



Barometers Behaving Badly I: Assessing the Influence of Analytical and Experimental Uncertainty on Clinopyroxene Thermobarometry Calculations at Crustal Conditions

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Abstract

The composition of clinopyroxene and clinopyroxene-liquid (Cpx-Liq) pairs are frequently used to calculate crystallization/equilibration pressures in igneous systems. While canonical uncertainties are often assigned to calculated pressures based on fits to calibration or test datasets, the sources of these uncertainties (and thus ways to reduce them) have not been rigorously assessed. We show that considerable uncertainties in calculated pressures arise from analytical error associated with Electron Probe Microanalyser (EPMA) measurements of Cpx. Specifically, low X-ray counts during analysis of elements with concentrations <1 wt% resulting from insufficient count times and/or low beam currents yield highly imprecise measurements (1σ errors of 10–40% for Na₂O).

Low analytical precision propagates into the calculation of pressure-sensitive mineral components such as jadeite. Using Monte Carlo approaches, we demonstrate that elemental variation resulting from analytical precision alone generates pressures spanning ~4 kbar (~15 km) for a single Cpx and ~6 kbar for a single Cpx-Liq pair using popular barometry expressions. In addition, analytical uncertainties in mineral compositions produce highly correlated arrays between pressure and temperature that have been previously attributed to transcrustal magma storage. Before invoking such geological interpretations, a more mundane origin from analytical imprecision must be ruled out. Most importantly, low analytical precision does not just affect the application of barometers to natural systems; it has also affected characterization of Cpx in experimental products used to calibrate and test barometers. The impact of poor precision on each individual measurement is often magnified by the small number of measurements made within experimental charges, meaning that low analytical precision and true variability in mineral compositions have not been sufficiently mediated by averaging multiple EPMA analyses. We compile the number of Cpx measurements performed in $N = 307$ experiments used to calibrate existing barometers, and $N = 490$ new experiments, finding ~45% of experiment charges were characterized by ≤ 5 individual Cpx analyses. Insufficient characterization of the true composition of experimental phases likely accounts for the fact that all Cpx-based barometers exhibit large errors (± 3 kbar) when tested using global experimental datasets.

We suggest specific changes to analytical and experimental protocols, such as increased count times and/or higher beam currents when measuring low concentration elements in relatively beam resistant Cpx in experiments and natural samples. We also advocate for increasing the number of analyses per experimental charge, resolving interlaboratory analytical offsets and improving data reporting. Implementing these changes is essential to produce a more robust dataset to calibrate and test the next generation of more precise and accurate Cpx-based barometers. In turn, this will enable more rigorous investigation of magma storage geometries in a variety of tectonic settings (e.g. distinguishing true transcrustal storage vs. storage in discrete reservoirs).

Keywords: Monte Carlo, experimental uncertainty, analytical uncertainty, thermobarometry, clinopyroxene

INTRODUCTION

Constraining the conditions under which magma is stored and transported within the crust is of critical importance to understanding volcanic systems (e.g. McGuire *et al.*, 2017). It has long been recognized, based on thermodynamic principles and

from chemical characterization of experimental products, that the composition of igneous minerals and co-existing liquids can be used to place constraints on the pressures, temperatures, and water concentrations at which these phases grew and equilibrated (e.g. Bacon & Carmichael, 1973; Brown & Parsons, 1981; Lindsley & Andersen, 1983; Putirka, 1999, 1997, 2008).

Many of the most commonly used mineral-based igneous thermobarometers revolve around clinopyroxene (Cpx), perhaps because this phase is relatively abundant in a wide variety of different volcanic systems and tectonic settings (e.g. mid-oceanic ridges, oceanic islands, and volcanic arcs), and in a wide range of lava compositions (basalts to rhyolites). Cpx-based equilibria are sensitive to pressure because there is a large change in volume associated with the exchange of Na and Al (the jadeite component, $\text{NaAlSi}_2\text{O}_6$) between liquid (Liq) and Cpx. There are also relatively large volume changes between different Cpx components (e.g. jadeite and diopside-hedenbergite, $\text{CaMgSi}_2\text{O}_6$ – $\text{CaFeSi}_2\text{O}_6$, Putirka, 2016; Putirka et al., 1996). The exchange of jadeite and diopside-hedenbergite between clinopyroxene and liquid is also sensitive to temperature, so the abundance of these components can be used as a thermometer (e.g. Putirka, 1999).

