

MODELLING MASS TRANSFER COEFFICIENTS AT LABORATORY SCALE FOR IN SITU SMARTSTRIPPING[®] UPSCALING

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INTRODUCTION



The groundwater remediation technology called SmartStripping[®] is an innovative process for groundwater in-situ remediation that reduces concentrations of Chlorinated Aliphatic Hydrocarbons (CAHs) dissolved in groundwater at industrial and civil sites and especially at sites with underground storage tanks.

The process can be defined as an innovative combination of Air Sparging (AS) and Soil Vapour Extraction (SVE): groundwater remediation occurs by enabling a transfer of contaminants from a saturated zone (groundwater) to an unsaturated zone (vadose) by blowing heated air from existing wells, which then enables groundwater stripping from the aquifer. The stripping allows the separation of CAHs from groundwater that vent up to the unsaturated zone which is under a continuous vacuum status, whereby the soil vapour is extracted. Vapours are treated with granular activated carbon (GAC) adsorption filters before being re-injected into the groundwater to start the stripping process again, through a continuous closed air-cycle system.

The application of this technology in each specific site needs the use of a comprehensive modelling and lab scale experiments for an optimal design of removal of volatiles as a function of operational parameters, from which air flow is the most relevant. In the present work, a combination of hydrodynamic and mass transfer model is developed and calibrated with specific laboratory tests.



EXPERIMENTAL SETUP



LAB SCALE SET UP



Experimental conditions tested:

- Sand (0.50-0.80 mm) saturated with water
- Air flow (stripping) = 1.3 L/min
- Several sampling points at 26 cm below surface
- Initial contaminant concentrations
 - Experiment 1: 55 mg/L Ethanol ۲
 - Experiment 2: 523 μ g/L mix-dichloroethylene (mix-DCE) \bullet



Container: sand saturated with water

MODELLING **INPUT PARAMETERS REQUIRES: MASS BALANCES INSIDE ZOI:** Diffusion Initial contaminant concentration in water from outside Contaminant properties: GAC filter the ZOI - Henry's law constant \longrightarrow Bibliographic data C: liquid concentration of contaminant, S: surface of the ZOI in dC $S \cdot K^* \cdot (C_o - C)$ r In the liquid phase: which lateral diffusion takes place, K*: liquid mass transfer Kglobal Parameters to be coefficient due to the lateral diffusion, C₀: Initial contaminant dt V_r·ε_L εL - K* concentration, V_r : Volume of the ZOI, ε_1 : water porosity in the ZOI calibrated dp r∙⁄R∙T G∙p p: contaminant partial pressure, G: air flow, R: ideal gas \succ Zone of influence data In the air phase: constant, T: temperature, ε_{G} : air porosity in the ZOI dt V_r.ε_A **ZONE OF** ε_A - Volume INLUENCE Obtained from - Surface (ZOI) hydrodynamic model r: rate of stripping, K_{G} : mass transfer coefficient in the gas developed with - Fraction of air phase, a: contact surface per unit of volume, p*: **STRIPPING:** ANCVC CEV® 110

| | contaminant partial pressure in the interface, p: contaminant partial pressure in the center of the bubble, K ₁ : | | -raction of water | ANSYS-CFX° V14.0 |
|--|---|-----|-------------------|------------------|
| | $r = K_G a(p^* - p) = K_L a(C - C^*) = K_{global}(C - p/H)$ contaminant partial pressure in the center of the bubble, K_L : mass transfer coefficient in the interface, C: concentration in | 1 C | | |
| | the bulk liquid, K _{global} : overall mass transfer coefficient, H: | 1 | | |
| | Henry's Law constant | 1.1 | | |
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PARAMETERS CALIBRATION



CONCLUSIONS

✓ The experimental setup designed and applied in this work is useful for calibration of parameters and proper simulation of the stripping process at lab scale.



