

# Another look at the colloid particle size-dependent dispersivity

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## Abstract

Laboratory and field studies have demonstrated that dispersion coefficients evaluated by fitting advectiondispersion transport models to nonreactive tracer breakthrough curves do not adequately describe colloid transport under the same flow field conditions. Here an extensive laboratory study was undertaken to assess whether the dispersivity, which traditionally has been considered to be a property of the porous medium, is dependent on colloid particle size and interstitial velocity. A total of 48 colloid transport experiments were performed in columns packed with glass beads under chemically unfavorable colloid attachment conditions. Nine different colloid diameters, and various flow velocities were examined. The breakthrough curves weresuccessfully simulated with a mathematical model describing colloid transport in homogeneous, water saturated porous media. The experimental data set collected in this study demonstrated that the dispersivity is positively correlated with colloid particle size, and increases with increasing velocity. The dispersivity values determined in this laboratory study were compared with 380 dispersivity values from earlier laboratory- and field-scale solute, colloid and biocolloid transport studies published in the literature.

## **Experimental Approach**

A total of 48 flowthrough experiments were conducted in glass The transport of colloids (including biocolloids) in one-dimensional, columns with diameter 2.5 cm and length L=15 or 30 cm, packed homogeneous, water-saturated porous media with first-order with spherical glass beads with diameter  $d_c=2$  mm. Each column attachment (or filtration) and inactivation is governed by (Sim and was packed with glass beads under standing distilled deionized Chrysikopoulos, 1995): water (ddH<sub>2</sub>O) to minimize air entrapment. The columns were placed horizontally to minimize gravity effects (Chrysikopoulos and Syngouna, 2014). Fluorescent polysterene microspheres with diameters d<sub>p</sub>=28, 300, 600, 1000, 1750, 2100, 3000, 5000, and 5500 nm were used as model colloid particles. Microsphere concentrations were measured by a fluorescence spectrophotometer. Straining and wedging are not considered important mechanisms of mass loss in the packed columns examined in this The rate of colloid attachment onto the solid matrix is described by study, because the colloid to collector diameter ratios  $(d_p/d_c)$  were the following first-order equation (Sim and Chrysikopoulos, 1998): well below the suggested threshold of 0.004 (Johnson et al., 2010) or 0.003 (Bradford and Bettahar, 2006) for all cases examined. In the experiments conducted only the two unknown model parameters U and D<sub>1</sub> were estimated by fitting the analytical For a semi-infinite one-dimensional porous medium in the solution. All fittings were conducted with the in-house developed presence of a colloid source in the form of an "instantaneous" nonlinear least squares regression software "ColloidFit," which pulse, the appropriate initial and boundary conditions are: incorporates the state of the art model-independent parameter estimation package "Pest" (Doherty et al., 1994). Furthermore given the flow rate of the pump along with the aquifer porosity, the target velocities were calculated and compared against the fitted velocities, indicating that no systematic error existed.



**Figure 1.** Breakthrough concentration data normalized with respect to M<sub>in</sub>, for the tracer (diamonds) and the colloids with  $d_p=5000$  nm (circles) as a function of time. The experiments were conducted in a 30-cm column. The curves represent fitted model simulations for the tracer (dotted curve) and the colloids (solid curve). Here  $\alpha_1 = 0.14$  cm for the tracer, and  $\alpha_1 = 0.54$  cm for the colloids

(1/mL)

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## **Mathematical Model**

$$\frac{\partial C(t,x)}{\partial t} + \frac{\rho_{b}}{\theta} \frac{\partial C^{*}(t,x)}{\partial t} = D_{L} \frac{\partial^{2} C(t,x)}{\partial x^{2}} - U \frac{\partial C(t,x)}{\partial x}$$
$$-\lambda C(t,x) - \lambda^{*} \frac{\rho_{b}}{\theta} C^{*}(t,x)$$

$$\frac{\rho_{b}}{\theta} \frac{\partial C^{*}(t,x)}{\partial t} = k_{c}C(t,x) - k_{r} \frac{\rho_{b}}{\theta}C^{*}(t,x) - \lambda^{*} \frac{\rho_{b}}{\theta}C^{*}(t,x)$$

$$\frac{\partial C(t,\infty)}{\partial x} = 0, \qquad C(0,x) = 0, \qquad -D_{L} \frac{\partial C(t,0)}{\partial x} + UC(t,0) = M_{\delta} \delta(t)$$



**Figure 2.** Breakthrough concentration data normalized with respect to M<sub>in</sub>, for the colloids with  $d_p = 1000$  nm (squares) and the colloids with  $d_p = 5500$  nm (circles) as a function of time. The experiments were conducted in a 30-cm column. The curves represent fitted model simulations for the colloids with d<sub>p</sub>=1000 nm (dotted curve), and  $d_p=5500$  nm (solid curve). Here,  $\alpha_L=0.36$  cm for the colloids with  $d_p=1000$  nm, and  $\alpha_1=0.57$  cm for the colloids with  $d_p=5500$  nm.







accessible by colloids (reduction of the colloid effective porosity). Under the experimental conditions of this study, the interstitial fluid has continuous access to the entire volume of void-space.

• Colloid dispersivity is not only a function of scale, as conventionally assumed, but also a function of colloid diameter and interstitial velocity. • It was shown that dispersivity increases linearly with increasing colloid particle size. This phenomenon was attributed to:

• Some of the colloids are getting slowed down having to pass through small pore spaces, where other colloids are managing to stay in fast flow paths • Fitted dispersion coefficients based on tracer data should not be used to analyze colloid experimental data.



Figure 7. Comparison between the target (based on Q) and fitted interstitial velocities: (a) U<sub>Target</sub> versus U<sub>Fitted</sub>, and (b) Histogram of U<sub>Fitted</sub>-U<sub>Target</sub>. The solid line represents the ideal one-to-one correlation ( $R^2=1$ ).

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Conclusions

- Size exclusion
- 2. Exclusion from the lower velocity regions
- 3. Possible existence of preferential flow paths

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Figure 6. Compilation of 432 longitudinal dispersivities as a function of length scale. Molecular sized solutes are represented by gray symbols, and colloids/biocolloids by various colored symbols. The solid line is a standard linear regression line...

### References

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