By parameterizing the relationship between phase composition and intensive parameters in experiments, measurements of natural crystals and co-existing equilibrium liquids can be used to calculate pressures and temperatures (e.g. Putirka, 1999, 2008; Neave & Putirka, 2017; Petrelli et al., 2020; Wang et al., 2021). The vast majority of published calibrations have a structure rooted in thermodynamics, both in terms of the form of the expression, and the compositional components included (e.g. jadeite in Cpx).

For example, both the Putirka et al. (1996) eqP1 and Putirka (2008) eq30 models for Cpx-Liq barometry have the same general form informed by thermodynamics:

$$P = a + \frac{bT}{10^4} + \frac{cT}{10^4} \ln \left(\text{Jd}^{\text{Cpx-Liq}} \right) + \dots \quad [\text{Equation 1}]$$

Additional empirical composition terms are added to improve regression statistics (Neave & Putirka, 2017), although sometimes these terms can also be approximately tied to thermodynamic reasoning. For example, the $X_{\text{Na}}^{\text{Liq}} X_{\text{Al}}^{\text{Liq}}$ term in Putirka et al. (1996) eqP1 implies identical activity coefficients for Na and Al in the liquid. In general, an increasing number of empirical terms have been added to barometry equations to improve the fit between calculated and experimental pressures in calibration datasets as more experimental data have become available. For example, while both eqP1 (P1996) and eq30 (P2008) have additional empirically derived terms for the sum of the cation fraction of Na and Al, eq30 has terms for the liquid Mg#, the diopside-hedenbergite component in the Cpx, and the log of liquid Fe, Mg and K cation fractions.

More recently, machine learning algorithms have been calibrated for equilibria involving Cpx (Petrelli et al., 2020; Higgins et al., 2022; Jorgenson et al., 2022). Instead of creating equations with specific terms informed by thermodynamics (as in Equation 1), these machine learning techniques simply input selected oxide data from a training dataset into the machine learning algorithm, along with the predictor variable (e.g. pressure or temperature; Petrelli et al., 2020).

When discussing errors and uncertainties associated with different thermobarometers in this manuscript, we follow the National Physics Laboratory terminology guidelines (Bell, 2001). Error is taken as the difference between the measured value and the true value; if a barometer calculates a pressure of 3 kbar but the experiment was performed at 2 kbar, the error is +1 kbar. Uncertainty quantifies the doubt about the measurement result, so an error with an unknown magnitude is referred to as an uncertainty. A random uncertainty or error means that if the measurement is repeated, a different value is obtained each time, but averaging of sufficient measurements converges on the true value. Random uncertainties may follow a normal or Poisson

distribution (or other unskewed distribution). However, the central limit theorem states that as the sample size grows, a distribution of sample means approximates a normal distribution (even if the data are not normally distributed). Systematic uncertainties or errors cause the measurement to be offset from the true value, so additional measurements do not help to converge on the correct answer, meaning that more measurements produce a more precise, but inaccurate result.

The success of a given barometer or thermometer is normally assessed by comparing predicted pressures and temperatures with experimentally determined pressures and temperatures. The goodness of fit is typically assessed using R^2 values and estimates of the standard error estimate (SEE, sometimes referred to as the root mean-square error, RMSE). In general, Cpx and Cpx-Liq thermometers have high R^2 values (>0.8) and relatively low SEE (e.g. ± 20 – 100 K; e.g. Putirka, 2008, Petrelli et al., 2020). These SEEs represents only a 2%–10% error considering the temperatures of most magmatic systems (~ 1000 – 1400 K). In contrast, barometers commonly have SEE of 1.4–5 kbar ($\sim \pm 6$ – 19 km, using a representative ρ of 2700 kg/m^3 throughout this paper), which corresponds to very large percentage errors given the depths of interest in many volcanic systems. For example, these large SEEs span the entire crustal column in many tectonic settings (e.g. 5–8.5 km in MORB, White et al., 1992; 14–24 km in Hawai'i, Leahy et al., 2010). This means that pressures calculated on individual Cpx analyses do not have sufficient precision/resolution to reliably identify upper, mid or lower crustal storage, or distinguish storage in distinct magma reservoirs separated by ~ 1 – 2 km (unlike melt inclusion saturation pressures, which can achieve such precision at relatively low pressures; Lerner et al., 2021; Wieser et al., 2022a; Wieser et al., 2021). Even in arcs with Moho pressures of 8 ± 3 kbar (based on $\sim 30 \pm 12$ km from Profeta et al., 2016), these SEE estimates can only just distinguish between upper, mid and lower crustal storage. As many of the uncertainties associated with thermobarometers are random uncertainties, averaging multiple experiments (or natural Cpx analyses) can result in significant improvements. For example, Putirka et al. (1996) showed that the SEE on individual experiments using their eqP1 is 1.36 kbar, but can be reduced to 0.32 kbar if they average experiments conducted at the same pressure.

