



## RESEARCH LETTER

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## Key Points:

- First geophysical monitoring of a neutrally buoyant and electrically conductive tracer
- Comparisons between neutrally buoyant and dense tracers highlight very different dynamics
- Neutrally buoyant tracers make geophysical monitoring experiments compatible with hydrogeological tracer tests

## Correspondence to:

A. Shakas,  
alexis.shakas@unil.ch

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Neutrally buoyant tracers in hydrogeophysics:  
Field demonstration in fractured rock

Alexis Shakas<sup>1</sup> , Niklas Linde<sup>1</sup> , Ludovic Baron<sup>1</sup>, John Selker<sup>2</sup>, Marie-Françoise Gerard<sup>3</sup>, Nicolas Lavenant<sup>3</sup>, Olivier Bour<sup>3</sup>, and Tanguy Le Borgne<sup>3</sup>

<sup>1</sup>Applied and Environmental Geophysics Group, Institute of Earth Sciences, University of Lausanne, Lausanne, Switzerland, <sup>2</sup>Department of Biological and Ecological Engineering, Oregon State University, Corvallis, Oregon, USA, <sup>3</sup>Géosciences Rennes, Université de Rennes 1, CNRS, UMR 6118, Rennes, France

**Abstract** Electrical and electromagnetic methods are extensively used to map electrically conductive tracers within hydrogeologic systems. Often, the tracers used consist of dissolved salt in water, leading to a denser mixture than the ambient formation water. Density effects are often ignored and rarely modeled but can dramatically affect transport behavior and introduce dynamics that are unrepresentative of the response obtained with classical tracers (e.g., uranine). We introduce a neutrally buoyant tracer consisting of a mixture of salt, water, and ethanol and monitor its movement during push-pull experiments in a fractured rock aquifer using ground-penetrating radar. Our results indicate a largely reversible transport process and agree with uranine-based push-pull experiments at the site, which is in contrast to results obtained using dense saline tracers. We argue that a shift toward neutrally buoyant tracers in both porous and fractured media would advance hydrogeophysical research and enhance its utility in hydrogeology.

**Plain Language Summary** Geophysical techniques allow for nondestructive monitoring of groundwater processes provided that contrasts in physical properties are available through time and space. To achieve this contrast, the common approach is to inject saline water (a so-called saline tracer) into the ground and to subsequently monitor its movement using geophysical measurements. Unfortunately, classical saline tracers are denser (thus heavier) than fresh groundwater, which leads to instabilities and downward movement of the tracer. This implies that the information gained from saline tracer tests might be unrepresentative of natural groundwater conditions. We introduce a saline tracer that has the same density as the ambient groundwater by mixing the saline tracer with ethanol. This enables, for the first time, geophysical imaging of groundwater flow and transport without associated density effects. By considering experiments performed in fractured rock, we demonstrate (1) that previously used dense saline tracers used for geophysical monitoring lead to very strong density effects, (2) that the new tracer solution can be imaged very well using geophysical techniques, and (3) that the recovered tracers (the so-called breakthrough curves) agree well with traditional tracer tests and hydrogeological theory.

## 1. Introduction

Geophysics enables remote monitoring and imaging of subsurface mass transfer at scales ranging from decimeters [e.g., Garré *et al.*, 2011; Beff *et al.*, 2013], to meters [e.g., Slater and Sandberg, 2000; Singha and Gorelick, 2005], kilometers [e.g., Falgàs *et al.*, 2009; Rosas-Carbajal *et al.*, 2015], and beyond [e.g., Zhdanov *et al.*, 2011]. The need for methodological developments that ensure appropriate integration of geophysical data in subsurface hydrology have given rise to the research field of hydrogeophysics [Hubbard and Linde, 2011; Binley *et al.*, 2015], which has had an impact on hydrology over the last decade [National Research Council, 2012]. In hydrogeophysics, geophysical experiments are made to support hydrological research and applications. This implies not only that geophysical data should be informative of the processes being studied but also that its acquisition should not perturb hydrological data or significantly affect the design of hydrological experiments.

