ORIGINAL PAPER

Crystal scale anatomy of a dying supervolcano: an isotope and geochronology study of individual phenocrysts from voluminous rhyolites of the Yellowstone caldera

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Received: 12 July 2011/Accepted: 31 January 2012 © Springer-Verlag 2012

Abstract A voluminous (>600 km³) and long-lived (\sim 520–75 ka) phase of rhyolitic eruptions followed collapse of the Yellowstone caldera 640 ka. Whether these eruptions represent a dying cycle, or the growth of a new magma chamber, remains an important question. We use new U-Th zircon ages and δ^{18} O values determined by ion microprobe, and sanidine Pb isotope ratios determined by laser ablation, to investigate the genesis of voluminous post-caldera rhyolites. The oldest post-caldera rhyolites, erupted between \sim 520 and 470 ka, exhibit extreme age and oxygen isotopic heterogeneity, requiring derivation from individual parcels of low- δ^{18} O melts. We find a progressive increase in zircon homogeneity for rhyolite eruptions from ~260 to 75 ka, with homogeneous low- δ^{18} O zircon values of 2.7–2.8‰ that are in equilibrium with low- δ^{18} O host melts for the majority of the youngest eruptions. New sanidine Pb isotope data define separate arrays for post-caldera rhyolites and the calderaforming tuffs that preceded them, indicating that they were not sourced from a mushy Lava Creek Tuff batholith that remained after caldera collapse. Rather, our new age and isotopic data indicate that the post-caldera rhyolites were generated by remelting of a variety of intracaldera source

Communicated by J. Hoefs.

Published online: 04 March 2012

Electronic supplementary material The online version of this article (doi:10.1007/s00410-012-0724-x) contains supplementary material, which is available to authorized users.

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A. K. Schmitt Department of Earth and Space Sciences, University of California, Los Angeles, CA 90095, USA rocks, consisting of pre-Lava Creek Tuff volcanic and plutonic rocks and earlier erupted post-Lava Creek Tuff rhyolites. Batch assembly of low- δ^{18} O melts starting at ~ 260 ka resulted in progressive homogenization, followed by differentiation and cooling up until the last rhyolite eruption ~ 75 ka, a trend that we interpret to be characteristic of a dying magma reservoir beneath the Yellowstone caldera.

Keywords Yellowstone · Caldera · Zircon · Rhyolite · Geochronology · Oxygen isotopes · Magma evolution

Introduction

Large caldera volcanoes, or "supervolcanoes," have produced some of the Earth's largest known eruptions of silicic magma. The processes and timescales by which large volumes of silicic magma are generated in the shallow crust, and the petrogenetic paths along which they evolve, whether toward explosive caldera-forming eruptions, effusive lava flows, or stalled plutons and batholiths, are important for hazard and resource assessments for caldera centers worldwide. The advent of microanalytical techniques that enable dating and isotopic measurements of individual crystals has greatly improved our understanding of magmatic systems (see Putirka and Tepley 2008 and references therein) and proven critical to deciphering the complex magmatic histories of caldera volcanoes, where multiple episodes of intrusion, assembly and melt extraction are retained in the crystal record (e.g., Vazquez and Reid 2004; Bacon and Lowenstern 2005; Charlier et al. 2005; Folkes et al. 2011).

One prime example of a caldera complex that has produced large volumes of silicic magma with diverse age and isotopic signatures is the Yellowstone Plateau volcanic field



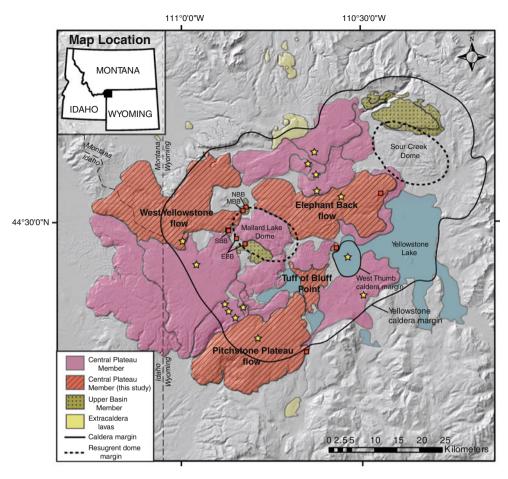


Fig. 1 Map showing the distribution of post-Lava Creek Tuff rhyolites in the Yellowstone caldera. The geologic map of Christiansen (2001) was used to define unit boundaries. The Central Plateau Member units analyzed in this study are indicated by the *bold labels*. *Squares mark* our sampling localities. The locations of the Biscuit Basin rhyolite flows, including North Biscuit Basin (NBB), Middle Biscuit Basin (MBB), South Biscuit Basin (SBB), and East Biscuit Basin (EBB), are also shown. NBB and SBB are Central Plateau

Member units, whereas MBB and EBB are Upper Basin Member units (see text for more detail). Stars mark the inferred vent sources for individual Central Plateau Member rhyolite eruptions (Christiansen 2001). The Yellowstone caldera margin, resurgent dome margins, and West Thumb caldera margin are taken from Christiansen (2001) and Christiansen et al. (2007). The *inset* shows the map location in the western USA

in western Wyoming (Fig. 1). In this study, we investigate magma genesis during the last caldera-forming cycle at Yellowstone by combining several crystal scale analytical approaches, focusing on dating and isotopic analyses of refractory zircon phenocrysts. Our new results enable us to evaluate whether Yellowstone's youngest episode of rhyolitic volcanism represents a rejuvenating or waning stage of magmatism, using single crystals as probes of magma evolution from the oldest to youngest rhyolite eruptions.

Background

Caldera cycles at Yellowstone

The Yellowstone Plateau volcanic field is the youngest caldera center in the 16-Ma track of the Yellowstone

hotspot (Pierce and Morgan 1992). It offers one of the best preserved records of caldera complex evolution above a continental hotspot, with extensive exposures of explosive and effusive deposits from three climactic caldera-forming cycles. These include the 2.06-Ma Huckleberry Ridge Tuff (HRT), 1.29-Ma Mesa Falls Tuff (MFT), and 0.64-Ma Lava Creek Tuff (LCT) (Lanphere et al. 2002). Estimated eruptive volumes for these tuffs are among the largest known for caldera volcanoes, at 2,450, 280, and 1,000 km³, respectively (Christiansen 2001). Ash fall deposits from the HRT and LCT eruptions once covered much of the North American continent, and the corresponding ash flow tuffs reached thicknesses of up to 500 m near the eruptive vents (Christiansen 2001).

Large caldera-forming eruptions at Yellowstone are bracketed by smaller, predominantly effusive eruptions of pre- and post-caldera lavas. The third LCT caldera cycle



resulted in the formation of the $\sim 70 \times 50$ km Yellow-stone caldera, which was subsequently filled by more than 600 km³ of post-caldera rhyolites that buried most of the caldera floor and the western caldera rim (Fig. 1). Many of these individual lava flows are quite large (>10 km³) and form well-defined lobate extrusions that cover broad geographic areas of the Yellowstone Plateau volcanic field (Fig. 1). These rhyolites represent either residual magma from the third LCT cycle, or perhaps the beginning of a fourth eruptive cycle, owing to their very large cumulative eruptive volume (Hildreth et al. 1984; Christiansen 2001; Christiansen et al. 2007).

Post-caldera rhyolites of the Yellowstone caldera

Post-caldera rhyolites of the LCT cycle are subdivided into two principal members based on location, age, and composition. The earliest erupted ($\sim 520-470 \text{ ka}$) rhyolites compose the Upper Basin Member (UBM) flows $(\sim 50 \text{ km}^3 \text{ total volume})$ located near the Mallard Lake and Sour Creek resurgent domes of the Yellowstone caldera (Fig. 1). UBM flows have more primitive compositions than younger post-LCT rhyolites (Christiansen et al. 2007; Girard and Stix 2009), and they have the lowest magmatic δ^{18} O signatures of all studied Yellowstone rhyolites (Hildreth et al. 1984; Bindeman and Valley 2001; Bindeman et al. 2008). Oxygen isotope studies of individual UBM units reveal distinct low- δ^{18} O obsidian values that range from ~ 0 to 2‰, significant phenocrystic δ^{18} O heterogeneity spanning an $\sim 8\%$ range, and zircons with up to 2.1 Ma older cores and inherited (pre-caldera) normal- δ^{18} O values (Bindeman and Valley 2001; Bindeman et al. 2008). This evidence is taken by the authors to indicate that UBM rhyolite lavas were derived from independent magma batches produced by bulk remelting of hydrothermally altered pre-caldera protoliths.

The youngest ($\sim 260-75$ ka) and most voluminous post-LCT eruptions compose the Central Plateau Member (CPM) units, which consist of at least 17 lava flows and two pyroclastic tuffs (Christiansen et al. 2007). Individual eruptive volumes of CPM rhyolites are typically on the order of $\sim 10 \text{ km}^3$, but several approach $\sim 50-70 \text{ km}^3$ (Christiansen et al. 2007). The 50-km³ Tuff of Bluff Point eruption was of sufficient volume and explosivity to generate a smaller caldera within the larger Yellowstone caldera, a ~ 10 -km-wide elliptical depression filled by the western portion of the Yellowstone Lake, known as the West Thumb caldera (Fig. 1). CPM rhyolites were erupted in discrete batches from vents along two northwest-striking lineaments that bisect the Yellowstone caldera (Fig. 1). However, the rhyolites display no apparent regional differences in composition for different vent localities (Girard and Stix 2010). Recent studies of several flows previously recognized as UBM units, including the Scaup Lake, South Biscuit Basin, and North Biscuit Basin flows, demonstrate that they have δ^{18} O ranges similar to CPM rhyolites and Ar–Ar eruption ages that are ~ 300 ky younger than UBM rhyolites (Christiansen et al. 2007; Bindeman et al. 2008), suggesting that they belong to the CPM rather than UBM group. Here, we include them with the "early" CPM group and suggest that they are important for understanding the more voluminous "later" CPM rhyolites.

Like UBM rhyolites, early CPM units exhibit heterogeneity in zircon age and δ^{18} O values, but have less extreme low- δ^{18} O obsidian values of $\sim 3-4\%$ (Bindeman and Valley 2001). The more voluminous later CPM rhyolites have low- δ^{18} O melt values of $\sim 3-4\%$ that match those of the early CPM units, but they possess less phenocrystic δ^{18} O heterogeneity than early CPM units (Bindeman and Valley 2001; Bindeman et al. 2008). U-Pb and U-Th zircon ages of later CPM rhyolites closely approximate their respective Ar-Ar eruption ages, and individual units reveal no or subtle zircon inheritance from pre-caldera source rocks (Vazquez and Reid 2002), with the possible exception of the voluminous Solfatara Plateau flow, which exhibits δ^{18} O zircon disequilibria (Bindeman et al. 2008) and possesses a Pb isotopic composition similar to that of HRT rocks (Vazquez et al. 2009).

Oxygen isotope insights into Yellowstone rhyolite petrogenesis

The high-latitude, high-altitude, intracontinental position of the Yellowstone caldera makes it possible to resolve meteoric water signatures in the oxygen isotopic compositions of erupted rhyolites. While it has long been recognized that large volumes of rhyolite produced by the Yellowstone caldera have drastic depletions in ¹⁸O that require significant involvement of meteoric water (Friedman et al. 1974; Hildreth et al. 1984; Taylor 1986; Hildreth et al. 1991), the processes by which such depletions occurred were unclear until age and oxygen isotopic investigation of Yellowstone rhyolites on the single crystal scale (Bindeman and Valley 2000, 2001; Bindeman et al. 2001). Earlier interpretations included direct influx of meteoric waters into silicic magmas (Hildreth et al. 1984) or assimilation of wall rocks altered by hydrothermal fluids (Hildreth et al. 1991). As addressed by Bindeman and Valley (2000, 2001), each of these scenarios is unrealistic based on heat and mass-balance constraints. For instance, even in an extreme case of assimilation, in which the host magma and bulk assimilant are mixed in equal parts, the bulk assimilant would have to possess an exceptionally low- δ^{18} O value (<-6‰) to produce the observed UBM melt δ^{18} O values of ~0–2‰. However, such ultra low- δ^{18} O rocks are rare at Yellowstone caldera. Isotopic mixing



models yield considerably higher estimates of $\sim -1\%$ for the average $\delta^{18}O$ composition of low- $\delta^{18}O$ Yellowstone source rocks (Watts et al. 2010).