In general, more recent calibrations of mineral-only and mineral-melt barometers state smaller SEEs (e.g. ± 1.4 kbar for Cpx-Liq from Neave & Putirka, 2017, ± 1.66 kbar for Cpx-only from Wang et al., 2021, vs. ± 3.6 kbar for equation 30 and ± 5 kbar for eq32c of Putirka, 2008). However, statistics quoted by different studies are not directly comparable. For example, the ± 1.4 kbar SEE commonly stated by petrological studies using the Neave & Putirka (2017) barometer describes the model fit to the calibration dataset of 113 experiments. When this thermobarometer is applied to their global compilation of experimental data that span a wider compositional range than was used in the calibration, the SEE is ± 3.6 – 3.8 kbar. When applied to test data with compositions more similar to the calibration dataset (i.e. tholeiitic basalts), the Neave & Putirka (2017) barometer return a SEE similar to the quoted SEE of 1.4 kbar. The ± 1.66 kbar SEE from Wang et al. (2021) describes the overall fit to 100 random splits into training and validation of the calibration dataset. However, when applied to global datasets (including data not used at any point during model tuning), this barometer has a SEE of ± 3.68 kbar. Finally, the Cpx-only and Cpx-Liq machine-learning barometer of Petrelli et al. (2020) have SEEs of ± 3.1 kbar and ± 3.2 kbar, respectively, when applied to a subset of $N = 119$ experiments not used for calibration. The 'global' statistics for newer barometers are more

similar to the larger quoted errors of Putirka (2008), who calculate statistics using a global calibration dataset.

It is also important to recognize that the quoted SEE describes the overall fit across the range of pressures or temperatures being considered. Close to the edges of the distribution (e.g. very shallow or very deep pressures), the error can be larger than the SEE, and toward the center of the distribution, the error can be smaller. Additionally, the SEE estimated from a barometer between 0 and 40 kbar (e.g. Putirka et al., 1996) is not directly comparable to one estimated between 0 and 20 kbar (e.g. Petrelli et al., 2020), which can also make comparison of statistics from different barometers challenging.

Despite the fact that it is difficult to directly compare statistics, we can say in general that the majority of Cpx-based barometers show large SEE (± 2 –5 kbar, 7.6–19 km) when applied to global datasets (Wieser et al., 2023). Given that significantly greater precision in crustal barometry is needed to address many of the key issues in igneous petrology (McGuire et al., 2017), we try to reconcile the source of these large errors, to identify ways in which to improve Cpx-based barometric methods in future.

