



TephraNZ: a major and trace element reference dataset for prominent Quaternary rhyolitic tephtras in New Zealand and implications for correlation

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Abstract

20 Although analyses of tephra-derived glass shards have been undertaken in New Zealand for
nearly four decades (pioneered by Paul Froggatt), our study is the first to systematically develop a
formal, comprehensive, open access, reference dataset of glass-shard compositions for New Zealand
tephras. These data will provide an important reference tool for future studies to identify and correlate
tephra deposits and for associated petrological and magma-related studies within New Zealand and
25 beyond.

Here we present the foundation dataset for “TephraNZ”, an open access reference dataset for
selected tephra deposits in New Zealand. Prominent, rhyolitic, tephra deposits from the Quaternary
were identified, with sample collection targeting original type sites or reference locations where the
tephra’s identification is unequivocally known based on independent dating or mineralogical
30 techniques. Glass shards were extracted from the tephra deposits and major and trace element
geochemical compositions were determined. We discuss in detail the data reduction process used to
obtain the results and propose that future studies follow a similar protocol in order to gain comparable
data. The dataset contains analyses of twenty-three proximal and twenty-seven distal tephra samples
characterising 45 eruptive episodes ranging from Kaharoa (636 ± 12 cal. yrs BP) to the Hikuroa Pumice
35 member (2.0 ± 0.6 Ma) from six or more caldera sources, most from the central Taupō Volcanic Zone.
We report 1385 major element analyses obtained by electron microprobe (EMPA), and 590 trace
element analyses obtained by laser ablation (LA)-ICP-MS, on individual glass shards.

Using PCA, Euclidean similarity coefficients, and geochemical investigation, we show that
chemical compositions of glass shards from individual eruptions are commonly distinguished by major
40 elements, especially CaO, TiO₂, K₂O, FeO, (Na₂O+ K₂O and SiO₂/K₂O), but not always. For those
tephras with similar glass major-element signatures, some can be distinguished using trace elements
(e.g. HFSEs: Zr, Hf, Nb; LILE: Ba, Rb; REE: Eu, Tm, Dy, Y, Tb, Gd, Er, Ho, Yb, Sm), and trace
element ratios (e.g. LILE/HFSE: Ba/Th, Ba/Zr, Rb/Zr; HFSE/HREE: Zr/Y, Zr/Yb, Hf/Y;
LREE/HREE: La/Yb, Ce/Yb).

45 Geochemistry alone cannot be used to distinguish between glass shards from the following
tephra groups: Taupō (Unit Y in the post-Ōruanui eruption sequence of Taupō volcano) and Waimihia
(Unit S); Poronui (Unit C) and Karapiti (Unit B); Rotorua and Rerewhakaaitu; and Kawakawa/Ōruanui,
Okaia, and Unit L (of the Mangaone subgroup eruption sequence). Other characteristics can be used to
separate and distinguish all of these otherwise-similar eruptives except Poronui and Karapiti.
50 Bimodality caused by K₂O variability is newly identified in Poihipi and Tahuna tephras. Using glass
shard compositions, tephra sourced from Taupō Volcanic Centre (TVC) and Mangakino Volcanic
Centre (MgVC) can be separated using bivariate plots of SiO₂/K₂O vs. Na₂O+K₂O. Glass shards from
tephras derived from Kapenga Volcanic Centre, Rotorua Volcanic Centre, and Whakamaru Volcanic
Centre have similar major- and trace-element chemical compositions to those from the MgVC, but can
55 overlap with glass analyses from tephras from Taupō and Okataina volcanic centres. Specific trace
elements and trace element ratios have lower variability than the heterogeneous major element and
bimodal signatures, making them easier to geochemically fingerprint.



1. Introduction

60 Tephrochronology is the principle by which volcanic ash (tephra) deposits are used as
stratigraphic isochronous marker horizons (isochrons) for correlating, dating, and synchronising
deposits and events in geologic, palaeoenvironmental, and archaeological records (Sarna-Wojcicki, 2000;
Shane, 2000, Dugmore et al., 2004; Lowe, 2011; Alloway et al., 2013). In regions where rates of
volcanism are high, and eruptive products are widespread, tephrochronology is an essential tool in many
65 aspects of geoscience and associated research. Geochemical fingerprinting of the glass shards within the
tephra deposits is one of the most common ways in which tephra are correlated. Traditionally, major
elements were used for correlations (e.g. Westgate and Gorton, 1981; Froggatt, 1983, 1992), but more
recent studies have included minor and trace element compositions as well (e.g. Westgate et al., 1994;
Pearce et al., 2002, 2004, 2007; Pearce, 2014; Knott et al., 2007; Allen et al., 2008; Denton and Pearce,
70 2008; Turney et al., 2008; Westgate et al., 2008; Kuehn et al., 2009; Hopkins et al., 2017; Lowe et al.
2017).

Trace elements are more strongly partitioned by fractional crystallisation processes that occur
during the formation of melts, and therefore have the potential to be unique for discrete eruption
episodes (e.g. Pearce et al., 2004). Specifically, a number of key trace elements have been identified as
75 important for the correlation of rhyolitic tephra, including the high field strength elements (HFSEs) Zr
and Nb; the large ion lithophile elements (LILE) Rb, Sr, and Ba; the heavy rare earth elements (HREE)
Gd, Yb, Sc, and Y; and the light rare earth elements (LREE) La and Nd. Also identified are important
trace element ratios, including: (1) HFSE/HREE – for example Zr/Y, Nb/Y, Hf/Y; (2) LILE/HFSE – for
example Ba/Th; (3) LREE/HFSE – for example Ce/Th, La/Nb; (4) LREE/HREE – for example La/Yb,
80 Ce/Yb; and (5) HFSE/HFSE – for example Zr/Nb, Zr/Th. Some studies have shown that trace elements
and trace element ratios can distinguish between tephra beds that have indistinguishable glass-shard
major element signatures and thus are a more robust way of providing accurate correlations (e.g.
Westgate et al., 1994; Pearce et al., 1996; 2002; 2004; Allan et al., 2008; Hopkins et al., 2017).

Tephra correlation is also increasingly being quantified through statistical approaches on
85 geochemical data (Lowe et al., 2017), but many of these approaches (supervised learning) often require
a robust (comprehensive) set of “known” reference data against which to test the analyses of
“unknown” samples. Statistics can also scale data to make them comparable, but they cannot account or
correct for inter-laboratory or historical variance in analyses. Therefore, incomplete datasets, or datasets
constructed from a range of data sources, will limit the ability to provide holistic statistical correlations
90 with accurate outputs. Therefore the formation of reference datasets that are run in one analytical
session, in one lab, with a consistent methodology are highly valuable but difficult to obtain. The
production of tephra databases is thus being recognised as an exceptionally useful tool internationally
(e.g. Lowe et al., 2017), made more obtainable with open access journals and online, effectively
limitless storage, leading to easier publication and maintenance of large data repositories. Ideally, a



95 global tephra database would exist, but at present this is beyond the scope and remit of any individual
researchers or institutes. Therefore regional databases for volcanically active and other regions are
becoming increasingly popular, such as TephraKam – Kamchatka (Portnyagin et al., 2019); TephraBase
– Europe (Newton, 1996); AntT tephra database – Antarctic ice cores (Kurbatov et al., 2014); Alaska
Tephra Database (Wallace, 2018); Klondyke Goldfields, Yukon (Preece et al., 2011); VOLCORE –
100 DSDP, ODP, and IODP marine tephra deposits (Mahony et al., 2020).

1.1. Geologic Setting

The volcanically active nature of New Zealand (Mortimer and Scott, 2020), and the longevity
and consistency of large-scale rhyolitic eruptions (Howorth, 1975; Froggatt and Lowe, 1990; Houghton
105 et al., 1995; Wilson et al., 1995a, 2009; Jurado-Chichay and Walker, 2000; Carter et al., 2003; Briggs et
al., 2005; Wilson and Rowland, 2016; Barker et al., 2021) mean the landscape currently has a very long,
detailed, and complex rhyolitic tephrostratigraphic framework that is used for a wide range of
applications. However, at present New Zealand tephra studies are lacking a comprehensive reference
dataset resource that has been developed in a systematic way.

110 The first large rhyolite-producing eruptions in the Quaternary in New Zealand were sourced
from the Coromandel Volcanic Zone (CVZ) (Carter et al., 2003; Briggs et al., 2005). At or after ~2 Ma,
volcanism moved into the Taupō Volcanic Zone (TVZ), currently the most active rhyolitic system on
Earth (Wilson et al., 1995a, 2009; Wilson and Rowland, 2016). Nine calderas are recognised within the
TVZ : Mangakino (1.6–1.53 Ma and 1.2–0.9 Ma); Kapenga (0.9–0.7 Ma, 0.3–0.2 Ma, and ~0.06 Ma);
115 Whakamaru (0.35–0.32 Ma); Reporoa (~0.23 Ma); Rotorua (~0.22 Ma); Ohakuri (~0.22 Ma); Maroa
(0.32–0.013 Ma); Taupō (0.32–0.0018 Ma); and Okataina (~0.6–0 Ma) (**Fig. 1B**; Houghton et al., 1995;
Wilson et al., 1995a, 2009; Gravely et al., 2006, 2007). The TVZ is further subdivided into the “old
TVZ”, which is defined as being active from inception to the Whakamaru eruptives (~0.34 Ma), and the
“young TVZ”, which is defined as being active from the Whakamaru eruptives to the present. “Modern
120 TVZ” is also used to describe the activity since the Rotoiti eruption (which includes the Rotoehu Ash
and Matahi Scoria members) ~45–47 ka (Danišik et al., 2012; Flude and Storey et al., 2016) to the
present (Wilson et al., 1995a, 2009). In addition to these rhyolitic caldera sources in the TVZ and CVZ,
the peralkaline rhyolitic Tuhua/Mayor Island (MI) volcano (**Fig. 1**), forming the Tuhua Volcanic Centre
(TuVC) (Froggatt and Lowe, 1990), is responsible for (amongst at least six other dispersed MI-derived
125 tephra; Shane et al., 2006) the Tuhua tephra (7637 ± 100 cal. yr BP; Lowe et al., 2019), a well-
recognised mid-Holocene rhyolitic marker horizon within the New Zealand geologic record due to its
distinctive peralkaline geochemistry and mineralogy (Buck et al., 1981; Hogg and McCraw 1983;
Froggatt and Lowe 1990; Wilson et al., 1995b; Lowe et al., 1999; Shane et al., 2006).



New Zealand sits in the path of predominantly westerly to southern-westerly winds, and
130 therefore the majority of tephra plumes are dispersed to the east of the volcanic zones (Barker et al.,
2019). However, tephra deposits from these rhyolitic eruptions are found in a range of different
environments, including:
(1) Marine (e.g. Nelson et al., 1985; Carter et al., 1995; Alloway et al., 2005; Allan et al., 2008; Lowe,
2014; Hopkins et al., 2020)
135 (2) Lacustrine (e.g. Lowe, 1988; Shane and Hoverd, 2002; Molloy et al., 2009; Shane et al., 2013;
Hopkins et al., 2015, 2017; Peti et al., 2020)
(3) Bog settings (e.g. Lowe, 1988; Newnham et al., 1995, 2007, 2019; Lowe et al., 1999, 2013; Gehrels
et al., 2006), or
(4) within terrestrially exposed (commonly marine or riverine) sediments, for example in the
140 - Whanganui Basin (e.g. Seward, 1976, Naish et al., 1996; Pillans et al., 2005; Rees et al., 2019),
- Wairarapa region (e.g. Shane and Froggatt, 1991; Shane et al., 1995; Nicol et al., 2002), or
- Hawke's Bay region (e.g. Erdman and Kelsey, 1992; Bland et al., 2007; Orpin et al., 2010;
Hopkins and Seward, 2019) (**Fig. 1**).

Because of their pervasive nature, high repose period, and high preservation potential, tephra
145 deposits are a common chronological aid in many studies in New Zealand. For example, the eruption of
Kaharoa (636 ± 12 cal. yr BP, Hogg et al., 2003) from Mt Tarawera in the Okataina Volcanic Centre
(OVC) has been used to date the arrival of Polynesians in northern New Zealand and map their
expansion and impact across the country (Newnham et al., 1998; Lowe and Newnham, 2004). The
Rerewhakaaitu eruption ($17,496 \pm 462$ cal. yr BP; Lowe et al., 2013), sourced from OVC, is used as a
150 marker horizon for the transition between the last glacial and present interglacial (Newnham et al.,
2003), and several other widespread late Quaternary tephra deposits form boundaries or key
stratigraphic markers in the New Zealand Climate Event Stratigraphy developed by the NZ-INTIMATE
community (e.g. Kawakawa/Oruanui tephra; Barrell et al., 2013; Lowe et al., 2013). Compositions of
glass and mineral components from rhyolitic tephra deposits have also been used to reconstruct changes
155 in magmatic systems, and give insight into the complexity of caldera-related eruption episodes (e.g.
Smith et al., 2002, 2005; Cooper et al., 2012; Barker et al., 2016, 2021; Wilson and Rowland, 2016).

Many of the commonly found rhyolitic tephra horizons in New Zealand are well studied, dated,
and geochemically and mineralogically characterised. However, often these studies have been eruption-,
source-, or depocentre-specific, and thus only provide a small, effectively piecemeal catalogue of tephra
160 geochemistry that is not necessarily comparable to those of other studies. In addition, data are not
usually published in their entirety, or not at all, meaning future studies cannot access nor use the data
for correlation techniques. Furthermore, Lowe et al. (1999) identified that differing procedural methods
employed at different institutes around New Zealand before and after 1995 produced variable elemental
concentrations for the same tephra (post-1995 SiO_2 values were lower by 0.5–1.0 wt.%, and all other



165 elements had slightly higher values). Therefore, it is likely that some of the older tephra compositions that have been relied upon in the past for correlative purposes are no longer appropriate.

It is therefore timely for a comprehensive, systematic, and accessible New Zealand tephra database to be initiated and developed. In this study we present “TephraNZ” as a foundation reference dataset of internally consistent, open access data for major and trace element compositions of glass shards from a selection of the most pervasive Quaternary tephra deposits in New Zealand (**Table 1**). This is by far the most complete dataset of New Zealand tephra-derived glass-shard compositions published to date. We discuss in detail the sample preparation, methods of analysis and data reduction processes used to obtain and interrogate the data, thereby providing a template for future studies to produce comparable datasets. Using the glass-shard data obtained, we present an overview of the geochemical variability for a range of rhyolitic tephtras of the TVZ, we suggest key geochemical parameters that can be used to identify the individual tephra layers, and apply common statistical principles to explore the data. Finally, we propose some future avenues of study, utilising these data, which would aid in the progression of a formal, holistic New Zealand tephrostratigraphical framework.

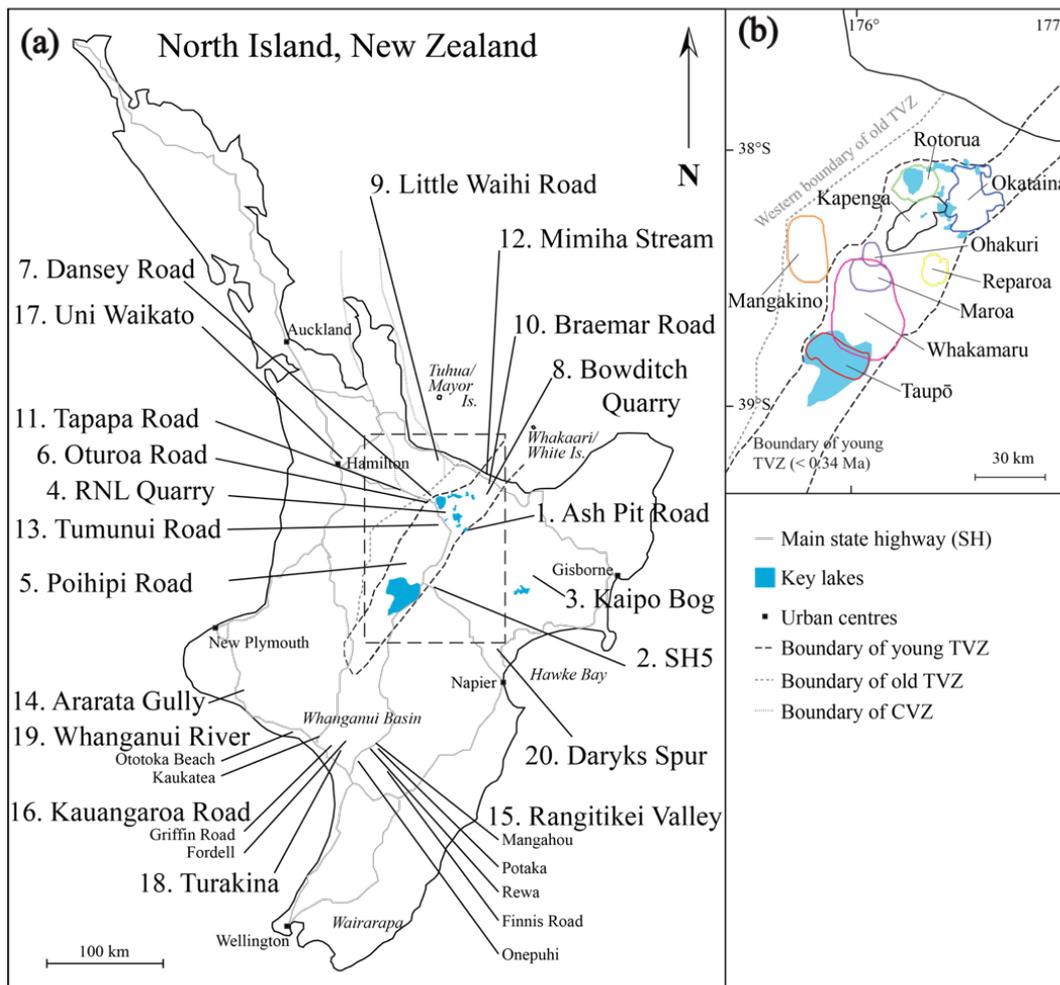
2. Methods

180 2.1. Sample collation and collection

Known tephra samples in personal collections were collated, prepared, and reanalysed for this study. Where samples were lacking for key tephra deposits, their type localities were found and samples obtained through field work (**Fig. 1**). **Table 1** provides full details of all the sample locations including their status as either proximal (0-10s km from source) or distal (10-100s km from source) and GPS coordinates for their exact sampling location. We note here that we have not attempted to sample multiple tephra beds from a single eruptive episode in proximal sequences, nor deposits of the same tephra at different azimuths, as has been undertaken in some more localised or petrologically-focussed studies (e.g. Shane et al., 2005, 2008). We recognise this limitation but instead have concentrated on analysing a wide range of pervasive rhyolitic tephtras, both proximal and distal, in a systematic and well-190 documented way so that future tephrostratigraphic studies will have a foundation of new, high-quality glass-shard compositional data for facilitating robust correlations and applications. Where we have both, we compare proximal and distal analyses of the same tephra and comment on similarities or differences. In addition, we have used statistical methods to demonstrate the integrity of our new datasets (and show how such methods can enable unknown tephtras to be classified).



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Figure 1. (a) Map of the North Island, New Zealand, detailing the samples sites where the reference tephra deposits for the TephraNZ database were collected. Outlines of CVZ (Coromandel Volcanic Zone) and TVZ (Taupō Volcanic Zone) are shown by dashed lines. Exact co-ordinates for all sample sites are detailed in Table 1. (b) Inset, outline shown in (a), the calderas of TVZ (details from Houghton et al., 1995; Wilson et al., 1995a, 2009; Gravely et al., 2006, 2007); outline colours of the calderas are used throughout this article in graphs to link the tephra data with their source caldera, if known.



2.2. Sample preparation

205 Bulk tephra samples were disaggregated in water for 1-5 min in an ultra-sonic water bath. Clays
and ultra-fines ($< 5 \mu\text{m}$) were rinsed off and samples were then wet sieved using disposable sieve cloths
to 125–250 μm shard size or, where necessary, 60–125 μm . Samples were then dried for 12–24 hr at 50°
C before mounting in epoxy resin. Seven samples were mounted into individual drill holes (4-mm
210 diameter) in 25 mm epoxy round blocks (a 4:1 ratio of EpoTek 301 resin [A]: hardener [B]). Individual
drill holes were then backfilled using the same epoxy mix (see Lowe, 2011, p. 124, for a schematic
illustration). Sample blocks were polished using the following sequence: ~ 3 min in a figure of eight
pattern on 800 grit paper with water lubricant to remove the epoxy and break through to the glass
shards, ~ 1 min on 1200 grit paper with water lubricant to remove any large scratches, and ~ 1 min on
2500 grit paper with water lubricant to begin to reveal the outline of the shards. Blocks were then
215 moved on to the diamond laps with their appropriate lubricant, all at 280 revolutions min^{-1} rotating the
block 90 degrees every 30 s followed by 2 min of ultrasonic bathing at $< 24^\circ\text{C}$ between each lap stage
to remove any loose material on the surface of the blocks: ~ 3 min on 6 μm , ~ 1 min at 3 μm , and ~ 1
min at 1 μm . Blocks were then carbon coated before loading in the electron microprobe system for
analysis (EMPA).