Individual zircon crystals from early UBM rhyolites of the Yellowstone caldera contain unequivocal age and oxygen isotopic evidence for inheritance from specific volcanic protoliths, supporting a magma genesis model of wholesale remelting of hydrothermally altered intracaldera blocks (Bindeman et al. 2008). Unique low- δ^{18} O signatures indicate that UBM rhyolites were generated by spatially discrete episodes of remelting, which yielded independent parcels of variably ¹⁸O-depleted melts (Bindeman et al. 2008). This study builds on these and other previous results by presenting new age and O isotope data obtained by ion microprobe for individual zircons and Pb isotope data obtained by laser ablation for individual sanidines, for the youngest erupted post-caldera rhyolites of the Yellowstone caldera, the CPM rhyolites. In addition, we report new isotopic and age data for the oldest known UBM rhyolite. the East Biscuit Basin flow. With the addition of these important new datasets, we are able to construct an integrated model of post-LCT rhyolite genesis.

Samples and methods

We report new data for four young large-volume CPM rhyolites, including three lava flows (Pitchstone Plateau, West Yellowstone, Elephant Back) and one explosive tuff (Tuff of Bluff Point), two older, small-volume CPM rhyolite lavas (North Biscuit Basin, South Biscuit Basin), and one newly defined unit that represents the earliest and most primitive UBM rhyolite, the East Biscuit Basin flow (Fig. 1). These units span the full eruptive age range of post-LCT rhyolites and have a broad spatial distribution within the Yellowstone caldera (Fig. 1). With the exception of the East Biscuit Basin flow, which is one of the few low-silica rhyolites at Yellowstone, all samples are highsilica rhyolites with $\sim 5-20$ vol% sanidine, quartz, clinopyroxene, orthopyroxene, Fe-Ti oxides and accessory phases (see Supplementary Material 1 for petrographic descriptions, Supplementary Material 2 for photomicrographs, and Supplementary Material 3 for whole-rock major and trace element geochemistry). Notably, only North Biscuit Basin, South Biscuit Basin, and East Biscuit Basin flows have plagioclase, and East Biscuit Basin lacks sanidine and quartz. Abundant glass reentrants, sieved plagioclase phenocrysts, and anhedral crystal clusters were observed in samples of South Biscuit Basin and East Biscuit Basin lavas, consistent with the descriptions of Girard and Stix (2009). Zircon phenocrysts commonly occur as isolated crystals in the groundmass glass or as inclusions within Fe-Ti oxide and pyroxene aggregates. Most zircon

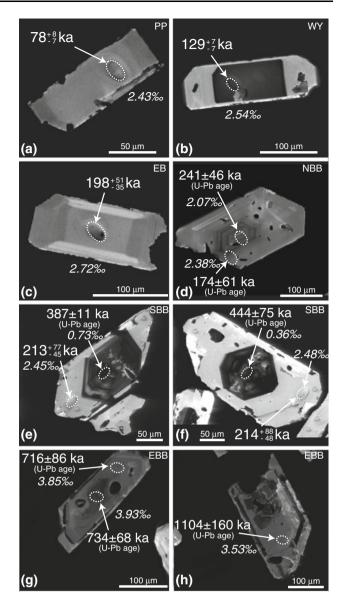


Fig. 2 Cathodoluminescence (CL) images of zircons dated for U-Th and U-Pb ages and analyzed for oxygen isotope compositions by ion microprobe. Analyzed spots are bounded by dashed ellipses. U-Th ages and uncertainties are shown in bold text; U-Pb ages are indicated by parentheses. Oxygen isotope compositions of dated zircon spots are shown in italic text. See Table 1 for a compilation of all ion microprobe data. a Pitchstone Plateau (PP) zircon, euhedral, and unzoned in CL. b West Yellowstone (WY) zircon, euhedral with distinct core-rim CL zoning. c Elephant Back (EB) zircon, euhedral with subtle core-rim CL zoning. d North Biscuit Basin (NBB) zircon exhibits core-rim zoning in CL, U-Pb age, and O isotope composition. e, f South Biscuit Basin (SBB) zircons, pronounced core-rim zoning in CL, U-Pb and U-Th age, and O isotope composition, cores are CL dark with subtle sector zoning. g East Biscuit Basin (EBB) zircon, zoned in CL and U-Pb age. h East Biscuit Basin (EBB) zircon has a U-Pb age that significantly predates the Lava Creek Tuff and post-Lava Creek Tuff eruptions

crystals are subhedral–euhedral, ~ 100 –200 μm in length, and many exhibit cathodoluminescence (CL) zoning (Fig. 2).



U–Th disequilibrium dating of zircons was performed with a CAMECA ims 1270 ion microprobe at UCLA using a $\sim\!30\times35~\mu m$ spot size, according to the protocols described by Schmitt et al. (2006). Analytical details and operating conditions can be found in Supplementary Material 4. A small percentage (<10%) of the analyzed CPM zircon grains were found to be in $^{238}U^{-230}$ Th secular equilibrium and were reanalyzed for $^{238}U^{-206}$ Pb concordia ages according to the techniques outlined in Schmitt et al. (2003). Two Biscuit Basin samples (East Biscuit Basin and North Biscuit Basin) were dated for U–Pb ages at Stanford University using a SHRIMP-RG ion microprobe. For these analyses, the analytical techniques and conditions of Barth and Wooden (2010) were used.

All zircons dated for U–Th and U–Pb ages were analyzed for δ^{18} O compositions in subsequent analytical sessions at the UCLA SIMS facility. Back-scattered electron and CL images of zircon grain mounts enabled us to target the dated ion probe spots. After repolishing our zircon mounts according to the protocols of Kita et al. (2009), we determined the δ^{18} O compositions of the dated zircon spots a few microns below the original ion probe pits. Corrections for instrumental mass fractionation (IMF) were made using zircon standard KIM-5 (Valley 2003), mounted in the center of each zircon grain mount. KIM-5 zircons were analyzed repeatedly during each analytical session, with external errors that were typically less than 0.22% (1 SD) for each sample block (15–20 analyzed spots per block).

Pb isotope analyses of sanidine phenocrysts and host groundmass glasses were performed with a Finnigan Neptune MC-ICP-MS laser ablation system at Washington State University using the techniques and operating conditions of Kent (2008), as described in Supplementary Material 4. We analyzed four post-LCT units, including an older CPM rhyolite (South Biscuit Basin) and three younger CPM rhyolites (Pitchstone Plateau, Elephant Back, and the Tuff of Bluff Point), all of which were dated for U-Th and U-Pb ages and analyzed for δ^{18} O compositions by us. In addition, we measured Pb isotope ratios for sanidine separates from several members of major calderaforming tuff eruptions at Yellowstone (HRT [members B and C] and LCT [member A]), which were previously studied for U-Pb ages and δ^{18} O compositions (Bindeman and Valley 2001; Bindeman et al. 2001, 2008).

Results

Results presented below include zircon crystallization ages and $\delta^{18}O$ compositions, and sanidine Pb isotope compositions. We also include whole-rock thermometry for all studied rhyolites and synthesize these new data with thermometry results from previous studies. The results in each

subsection are organized according to eruption age (oldest to youngest).

Zircon crystallization ages and $\delta^{18}O$ compositions

Earliest Upper Basin Member rhyolite

Zircon dating of the most primitive rhyolite to erupt within the Yellowstone caldera, the East Biscuit Basin flow, indicates that it is also the oldest, with an error-weighted mean U-Pb age (781 \pm 76 ka. n = 9, mean square of weighted deviates (MSWD) = 1.0). This age falls between the Ar-Ar eruption ages of the MFT and LCT (Fig. 3). No sanidine is present in the East Biscuit Basin flow, and our attempts to perform 40Ar/39Ar dating on plagioclase were unsuccessful, so we are unable to report an Ar-Ar eruption age for this unit. However, eruption of the East Biscuit Basin flow within the Yellowstone caldera near the Mallard Lake Resurgent dome (Fig. 1) indicates that it is a post-LCT rather than pre-LCT flow. The U-Pb zircon age range published by Bindeman et al. (2001) for the Middle Biscuit Basin flow overlaps the broad age range that we find for the East Biscuit Basin flow, and isotopically, the two units are similar (this study, Pritchard and Larson in review), suggesting that East Biscuit Basin and Middle Biscuit Basin flows may be related UBM units. Some East Biscuit Basin zircons are zoned in CL and U-Pb age (Fig. 2g), and the oldest zircons are generally present as fragments rather than whole grains (Table 1).

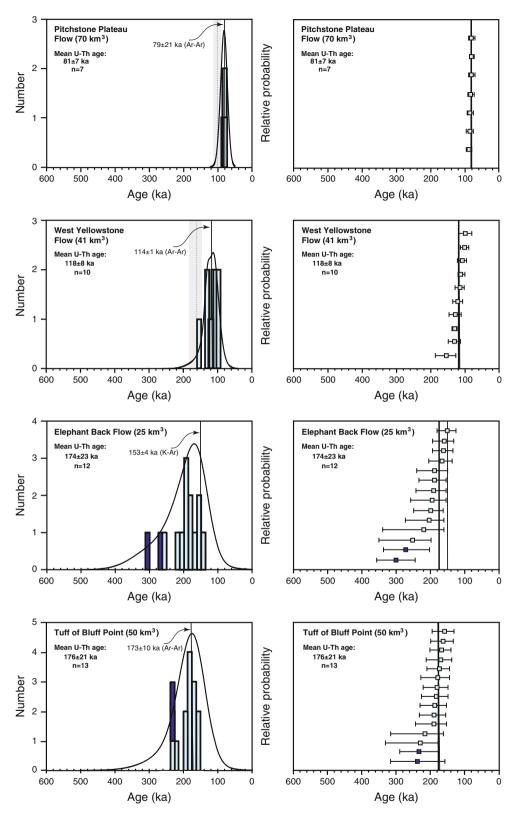
Complementary oxygen isotope data for dated East Biscuit Basin zircon spots reveal that it has zircon δ^{18} O values that are for the most part higher than LCT zircons, in contrast to other analyzed post-LCT units, which typically have lower zircon δ^{18} O values (c.f., Bindeman et al. 2008). However, East Biscuit Basin does have host glass and plagioclase with low- δ^{18} O values that are distinctive of post-LCT UBM units, 1.0 and 1.4%, respectively. The average zircon δ^{18} O value of 4.3 \pm 0.3% is more than 5% higher than the calculated equilibrium zircon value for East Biscuit Basin glass (Fig. 4) and is therefore one of the most extreme cases of oxygen isotopic disequilibrium documented for Yellowstone rhyolites. Unlike other UBM units, there is no clear core-rim δ^{18} O zoning in East Biscuit Basin zircons, and the δ^{18} O range of zircon cores and rims is relatively narrow ($\sim 2\%$) compared to the exceptionally large ($\sim 6-8\%$) range found for zircons from the Middle Biscuit Basin flow, Dunraven Road flow, and Canyon flow (Bindeman et al. 2008).

