

Geogenic radon emissions affected by atmospheric pressure - evidence from a breathing well

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Breathing water wells are usually identified by subsurface gases that are exhaled during periods of decreasing atmospheric pressure (P_{ATM}) and inhalation of air during when P_{ATM} is increasing. An 11-day monitoring program in a breathing water well identified in the Calgary (AB, Canada) region showed consistent variations in radon concentrations ($[Rn]$) inside the well casing over four P_{ATM} cycles (with observed changes in P_{ATM} between 0.5 and 1.8 kPa in magnitude; Figure 1). Decreasing P_{ATM} caused high $[Rn]$ gases to enter the well from the subsurface, with peak $[Rn] > 60,000$ Bq/m³. During periods of increasing P_{ATM} , $[Rn]$ decreased to background levels (< 100 Bq/m³) as atmospheric air was 'inhaled' into the well.

Gas and isotopic composition of daily grab gas samples collected from the well casing showed two endmember mixing, with a high $[Rn]$ endmember associated with i) elevated $[CO_2]$ and $[Ar]$, ii) decreased $[O_2]$ and $[CH_4]$, iii) decreased $^{13}C-CO_2$ values, iv) decreased R/R_A values for He, and v) Ne isotope ratios consistent with a deeper geogenic component. The deeper origin of the high $[Rn]$ endmember gases observed during exhalation combined with the short Rn half-life (3.82 days) suggest rapid transport to the ground surface, perhaps as a buoyant free phase gas.

Rapid buoyant transport of high $[Rn]$ gas from the subsurface could be a previously unidentified source of radon in indoor air in non-tectonic regions.

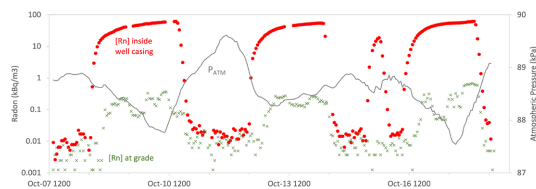


Figure 1: Time series of i) atmospheric pressure (grey line), and radon concentration measured ii) inside the wellhead of the breathing water well (red), and iii) in air sampled at grade about 2.5m from the wellhead.