A new test dataset—ArcPL

The majority of Cpx-based barometers were calibrated using experiments present in the LEPR dataset compiled in 2008 (Library of Experimental Phase Relationships; Hirschmann et al., 2008, e.g. $N=850$ experiments in Petrelli et al., 2020). For clarity, in this paper the word experiment is used to represent a single experimental run/capsule/charge. In addition to using experimental data within LEPR, we also compile Cpx-bearing experiments on variably hydrous compositions ranging from basalt to rhyolite at crustal conditions (<13 kbar) that were not included in LEPR. Most of the studies represented in this new dataset were published after LEPR, and have not been added to it yet (Costa, 2004; Bogaerts et al., 2006; Pichavant & Macdonald, 2007; Hamada & Fujii, 2008; Mercer & Johnston, 2008; Alonso-Perez et al., 2009; Feig et al., 2010; Parman et al., 2011; Krawczynski et al., 2012; Almeev et al., 2013; Blatter et al., 2013, 2017; Rader & Larsen, 2013; Cadoux et al., 2014; Mandler et al., 2014; Nandedkar et al., 2014; Parat et al., 2014; Andújar et al., 2015; Melekhova et al., 2015; Riker et al., 2015; Erdmann et al., 2016; Erdmann & Koepke, 2016; Husen et al., 2016; Ulmer et al., 2018; Firth et al., 2019; Neave et al., 2019; Waters et al., 2021; Nakatani et al., 2022; Marxer et al., 2022). We also include a small number of pre-2008 experiments, which were not included in LEPR (Rutherford et al., 1985; Berndt, 2004; Sisson et al., 2005). We call our new dataset ArcPL (Arc post-LEPR, $N=490$). We also consider a subset of the LEPR database relevant to arc magmas (ArcLEPR, $N=307$), as well as the entire LEPR dataset used to calibrate the thermobarometers of Putirka (2008) and Petrelli et al. (2020) at various points in this manuscript. One advantage of ArcPL is that many of the experiments were conducted relatively recently, meaning that it was easier to obtain additional analytical information from the authors, and more information on analytical conditions was generally presented in the text and supporting information. We use these different datasets of experiments to place quantitative constraints on sources of uncertainty when calibrating and testing Cpx-based barometers. The relative performance of each barometer is evaluated in Wieser et al., 2023.

Sources of uncertainty

Thermobarometer calibrations based on experimental products are subject to three main sources of uncertainty:

(1) Uncertainties relating to the regression method used to calibrate thermobarometry equations and models (e.g. extrapolation, overfitting, incomplete data; Fig. 1a).

(2) Analytical uncertainties associated with measurements of minerals and glasses, normally by electron probe microanalyzer (EPMA, e.g. insufficient counts, heterogeneity in primary standards within a single chip and between chips, beam damage, instrument miscalibration; Fig. 1b).

(3) Experimental errors (e.g. crystallization under disequilibrium conditions, uncertainty in the pressure in a piston-cylinder; Fig. 1c), as well as additional issues associated with analyzing small experimental products (Fig. 1d).

Each of these are discussed in detail below.

Model formulation and calibration

One unavoidable reason why barometers have higher percentage errors than thermometers relates to thermodynamics, namely major and minor element variations in available igneous phases are simply more sensitive to temperature than pressure, because relevant reactions have larger changes in entropy (relating to T) than volume (relating to P). This is in contrast to subsolidus metamorphic systems where a relatively large number of very P -sensitive phase transitions and chemical reactions are available (Powell & Holland, 2008). The paucity of P -sensitive reactions between solid mineral phases and silicate melts reflects the ability of liquid-rich igneous systems to absorb changes in volume (and to a lesser extent temperature and composition) without drastically changing the composition of the coprecipitating phases (Putirka, 2008). Thus, it is expected that regressions of experimental Cpx compositions against pressure will have relatively poor predictive power (Fig. 1a), as many variables other than pressure have more influence on the reaction (e.g. temperature, water content, melt composition).

However, there are a number of sources of uncertainty that can be introduced during the calibration of barometric models, which decrease the accuracy and precision of barometers beyond some theoretical limit determined by the thermodynamics of each reaction. Identifying these is vital to produce the best possible barometers for igneous systems.

Firstly, the calibration dataset may span a restricted compositional, pressure or temperature range, and empirical fits developed on this dataset may return poor results when extrapolated outside this range (Fig. 1a). For example, Neave & Putirka (2017) cautioned users about the limited applicability of their model in highly oxidized and alkaline systems, as it was predominantly calibrated using tholeiites. Similarly, Wang et al. (2021) attributed the poorer performance of their barometer at pressures >12 kbar to the lack of high-pressure experiments in their calibration dataset. Models can also be overfitted to calibration data. This is a particular problem for certain machine-learning algorithms such as extra tree regressors, which can perfectly fit the training data, but produce erratic results on test data.