Early Central Plateau Member rhyolites

Zircon age data for the North Biscuit Basin and South Biscuit Basin flows corroborate earlier interpretations that



Fig. 3 U-Th and U-Pb ages of individual Yellowstone zircons determined by ion microprobe (see Table 1 for data). Left panels show error-weighted probability distribution curves for zircon analyses. U-Th zircon ages are shown by the light bars; zircons in or close to U-Th secular equilibrium were dated for U-Pb ages, shown by the dark bars. Ar-Ar eruption ages are shown by thin vertical lines (Gansecki et al. 1996; Lanphere et al. 2002; Christiansen et al. 2007, this study). The K-Ar eruption age for the Elephant Back flow is from Obradovich (1992). For the Pitchstone Plateau flow and West Yellowstone flow, we include the mean U-Th ages (thin dashed lines) and error fields (shaded) published by Vazquez and Reid (2002) for these units. Right panels show individual zircon analyses (light squares for U-Th ages and dark squares for U-Pb ages) are shown with their error bars $(\pm 1\sigma)$. Bold vertical lines show the errorweighted mean U-Th and U-Pb zircon ages. Thin vertical lines show the eruption ages (as indicated and labeled in the left

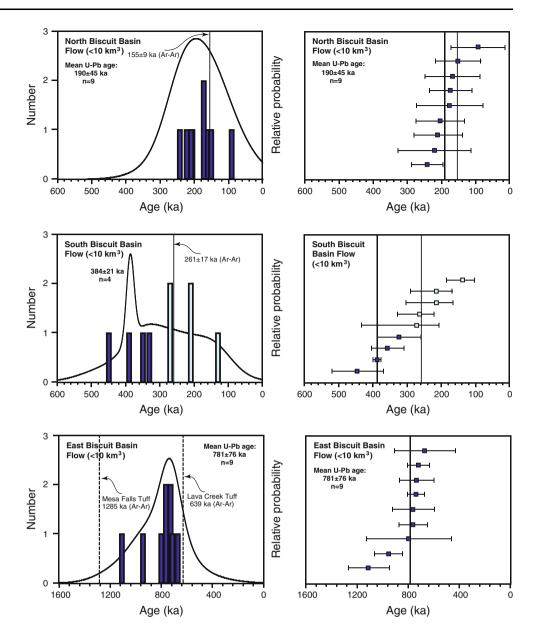


they are distinct units rather than a single Biscuit Basin rhyolite (Bindeman et al. 2008). The North Biscuit Basin flow has an error-weighted mean U-Pb age (190 \pm 45 ka, n=9, MSWD = 0.4) that is unimodal and normally

distributed (Fig. 3), consistent with our findings for CPM rhyolites (see next section). The 40 Ar/ 39 Ar sanidine eruption age determined by us for North Biscuit Basin (155 \pm 9 ka) is within the 170–75 ka age range of CPM



Fig. 3 continued



rhyolite eruptions (Christiansen et al. 2007), indicating that it is an early CPM unit. Core-rim zoning of North Biscuit Basin zircons is apparent in CL and U-Pb age (Fig. 2d), with the oldest core ages closely approximating the Ar-Ar eruption age of the oldest CPM unit, the Scaup Lake flow (257 \pm 13 ka; Christiansen et al. 2007).

South Biscuit Basin flow has a non-normally distributed zircon age population that spans a broad age range (Fig. 3). The error-weighted mean U-Pb age of the oldest South Biscuit Basin age peak (384 \pm 21 ka, n=4, MSWD = 0.7) significantly predates the onset of CPM eruptions (\sim 260 ka). South Biscuit Basin zircons exhibit distinct core-rim zoning in both CL and age (Fig. 2e, f). U-Th ages of zircon rims typically overlap the 40 Ar/ 39 Ar sanidine eruption age within error (261 \pm 17 ka; Fig. 3). This evidence suggests that the South Biscuit Basin flow contains

antecrystic zircon crystals, with cores that skew the apparent U-Pb age to older values, and overgrowth rims that reflect younger zircon growth prior to eruption.

Like the zircon age data, oxygen isotope data for dated zircon spots require that the North Biscuit Basin and South Biscuit Basin flows be treated as separate Biscuit Basin units (Fig. 4). North Biscuit Basin has an average zircon δ^{18} O value of $2.7 \pm 0.2\%$, which is identical to the average zircon δ^{18} O values determined for the CPM Tuff of Bluff Point and Elephant Back flow (see next section). North Biscuit Basin zircon rims fall within the same δ^{18} O range as zircon cores, and there is no discernable core-rim zoning; however, the calculated equilibrium zircon δ^{18} O value with host glass is about 0.5% lower than the average zircon value, indicating subtle oxygen isotopic disequilibrium. South Biscuit Basin exhibits more extreme δ^{18} O



Table 1 U-Th and U-Pb ages, δ^{18} O values, and sizes of zircons in studied Yellowstone rhyolites

Sample	Core- rim	Model U-Th age (ka) ^a	$+1\sigma$	-1σ	U-Pb concordia age (ka)	±1σ	δ^{18} O (‰) (same spot)	δ^{18} O (‰) (adjacent)	$\pm 1\sigma$ ext	Length (μm)	Width (µm)
Pitchstone Plate	au flow (YL02-1)									
YL02-1_g1	C	83	9	-9	_	_	3.16	3.02	0.22	133	77
YL02-1_g2	C	82	10	-9	_	_	3.01	3.23	0.22	117	51
YL02-1_g3	C	78	8	-7	_	_	2.43	_	0.22	144	54
YL02-1_g5	C	84	11	-10	_	_	2.61	3.01	0.22	218	60
YL02-1_g6	C	87	9	-8	_	_	2.44	2.61	0.22	108	56
YL02-1_g7	C	78	10	_9	_	_	2.85	_	0.22	82	49
YL02-1_g8	C	79	9	_9	_	_	2.14	2.90	0.22	104	61
West Yellowsto	ne flow (YL96-1)									
YL96-1_g1	C	129	7	- 7	_	_	2.54	3.08	0.22	231	73
YL96-1_g2	C	126	19	-16	_	_	2.92	2.85	0.22	175	104
YL96-1_g3	C	153	34	-26	_	_	2.94	2.67	0.22	243	110
YL96-1_g4	C	98	21	-17	_	_	2.38	2.41	0.22	188	103
YL96-1_g5	C	120	15	-13	_	_	3.08	_	0.22	154	66
YL96-1_g6	C	110	10	- 9	_	_	2.95	_	0.22	204	83
YL96-1_g7	C	100	13	-12	_	_	2.69	3.18	0.22	203	90
YL96-1_g8	C	114	15	-13	_	_	2.58	2.72	0.22	189	75
YL96-1_g9	C	129	20	-17	_	_	3.11	3.04	0.22	169	79
YL96-1_g10	C	107	14	-13	_	_	2.90	_	0.22	178	72
Elephant Back f											
YS-4_g1	C	345	∞	-124	271	67	3.26	_	0.22	219	104
YS-4_g2	C	187	54	-36	_	_	3.57	_	0.22	216	84
YS-4_g3	C	189	54	-36	_	_	2.87	2.51	0.22	197	77
YS-4_g4	C	198	51	-35	_	_	2.72	3.21	0.22	179	79
YS-4_g5	C	149	32	-24	_	_	2.64	-	0.22	220	86
YS-4_g6	C	250	101	-52	300	56	2.83	2.52	0.22	245	94
YS-4_g8	C	194	65	-40	_	-	2.48	_	0.22	207	97
YS-4_g9	C	165	40	-29	_	_	2.40	3.21	0.22	160	74
YS-4_g10	C	187	48	-33	_	_	2.42	-	0.22	186	71
YS-4_g11	C	203	71	-43	_	_	3.07	_	0.22	248	82
YS-4_g12	C	218	123	-56	_	_	2.34	_	0.22	153	89
YS-4_g13	C	160	36	-27	_	_	2.59	2.51	0.22	131	60
YS-4_g14	C	160	34	-26	_	_	2.15	_	0.22	304	114
Tuff of Bluff Po			51	20			2.13		0.22	501	111
YS-2_g1	C C	368	∞	-132	237	79	2.36	2.60	0.22	371	142
YS-2_g1 YS-2_g2	C	189	54	-36	_	-	2.74	2.86	0.22	256	91
YS-2_g2 YS-2_g3	C	170	34 44	-30	_	_	2.74	2.84	0.22	300	112
YS-2_g3 YS-2_g4	C	159	37	-31 -28	_	_	2.67	2.70	0.22	338	131
YS-2_g5	C	179	50	-26 -34	_	_	2.84	2.49	0.22	397	119
YS-2_g5 YS-2_g6	C	188	44	-34 -31	_	_	2.94	2.52	0.22	263	107
YS-2_g0 YS-2_g7	C	173	39	-31 -28	_	_	2.94	2.74	0.22	182	107
YS-2_g8	C	186	45	-26 -32	_	_	2.65	2.74	0.22	277	120
YS-2_g8 YS-2_g9	C	167	35	-32 -26			2.85	2.29	0.22	277	90
_					_	_					
YS-2_g10	C	162	37	-28	- 222	- 57	2.25	2.30	0.22	245	106
YS-2_g11	C	229	103	-52	232	57	2.98	2.72	0.22	276	111
YS-2_g12	C	180	42	-30	_	_	2.56	2.44	0.22	247	118
YS-2_g13	С	215	102	-52	_	_	2.60	3.09	0.22	203	102
YS-2_g14	C	182	45	-32	_	_	2.39	2.73	0.22	237	140



Table 1 continued

Sample	Core- rim	Model U-Th age (ka) ^a	$+1\sigma$	-1σ	U-Pb concordia age (ka)	$\pm 1\sigma$	δ^{18} O (‰) (same spot)	δ^{18} O (‰) (adjacent)	$\pm 1\sigma$ ext	Length (μm)	Width (µm)
North Biscuit Ba	sin flow	(08YS-14)									
KWBBRR-2.1	C	_	_	_	92	79	3.22	3.22	0.35	357	214
KWBBR-4.1	C	_	_	_	151	66	2.71	6.74	0.35	Fragment	Fragment
KWBBR-3.1	C	_	_	_	168	80	2.61	2.71	0.35	Fragment	Fragment
KWBBR-8.1	R	_	_	_	175	97	2.82	2.52	0.35	276	149
KWBBRR-9.1	R	_	_	_	174	61	2.38	_	0.35	_	_
KWBBR-7.1	C	_	_	_	204	71	2.70	2.74	0.35	229	85
KWBBR-2.1	C	_	_	_	210	72	2.90	_	0.35	328	162
KWBBR-5.1	R	_	_	_	221	106	3.29	2.30	0.35	Fragment	Fragment
KWBBRC-9.2	C	_	_	_	241	46	2.07	_	0.35	288	114
South Biscuit Ba	sin flow	(YL96-2)									
YL2_58-3	R	213	77	-45	_	_	2.45	_	0.20	_	_
YL2_52-1	C	137	50	-34	_	_	-0.20	_	0.20	300	105
YL2_5-2	R	262	65	-40	_	_	2.13	_	0.24	_	_
YL2_52-3	R	270	164	-63	_	_	2.39	_	0.20	_	_
YL2_59-2	R	214	88	-48	_	_	2.48	_	0.24	_	_
YL2_6-2	R	356	∞	-134	322	64	2.37	_	0.24	_	_
YL2_58-1	C	321	306	-72	387	11	0.73	_	0.20	260	110
YL2_59-1	C	449	∞	-123	444	75	0.36	_	0.24	300	140
YL2_6-1	C	326	∞	-103	357	46	1.55	_	0.24	215	70
East Biscuit Basi	n flow (0	08YS-15b)									
KWEBB-2.1	C	_	-	_	667	236	5.13	4.90	0.35	135	144
KWEBB-9.1	R	_	-	_	716	86	3.85	4.07	0.35	_	-
KWEBB-8.1	C	_	-	_	734	68	3.93	_	0.35	257	89
KWEBB-6.1	C	_	_	-	733	137	4.05	-	0.35	204	86
KWEBB-4.1	R	_	_	-	757	166	3.90	4.41	0.35	220	129
KWEBB-7.1	C	_	_	-	761	110	4.84	4.99	0.35	Fragment	Fragment
KWEBB-1.1	C	_	_	-	793	334	3.98	-	0.35	Fragment	Fragment
KWEBB-3.1	C	_	_	-	949	111	4.94	-	0.35	Fragment	Fragment
KWEBB-5.1	C	_	_	_	1,104	160	3.53	3.63	0.35	Fragment	Fragment

 $[^]a$ Model ages were calculated using zircon–vitrophyre isochrons. Published thermal ionization mass spectrometry (TIMS) data for U and Th isotopes of CPM vitrophyres (Vazquez and Reid 2002) were used to estimate the initial 230 Th abundances of CPM zircons; average activity ratios of (230 Th)/(232 Th) = 0.78 \pm 0.05 and (238 U)/(232 Th) = 0.72 \pm 0.05 were used for CPM units without published TIMS data. Decay constants: λ_{230} : 9.158 \times 10 $^{-6}$ year $^{-1}$; λ_{232} : 4.9475 \times 10 $^{-11}$ year $^{-1}$; λ_{238} : 1.55125 \times 10 $^{-10}$ year $^{-1}$

disequilibrium, with zircons that are up to 2% lower than the equilibrium value, though zircon rims more closely approximate an equilibrium value (Fig. 4). The calculated equilibrium zircon δ^{18} O value for South Biscuit Basin is about 1% lower than equilibrium values for other CPM units, but clearly more similar in oxygen isotopic composition to CPM than to UBM rhyolites.

