TephraKam: geochemical database of glass compositions in tephra and welded tuffs from the Kamchatka volcanic arc (northwestern Pacific)

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Abstract. Tephra layers produced by volcanic eruptions are widely used for correlation and dating of various deposits and landforms, for synchronization of disparate paleoenvironmental archives, and for reconstruction of magma origin. Here we present our original database TephraKam, which includes chemical compositions of volcanic glass in tephra and welded tuffs from the Kamchatka volcanic arc. The database contains 7049 single-shard major element analyses obtained by electron microprobe and 738 trace element analyses obtained by laser ablation inductively coupled plasma mass spectrometry on 487 samples collected in close proximity to their volcanic sources in all volcanic zones in Kamchatka. The samples characterize about 300 explosive eruptions, which occurred in Kamchatka from the Miocene up to recent times. Precise or estimated ages for all samples are based on published 39Ar/40Ar dates of rocks and 14C dates of host sediments, statistical age modeling and geologic relationships with dated units. All data in TephraKam are supported by information about source volcanoes and analytical details. Using the data, we present an overview of geochemical variations in Kamchatka volcanic glasses and discuss applications of these data for precise identification of tephra layers, their source volcanoes, and temporal and spatial geochemical variations in pyroclastic rocks in Kamchatka. The data files described in this paper are available on ResearchGate at https://doi.org/10.13140/RG.2.2.23627.13606 (Portnyagin et al., 2019).

1 Introduction

Tephra layers are widely used for correlation and dating of various deposits and landforms, for the synchronization of disparate paleoenvironmental archives, and for reconstruction of magma origin and temporal evolution. These applications are in high demand in paleoclimatology, paleoseismology, archaeology, and other Quaternary science disciplines (e.g., Lowe, 2011), as well as in petrology and geochemistry (e.g., Cashman and Edmonds, 2019; Ponomareva et al., 2015a; Straub et al., 2015). Tephra is composed of minerals, volcanic glass (melt rapidly quenched upon eruption), and rock fragments in different proportions. A major modern approach for correlation of tephra layers between different locations is using major and trace element composition of volcanic glass (e.g., Cashman and Edmonds, 2019; Lowe, 2011; Ponomareva et al., 2015a). The composition of volcanic glasses has been shown to vary significantly on spatial scales ranging from volcanic region to a single volcano, reflecting a large variability of thermodynamic conditions of magma storage and fractionation and the composition of crustal and mantle sources of magmas (e.g., Bachmann and...
Tephra often dominates the erupted products in terms of volume, eruption frequency, and variety of compositions, some of which may never occur in lava. It is particularly true for highly explosive volcanic arcs where the vast majority of the magma is erupted as tephra (e.g., Kutterolf et al., 2008). Therefore, tephra studies have a large, still only partly explored, potential to trace temporal and spatial variations in magma compositions in volcanic arcs (e.g., Clift et al., 2005; Kimura et al., 2015; Straub et al., 2004, 2015).

The Kamchatka Peninsula (Fig. 1) hosts more than 30 recently active large volcanic centers and a few hundred monogenetic vents, comprising the northwestern segment of the Pacific Ring of Fire. Kamchatka volcanism is highly explosive. According to some estimates, Kamchatka has the largest number of Quaternary calderas per unit of arc length in the world (Hughes and Mahood, 2008). Kamchatka tephra layers provide chronological control for deposits and events over large areas, both in Kamchatka and farther afield, up to Greenland and North America, which is critical for many studies (e.g., Cook et al., 2018; Hulse et al., 2011; Kozhurin et al., 2014; Mackay et al., 2016; Pembeda et al., 2016; Pinegina et al., 2013, 2014, 2012; Plunkett et al., 2015; van der Bilt et al., 2017). However, geochemical characterization of Kamchatka volcanic glasses is still in a developing phase. In the Kamchatka volcanic arc, the Holocene tephrochronological framework (until recently) has been based mainly on direct tracing of tephra layers, bulk composition of tephra, and bracketing radiocarbon dates (e.g., Bazanova et al., 2005; Braitseva et al., 1998, 1996, 1995, 1997; Pevzner, 2010; Pevzner et al., 1998, 2006). Significant progress towards creating geochemical database of Kamchatka tephras has been achieved in the past 10 years (Dirksen et al., 2011; Kyle et al., 2011; Plunkett et al., 2015; Ponomareva et al., 2013a, b, 2017, 2015b). However, the published geochemical data are mostly restricted to the Holocene and do not include data on trace element composition of volcanic glasses.

In this paper, we present TephraKam – our original, internally consistent and, so far, most complete database of single-shard glass composition from tephras and welded tuffs of Kamchatka volcanoes, covering the period from the Miocene until the present (Portnyagin et al., 2019). The data have been collected during the past 10 years and include major element compositions obtained by electron microprobe and trace element compositions of representative samples by laser ablation inductively coupled plasma mass spectrometry (LA-ICP-MS). Ages based on published radiocarbon and $^{39}$Ar/$^{40}$Ar dates, as well as on the age models and stratigraphy, are provided for all samples. Using these data, we present an overview of geochemical variations in Kamchatka volcanic glasses and suggest some key geochemical parameters and diagrams, which permit precise identification of tephra layers and their sources and assessment of regional geochemical variations in Kamchatka.
2 Volcanoes of Kamchatka and studied samples

The Kamchatka Peninsula overlies the northwestern margin of the subducting Pacific plate and is one of the most volcanically and tectonically active regions in the world (e.g., Gorbatov et al., 1997). Kamchatka hosts more than 30 large active volcanoes, 40 calderas, and hundreds of monogenetic vents grouped into two major volcanic belts running northeast–southwest along the peninsula, which are the Eastern Volcanic Belt and the Sredinny Range (SR) (Fig. 1). The Eastern Volcanic Belt includes the volcanic front (VF) and rear arc (RA) in the southern (51–53° N) and central (53–55° N) segments, and the volcanic zone of the Central Kamchatka Depression (CKD) in the northern segment (55–57° N). Definition of VF and RA volcanoes varies in published studies. In this work, VF volcanoes are defined as those located at the closest distance to the deep-sea trench along the volcanic arc. RA volcanoes are located behind the frontal volcanoes. The current configuration of the volcanic belts is believed to have existed since ca. 2.5 Ma (Avdeiko et al., 2007; Lander and Shapiro, 2007; Legler, 1977; Volynets, 1994).