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2.3. EPMA method and data reduction

Major element analysis of glass shards was undertaken at Victoria University of Wellington
(VUW) by wavelength dispersive X-ray spectroscopy (WDS) on a JEOL JXA8230 Superprobe electron
probe microanalyser (EPMA). Broadly the method follows that espoused by Kuehn et al. (2011).
225 Backscatter electron images of each sample were taken and used as block maps to allow the location of
EPMA analyses to be replicated for trace element analysis. A defocused circle beam 10 μm in diameter
was used at 8 nA to analyse all major elements as oxides (SiO_2 , TiO_2 , Al_2O_3 , FeO , MnO , MgO , CaO ,
 Na_2O , and K_2O). During standardisation, Na_2O was run twice, the second time skipping the peak search
to reduce the volatilisation of the element, with the second standardisation value then used. **Table 2**
230 shows the EPMA set up and run times. During the analysis, VG-568 was run as a calibration standard,
and VG-A99 and ATHO-G were run as secondary standards, with two of each standard (calibration and
secondary) analysed between ten sample analyses to monitor machine drift.

Initial concentrations were determined using the ZAF correction method, with secondary offline
data reduction undertaken to correct for variability in VG-568. Internal correction values were
235 calculated using the GeoREM reference values of VG-568 (**Eq. 1**) and applied to all the data (**Eq. 2**).
Following this, samples were corrected for deviations from 100 wt.% total, this assumes any variation is



due mostly to magmatic water, with a very small amount of minor and trace elements (Froggatt, 1983; Lowe, 2011) that are not analysed by the EPMA (Eq. 3). The difference is reported as “H₂O_D” in all data tables to allow back calculation to original data values including totals. Accuracy and analytical
240 precision of the standards were calculated, where accuracy is the percentage offset from the reference value for the secondary standards (Eq. 4), and precision is the standard deviation of all of the measured secondary standards throughout a run, reported at 2 standard deviations (sd) to represent a 95% variability.

245 **Eq. (1)** $Internal\ correction\ value = average(X_m^p/X_r^p)$

where X_m^p = measured concentration of element X of the calibration standard, and X_r^p = reference concentration for element X of the calibration standard (reference values taken from GeoRem preferred values <http://georem.mpch-mainz.gwdg.de/>).

250

Eq. (2) $corrected\ data = X_m^i / internal\ correction\ value$

where X_m^i = measured concentration for element X of any sample or standard.

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Eq. (3) $Secondary\ hydration\ corrected\ data = ((corrected\ data\ (Eq.2)/total\ for\ that\ sample) \times 100)$

260 **Eq. (4)** $\% \text{ offset from standard (accuracy)} = (((X_r^s - averageX_m^s) / X_r^s) \times 100)$

where X_r^s = reference concentration for element X of the secondary standard (GeoRem preferred value), and $averageX_m^s$ = average concentration measured for element X of all analyses)

265 2.4. LA-ICP-MS method and data reduction

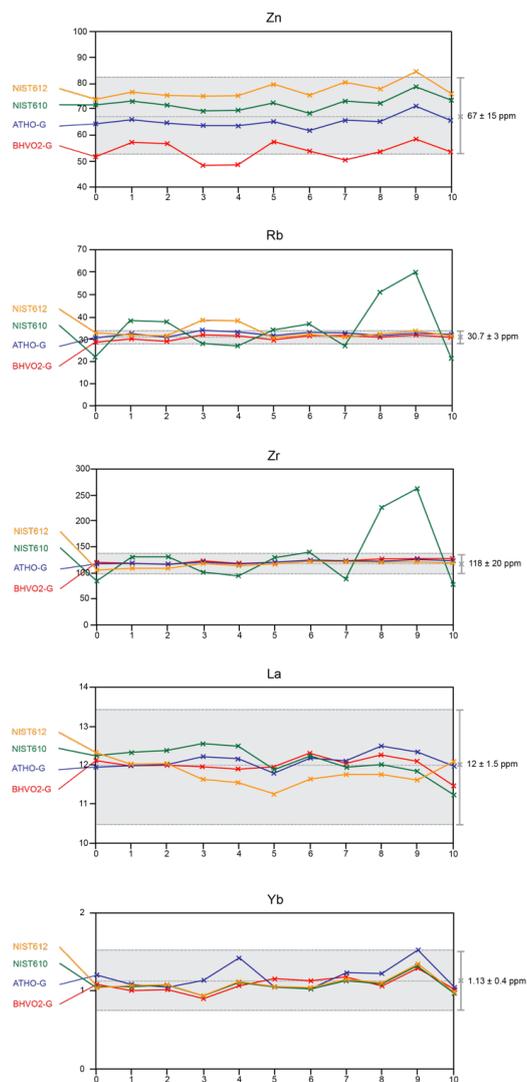
In situ trace element analysis was undertaken at VUW using laser-ablation inductively-coupled-plasma mass-spectrometry (LA-ICP-MS) where a RESolution S155-SE 193 nm ArF excimer laser system was coupled with an Agilent 7900 quadrupole ICP-MS. Data for 43 trace elements were acquired using a static spot method, with a 25 μm spot size, ablation time of 30 s, repetition rate of 5 Hz
270 power (see **Supplementary Material Table 1** for full LA-ICP-MS set up details). Synthetic glass standards NIST-612 and NIST-610 were used to tune the ICP-MS and obtain the P/A factors, at a range of spot sizes and laser powers. During the analysis a full range of standards were analysed to determine which produced the most accurate and precise results as a calibration standard, including NIST-612,



275 NIST-610, BHVO2-G, and ATHO-G. StHS6/80-G was analysed as a secondary standard throughout,
and all standards (calibration and secondary) were analysed twice every ten samples. All data were
reduced offline using Iolite v.3TM software (Paton et al., 2011), using ⁴³Ca as the internal standard value
(index channel) and the “*Trace_Elements_IS*” data reduction scheme (DRS). The data were reduced
against ATHO-G as the calibration standard. No post-processing data reduction was necessary for the
trace elements data; precision and accuracy were calculated on STHS6/80-G as described above (**Eq.**
280 **4**).

2.5. Standardisation method

Multiple calibration standards with different trace element concentrations were analysed to
determine which would be most suitable for trace element data reduction. Potential calibration standards
included NIST-612, NIST-610, BHVO2-G, and ATHO-G. These were each run twice every ten
285 samples, along with secondary standard STHS6/80-G. **Figure 2** shows the STHS6/80-G results of a
range of selected, commonly-used trace elements, including Zn (transition metal), Rb (LILE), Zr
(HFSE), La (LREE), Yb (HREE), normalised using each of the calibration standards. Overall, the
results show that for the lighter masses (e.g. Zn) there is a large variability in the measured STHS6/80-
G values across the different standards, but all except BHVO2-G sit within error (2 sd) of the reference
290 value (**Fig. 2**). For the heavier masses, the variation from the reference value observed within the
analysed values decreases, except for NIST-610, which remains highly variable in the middle masses
(Rb, Zr, **Fig. 2**), with variability reducing in the heavier masses (La, Yb, **Fig. 2**). The data show that the
use of ATHO-G as the calibration standard (for data reduction of rhyolites) produces the most accurate
and precise data for the secondary standard, for all except the elements with the heaviest masses, and
295 smallest concentrations (e.g. Yb).



300

Figure 2. Compilation of trace element standard data produced during the first run of glass shard analyses. These data show selected element concentrations of secondary standard STHs6/80 normalised using difference calibration standards (NIST-612 – orange; NIST-610 – green, ATHO-G – blue, and BHVO2-G – red). The grey shaded area shows the preferred GeoREM reference value (<http://georem.mpch-mainz.gwdg.de/>) error margin reported for each element for STHs6/80.



2.6. Statistical methods

2.6.1. Principal Component Analysis

To determine the elements that show the most variance with the reference dataset, and therefore
305 the most appropriate (optimum) for compositional separation, we have used principal component
analysis (PCA). PCA analysis was run in the coding platform R and RStudio using packages
“factoextra”, “ggbiplot”, “vegan”, “cowplot”, “rioja”, and “ggrepel”. Data for Tuhua tephra were
removed as these would unnecessarily skew the results due to their distinct geochemistry. Only element
values were used (no ratios [e.g. $\text{SiO}_2/\text{K}_2\text{O}$] or sums [e.g. $\text{Na}_2\text{O} + \text{K}_2\text{O}$]). All element values were
310 centred (column mean subtracted from each value) and scaled (value divided by the standard deviation
of the column) to allow their variability to be comparable, even when their absolute values are not. PCA
was run using the “prcomp” function, and PCA contributions were calculated using “fiz_comp”
function. A template of the coding script used can be found in **Supplementary Material 1**.

2.6.2. Euclidean similarity coefficients

To identify the tephra samples that were most similar, and could therefore pose problems in
315 unique fingerprinting, we ran euclidean similarity coefficients (ESC) analysis. ESC was run on the
coding platform R and RStudio using the package “stats”. Following the guidelines of Hunt et al. (1995)
for ESC analysis, we used non-normalised, mean concentrations of the elements highlighted by the
PCA to be the most indicative of variance in the dataset. These values were input as comparison values,
320 and the function “as.matrix.dist” was used to run the “euclidian” statistical method. This method
calculates the similarity of samples based on an infinite number of comparison input values. A template
of the coding script used can be found in **Supplementary Material 1**, the output table was manipulated
post-production to provide the colour formatting shown in **Figure 14**.

3. Results

325 The averages and their standard deviations for all samples are reported in **Table 3**; the full
reference dataset can be found in **Supplementary Material Table 2**. All reported values in the text and
figures (unless stated otherwise) are recalculated (normalised) to 100% on a volatile-free basis
(following Lowe et al., 2017) with the difference between the raw total and 100% being reported as
“ H_2O_D ” (**Table 3**). For best correlation results, we recommend that the full dataset is used in order to
330 see the trends in the geochemical data rather than just the means and standard deviations.



3.1. Major element results

All glass shards analysed are characterised as rhyolitic according to the classification of Le Maitre (1984) (**Fig. 3**), with SiO₂ concentrations ranging from 72.5 wt.% to 79.8 wt.% (with the majority 74-79 wt.%), and Na₂O + K₂O ranging from 5.2 wt.% to 9.8 wt.%. Three compositional regions with high concentrations of samples are evident within **Figure 3**. These show a negative trend between SiO₂ and Na₂O + K₂O, with each region separated by differing SiO₂ values – for example, SiO₂ = 76-77 wt.%, 77-78 wt.%, and 78-79 wt.%. Glass samples from the peralkaline Tuhua tephra (Tuhua/Mayor Island, TuVC) are identifiable because of their unique (peralkaline) geochemistry, with much higher Na₂O + K₂O (≥ 9 wt.%) for equivalent SiO₂ (= 73.5-75 wt.%; Lowe, 1988) in comparison to those of the rhyolitic TVZ-sourced deposits (Na₂O + K₂O ≤ 8.5 wt.%). Tuhua-tephra-derived glasses also have higher FeO (≥ 5.6 wt.%) and Na₂O (≥ 4.7 wt.%), but lower CaO (≤ 0.8), and Al₂O₃ (≤ 10.1) in comparison to the analyses for the rest of the samples (FeO = 0.2-2.8 wt.%, Na₂O = 2.6-5.1 wt.%, CaO = 0.5-2.6 wt.%, and Al₂O₃ = 11.8-15.2 wt.%; **Fig. 4**). For all other major elements, the compositional variation of the Tuhua tephra samples sits within the overall range for the other samples, with TiO₂ = 0.02-0.55 wt.%, MnO = 0.01-0.2 wt.%, MgO = 0.01- 0.63 wt.%, K₂O = 1.8-6.0 wt.%, and Cl = 0.01-0.72 wt.% (**Fig. 4**).

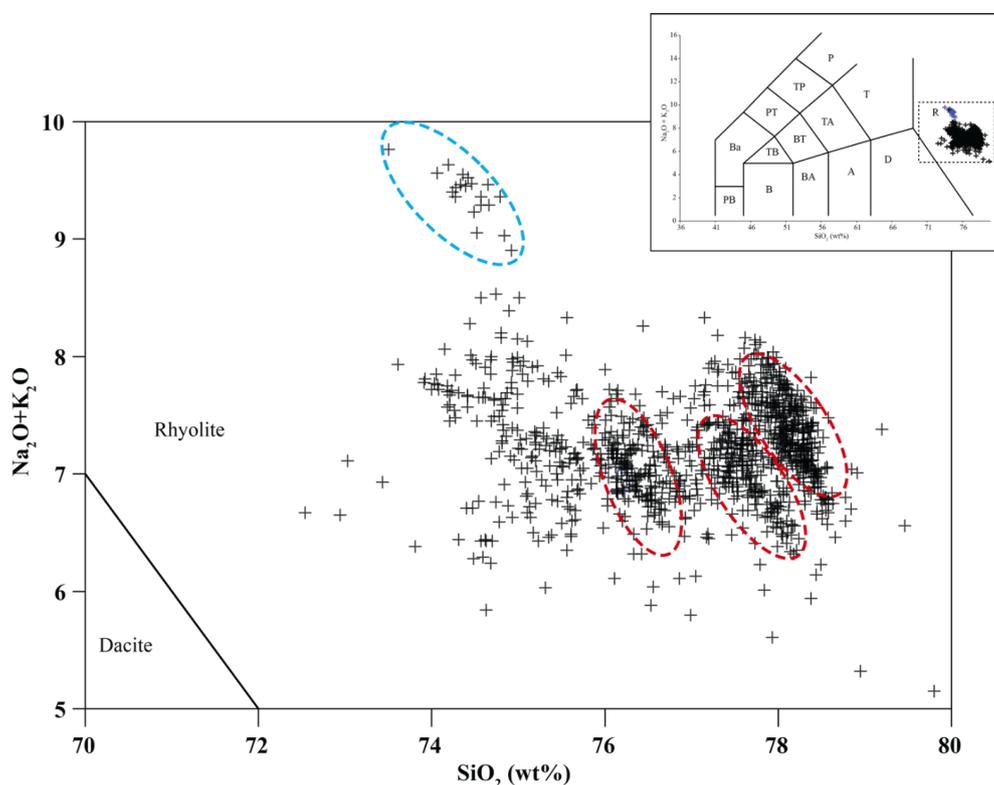
Of the 45 tephra samples, 22 have a ‘homogeneous signature’, homogeneity being defined here where the standard deviation of the sample is equal to or less than analytical error (2sd of secondary standard, e.g. for FeO = ± 0.23 wt.%, CaO = ± 0.10 wt.%). The majority (~64%) of the samples that have a homogeneous signature are from OVC (e.g. Whakatane, Mamaku, Rotoma) or from calderas older than OVC (~32%), for example: Upper Griffins Road tephra, a correlative of the Whakamaru eruptives, Whakamaru Volcanic Centre (WVC), and Mangapipi tephra, a correlative to deposits of Mangakino Volcanic Centre (MgVC; **Fig. 5a**). Ten samples show a heterogeneous signature (where standard deviations for both FeO and CaO are greater than analytical errors), with most from a proximal source (~30%), or from tephtras deposited in the Whanganui Basin area (40%), and with the remainder being from the Mangaone Subgroup eruptives from the OVC: Hauparu, Maketu, and Ngamotu (**Fig. 5b**).

Glass shards from four tephra samples show a bimodal signature in major and trace elements, where the populations split into two distinct groups. Tephtras showing this phenomenon include Rotorua (OVC), Rerewhakaaitu (OVC), Poihipi (TVC), and Tahuna (TVC). The bimodal signatures of Rerewhakaaitu and Rotorua are well documented (Shane et al., 2008), whereas those of Poihipi and Tahuna are newly identified here (**Fig. 6**). All four of these tephra horizons have their glass-shard bimodal signatures produced predominantly by K₂O concentrations, into high (≥ 3.8 wt.%) and low (≤ 3.6 wt.%) populations (**Fig. 6**) linked to the crystallisation of biotite minerals.

For five of the tephtras, we undertook analyses from both proximal and distal samples. These tephtras included Whakatane, Rotoma, Waiohau, Rotorua, and Rerewhakaaitu, which are all derived



370 from OVC (**Table 1**). For Rotoma, Rerewhakaaitu, and Waiohau, the signatures of the proximal and distal deposits are indistinguishable, whereas, for Whakatane and Rotorua the proximal signature is highly variable, and the distal signature is homogeneous but overlapping with part of the extent of the proximal signature (**Fig. 7**). Similar findings are reported and discussed in more detail for Whakatane tephra in Kobayshi et al. (2005) and Holt et al. (2011); and for Rotorua tephra in Shane et al. (2003a) and Kilgour and Smith (2008).



375 Figure 3. Total alkali ($\text{Na}_2\text{O} + \text{K}_2\text{O}$) vs. SiO_2 (TAS) plot for glass compositions for all reference data (presented on a normalised basis). Identified and highlighted by blue dashed outline are the glass shard compositions for the Tuhua tephra (Mayor Island; MI), and highlighted by the red dashed outlines are the regions on the TAS diagram that show the highest density of samples. The inset shows a full TAS diagram (always on an anhydrous basis) to provide context for the enlarged figure. Regions of the TAS diagram follow the nomenclature of Le Maitre (1984) are: A – andesite, B – basalt, Ba – basanite, BA – basaltic andesite, BT – basaltic-trachyte, D – dacite, P – phonolite, PB – picro-basalt, PT – phonotephrite, R – rhyolite, T – trachyte, TA – trachy-andesite, TB – trachy-basalt, TP- tephriphonolite.

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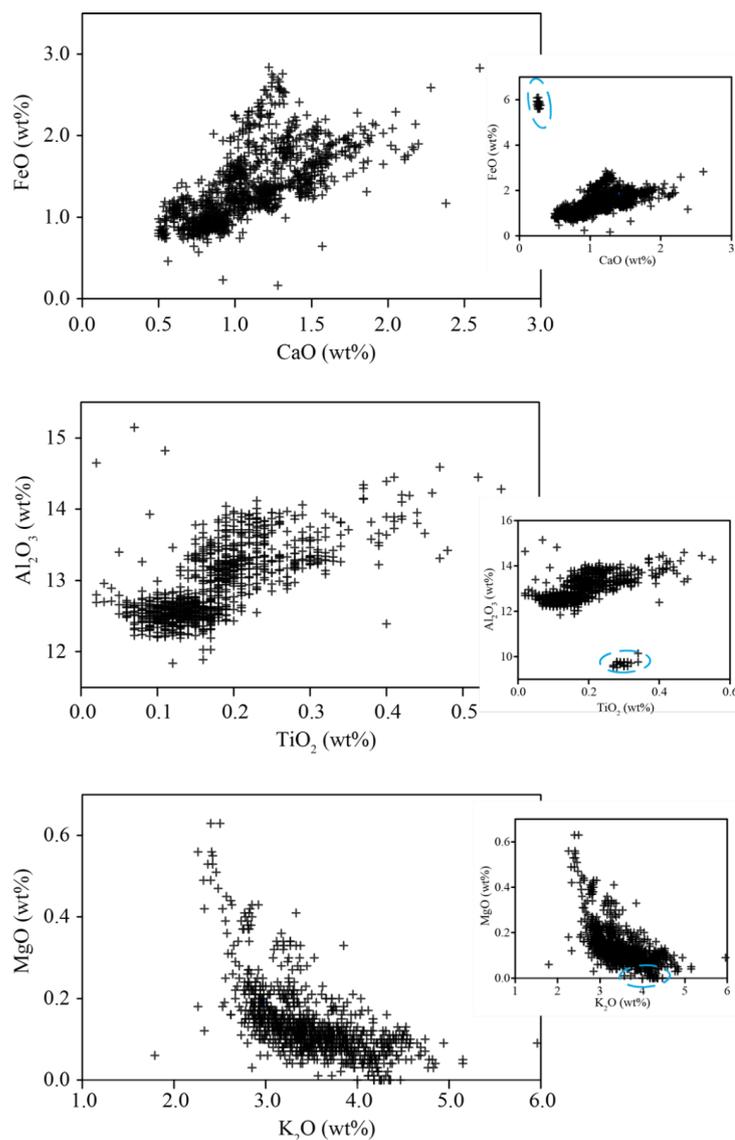
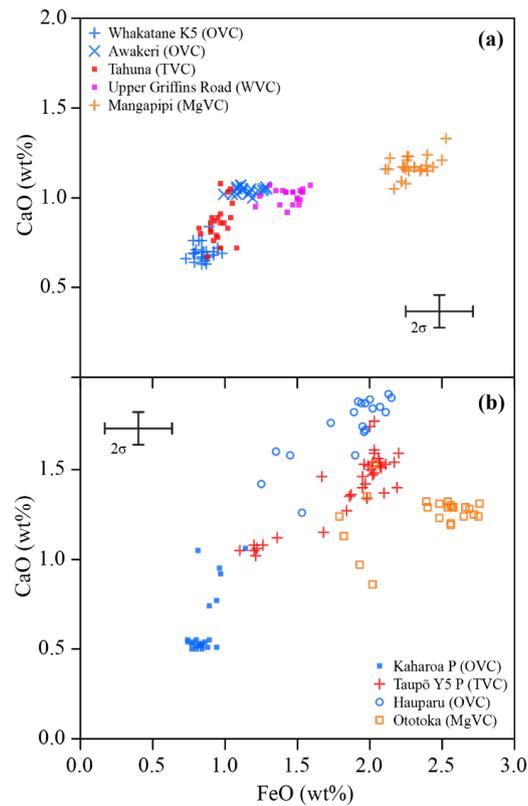


Figure 4. Major element bivariate plots of glass shard compositions for all reference data (presented on a normalised basis). Highlighted in the insets by blue dashed lines are the Tuhua tephra samples. These are removed from the enlarged figure to allow the detail of the majority of the samples to be seen more clearly. Total iron expressed as FeO.