Later Central Plateau Member rhyolites

Zircon ages for studied CPM rhyolite samples are unimodal and do not exhibit xenocrystic or antecrystic age inheritance (Fig. 3), in contrast to the significant zircon age heterogeneity documented for UBM samples (Bindeman et al. 2001). Error-weighted mean U-Th ages for the Pitchstone Plateau flow (81 \pm 7 ka, n=7, MSWD = 0.2), West Yellowstone flow (118 \pm 8 ka, n=10, MSWD = 1.1), and Tuff of Bluff Point (176 \pm 21 ka, n=13, MSWD = 0.1) are identical to 40 Ar/ 39 Ar sanidine eruption ages published by Christiansen et al. (2007) for these units (Fig. 3). Mean U-Th zircon ages previously reported by Vazquez and Reid (2002) for the Pitchstone Plateau and West Yellowstone flows (dashed lines and shaded error fields in Fig. 3) are slightly older than our mean U-Th age estimates, owing to the fact that these authors found and included some antecrysts in their age averages,



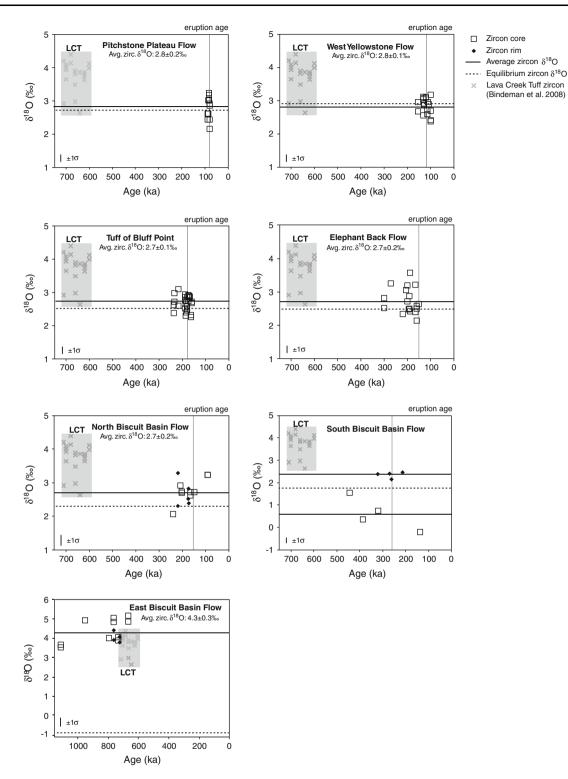


Fig. 4 Oxygen isotope compositions of individual Yellowstone zircons determined by ion microprobe (see Table 1 for data). Oxygen isotope compositions are plotted against U-Th and U-Pb ages for individual zircon spots. Average zircon δ^{18} O values for each unit are shown by the *solid horizontal lines*. For the South Biscuit Basin flow, we include two average zircon δ^{18} O values (zircon cores and their rims). Equilibrium zircon δ^{18} O values, calculated using host glass δ^{18} O values and Δ^{18} O melt-zircon = 1.9% (applicable for liquidus

temperatures of $\sim 800^{\circ}$ C; Bindeman and Valley 2002), are shown by the *dashed horizontal lines*. For reference, we include the age and oxygen isotope field for Lava Creek Tuff (LCT) zircons (*shaded* Bindeman et al. 2008) and eruption ages of the studied post-caldera rhyolites (*vertical gray lines* see Fig. 4 for references). Uncertainties (1 σ) for individual zircon δ^{18} O measurements are shown in the *bottom left corner* of *each panel*



whereas no antecrysts were apparent by our analyses. The error-weighted mean U-Th age for the Elephant Back flow (174 \pm 23 ka, n = 12, MSWD = 0.3) is 21 ky older than this unit's published K-Ar sanidine eruption age (Obradovich 1992) (Fig. 3). A few older zircon crystals in the Elephant Back flow and the Tuff of Bluff Point were found to be in ²³⁸U-²³⁰Th secular equilibrium and redated with U-Pb (dark bars in Fig. 3). However, these older zircon grains are within uncertainty of the U-Th zircon age dating methods. Vazquez and Reid (2002) reported the presence of older zircon grains in several CPM units (Pitchstone Plateau flow, Solfatara Plateau flow, West Yellowstone flow, and Dry Creek flow) that match the $\sim 250-300$ ka ages of the older grains that we find for the Elephant Back flow and Tuff of Bluff Point.

Oxygen isotope ratios determined by ion microprobe for dated CPM zircon spots generally overlap within uncertainty (Table 1) and are about 1% depleted relative to LCT zircons (Fig. 4). Pitchstone Plateau flow and West Yellowstone flow have identical average zircon δ^{18} O values of 2.8 ± 0.2 and $2.8 \pm 0.1\%$, respectively, which closely approximate the calculated equilibrium zircon δ^{18} O values based on laser-fluorination data for host glass and major phenocryst phases of quartz and sanidine. The Tuff of Bluff Point and Elephant Back flow have slightly lower, but overlapping, average zircon δ^{18} O values of 2.7 \pm 0.1 and $2.7 \pm 0.2\%$, respectively (Fig. 4). Calculated equilibrium zircon δ^{18} O values for the Tuff of Bluff Point and Elephant Back flow are about 0.25% lower than the average zircon δ^{18} O values obtained by ion microprobe, but overlap the equilibrium estimates within error (Fig. 4).

Comparison of oxygen isotope data for all studied Yellowstone rhyolites

When our new zircon data are combined with all published ion microprobe oxygen isotope data for Yellowstone rhyolites (Fig. 5a, arrows indicate the new units presented in this study), the following features are apparent: (1) East Biscuit Basin flow has zircons with oxygen isotope values that are generally higher than other post-LCT units, but has a low- δ^{18} O glass value that is distinctive of UBM lavas, (2) North Biscuit Basin zircons fall within the oxygen isotope range defined by CPM lavas, (3) CPM rhyolites have zircons with more homogeneous oxygen isotope compositions than older Yellowstone rhyolites, (4) CPM zircons generally have oxygen isotopic compositions that are in equilibrium with coexisting phenocrysts and groundmass glasses, and (5) zircon compositions of individual CPM units overlap within a relatively narrow δ^{18} O range. Taken together, this new evidence indicates that CPM rhyolites can be distinguished from older Yellowstone rhyolites by their relatively homogeneous low- δ^{18} O zircon oxygen isotope compositions that are in equilibrium with host melts, whereas UBM rhyolites are more complex, retaining significant zircon oxygen isotope heterogeneity and disequilibria in low- δ^{18} O melts. In addition to a trend of increasing homogeneity through time (Fig. 5a), we also observe a positive correlation between zircon homogeneity and eruptive volume, in which more voluminous post-caldera rhyolites tend to have narrower zircon δ^{18} O ranges (Fig. 5b).

Lead isotope compositions of individual sanidines

Major caldera-forming tuffs

Lead isotope compositions measured for individual sanidine crystals from HRT and LCT rocks define separate arrays that can be used to differentiate individual units (Fig. 6a). Similar results were obtained for groundmass glasses of HRT, MFT, and LCT, in which case Pb isotopes can be used to distinguish different compositional fields of tuff members (Doe et al. 1982; Hildreth et al. 1991, light gray ellipses in Fig. 6a). Our new Pb isotope data for individual sanidines show that these isotopic differences also exist on the crystal scale, and we find that sanidines from two different members of the first major caldera-forming eruption at Yellowstone, HRT members B and C, bracket the entire range of Yellowstone rhyolite Pb isotope compositions (Fig. 6a). HRT member C has a bimodal Pb isotope composition, with one population that possesses an exceptionally radiogenic Pb isotope signature, while HRT member B has the least radiogenic Pb isotope signature of the units analyzed in this study, falling within the compositional field of Yellowstone basalts (dark gray field in Fig. 6a). In addition, HRT member B has more homogeneous sanidine Pb isotope compositions than HRT member C or LCT member A (Fig. 6a), perhaps indicating less crustal contamination in HRT member B relative to other tuff members.

Central Plateau Member rhyolites

The isotopic array defined by CPM sanidines is distinct from those of the caldera-forming tuffs that predate them (Fig. 6a). The fact that the CPM sanidine field does not intersect the LCT field (Fig. 6a) indicates that CPM rhyolites cannot be magmatic remnants of the LCT batholith, in contrast to the early interpretations of Hildreth et al. (1984). We observe that CPM lead isotope ratios fall between the HRT and LCT fields (Fig. 6a), consistent with the interpretations of Bindeman et al. (2008) for derivation of post-LCT rhyolites from



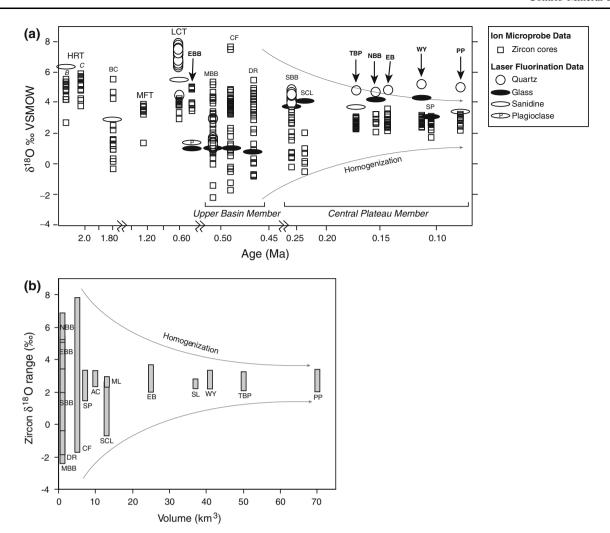


Fig. 5 a Ion microprobe analyses of δ^{18} O in zircon (*squares*) plotted against eruption ages for Yellowstone rhyolites. Laser-fluorination δ^{18} O data for quartz (*circles*), sanidine (*open ovals*), and glass (*filled ovals*) are also included. Our new oxygen isotope data for six post-Lava Creek Tuff units (denoted by *arrows*) are synthesized with data published by Bindeman et al. (2008). Our new data reveal a trend of increasing homogenization of zircon δ^{18} O compositions with decreasing eruption age (trend shown with *light gray arrows*). Unit abbreviations are as follows: Huckleberry Ridge Tuff (HRT) members B and C; Blue Creek flow (BC); Mesa Falls Tuff (MFT); Lava Creek Tuff (LCT); East Biscuit Basin flow (EBB); Middle Biscuit Basin flow (MBB); Canyon flow (CF); Dunraven Road flow (DR); South Biscuit Basin flow (SBB); Scaup Lake flow (SCL); Tuff of Bluff Point (TBP); North Biscuit Basin flow (NBB); Elephant Back flow (EB); West Yellowstone flow (WY); Solfatara Plateau flow (SP);

Pitchstone Plateau flow (PP). Eruption ages are from Obradovich (1992), Gansecki et al. (1996), Lanphere et al. (2002), Christiansen et al. (2007), Bindeman et al. (2008), and this study. No eruption age is available for the post-LCT EBB flow; it is plotted immediately after LCT. Symbols for SBB and SCL, and NBB and EB were slightly offset from one another to avoid overlap. Modified after Bindeman et al. (2008). **b** Zircon δ^{18} O ranges plotted against eruptive volumes for post-Lava Creek Tuff rhyolites. Zircon δ^{18} O ranges are from Bindeman and Valley (2001), Bindeman et al. (2008) and this study. Eruptive volumes are from Christiansen et al. (2007). Our new data reveal a trend of increasing homogenization of zircon δ^{18} O compositions with increasing eruptive volume (trend shown with light gray arrows). Unit abbreviations are the same as those used in **a**, with the addition of post-Lava Creek Tuff units Aster Creek flow (AC), Mallard Lake flow (ML), and Summit Lake flow (SL)

remelting of HRT, LCT, and pre-LCT intracaldera rocks; however, we note that other isotopic parameters necessitate contribution from additional sources, and therefore, it is not possible to derive post-LCT rhyolites through simple binary mixing of these two end members (see next section). Similar to the results for Pb isotope compositions of CPM groundmass glasses (Doe et al. 1982; Vazquez et al. 2009, light gray ellipses in Fig. 6a, b), the

