

# LINKING SURFACE 3H TO 3H-3HE GROUNDWATER AGE BY GAS-LIQUID PHASE TRANSPORT MODELING

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The tritium and helium-3 ( $^3\text{H}$ - $^3\text{He}$ ) isotope groundwater age dating method provides a valuable tool for estimating groundwater residence times of 50 years or less. However, interpretation of  $^3\text{H}$ - $^3\text{He}$  data without consideration of unsaturated zone processes leads to difficulties. The negative effects of dispersivity and diffusivity on uncertainty of calculated  $^3\text{H}$ - $^3\text{He}$  groundwater ages are exaggerated without consideration of  $^3\text{H}$  dispersion and  $^3\text{He}$  gas partitioning in the unsaturated zone. Surface concentration of  $^3\text{H}$  inferred from  $^3\text{H}$ - $^3\text{He}$  age and  $^3\text{H}$  groundwater concentration will be underestimated without consideration of vadose zone residence time. To address these difficulties, a coupled gas-liquid phase transport model offers improved linking of surface water  $^3\text{H}$  activity to  $^3\text{H}$ - $^3\text{He}$  age in groundwater.

Importantly, the gas-liquid phase model includes simulation of  $^3\text{H}$  and  $^3\text{He}$  transport in the unsaturated zone, where  $^3\text{He}$  preferentially partitions into the gas phase. By doing so, the gas-liquid phase model confronts a recurring problem in  $^3\text{H}$ - $^3\text{He}$  age interpretation: assuming the  $^3\text{H}$  activity in surface water (generally inferred from IAEA precipitation records) is the same as the initial activity for the  $^3\text{H}$ - $^3\text{He}$  groundwater age dating chronometer. This problem arises because the  $^3\text{H}$  concentration history reaching the water table, the actual location of recharge to groundwater, can be significantly different than the  $^3\text{H}$  concentration history measured in surface water.

Conventional groundwater age interpretation of  $^3\text{H}$ - $^3\text{He}$  data from shallow irrigation wells in the San Joaquin Valley, California suggests  $^3\text{H}$  activities in recharging groundwater at the water table are higher than seen in precipitation for the calculated recharge year. Gas-liquid phase  $^3\text{H}$  and  $^3\text{He}$  modeling results, however, show that accounting for travel time and dispersion from transport through a 5-meter thick vadose zone completely resolves this unrealistic discrepancy and confirms the primary recharge source since agricultural development is irrigation return and not precipitation. In the larger realm of isotopic data interpretation, the gas-liquid phase transport modeling approach enables direct coupling of isotopic transport processes between atmospheric, surface water, vadose zone, and groundwater systems.

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