The products of the continuous explosive volcanism in Kamchatka during the last 2.5 Ma are not equally represented in the depositional record. Holocene tephra layers mantle the topography and, being interlayered with paleosol or peat horizons, form a sequence that provides a nearly continuous record of the Holocene explosive activity (e.g., Bazanova and Pevzner, 2001; Braitseva et al., 1998, 1996, 1995, 1997; Kyle et al., 2011; Pevzner, 2010; Pevzner et al., 1998; Ponomareva et al., 2015a, b, 2017). Earlier pre-Holocene pyroclastic products are mostly ignimbrite (pumiceous or welded tuffs), which survived through glacial stages better than loose pyroclastics and in many cases experienced alteration (Bindeman et al., 2019, 2010; Ponomareva et al., 2018; Seligman et al., 2014). These deposits are partly eroded by glacial processes, buried by younger deposits, and/or covered with dense vegetation, which hampers their identification.

TephraKam database provides data on volcanic glass composition from 65 volcanic centers in Kamchatka. Of these centers, 43 were active in the Holocene and the remaining 22 centers ceased their activity prior to the Holocene. Some volcanic centers are individual volcanic cones (e.g., Ilinsky), calderas (e.g., Kurile Lake caldera), monogenetic lava fields (e.g., Tolbachik lava field), or monogenetic vents (cinder cones and craters), while other centers combine several volcanoes and/or calderas (e.g., Karymsky center). The latter approach was used in cases when thick local pyroclastic deposits could not be unambiguously assigned to a certain volcano within the volcanic cluster. We define the source vent within the volcanic center where possible (e.g., Karymsky/Polovinka caldera). Eight ignimbrite units come from unknown sources so we use coordinates of their samples instead of vent coordinates.

We have analyzed glass from 487 samples, including 11 replicate samples marked as "--rep" in the database. Overall, our samples characterize about 300 individual explosive eruptions. A total of 298 samples come from tephra fall deposits, 187 are from ignimbrite units (42 of them welded), and 2 are from lava. Our sampling covers all the Quaternary volcanic belts: 25% of the samples are from VF, 27% are from RA, 40% are from CKD, and 8% are from the Sredinny Range. The coverage among the volcanic centers is not uniform: some volcanoes are characterized by only one sample while others are densely sampled and analyzed. The sampling density partly reflects the amount of large explosive eruptions from a certain volcano. The analyzed samples span an age interval from the Miocene (ca. 6 Ma) to recent times (Fig. 2). About 60% of samples and data presented in this database are from the Holocene and characterize all large explosive eruptions in Kamchatka during this time (Braitseva et al., 1995, 1997; Ponomareva et al., 2013b, 2015b), as well as many moderate-size eruptions. Ages, tephra dispersal areas, and volumes for most of the Holocene eruptions have already been published (Braitseva et al., 1998, 1996, 1997; Kyle et al., 2011; Ponomareva et al., 2017; Zaretskaya et al., 2007). The pre-Holocene record of explosive eruptions is spotty, and its representativeness decreases with increasing rock age (Fig. 2). Most of the pre-Holocene samples characterize ignimbrites associated with large (diameter 5–40 km) collapsed calderas. A total of 69 of the ~200 Pleistocene samples characterize 38 eruptions dated with the help of radiocarbon, 39Ar/40Ar, or age modeling while the rest of the samples provide information about earlier unreported eruptions. Analyzed samples were collected between 1975 and 2016 by 22 contributors; the largest collections come from the authors of this paper.

3 Methods and database structure

3.1 Sample preparation

Tephra samples have been prepared using our original technique, developed in the past 20 years at GEOMAR (Kiel) and the Vernadsky Institute (Moscow). The technique aims for uncomplicated, time- and material-effective preparation of many tephra samples for microanalytical work. The technique applies no strong heating (>50°C) and no acid leaching, which may cause chemical modification of tephra glasses (e.g., Hunt and Hill, 1996). All material used for preparation were thoroughly tested to exclude those causing chemical contamination of polished glass surface.

The samples have been cleaned in water to wash out clay and the finest (<5 µm) fraction and then dried. Fine- and medium-grained ash has been mounted without splitting into fractions and additional crushing. Lapilli and welded tuffs have been carefully crushed in a hand mortar and then mounted. The samples were mounted in 25 mm diameter, 4 mm thick Plexiglas holders with 12 and 16, respectively,
3.2 Electron probe microanalysis (EPMA)

In this study, EPMA data have been obtained following published recommendations for the analytical conditions, primary and secondary reference materials, the number of analyses, and other factors and procedures, which may influence the quality of EPMA data for tephra glasses and their interpretation (Froggatt, 1992; Morgan and London, 1996, 2005; Hunt and Hill, 2001; Turney et al., 2004; Kuehn et al., 2011; Lowe et al., 2017). A particularly influential study was an intercomparison of electron microprobe data for volcanic glasses between different labs (Kuehn et al., 2011), which confirmed a high quality of our data (GEOMAR lab is no. 12 in Kuehn et al., 2011) and allowed us to further improve our EPMA protocol.