CPM sanidines analyzed in this study have overlapping Pb isotope compositions that define a unique array. Glass analyses from our rhyolite mounts (open symbols) fall within the Pb isotope fields defined by the sanidines (filled symbols) for each CPM rhyolite (Fig. 6b). One notable outlier of the CPM group is the Solfatara Plateau flow ("SP" in Fig. 6a), which has a glass composition that falls outside of the defined CPM lead isotope field



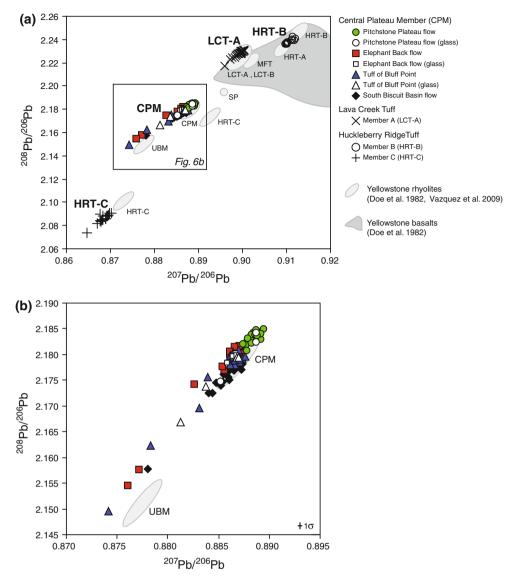


Fig. 6 a Lead isotope compositions of individual Yellowstone sanidines determined by laser ablation (see Table 2 for data) and published whole-rock and glass compositions of Yellowstone rhyolites (*gray ellipses*) and basalts (*dark gray field*). Host glass analyses for the Central Plateau Member (CPM) rhyolites analyzed in this study are shown by the *open symbols*. Distinct lead isotope compositions are apparent for Huckleberry Ridge Tuff members B and C (HRT-B and HRT-C), Lava Creek Tuff member A (LCT-A), and CPM rhyolites. Note that post-LCT CPM rhyolites do not overlap the LCT compositional array and therefore cannot be remnants of LCT magma that remained after caldera collapse. Abbreviations for Yellowstone units from Doe et al. (1982) and Vazquez et al. (2009)

(Vazquez et al. 2009). We also observe that the Elephant Back flow, Tuff of Bluff Point, and South Biscuit Basin flow have heterogeneous sanidine compositions, some of which more closely approximate the Pb isotopic composition of the UBM rather than CPM array (Fig. 6b). In contrast, the Pitchstone Plateau flow sanidines are the most homogeneous and form a less radiogenic Pb isotopic end member (Fig. 6b).

are as follows: Huckleberry Ridge Tuff member A (HRT-A), Mesa Falls Tuff (MFT), Lava Creek Tuff member B (LCT-B), Solfatara Plateau flow (SP), Upper Basin Member (UBM). **b** Higher resolution version of **a**, showing lead isotope variation within the CPM rhyolite group. Note that the Pitchstone Plateau flow sanidines and glasses (filled and open circles) form a distinct lead isotope end member in the CPM array and have more homogeneous lead isotope compositions than older CPM units. Older CPM units exhibit more heterogeneity and have lead isotopic compositions that are more similar to those for UBM rhyolites. The average external error $(\pm 1\sigma)$ for lead isotope analyses is shown in the bottom right corner

Comparison with published radiogenic isotope datasets

It is apparent from published Sr, Nd, and Pb isotope datasets for CPM whole-rocks and glasses that multiple sources, including Yellowstone basalts and highly radiogenic Archean crust, must be accounted for in any realistic isotopic mixing models for Yellowstone rhyolites (Hildreth et al. 1991; Vazquez et al. 2009; Watts et al.



Table 2 Pb isotopes of sanidines in studied Yellowstone rhyolites

Sample 2 Pb isotop	²⁰⁶ Pb/ ²⁰⁴ Pb	$\pm 1\sigma$ ext	²⁰⁷ Pb/ ²⁰⁴ Pb	$\pm 1\sigma$ ext	²⁰⁸ Pb/ ²⁰⁴ Pb	$\pm 1\sigma$ ext	²⁰⁷ Pb/ ²⁰⁶ Pb	$\pm 1\sigma$ ext	²⁰⁸ Pb/ ²⁰⁶ Pb	$\pm 1\sigma$ ext
Pitchstone Plateau	flow (YL02-1									
YL02-1_1.1	13.37	0.32	11.89	0.29	29.07	0.72	0.8889	0.0006	2.1844	0.0009
YL02-1_1.2	13.02	0.32	11.58	0.29	28.33	0.72	0.8883	0.0006	2.1837	0.0009
YL02-1_1.3	13.46	0.32	11.96	0.29	29.25	0.72	0.8887	0.0006	2.1838	0.0009
YL02-1_1.4	13.51	0.32	11.99	0.29	29.36	0.72	0.8884	0.0006	2.1839	0.0009
YL02-1_2.1	14.27	0.32	12.67	0.29	31.02	0.72	0.8884	0.0006	2.1830	0.0009
YL02-1_2.2	14.07	0.32	12.50	0.29	30.62	0.72	0.8884	0.0006	2.1841	0.0009
YL02-1_2.3	14.24	0.32	12.66	0.29	30.99	0.72	0.8889	0.0006	2.1836	0.0009
YL02-1_4.1	11.88	0.32	10.52	0.29	25.76	0.72	0.8878	0.0006	2.1823	0.0009
YL02-1_4.2	12.62	0.32	11.20	0.29	27.39	0.72	0.8882	0.0006	2.1831	0.0009
YL02-1_4.3	12.28	0.32	10.91	0.29	26.68	0.72	0.8882	0.0006	2.1837	0.0009
YL02-1_5.1	12.13	0.32	10.77	0.29	26.30	0.72	0.8892	0.0006	2.1830	0.0009
YL02-1_5.2	8.24	0.32	7.31	0.29	17.69	0.72	0.8887	0.0006	2.1846	0.0009
YL02-1_5.3	11.75	0.32	10.41	0.29	25.47	0.72	0.8888	0.0006	2.1837	0.0009
YL02-1_6.1	11.37	0.32	10.09	0.29	24.60	0.72	0.8877	0.0006	2.1818	0.0009
YL02-1_6.2	11.83	0.32	10.50	0.29	25.63	0.72	0.8883	0.0006	2.1822	0.0009
YL02-1_6.3	11.74	0.32	10.43	0.29	25.43	0.72	0.8886	0.0006	2.1837	0.0009
YL02-1_7.1	10.97	0.32	9.72	0.29	23.78	0.72	0.8879	0.0006	2.1832	0.0009
YL02-1_7.2	11.61	0.32	10.29	0.29	25.15	0.72	0.8874	0.0006	2.1820	0.0009
YL02-1_7.3	12.55	0.32	11.13	0.29	27.21	0.72	0.8878	0.0006	2.1808	0.0009
YL02-1_8.1	12.48	0.32	11.07	0.29	27.09	0.72	0.8895	0.0006	2.1849	0.0009
YL02-1_8.2	12.55	0.32	11.14	0.29	27.23	0.72	0.8892	0.0006	2.1839	0.0009
YL02-1_8.3	12.77	0.32	11.34	0.29	27.72	0.72	0.8888	0.0006	2.1837	0.0009
YL02-1_glass-1	13.75	0.32	12.22	0.29	29.90	0.72	0.8887	0.0006	2.1823	0.0009
YL02-1_glass-2	12.48	0.32	11.04	0.29	26.99	0.72	0.8853	0.0006	2.1748	0.0009
YL02-1_glass-3	12.70	0.32	11.24	0.29	27.53	0.72	0.8887	0.0006	2.1841	0.0009
Elephant Back flow	v (06YS-4)									
06YS-4_2.1	11.28	0.18	9.88	0.16	24.13	0.38	0.8772	0.0004	2.1575	0.0012
06YS-4_2.2	11.52	0.18	10.20	0.16	24.93	0.38	0.8870	0.0004	2.1816	0.0012
06YS-4_2.3	11.65	0.18	10.20	0.16	24.93	0.38	0.8760	0.0004	2.1545	0.0012
06YS-4_2.4	11.40	0.18	10.08	0.16	24.63	0.38	0.8868	0.0004	2.1798	0.0012
06YS-4_3.1	8.79	0.18	7.75	0.16	18.86	0.38	0.8865	0.0004	2.1801	0.0012
06YS-4_3.2	8.16	0.18	7.18	0.16	17.46	0.38	0.8862	0.0004	2.1801	0.0012
06YS-4_3.3	10.43	0.18	9.22	0.16	22.49	0.38	0.8869	0.0004	2.1798	0.0012
06YS-4_4.1	11.08	0.18	9.79	0.16	23.91	0.38	0.8867	0.0004	2.1796	0.0012
06YS-4_4.2	11.82	0.18	10.47	0.16	25.56	0.38	0.8874	0.0004	2.1802	0.0012
06YS-4_4.3	12.59	0.18	11.15	0.16	27.25	0.38	0.8870	0.0004	2.1792	0.0012
06YS-4_5.1	12.38	0.18	10.96	0.16	26.80	0.38	0.8865	0.0004	2.1795	0.0012
06YS-4_5.2	12.77	0.18	11.31	0.16	27.67	0.38	0.8867	0.0004	2.1806	0.0012
06YS-4_5.3	13.18	0.18	11.68	0.16	28.58	0.38	0.8868	0.0004	2.1794	0.0012
06YS-4_6.1	8.96	0.18	7.87	0.16	19.19	0.38	0.8827	0.0004	2.1742	0.0012
06YS-4_6.2	10.61	0.18	9.37	0.16	22.87	0.38	0.8856	0.0004	2.1769	0.0012
06YS-4_6.3	11.70	0.18	10.32	0.16	25.28	0.38	0.8854	0.0004	2.1776	0.0012
06YS-4_7.1	10.36	0.18	9.16	0.16	22.36	0.38	0.8866	0.0004	2.1815	0.0012
06YS-4_7.2	13.19	0.18	11.70	0.16	28.60	0.38	0.8873	0.0004	2.1793	0.0012
06YS-4_7.3	9.93	0.18	8.78	0.16	21.40	0.38	0.8861	0.0004	2.1805	0.0012
06YS-4_glass-1	11.41	0.18	10.07	0.16	24.65	0.38	0.8859	0.0004	2.1784	0.0012
06YS-4_glass-2	14.42	0.18	12.79	0.16	31.33	0.38	0.8867	0.0004	2.1801	0.0012



