

Methodology of laboratory test for the description and comparison of the reactive and migration properties of new types of iron nanoparticles



Pešková Kristýna, Nosek Jaroslav, David R. Fargas, Černík Miroslav

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### Introduction

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Research is underway in the framework of the project NANOREM, aiming to develop new types of iron nanoparticles for the remediation of the rock environment that are stable in air and do not have to be stored in an inert atmosphere. This greatly simplifies handling of the material during remediation (transport and preparation onsite). The pyrophoric properties of these nanoparticles are suppressed by stabilizing their surface using an inorganic shell, which prevents rapid oxidation (degradation) on contact with air. The aim of presenting methodologies for laboratory tests is to confirm the influence of activation of new type of iron nanoparticles (NANOFER STAR) on their reactive and migration properties. The activation process involves leaving a prepared stock slurry in a concentrated state (usually 20% wt.) for 48 hours and causes disintegration of the inorganic surface shell.

## Methodology

**Migration tests:** details of methodology see Fig. 1. Vertical orientation and washing against the direction of gravity. Concentration of nZVI at the input to the column ca. 1,2 g/L. Concentration of total Fe in time measured in output water. pH and ORP measured continuously. At the end of test measured the concentration of total Fe in whole column. Breakthrough curves of each type of nZVI and distribution curves of the nZVI in the columns evaluated and compared with each other.

**Reactivity tests:** using reagent 100, 250, 500 ml bottles with a Teflon septum. Each sample container used for a single sample and therefore constitutes one time step of the kinetics. The reactor tests - an extension of the batch tests. Using a working stirred reactor with 2.5 litres of contaminated water into which the nZVI dosed. The decrease in the concentration of contamination after the application of the and changes in the pH, ORP monitored (CHC: GC/MS - Varian 3800/Saturn 2200; pH, ORP: WTW 3430i). Concentration and kinetic tests with water from sites.

celle ICP OES ORP Conductivity Fig. 1: Block diagram of the column test



Fig. 2: Left – Reagent bottles for kinetic batch tests; Right – reactors

# Results

### **Column tests:**

Type of nZVI	NF STAR 197 non-activated	NF STAR 197 activated
Fein (mg/l)	1890 mg/l	1713 mg/l
Total Fe on the output (mg) (%)	47 mg <i>(2 %)</i>	72 mg <i>(3%)</i>
Total Fe in the column (mg) (%)	2421 mg <i>(98 %)</i>	2592 mg <i>(97%)</i>
Length of sand bed (cm)	21,3	21
Pore volume of the column (ml)	25,2	27,0

#### Fig. 3: Overall balance of nZVI before and after activation process



#### **Batch tests:**





Fig. 6: Kinetic test with CHC using activated NF STAR with concentration 2 g/L and changes in phys.-chem. parameters

Fig. 4: Comparison of migration properties of non-activated and activated NF STAR and changes in physical-chemical parameters



Concentration of nZVI (g·l<sup>-1</sup>)



Fig. 7: Reactivity tests of non-activated and activated NF STAR through Cr(VI) to Cr(III) conversion and changes in phys.-chem. parameters

• The results of the performed tests show that the activated NF STAR iron nanoparticles are significantly more reactive and achieve better migration parameters. The disintegration of the inorganic surface shell causes that the particles have more negative Zeta-potential. This fact improve their migration and contribute to better distribution in the rock environment.



- The pilot and field tests confirmed that dilution Fig. 8: Changes in Zeta-potential to low application concentrations and dilution with the groundwater during transport in the rock environment does not lead to the

initial

Fig. 5: Reactivity tests with CHC using different concentrations of non-activated

and activated NF STAR and changes in phys.-chem. parameters

Concentration of nZVI (g·l<sup>-1</sup>)



disintegration of the inorganic surface shell (to the activation of the nanoparticles). This causes lower reactivity and increase of the remediation costs.

<u>Kristýna Pešková<sup>1</sup></u>, Jaroslav Nosek<sup>1</sup>, David R. Fargas<sup>2</sup>, Miroslav Černík<sup>1</sup>

kristyna.peskova@tul.cz, jaroslav.nosek1@tul.cz

<sup>1</sup>Technical University of Liberec, Institute for Nanomaterials, Advanced Technologies and Innovation (CxI),

Bendlova 1409/7, 461 17 Liberec, CZ

<sup>2</sup>Universtitat Politecnica de Catalunya, Calle Jordi Girona 31, Barcelona, Spain



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