## Reconstructing final H<sub>2</sub>O contents of hydrated rhyolitic glasses: Insights into H<sub>2</sub>O degassing and eruptive style of silicic submarine volcanoes

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H<sub>2</sub>O degassing during magma ascent plays a fundamental role in determining eruption style and deposit characteristics, hence volcanic hazard. H<sub>2</sub>O contents of pyroclast glasses offer insight into eruptive processes, but secondary hydration (i.e. the addition of H<sub>2</sub>O from the surrounding environment at low temperature in the time following deposition) can alter this record. In order to obtain meaningful final H<sub>2</sub>O data from hydrated glasses it is necessary to distinguish between the original final dissolved H<sub>2</sub>O content and the  $H_2O$  added subsequently during hydration. Since  $H_2O$  added during hydration is added as molecular H2O (H2Om), and the species interconversion between  $H_2O_m$  and hydroxyl (OH) species is negligible at ambient temperature, the final OH content of the glass remains unaltered during hydration. By using H<sub>2</sub>O speciation models to find the original  $H_2O_m$  content that would correspond to the measured OH content of the glass, the original total H<sub>2</sub>O (H<sub>2</sub>O<sub>1</sub>) content of the glass prior to hydration can be reconstructed. These  $H_2O$  speciation data are obtained using FTIR spectroscopy. In many cases OH cannot be measured directly and instead is calculated indirectly as  $OH = H_2O_t - H_2O_m$ . Here we demonstrate the importance of using a speciesdependent H<sub>2</sub>O<sub>t</sub> molar absorptivity coefficient to obtain accurate  $H_2O_t$  and  $H_2O$  speciation data and outline a methodology for calculating such а coefficient for both hydrated and unhydrated rhyolite and andesite glasses. Using this method we present reconstructed final H2Ot contents of hydrated silicic pumice from submarine volcanoes in the Japanese Izu-Bonin Arc and use these data to investigate submarine silicic eruptions and the processes that produce submarine pumice, in order to understand their associated hazards. In particular, we show that pre-hydration final  $H_2O_t$  contents for pumice from Kurose Nishi and Oomurodashi volcanoes were typically <0.3 wt%  $H_2O_t$ , equivalent to quench depths of <50 mbsl, and that pre-hydration final  $H_2O_t$ contents vary with pumice texture.