Table 2 continued

Sample	²⁰⁶ Pb/ ²⁰⁴ Pb	$\pm 1\sigma$ ext	²⁰⁷ Pb/ ²⁰⁴ Pb	$\pm 1\sigma$ ext	²⁰⁸ Pb/ ²⁰⁴ Pb	$\pm 1\sigma$ ext	²⁰⁷ Pb/ ²⁰⁶ Pb	$\pm 1\sigma$ ext	²⁰⁸ Pb/ ²⁰⁶ Pb	$\pm 1\sigma$ ext
06YS-4_glass-3	8.54	0.18	7.53	0.16	18.31	0.38	0.8864	0.0004	2.1796	0.0012
06YS-4_glass-4	11.07	0.18	9.79	0.16	23.90	0.38	0.8868	0.0004	2.1797	0.0012
Tuff of Bluff Poin	t (06YS-2)									
06YS-2_1.1	12.53	0.18	11.09	0.16	27.13	0.38	0.8863	0.0004	2.1796	0.0012
06YS-2_1.2	12.36	0.18	10.95	0.16	26.75	0.38	0.8871	0.0004	2.1793	0.0012
06YS-2_1.3	12.88	0.18	11.40	0.16	27.90	0.38	0.8862	0.0004	2.1797	0.0012
06YS-2_1.4	12.76	0.18	11.30	0.16	27.65	0.38	0.8869	0.0004	2.1809	0.0012
06YS-2_3.1	7.42	0.18	6.53	0.16	15.83	0.38	0.8865	0.0004	2.1804	0.0012
06YS-2_3.2	7.46	0.18	6.56	0.16	15.91	0.38	0.8867	0.0004	2.1798	0.0012
06YS-2_3.3	8.12	0.18	7.16	0.16	17.36	0.38	0.8870	0.0004	2.1786	0.0012
06YS-2_3.4	8.16	0.18	7.19	0.16	17.45	0.38	0.8871	0.0004	2.1793	0.0012
06YS-2_4.1	8.24	0.18	7.26	0.16	17.64	0.38	0.8864	0.0004	2.1796	0.0012
06YS-2_4.2	8.76	0.18	7.72	0.16	18.77	0.38	0.8866	0.0004	2.1779	0.0012
06YS-2_4.3	8.46	0.18	7.47	0.16	18.13	0.38	0.8874	0.0004	2.1793	0.0012
06YS-2_5.1	11.23	0.18	9.92	0.16	24.23	0.38	0.8861	0.0004	2.1779	0.0012
06YS-2_5.2	11.18	0.18	9.89	0.16	24.14	0.38	0.8866	0.0004	2.1789	0.0012
06YS-2_5.3	11.81	0.18	10.45	0.16	25.53	0.38	0.8871	0.0004	2.1790	0.0012
06YS-2_6.1	6.42	0.18	5.57	0.16	13.45	0.38	0.8742	0.0004	2.1495	0.0012
06YS-2_6.2	7.17	0.18	6.29	0.16	15.22	0.38	0.8831	0.0004	2.1696	0.0012
06YS-2_6.3	11.34	0.18	10.01	0.16	24.46	0.38	0.8839	0.0004	2.1755	0.0012
06YS-2_6.4	8.53	0.18	7.46	0.16	18.15	0.38	0.8783	0.0004	2.1623	0.0012
06YS-2_9.1	10.37	0.18	9.17	0.16	22.35	0.38	0.8876	0.0004	2.1795	0.0012
06YS-2_glass-1	8.44	0.18	7.44	0.16	18.07	0.38	0.8868	0.0004	2.1795	0.0012
06YS-2_glass-2	5.37	0.18	4.70	0.16	11.29	0.38	0.8870	0.0004	2.1794	0.0012
06YS-2_glass-3	4.56	0.18	3.95	0.16	9.45	0.38	0.8813	0.0004	2.1668	0.0012
06YS-2_glass-4	5.41	0.18	4.70	0.16	11.34	0.38	0.8837	0.0004	2.1737	0.0012
South Biscuit Basi	n flow (YL96	-2)								
YL96-2_1.1	11.18	0.18	9.90	0.16	24.17	0.38	0.8872	0.0004	2.1811	0.0012
YL96-2_1.2	14.02	0.18	12.44	0.16	30.43	0.38	0.8867	0.0004	2.1786	0.0012
YL96-2_1.3	14.16	0.18	12.55	0.16	30.69	0.38	0.8863	0.0004	2.1779	0.0012
YL96-2_1.4	14.42	0.18	12.80	0.16	31.33	0.38	0.8869	0.0004	2.1782	0.0012
YL96-2_2.1	14.40	0.18	12.77	0.16	31.25	0.38	0.8865	0.0004	2.1769	0.0012
YL96-2_2.2	14.87	0.18	13.17	0.16	32.25	0.38	0.8861	0.0004	2.1770	0.0012
YL96-2_2.3	14.99	0.18	13.28	0.16	32.54	0.38	0.8858	0.0004	2.1768	0.0012
YL96-2_3.1	14.20	0.18	12.56	0.16	30.76	0.38	0.8848	0.0004	2.1745	0.0012
YL96-2_3.2	14.52	0.18	12.86	0.16	31.48	0.38	0.8852	0.0004	2.1739	0.0012
YL96-2_3.3	15.34	0.18	13.59	0.16	33.28	0.38	0.8854	0.0004	2.1747	0.0012
YL96-2_4.1	14.56	0.18	12.90	0.16	31.58	0.38	0.8860	0.0004	2.1757	0.0012
YL96-2_4.2	14.90	0.18	13.22	0.16	32.36	0.38	0.8869	0.0004	2.1778	0.0012
YL96-2_4.3	15.18	0.18	13.45	0.16	32.95	0.38	0.8863	0.0004	2.1773	0.0012
YL96-2_5.1	11.22	0.18	9.92	0.16	24.21	0.38	0.8866	0.0004	2.1778	0.0012
YL96-2_5.2	12.25	0.18	10.85	0.16	26.49	0.38	0.8869	0.0004	2.1778	0.0012
YL96-2_5.3	11.48	0.18	10.15	0.16	24.86	0.38	0.8860	0.0004	2.1751	0.0012
YL96-2_6.1	14.29	0.18	12.66	0.16	30.99	0.38	0.8864	0.0004	2.1772	0.0012
YL96-2_6.2	14.68	0.18	13.03	0.16	31.87	0.38	0.8860	0.0004	2.1770	0.0012
YL96-2_6.3	14.61	0.18	12.95	0.16	31.73	0.38	0.8863	0.0004	2.1779	0.0012
YL96-2_7.1	10.25	0.18	8.98	0.16	21.89	0.38	0.8780	0.0004	2.1578	0.0012
YL96-2_7.2	10.40	0.18	9.18	0.16	22.38	0.38	0.8840	0.0004	2.1724	0.0012



Table 2 continued

Sample	²⁰⁶ Pb/ ²⁰⁴ Pb	$\pm 1\sigma$ ext	²⁰⁷ Pb/ ²⁰⁴ Pb	$\pm 1\sigma$ ext	²⁰⁸ Pb/ ²⁰⁴ Pb	$\pm 1\sigma$ ext	²⁰⁷ Pb/ ²⁰⁶ Pb	$\pm 1\sigma$ ext	²⁰⁸ Pb/ ²⁰⁶ Pb	$\pm 1\sigma$ ext
YL96-2_7.3	10.89	0.18	9.61	0.16	23.44	0.38	0.8844	0.0004	2.1725	0.0012
YL96-2_8.1	13.25	0.18	11.74	0.16	28.69	0.38	0.8855	0.0004	2.1762	0.0012
YL96-2_8.2	13.78	0.18	12.22	0.16	29.88	0.38	0.8873	0.0004	2.1771	0.0012
YL96-2_8.3	14.12	0.18	12.52	0.16	30.63	0.38	0.8874	0.0004	2.1782	0.0012
Lava Creek Tuff,	Member A (Le	CT-3A)								
LCT-3A_1.1	16.20	0.32	14.57	0.29	36.01	0.72	0.8996	0.0006	2.2298	0.0009
LCT-3A_1.2	16.13	0.32	14.50	0.29	35.84	0.72	0.8989	0.0006	2.2279	0.0009
LCT-3A_1.3	16.66	0.32	14.99	0.29	37.02	0.72	0.8986	0.0006	2.2261	0.0009
LCT-3A_1.4	16.50	0.32	14.85	0.29	36.67	0.72	0.8996	0.0006	2.2273	0.0009
LCT-3A_2.1	16.52	0.32	14.87	0.29	36.79	0.72	0.8998	0.0006	2.2298	0.0009
LCT-3A_2.2	16.68	0.32	15.02	0.29	37.12	0.72	0.9004	0.0006	2.2312	0.0009
LCT-3A_2.3	16.70	0.32	15.03	0.29	37.20	0.72	0.8999	0.0006	2.2299	0.0009
LCT-3A_3.1	16.59	0.32	14.91	0.29	36.87	0.72	0.8992	0.0006	2.2277	0.0009
LCT-3A_3.2	16.75	0.32	15.07	0.29	37.26	0.72	0.8995	0.0006	2.2289	0.0009
LCT-3A_3.3	16.80	0.32	15.14	0.29	37.39	0.72	0.8998	0.0006	2.2293	0.0009
LCT-3A_4.1	16.68	0.32	14.97	0.29	37.01	0.72	0.8972	0.0006	2.2229	0.0009
LCT-3A_4.2	16.80	0.32	15.12	0.29	37.32	0.72	0.8992	0.0006	2.2279	0.0009
LCT-3A_4.3	16.86	0.32	15.19	0.29	37.50	0.72	0.8994	0.0006	2.2283	0.0009
LCT-3A_5.1	16.69	0.32	15.01	0.29	37.12	0.72	0.8997	0.0006	2.2292	0.0009
LCT-3A_5.2	16.77	0.32	15.11	0.29	37.34	0.72	0.9004	0.0006	2.2304	0.0009
LCT-3A_5.3	16.84	0.32	15.17	0.29	37.48	0.72	0.9004	0.0006	2.2306	0.0009
LCT-3A_6.1	16.75	0.32	15.08	0.29	37.27	0.72	0.8999	0.0006	2.2279	0.0009
LCT-3A_6.2	16.74	0.32	15.08	0.29	37.24	0.72	0.9002	0.0006	2.2290	0.0009
LCT-3A_6.3	16.88	0.32	15.18	0.29	37.56	0.72	0.8997	0.0006	2.2281	0.0009
LCT-3A_7.1	16.60	0.32	14.90	0.29	36.74	0.72	0.8959	0.0006	2.2175	0.0009
LCT-3A_7.2	16.75	0.32	15.06	0.29	37.24	0.72	0.8977	0.0006	2.2244	0.0009
LCT-3A_7.3	16.74	0.32	15.03	0.29	37.14	0.72	0.8983	0.0006	2.2249	0.0009
LCT-3A_8.1	17.06	0.32	15.34	0.29	37.93	0.72	0.8988	0.0006	2.2270	0.0009
LCT-3A_8.2	17.05	0.32	15.38	0.29	38.03	0.72	0.9002	0.0006	2.2312	0.0009
LCT-3A_8.3	16.85	0.32	15.18	0.29	37.53	0.72	0.8998	0.0006	2.2293	0.0009
Huckleberry Ridge	e Tuff, Membe	er B (HRT-	1)							
HRT-1_1.1	16.14	0.32	14.65	0.29	35.90	0.72	0.9099	0.0006	2.2361	0.0009
HRT-1_1.2	16.40	0.32	14.94	0.29	36.63	0.72	0.9101	0.0006	2.2372	0.0009
HRT-1_1.3	16.62	0.32	15.13	0.29	37.11	0.72	0.9121	0.0006	2.2405	0.0009
HRT-1_2.1	16.12	0.32	14.64	0.29	35.94	0.72	0.9097	0.0006	2.2359	0.0009
HRT-1_2.2	16.25	0.32	14.78	0.29	36.24	0.72	0.9100	0.0006	2.2366	0.0009
HRT-1_2.3	16.05	0.32	14.63	0.29	35.90	0.72	0.9118	0.0006	2.2391	0.0009
HRT-1_3.1	15.95	0.32	14.52	0.29	35.59	0.72	0.9102	0.0006	2.2356	0.0009
HRT-1_3.2	16.17	0.32	14.73	0.29	36.11	0.72	0.9115	0.0006	2.2392	0.0009
HRT-1_3.3	16.30	0.32	14.82	0.29	36.34	0.72	0.9103	0.0006	2.2364	0.0009
HRT-1_4.1	16.21	0.32	14.75	0.29	36.17	0.72	0.9105	0.0006	2.2376	0.0009
HRT-1_4.2	16.20	0.32	14.75	0.29	36.19	0.72	0.9113	0.0006	2.2401	0.0009
HRT-1_4.3	16.24	0.32	14.78	0.29	36.26	0.72	0.9116	0.0006	2.2422	0.0009
Huckleberry Ridge	e Tuff, Membe	er C (HRT-	C)							
HRT-C_3.1	17.40	0.32	15.18	0.29	36.46	0.72	0.8693	0.0006	2.088	0.0009
HRT-C_3.2	17.47	0.32	15.27	0.29	36.69	0.72	0.8702	0.0006	2.090	0.0009
HRT-C_3.3	17.08	0.32	14.90	0.29	35.78	0.72	0.8695	0.0006	2.089	0.0009
HRT-C_5.1	17.72	0.32	15.44	0.29	37.09	0.72	0.8682	0.0006	2.085	0.0009