The glasses were analyzed at GEOMAR (Kiel, Germany) using a JEOL JXA 8200 electron microprobe equipped with five wavelength dispersive spectrometers, including three high-sensitivity instruments (two PETH and a TAPH). The analytical conditions were 15 kV accelerating voltage, 6 nA current, and 5 µm electron beam size for all analyses. The current and size conditions correspond to the current density of 0.076 nA µm$^{-2}$, which is within the recommended range ($< 0.1$ nA µm$^{-2}$) to minimize the Na loss during analysis, especially in a combination with short counting time for Na (Morgan and London, 1996, 2005; Kuehn et al., 2011). Counting times in the latest version of the program are 5 and 10 s (peak and background, respectively) for Na; 20 and 10 s for Si, Al, Mg, Ca, and P; 30 and 15 s for Fe, K, Ti Cl, and S; 40 and 20 s for F; and 60 and 20 s for Mn. The counting times have been optimized several times in the course of this study. The changes are reported in TephraKam Table 1a. The changes have not affected the data accuracy for most elements, but the precision of single-point analyses has improved. Decreasing counting time for Na from 20 to 5 s in 2010, which was aimed at minimizing loss of this element during analysis, resulted in analytical precision that was about 2 times lower for single-point Na analyses (Morgan and London, 1996), but the accuracy of Na analysis was improved and became less dependent on the reference material for standardization.

Basaltic glass (USNM 113498/1 VG-A99) for Ti, Fe, Mg, Ca, and P; rhyolitic glass (USNM 72854 VG568) for Si, Al, Na, and K; scapolite (USNM R6600-1) for S and Cl (all from the Smithsonian collection of natural reference materials; Jarosewich et al., 1980); comendite obsidian KN-18 (Nielsen and Sigurdsson, 1981; Mosbah et al., 1991) for F; and synthetic rhodonite for Mn were used for calibration and monitoring of routine measurements. Two to three analyses of all standard glasses and scapolite were performed at the beginning of analytical session, after every 50–60 analyses and at the end. The data reduction included online CITZAF correction and small drift correction for systematic deviations (if any) from the reference values obtained on standard materials. The latter correction has not exceeded a few relative
percent for all elements and allowed us to achieve the best possible accuracy and precision. The correction resulted in a very minor change of the mean concentrations but allowed 20–40% improvement of the analytical precision (2 SD; SD is sample standard deviation) and the shape of data distribution, making it closer to Gaussian distribution (see the Supplement to Ponomareva et al., 2017).

The glass analyses used in this study were obtained in the period from 2009 to 2019. The summary of data for reference materials collected over this period of time is presented in TephraKam Table 1b (Portnyagin et al., 2019). The data include results obtained on major reference glasses and minerals, which were used in calibration of quantitative microprobe measurements (USNM 72854 VG568, USNM 113498/1 VG-A99, USNM R6600-1, KN18), and also results obtained for other reference materials analyzed as unknown in the course of our study. The latter include natural glasses USNM 111240/52 VG-2 (Jarosewich et al., 1980); Lipari obsidian (Hunt and Hill, 1996); Mt. Ediza Sheep Track tephra; Laki 1783 AD tephra; Old Crow tephra (Kuehn et al., 2011); and glasses made of natural rock powders ATHO-G, BM90/21-G, GOR128-G, KL2-G, StHslo60/8-G, ML3B-G (Jochum et al., 2006), and artificial glass NIST SRM 612 (Jochum et al., 2011). The data demonstrate remarkable agreement with recommended concentrations for all elements and thus excellent accuracy of our data, which reproduces reference concentrations within the reported 2 SD in nearly all cases. The latter is also true for concentrations, which significantly exceed the concentrations in reference glasses used for calibration. This is illustrated, for example, by analyses of Na$_2$O in NIST SRM 612 glass (13.70±0.30 wt % recommended vs. 13.73±0.40 wt % measured) and MgO in GOR128-G glass (26.00±0.30 wt % recommended vs. 25.66±0.68 wt % measured).

Precision of single-point analyses depends on the element concentration and analytical conditions for every element. Assuming that the reference materials used in this study were perfectly homogeneous (which may be not true for natural glasses containing microlites of minerals), the precision of single-point analysis of typical rhyolite can be assessed from 2 SD of the long-term mean concentrations obtained for glass USNM 72854 VG568 or Lipari obsidian. Precision of single-point basaltic glass analysis can be evaluated from the data on glasses USNM 113498/1 VG-A99 or USNM 111240/52 VG-2. For more precise determination of a single-point analytical precision, we provide TephraKam Table 1c (Portnyagin et al., 2019), where the precision for every element is calculated based on element concentration in glass, taking into account long-term reproducibility of reference materials. Correlation of the oxide concentrations in reference materials plotted against long-term relative standard deviation (2 RSD, %) allows for estimating analytical detection limits and finding element concentrations at which 2 RSD approaches 100% (TephraKam Table 1c).

During the subsequent data reduction, we excluded analyses with the totals lower than 90 wt %, which resulted from possible unevenness of sample surface, entrapment of voids, or epoxy during analysis of very small glass fragments. The latter has been also identified by unusually high measured chlorine concentrations, resulting from entrapment of epoxy resin during analysis (see Sect. 3.1). Analyses contaminated by occasional entrapment of crystal phases, usually microlites of plagioclase, pyroxene, or Fe-Ti oxides, were mostly identified and excluded on the basis of excessive concentrations of Al$_2$O$_3$, CaO, or FeO (and TiO$_2$), respectively, compared to the prevailing composition of glasses in every sample. Because volcanic glasses can be hydrated over time during post-magmatic interaction with meteoric water or seawater or contain significant but variable amounts of H$_2$O that were not completely degassed during the eruption, all analyses were then normalized to an anhydrous basis, accounting for the presence of halogens (Cl, F) substituting for O.