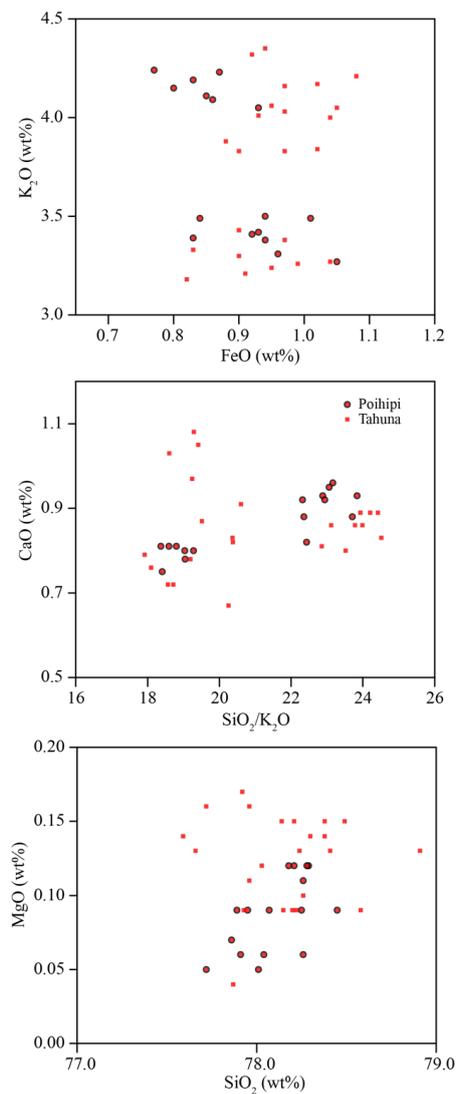
390 **Figure 5. Major element bivariate plots examples for glass shard analyses of tephras (presented on a normalised basis) which show (a) a homogeneous signatures, where the standard deviation of the analysis is less than the analytical error (shown as 2σ); and heterogeneous signatures, where the standard deviation of the analysis is greater than the analytical error. Different colours indicate the differing caldera sources (shown on Fig. 1) and different symbols show the different tephras. P = proximal sample (see Table 1). Total iron expressed as FeO.**

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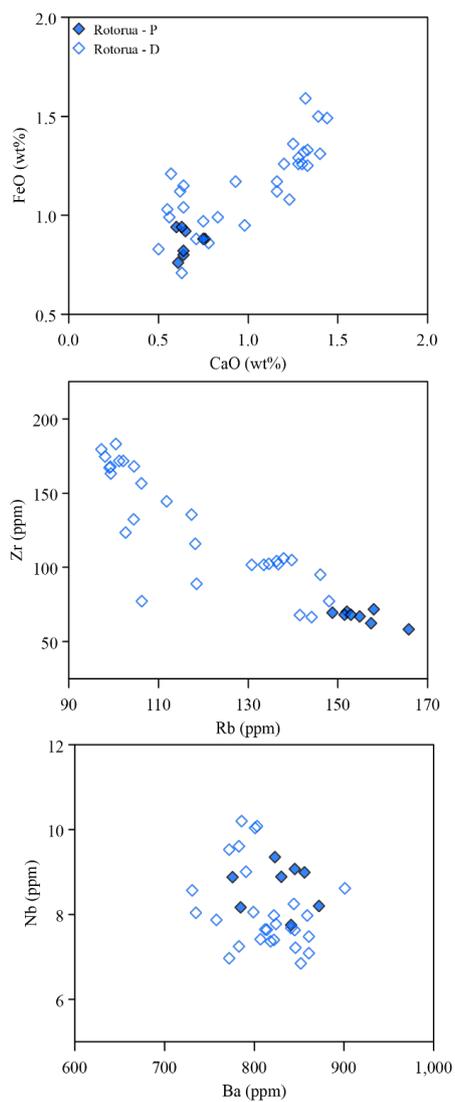
3.2. Trace element results

Figure 8 shows a chondrite-normalised spider plot of all the trace element data for the glass shards analysed. The majority of the data plots along a common pattern of variable concentrations of HFSE, LILEs and LREEs, but they show more consistent concentrations of HREEs. Of note are peaks in Ba, Nd, Pr, and a negative Sm anomaly. Sr shows the largest variability in concentrations from < 1 to ≤ 100 ppm, likely caused by a variability in feldspar crystallisation (Pearce et al., 2004). Several different patterns are observable within this full data suite pertaining to individual samples. The obviously different signature is that for glass from Tuhua tephra which shows a low concentration of Ba (< 10 ppm) and Sr (< 1 ppm) in comparison with values for the rest of the samples, and with high concentrations of all other elements, especially the REEs (**Fig. 8**). Analyses of glass shards from the Maketu tephra can also be identified by their very high Ba values (> 1000 ppm), mid-range Nb values (between those of Tuhua and the general trend), and much higher concentrations of all other elements (**Fig. 8**). We also note Er and Lu peaks, which pertain to glasses from the Te Rere tephra, that sit at the higher concentration levels of the general trend (**Fig. 8; Table 3**), and samples from Ngamotu, Rotoehu/Rotoiti and Earthquake Flat that sit at the lower overall trace element concentration levels of the general trend, but with high Ba values (**Fig. 8**). For the tephtras where both proximal and distal samples of glass have been analysed for trace elements, the HFSEs (including Zr, Hf, Th, and Ti) and LILEs (including Rb, Sr, and Cs) can be used to maintain the heterogeneity between the proximal and distal samples, whereas the HREE and the LREE tend to have a lower variability (**Fig. 7**).



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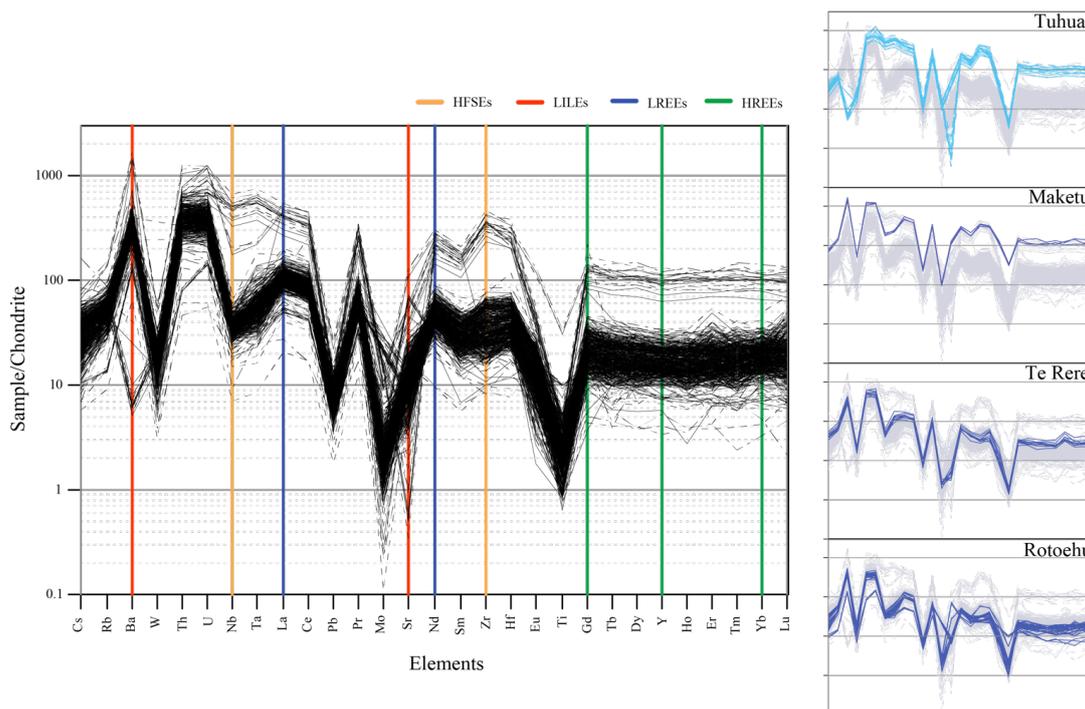
Figure 6. Selected major element biplots of glass analyses (presented on a normalised basis) of samples from Poihipi and Tahuna tephtras (both TVC sourced) that exhibit a bimodal signature. This bimodality is identified as being caused by K_2O concentration (e.g. see Lowe et al., 2008; Shane et al., 2008), and therefore plots with other elements (major or trace) do not show this bimodality. Total iron expressed as FeO .



425

Fig. 7. Major and trace element biplots showing the glass-shard-derived geochemical relationship of Rotorua (OVC) proximal (P) and distal (D) tephra deposits (presented on a normalised basis, total iron expressed as FeO). Distal deposits may have a signature with lower geochemical variability which overlaps within the spread of the heterogeneous proximal signatures. This variation can often be resolved by using trace element plots of selected elements – see text for discussion.

430



435 **Figure 8.** Chondrite normalised (McDonough and Sun 1995) trace element spider plot for glass analyses for all reference samples. Highlighted are key elements discussed in the text coloured by their characteristics including HFSE, LILE, LREE, and HREE. The full plot is presented to show the density of data with the dominant trend line plus the obvious deviations from this. The samples which correspond to these deviations are shown in the smaller plots at right, including analyses on glass from Tuhua (MI), Maketu, Te Rere, and Rotoehu (OVC) tephra. The Rotoehu Ash signature is also similar to that for the Rotoiti Ignimbrite (which are coeval deposits; Nairn, 1972), the Earthquake Flat tephra (Kapenga VC; Nairn and Kohn, 1973), and the Ngamotu tephra (OVC; Jurado-Chichay and Walker, 2000).

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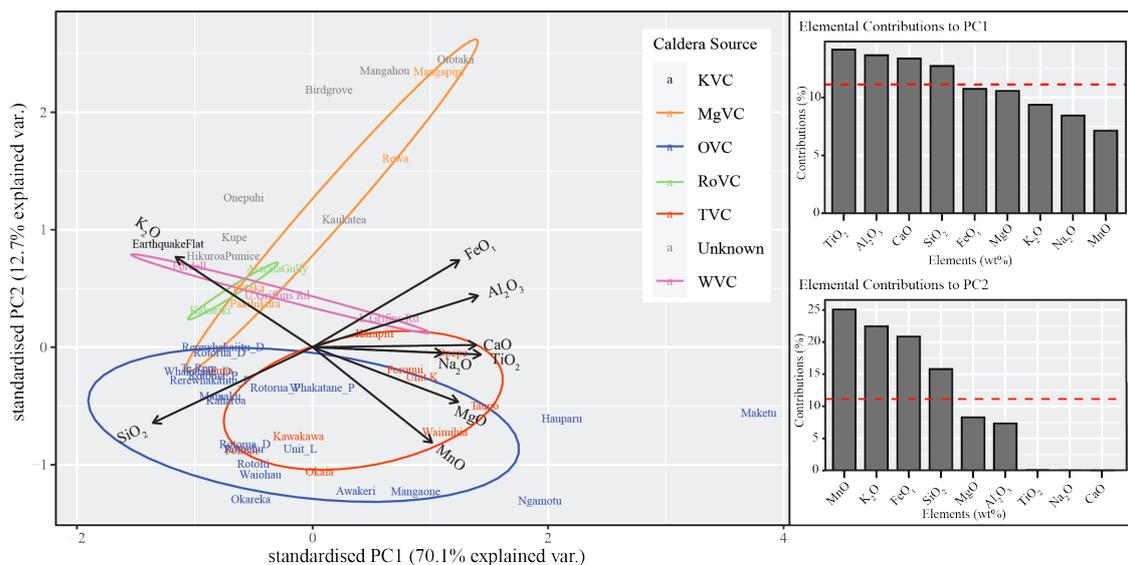
4. Discussion

4.1. Distinguishing geochemical characteristics

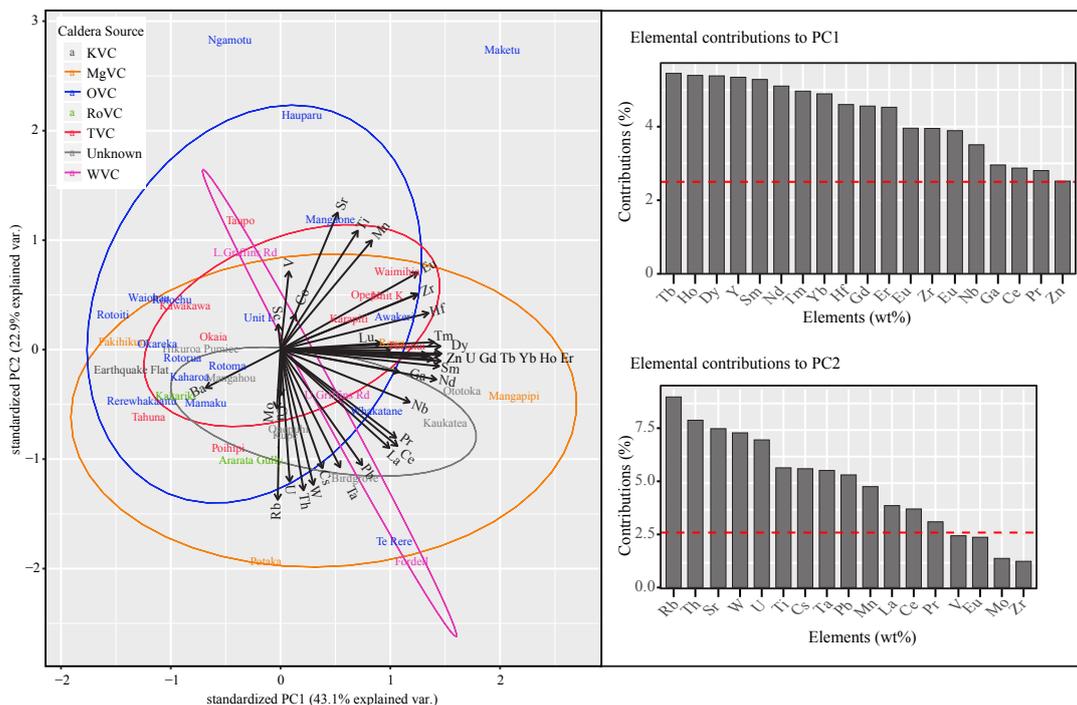
445 4.1.1. Major and trace elements in general

In many cases, the major element concentrations in glass are sufficient to allow different tephra to be distinguished, a result consistent with the findings from much previous work both in New Zealand and elsewhere (e.g. Lowe et al., 2017). PCA results for the glass-shard major elements (**Fig. 9**) show that PC1 and PC2 explain 82.7% of the variance within the data. When scaled, concentrations of TiO₂, Al₂O₃, CaO, and SiO₂ for PC1 (**Fig. 9**), and MnO, K₂O, FeO, and SiO₂ for PC2 (**Fig. 9**) are shown to have the highest contribution to the variance and are therefore most appropriate for distinguishing between tephra deposits for the reference dataset as a whole (**Fig. 9**). These major elements, especially CaO, FeO, and K₂O, have long been recognised as being useful to distinguish many New Zealand late Quaternary tephra from one another (e.g. Lowe, 1988; Shane, 2000; Alloway et al., 2013), the presence of TiO₂, Al₂O₃, and MnO are somewhat unusual. In a number of cases (discussed below) however, major element concentrations are shown to overlap for certain tephra horizons, and thus trace elements and trace element ratios are investigated to provide additional variables to use as discriminants. PCA was also applied to scaled trace elements with the results suggesting that PC1 and PC2 could explain 66.0 % of the variability in the trace element data with Tb, Ho, Dy, Y, Sm, Nd, Tm, Yb, Hf, and Gd the ten highest contributors to PC1, and Rb, Th, Sr, U, Cs, Ta, Pb, La, Ce, and Pr highlighted as the ten highest contributors the variance of PC2 (**Fig. 10**). Therefore, statistically these trace element concentrations and ratios of these trace elements have the potential to be the most useful in distinguishing the individual tephra horizons when using their glass-shard compositions alone.

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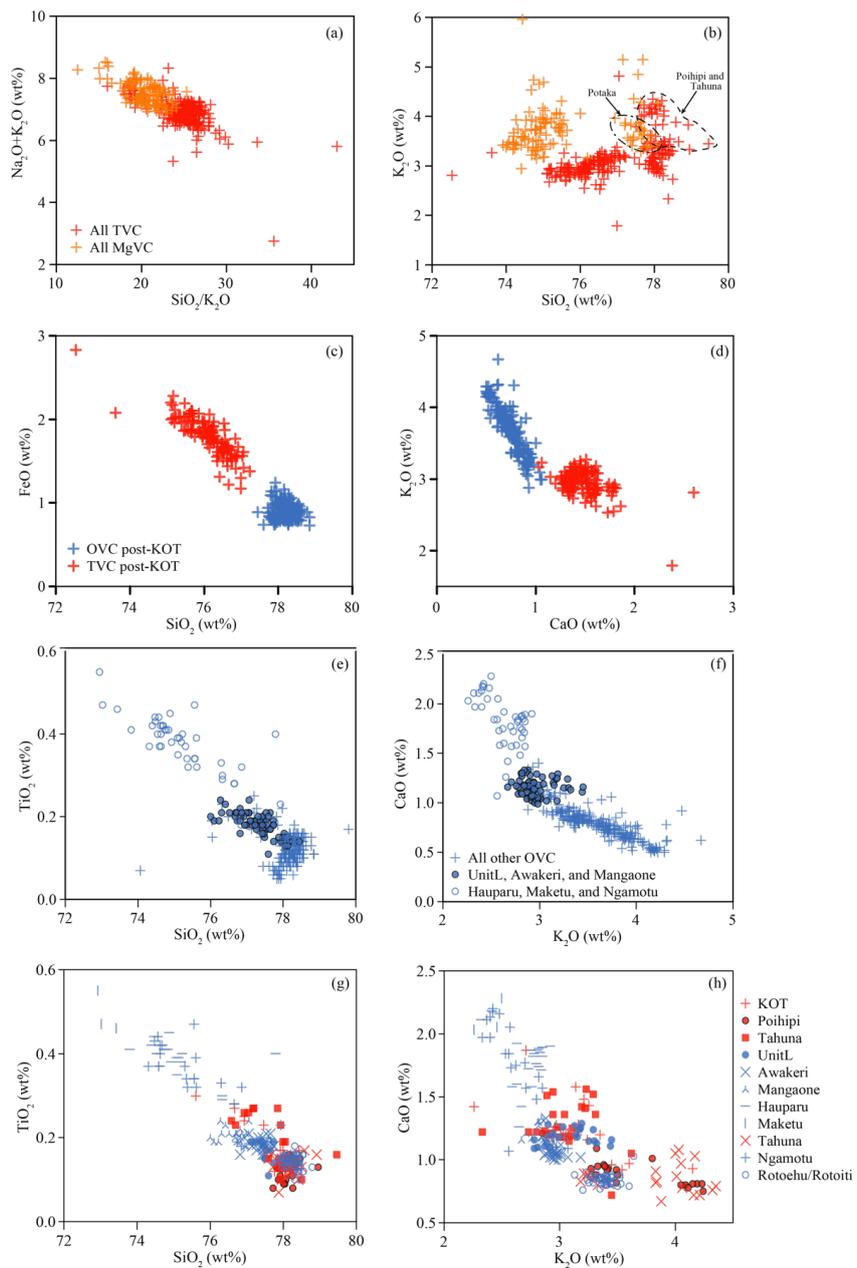


470 **Figure 9.** Results of PCA analysis on all TephraNZ major element reference data for glass (normalised). Data are scaled to allow comparison within the PCA analysis. Tephra names are coloured as per their source centre. PCA analysis was performed in R (see SM1 for R script). Bar plots highlight the top elemental contributions for PC1 and PC2. The red dashed line on the elemental contribution plots indicate the expected average contribution; if the contribution by each element was uniform, the expected value would be $1/\text{no. variables}$ (e.g. $1/9 = \sim 11\%$). Therefore, a variable with a contribution larger than this cut off line ($\sim 11\%$) is considered important in the contributing to the component.