Table 2 continued

Sample	²⁰⁶ Pb/ ²⁰⁴ Pb	$\pm 1\sigma$ ext	²⁰⁷ Pb/ ²⁰⁴ Pb	$\pm 1\sigma$ ext	²⁰⁸ Pb/ ²⁰⁴ Pb	$\pm 1\sigma$ ext	²⁰⁷ Pb/ ²⁰⁶ Pb	$\pm 1\sigma$ ext	²⁰⁸ Pb/ ²⁰⁶ Pb	$\pm 1\sigma$ ext
HRT-C_5.2	17.79	0.32	15.49	0.29	37.23	0.72	0.8690	0.0006	2.086	0.0009
HRT-C_5.3	17.71	0.32	15.44	0.29	37.10	0.72	0.8688	0.0006	2.086	0.0009
HRT-C_6.1	17.56	0.32	15.30	0.29	36.74	0.72	0.8682	0.0006	2.085	0.0009
HRT-C_6.2	17.58	0.32	15.31	0.29	36.76	0.72	0.8676	0.0006	2.084	0.0009
HRT-C_6.3	17.70	0.32	15.42	0.29	37.03	0.72	0.8679	0.0006	2.083	0.0009
HRT-C_7.1	14.30	0.32	12.44	0.29	29.89	0.72	0.8682	0.0006	2.088	0.0009
HRT-C_7.2	16.96	0.32	14.79	0.29	35.55	0.72	0.8697	0.0006	2.090	0.0009
HRT-C_7.3	15.59	0.32	13.56	0.29	32.63	0.72	0.8677	0.0006	2.090	0.0009
HRT-C_8.1	16.87	0.32	15.20	0.29	37.36	0.72	0.9017	0.0006	2.221	0.0009
HRT-C_8.2	16.95	0.32	15.29	0.29	37.59	0.72	0.9020	0.0006	2.221	0.0009
HRT-C_8.3	16.86	0.32	15.21	0.29	37.38	0.72	0.9016	0.0006	2.220	0.0009
HRT-C_9.1	17.49	0.32	15.24	0.29	36.63	0.72	0.8691	0.0006	2.088	0.0009
HRT-C_10.1	18.22	0.32	15.81	0.29	37.94	0.72	0.8647	0.0006	2.073	0.0009
HRT-C_11.1	18.01	0.32	15.67	0.29	37.63	0.72	0.8671	0.0006	2.082	0.0009
HRT-C_12.1	17.78	0.32	15.50	0.29	37.25	0.72	0.8694	0.0006	2.088	0.0009

Uncertainties of Pb isotope ratios are $\pm 1\sigma$ external errors for a NIST 612 glass standard (Baker et al. 2004)

We note that HRT-C grain 8 is an outlier and not included in Fig. 6a

2010; Pritchard and Larson in review). Our new Pb isotope data represent a departure from prior work in that they were acquired for single sanidine crystals, rather than whole-rock or glass averages, and with a different technique, LA-ICP-MS for single spot analyses rather than TIMS for bulk analyses. More high-resolution single crystal analyses are needed for detailed assessments of radiogenic source components in Yellowstone magmas. However, using our existing sanidine lead isotope dataset, we are able to make several important observations about CPM rhyolites in the context of published radiogenic isotope datasets: (1) Compared to individual tuff members of Yellowstone's caldera-forming eruptions, post-caldera CPM rhyolites define a larger range in radiogenic isotope space, (2) The CPM array is unique and does not overlap any of the established radiogenic isotope fields for Yellowstone rhyolites or basalts, (3) Crystals within CPM rhyolites have heterogeneous radiogenic isotope composition that become progressively more homogenous in younger units, and (4) A significant proportion of CPM crystals have radiogenic isotope compositions that are characteristic of UBM rather than CPM units.

Thermometry of analyzed rhyolites

We determined zircon saturation temperatures and liquidus temperatures for all analyzed rhyolites and synthesized this new data with temperature data published by Bindeman and Valley (2001) and Vazquez et al. (2009) (Fig. 7). When temperatures are plotted against melt δ^{18} O values for

individual UBM units, we find that they have similar oxygen isotope compositions and temperature ranges (~ 820 –900°C) and that the oldest and most primitive (~ 70 wt% SiO₂) East Biscuit Basin flow also has the highest liquidus temperature (~ 940 °C; Fig. 7a). Temperatures and melt δ^{18} O values for CPM rhyolites are more variable than UBM rhyolites, with temperatures that range from ~ 750 to 900°C (Fig. 7a). We observe a trend of decreasing temperature with increasing melt δ^{18} O values for CPM rhyolites, but the temperature ranges for individual CPM units are often quite large due to discrepancies between the different geothermometers. Therefore, we interpret this trend with caution; more precise thermometry data are needed before this trend can be confirmed or refuted.

When temperatures of CPM rhyolites are plotted against their eruption ages (Fig. 7b), it appears that they become progressively cooler through time. Vazquez et al. (2009) first described this trend using a smaller dataset that is now synthesized with our new data in Fig. 7b. Using a cumulative eruptive volume curve constructed with published eruptive volumes and eruption ages for CPM units (Christiansen et al. 2007), we observe an inverse correlation between temperature and eruptive volume (Fig. 7b). A spike in the cumulative eruptive volume curve, following eruption of the very voluminous 114-ka West Yellowstone flow, marks the point at which temperatures of CPM rhyolites begin to decrease. However, as noted previously, the large temperature ranges between the various thermometers and their inherent uncertainties warrant caution in these observations.



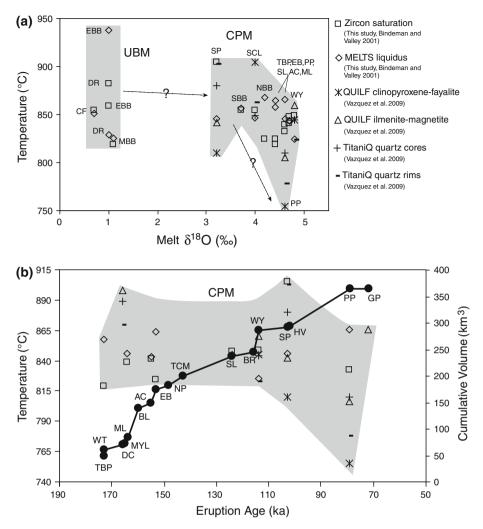


Fig. 7 a Temperatures of Upper Basin Member (UBM) and Central Plateau Member (CPM) rhyolites plotted against melt δ^{18} O values. *Symbols* show the temperatures derived from different thermometry datasets, synthesized from Bindeman and Valley (2001), Vazquez et al. (2009) and this study (see Supplementary Material 3 for temperature data). Melt δ^{18} O values are taken from Bindeman and Valley (2001), Bindeman et al. (2008) and this study. Temperature-melt δ^{18} O fields for UBM and CPM rhyolites are *shaded*. Unit abbreviations are the same as those used in Fig. 5. We tentatively indicate a trend of decreasing temperatures with increasing melt δ^{18} O values (*dashed arrows*); however, the temperature ranges for individual units are often quite large, making this trend difficult to confirm. **b** Temperature and cumulative eruptive volume of Central

Discussion

In the following discussion, we focus on our new age (U-Th and U-Pb) and isotopic (O and Pb) data for post-LCT rhyolites of the Yellowstone caldera. We synthesize our new data with previously published age and geochemical datasets for post-LCT rhyolites to create a unified model of magma genesis. Our discussion is subdivided into three main parts: (1) Model of post-LCT (UBM and CPM) magma genesis, (2) Petrologic, thermal, and temporal

Plateau Member (CPM) rhyolites plotted against eruption age. Thermometry symbols are the same as those used in **a**. Volumes and eruption ages are from Christiansen et al. (2007). The cumulative eruptive volume curve is shown as a black line connected by individual units (black circles). Unit abbreviations are the same as those used in Fig. 5, with the addition of CPM units West Thumb flow (WT), Dry Creek flow (DC), Mary Lake flow (MYL), Buffalo Lake flow (BL), Nez Perce Creek flow (NP), Tuff of Cold Mountain Creek (TCM), Bechler River flow (BC), Hayden Valley flow (HV), and Grants Pass flow (GP). Note that the temperature curve is negatively correlated with the cumulative eruptive volume curve for CPM rhyolites, interpreted here as cooling of the CPM magma reservoir as eruptions proceeded in the Yellowstone caldera

considerations for CPM magmas, and (3) Similarities and differences between CPM magmas and voluminous late-stage magmas of the pre-Yellowstone Heise volcanic field.

Model of post-LCT magma genesis

Our new age and isotopic datasets for CPM rhyolites require several key changes to published post-LCT magma genesis models (Bindeman and Valley 2000, 2001; Vazquez and Reid 2002; Bindeman et al. 2008). The most



salient new result of this study is that unlike UBM rhyolites, which were generated from independent parcels of melt with unique oxygen isotopic compositions (Bindeman et al. 2008), many of the voluminous CPM rhyolites have more homogeneous oxygen isotopic compositions that permit derivation from a common magma source. There are notable exceptions to this rule, for instance the Scaup Lake flow, South Biscuit Basin flow, and Solfatara Plateau flow. However, as we discuss below, the ~ 260 -ka Scaup Lake and South Biscuit Basin flows retain important age and isotopic information about the early stages of CPM magma assembly prior to the loss of crystal evidence by convection and homogenization of a large CPM magma reservoir. We discuss our new model for post-LCT magma genesis chronologically, starting with discussion of the early UBM rhyolite eruptions (\sim 520–470 ka) followed by discussion of the later (~260-75 ka) CPM rhyolite eruptions.

Previous studies of UBM rhyolites (Canyon flow, Dunraven Road flow, Middle Biscuit Basin flow) have shown that they contain a large proportion of zircon xenocrysts with ages and normal- δ^{18} O values that correspond to HRT, LCT, and pre-LCT units (Bindeman et al. 2001, 2008). These zircons are not in oxygen isotopic equilibrium with their host low- δ^{18} O melts and provide evidence for wholesale remelting of hydrothermally altered (¹⁸Odepleted in the groundmass) intracaldera rocks of HRT, LCT, and pre-LCT protoliths (Bindeman et al. 2008; Fig. 8). Our new results for the East Biscuit Basin flow indicate that like other UBM units, it contains zircons with a broad age spectrum and normal- δ^{18} O values that are not in equilibrium with the low- δ^{18} O host melt; however, unlike other UBM units, East Biscuit Basin flow does not contain any zircons with low- δ^{18} O values or younger (eruption) ages. The East Biscuit Basin flow also has a unique mineralogy and major and trace element geochemistry relative to other UBM units, for instance its lack of sanidine and quartz, lower-silica composition, and higher Sr concentration, indicating that it was the first, most primitive rhyolite to erupt with the Yellowstone caldera (Girard and Stix 2009). Thermometry results for the East Biscuit Basin flow show that it also had the hottest liquidus temperature of any post-LCT rhyolite (Fig. 7a). Our new zircon data indicate that the East Biscuit Basin flow was generated by rapid remelting of pre-LCT intracaldera protoliths and erupted before a new phase of (low- δ^{18} O) zircon saturation and crystallization.

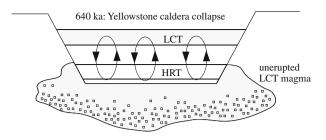
Like UBM rhyolites, the \sim 260-ka Scaup Lake flow has a heterogeneous zircon population with δ^{18} O values that are out of equilibrium with the low- δ^{18} O host melt (Bindeman et al. 2008). Disequilibrium textures of major phenocryst phases, less-evolved mineral and whole-rock geochemistry, and a lower whole-rock Rb/Sr ratio set this

unit apart from other CPM rhyolites (Vazquez and Reid 2002; Vazquez et al. 2009; Girard and Stix 2010). Zircons from the South Biscuit Basin flow have comparably heterogeneous δ^{18} O values, and both units have zircon cores with the distinctive low- δ^{18} O signature of UBM rhyolites, suggesting that they represent remelts of intracaldera UBM intrusions (Bindeman et al. 2008). Our new Pb isotope data reveal that South Biscuit Basin flow also contains sanidine crystals that are compositionally more similar to UBM rather than CPM rhyolites (Fig. 6). Mineral and whole-rock geochemistry of the Scaup Lake flow and South Biscuit Basin flow are nearly identical and define a unique end member composition of CPM rhyolite eruptions (Girard and Stix 2010). Taking this combined evidence into account, we suggest that the Scaup Lake and South Biscuit Basin flows are important bridging links between the UBM and CPM eruptions, representing the oldest, least-evolved CPM magma batches that retain a genetic connection to the plutonic UBM protoliths that sourced them (Fig. 8).