In order to image a tracer (or contaminant plumes) with geophysics, there must exist a naturally occurring or imposed contrast in physical properties between the tracer (contaminant) and the surrounding formation water. When such a contrast is present, geophysical imaging can provide insight into the transport

processes that take place in the hydrogeological system. In situ imaging of transport processes with geophysics may thus help to unravel complex processes, such as anomalous transport, dual-domain mass transfer, or reversible/irreversible dispersion, that are often difficult to infer from breakthrough curve analysis alone [Swanson *et al.*, 2012, 2015]. In groundwater geophysics, the contrast agent for tracer tests is often dissolved table salt (NaCl) [Day-Lewis *et al.*, 2003; Singha and Gorelick, 2005; Doetsch *et al.*, 2012; Shakas *et al.*, 2016].

Salt increases the electrical conductivity and enables tracking of tracer plumes using electrical [Kemna *et al.*, 2002; Singha and Gorelick, 2005], induction-based electromagnetic [e.g., Falgàs *et al.*, 2009; Rosas-Carbajal *et al.*, 2015], or high-frequency electromagnetic [Day-Lewis *et al.*, 2003; Tsoflias and Becker, 2008] methods. Note that the studies mentioned above have been conducted in both fractured and porous-media systems. The salinity contrast needed for reliable geophysical imaging implies that the saline solution is significantly denser than the surrounding water, which results in buoyancy-induced tracer movement; this has been verified in both laboratory experiments [e.g., Istok and Humphrey, 1995] and numerical tests [e.g., Beinhorn *et al.*, 2005; Kemna *et al.*, 2002]. Doetsch *et al.* [2012] provide a field demonstration of density effects using time-lapse electrical resistivity tomography. In accordance with numerical modeling, they found that a tracer injected in a gravel aquifer rapidly plunged to the underlying clay aquitard. Previous field experiments with lower salinity contrasts (and less density effects) at the site did not enable reliable time-lapse inversion results.

In the hydrogeophysics literature, it is rare to find field-based studies in which density effects are assessed or accounted for [e.g., Doetsch *et al.*, 2012; Shakas *et al.*, 2016; Haaken *et al.*, 2016]. Even if density effects are considered in the modeling and associated inversion, the ubiquitous use of dense saline tracers in hydrogeophysics is problematic as they change the system dynamics [Tenchine and Gouze, 2005]. That is, the use of geophysics imposes constraints on experimental design that might be unacceptable for field hydrogeologists. This implies (1) that hydrogeologists might be reluctant to consider geophysics in their work if they feel that hydrological experiments will be compromised by using dense tracers, (2) that comparisons between hydrogeophysical results and hydrogeological experiments using nonsalt tracers are difficult, and (3) that the inferred system properties and hydrological processes might have low predictive capacity in describing natural flow dynamics.

In this work, we introduce a neutrally buoyant tracer based on a water-salt-ethanol mixture that we refer to as wethanalt. Ethanol is fully miscible and has a similar viscosity as water. It is less dense than water and can be used to ensure that the resulting tracer solution is neutrally buoyant, while it maintains a high electrical conductivity with respect to the formation water. Furthermore, ethanol has the distinct advantage of being nontoxic [Thakker, 1998] and biodegradable [Schaefer *et al.*, 2010].

We present two field experiments in which we demonstrate the value of using neutrally buoyant and electrically conductive tracers for imaging transport processes. We consider push-pull experiments at a well-characterized fractured rock site with geophysical monitoring using single-hole ground-penetrating radar [Dorn *et al.*, 2011, 2012]. Our results are compared with previous experiments at the site that were carried out using dense saline solutions [Shakas *et al.*, 2016] and a traditional push-pull experiment using a fluorescent tracer, without geophysical monitoring [Kang *et al.*, 2015].

## 2. Methodology

The properties of ethanol-water and salt-water mixtures have been tabulated [Haynes, 2016]. To the best of our knowledge, such a laboratory study does not exist for wethanalt (ethanol-water-salt mixtures). Here we present a practical method for obtaining a neutrally buoyant wethanalt mixture by utilizing the existing tables as a guide and further fine-tuning the density in the field. In the following section, where not otherwise noted, the material properties of ethanol, water, and salt are taken from Haynes [2016].