A second source of uncertainty relates to the fact that not all experimental studies report the concentrations of all the major and minor elements in Cpx and Liq used when performing thermobarometry calibrations. This is problematic for several reasons. Firstly, missing elements from the quantification routine during EPMA analysis can affect the concentration of elements, which are measured. For example, EPMA analyses calculating water by difference without a subsequent iterative matrix correction produce water estimates that are low by as much as 1 to 2 wt% absolute (Roman et al., 2006). This is equally true for other unanalyzed elements (Fournelle et al., 2020). Missing elements also

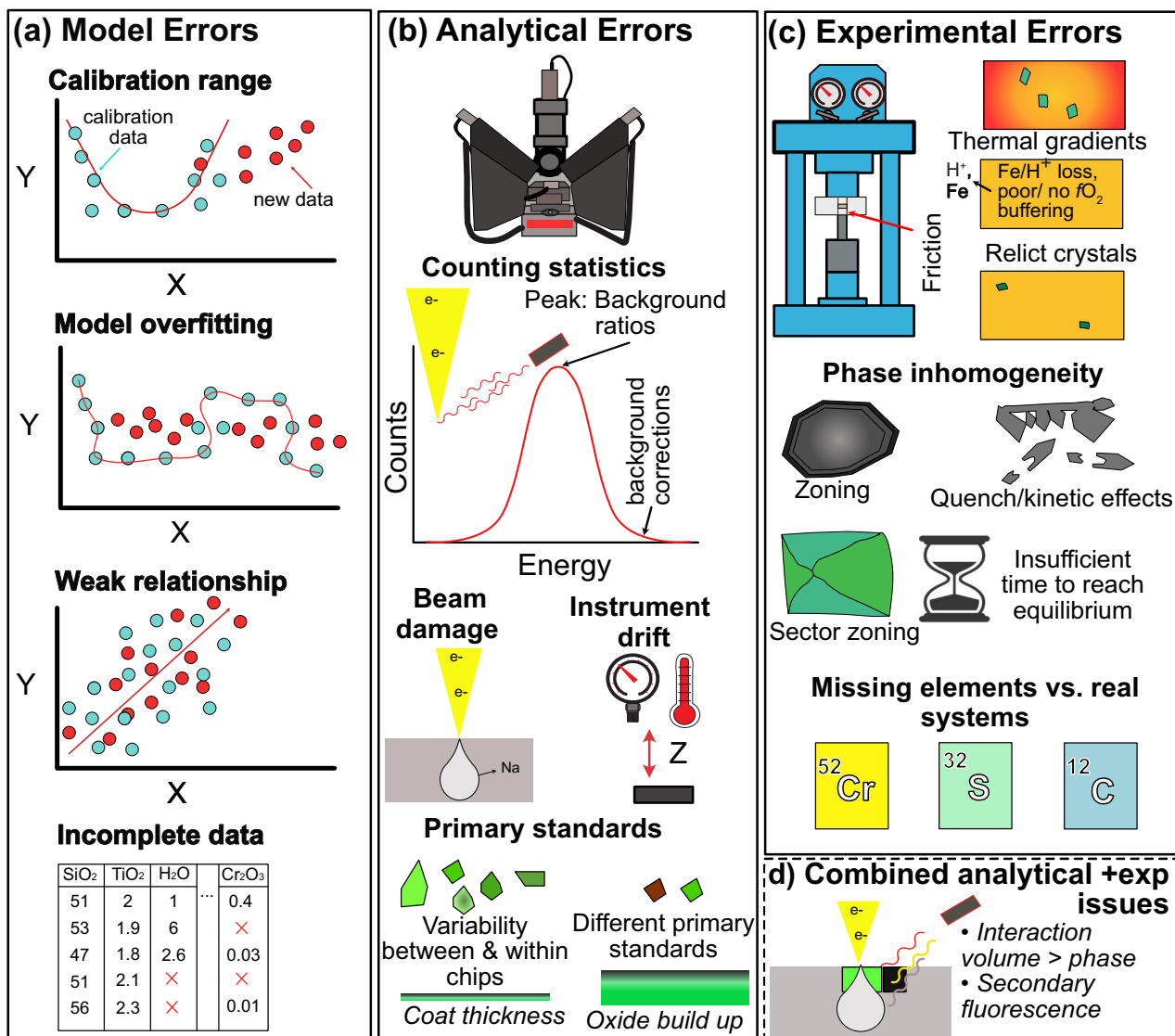


Fig. 1. Schematic diagram showing the three main categories of uncertainty associated with thermobarometry.