475 **Figure 10. Results of PCA analysis on all TephraNZ reference trace element data for glass. Data are scaled to allow comparison within the PCA analysis. Tephra names are coloured as per their source centre. PCA analysis was performed in R (see SM1 for R script). Bar plots highlight the top elemental contributions for PC1 and PC2. The red dashed line on the elemental contribution plots indicate the expected average contribution, a variable with a contribution larger than this cut off line is considered important in the contributing to the component.**

480





485 **Figure 11. Major element biplots to distinguish between caldera sources of tephra based on their glass major element compositions**
(presented on a normalised basis; total iron is expressed as FeO). (a) & (b) show a comparison between all glass shard analyses for
the TVC and MgVC sourced tephra. (c) & (d) Indicate the distinction in glass compositional signature for the eruptives from the
OVC and TVC that post-date the Kawakawa/Oruanui (KOT) eruption. (e) & (f) These plots distinguish the glass analyses for tephra
from the OVC into component eruptive time periods, with tephra from the Mangaone subgroup (Jurado-Chichay and Walker, 2000;
Smith et al., 2002) distinguishable from all other tephra from the OVC. (g) & (h) Show the similarity in the geochemical compositions
490 for the tephra from the OVC and TVC for the eruptions prior to, and including, the KOT eruption. Colours are consistent for each
caldera source, symbols are representative of different groups of tephra defined in the keys for each set of plots.

4.1.2 Source-specific major and trace elements

The central TVZ contains nine recognised calderas, each with different eruption histories, but all
495 having produced large magnitude/volume tephra-producing rhyolitic eruptives. Some of the calderas are
attributed to single caldera collapse events (Rotorua, Reporoa, and Ohakuri), others to composite
collapse events that overlap spatially but not temporally (Mangakino and Kapenga), but the majority to
multiple collapse events over an extended period of time (Maroa, Okataina, Taupō, and Whakamaru)
(**Fig. 1**; Wilson et al., 1995a, 2009; Barker et al., 2021). Although the calderas are mostly discrete in
500 space, evidence from multiple eruptions has shown their plumbing systems may be linked tectonically
(e.g. Wilson et al., 2009; Allan et al., 2012). Hence, the ability to trace a tephra deposit to a caldera
source through glass-shard geochemistry alone could be challenging.

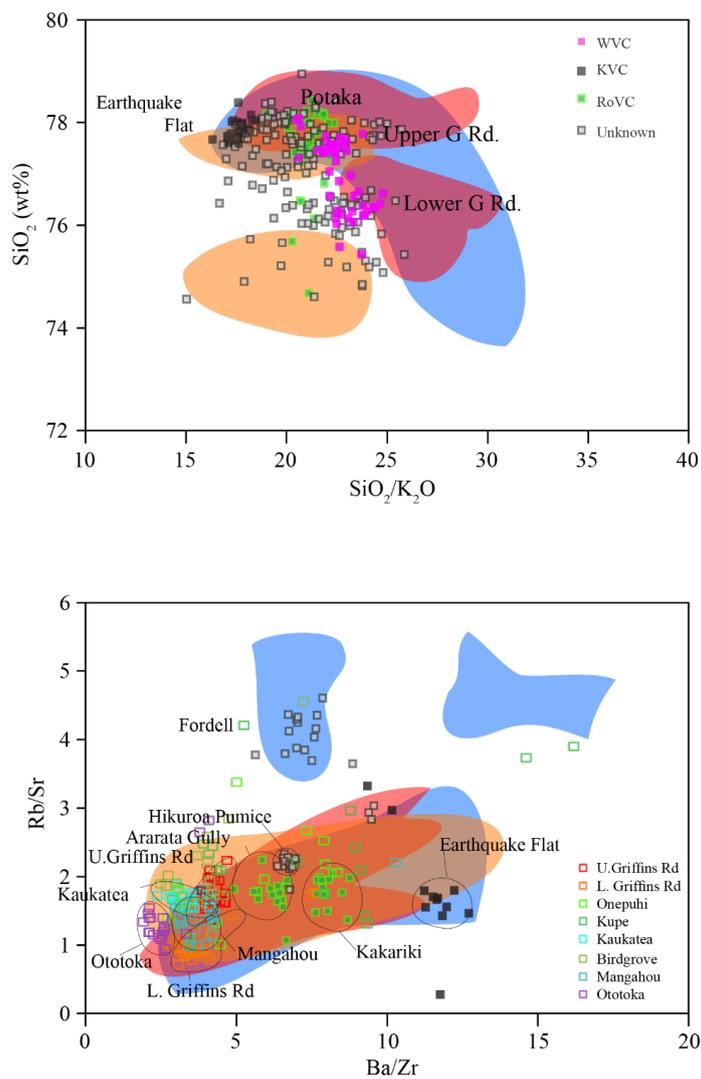
The results of the PCA analysis suggest that tephra sourced from the TVC can be distinguished
from those of a proposed Mangakino source (MgVC) (**Fig. 9**). Using $\text{SiO}_2/\text{K}_2\text{O}$ vs. $\text{Na}_2\text{O}+\text{K}_2\text{O}$ ratios,
505 the glass shards of the TVC tephra generally have higher $\text{SiO}_2/\text{K}_2\text{O}$ and lower $\text{Na}_2\text{O}+\text{K}_2\text{O}$ ratios in
comparison to those of the equivalent oxides for MgVC-sourced tephra (**Fig. 11a**). This information is
important, but because of the age differences for the calderas (TVC ~ 0.32 Ma to present, and MgVC
~1.6 Ma to 1.53 Ma and ~1.2 Ma to 0.95 Ma 1995), the use of this distinction is likely more important
for discussions on mantle source dynamics rather than for geochemical correlation of tephra deposits.

510 Previous studies have suggested that the geochemical characteristics of glass shards from TVC
and OVC tephra deposits can be distinguished from after the eruption of the Kawakawa/Oruanui (KOT)
to the present day using $f\text{O}_2$ of Fe-Ti oxides and minerals (Shane, 1998), pumice and lava compositions
(Sutton et al., 2000), and glass chemistry (Froggatt and Lowe, 1990). Our results also show there is a
bimodality in the TVC glass-shard data as a whole and that the post-KOT tephra deposits from the TVC
515 and OVC are quite different whereas the pre-KOT tephra from OVC and TVC are similar (**Fig. 11c&d**).
Most glass shards erupted after the KOT event from the TVC have low SiO_2 (≤ 77 wt.%) and less
variable K_2O (~3 wt.%), and higher values for all other major elements in comparison to those of the
glass shards erupted from the OVC (**Fig. 11c&d**). In comparison, tephra erupted from the TVC and
OVC prior to, and including the KOT, do show a large amount of overlap in their glass geochemical
520 signatures. For OVC, there is a high density of samples that have their SiO_2 concentrations at ~78 wt.%;



525 however, there is a high variability in SiO₂ overall, with Maketu, Hauparu, and Ngamotu of the
Mangaone subgroup plotting with SiO₂ concentrations $\leq \sim 76$ wt.%, and the remaining Mangaone
subgroup samples (Unit L, Awakeri, and Mangaone) clustering at SiO₂ = 76-77.5 wt.% (FeO \approx 1.2
wt.%, K₂O \approx 2.8 wt.%, Al₂O₃ \approx 13 wt.%, CaO \approx 1.2 wt.%), a finding consistent with those of Smith
et al. (2005) who divided the Mangaone subgroup into 'old' and 'young' eruptives on the basis of low
and high SiO₂, respectively, and also unlike the other OVC sourced samples that plot around SiO₂
 \approx 77.5-79 wt.%, (FeO \approx 0.8-0.9 wt.%, K₂O = 2.75-4.5 wt.%, Al₂O₃ \approx 12-13 wt.%, CaO \approx 0.5-1.0
wt.%; **Fig. 11c&d**). Analyses from the Rotoehu/Rotoiti tephra deposits plot independently from those
of other OVC eruptives for this time period. However, they overlap with those of some TVC-tephra-
530 derived glass compositions (Poihipi, Tahuna, Okaia, and KOT). The Rotoehu/Rotoiti tephra deposits
have a markedly homogeneous geochemical signature, and are also much older than TVC eruptions
(**Table 1**). Hence, coupled with the thickness of the deposits, it is likely that a tephra linked to the
Rotoehu/Rotoiti eruption would be obvious to distinguish through stratigraphy and age combined with
the geochemistry.

535 The TephraNZ dataset presented here also includes analyses of glass of samples from tephra
erupted from the Kapenga Volcanic Centre (KVC; Earthquake Flat eruption), Rotorua Volcanic Centre
(RoVC), and Whakamaru Volcanic Centre (WVC). In addition, some older tephra deposits have been
recorded in the Whanganui Basin and elsewhere. These are well-known beds but their caldera sources
are not yet defined (Alloway et al., 1993; Pillans et al., 1994, 2005; Shane et al., 1996; Rees et al., 2018;
540 2019). **Figure 12** shows a comparison plot for the data from KVC, RoVC, and WVC with those regions
populated by glass data from samples from the OVC, TVC and MgVC sources. Overall, the samples
plot with a lower SiO₂/K₂O ratio ($\leq \sim 25$), similar to that of the MgVC-sourced tephra, which seems to
be indicative of older sources in comparison to the OVC and TVC. The samples potentially linked to
RoVC (Bussell, 1986; Bussell and Pillans, 1997) show different geochemical compositions. For
545 example, Kakariki-tephra-derived glass has slightly higher SiO₂ ≥ 78 wt.% in comparison to that of the
Ararata Gully tephra (SiO₂ ≤ 77 wt.%), suggesting that they are likely derived from different eruptions,
but potentially the same source (Mamaku Ignimbrite reportedly has variable geochemical phases;
Milner et al., 2003). Glass from the KVC sample (Earthquake Flat tephra) has a very homogeneous
signature in the major elements, but a more variable signature in the trace elements, both of which
550 overlap with OVC- and TVC-source signatures. There is a very large spread for the data from the
unknown samples, precluding the ability to specify their source based simply on major and trace
elements alone. Nevertheless, their glass compositional signatures are clearly more similar to those of
the older MgVC sourced tephra, in comparison to those of the younger TVC and OVC deposits, as
would be expected based on their known age range (**Table 1**).



555

Figure 12. Major and trace element biplots of indicative elements in glass to show the relationships between the tephras from Mangakino Volcanic Centre (MgVC – orange shaded regions), Taupō Volcanic Centre (TVC – red shaded regions), and Okataina Volcanic Centre (OVC- blue shaded regions), and the tephras from known and unknown sources within the TephraNZ data base.

560



4.1.3 Homogeneous, heterogeneous, and bimodal samples

Fingerprinting of glass shards for correlation relies on the ability to distinguish between different deposits and therefore a homogeneous signature that is distinct from all other samples is the ideal ‘fingerprint’. However, sometimes there is more complexity in the geochemical data and heterogeneity can develop in a tephra deposit through a number of mechanisms (Lowe, 2011):

- (1) Variability in the magma body itself (e.g. Nairn, 1992; Nairn et al., 2004; Smith et al., 2004; Kobayashi et al., 2005; Shane et al., 2008; Charlier and Wilson, 2010; Klemetti et al., 2011; Cole et al., 2014);
- (2) Proximal vs. distal complexity, linked to (1) (e.g. Manning, 1996; Shane et al., 2003a; Holt et al., 2011);
- (3) Post- or syn- depositional reworking (e.g. Schneider et al., 2001)

For example, the heterogeneous signature identified for the Kaharoa tephra agrees with previous findings for this eruption. Nairn et al. (2004) and Sahetapy-Engel et al. (2014) reported that tephra compositional variability within the Kaharoa deposits shows sequential tapping of a stratified magma body coupled with syn-eruptive changes in dispersal patterns. In general, this is likely one of the reasons why some of the proximal tephra deposits analysed in this study have a more variable geochemical signature in comparison to those of their distal counterparts (**Fig. 7**). Although the proximal deposits record the detail in the eruption progression, the distal deposits tend to record the very largest phase of the eruption (e.g. Walker, 1980) but differences can be expected to occur according to the azimuths of wind direction during an eruption and the number and degree of interconnectedness of magma bodies involved in the eruption (e.g. Walker, 1981; Kilgour and Smith, 2008; Sahetapy-Engel et al., 2014; Storm et al. 2014; Rubin et al., 2016).

The tephrochronological principle is much more likely to utilise distal unknown deposits, and therefore we suggest that using the distal signature (or signatures) maybe more appropriate for correlation. In general, distal tephtras are more chemically homogeneous – but with some notable and well-documented exceptions – and this attribute therefore allows them to be traced over large areas (Manning, 1996). Alternatively, the identification of heterogeneity or bimodality in distal tephtras, once recognised, can be an additional useful characteristic for fingerprinting (e.g. Shane et al., 2003a, 2008; Lowe et al., 2017). These statements, however, rely on the tephra being identified as a primary deposit, and not reworked. Reworking is commonly seen in paleofluvial deposits, for example those in the Whanganui Basin, and in other environments prone to mixing such as in surficial soils. This reworking can mix tephra from multiple eruptions, and can cause highly variable glass chemistry within a single deposit (e.g. Shane et al., 2005, 2006). Fluvial reworking can be commonly identified by sedimentary structures within the deposit, for example, ripples or cross bedding indicative of fluvial transport and deposition (e.g. Shane, 1994; Schneider et al., 2001), over thickening of deposits (e.g. Vucetich and



Pullar, 1969; Lowe, 2011), or through shard morphology, for example anomalously large shards or rounding of shards (e.g. Leaphy, 1997).

Heterogeneous signatures (where the standard deviation of the analyses is greater than the analytical error) in major element compositions were identified for ten of the tephra deposits: Kaharoa, Taupō Y5 Proximal (P), Whakatane P, Hauparu, Maketu, Ngamotu, Fordell, Onepuhi, Birdgrove, and Ototoka. Our data show that for some samples, specific trace elements and trace element ratios have lower geochemical variability (**Fig. 13a**). The elements that work best to separate out the individual units within a deposit with a heterogeneous signature reflect the minerals that have formed during fractional crystallisation of the melt. Because of this, different elements or element ratios work for different tephtras. For example, for Kaharoa, Sr acts to effectively reduce the variability of the signature, whereas for Taupō Sr can be used to maintain the heterogeneity in the sample (**Fig. 13a**).

Bimodality was identified for four of the tephra horizons analysed: Rotorua (OVC), Rerewhakaaitu (OVC), Poihipi (TVC), and Tahuna (TVC). For all four of these, K₂O concentration causes the bimodality, and therefore trace elements with similar chemical properties reinforce the bimodality (for example, LILEs Rb, Sr, and Cs; HFSEs Zr, or REE Eu), whereas most other trace elements do not show this bimodal signature (**Fig. 13b**).

4.2. Indistinguishable tephtras

Euclidean similarity coefficient (ESC) analysis was used on all glass-shard reference data for tephtras from Rotoiti/Rotoehu to Kaharoa in addition to the PCA and geochemical investigation to determine those samples that have indistinguishable element concentrations at similar ages (**Fig. 14**). **Figure 14** shows similarities (similarity coefficient values (SC) are observed in the major element signatures between Waimihia and Unit K (SC=0.113); Mamaku and Rotoma-D (SC=0.170), Rotoiti/Rotoehu and Rotoma-D (SC= 0.105 and 0.126, respectively); KOT and Okaia (SC=0.141); KOT and Unit L (SC=0.137); and Poihipi and Tahuna (SC=0.120). When the key trace element are analysed, only Poihipi and Tahuna (SC = 15) and Mamaku and Rotoma-D (SC=3) come up with significantly low similarity coefficients, hence suggesting that these samples will be indistinguishable in both major and trace elements. When trace element ratios are run through the SC analysis, Waimihia and Unit K (SC=6), Waimihia and Poronui (SC=9), Unit K and Poronui (SC=5), and Karapiti and Waimihia (SC=7) show significantly low similarity coefficients. In addition, when simple geochemical assessment is applied, similarities are observed between Taupō and Waimihia, and Waiohau, Rotorua, and Rerewhakaaitu (**Table 5**).

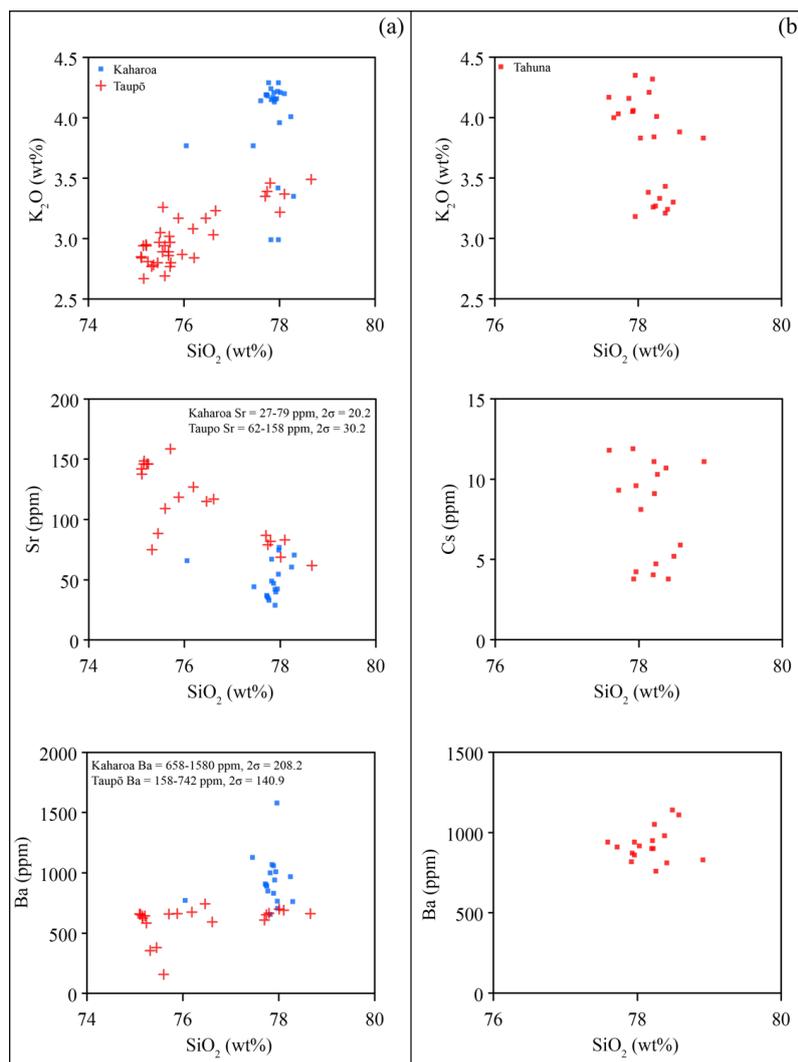
These results suggest that for Poihipi and Tahuna, and Mamaku and Rotoma tephtras, trace element ratios in glasses could enable them to be distinguished. **Figure 15a** shows that for Poihipi and Tahuna the best separation (although some overlap remains) is seen in the ratios La/Yb vs. Ba/Y; in addition, Tahuna also shows a bimodality in Ba/Th ratio which is not seen for Poihipi. For Rotoma and



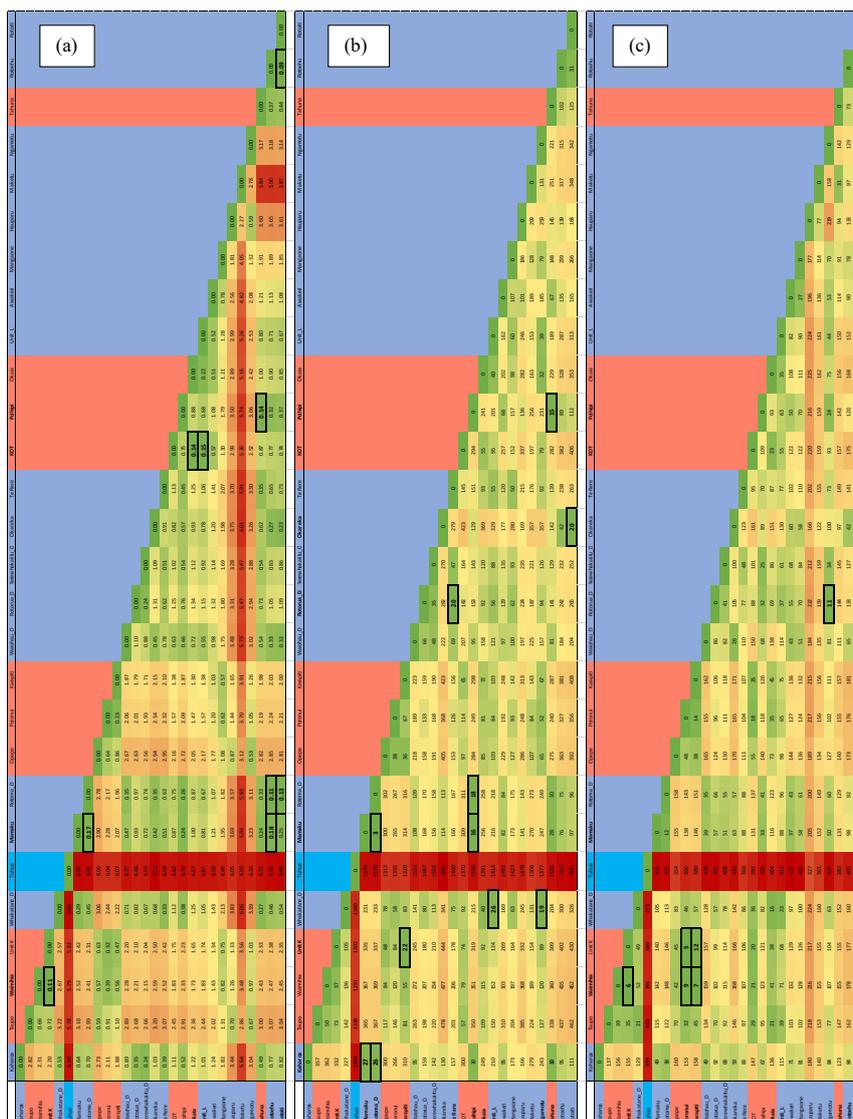
Mamaku, the tephra can be separated (although some overlap remains) using Ba/Th vs. Rb/Sr and Rb/Zr vs. Rb/Sr (**Fig. 15b**). Rotoma and Rotoehu/Rotoiti are very similar in their glass-shard major elements, but can be distinguished using specific, but a wide range, of trace elements (**Fig. 15c**). They are also very different in age, hence should not be too difficult to distinguish on the basis of stratigraphy or dating.