### 2.1. Wethanalt Properties

Ethanol ( $C_2H_6O$ ) has a density of  $0.789\text{ g cm}^{-3}$ , relative electrical permittivity of 25.3 (zero-frequency limit), and dynamic viscosity of  $1.203\text{ mPa s}$  at  $20^\circ\text{C}$ . At the same temperature, demineralized water ( $H_2O$ ) has a density of  $1\text{ g cm}^{-3}$ , relative electrical permittivity of 81 and dynamic viscosity of  $1.004\text{ mPa s}$ . Both liquids are electrically resistive, and it is the addition of salt to the mixture that will determine the electrical conductivity. The most common choice of salt in hydrogeophysical applications is sodium chloride (NaCl), that dissociates into  $Na^+$  and  $Cl^-$  ions when dissolved in water.

Ethanol and water are miscible in all proportions, and their mixing results in an exothermic reaction [Peeters and Huyskens, 1993] which leads to an increase in temperature when the mixture is prepared. Another property of ethanol-water mixtures is that the dynamic viscosity of the mixture is increased compared with the constituents. The maximum viscosity of the mixture is 2.85 mPa s (20°C) when the mass proportion of ethanol to water is 0.42:0.58. Ethanol does not pose any health risks when diluted with water [Thakker, 1998] and is biodegradable [Schaefer et al., 2010], which implies that it may increase microbial activity when used as a tracer.

## 2.2. Wethanalt Preparation

The preparation of a neutrally buoyant wethanalt solution is complicated by the facts (1) that the density of an ethanol-water mixture does not average arithmetically when adding salt and (2) that the necessary precision in density needs to be sufficiently low (for instance, Istok and Humphrey [1995] perform a laboratory study and report density effects for density variations ( $\Delta\rho$ ) in the range  $0.0075\% \leq \Delta\rho \leq 0.15\%$ ). To achieve this, our preferred field procedure is to first rely on tabulated values in Haynes [2016] to obtain a desired density of an initial ethanol-water mixture and then assume arithmetic averaging to predict the necessary amount of salt to add in order for the density to be equal to that of the formation water. We then prepare a wethanalt mixture with 5% less salt than predicted with this simple model. For our experiments we first mixed 85 L of demineralized water with 25 L of (99.98%) ethanol; this resulted in an increase in temperature of 8°C. The viscosity of the mixture was 2.26 mPa s according to Haynes [2016]. We then pumped formation water through a plastic tube that was coiled in the ethanol-water container, in order to reduce the temperature of the mixture to the ambient water temperature. Using this approach, we successfully reduced the temperature of the tracer mixture in all wethanalt experiments from  $\sim 24^\circ\text{C}$  to  $\sim 18^\circ\text{C}$ , while the ambient water temperature is  $16^\circ\text{C}$ .

To achieve a neutrally buoyant solution, we relied on Archimedes' principle, namely, that "Any object, wholly or partially immersed in a fluid, is buoyed up by a force equal to the weight of the fluid displaced by the object" [Archimedes (250 BC), 1897]. To do so, we used two containers: the first filled with the wethanalt mixture and the second containing formation water. In the second container, we submerged a balloon that we carefully filled with formation water and allowed for any air bubbles to escape. The balloon weighed 3.5 kg and was slightly positively buoyant, so we further adjusted its weight with plastic O rings (3 g each with a density of  $1.1\text{ g cm}^{-3}$ , resulting in a net submerged weight of 0.3 g per O ring) until we reached neutral buoyancy (i.e., not observing any vertical movement of the balloon when suspended in the middle of the water-filled container). The weight adjustments were made to a precision below  $0.1\text{ g kg}^{-1}$ . We then transferred the balloon with the attached O rings to the wethanalt container, in which we had mixed an initial amount (4 kg) of salt in the ethanol-water mixture, well below the amount (4.22 kg) predicted from arithmetic averaging. We then proceeded to add salt in increments between 20 and 80 g, until the balloon was neutrally buoyant in the wethanalt mixture. In the final stage, a total of 4.44 kg of salt was added and the mixture was stirred with a mixing propeller to ensure well-mixed conditions. This procedure allows us to obtain a wethanalt solution that is at most 0.01% different in density than the formation water. While ethanol is biodegradable [Schaefer et al., 2010], it also does not pose any substantial health risks after sufficient dilution [Thakker, 1998].