cause issues during regression analysis. It is generally considered that up to 5% missing data on each attribute (i.e. a single oxide) are manageable during regression, 5%–15% requires use of more sophisticated approaches, and >15% can severely impact the prediction accuracy of the model on unseen data (Pyle, 1999; Twala, 2009). Unfortunately, the percentage of missing data in experimental compilations used to calibrate barometers is well within the severe category. For example, 66% of experiments in the LEPR dataset used by Petrelli et al. (2020) and 73% in Putirka (2008) do not report H₂O contents in the glass phase, and 42% of Putirka (2008) do not report Cr₂O₃ in the Cpx. Yet, both these elements feature heavily in parameterizations of Cpx-only and Cpx-Liq barometry, and the presence/absence of Cr actually affects phase stability (discussed in detail below, Voigt et al., 2017).

There is abundant literature, but little consensus, about how to deal with missing values during regression analysis (Emmanuel et al., 2021). The simplest way is list-wise deletion, where experiments with any missing values for any oxides are simply removed from the calibration dataset. However, in the experimental datasets used to calibrate thermobarometers, which have large amounts of missing data, list-wise deletion can

result in a dramatic reduction in the number of observations used for calibration. For example, 54% of experiments in the calibration dataset of Petrelli et al. (2020) do not report P₂O₅ in glass, 61% do not report Cr₂O₃ in glass, and 71% do not report K₂O in Cpx. List-wise deletion of all oxides used in the regression model would leave only 7/850 experiments for Cpx-Liq and 80/850 for Cpx-only, both of which are very small calibration datasets. The software JMP used by Putirka (2008, and previous papers) uses list-wise deletion by default; the shrinking dataset issue was mitigated by excluding elements from the calibration, which results in lots of rows being deleted, but likely do not have a major influence on the equilibria of interest (e.g. MnO in Liq, K₂O in Cpx).

List-wise deletion can be very problematic if missing values reflect an underlying bias in the measurement procedure (termed ‘missing not at random’ by Rubin, 1976). For example, it is far more likely that H₂O-poor or nominally anhydrous experiments do not report H₂O in experimental glasses compared with very water-rich experiments. Removing experiments with no reported water could easily bias the remaining dataset toward compositions and phase assemblages found in more hydrous magmas, meaning the resulting model would be poorly calibrated for relatively anhy-

drous melts. For H₂O-poor experiments, it is far better to enter H₂O = 0 into the dataset than to remove the experiment. Missing H₂O data has been dealt with in several ways. Petrelli et al. (2020) fill all missing values for H₂O (and any other elements) with zeros.

Alternatively, missing values can be imputed (i.e. estimated). For example, Putirka (2008) replaced missing H₂O data with 100 minus the sum of major element totals (i.e. volatiles by difference). While this method is associated with large uncertainty unless calibrated using hydrous standards (Hughes et al., 2019), it is likely better than replacing missing values with the mean or median of the dataset (which can create issues when data are not missing at random, as discussed above for H₂O; Twala, 2009).

The presence/absence of reported Cr₂O₃ (or Cr) data in Cpx demonstrates one challenge associated with imputation for experimental datasets. In many cases, experiments on synthetic starting materials do not report Cr concentrations in Cpx or Liq because no Cr was added to the starting material (e.g. Hamada & Fujii, 2008), whereas in other studies, Cr was likely present but not reported. For example, Parat et al. (2014) use a natural starting material with ~187 ppm Cr, but report no Cr₂O₃ data in Cpx or glasses. Using an approximate partition coefficient (~10, Hart & Dunn, 1993), their Cpx could contain ~0.1–0.2 wt% Cr₂O₃. Finally, there are experiments where it is unclear whether Cr is present or not. For example, the natural starting materials of Cadoux et al. (2014) are sufficiently evolved (67–71 wt% SiO₂) that it is unlikely there is very much Cr left, but the actual Cr content of the starting material (and experimental Cpx) are not reported.

