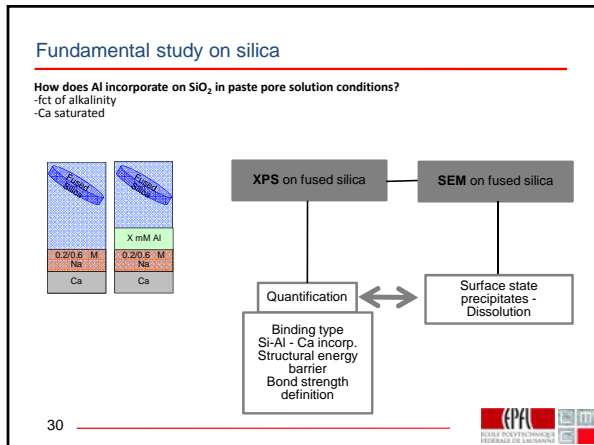
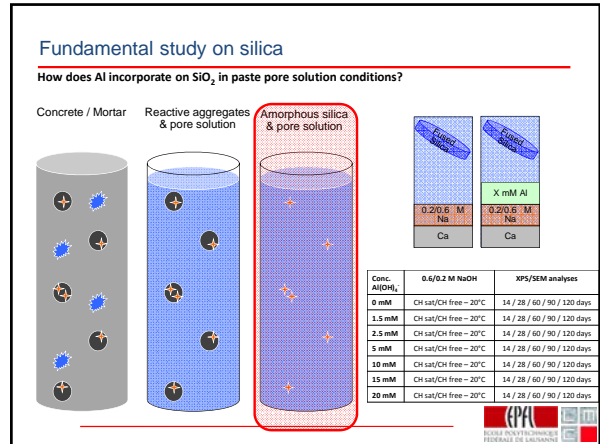
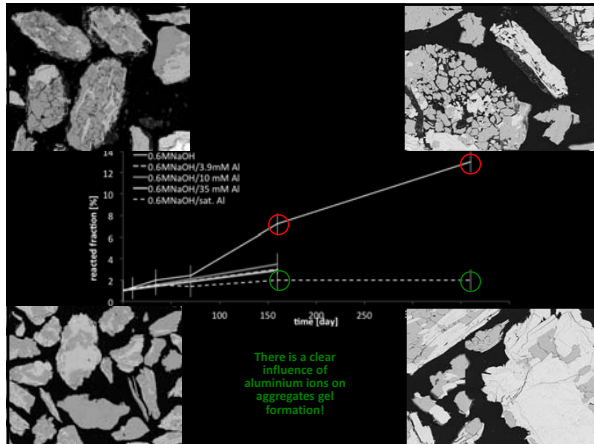


Percentage of gel pocket surface in function of total aggregate surface



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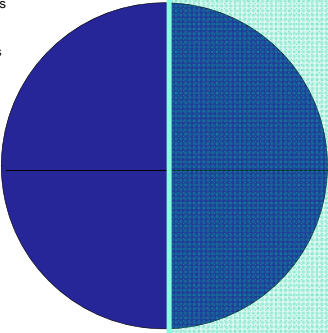





The effect of aluminium on silica dissolution – « 4 sides test »

The aim of this test is to provide the same initial surface state for all the treatments

Differences on the 4 sides were visible only after 90 days



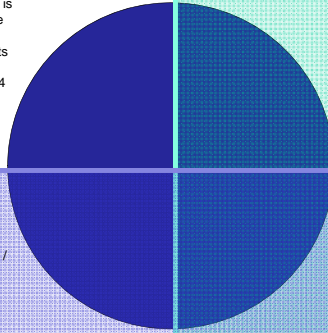
Step 1:
Pre treatment:
30 mM Al / 0.2M Na solution
90 days



The effect of aluminium on silica dissolution – « 4 sides test »


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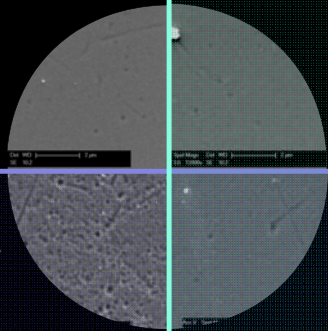


Step 1:
Pre treatment:
30 mM Al / 0.2M Na solution
90 days

Step 2:
Post treatment: 28 / 60 / 90 / 120 days
0.6M Na solution




90 days post treatment:



Step 1:
Pre treatment:
30 mM Al / 0.2M Na solution
90 days

Step 2:
Post treatment: 28 / 60 / 90 / 120 days
0.6M Na solution



Conclusion

It was showed that the **aluminium present in the C-S-H** of blended pastes **doesn't decrease the alkalinity** of the system

Al incorporation and effect was successfully showed for concrete condition



The mechanism controlling silica dissolution in presence of Al was partially explained and confirmed

Al is **incorporated in the silica framework** and probably stabilizes the step retreat dissolution of the Q2 species

The **alkalinity has an influence** on the aluminium incorporated fraction. The lower alkalinity present a higher aluminium incorporation
→ Aluminium effect has to be combined with an alkalinity reduction to be more efficient (visible on expansion curves?)

Approximation of dissolution rate? → prediction of gel formation rate → expansion!

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Acc.V Spot Magn Dat WD |-----| 2 µm
3.00 kV 3.0 12000x SE 10.2

Thank you

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