Waimihia and Unit K (Taupō Subgroup) tephra are very difficult to distinguish, and their relative similarity in age (3382 ± 50 and 5088 ± 73 cal. yr BP, respectively; Lowe et al., 2013) and mineralogy could see them misidentified if dates were unavailable or imprecise. Geochemical investigation beyond the PCA and SC analyses of glass shows that Lu, Sc, Mn, and Co can be used to geochemically distinguish these two tephra (**Fig. 15d**), indicative of fractional crystallisation of differing amounts of clinopyroxene, plagioclase, and amphibole during the eruptive events. Although not identified by the SC analysis directly, Poronui (11,195- 51 cal yr BP) and Karapiti (11,501 \pm 104 cal. yr BP) tephra also have comparable age, geochemistry, and mineralogy; thus using major, trace, and trace element ratios these two tephra remain indistinguishable. Glass shards from the three Holocene tephra, Waimihia, Poronui, and Karapiti, also have very similar trace element and trace-element ratios but, as for Waimihia and Unit K, they can be distinguished with Lu, Sc, Mn, and Co, where Waimihia has higher Sc, Lu, and Mn, but lower Co in comparison to those of the Poronui and Karapiti tephra. They can also be distinguished simply with a biplot of FeO vs. CaO, or Na₂O+K₂O or SiO₂/K₂O, or SiO₂, where the Waimihia samples in general have lower FeO, Na₂O+K₂O, SiO₂/K₂O, and higher CaO, and SiO₂ in comparison to the equivalent values for Poronui and Karapiti samples (**Fig. 15e, Table 5**).

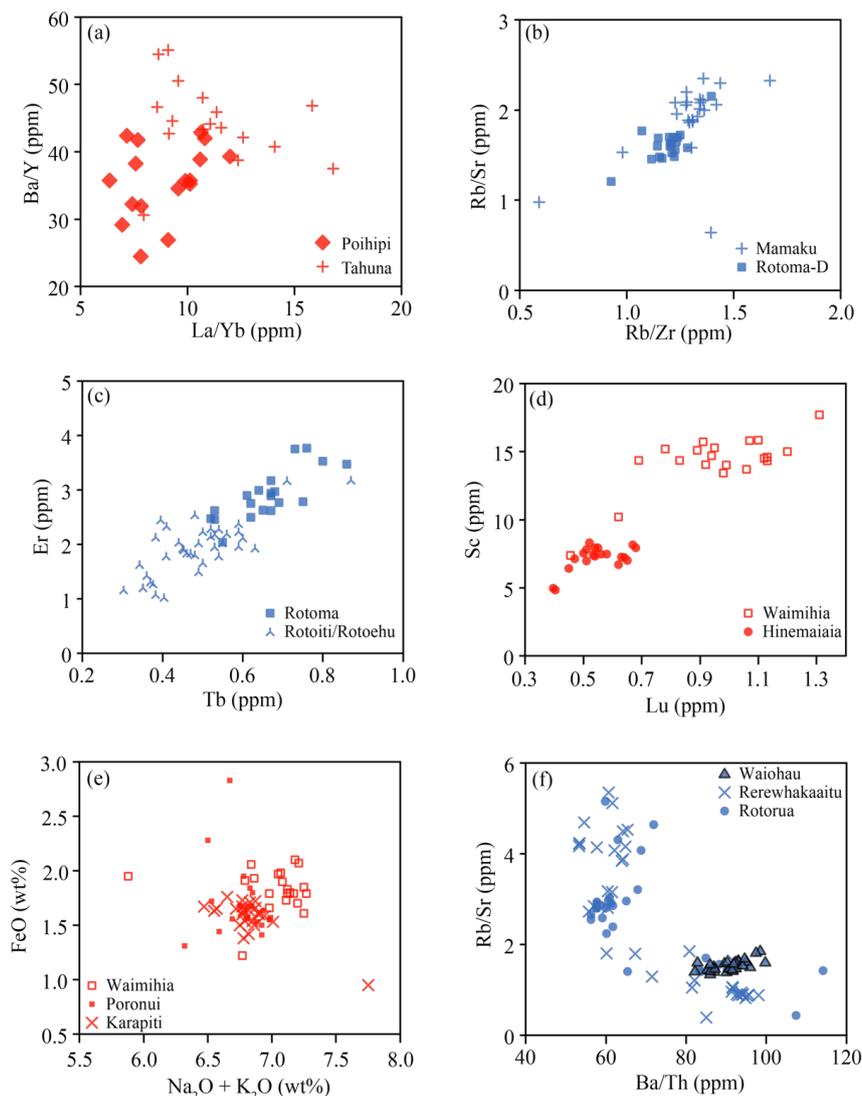
Geochemical investigation and PCA analysis also highlights the similarity of the Waiohau, Rotorua, and Rerewhakaaitu tephra. There is added complexity with these samples as we have both proximal and distal deposits to compare, where, as discussed previously, the proximal samples will likely be more heterogeneous. Glass analyses of the Waiohau tephra show it can be distinguished from those for the Rotorua and Rerewhakaaitu tephra using a range of trace elements and trace-element ratios. In addition, the Rotorua and Rerewhakaaitu tephra are observed to be bimodal for some elements. The Waiohau also has different mineralogy from that of Rotorua and Rerewhakaaitu tephra (Froggatt and Lowe, 1990; Lowe et al., 2008). Conversely, the Rotorua and Rerewhakaaitu tephra are usually indistinguishable in geochemistry and mineralogy, and therefore accurate dating and stratigraphic super-positioning would have to be relied upon to distinguish them (**Fig. 15f, Table 5**).



665 **Figure 13. Biplots to show examples of how trace elements in glass enable manipulation of heterogeneous and bimodal geochemical data. Panel (a) shows analyses of glass from Kaharoa and Taupō tephra, both of which show a heterogeneous signature with most major elements (presented on a normalised basis). Sr has a low variability for Taupō, but does not for Kaharoa tephra, conversely, Ba has a low variability for Kaharoa, but does not for Taupō. Panel (b) shows the bimodal signature created for Tahuna tephra using K₂O composition; this is also seen for Cs, but is not for Ba.**



670 **Figure 14. Results of Euclidean similarity coefficient (SC) calculations for major (a), trace (b), and trace ratio (c) concentrations in glass from all the tephras analysed. See Supplementary Material for R code used for these calculations. Colour coding shows SC values: green shows the smaller the value (similar compositions); through to red showing the larger the values (different compositions). The lowest values, and hence the most similar tephras compositionally, based on all geochemical data, are highlighted with bold outlines and text.**



675 **Figure 15.** Biplots for glass analyses for specific tephras which have very similar compositions and similar ages (see text for discussion
 and Table 4 for alternative elements). Plots show examples of the elements that enable these tephras to be separated (a) Poihipi and
 Tahuna (from TVC); (b) Mamaku and Rotoma (from OVC); (c) Rotoma and Rotoiti/Rotoehu (from OVC); (d) Waimihia and Unit
 K (from TVC); (e) Waimihia, Poronui and Karapiti – note that Poronui and Karapiti are indistinguishable using glass-chemistry;
 680 (f) Waiohau, Rerewhakaaitu and Rotorua – note that Rerewhakaaitu and Rotorua are indistinguishable using glass chemistry. All
 major element data presented on a normalised basis, and total iron is expressed as FeO.



685 KOT, Okaia, and Unit L (Mangaone Subgroup) show indistinguishable major elements in their constituent glass shards, and very similar trace elements. The TephraNZ samples have been compared to existing published data and are complementary in major elements (e.g. Sandiford et al., 2002; Shane et al., 2002; Smith et al., 2002; 2005; Lowe et al., 2008; Allan et al., 2008; Molloy, 2008; 20). This is the first time trace element glass data have been published for Unit L and Okaia tephtras. Our results show that Unit L glass shows bimodality in Rb/Zr, Ba/Th, Ce/Th and Y/Th and in this way it can be distinguished from the KOT and Okaia tephtras (**Table 5**).

690 4.3. Proposed future research

This foundation dataset, derived in a formalised way, is unique in New Zealand and provides researchers with new avenues of research. It is our hope that the foundation dataset can be improved and expanded with analyses of other known deposits, and that a subsidiary catalogue of accurately correlated geochemical samples can be added to bolster the dataset. As noted earlier, it is beyond the scope of this paper to dive too deeply into the detail of the data but we feel that it will provide the basis for countless projects in the future. Below we highlight some of the current gaps which we think would benefit from further research.

4.3.1 Further statistical analysis

700 We have applied simple ordination and statistical analyses to this dataset; however, we believe that further rigorous statistical analysis could be applied. Firstly, the analyses we present in this publication have been applied to mean values for each of the tephra samples (e.g. data from **Table 3**); there is no reason why these simple tests could not be applied to the full dataset, using all the individual values analysed for each sample. Secondly, for simplicity we chose to split up the assessment of major and trace elements, these could be run concurrently. Third, we chose very basic tests (PCA and ESC) to fit with our requirements, however there is likely some more appropriate statistical test that could be applied to get the most out of this exceptional dataset. For example, (extended) Canonical Variates Analysis (CVA); applying CVA to PCA results could determine optimal discrimination between multivariate data for single tephra deposits. This discrimination will increase the ability to identify an unknown tephra based on its similarity to known signatures plotted in multivariate space (e.g. discriminant function analysis; Tyron et al., 2009; 2010; Lowe et al., 2017; Bolton et al., 2020).



4.3.2 Whanganui Basin correlatives

A number of the tephras reported in this research were sampled from the Whanganui Basin, an
715 uplifted Plio-Pleistocene basin margin sequence that preserves as many as 45-superposed cyclothem
deposited since ~3 Ma (Naish and Kamp, 1997; Naish et al., 1996, 2005; Carter and Naish, 1998; Carter
et al., 1999; Pillans, 2017; Grant et al., 2018, 2019; Tapia et al., 2019). The tephra deposits within the
basin contribute to the robust chronological framework that has been constructed for this region
(Seward, 1976; Beu and Edwards, 1984; Alloway et al., 1993; Naish and Kamp, 1995; Shane et al.,
720 1996; Saul et al., 1999; Pillans et al., 1994, 2005; Naish et al., 1996, 2005; Rees et al., 2018, 2019).
These tephras also record a critical time in New Zealand's volcanological history – the transfer between
activity from the Coromandel Volcanic Zone to the Taupō Volcanic Zone (Briggs et al., 2005). Deposits
from this period are generally poorly exposed at source, and thus distal tephras could provide an insight
into the eruptive history, geochemical evolution, and potentially even caldera evolution during this
725 period (Houghton et al., 1995). Most of the tephras reported in this research are well known and well
dated, which is why they were included in the study. However, most do not have a known source
caldera or source eruptives, or have only been variably correlated to other deposits in New Zealand (e.g.
Lowe et al., 2001; Pearce et al., 2008). There are also numbers of tephra deposits in the Whanganui
Basin that have yet to be studied, and thus a research project that is tephra focused, rather than using it
730 as an accessory to a different line of enquiry, is timely.

4.3.3 IODP and ODP correlatives

At present there is a wealth of information that has yet to be fully investigated in the tephra
deposits in ODP Leg 181 Sites 1122, 1123, 1124, 1125 (Carter et al., 2003, 2004; Alloway et al., 2005;
735 Allan et al., 2008) and IODP Expedition 372 and 375 sites U1517 and U1520 (Pecher et al., 2018;
Saffer et al., 2018). Pioneering work includes that undertaken by Watkins and Huang (1977) and Nelson
et al. (1985) and findings from more 'local' marine coring expeditions include those reported by Shane
et al. (2006). The new reference material built by this project will allow more definitive identification
and correlation of tephras within these cores, specifically post-2 Ma. However, the reports currently
740 published on these deposits suggest that there are many more tephra deposits to be found in these
marine and offshore sites than we have in the TephraNZ dataset (Carter et al., 2003; Alloway et al.,
2005; Holt et al., 2010, 2011). The TephraNZ dataset can provide a formalised correlation framework
from which other unknown deposits can be determined, characterised, and integrated into a holistic
tephrostratigraphic reconstruction. Allan (2008) and Allan et al. (2008) reported the major and trace
745 element geochemistry of glass shards for tephra deposits dating from ~1.65 Ma in the ODP 1123 core.
They also give orbitally-tuned ages for these tephras. However, of the 38 identified tephras only seven
were correlated to onshore equivalents. In addition, Alloway et al. (2005) reported over 100 tephra



layers in the four ODP Leg 181 cores, dating back through orbital tuning (astrochronology) to 1.81 Ma. Using major element chemistry of glass, 13 tephra were correlated to equivalent onshore tephra including KOT, Omataroa, Rangitawa/Onepuhi, Kaukatea, Kidnappers-B and -A/Potaka, Unit D/Ahuroa, Ongatiti, Rewa, Sub-Rewa, Pakihikura, Ototoka and Table Flat. Analyses of glass from some of these are currently not in the TephraNZ database but could be easily added if the appropriate reference samples were available and capacity to analyse them were available. Alloway et al. (2005) reported an additional six tephra deposits that are correlated between the cores, but not to onshore equivalents, leaving potentially ~81 tephra horizons within the ODP cores that are uncorrelated. This information could provide a detailed investigation into the timing and evolution of the TVZ eruptions that is unobtainable from onshore deposits.

4.3.4 Mineral compositions

The TephraNZ reference dataset is only populated by glass major and trace element analyses at present. This is because glass geochemistry is one of the most frequently used and accessible tools for tephra correlation. Aerodynamic sorting of tephra componentry through transportation adds to the favourability of glass shards as the dominant tool because glass shards tend to be the only phase that is found at both proximal and distal sites. However, previous New Zealand-based studies have specified how mineral assemblages and their geochemical compositions can be used to distinguish certain tephra and their source (e.g. Nairn and Kohn, 1973; Lowe, 1988; Froggatt and Lowe, 1990; Froggatt and Rogers, 1990; Shane, 1998; Shane et al., 2003b; Allan et al., 2008; Lowe et al., 2008; Lowe, 2011). For example, the mineral cummingtonite, where predominant, is a known identifier for tephra from the Haroharo complex of the OVC (Whakatane, Rotoma, Rotoehu/Rotoiti (**Table 4**); Ewart, 1968; Lowe, 1988; Froggatt and Lowe, 1990). At present, ferromagnesian mineralogical assemblages (following Froggatt and Lowe, 1990; Smith et al., 2005; Lowe et al., 2008) for all the TephraNZ samples younger than and including Rotoehu/Rotoiti have been published (see Table 4). Extending this tabulation to include the older samples would add another useful criterion to the correlation toolbox.

Additionally, the fractional crystallisation of plagioclase, biotite, amphibole, zircon, hydrous mineral phases, or Fe-Ti oxides has been shown to be the key impactor on the trace element chemistry (Shane, 1998; Allan, 2008; Turner et al., 2009, 2011). Thus the prevalence of these minerals is also an important potential fingerprinting tool. The information on the mineralogy of the tephra is not only useful for fingerprinting but also can be used in determining the characteristics of the magma source components, and potentially provide estimates for the temperature, pressure, and oxidation states of the magmatic system before eruption (e.g. Lowe, 1988, 2011; Shane, 1998). Thus, this information can allow hypotheses to be developed on the reactivation and triggering of these large-scale eruptions, an important step for hazard and risk monitoring.



4.3.5 The New Zealand tephra “Bermuda Triangle”

785 At present the TephraNZ database is very well populated for samples from the Rotoiti/Rotoehu
through to Kaharoa eruption. It also has a high number of samples, but not an exhaustive list, from
Mamaku ignimbrite (~0.22-0.23 ka) to the Hikuroa Pumice (2 Ma). There is a stark deficit in tephtras
between the Rotoiti/Rotoehu eruption and Mamaku ignimbrite (**Table 1**). This ~150 kyr gap in the
790 volcanic record (~220 ka to 45 ka) is intriguing as there is proximal evidence for activity during this
period. For example, Rosenberg et al. (2020) reported the occurrence of volcanic formations in cores
from the Taupō region in the age range of ~168 to 92 ka, including the Huka Falls formations,
Racetrack rhyolites, and the Te Mihi rhyolites. Tephra deposits, in some cases strongly weathered
successions of multiple units broadly lumped together as a ‘formation’, such as the so-called Hamilton
Ash Formation, have been reported during this time period both terrestrially and in marine and
795 lacustrine sediment cores (Ward, 1967; Pain, 1975; Vucetich et al., 1978; Iso et al., 1982; Froggatt,
1983; Manning, 1996; Lowe et al., 2001; Newnham et al., 2004; Allan et al., 2008; Briggs et al., 2006;
Lowe, 2019; B. Laeuchli *pers. comms.* 2020). However, at present the authors are not aware of a
detailed, up-to-date study into the primary compositions of these tephra deposits. The key deposits
identified during this time period include (but are not limited to) Kaingaroa Ignimbrite (~0.18 Ma;
800 Froggatt, 1983), Tablelands Tephra Formation (~0.21–0.18 Ma; Iso et al., 1982, 0.34-0.39 Ma;
Manning, 1996), Hamilton Ash Formation (0.125–0.34 Ma; Lowe, 2019); Kutarere tephra (= Mamaku
ignimbrite 0.22-0.23 Ma; Shane et al., 1994; Houghton et al., 1995; Black et al., 1996; Tanaka et al.,
1996; Milner et al., 2003), Kukumoa Subgroup (~0.22-0.05 Ma; Manning, 1996), and Tikotiko Ash
(~0.125 ka; Lowe, 2019). A number of these studies are outdated, and with improved methodologies
805 (major and trace element analysis, potentially of melt inclusions where preserved, dating techniques,
and other measures to help construct time frames such as via phytolith studies to determine glacial vs
interglacial periods) it could be timely to further investigate this period of (apparent) deficit.

5. Conclusions

Major and trace element geochemical compositions of glass shards for a large suite of
810 prominent, widespread New Zealand rhyolitic tephtras have been analysed systematically and published
for the first time as “TephraNZ”. TephraNZ is a foundation dataset for collating geochemical data about
New Zealand tephtras. The foundation reference dataset is made up of known deposits that have their
ages quantified through independent methods, or are from the type sites where tephtras were first
defined, or well-documented reference sections. Detailed methodology is reported to allow subsequent
815 research to acquire comparable data to those in this database. Principal component analysis indicates



that for the TephraNZ database, as a whole, major elements CaO, TiO₂, K₂O, and FeO_t are responsible for the variability in PC1 and PC2 space. For trace elements, Tb, Ho, Dy, Y, Sm, Nd, Tm, Yb, Hf, Gd and Rb, Th, Sr, U, Cs, Ta, Pb, La, Ce, Pr are responsible for ~ 69% of the variability, and trace element ratios, Ba/Zr, Ba/Hf, Ba/Eu, Zr, Nb, Rb/Zr, Ba/Y, Zr/Th, Zr/ Yb, Nb/Y, Zr/Y, and Ba/Th, Ce/Th Y/Th, 820 Ba/La, Ba/Ce, Zr/Th and Sr/Nb are responsible for ~73 % of the variance. Euclidean similarity coefficients can also be used to distinguish between some geochemically similar glass analyses. However, further detailed geochemical investigation is required to distinguish others. Geochemically indistinguishable tephtras (on the basis of both major and trace element glass-shard compositions) are identified as Taupō and Waimihia; Poronui and Karapiti; Rotorua and Rerewhakaaitu; and KOT and 825 Okaia. Only Poronui and Karapiti are noted as entirely indistinguishable, with other methods of characterisation listed as alternative options, including mineralogy, age, and stratigraphic relationships.

6. Author contribution

JLH and RJW designed the project. DJL and BJP contributed samples from previous field campaigns, and DJL provided guidance on new and existing field locations for sample collection. JLH and JEB 830 undertook the field work, lab work, analysis and data reduction. ABHR advised on statistical analysis and R-coding. LA supervised and helped JEB develop LA-ICP-MS analysis and data reduction. FT supervised and helped JEB develop sample mounting and polishing procedures. JLH wrote the manuscript with contributions from all co-authors.