### 3.3 Laser ablation inductively coupled plasma mass spectrometry (LA-ICP-MS)

In the past 25 years LA-ICP-MS became a common technique to quantify concentrations of a wide range of trace elements in tephra glasses (e.g., Westgate et al., 1994; Pearce et al., 1996, 2007, 2014; Tomlinson et al., 2010; Kimura and Chang, 2012; Maruyama et al., 2016; Lowe et al., 2017). The LA-ICP-MS technique adopted for this study and its development were based on the principal results and recommendations from the previous works and our own experimental results. All the trace element analyses were obtained at the Institute of Geosciences, CAU Kiel, Germany. Conditions of analysis are summarized in TephraKam Table 1d (Portnyagin et al., 2019).

Before 2017, analyses were performed using a quadrupole-based ICP-MS (Agilent 7500s) and a Coherent GeoLas ArF 193 nm Excimer LA system. In situ microsampling was done with 24–50 μm pit size and 10 Hz pulse frequency at 5–10 J cm$^{-2}$ fluence. Analyses were performed using a large volume ablation cell (ETH Zürich, Switzerland) (Fricker et al., 2011). The generated aerosol was transported with 0.75 L min$^{-1}$ He and mixed with 0.6 L min$^{-1}$ Ar prior to introduction into the ICP. The ICP-MS was operated under standard conditions at 1500 W and optimized for low oxide formation (typically Th$_3$O$_7$ ≤ 0.4 %) and robust plasma (U/Th ≈ 1). Ca, Ti, Si, and 30 trace elements were analyzed. The calibration was based on NIST SRM 612 glass standard (Jochum et al., 2011) and matrix-corrected using ATHO-G and KL2-G glasses (Jochum et al., 2006). The measured intensities were converted to element concentrations using a conventional approach (Longerich et al., 1996; Pearce et al., 1996), with $^{43}$Ca as the internal standard and the anhydrous normalized CaO from EPMA data. Initial data reduction was performed in Glitter software (Griffin et al., 2008) setting integration...
windows of time-resolved data for background (20 s) and variable times depending on signal duration from individual tephra shards. Si and Ti concentrations obtained by LA-ICP-MS were compared to microprobe data, and the LA-ICP-MS analyses that deviate from EPMA data for these elements by more than 20% relative were rejected. The threshold value of 20% corresponds to ~3 RSD obtained for Si and Ti on reference glasses using 24 µm pit size (TephraKam Table 1e).

Beginning from January 2017, the analyses were obtained using a new ICP-MS Agilent 7900s and a Coherent GeoLas ArF 193 nm Excimer LA system operated with a fluence of 5 J cm⁻², at a repetition rate of 10 Hz and a 15–24 µm ablation craters. Analyses were performed using the same large volume ablation cell as before (ETH Zürich, Switzerland) but modified for fast washout. Helium (0.7 L min⁻¹) with addition of 14 mL min⁻¹ H₂ was used as carrier gas. The carrier gas was mixed with Ar (~1 L min⁻¹) prior to introduction to the ICP-MS. A total of 10 major elements (Si, Ti, Al, Fe, Mn, Mg, Ca, Na, K, P) and 31 trace elements were analyzed. Analyses included 20 s background (laser off) and 30 s signal (laser on) measurements. Dwell time for different elements varied from 5 to 20 ms depending on their abundance. One complete measurement cycle lasted 0.607 ms, and initial data reduction was performed in Glitter software (Griffin et al., 2008), which included manual selection of integration windows and preliminary calibration. The typical integration intervals for tiny tephra shards were 6–10 s and included 10–17 cycles. The intensities corrected for background and averaged over the selected intervals were normalized to the intensity of ⁴³Ca isotope and converted to concentrations by matching the sum of major element oxides to 100 wt % (Liu et al., 2008; Pettke et al., 2004). The calibration and correction of instrumental drift used data on ATHO-G reference glass, which was measured in duplicate after every 18 points on unknown samples. The reference concentrations of all elements, except Na, in ATHO-G were used following Jochum et al. (2006). Na concentrations was accepted to be 4.1 wt %, ca. 10% higher, relative to those reported by Jochum et al. (2006) to reproduce Na in other reference glasses analyzed by LA-ICP-MS in this study (TephraKam Table 1d, e) and better comply with Na obtained by electron microprobe (TephraKam Table 1b). Sc concentrations were corrected for SiO⁺⁺ interference using reference glasses with known Sc and variable SiO²⁻ content. The data were further filtered for inclusion of phenocryst phases by comparison of major element concentrations with those obtained by EPMA, and obvious outliers were rejected to leave only glass analyses.

During all periods of data collection from 2011 to 2019, BCR2-G, KL2-G, and STHS60/8-G glasses (Jochum et al., 2006) were analyzed as unknown in one series with the samples (TephraKam Table 1e). The data confirms good consistency of the entire dataset and no bias related to periodic instrumental upgrades. Based on these data, the analytical precision and accuracy are typically between ±2% and 8% for 20 s long analyses, but the precision might be reduced for very short analyses of tiny glass shards and for elements occurring at concentrations below 0.1 ppm.

