INTRODUCTION

Rock forming minerals may be crudely classified into two groups: (1) hydrous minerals in which, intrinsic to the mineral structure, hydrogen appears explicitly in the chemical formula, and (2) anhydrous minerals containing no essential hydrogen in their chemical makeup (Aines and Rossman 1984). Numerous previous studies have however revealed the widespread existence of diverse water species, including molecular water (H$_2$O), hydroxyl groups (OH$^-$), and structurally incorporated hydrous complexes (e.g., Si-OH or Al-OH), in minerals considered stoichiometrically anhydrous (Aines and Rossman 1984). Substitutional hydroxyl in nominally anhydrous minerals has been shown to potentially contribute significant quantities of water to the upper mantle (Bell and Rossman 1992; Ingrin and Skogby 2000).

The high-temperature behavior of defect hydrogen species in quartz: Implications for hydrogen isotope studies

KEVIN GRANT,1,* SARAH A. GLEESON,2,† AND STEVE ROBERTS1

1School of Ocean and Earth Science, Southampton Oceanography Centre, University of Southampton, Empress Dock, Southampton SO14 3ZH, U.K.
2School of Earth Sciences, University of Leeds, Leeds LS2 9JT, U.K.

ABSTRACT

A micro-infrared spectroscopic study of hydrothermal vein quartz known to have anomalous $\delta^{18}D$ signatures has identified two hydrogen reservoirs. In samples that generate an isotopic signature in accordance with that anticipated under the accepted model of quartz crystallization, submicroscopic aggregates of liquid water are the dominant hydrous species. Samples which generate an anomalous $\delta^{18}D$ signature contain, in addition to liquid water, structurally incorporated hydrous species associated with impurity cations.

Infrared spectra obtained during in situ stepped heating experiments, coupled with infrared analysis at 25 $^\circ$C, demonstrate that hydrogen liberated between 300 and 500 $^\circ$C is chiefly molecular, liquid water. Hydrogen liberated at temperatures greater than 500 $^\circ$C is dominantly that associated with specific structurally incorporated cation defects. Since both defect hydrogen and molecular water are contemporaneously incorporated from the precipitating medium during crystallization, we propose that irregular $\delta^{18}D$ signatures, released following decrepitation at temperatures greater than 500 $^\circ$C, are due to isotopically fractionated hydrogen released from interstitial OH defect sites in the quartz structure.

$\delta^{18}D$ signatures obtained from stoichiometrically anhydrous minerals are generally interpreted under the assumption that the hydrogen measured comes uniquely from decrepitated fluid inclusions. Instead, we suggest that $\delta^{18}D$ ratios obtained from hydrothermal quartz reflect a sum of the contributions made by individual hydrogen reservoirs, each with a potentially distinctive $\delta^{18}D$ signature. Thus, if the overall $\delta^{18}D$ signature is attributed entirely to fluid inclusion phases, the nature of the precipitating fluid may be misinterpreted. Hydrogen extracted as molecular water at between 300 and 500 $^\circ$C provides a true reflection of the hydrothermal solution associated with crystal precipitation.

Stable isotope ratios of O atoms ($\delta^{18}O$) and hydrogen ($\delta^{18}D$) are extremely important geochemical tracers used to determine