## 2.3. Setup of the Field Experiment

We performed the wethanalt push-pull experiments in a well-characterized fractured granitic aquifer located in Brittany, France (<http://hplus.ore.fr/en>). Previous studies indicate that flow at the site is dominated by a few, highly transmissive fractures [Le Borgne et al., 2007; Dorn et al., 2011]. All the experiments were performed in two adjacent boreholes, B1 and B2, that are separated by  $\sim 6\text{ m}$ . A double-packer system isolated a fracture intersecting the B1 borehole at 77.8 m, in which we injected the tracer followed by an almost equal volume of formation water (chaser). We then reversed the flow in order to retrieve the tracer, either immediately (push-pull) or after a waiting period (push-wait-pull) during which the pumps were off.

Ground-penetrating radar (GPR) monitoring took place at 5 cm intervals between 60 m and 85 m depth along borehole B2, which was isolated with a borehole liner [Shakas et al., 2016]. In an effort to further separate the direct wave from the reflections of interest, we reduced the transmitter-receiver offset from 4 m by Shakas et al. [2016] to 3.2 m. The final images account for this offset during migration.

## 2.4. Data Processing

### 2.4.1. Tracer Breakthrough Curves

We transformed electrical conductivity values, measured using a conductivity-temperature-depth (CTD) diver located at the outlet of the pump used for pulling, into salt concentration [Sen and Goode, 1992]. We then removed the background concentration and normalized the data to the injected tracer concentration, also with the background concentration removed. We also shifted the breakthrough data to account for the time it takes for the water to flow through the tubing from the packer to the outflow location. The experiments using the dense saline tracer published by Shakas *et al.* [2016] were performed in 2014 and the wethanalt experiments in 2016. For both field campaigns, we performed a series of tracer experiments. This implies that any unrecovered salt may lead to an increasing background concentration over time. To account for this, we present the uncertainty in the breakthrough curves (BTCs) with a thickness (see Figures 1i, 2i, and 3) obtained by varying the considered background concentration between a minimum (ambient concentration at the beginning of the field campaign) and a maximum (initial concentration at the beginning of each experiment). In order to compare experiments with different tracer/chaser volumes and flow rates, we normalize the time of each experiment with the theoretical peak arrival time. In an ideal push-pull experiment [Neretnieks, 2007], the theoretical peak arrival time ( $t_{\text{peak}}$ ) measured from the onset of pulling depends on the duration of both injection ( $t_{\text{injection}}$ ) and chasing ( $t_{\text{chasing}}$ ) and is given by  $t_{\text{peak}} = t_{\text{injection}}/2 + t_{\text{chasing}}$ . The same formula applies to push-wait-pull experiments in the absence of ambient flow.