Overall, the data obtained since 2017 using the very sensitive modern instrument Agilent 7900 and after implementation of additional improvements (modified cell, addition of H₂ in carrier gas) are more accurate compared to earlier data for the same spot size of 24 µm. Older data obtained with 50 µm spot have comparable precision to the most recent data. However, some former data can be affected by entrapment of crystal phases (e.g., Pearce et al., 2014) that was impossible to identify by only using data on Ti and Si concentrations. Thus, outliers in pre-2017 data should be considered with care. Besides a smaller laser beam, the post-2017 data were quantified using a more efficient approach by normalizing oxides to 100%. These data are directly comparable with EPMA data for all elements except volatiles F, Cl, and S. Thus, contamination of these analyses by occasional entrapment of crystal phases is excluded. The recent LA-ICP-MS data also provide accurate concentrations of Ti, Mn, and P, occurring in silicic glasses in concentrations approaching and below the detection limit of our EMP analyses (0.02 wt%–0.03 wt % for these elements).

3.4 Tephra ages

Knowledge of tephra ages (or at least approximate age ranges) is crucial for their use as marker horizons. For many tephras in our database, the age estimates are available from published data (Auer et al., 2009; Bazanova and Pevzner, 2001; Bazanova et al., 2019; Bindeman et al., 2019, 2010; Braitsvea et al., 1998, 1991, 1995; Churikova et al., 2015; Cook et al., 2018; Dirksen, 2009; Dirksen and Bazanova, 2009; Dirksen and Melekestsev, 1999; Florensky, 1984; Kozhurin et al., 2006; Masurenkov, 1980; Melekestsev et al., 1992, 1995; Pevzner, 2015; Plechova et al., 2011; Ponomareva et al., 2018, 2013b, 2017, 2015b, 2006; Ponomareva, 1990; Seligman et al., 2014; Volynets et al., 1998; Zaretskaia et al., 2001; Zaretskaya et al., 2007; Zelenin et al., 2020).

In this case, we report bibliographical references and details on the age estimates and dating techniques. The majority of previously reported tephra ages were obtained by radiocarbon dating of host sediments. The ages are usually published as uncalibrated ¹⁴C dates. In the TephraKam database (Portnyagin et al., 2019), the published ¹⁴C dates have been recalculated to calibrated ages before present (cal yr BP) with 95% error interval using the most recent IntCal13 calibration curve (Reimer et al., 2013). Some calibrated ages are based on poorly documented ¹⁴C dates and reported as approximate ages (~ symbol) or as an age range. Holocene tephras in northern Kamchatka, as well as some Holocene marker tephras, were dated with the help of Bayesian age model combining 223 individual ¹⁴C dates (Ponomareva et al., 2017). One Holocene tephra (KHG from Khangar vol-
A total of 21 welded tuff units were dated by 39Ar/40Ar (Bindeman et al., 2019, 2010; Seligman et al., 2014). For undated samples, the age estimates were derived from their stratigraphic relationships with the dated ones. Designated age group is provided for all samples according to the geologic timescale.

3.5 Database structure

The TephraKam database is provided in Excel 2016 file (.xlsx) and consists of six folders (TephraKam Tables 2a–f, Portnyagin et al., 2019): (a) comments, (b) volcanoes, (c) sample description, (d) major elements, (e) trace elements, and (f) discrimination diagrams. Table 2a, comments, explains abbreviations of columns in the data tables. Table 2b, volcanoes, contains information about volcanic centers of Kamchatka, from which volcanic glass data exist and are presented in the database. Table 2c, sample description, includes coordinates, information of sample age, outcrop, type of material (ash, pumice, and welded tuff), collector’s name, and other information including data for source volcano via link to Table 2b. Table 2d, major elements, contains EPMA data on individual glass shards from samples studied and related information. Table 2e, trace elements, contains LA-ICP-MS major and trace element data on single glass shards, information about the dates and conditions of LA-ICP-MS analysis, trace element concentrations normalized to mantle composition, and some element ratios for plotting the data. The tables are linked to each other so that any changes in volcano or sample description will be seen in geochemical data tables. Table 1f, discrimination diagrams, contains sample plots and coordinates of corner points to draw compositional fields of the modern volcanic zones in Kamchatka using coordinates Nb/Y vs. La/Y and Nb/Y vs. Th/Y.

4 Data overview

4.1 Spatial and temporal variations in volcanic glass compositions

Major element data are available for all samples and are comprised of 7049 individual analyses. Trace elements are available for 114 samples and include 738 individual analyses. About 30% of the major element data have already been published, e.g., for the Shiveluch eruptions (Ponomareva et al., 2015b); the Ushkovsky eruptions (Ponomareva et al., 2013b); and a part of the Bezymianny and Tolbachik eruptions (Ponomareva et al., 2017). The majority of the trace element data are presented here for the first time.

An overview of the available major element data is shown in Fig. 3, a common classification diagram for island arc rocks in coordinates SiO2 vs. K2O (Gill, 1981; Le Maitre et al., 2002; Peccerillo and Taylor, 1976). In this diagram, lines dividing low-K2O, medium-K2O, and high-K2O compositions are drawn along typical trends of magma fractionation from basalts to rhyolites. Thus, this diagram is useful to access the extent of magma fractionation and relative enrichment in K2O of parental magma and/or source rock. Basaltic glasses are very rare in tephra from Kamchatka. The vast majority of glasses have basaltic andesite, andesite, dacite, and rhyolite low- to high-K2O compositions. The compositions are not uniformly distributed in SiO2-K2O coordinates. Some compositions are more common; the others are rather rare. For example, dacite and rhyolite tephra glasses with K2O ∼ 2 wt% and SiO2 of 68 wt%–72 wt%, very low-K2O rhyolites, and alkali rhyolites with K2O > 5% are extremely rare or unknown in Kamchatka. In turn, medium-K2O rhyolite glasses with K2O ∼ 3 wt% and SiO2 of ∼ 75 wt% are very common and characterize many eruptions from all volcanic zones.