The paucity of Cr data (resulting from Cr-free starting materials and the lack of Cr measurements in Cr-bearing starting materials) has been discussed in detail in the context of MORB differentiation and liquid-barometry by Voigt et al. (2017). Voigt et al. (2017) showed that Cr-containing experiments stabilize Cpx at higher temperatures than Cr-free experiments. Pressures calculated using their melt geobarometer for Cr = 0 versus true Cr contents in MORBs differ by up to 1.5 kbar. Thus, replacing missing values with the mean of the dataset, or any other prediction, would fail to capture the fact that there truly was no Cr, which has likely influenced Cpx stability. We quantitatively evaluate the effect of missing Cr data on the Cpx-only and Cpx-Liq thermobarometers in the discussion section.

Analytical uncertainty

In addition to thermodynamic considerations (larger changes in entropy compared with volume), the larger percentage errors associated with barometers compared with thermometers may result from the lower concentration of elements that are important in barometry compared with thermometry, meaning analytical errors are larger (e.g. Na in Cpx for pressure vs. Mg-Fe for temperature). A number of random and systematic uncertainties can arise during EPMA analyses. Each wavelength dispersive spectrometer (WDS) is calibrated for a specific element using the relationship between the peak-background ratio and the concentration of that element in a primary standard material. In turn, this relationship is used to predict the element concentration in an unknown material based on the measured peak-background ratio. A matrix correction is applied to account for the influence of the specific material analyzed on the intensity of measured X-rays (e.g. atomic number effects, absorption, and fluorescence).

The generation of X-rays from a sample excited by an electron beam is a random process (like radioactive decay), meaning that the emission of any given X-ray cannot be predicted. However, if X-ray arrivals are collected over a long enough time interval, the average number of X-ray arrivals per unit time is a function

of the rate of X-ray production. X-ray production is determined by the specific element and electron shell ionization efficiencies, element concentration, beam current, beam voltage and detector take off angle. The instrumental efficiency of X-ray detection is controlled by the WDS crystal size/geometric efficiency and detector efficiency (as well as the vacuum). X-ray intensities are then normalized to beam current, yielding counts per second per nA. Instrumental specific effects are accounted for when converting counts into concentration by performing the X-ray measurements using the same beam energy and spectrometer on both the primary standard and the unknown material to produce a k-ratio, which should be identical within measurement error for all instruments using the same element, X-ray emission line, beam energy and takeoff angle (assuming the high voltage calibration and effective take off angle of the spectrometers are sufficiently similar).

The precision of an EPMA measurement is determined by the total accumulated number of X-rays counted by the WDS detector at the peak and background positions. Errors directly resulting from the fundamentally random process of X-ray generation and detection are termed counting statistics. When averaged over sufficient time period, X-ray counts follow a Poisson distribution, which approximates a normal distribution at sufficiently high counting rates (i.e. the central limit theorem). Errors resulting from counting statistics can be estimated using two main methods.

Equation 2 can be used to predict the relative error (100*standard deviation (σ)/concentration) for a given element in an unknown sample using information from count rates in the primary standard (Weill et al., 2004), the relative concentration of the element in the primary standard and sample, and the time spent counting:

$$\text{Error (\%)} = 100 \times \sqrt{\frac{X_{\text{sample}}(P_{\text{std}} - B_{\text{std}})}{X_{\text{std}}} + 2B_{\text{std}}}{\left(\frac{X_{\text{sample}}(P_{\text{std}} - B_{\text{std}})}{X_{\text{std}}}\right)^2 t}}, \quad (\text{Equation 2})$$

where P_{std} and B_{std} are the counts per second (cps) of the peak and background for the element in the primary standard, X_{std} is the element concentration in the primary standard (wt%), X_{sample} is the concentration in the unknown sample (wt%), and t is the count time on the unknown sample (s).

Equation 2 assumes that the total count times on the peak and background are the same and does not account for the different matrices of the primary standard and the sample. It can be made more versatile by including a factor for the probe current (I), substituting counts per second on the primary standard with counts per second per nA ($P_{I, \text{std}}$ and $B_{I, \text{std}}$):

$$\text{Error (\%)} = 100 \times \sqrt{\frac{X_{\text{sample}}(I P_{I, \text{std}} - I B_{I, \text{std}})}{X_{\text{std}}} + 2 I B_{I, \text{std}}}{\left(\frac{X_{\text{sample}}(I P_{I, \text{std}} - I B_{I, \text{std}})}{X_{\text{std}}}\right)^2 t}}. \quad (\text{Equation 3})$$