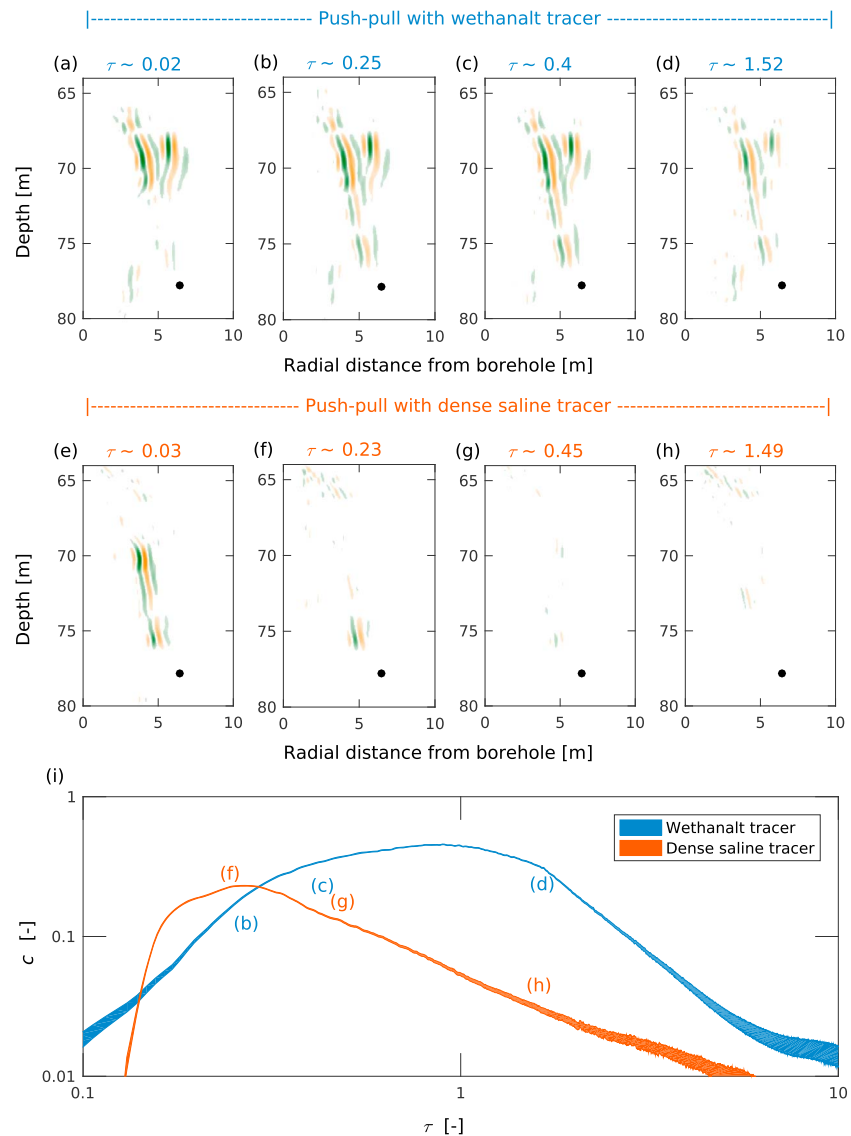
### 2.4.2. Migrated GPR Difference Sections

The processing of the GPR data is described in Shakas *et al.* [2016]. To each GPR trace, we apply a bandpass filter with a frequency window between 20 and 200 MHz (the emitted signal is centered at 100 MHz) followed by minor time shifts to align collocated traces. Compared with Shakas *et al.* [2016], the only difference in the processing was the use of singular value decomposition (SVD) to remove the direct wave. We accomplish this by decomposing each GPR section that we then reconstruct without the first singular value, which corresponds mainly to the direct wave. We then take the difference of each section and the reference (taken before the initiation of the push-pull experiment), apply a time gain, and finally use a Kirchhoff migration algorithm with a constant velocity model of  $v = 0.12 \frac{\text{m}}{\text{ns}}$ . This results in migrated GPR difference sections where the presence of the conductive tracer manifests itself as alternating (green-orange) stripes, whose horizontal extent is caused by the finite size of the source wavelet [Shakas *et al.*, 2016]. For visualization purposes, we suppress any reflections that are below the estimated noise level of our GPR data set (computed as 15% of the maximum amplitude).