The compositions of glasses are grouped in Fig. 3 according to their source volcano location in Kamchatka (Fig. 3a), age (Fig. 3b), type of volcano (Fig. 3c), and type of sample (Fig. 3d). Figure 3a–c show glasses taken only from tephra; Fig. 3d compares glasses taken from tephra and welded tuffs. Glasses from VF are represented by a full range of compositions from basaltic andesites to rhyolites and belong to low-K2O and medium-K2O series. RA glasses have medium- and high-K2O compositions and overlap only marginally with VF glasses. CKD glasses have similar range of compositions with RA glasses, although medium-K2O rhyolite glasses similar to VF glasses are abundant in CKD (e.g., Shiveluch volcano). SR glasses exhibit compositional bimodality. Compositions with SiO2 from 65 wt% to 72 wt% are not known in the SR. The glasses have medium-K2O (some SR rhyolites) and predominantly high-K2O compositions. The glass compositions do not exhibit clear temporal variability, suggesting similar composition of erupted magmas since at least Middle Pleistocene (Fig. 3b). The compositions of tephra glasses from complex volcanoes and calderas are similarly variable and cover all compositional ranges. Tephra glasses from monogenetic volcanoes tend to have either the most mafic basaltic andesite compositions (basaltic cinder cones) or rhyolitic compositions (explosive craters). Some intermediate compositions are also known but are not as abundant as previously mentioned types (Fig. 3c). In addition to tephra glasses, the database includes glasses from welded tuffs and obsidian, which are important for characterizing the oldest explosive eruptions in Kamchatka. In comparison with tephra, welded tuff glasses tend to have SiO2-rich dacite and rhyolite compositions, although samples with some andesitic glasses are also present (Fig. 3d). Many welded tuffs have compositions with K2O > 5 wt%, which is higher than in the majority of tephra glasses. This K2O enrichment is most likely related to secondary alteration of glass, as discussed in more detail in Sect. 4.3.
Data on concentrations of trace elements in glasses add significant information, which are highly valuable for precise identification of volcanic sources, as well as for petrological and geochemical applications of this database. The data for Ti, Mn, and P obtained by LA-ICP-MS are generally of higher precision in comparison to EPMA data, particularly for glasses with concentrations of these elements below 500 ppm (0.05 wt %), approaching the detection limit of EPMA.

Trace elements provided in this database belong to different groups with contrasting geochemical properties in magmatic systems and therefore provide different geochemical information. Behavior of Sr, Ti, V, Sc, P, Zr, Hf, and heavy rare earth elements (heavy REEs: Eu, Gd, Tb, Dy, Ho, Er, Tm, Yb, Lu) is strongly controlled by solid crystalline phases. When plagioclase (Sr, Eu), Fe-Ti oxides (Ti, V), pyroxene (Sc), apatite (P), zircon (Zr, Hf), and amphibole (heavy REEs) crystallize from magmas, these elements behave as “compatible elements”, and their concentrations decrease in residual melts. In contrast, elements Rb, Ba, Th, Nb, Ta, Pb, and light REEs (La, Ce, Pr, Nd, Sm) behave as “incompatible elements” in most magmas of Kamchatka because they are not concentrated in solid phases and enriched in residual melt. Systematics of incompatible elements can be informative of the magma source composition and subduction-related parameters, such as, for example, the distance from volcano to subducting plate (e.g., Volynets et al., 1994; Churikova et al., 2001; Duggen et al., 2007). The ratios between incompatible elements do not change as magma fractionates and are instructive for identifying the source volcano of variably fractionated melts. This information is quite unique in comparison to the systematics of major elements, which, in contrast to incompatible trace elements is largely related to the conditions of magma storage and also to syneruptive crystallization and magma mixing (Cashman and Edmonds, 2019; Ponomareva et al., 2015a).

Although detailed evaluation of trace element systematics in Kamchatka glasses is beyond the scope of this work, in Fig. 4 we illustrate some regularity in trace element composition of tephra glasses from different volcanic zones in Kamchatka, which can help identify source volcano or at least volcanic zone for tephra of unknown provenance. In this diagram, we show only Holocene and Late Pleistocene samples as their source volcanoes are reliably constrained, and the available data are the most representative compared to older volcanic rocks.
Figure 4. Trace element variations in Holocene and late Pleistocene tephra glasses south to north (left) and across (right) the Kamchatka volcanic belts. Trace element ratios are normalized to primitive mantle values (McDonough and Sun, 1995). Nb/Y and La/Yb ratios reflect mantle source depletion and enrichment (e.g., Pearce et al., 1995) and also the extent of amphibole fractionation (e.g., Brophy, 2008). Ba/Th is a function of slab-derived fluid contribution to the source of magmas. Th/La is related to the amount of subducted sediments involved in magma generation (e.g., Elliott et al., 1997; Pearce et al., 1995; Plank, 2005).

(Nb/Y)_N and (La/Yb)_N ratios in glasses (N denotes values normalized to primitive mantle following McDonough and Sun, 1995) reflect source enrichment in highly incompatible elements (La, Nb) relative to less incompatible elements (Yb, Y) (Pearce et al., 1995, 1984). These ratios are also strongly influenced by amphibole crystallization in evolved magmas, which are an important host for heavy REE and Y but not for La and Nb (Brophy, 2008). Tephra glasses from frontal Kamchatka volcanoes have relatively low (Nb/Y)_N < 1.3 and (La/Yb)_N < 5. This is a distinctive compositional feature of VF tephra in comparison to glass compositions from the other volcanic zones in Kamchatka. RA glasses have (Nb/Y)_N = 1.1–3.6 and (La/Yb)_N = 3.1–10.9. Glasses from the SR tephra have even higher (Nb/Y)_N > 4.5 and (La/Yb)_N = 5.5–11.8, mostly overlapping with RA compositions. Both ratios increase with increasing distance from the deep-sea trench. CKD glasses have (Nb/Y)_N similar to those in RA glasses and (La/Yb)_N > 3 overlapping with RA and SR glasses.