Younger CPM units (Pitchstone Plateau flow, West Yellowstone flow, Elephant Back flow, Tuff of Bluff Point and North Biscuit Basin flow) are distinctly more homogenous in their zircon δ^{18} O values and have mineral assemblages that are in oxygen isotopic equilibrium with low- δ^{18} O groundmass glasses. Our data indicate that these CPM units were derived from batch assembly and homogenization of heterogeneous pockets of low- δ^{18} O melts, such as the older Scaup Lake and South Biscuit Basin flows, which tapped UBM intrusions (Fig. 8). The fact that CPM rhyolites have zircon populations that fall within a narrow δ^{18} O range that approximates the homogenized UBM zircon average (Fig. 5a) supports this interpretation. Furthermore, several CPM units dated by us (Tuff of Bluff Point, Elephant Back flow and North Biscuit Basin flow) and some of those dated by Vazquez and Reid (2002) (Pitchstone Plateau flow, Solfatara Plateau flow, West Yellowstone flow and Dry Creek flow) have older zircon crystals with $\sim 250-300$ ka ages that match those of the Scaup Lake and South Biscuit Basin flows. In addition to zircon crystal evidence for a link between UBM and CPM magmas, we also find that a subset of sanidine crystals within the Tuff of Bluff Point and Elephant Back flow point to a UBM rather than CPM heritage (Fig. 6).

Similar to observations made using whole-rocks and phenocrysts of post-LCT rhyolites for progressively more-evolved, more-juvenile, and cooler CPM magmas through time (Hildreth et al. 1984, 1991; Vazquez et al. 2009; Girard and Stix 2010), our new isotope and age data for single crystals demonstrate that younger CPM rhyolites tend to have less heterogeneous zircon age populations (Fig. 3), slightly higher average zircon δ^{18} O values (Fig. 4), and less heterogeneous, but more-juvenile sanidine lead isotope compositions (Fig. 6). Taken together,





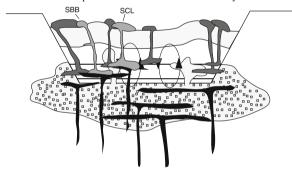
unexposed intracaldera intrusions

-520-470 ka: Upper Basin Member rhyolites

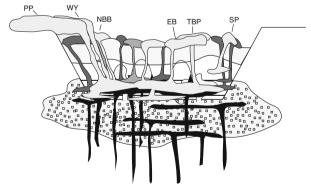
Sour Creek dome

unexposed intracaldera intrusions

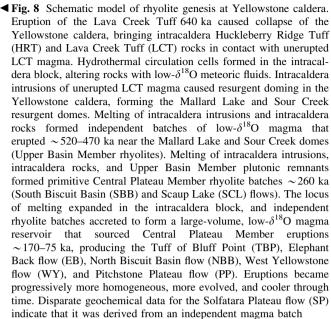




~170-75 ka: Central Plateau Member rhyolites



this evidence indicates that the CPM rhyolites tapped an evolving magma reservoir in which antecrystic and xenocrystic heterogeneity of discrete magma batches was progressively annealed by convection, homogenization, and dissolution–reprecipitation (Fig. 8). Subtle isotopic differences in younger units suggest open-system processes such as assimilation and recharge within the differentiating CPM magma chamber.



Petrologic, thermal, and temporal considerations for CPM eruptions

The sequential nature of CPM rhyolite eruptions and detailed constraints on eruption ages and eruptive volumes for CPM units (Christiansen et al. 2007) make it possible to evaluate magma genesis from the onset of eruptions $\sim\!260~\rm ka$ to the last most evolved eruption $\sim\!75~\rm ka$. In this section, we discuss petrologic, thermal, and temporal considerations for the model of post-LCT magma genesis presented in the previous section.

The earliest erupted CPM rhyolites, the ~ 260 -ka Scaup Lake and South Biscuit Basin flows, define the leastevolved end member composition of CPM rhyolite eruptions in terms of common fractionation indices such as Al₂O₃ (plagioclase and sanidine), TiO₂ (Fe-Ti oxides), Rb/Sr (plagioclase and sanidine), and Zr/Nb (zircon; see Supplementary Material 5). We estimate $\sim 25-30\%$ fractional crystallization (30% plagioclase, 60% sanidine, 10% quartz) to derive the youngest and most evolved Pitchstone Plateau flow from the more-evolved Scaup Lake and South Biscuit Basin flows (see Supplementary Material 6 for fractional crystallization calculations). We note that this estimate is significantly less than the original estimate of >40% fractional crystallization by Vazquez and Reid (2002), which was based on a higher Rb/Sr ratio that is revised to a lower ratio in Vazquez et al. (2009). In addition to the geochemical disparities between old and young CPM rhyolites, distinct petrographic differences are also apparent. Disequilibrium mineral textures are commonly observed in thin sections of the Scaup Lake and South Biscuit Basin flows, indicating incomplete dissolution of



crystals, whereas progressively younger CPM units have more euhedral phenocrysts with fewer disequilibrium mineral textures (see Supplementary Material 2 for photomicrographs).

If younger CPM rhyolites were indeed assembled from batches of early CPM magmas, like the Scaup Lake and South Biscuit Basin flows, crystal dissolution would have had to have been complete by the time of eruption of the Tuff of Bluff Point, North Biscuit Basin flow, and Elephant Back flow (~170-155 ka), which have distinct and homogeneous zircon ages and O isotope compositions relative to the earliest CPM eruptions (Fig. 5a). For a liquidus temperature of 850°C and a Zr undersaturation of 40 ppm, ~ 30 ky is required for complete dissolution of a spherical zircon phenocryst 100 µm in diameter (Watson 1996). This time requirement is consistent with the >90 kyage gap between early ($\sim 260 \text{ ka}$) and late ($\sim 170\text{--}75 \text{ ka}$) CPM eruptions. Based on the exceptionally large cumulative eruptive volume of CPM rhyolites (>600 km³), batch assembly would have had to proceed at a rate of at least 3–4 km³/ky. Assuming basalt can melt 1–2 times the volume of rhyolite, this corresponds to a basaltic input rate of 1–4 km³/ky. Interestingly, this assembly process seems to have been thermally buffered until ~110 ka, based on the similarity of thermometry results for CPM rhyolites up until this point (Fig. 7b). After the eruption of the very voluminous West Yellowstone flow (41 km³), CPM magmas appear to become cooler through time (Fig. 7b). If this trend is real (despite the significant spread in the compiled thermometry data), then this may mark the point in time at which batch assembly stopped beneath the Yellowstone caldera. In this scenario, the negative correlation between cumulative eruptive volume and temperature reflects cooling of the assembled CPM magma reservoir as recorded by sequential rhyolite eruptions. The youngest Pitchstone Plateau flow represents one of the last (and largest) extractions of magma from this assembled source.

Analogy between CPM eruptions and voluminous late-stage magmas of the extinct Heise supervolcano

The Heise caldera complex in eastern Idaho is Yellowstone's most immediate predecessor in the Yellowstone hot spot track (Pierce and Morgan 1992). By analogy with Yellowstone, the terminal stage of caldera cluster evolution at Heise produced the large-volume (>1,800 km³) low- $\delta^{18}{\rm O}$ Kilgore Tuff and post-Kilgore rhyolites, comparable in timing, magnitude of $\delta^{18}{\rm O}$ depletion and cumulative eruptive volume to the CPM phase of rhyolitic eruptions at Yellowstone (Watts et al. 2011). While individual Kilgore Tuff zircons exhibit a wide range in $\delta^{18}{\rm O}$ values, from -1.3 to 6.1‰, bulk zircon and less refractory phenocrysts phases of quartz and sanidine have homogeneous $\delta^{18}{\rm O}$ values that correspond to a low- $\delta^{18}{\rm O}$ magma value of 3.4 \pm 0.1‰ for samples erupted >100 km apart, providing a compelling case for batch assembly and homogenization of heterogeneous melts on a caldera-wide scale prior to the Kilgore Tuff eruption (Watts et al. 2011).

In comparison, most CPM rhyolites are homogeneous in δ^{18} O even on the single crystal scale, with average zircon values that fall within a narrow low- δ^{18} O range of 2.7-2.8% (Figs. 4, 5a). However, the oldest CPM rhyolites, the South Biscuit Basin and Scaup Lake flows, exhibit much greater zircon heterogeneity (Fig. 5a). We propose that the homogenization through time effect that we observe for CPM rhyolites (Fig. 5a), whereby they become progressively more homogeneous and retain fewer inherited crystals in younger eruptions, is an expected outcome for caldera cluster evolution: heterogeneities of the intracaldera block are progressively annealed by successive episodes of bulk remelting. While at Yellowstone, these late-stage magmas erupted as sequential "snapshots" over the past 260 ky, the Kilgore Tuff erupted in a single, climactic pulse that tapped an assembled, but not fully annealed, magma chamber.

Concluding remarks

The results of the present study provide high-resolution, crystal scale insights into magma genesis under one of the world's largest supervolcanoes. This study builds on previous studies of Yellowstone and pre-Yellowstone supereruptions in the Yellowstone hotspot track that reveal diverse paths of large-volume rhyolite genesis. In some cases, magma accumulation leads to the formation of a large isotopically and chemically homogeneous magma reservoir, with few zircon xenocrysts of diverse δ^{18} O compositions (e.g., Yellowstone's MFT and LCT), while in other cases, significant zircon δ^{18} O heterogeneity persists even after segregation of >1,800 km³ of magma, as in the case of Heise's Kilgore Tuff.

Our study of voluminous CPM lavas reveals the anatomy of a failed supereruption. With their young ages that permit detailed U-Th zircon geochronology, CPM lavas can be used to gain insight into crystal scale records of large silicic magmatic systems. Based on the results of our



¹ This calculation is based on 1.5 kJ/kg K heat capacity of basalt, cooling from 1,250°C liquidus to 550°C ambient temperature, with 400 kJ/kg latent heat of crystallization, assuming it is rapidly quenched and 100% crystalline (diabase) with a 40% heat transfer efficiency (Dufek and Bergantz 2005), yielding 580 kJ/kg. To melt a rhyolitic rock by 50%, heating from 550°C ambient temperature to 850°C liquidus, with 0.7 kJ/kg K heat capacity and 300 kJ/kg latent heat of fusion would require 360 kJ/kg. Thus, under the specified conditions, basalt can melt 1–2 times the volume of rhyolite.

present study, it appears that progressive stages of $>600 \text{ km}^3$ of rhyolitic magma accumulation occurred from ~ 260 to 155 ka beneath the Yellowstone caldera. This time was sufficiently long to form a homogenous and voluminous magma reservoir that began cooling and erupting large-volume rhyolitic lavas. Annealing of the preexisting crystal cargo, including diverse zircon, quartz, and sanidine crystals, proceeded until all age and isotopic evidence of inheritance was lost by the time of the most evolved and coolest rhyolite eruption ~ 75 ka.

Given that voluminous CPM lavas are low- δ^{18} O rhyolites with isotopic and age evidence for a genetic connection to buried intracaldera volcanic and plutonic rocks from previous eruptions, we speculate that the evolution of caldera volcanism from the Yellowstone caldera block is complete. The remelting of rocks from the uppermost, fusible parts of crust signifies a terminal stage of caldera cluster evolution. These final eruptions provide crystal scale records of a dying magmatic cycle rather than a newly formed or rejuvenated magma chamber. If we can use Heise as a guide, where about 2.5 million years elapsed between the final Kilgore caldera cycle and the onset of eruptions at Yellowstone, more than 2 million years of heating, melting, assembling, and distilling thousands of km³ of high-silica rhyolite are necessary for renewed supereruptions in the hotspot track. Our in-depth study of isotopic and age variations in single crystals may shed light on similar processes occurring in large caldera centers elsewhere on Earth, and perhaps provide a viable approach for their investigation.

Acknowledgments We thank Jorge Vazquez for performing U–Pb dating of Biscuit Basin zircons at the USGS-Stanford SHRIMP Lab, Guillaume Girard for advice on Yellowstone sampling localities, Luke Sitts for assistance during analytical sessions at the UCLA SIMS facility, and Rachel Weber for fieldwork assistance. Guillaume Girard and two anonymous reviewers are thanked for their constructive feedback on the manuscript. This work was supported by a NSF grant EAR/CAREER-844772 to I.N. Bindeman, a NSF grant IF-0732691 to the UCLA SIMS facility, and a University of Oregon Staples grant to K.E. Watts.

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