## 3. Results

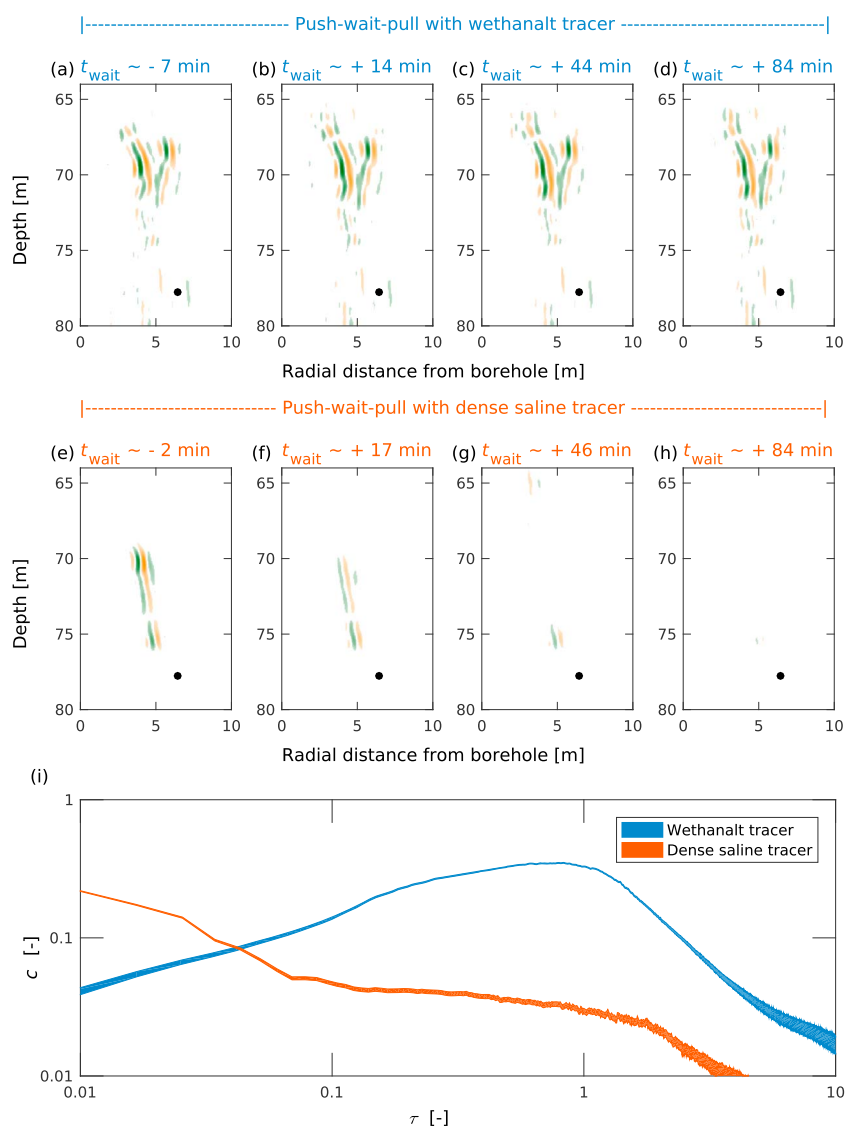
We now compare the BTCs and the migrated GPR difference sections obtained from the combined experiments with a saline tracer and wethanalt. Here we consider both push-pull and push-wait-pull setups. For the BTCs, we use a normalized time,  $\tau = t/t_{\text{peak}}$ , where  $t$  corresponds to the time after the onset of pulling and  $t_{\text{peak}}$  is the theoretical peak arrival time. We also use a normalized concentration,  $c = C/C_0$  that corresponds to the measured concentration ( $C$ ) divided by the initial tracer concentration ( $C_0$ ), after removing the background concentration from both. It takes about 3 min to acquire GPR data over the considered depth range, so each GPR section is indicated by an approximate time. The main experimental parameters are listed in Table 1.

A comparison between the push-pull experiments is made in Figure 1, where all time references are made (in normalized time) to the onset of the pulling phase. Figure 1 displays representative migrated GPR difference sections for the wethanalt (Figures 1a–1d) and saline (Figures 1e–1h) push-pull experiments, and the BTCs are plotted in Figure 1i. At the onset of pulling, the wethanalt tracer (Figure 1a) is localized within a depth range of 67 m to 72 m, while the saline tracer (Figure 1e) is found within 69 m to 76 m. At  $\tau$  close to 0.25, the wethanalt tracer (Figure 1b) is found over the same depth range while the dense saline tracer (Figure 1f) has quickly migrated down toward the injection location. These results are in accordance with the peaks of the BTCs that occur with  $\tau = 0.9$  and  $c = 0.45$  for wethanalt, while  $\tau = 0.27$  and  $c = 0.23$  for the dense saline tracer. This indicates that the measured peak arrival is close to the theoretical peak when using wethanalt, while it is much smaller when using a dense saline tracer. Shakas *et al.* [2016] demonstrated through modeling that this early peak arrival of the dense saline tracer was a consequence of density effects and the geometry of the fracture network. Reflections from the wethanalt tracer remain visible, from 67 m to 77 m, at much later times (Figure 1d) than the peak arrival measured in the borehole.



**Figure 1.** Results from two separate push-pull experiments using either wethanalt or a dense saline tracer (experiments a and c in Table 1, respectively). The migrated GPR difference sections for (a–d) wethanalt and (e–h) dense saline tracers are presented at similar normalized acquisition times ( $\tau$ ), where the black circle corresponds to the tracer injection location. The corresponding breakthrough curves are plotted in logarithmic scale (i) as a function of normalized time and normalized concentration ( $c$ ). The resulting uncertainty due to the background salt concentration is indicated by the thickness of each curve.