7. Competing interest

835 The authors declare that they have no conflict of interest.

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Table 1. Overview of the foundation tephra included in TephraNZ database

Tephra name	Alternative name(s)	Caldera source	Age (cal. yr BP, 2sd) (unless otherwise shown)	Dating method*	Age reference	Proximal/d istal	Easting	Southing	Site number (Fig. 1)	Site description (Fig. 1)
Kaharoa		Okataina	636 +/- 12	C14 wiggle match	1	P	176°31'81.7"	38°18'56.8"	1	Ash Pit Road
Taupō	Unit Y	Taupo	1718 +/- 10	C14 wiggle match	2, 3	P	176°11'58.5"	38°44'50.5"	2	SH 5
Waimihia	Unit S	Taupo	3382 +/- 50	C14 Bayes model 1	4	D	177°18'24.5"	38°68'35.8"	3	Kaipō Bog
Unit K (Taupō series)		Taupo	5088 +/- 73	C14 Bayes model 1	4	D	177°18'24.5"	38°68'35.8"	3	Kaipō Bog
Whakatane		Okataina	5542 +/- 48	C14 Bayes model 1	4	D	177°18'24.5"	38°68'35.8"	3	Kaipō Bog
Tuhua		Mayor Island	7637 +/- 100	C14 Bayes model 2	5	D	177°18'24.5"	38°68'35.8"	3	Kaipō Bog
Mamaku		Okataina	7992 +/- 58	C14 Bayes model 1	4	D	177°18'24.5"	38°68'35.8"	3	Kaipō Bog
Rotoma		Okataina	9472 +/- 40	C14 Bayes model 1	4	P	176°28'82.3"	38°17'90.5"	1	Ash Pit Road
Opepe	Unit E	Taupo	10,004 +/- 122	C14 Bayes model 1	4	D	177°18'24.5"	38°68'35.8"	3	Kaipō Bog
Poronui	Unit C	Taupo	11,195 +/- 51	C14 Bayes model 1	4	D	177°18'24.5"	38°68'35.8"	3	Kaipō Bog
Karapiti	Unit B	Taupo	11,501 +/- 104	C14 Bayes model 1	4	D	177°18'24.5"	38°68'35.8"	3	Kaipō Bog
Waiohau		Okataina	14,018 +/- 91	C14 Bayes model 1	4	P	176°19'29.3"	38°10'11.8"	4	RNL pumice quarry
Rotorua		Okataina	15,738 +/- 263	C14 Bayes model 1	4	D	177°18'24.5"	38°68'35.8"	3	Kaipō Bog
Rerewhakaaitu		Okataina	17,209 +/- 249	C14 Bayes model 1	4	D	177°18'24.5"	38°68'35.8"	3	Kaipō Bog
Okareka		Okataina	21,858 +/- 290	C14 Bayes model 3	4	P	176°19'29.3"	38°10'11.8"	4	RNL pumice quarry
Te Rere		Okataina	25,171 +/- 964	C14 Bayes model 3	4	P	176°19'29.3"	38°10'11.8"	4	RNL pumice quarry
Kawakawa/Oruanui		Taupo	25,358 +/- 162	C14 Bayes model 3	6	P	175°53'31.2"	38°37'19.2"	5	Poihipi Road
Poihipi		Taupo	28,446 +/- 670	C14 Bayes model 3	4	P	176°10'50.5"	38°02'36.2"	6	Oturoa Road
Ohaia		Taupo	28,621 +/- 1428	C14 Bayes model 3	4	P	175°53'31.4"	38°37'12.9"	5	Poihipi Road
Unit L (Mangaone Sgp series)		Okataina	30,900 +/- 1500	Bayes model	9	P	176°08'22.2"	38°05'39.5"	7	Dansey Road
Awakeri	Unit J	Okataina	30,900 +/- 560	C14 + ZDD	9	P	176°43'16.9"	38°01'43.9"	8	Bowditch Quarry SH30
Mangaone	Unit I	Okataina	31,500 +/- 520	C14 + ZDD	9	P	176°43'16.9"	38°01'43.9"	8	Bowditch Quarry SH30
Hauparu	Unit F	Okataina	36,165 +/- 450	C14	8	P	176°14'43.9"	37°46'13.1"	9	Little Waihi Road, Maketu
Maketu	Unit D	Okataina	36,100 +/- 440	C14 + ZDD	9	P	176°14'43.9"	37°46'13.1"	9	Little Waihi Road, Maketu
Tahuna		Taupo	38,400 +/- 1700	Bayes model	9	P	176°44'56.7"	37°56'58.1"	10	Braemar Road
Ngamotu	Unit B	Okataina	39.0 ka	7	P	176°44'56.7"	37°56'58.1"	10	Braemar Road	
Earthquake Flat Ig		Kapenga	45,160 +/- 2900	(U-Th)/He	10	P	176°15'02.9"	38°16'52.9"	13	Tumunui Road
Rotoehu Ash		Okataina	45,170 +/- 3300	(U-Th)/He	10	P	175°53'04.2"	37°59'47.5"	11	Tapapa Road
Rotoiti Ignimbrite		Okataina	45,170 +/- 3300	(U-Th)/He	10	P	176°43'33.8"	37°52'42.2"	12	Mimihi Stream, SH2
Ararata Gully 318	? Mamaku Ig	? Rotorua	0.235 Ma	MIS7c/b boundary	11	D	174°21'26.9"	39°33'17.2"	14	Ararata Gully
Kakariki 272	< Rangitawa = ? Mamaku	0.235 - 0.3 Ma	Stratigraphy	12, 13	D	175°44'51.0"	40°09'52.4"	15	Rangitikei Valley	
Fordell 449	? Whakamaru Ig	? Whakamaru	0.31 Ma	MIS9a	13	D	175°15'17.8"	39°56'33.3"	16	Kauangaroa Road
Rangitawa	Mt Curl, Mangaroa, Ohariu, Haywards ash, Lower Finnis Road, Ohinewai Tephra, Hamilton Ash H1	Whakamaru	345 +/- 12 ka (1sd)	glass-4PFPT	14	D	175°19'01.9"	37°47'26.5"	17	Univ of Waikato, Hamilton
Upper Griffin Road 307	? Whakamaru Ig	? Whakamaru	0.3 - 0.4 Ma	Stratigraphy	13	D	175°13'18.4"	39°58'02.4"	16	Kauangaroa Road
Lower Griffin Road 309	? Whakamaru Ig	? Whakamaru	0.3 - 0.4 Ma	Stratigraphy	13	D	175°13'18.4"	39°58'02.4"	16	Kauangaroa Road
Onepuhi 267		Unknown	0.57 Ma	Astronomical	15	D	175°28'38.9"	40°04'32.2"	15	Rangitikei Valley
Kupe 481		Unknown	0.63 +/- 0.08 Ma (1sd)	glass-4PFPT	15, 16	D	175°18'09.5"	40°00'12.9"	18	Turakina
Kauakatea 232		Unknown	0.86 +/- 0.08 Ma (1sd)	glass-4PFPT	15, 16	D	175°06'09.5"	39°54'07.2"	19	Wanganui River
Potaka 305	Kidnappers	Mangakino	1.00 +/- 0.05 (1sd)	40Ar/39Ar	17	D	175°38'36.0"	39°59'49.6"	15	Rangitikei Valley
Rewa 304	Ahuroa?	? Mangakino	1.20 +/- 0.14 Ma (1sd)	glass-4PFPT	15, 16	D	175°35'04.5"	39°59'31.8"	15	Rangitikei Valley
Mangapipi 510	? Ig B	Mangakino	1.51 +/- 0.16 Ma (1sd)	glass-4PFPT	15, 16	D	175°20'35.2"	39°55'14.7"	18	Turakina
Pakihikura 303	? Ngaroma	Mangakino	1.58 +/- 0.16 Ma (1sd)	glass-4PFPT	15	D	175°21'16.8"	39°54'57.6"	18	Turakina
Birdgrove 511		Unknown	1.6 Ma	Astronomical	15	D	175°21'20.9"	39°54'54.3"	18	Turakina
Mangahou 302		Unknown	1.63 Ma	Astronomical	15	D	175°38'42.1"	39°57'00.8"	15	Rangitikei Valley
Ototoka 521		Unknown	1.72 +/- 0.32 Ma (1sd)	glass-4PFPT	15	D	174°50'22.9"	39°52'10.6"	19	Ototoka Beach
Hikuroa Pumice member		Unknown	2.0 +/- 0.6 Ma	zircon-4PFPT	18	D	176°49'36.8"	39°14'37.2"	20	Darkys Spur

Age references

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Dating Methods

- Bayesian model 1 = OxCal P_sequence
 Bayesian model 2 = OxCal P_sequence Isochron informed
 Bayesian model 3 = OxCal Tau-boundary



Table 2. Set up for electron microprobe analysis

Channel 1			Channel 2			Channel 3			Channel 4			Channel 5		
Element	Mineral	Time												
Ca	VG-A99	30/15	Na*	VG-568	10/5	Si	VG-568	30/15	K	VG-568	30/15	Cl	VG-568	30/15
Ti	TiO2	30/15	Mg	VG-A99	30/15	Al	VG-568	30/15	Fe	VG-A99	30/15	Mn	MnO	30/15

* Na is run twice, on the second run the peak search is skipped to reduced the volatilisation of the element



Table 3. Average and standard deviation for all tephra, see SM Table 1 for full dataset.

Tephra name	SiO ₂ (wt%)	TiO ₂ (wt%)	Al ₂ O ₃ (wt%)	FeO _T (wt%)	MnO (wt%)	MgO (wt%)	CaO (wt%)	Na ₂ O (wt%)	K ₂ O (wt%)	Cl (wt%)	H ₂ O*	SiO ₂ /K ₂ O	Na ₂ O + K ₂ O
Kaharoa	Average 77.8	0.09	12.6	0.92	0.07	0.10	0.63	3.75	3.98	0.17	3.66	19.5	7.73
n=25 (18)	0.41	0.05	0.20	0.35	0.03	0.18	0.18	0.18	0.39	0.03	1.45	1.06	0.56
Taupo Y5	Average 75.6	0.25	13.4	1.83	0.10	0.19	1.41	4.25	2.99	0.16	1.39	25.3	7.11
n=25 (20)	0.67	0.07	0.45	0.53	0.04	0.08	0.13	0.30	0.16	0.05	4.28	1.65	0.34
Waimihia K3	Average 76.2	0.21	13.2	1.83	0.10	0.15	1.33	4.11	2.92	0.18	2.61	26.1	7.02
n=22 (20)	0.56	0.03	0.24	0.19	0.03	0.02	0.13	0.26	0.14	0.02	1.53	3.86	0.40
Unit K	Average 76.3	0.20	13.1	1.77	0.08	0.14	1.31	4.11	2.99	0.15	1.56	25.5	7.10
n=24 (20)	0.29	0.02	0.13	0.13	0.02	0.03	0.11	0.10	0.10	0.03	1.71	2.87	0.20
Whakatane K5	Average 78.3	0.11	12.5	0.84	0.04	0.10	0.69	3.58	3.85	0.16	1.97	20.4	7.43
n=21	0.18	0.02	0.08	0.05	0.02	0.01	0.04	0.13	0.09	0.02	1.49	2.0	0.23
Whakatane - P	Average 77.5	0.17	12.7	1.43	0.06	0.12	1.02	3.68	3.29	0.16	2.10	23.5	6.97
n=25 (21)	0.78	0.04	0.23	0.35	0.02	0.03	0.21	0.51	0.36	0.02	2.18	2.20	0.86
Tuhua K6	Average 74.4	0.30	9.7	5.80	0.12	0.01	0.27	5.18	4.19	0.23	0.95	17.8	9.37
n=23 (22)	0.31	0.02	0.14	0.15	0.02	0.01	0.02	0.18	0.16	0.01	0.52	1.98	0.33
Mamaku K7	Average 78.3	0.12	12.6	0.88	0.04	0.11	0.79	3.61	3.59	0.16	2.14	21.8	7.20
n=23 (23)	0.18	0.02	0.15	0.05	0.02	0.02	0.06	0.11	0.13	0.01	1.54	1.33	0.25
Rotoma - P	Average 78.2	0.11	12.5	0.89	0.04	0.11	0.73	3.66	3.74	0.15	0.68	20.9	7.40
n=24	0.17	0.01	0.09	0.08	0.02	0.01	0.03	0.10	0.14	0.01	0.66	1.2	0.24
Rotoma - K8 - D	Average 78.2	0.12	12.6	0.87	0.05	0.13	0.84	3.73	3.45	0.15	0.83	22.7	7.19
n=23 (20)	0.18	0.01	0.08	0.05	0.01	0.02	0.07	0.13	0.24	0.01	0.75	0.75	0.37
Opepe K9	Average 76.0	0.23	13.4	1.80	0.07	0.17	1.74	3.77	2.76	0.16	4.02	27.6	6.53
n=14 (8)	0.43	0.05	0.16	0.19	0.03	0.04	0.20	0.16	0.30	0.03	1.50	1.42	0.46
Poronui K10	Average 76.4	0.21	13.2	1.64	0.07	0.16	1.53	3.75	3.02	0.14	1.46	25.3	6.77
n=18 (15)	0.49	0.08	0.21	0.22	0.03	0.04	0.14	0.11	0.16	0.02	1.47	3.15	0.27
Karapiti K11	Average 76.6	0.19	13.1	1.58	0.06	0.14	1.48	3.67	3.16	0.14	1.84	24.3	6.82
n=25 (11)	0.26	0.02	0.11	0.16	0.02	0.02	0.14	0.18	0.36	0.02	1.27	0.72	0.54
Waiohau - P	Average 78.0	0.12	12.5	0.96	0.06	0.14	0.84	3.89	3.31	0.14	3.01	23.6	7.20
n=19 (19)	0.27	0.02	0.26	0.13	0.01	0.03	0.07	0.19	0.14	0.03	2.33	1.06	0.28
Waiohau - K14b D	Average 78.1	0.12	12.4	1.03	0.05	0.16	0.88	3.87	3.23	0.14	1.02	24.2	7.10
n=10 (10)	0.26	0.02	0.17	0.05	0.02	0.01	0.03	0.26	0.09	0.02	1.02	0.70	0.27
Rotorua - P	Average 77.6	0.14	12.5	1.15	0.05	0.14	1.00	3.75	3.51	0.15	4.58	22.5	7.26
n=29 (28)	0.57	0.06	0.28	0.21	0.02	0.07	0.33	0.31	0.47	0.04	2.18	2.79	0.45
Rotorua - K15 D	Average 77.5	0.08	12.5	0.91	0.06	0.09	0.73	3.80	4.13	0.14	0.64	19.0	7.93
n=10 (10)	0.35	0.05	0.19	0.15	0.01	0.07	0.21	0.23	0.43	0.02	0.63	2.47	0.34
Rerewhakaaitu - P	Average 78.0	0.09	12.3	0.98	0.05	0.10	0.77	3.61	3.84	0.15	3.06	20.5	7.45
n=21 (19)	0.02	0.20	0.10	0.29	0.03	0.10	0.26	0.39	0.03	0.03	1.97	2.31	0.28
Rerewhakaaitu - K17 D	Average 77.7	0.08	12.6	0.94	0.05	0.10	0.79	3.71	3.94	0.15	3.31	19.9	7.65
n=11 (10)	0.30	0.02	0.29	0.07	0.02	0.06	0.29	0.29	0.36	0.02	1.98	1.99	0.43
Okareka	Average 78.5	0.14	12.6	0.87	0.06	0.12	0.91	3.57	3.24	0.20	4.93	24.2	6.81
n=22 (22)	0.20	0.02	0.11	0.06	0.02	0.02	0.11	0.18	0.02	0.11	0.73	1.09	0.29
Te Rere	Average 78.1	0.09	12.4	1.05	0.06	0.06	0.64	3.62	3.97	0.22	3.80	19.7	7.59
n=24 (20)	0.18	0.02	0.10	0.08	0.03	0.02	0.09	0.16	0.24	0.02	1.18	0.73	0.41
Kawakawa/Oruanui	Average 77.9	0.17	12.8	1.27	0.06	0.14	1.24	3.34	3.14	0.17	5.18	24.8	6.48
n=25 (23)	0.82	0.05	0.32	0.15	0.03	0.04	0.21	0.60	0.34	0.02	1.01	2.42	0.94
Pohiipi	Average 78.1	0.12	12.6	0.96	0.07	0.11	0.90	3.52	3.65	0.50	5.71	21.4	7.17
n=20 (18)	0.30	0.02	0.17	0.17	0.03	0.08	0.11	0.39	0.41	1.55	16.69	0.73	0.80
Okaia	Average 77.8	0.18	12.8	1.31	0.07	0.15	1.28	3.43	3.04	0.18	5.34	25.6	6.47
n=23 (23)	0.64	0.06	0.26	0.17	0.03	0.05	0.18	0.24	0.26	0.06	1.89	2.45	0.50
Unit L	Average 77.8	0.15	12.7	1.18	0.06	0.14	1.16	3.62	3.16	0.22	4.48	24.6	6.78
n=23 (20)	0.34	0.03	0.28	0.08	0.02	0.03	0.13	0.32	0.41	0.22	6.31	0.83	0.72
Awakeri	Average 77.4	0.18	13.0	1.16	0.07	0.18	1.07	3.99	2.97	0.15	3.57	26.1	6.95
n=21 (19)	0.22	0.01	0.16	0.11	0.02	0.02	0.07	0.13	0.07	0.12	1.29	3.08	0.20
Mangaone	Average 76.7	0.20	13.3	1.25	0.08	0.22	1.20	4.14	2.86	0.18	2.56	26.8	7.00
n=24 (19)	0.38	0.02	0.18	0.08	0.02	0.02	0.08	0.16	0.08	0.01	1.32	4.83	0.24
Hauparu	Average 75.2	0.38	13.7	1.86	0.08	0.39	1.78	3.87	2.73	0.15	3.05	27.6	6.60
n=21 (12)	0.90	0.05	0.39	0.26	0.02	0.06	0.21	0.16	0.15	0.02	1.57	6.14	0.31
Maketu	Average 73.1	0.49	14.4	2.32	0.10	0.56	2.12	4.49	2.41	0.16	4.29	30.4	6.90
n=3 (1)	0.26	0.05	0.20	0.26	0.01	0.06	0.14	0.35	0.13	0.03	6.25		
Ngamotu	Average 75.7	0.34	13.7	1.54	0.09	0.36	1.73	3.92	2.62	0.15	3.93	28.8	6.54
n=21 (19)	1.08	0.09	0.51	0.34	0.04	0.15	0.37	0.24	0.22	0.02	2.90	5.03	0.46
Tahuna	Average 78.2	0.13	12.5	0.94	0.04	0.12	0.86	3.49	3.75	0.18	3.84	20.8	7.24
n=23 (17)	0.31	0.02	0.19	0.07	0.02	0.03	0.10	0.28	0.41	0.02	0.69	0.76	0.69
Earthquake Flat Ig	Average 77.9	0.11	12.5	0.95	0.03	0.10	0.79	3.23	4.43	0.21	3.83	17.6	7.65
n=24 (11)	0.20	0.02	0.08	0.09	0.01	0.01	0.03	0.11	0.11	0.03	0.71	1.76	0.22
Rotoehu tephra	Average 78.3	0.14	12.6	0.91	0.05	0.13	0.83	3.66	3.42	0.16	3.57	22.9	7.08
n=21 (14)	0.18	0.02	0.06	0.05	0.02	0.01	0.03	0.14	0.08	0.01	0.80	2.28	0.22
Rototi Ig	Average 78.3	0.14	12.6	0.87	0.05	0.14	0.86	3.72	3.35	0.21	6.00	23.4	7.08
n=21 (19)	0.16	0.02	0.13	0.12	0.02	0.02	0.05	0.22	0.11	0.08	4.52	1.44	0.34
Ararata Gully 318	Average 77.2	0.15	12.8	1.29	0.03	0.13	1.03	3.64	3.65	0.17	5.61	21.2	7.29
n=24 (15)	0.70	0.04	0.30	0.19	0.01	0.05	0.17	0.09	0.09	0.02	0.79	7.60	0.18
Kakariki 272	Average 78.1	0.13	12.5	0.98	0.02	0.11	0.91	3.50	3.74	0.16	4.99	20.9	7.24
n=25 (18)	0.13	0.02	0.07	0.10	0.02	0.02	0.04	0.09	0.14	0.01	0.38	0.87	0.24
Fordell 449	Average 78.0	0.11	12.4	1.07	0.02	0.07	0.64	3.74	3.97	0.20	5.16	19.7	7.70
n=25 (19)	0.44	0.04	0.19	0.25	0.01	0.06	0.16	0.13	0.18	0.02	0.59	2.42	0.31
Upper Griffin Road 307	Average 77.5	0.17	12.5	1.44	0.03	0.11	1.01	3.69	3.47	0.19	5.46	22.4	7.16
n=21 (17)	0.19	0.02	0.09	0.10	0.02	0.02	0.04	0.11	0.10	0.02	0.47	1.93	0.21
Lower Griffin Road 309	Average 76.5	0.28	13.2	1.30	0.03	0.28	1.39	3.72	3.32	0.17	5.14	23.1	7.04
n=25 (21)	0.62	0.05	0.31	0.12	0.01	0.07	0.20	0.15	0.20	0.02	0.45	3.10	0.35
Onepuhi 267	Average 77.1	0.14	12.8	1.32	0.03	0.09	0.96	3.49	4.04	0.24	5.20	19.1	7.53
n=23 (10)	1.08	0.05	0.45	0.41	0.02								



Table 3 (cont.) Average and standard deviation for all tephra, see SM Table 1 for full dataset.