(Ba/Th)_N and (Th/La)_N ratios are indicative of magma source composition and related, respectively, to contributions from slab fluid and sediment melts in source of magmas (e.g., Elliott et al., 1997; Pearce et al., 1995; Plank, 2005). Both ratios exhibit significant variations along the volcanic arc and range from mantle-like values ~1 up to 4–6 times higher than in primitive mantle. On a regional scale, VF tephras tend to have higher (Ba/Th)_N and lower (Th/La)_N than RA tephra: (Ba/Th)_N = 1.5–6.2 and 1.2–3.3, (Th/La)_N = 0.8–2.7 and 1.0–4.4 in VF and RA, respectively. At a given distance along the volcanic arc, VF tephras generally have higher (Ba/Th)_N and lower (Th/La)_N in comparison with RA tephras. SR tephra have a relatively low (Ba/Th)_N < 2 and high (Th/La)_N > 2 in the range of RA tephra. CKD tephra has ratios similar to VF tephra.
To sum up the overview of geochemical data, compositions of glass in Kamchatka tephra are very variable, enabling robust correlation of tephra layers as well as identification of source volcano and volcanic zone, from which unknown tephra could come from.

4.2 Using composition of glasses for fingerprinting ash layers in Kamchatka

TephraKam was initially created for tephrochronological needs to enable reliable identification and dating of tephra layers in Kamchatka and neighboring areas and for identification of their sources. The data have been used in a number of publications (Cook et al., 2018; Derkachev et al., 2019; Plunkett et al., 2015; Ponomareva et al., 2018, 2013a, b, 2017, 2015b; Zelenin et al., 2020). Our experience showed that ash layers produced by the largest explosive eruptions in Kamchatka can be recognized using major element systematics of tephra glasses. A diagram of SiO₂ vs. K₂O is useful for primary identification (Fig. 3) because it utilizes elements of contrasting geochemical properties, reflecting mantle or crustal source enrichment in incompatible elements (K₂O) and the extent of magma fractionation (SiO₂). In this respect, the diagram is more informative compared to other diagrams widely used in tephrochronology such as FeO vs. CaO or FeO vs. TiO₂, utilizing elements whose concentrations are largely controlled by crystallization processes and strongly correlate with each other. More detailed discrimination of ash layers requires additional geochemical constraints. For example, low-K₂O tephra glasses from Ksudach and Avachinsky volcanoes are well distinguished using CaO vs. SiO₂ systematics; medium-K₂O glass from Bezimianny volcano tephra has lower Na₂O compared to Shiveluch glasses; high-K₂O glass from Ushkovsky tephra has distinctly higher P₂O₅ compared to glass from high-K₂O basaltic tephra SH#28 from Shiveluch (Ponomareva et al., 2017).

In rare cases, tephra glasses from different volcanoes have hardly distinguishable major element composition. In this case minor elements determined by EPMA (P, Cl) and trace elements by LA-ICP-MS are useful to identify source volcanoes. In Fig. 5, we illustrate this case using compositions of tephra glasses from the Baranii Amphitheater Crater at the foot of Opala volcano (eruption OP 1356BP) and Khangar volcano (eruption KHG6600 7490BP). Although these tephra have very different ages, this comparison is instructive to illustrate the value of minor and trace element data to distinguish compositionally similar tephras. The difference in major elements is very subtle and mostly within long-term analytical uncertainty: Khangar glass has about 0.5 wt% lower Al₂O₃; ≤0.2 wt% higher CaO; ≤0.5 wt% higher K₂O at given SiO₂; and otherwise completely overlapping TiO₂, FeO, CaO, and Na₂O contents (Fig. 5a–c). The two tephra, however, have clearly different Cl content (Fig. 5d) and very different shapes of normalized trace element patterns (Fig. 5e), enabling clear discrimination between source volcanoes. Summarizing our experience to date, all tephra from different volcanic sources in Kamchatka have characteristic and unique chemical compositions.

Distinguishing tephra layers from the same volcano is a more difficult task. However, there is a number of examples from Kamchatkan volcanoes where tephra compositions are different even over short intervals of time (e.g., Kyle et al., 2011; Ponomareva et al., 2013b, 2017, 2015b). For example, Ponomareva et al. (2015b) showed that even compositionally similar tephra layers of the frequently erupting Shiveluch volcano can be distinguished using major element systematics in tephra glasses, particularly when the time period of eruption can be narrowed using marker layers from other volcanoes. The cases of compositionally identical products of different eruptions from the same volcanic center are also known. For example, Derkachev et al. (2020) reported two late Pleistocene layers produced by large eruptions from Gorely caldera, which have barely distinguishable major and trace element composition of glass. On a longer timescale of
100,000 years, the products of the Gorely caldera eruptions are more variable (Seligman et al., 2014), enabling their identification using glass composition in tephra and welded tuffs.