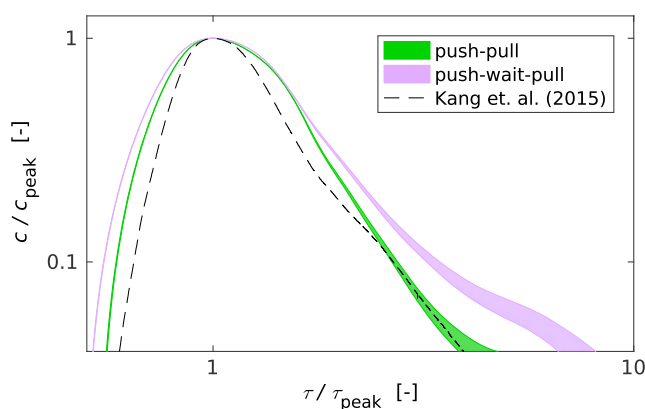
The push-wait-pull experiments are presented in Figure 2. All migrated GPR difference sections shown were acquired during the waiting time, so their acquisition times are referenced in minutes from the onset of waiting ( $t_{\text{wait}}$ ). After the pushing phase (Figures 2a and 2e), the distribution of both tracers is similar to the push-pull experiment (cf. Figures 1a and 1e), thereby indicating a strong reproducibility of the experiments. After 17 min, the saline tracer has sunk considerably (Figure 2f) due to its high density and is thereafter hardly detectable (Figures 2g and 2h). On the contrary, the migrated GPR difference sections for the wethanalt tracer remain almost identical during the whole waiting time (Figures 2a–2d). Once more, the peak arrivals of the BTCs support these observations with  $\tau = 0.83$  and  $c = 0.35$  for wethanalt, while  $\tau = 0.008$  and  $c = 0.05$  for the dense saline tracer. Again, the peak arrival of the dense tracer occurs much earlier than predicted from the theoretical peak arrival time. Later GPR difference sections (not shown) confirm that the wethanalt tracer is still visible in the migrated GPR difference sections at times twice as long as the theoretical peak arrival.



**Figure 2.** Results from two separate push-wait-pull experiments using either wethanalt or a dense saline tracer (experiments b and d in Table 1, respectively). The migrated GPR difference sections for (a–d) wethanalt and (e–h) dense saline tracers are presented at similar acquisition times referenced from the waiting phase ( $t_{\text{wait}}$ ), where the black circle corresponds to the tracer injection location. The corresponding breakthrough curves are plotted in logarithmic scale (i) as a function of normalized concentration ( $c$ ) and normalized time ( $\tau$ ) since the initiation of pulling. The resulting uncertainty due to the background salt concentration is indicated by the thickness of each curve.

#### 4. Discussion

The migrated GPR difference sections of the two push-pull experiments (Figure 1) suggest that different spatial regions of the fractured system are probed when using a neutrally buoyant or a dense saline tracer. The saline tracer remains closer to the injection location and is not pushed much further by chasing. This is expected since we are trying to displace a dense tracer with lighter formation water. Moreover, it is also in accordance with flow and transport simulations [Shakas et al., 2016; Haaken et al., 2016]. When wethanalt is used as a tracer, the chaser effectively pushes the tracer away from the injection location (Figure 1a) and into upper regions of the fractured system. By doing so, the migrated GPR difference sections from the wethanalt tracer probe an additional fracture that appears above 70 m depth and beyond 5 m radius (Figures 1a–1d). This fracture is not present in the migrated GPR difference sections acquired with the dense saline tracer (Figures 1e–1h).



**Figure 3.** Inferred impulse-response breakthrough curves for the wethanalt experiments (experiments a and b in Table 1) obtained by deconvolution. The uncertainty associated with the unknown background salinity is represented by the thickening of the lines at late times. For comparison purposes, we also plot the breakthrough curve from the push-pull experiment by *Kang et al.* [2015] described in Table 1.