Tephra name	Sc (ppm)	Ti (ppm)	V (ppm)	Mn (ppm)	Co (ppm)	Cu (ppm)	Zn (ppm)	Ga (ppm)	Rb (ppm)	Sr ⁸⁶ (ppm)	Sr ⁸⁸ (ppm)	Y (ppm)	Zr ⁹⁰ (ppm)
Kaharoa	Average 11.5	528	1.80	453	0.28	1.49	40.3	14.4	133	50.3	45.6	33.0	961
n=25 (18)	2sd 6.36	165	1.00	84.9	0.17	1.22	11.1	3.11	37.7	15.3	20.2	29.8	3753
Taupo Y5	Average 12.6	1175	1.27	531	0.67	2.85	49.3	13.7	110	112	111	24.2	162
n=25 (20)	2sd 3.65	448	0.87	210	0.72	4.08	15.3	3.41	30.0	31.3	30.2	7.16	59.8
Waimihia K3	Average 14.3	1189	3.03	731	0.43	7.11	71.8	18.8	102	112	113	30.7	211
n=22 (20)	2sd 2.14	239	9.89	151	0.12	17.09	23.9	7.10	18.4	23.6	18.6	6.71	30.9
Unit K	Average 7.19	1188	0.78	595	0.67	5.37	58.4	17.1	110	119	118	32.8	218
n=24 (20)	2sd 0.92	144	0.16	64.9	0.12	9.76	14.9	1.99	14.9	12.8	11.8	3.81	22.5
Whakatane K5	Average												
n=21	2sd												
Whakatane - P	Average 6.57	985	1.01	528	0.55	2.67	41.6	17.3	123	80.0	81.9	31.5	173
n=25 (21)	2sd 1.90	262	0.72	108	0.20	1.63	19.3	2.99	25.4	26.1	24.3	8.39	67.9
Tuhua K6	Average 3.40	1677	0.71	1091	0.57	5.43	215	44.1	150	21.1	11.7	131	1171
n=23 (22)	2sd 0.64	449	0.44	318	0.83	4.28	99.9	13.66	25.0	27.0	22.4	48.3	481
Mamaku K7	Average 4.39	755	1.13	436	0.49	2.76	28.1	15.1	127	68.8	65.0	25.9	99.4
n=23 (23)	2sd 0.80	133	0.26	64.9	0.10	1.95	7.3	2.23	20.0	11.9	10.2	3.49	17.0
Rotoma - P	Average												
n=24	2sd												
Rotoma - K8 - D	Average 4.08	776	1.27	464	0.45	1.25	37.9	14.0	115	71.2	71.6	27.1	96.9
n=23 (20)	2sd 0.44	79	0.27	37.4	0.11	0.42	7.2	1.19	13.3	8.21	6.32	2.40	9.71
Opepe K9	Average 8.02	1364	4.01	529	1.70	6.13	43.7	18.6	123	126	130	33.0	234
n=14 (8)	2sd 0.31	131	2.01	66.4	0.21	3.92	13.8	1.76	11.7	10.9	12.9	3.00	20.7
Poronui K10	Average 8.01	1297	2.87	463	1.62	4.23	46.9	17.1	126	118	118	32.9	230
n=18 (15)	2sd 1.51	253	1.00	92.3	0.34	1.96	15.4	2.71	23.5	20.4	19.3	6.59	33.8
Karapiti K11	Average 7.05	1162	2.60	395	1.57	2.93	44.1	16.3	109	109	109	112	30.6
n=25 (11)	2sd 1.32	242	0.84	79.3	0.41	2.02	11.0	2.80	18.9	14.2	14.6	5.49	38.7
Waiohau - P	Average 13.4	652	1.35	443	0.12	0.73	39.1	14.2	110	68.7	69.3	21.5	83.6
n=19 (19)	2sd 2.47	56	1.22	14.1	0.45	0.71	6.3	2.04	5.2	5.43	4.87	1.62	7.95
Waiohau - K14b D	Average 17.0	760	1.88	406	0.40	1.31	39.3	12.5	112	77.9	77.9	21.5	93.4
n=10 (10)	2sd 1.94	28	0.81	16.7	0.37	0.80	8.1	0.88	2.9	4.78	2.40	0.90	1.53
Rotorua - P	Average 13.8	903	2.89	424	0.53	1.19	53.6	14.3	118	74.4	76.1	23.6	127
n=29 (28)	2sd 2.83	356	2.20	65.6	0.45	1.00	26.8	2.04	18.0	34.0	33.3	5.63	37.7
Rotorua - K15 D	Average 17.9	470	1.79	483	0.44	2.63	30.7	13.7	146	76.5	73.7	20.9	74.3
n=10 (10)	2sd 2.32	221	1.23	45.4	0.36	1.89	9.5	2.65	20.3	92.5	86.4	3.38	32.1
Rerewhakaaitu - P	Average 13.9	720	2.51	386	0.29	2.49	35.3	13.3	134	63.3	64.1	22.1	82.9
n=21 (19)	2sd 2.64	446	3.30	67.7	0.67	3.41	9.6	2.83	18.1	43.8	43.7	6.97	20.5
Rerewhakaaitu - KJ	Average 18.5	499	2.54	481	0.56	2.05	34.0	13.9	142	60.3	58.1	18.9	75.3
n=11 (10)	2sd 4.02	119	3.69	51.4	0.51	1.24	9.3	1.30	16.9	13.1	12.1	2.32	12.3
Olarereia	Average 5.21	895	2.72	454	0.80	5.11	24.9	14.9	113	70.8	68.7	22.4	100
n=22 (22)	2sd 0.97	100	0.49	44.3	0.18	0.57	7.5	1.65	11.7	8.45	7.36	2.60	11.4
Te Rere	Average 5.86	698	1.56	367	0.75	3.14	41.5	17.2	139	34.7	34.2	37.9	123
n=24 (20)	2sd 1.68	150	2.05	62.0	0.69	1.83	10.8	2.71	28.2	9.56	8.81	6.85	18.6
Kawakawa/Oruan	Average 4.92	1003	3.33	404	1.27	5.73	32.0	16.6	124	95.5	94.9	23.1	145
n=25 (23)	2sd 1.36	222	3.09	76.6	0.51	7.60	9.9	5.71	15.8	18.1	17.3	3.16	31.7
Poihipi	Average 4.36	915	3.38	438	0.73	3.20	32.5	17.7	146	75.2	72.2	25.8	111
n=20 (18)	2sd 1.17	369	6.28	82.0	0.53	2.58	9.5	9.58	50.7	13.8	14.0	5.47	28.3
Okaiia	Average 5.21	1114	3.82	417	1.40	4.53	30.3	15.5	132	106	102	24.3	152
n=23 (23)	2sd 0.88	333	2.44	72.5	0.70	2.78	10.2	2.32	19.9	23.7	19.9	3.92	33.1
Unit L	Average 4.96	1061	2.19	469	0.90	3.94	29.6	15.3	116	104	100	27.9	148
n=23 (20)	2sd 0.81	167	0.68	121	0.36	2.22	9.4	1.87	19.7	18.0	17.8	7.78	15.0
Awakeri	Average 6.23	1216	1.48	630	0.40	2.18	45.8	17.7	103	107	107	35.3	163
n=21 (19)	2sd 1.52	297	0.48	156	0.12	1.18	16.5	3.85	24.3	15.7	15.6	7.89	37.1
Mangaone	Average 4.96	1233	1.85	669	0.43	2.08	33.3	16.1	88.0	114	118	32.9	173
n=24 (19)	2sd 0.48	94	0.23	58.7	0.07	1.34	10.4	1.61	7.1	9.37	10.6	2.78	16.2
Hauparu	Average 6.57	2254	12.1	637	1.73	3.40	40.4	15.8	82.0	155	153	30.3	239
n=21 (12)	2sd 1.11	240	1.87	64.2	0.38	1.82	18.7	2.06	11.5	29.4	28.4	3.37	28.5
Maketu	Average 8.80	2890	12.6	940	1.84	2.96	29.5	19.6	79.0	188	196	42.4	298
n=3 (1)	2sd												
Ngamotu	Average 4.70	2065	16.8	646	3.21	6.27	42.5	16.0	72.0	193	189	23.5	179
n=21 (19)	2sd 0.63	796	26.7	87.9	6.03	11.8	14.9	2.97	17.1	107	107	5.39	29.6
Tahuna	Average 4.05	895	2.61	442	1.03	3.62	25.9	18.2	136	61.4	61.5	21.0	102
n=23 (17)	2sd 0.61	146	0.91	86.7	0.35	1.65	7.4	7.01	38.1	9.56	10.4	3.13	18.3
Earthquake Flat Ig	Average 4.14	884	2.40	490	0.84	2.14	19.6	15.8	122	67.4	64.9	20.5	93.1
n=24 (11)	2sd 0.43	66	0.50	38.2	0.17	1.01	3.0	1.45	30.6	7.85	5.56	2.10	7.87
Rotoehu tephra	Average 4.33	1109	3.84	536	0.94	2.75	32.4	14.9	100	81.6	82.4	22.3	113
n=21 (14)	2sd 0.62	564	3.50	141	0.30	0.99	9.5	2.15	11.9	50.4	48.8	3.43	46.8
Rototi Ig	Average 4.10	880	2.92	453	0.81	3.39	27.1	14.4	103	64.9	64.8	20.0	90.6
n=23 (19)	2sd 0.46	84	0.27	47.8	0.27	1.98	8.3	1.90	15.6	8.09	6.81	2.50	10.1
Ararata Gully 318	Average 5.42	945	2.77	415	0.97	10.2	28.7	18.1	144	82.6	84.9	26.0	147
n=24 (15)	2sd 0.67	259	2.69	52.8	0.66	13.7	8.8	2.16	17.0	21.7	19.1	2.20	28.7
Kakariki 272	Average 4.43	848	4.14	366	1.29	18.0	28.2	17.3	133	75.8	77.8	21.9	107
n=25 (18)	2sd 0.59	146	4.56	40.1	0.97	22.0	7.6	2.94	20.5	10.7	7.75	2.53	11.3
Fordell 449	Average 6.68	743	3.78	428	0.67	15.0	47.8	18.4	162	41.0	41.1	39.3	135
n=25 (19)	2sd 1.24	174	3.81	85.8	0.37	22.7	10.5	3.87	36.6	12.4	12.0	8.71	30.2
Upper Griffin Roac	Average 5.84	1107	2.51	394	1.05	5.52	50.9	16.1	132	70.8	75.7	35.2	154
n=21 (17)	2sd 2.12	234	2.51	43.4	0.28	5.89	8.4	1.84	15.7	10.1	10.9	7.56	8.35
Lower Griffin Roac	Average 5.21	1792	17.6	492	1.71	23.5	47.5	18.7	115	114	118	24.3	195
n=25 (21)	2sd 1.36	948	35.1	95.4	2.72	68.1	16.3	12.4	9.9	22.9	25.0	2.49	53.4
Onepuhi 267	Average 5.27	1135	16.6	435	2.40	42.7	49.3	17.9	150	75.0	77.3	28.1	163
n=23 (10)	2sd 0.87	428	27.0	83.8	3.18	51.6	18.7	2.88	35.9	25.6	24.8	9.69	79.9
Kupe 481	Average 5.38	973	3.38	450	3.94	21.9	37.4	17.2	152	83.1	65.6	29.6	144
n=24 (17)	2sd 1.37	169	4.37	359	10.68	30.9	10.8	4.04	57.4	30.3	33.4	10.0	33.2
Kaukatua 232	Average 6.12	1099	2.16	505	0.81	8.93	59.3	19.0					



Table 3 (cont.) Average and standard deviation for all tephra, see SM Table 1 for full dataset.

Tephra name	Zr ⁹¹ (ppm)	Nb (ppm)	Mo (ppm)	Cs (ppm)	Ba ¹³⁷ (ppm)	Ba ¹³⁸ (ppm)	La (ppm)	Ce (ppm)	Pr (ppm)	Nd (ppm)	Sm (ppm)	Eu ¹⁵¹ (ppm)	Eu ¹⁵³ (ppm)
Kaharoa	Average 850	7.95	1.51	5.38	945	933	21.0	46.4	5.08	19.0	3.99	0.55	0.59
n=25 (18)	2sd 3282	1.99	0.65	1.81	243	208	4.18	9.23	0.97	3.89	1.35	0.20	0.20
Taupo Y5	Average 168	7.17	1.07	6.07	609	601	21.3	45.4	5.25	20.4	4.24	0.88	0.92
n=25 (20)	2sd 61.6	1.97	0.30	1.69	141	141	5.37	11.6	1.45	5.65	1.22	0.31	0.31
Waimihia K3	Average 222	8.53	1.28	5.49	619	625	25.6	61.2	6.55	25.7	5.36	1.10	1.19
n=22 (20)	2sd 33.6	1.82	0.22	1.41	116	123	4.42	11.3	1.19	5.27	1.19	0.21	0.20
Unit K	Average 219	8.99	1.67	6.24	655	653	26.4	56.2	6.37	26.0	5.76	1.12	1.14
n=24 (20)	2sd 25.4	1.14	0.53	1.46	75.8	71.6	2.95	6.29	0.84	3.76	0.86	0.15	0.12
Whakatane K5	Average												
n=21	2sd												
Whakatane - P	Average 175	8.83	1.75	8.76	750	737	31.6	66.7	7.66	30.2	6.18	1.00	0.96
n=25 (21)	2sd 69.7	1.42	0.44	1.61	149	141	20.8	49.4	5.72	23.7	5.13	0.32	0.29
Tuhua K6	Average 1178	102	10.6	6.08	161	158	85.6	182	21.6	86.8	20.1	2.13	1.90
n=23 (22)	2sd 486	41.5	4.15	1.29	356	349	27.8	51.7	7.42	30.5	6.93	0.67	0.61
Mamaku K7	Average 101	8.65	2.24	6.88	968	959	26.5	53.5	6.11	21.9	3.84	0.71	0.73
n=23 (23)	2sd 18.3	1.28	0.95	1.34	161	159	3.4	7.00	0.94	2.64	0.61	0.20	0.18
Rotoma - P	Average												
n=24	2sd												
Rotoma - K8 - D	Average 98.6	8.84	1.81	5.68	970	940	26.1	54.7	6.22	23.8	4.54	0.77	0.82
n=23 (20)	2sd 10.5	0.95	0.25	0.61	100	88.3	2.23	4.97	0.71	2.53	0.51	0.10	0.13
Opepe K9	Average 236	8.82	1.94	8.13	700	674	26.1	56.4	6.38	25.6	5.15	1.01	1.05
n=14 (8)	2sd 25.0	0.57	0.33	1.16	69.1	66.3	3.08	5.72	0.64	2.65	0.65	0.18	0.08
Poronui K10	Average 231	9.16	1.83	7.48	738	737	28.1	57.9	6.65	26.7	5.41	1.06	0.98
n=18 (15)	2sd 33.8	1.88	0.27	1.30	149	155	5.71	11.1	1.32	6.15	1.26	0.28	0.18
Karapiti K11	Average 189	8.41	1.73	7.99	675	688	25.8	55.2	6.03	25.4	5.19	0.99	1.00
n=25 (11)	2sd 37.6	1.62	0.38	1.86	135	135	4.61	9.68	1.11	4.94	1.03	0.16	0.15
Waiohau - P	Average 84.3	7.71	1.38	4.77	858	860	22.5	47.5	5.04	19.4	4.12	0.69	0.68
n=19 (19)	2sd 5.73	0.30	0.50	0.50	38.6	48.5	1.41	2.77	0.36	1.54	0.61	0.13	0.20
Waiohau - K14b D	Average 92.6	7.75	1.82	4.53	867	861	22.4	47.6	5.10	19.6	3.76	0.73	0.81
n=10 (10)	2sd 5.46	0.36	1.24	0.22	18.4	18.4	0.72	0.99	0.25	0.73	0.68	0.12	0.25
Rotorua - P	Average 132	8.12	2.20	5.31	793	812	23.4	49.3	5.30	20.1	4.05	0.62	0.68
n=29 (28)	2sd 41.9	0.98	2.84	1.11	36.0	39.8	3.43	7.96	0.99	3.68	1.18	0.21	0.29
Rotorua - K15 D	Average 79.4	8.30	2.47	7.15	821	819	25.9	53.0	5.38	18.8	3.53	0.45	0.63
n=10 (10)	2sd 36.3	0.93	1.13	1.17	43.2	44.7	3.19	6.48	0.74	2.76	0.50	0.22	0.41
Rerewhakaaitu - P	Average 89.3	7.69	2.31	5.89	794	798	24.1	50.3	5.25	19.5	4.04	0.45	0.54
n=21 (19)	2sd 29.0	1.11	2.02	1.08	64.3	65.8	3.11	7.43	0.90	5.12	1.49	0.16	0.26
Rerewhakaaitu - K	Average 76.1	7.94	2.18	7.73	849	836	24.9	51.2	5.00	18.2	2.70	0.30	0.51
n=11 (10)	2sd 14.2	1.24	1.61	3.07	50.9	43.8	1.29	2.88	0.24	2.22	0.80	0.30	0.37
Okareka	Average 115	8.58	1.92	4.78	1083	1036	25.6	51.1	5.79	19.5	3.72	0.51	0.55
n=22 (22)	2sd 20.4	0.75	0.39	0.64	112	108	2.81	4.95	0.69	1.01	0.55	0.10	0.12
Te Rere	Average 132	11.1	2.32	7.41	805	785	30.9	66.5	7.71	28.9	6.09	0.66	0.61
n=24 (20)	2sd 48.9	1.90	0.31	1.14	119	119	4.80	11.2	1.64	4.83	1.14	0.19	0.11
Kawakawa/Oruanu	Average 147	7.55	1.77	7.48	663	653	22.9	45.4	5.01	18.6	3.72	0.68	0.62
n=25 (23)	2sd 49.6	1.00	0.41	1.40	70.0	74.0	2.83	5.66	0.62	2.71	0.75	0.20	0.17
Pohipi	Average 111	9.62	2.05	7.12	962	949	27.6	60.3	6.20	22.4	4.42	0.74	0.74
n=20 (18)	2sd 29.1	1.76	0.48	1.51	232	212	3.44	25.2	1.08	3.89	0.96	0.45	0.34
Okiaia	Average 153	8.30	1.91	8.44	717	721	24.7	46.5	5.29	20.3	3.99	0.71	0.69
n=23 (23)	2sd 33.7	1.46	0.52	5.01	94.5	93.5	3.39	6.42	0.90	3.93	0.97	0.14	0.19
Unit L	Average 146	8.55	1.92	6.26	758	762	25.2	51.2	5.84	22.0	4.51	0.84	0.90
n=23 (20)	2sd 15.0	1.83	0.42	1.70	109	101	3.64	8.47	1.22	5.05	1.14	0.31	0.29
Awakeri	Average 163	10.59	2.05	4.45	918	894	29.0	62.9	7.24	28.4	5.95	1.22	1.14
n=21 (19)	2sd 39.1	2.83	0.53	1.30	209	205	7.11	14.7	1.66	6.63	1.62	0.48	0.38
Mangaone	Average 171	9.54	1.76	3.88	812	799	25.9	53.1	6.24	25.1	5.23	1.11	1.13
n=24 (19)	2sd 15.6	0.77	0.31	0.79	50.8	47.7	1.86	2.93	0.37	2.04	0.57	0.13	0.09
Hauparu	Average 237	9.15	1.65	3.83	986	979	24.1	46.6	5.48	22.7	4.76	1.13	0.99
n=21 (12)	2sd 29.8	1.09	0.32	1.14	794	765	2.87	5.23	0.73	2.60	0.89	0.17	0.14
Maketu	Average 304	10.8	1.96	2.66	786	782	27.4	51.8	5.97	29.0	6.14	1.58	1.45
n=3 (1)	2sd												
Ngamotu	Average 184	7.85	1.80	2.62	734	732	20.3	43.5	4.81	19.8	4.08	0.98	0.93
n=21 (19)	2sd 31.3	0.93	0.43	0.61	169	164	3.76	8.31	0.92	3.64	0.75	0.16	0.11
Tahuna	Average 102	8.42	2.15	7.92	941	923	26.4	50.8	5.31	18.5	3.42	0.42	0.50
n=23 (17)	2sd 17.0	1.07	0.71	3.11	108	105	7.05	10.88	1.71	7.82	1.71	0.11	0.11
Earthquake Flat Ig	Average 96.45	8.94	2.04	4.87	1064	1053	26.0	51.7	5.58	19.4	3.66	0.51	0.53
n=24 (11)	2sd 8.53	0.90	0.43	1.74	87.0	80.1	2.69	3.72	0.55	1.49	0.51	0.10	0.12
Rotoehu tephra	Average 116	8.73	1.88	3.64	1043	1056	25.6	51.8	5.54	20.6	3.81	0.64	0.66
n=21 (14)	2sd 49.2	0.80	0.46	1.25	121	121	2.75	6.00	0.62	2.10	0.43	0.24	0.25
Rotoiti Ig	Average 91.6	8.42	1.77	3.82	1065	1066	25.2	49.8	5.21	19.4	3.57	0.49	0.51
n=23 (19)	2sd 11.1	1.02	0.40	0.65	128	123	3.09	5.15	0.73	2.92	0.65	0.12	0.14
Ararata Gully 318	Average 145	9.62	3.80	7.96	827	829	28.3	58.4	6.25	23.9	4.27	0.71	0.58
n=24 (15)	2sd 30.2	0.98	2.10	1.15	96.6	91.7	2.43	4.73	0.65	2.33	0.61	0.10	0.16
Kakariki 272	Average 106	8.52	2.59	5.68	869	878	25.3	50.1	5.42	19.8	3.99	0.63	0.65
n=25 (18)	2sd 12.4	0.81	0.88	0.86	245	246	3.03	4.75	0.59	2.61	0.62	0.12	0.13
Fordell 449	Average 138	11.8	2.04	8.60	874	884	33.7	71.1	8.21	31.1	6.53	0.73	0.75
n=25 (19)	2sd 32.5	2.67	0.78	2.20	194	189	7.59	15.7	1.85	6.76	1.53	0.18	0.28
Upper Griffin Roat	Average 164	9.16	1.87	7.66	714	658	29.3	57.3	6.82	24.8	4.72	0.77	0.86
n=21 (17)	2sd 55.8	1.19	0.32	0.76	131	52.9	6.80	4.18	0.75	2.74	0.80	0.08	0.11
Lower Griffin Roat	Average 193	8.25	2.07	5.61	743	749	23.6	44.9	5.02	18.8	3.82	0.84	0.91
n=25 (21)	2sd 53.0	2.41	0.31	0.83	170	183	3.06	0.82	1.19	3.10	0.77	0.19	0.22
Onepuhi 267	Average 162	9.67	2.41	8.38	790	783	26.3	52.5	5.99	22.9	4.68	0.78	0.80
n=23 (19)	2sd 79.4	2.65	0.58	2.03	208	211	5.88	13.4	1.39	7.48	1.22	0.22	0.25
Kupe 481	Average 144	9.58	2.18	7.69	820	1055	26.4	59.7	7.13	24.0	4.78	0.70	0.72
n=24 (17)	2sd 36.3	2.63	0.58	2.50	240	580	8.18	15.2	4.08	7.98	1.51	0.17	0.16



Table 3 (cont.) Average and standard deviation for all tephra, see SM Table 1 for full dataset.