4.3 Effects of alteration on major and trace elements in glass from welded tuffs

TephraKam contains abundant data for glasses in welded tuffs of the Miocene to Pleistocene from different parts of Kamchatka. These rocks clearly represent products of large caldera-forming eruptions during the history of Kamchatka. Identification of their sources, age, and ash distribution is of great interest. However, some welded tuff glasses in the database have signs of secondary alteration that hamper their direct interpretation as compositions representative for native glass, i.e., quenched melt. Typically, the alteration results in characteristic “spaghetti”-like textures, precipitates of tiny magnetite crystals in glassy matrix followed by complete glass replacement by microcrystalline aggregate, and development of concentric perlite texture (Fig. 6). The process of devitrification is also associated with chemical modification of welded tuffs. Spot analyses usually reveal a large and correlated variability of alkalis and alkali earth elements within single samples, which typically is not observed in volcanic glasses from pumice fragments or ash layers. Representative trace element composition of variably altered glasses from the same unit of Karymsky/Stena-Sobolinaya caldera welded tuffs is shown in Fig. 7. A major feature of the glass alteration is enrichment in $K_2O$, $Rb$, and $Li$ (all are monovalent alkaline elements) that is inversely correlated with depletion in $Na_2O$, $CaO$, and $Sr$. Elements $Ba$, $U$, $Th$, $Pb$, $Nb$, $Ta$, $Zr$, $Hf$, $Ti$, $P$, and $Y$ and REEs exhibit small variability and are relatively immobile during alteration. Concentrations of the immobile elements are informative of the initial concentrations in glass and can be used for correlations between different ignimbrite units and with pristine tephra glasses. An example of geochemical fingerprinting of completely devitrified samples from Pauzhetka caldera in southern Kamchatka is provided by Ponomareva et al. (2018). These authors noted that elements $B$, $Ba$, $Eu$, and $V$ also reveal mobility in strongly altered tuffs and should be interpreted with caution.

4.4 Discrimination of glasses from different volcanic zones in Kamchatka using immobile trace elements

In comparison with major elements, concentrations of some trace elements in Kamchatka glasses are more variable and exhibit characteristic regional distribution (Fig. 4).
which should be related to systematically changing conditions of magma generation with increasing distance from volcano to the deep-sea trench and subducting plate. Glasses from the Central Kamchatka Depression volcanoes largely overlap with rear-arc glasses and partly overlap with volcanic front samples, which is in agreement with the geodynamic position of CKD (Fig. 1). Figure 8c and d show that glasses of the middle and early Pleistocene have compositions falling closely within the corresponding fields of different volcanic zones. Although Th/Y and La/Y are somewhat scattered at given Nb/Y, the volcanic zones can still be precisely identified in most cases. This consistency of compositions suggests that Kamchatka has not been affected by major tectonic reorganization, at least during Pleistocene and Holocene (ca. 2.5 Ma), and the present position of ancient volcanic centers corresponds closely to their initial position. Older Miocene–Pliocene samples are relatively rare in this database (Fig. 8e–f). However, we notice that glasses in Neogene rocks from the Sredinny Range have distinctive compositions that plot outside the range of Pleistocene–Holocene SR compositions and are more similar to modern rear arc and volcanic front samples. This might indicate that conditions of magma generation under Sredinny Range during the Pliocene were more similar to the present-day volcanic front and rear arc and that Sredinny Range volcanoes were located closer to the deep-sea trench and subducting plate at that time. This conclusion is in agreement with the proposed major tectonic reorganization in Kamchatka in the Neogene and shifting of volcanic front from the Sredinny Range to its present position in eastern Kamchatka (Avdeiko et al., 2007; Lander and Shapiro, 2007; Legler, 1977; Volynets, 1994). Thus, the diagrams in Fig. 8 can be successfully used to identify tephra erupted from different volcanic zones during the Holocene and Pleistocene. The proposed criteria are likely not valid for the Neogene, when the Kamchatka subduction zone had a different configuration. The diagrams can be particularly useful for identifying the provenance of distal tephras in marine sediments offshore of Kamchatka, for analysis of synchronicity of activity in different zones, and for analysis of temporal geochemical variability of volcanism.

5 Data availability

The archive zip file containing tables of this database is available on ResearchGate: https://doi.org/10.13140/RG.2.2.23627.13606 (Portnyagin et al., 2019). Updates will be available under the same web address or can be requested directly from the first two authors of this paper. A web-based version of this database will be released in the near future.
6 Conclusions

TephraKam is the largest and most comprehensive collection of internally consistent high-quality chemical analyses of major and trace elements in glasses of pyroclastic rocks of Kamchatka volcanoes. Precise or estimated ages are provided for every sample. Use of this database opens the possibility for reliable identification and correlation of tephra layers in Kamchatka and neighboring areas, enables dating of sedimentary archives onshore and offshore of Kamchatka and allows the multicomponent petrological and geochemical analysis of composition and origin of magmatic melts, preserved as quenched glass in tephra. The latter application is straightforward for rhyolite glasses, which have been shown to preserve the composition of magmas at depth (except for volatiles) and thus are informative of magma composition and its storage conditions at depth in Kamchatka (Ponomareva et al., 2015a). The amount of presented data is comparable and exceeds that available from published sources on the composition of volcanic rocks in Kamchatka.
chatka (e.g., GEOROC database). For silicic compositions, this database is a major source of information.

**Author contributions.** MVP and VVP designed the study and wrote the paper. MVP, VVP, LIB, MMP, and ANR conducted the fieldwork and collected most of the samples. MVP developed protocols of the EPM and LA-ICP-MS analyses and designed the database structure. MVP, VVP, AAP, and ANR produced the EPM and LA-ICP-MS data. EAZ compiled all the age data. DGS managed the LA-ICP-MS analyses. All authors discussed the results and participated in preparation of the paper.

**Competing interests.** The authors declare that they have no conflict of interest.

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