In the push-wait-pull experiments (Figure 2), the impact of density is even more evident. While the tracer distribution after pushing is similar to their push-pull counterparts (compare Figures 1a with 2a and 1e with 2e), the dense saline tracer quickly migrates toward the injection location during the waiting period (Figures 2f–2h). On the contrary, the wethanalt tracer provides consistent, high-amplitude reflections throughout the waiting period (Figures 2a–2d), suggesting that the tracer distribution remains the same during this time. This indicates that the sinking observed when using a dense saline tracer is primarily due to density effects and not to ambient flow, as the ambient flow should also affect the wethanalt tracer. This finding is further supported by the similar peak arrival times of the wethanalt tracer in the push-pull and push-wait-pull configurations ( $\tau = 0.9$  and  $\tau = 0.83$ , respectively). Nevertheless, the waiting period allows for more diffusion of the wethanalt tracer and possibly some ambient flow effects that manifest as a decrease of the peak concentration in the BTC between the push-pull ( $c = 0.45$ ) and push-wait-pull ( $c = 0.35$ ) configurations.

To further assess the suitability of wethanalt as a neutrally buoyant tracer, we compare our BTCs with the push-pull BTC by *Kang et al.* [2015], performed at the same fracture location but with an almost instantaneous injection of a neutrally buoyant fluorescein tracer (see Table 1 for parameters). To remove the imprint of the injection period from the wethanalt BTCs, we model the push-pull experiment as the convolution of a linear, time-invariant source operator with the impulse-response of the system. We infer the impulse-response using an iterative least squares inversion [*Menke, 2012*] with smoothness and positivity constraints [*Cirpka et al., 2007*]. Convergence is reached when the positivity constraints stop changing. In order to compare the late-time slope of the BTCs, we normalize concentration to the peak concentration and time to the peak arrival time and plot the BTCs in Figure 3. The push-pull experiment shows a late-time tailing comparable to the BTC from *Kang et al.* [2015], while the push-wait-pull experiment indicates a smaller slope. This indicates that the wethanalt BTCs are consistent with the fluorescein BTCs, in particular regarding the late-time concentration decay, which is important for investigating anomalous transport and dual-domain mass transfer processes.

**Table 1.** Experimental Parameters for (a) Push-Pull and (b) Push-Wait-Pull With Wethanalt, (c) Push-Pull and (d) Push-Wait-Pull With a Saline Tracer, and (e) the Fluorescein-Based Push-Pull Experiment by *Kang et al.* [2015]

Tracer ID	Tracer Volume (L)	Tracer Salinity (g kg <sup>-1</sup> )	Ethanol (%)	Tracer Density (kg m <sup>-3</sup> )	Chaser Volume (L)	Waiting Time (min)	Pumping Rate (L min <sup>-1</sup> )	Mass Recovery (%)
a	106	40	22.73	1000	113	0	2.9	92 ± 3
b	106	40	22.73	1000	108	120	2.9	78 ± 3
c	100	44	0	1044	90	0	3.3	71 ± 9
d	100	42	0	1042	90	227	3.3	50 ± 10
e	1	0	0	1000	400	0	5	87

## 5. Conclusions

Our results suggest that wethanalt, a mixture of saline water and ethanol, is a suitable tracer for conducting geophysical monitoring using electrical or electromagnetic methods, when density effects are undesirable. Tracer test experiments conducted in push-pull and push-wait-pull configurations, in conjunction with single-hole GPR monitoring, confirm that wethanalt provides a strong GPR signal and does not exhibit the density-driven downward flow observed in our past experiments with dense saline tracers [Shakas *et al.*, 2016]. Therefore, wethanalt may significantly improve our ability to monitor flow and transport processes in situ with hydrogeophysical methods, without the complications of density-driven flow and instabilities. Indeed, our results suggest that if a dense saline tracer is used, it is possible that observations (and inferences) made about the hydrogeological system are unrepresentative of the ambient conditions and may differ significantly if a neutrally buoyant tracer is used instead. We also propose a practical way to prepare a wethanalt mixture with a high electrical conductivity at ambient density for any freshwater hydrogeological application. Additionally, wethanalt is biodegradable, comparatively cheap to produce, and does not pose any health risks. We anticipate that wethanalt or other neutrally buoyant saline tracers will play an important role in advancing hydrogeophysics and in situ monitoring of transport processes. Moreover, since the buoyancy of wethanalt can be adjusted, wethanalt mixtures open a new window on the use of buoyant and nonbuoyant tracers for studying density effects.

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