Tephra name	Gd (ppm)	Tb (ppm)	Dy (ppm)	Ho (ppm)	Er (ppm)	Tm (ppm)	Yb (ppm)	Lu (ppm)	Hf (ppm)	Ta (ppm)	W (ppm)	Pb (ppm)	Th (ppm)	U (ppm)
Kaharoa	Average 3.95	0.64	5.00	1.19	3.97	0.75	6.15	1.58	18.4	0.75	1.55	17.7	12.3	5.27
n=25 (18)	2sd 1.32	0.18	3.47	1.31	5.31	1.27	11.5	3.11	65.3	0.18	0.54	4.48	5.18	8.96
Taupou Y5	Average 4.31	0.66	4.21	0.86	2.72	0.40	2.94	0.74	4.35	0.53	1.45	17.3	9.16	2.42
n=25 (20)	2sd 1.28	0.21	1.43	0.29	0.83	0.14	1.05	0.22	1.54	0.15	0.46	4.67	2.48	0.63
Waimihia K3	Average 5.18	0.84	5.10	1.13	3.30	0.52	3.46	0.95	5.69	0.66	1.80	19.9	10.6	2.59
n=22 (20)	2sd 1.25	0.20	1.01	0.24	0.63	0.11	0.80	0.21	0.89	0.13	0.53	3.29	1.64	0.46
Unit K	Average 5.61	0.86	5.61	1.18	3.41	0.50	3.60	0.55	6.41	0.70	1.70	20.5	10.9	2.89
n=24 (20)	2sd 0.91	0.12	0.66	0.16	0.46	0.08	0.59	0.08	0.78	0.11	0.37	5.00	1.39	0.41
Whakatane K5	Average													
n=21	2sd													
Whakatane - P	Average 5.83	0.88	5.31	1.16	3.20	0.50	3.40	0.58	5.07	0.73	1.77	21.1	11.6	2.96
n=25 (21)	2sd 2.86	0.36	1.67	0.31	0.99	0.18	0.92	0.18	2.27	0.18	0.72	4.94	2.52	0.54
Tuhua K6	Average 21.3	3.41	23.0	4.81	14.2	2.24	14.0	2.07	23.6	6.69	2.01	28.0	17.7	5.89
Mamaku K7	Average 3.87	0.64	4.18	0.85	2.69	0.40	3.05	0.45	3.32	0.81	1.68	16.5	11.9	2.94
n=23 (23)	2sd 0.98	0.18	0.81	0.14	0.39	0.10	0.51	0.11	0.47	0.09	0.42	3.46	1.54	0.47
Rotoma - P	Average													
n=24	2sd													
Rotoma - K8 - D	Average 4.15	0.66	4.17	0.92	2.90	0.42	3.24	0.48	3.31	0.77	1.47	16.1	11.5	2.86
n=23 (20)	2sd 1.63	0.09	0.50	0.11	0.45	0.07	0.38	0.07	0.51	0.10	0.25	2.29	1.13	0.39
Opepe K9	Average 5.78	0.86	5.20	1.09	3.57	0.49	3.27	0.53	5.89	0.69	1.55	18.9	11.1	2.88
n=14 (8)	2sd 1.17	0.11	0.27	0.17	0.11	0.09	0.09	0.09	0.57	0.10	0.31	1.96	2.15	0.37
Poronui K10	Average 9.40	0.89	5.69	1.24	3.53	0.56	3.89	0.57	6.13	0.73	1.79	21.2	12.3	3.03
n=18 (15)	2sd 9.69	0.19	1.18	0.23	0.83	0.14	0.72	0.13	0.94	0.16	0.39	3.82	2.68	0.61
Karapiti K11	Average 5.48	0.78	5.18	1.07	4.78	0.53	3.40	0.52	5.70	0.66	1.66	19.6	12.0	2.96
n=25 (11)	2sd 1.25	0.15	1.07	0.20	0.98	0.11	0.66	0.09	1.09	0.16	0.35	3.75	2.13	0.59
Waiohau - P	Average 3.53	0.58	3.74	0.78	2.34	0.38	2.56	0.36	2.69	0.62	1.36	15.3	9.37	2.37
n=19 (19)	2sd 0.58	0.12	0.50	0.12	0.26	0.06	0.40	0.06	0.32	0.10	0.28	0.88	0.65	0.18
Waiohau - K14b D	Average 3.17	0.55	3.72	0.71	2.40	0.31	2.60	0.40	3.08	0.65	1.50	14.9	9.72	2.47
n=10 (10)	2sd 0.58	0.12	0.43	0.10	0.23	0.05	0.38	0.08	0.24	0.09	0.30	0.91	0.47	0.10
Rotorua - P	Average 3.65	0.64	4.09	0.87	2.47	0.40	2.77	0.38	3.70	0.68	1.52	16.1	10.8	2.73
n=29 (28)	2sd 1.11	0.23	1.16	0.21	0.72	0.11	0.58	0.14	0.70	0.13	0.36	2.27	1.0	0.40
Rotorua - K15 D	Average 2.83	0.49	3.39	0.70	2.14	0.33	2.44	0.37	2.69	0.81	2.14	18.0	13.4	3.38
n=10 (10)	2sd 0.88	0.10	0.51	0.10	0.43	0.10	0.48	0.09	0.80	0.15	0.47	1.68	1.25	0.47
Rerewhakaaitu - P	Average 3.25	0.47	3.59	0.80	2.32	0.38	2.68	0.38	2.85	0.73	1.70	16.9	11.6	3.05
n=21 (19)	2sd 1.13	0.22	1.31	0.33	0.79	0.14	0.75	0.12	0.72	0.13	0.42	2.37	2.19	0.59
Rerewhakaaitu - K1	Average 2.90	0.51	3.07	0.73	2.03	0.39	2.15	0.35	2.74	0.77	1.90	18.8	13.2	3.32
n=11 (10)	2sd 0.43	0.20	0.63	0.25	0.38	0.12	0.56	0.09	0.33	0.12	0.24	3.49	1.51	0.41
Okareka	Average 3.64	0.54	3.33	0.75	2.93	0.37	2.85	0.41	3.07	0.81	1.37	13.2	10.7	2.47
n=22 (22)	2sd 0.59	0.10	0.45	0.12	1.12	0.07	0.52	0.08	0.44	0.16	0.38	1.39	1.24	0.33
Te Rere	Average 5.87	0.97	6.14	1.26	4.51	0.64	4.34	0.62	4.32	1.25	1.74	19.7	14.0	3.55
n=24 (20)	2sd 1.10	0.16	1.10	0.21	1.90	0.16	0.90	0.14	1.05	0.54	0.23	2.75	1.91	0.57
Kawakawa/Oruanu	Average 3.59	0.60	3.54	0.75	2.34	0.38	2.52	0.39	4.15	0.62	1.59	15.3	11.2	2.85
n=25 (23)	2sd 0.70	0.12	0.71	0.12	0.71	0.07	0.46	0.10	0.81	0.16	0.37	2.10	1.59	0.40
Poihipi	Average 3.75	0.64	3.97	0.86	2.54	0.40	3.14	0.42	3.64	0.91	1.85	21.5	13.3	3.34
n=20 (18)	2sd 0.85	0.18	1.18	0.19	0.49	0.10	0.75	0.10	0.75	0.24	0.57	12.0	2.99	0.73
Okiaia	Average 3.95	0.58	4.00	0.80	2.59	0.39	2.88	0.41	4.20	0.71	1.66	16.9	12.0	3.11
n=23 (23)	2sd 0.71	0.13	0.82	0.16	0.66	0.13	0.56	0.06	0.73	0.13	0.37	2.93	1.90	0.58
Unit L	Average 4.34	0.68	4.34	0.94	3.06	0.47	3.22	0.49	4.32	0.74	1.54	16.0	11.7	2.90
n=23 (20)	2sd 1.20	0.21	1.50	0.30	0.98	0.13	0.88	0.17	0.58	0.15	0.25	2.74	1.85	0.47
Awakeri	Average 5.60	0.93	5.69	1.17	3.68	0.59	3.93	0.59	4.69	0.79	1.48	17.6	10.5	2.56
n=21 (19)	2sd 1.51	0.26	1.26	0.30	0.87	0.16	0.91	0.16	1.00	0.23	0.41	4.30	2.25	0.61
Mangaone	Average 5.19	0.81	5.23	1.09	3.40	0.54	3.68	0.56	4.79	0.71	1.20	14.3	8.94	2.16
n=24 (19)	2sd 0.64	0.11	0.63	0.14	0.46	0.07	0.39	0.08	0.54	0.07	0.20	1.54	0.85	0.18
Hauparu	Average 4.41	0.76	4.89	1.01	3.03	0.49	3.34	0.48	5.58	0.61	1.12	12.6	8.08	1.95
n=21 (12)	2sd 0.89	0.10	0.61	0.13	0.38	0.08	0.56	0.12	0.90	0.15	0.19	2.40	1.22	0.41
Maketu	Average 6.15	1.05	7.60	1.33	4.20	0.76	4.27	0.55	7.38	0.72	1.13	13.9	7.60	1.71
n=3 (1)	2sd													
Ngamotu	Average 3.87	0.62	3.78	0.78	2.50	0.38	2.87	0.45	4.58	0.58	0.99	11.8	7.02	1.86
n=21 (19)	2sd 0.89	0.13	0.84	0.16	0.57	0.11	0.71	0.11	0.83	0.10	0.17	1.61	1.89	0.38
Tahuna	Average 3.11	0.48	3.06	0.67	2.18	0.34	2.49	0.38	3.12	0.86	1.55	16.4	13.0	3.26
n=23 (17)	2sd 0.49	0.10	0.45	0.10	0.45	0.06	0.45	0.07	0.69	0.17	0.43	1.09	0.67	0.10
Earthquake Flat Ig	Average 3.05	0.53	3.08	0.61	2.08	0.32	2.31	0.38	2.71	0.76	1.50	14.8	11.4	2.93
n=24 (11)	2sd 0.47	0.09	0.46	0.11	0.44	0.11	0.34	0.07	0.50	0.19	0.61	2.79	3.42	0.87
Rotoehu tephra	Average 3.63	0.58	3.60	0.77	2.21	0.38	2.75	0.45	3.21	0.72	1.40	13.8	9.95	2.56
n=21 (14)	2sd 0.76	0.11	0.65	0.15	0.45	0.09	0.48	0.08	1.06	0.10	0.24	2.36	1.41	0.41
Rotoiti Ig	Average 3.06	0.46	3.11	0.63	1.96	0.30	2.57	0.34	2.67	0.70	1.33	12.9	9.52	2.36
n=23 (19)	2sd 0.75	0.07	0.41	0.11	0.34	0.06	0.33	0.06	0.43	0.16	0.38	1.96	1.25	0.34
Ararata Gully 318	Average 4.08	0.60	4.26	0.85	2.53	0.40	2.78	0.41	4.04	0.85	1.76	22.4	13.5	3.37
n=24 (15)	2sd 0.72	0.08	0.53	0.10	0.26	0.06	0.50	0.07	0.52	0.15	0.38	3.96	1.26	0.30
Kakariki 272	Average 3.39	0.55	3.48	0.72	2.22	0.37	2.41	0.40	3.39	0.77	1.51	19.5	13.7	3.06
n=25 (18)	2sd 0.58	0.11	0.54	0.11	0.37	0.07	0.28	0.06	0.37	0.15	0.34	2.29	1.86	0.50
Fordell 449	Average 6.38	1.05	6.69	1.35	3.93	0.62	4.30	0.66	4.74	1.08	2.03	24.9	14.9	3.68
n=25 (19)	2sd 1.44	0.24	1.98	0.32	0.91	0.18	1.07	0.17	1.12	0.27	0.58	4.97	3.40	0.81
Upper Griffin Roac	Average 5.04	0.82	5.23	1.23	3.64	0.53	3.77	0.57	4.82	0.76	1.67	17.3	12.3	3.10
n=21 (17)	2sd 1.38	0.09	1.37	0.14	0.38	0.08	0.77	0.07	0.37	0.12	0.28	5.29	1.01	0.31
Lower Griffin Roac	Average 3.64	0.58	4.06	0.82	2.46	0.37	2.73	0.43	5.37	0.74	1.40	17.4	12.6	2.97
n=25 (21)	2sd 0.56	0.11	0.69	0.13	0									



Table 4. Dominant ferroganessian mineral assemblages for late Quaternary silicic tephra deposits updated from Froggatt and Lowe 1990

Assemblage 1 Hyp +/- aug +/- hbl	Assemblage 2 Hyp + hbl +/- aug	Assemblage 3 Hyp + hbl + bio	Assemblage 4 Hyp + cgt +/- hbl	Assemblage 5 Hyp + aug +/- hbl	Assemblage 6 Aegirine*
Taupō VC	Okataina VC	Okataina VC	Okataina VC	Okataina VC	Tuhua VC (Mayor Is)
Taupō (all)	Mamaku	Kaharoa	Whakatane	Hauparu	Tuhua
(Mapara)	Waiohau	Rotorua (upper)	Rotoma	Te Mahoe	
(Whakaipo)	Rotorua (lower)	Rerewhakaaitu	Rotoiti/Rotoehu (all)	Maketu	
Waimihia	Unit L	Okareka			
Unit K	Te Rere	Rotoiti/Rotoehu (upper)			
Motutere (all?)	(Omatoroa)				
Opepe	Awakeri	Kapenga VC			
Poronui	Mangaone	Earthquake Flat			
Karapiti	Tahuna				
	Ngamotu	Maroa VC			
		Puketarata			
	Taupō VC				
	Kawakawa (all)				
	Poihipi				
	Okaia				
	(Tihoi)				
	(Waihoroa)				
	(Otake)				

The assemblages are listed with mineral species in order of abundance, the diagnostic mineral in each assemblage is in bold, tephra are listed multiple times if their mineral assemblage changes through the eruption sequence, and deposits in brackets are not included in the TephraNZ database

*Assemblage 6 Aegirine +/- riebeckite +/- aenigmatite +/- olivine +/- tuhualite

Motutere was listed in Froggatt and Lowe 1990 as a single unit with Assemblage 1 mineralogy, but this has subsequently been redefined by Wilson 1993 into two subunits G&H, which do not have their independent assemblages defined.



Table 5. Geochemically similar tephra and their distinctions

Tephra	Ages (ka) ^a	Magma volume (km ³) [*]	Geochemical distinction	Other distinctions
Taupo (Unit Y, TVC)	1,718 +/- 10	13.4		age, volume, stratigraphic relationship
Waimihia (Unit S, TVC)	3,382 +/- 50	5.1	Indistinguishable	
Waimihia (Unit S, TVC)	3,382 +/- 50	5.1	Lu, Sc, Mn, Co, Ba	age, volume, stratigraphic relationship to Stent Tephra
Unit K (TVC)	5,088 +/- 73	0.12		
Waimihia (Unit S, TVC)	3,401 +/- 108	5.1	FeO vs. CaO, Na ₂ O+K ₂ O, Lu, Sc, Mn, Co,	age, volume, stratigraphic relationships
Poronui (Unit C, TVC)	11,159 +/- 51	0.23		
Waimihia (Unit S, TVC)	3,382 +/- 50	5.1	FeO vs. CaO, Na ₂ O+K ₂ O, Lu, Sc, Mn, Co,	age, stratigraphic relationships
Karapiti (Unit B, TVC)	11,501 +/- 104	0.42		
Unit K (TVC)	5,088 +/- 73	0.12	FeO vs. CaO, Na ₂ O+K ₂ O, Lu, Sc, Mn, Co,	age, stratigraphic relationships
Poronui (Unit C, TVC)	11,170 +/- 115	0.23		
Unit K (TVC)	5,088 +/- 73	0.12	FeO vs. CaO, Na ₂ O+K ₂ O, Lu, Sc, Mn, Co,	age, stratigraphic relationships
Karapiti (Unit B, TVC)	11,501 +/- 104	0.42		
Mamaku	7,992 +/- 58	13.0	Ba/Th vs. Rb/Sr, Rb/Zr vs. Rb/Sr	volume, chemistry
Rotoma	9,472 +/- 40	8.0		
Poronui (Unit C, TVC)	11,195 +/- 51	0.23	Indistinguishable	
Karapiti (Unit B, TVC)	11,501 +/- 104	0.4		
Waiohau	14,018 +/- 91	3.3	Ba, Hf, Ba/Zr vs. Ba/Th, Rb/Sr, Rb/Zr; Rotorua is bi-modal	volume, chemistry, mineralogy
Rotorua	15,738 +/- 263	1.0		
Waiohau	14,018 +/- 91	3.3	Cs, La, Ce, Nd, Eu, Rb/Zr, Ba/Th; Rerewhakaaitu is bimodal	volume, chemistry, mineralogy
Rerewhakaaitu	17,209 +/- 249	5.0		
Rotorua	15,738 +/- 263	1.0	Indistinguishable	age
Rerewhakaaitu	17,209 +/- 249	5.0		
Poihipi	28,446 +/- 670	0.5	La/Yb vs. Ba/Y, Rb/Zr vs. Ba/Th	age
Tahuna	39,268 +/- 1193	2.0 ^b		
Rotoma	9,472 +/- 40	8.0	Ba, Cs, Y, Sm, Nd, Pr, Er, Ho, Dy, Tb, Eu, Tm, Yb, Pb, U, Th	age, volume, stratigraphic relationships
Rotoehu	45,170 +/- 3300	90 ^c		
KOT	25,358 +/- 162	530.0	Indistinguishable	volume, shard morphology
Okaia	28,621 +/- 1428	3.0		
KOT	25,358 +/- 162	530.0	Unit L bimodal in Rb/Zr, Ba/Th, Ce/Th, Y/Th	volume, mineral geochemistry (Smith et al., 2002)
Unit L (Mangaone sgp OVC)	31.4 ka	c. 7.0 ^d		

^a Ages from Table 1

^{*} Volumes from Lowe et al., 2008 and references therein, except for Tahuna, Rotoiti/Rotoehu, and Unit L (Mangaone Subgroup)

^b Tahuna airfall volume estimate in km³ from Froggatt and Lowe 1990

^c Rotoehu airfall volume in km³ from Froggatt and Lowe 1990

^d Unit L airfall volume in km³ from Jurado-Chichay and Walker 2000