When the measured total counts ($P_{\text{Tot, sample}}$) on the peak and background ($B_{\text{Tot, sample}}$) are known, a more precise way of calculating the relative error for a specific element is given by

$$\text{Error (\%)} = 100 \times \sqrt{\frac{\frac{P_{\text{Tot, sample}}}{t^2} + \frac{B_{\text{Tot, sample}}}{t^2}}{\left(\frac{P_{\text{Tot, sample}}}{t} - \frac{B_{\text{Tot, sample}}}{t}\right)}}. \quad (\text{Equation 4})$$

Equations 2–4 yield very similar results (often within ~1%), with slight discrepancies resulting from the fact that Equation 4 accounts for differences in the matrix of the sample and standard,

which affects the intensity of generated X-rays, and slight differences in the background count rate on the sample and standard. The main advantage of Equations 2–3 is that they can be used predictively to assess the approximate precision on an unknown using count rates from primary standards on a given spectrometer and crystal. Equation 4 is best applied after an analysis has been conducted, and is what most commercial EPMA software uses to output 1 sigma errors for a specific analysis.

Uncertainties resulting from counting are almost always random (i.e. following a Poisson distribution), whereas instrument calibration can introduce systematic errors. Historically, because the accuracy of the physical models used for performing matrix corrections were limited (though today that is generally no longer the case), EPMA analyses of geological materials have tended to use primary standards with similar matrices to the target analytes (e.g. Na in Albite for analysis of Na in feldspar, Na in jadeite for analysis of Na in pyroxene). The most common primary standards are from the Smithsonian National Museum of Natural History (NMNH) collections (Jarosewich *et al.*, 1980; Jarosewich, 2002). However, it is well documented that different chips of the same natural NMNH standard may differ from published compositions determined by wet chemistry, and that heterogeneity can also exist within a single chip (Jarosewich *et al.*, 1980; Rose *et al.*, 2008; Fournelle, 2012). As each EPMA laboratory has its own set of primary standards, for any given named standard (e.g. San Carlos olivine), heterogeneity between the different chips in different laboratories can result in systematic offsets between measurements made in these laboratories. Different laboratories also rely on different primary standards to calibrate a specific element, increasing the variability. Some laboratories also utilize 'home grown' materials obtained from their mineral collections or synthetically produced 'in-house' standards, and some commercially available standard materials may not be what are advertised. For example, some commercial 'San Carlos olivine' has not been split from the Smithsonian mineral collection, but rather was merely collected from the same geographical vicinity.

The resulting systematic offsets in instrument calibration can be identified using round-robins, where the same material is analyzed as an unknown in different laboratories using their routine calibration strategy. Hunt *et al.* (1998) distributed a Lipari Obsidian sample to five laboratories, and Kuehn *et al.* (2011) distributed three different glass compositions (rhyolite, phonolite, basalt) to 27 laboratories. The Kuehn round-robin identified a number of outlier laboratories with large systematic errors, especially with regards to the correction of time dependent intensity (TDI) effects, sometimes referred to as volatile element or beam damage effects. These changes in emitted intensity are usually due to ion mobility within the interaction volume (Morgan & London, 2005; Hughes *et al.*, 2019) and are observable on not only highly mobile alkali elements such as Na and K (which can dramatically decrease in intensity during the measurement and therefore must be extrapolated back to their zero time intensities), but also for less mobile elements such as Si and Al, whose apparent intensities increase as the alkali element intensities decrease due to migration toward subsurface charge accumulation. Such TDI corrections can range from under a few percent (relative) to 30%–100%. Thus, changes in counts during beam exposure should be carefully inspected for each element in representative samples when performing a study on hydrous/alkaline melt compositions and used to inform decisions about which elements to perform TDI corrections for. As many Cpx-Liq thermobarometers are parameterized in terms of Na and K (and Si and Al) in the liquid (e.g. Neave & Putirka, 2017, eq30–31 of Putirka, 2008), different TDI correction routines

used to measure different experiments could add significant scatter to an experimental dataset compiled from many different laboratories.

At various points over the last few decades, authors have used smaller round-robins to identify systematic offsets in glass measurements and have corrected data compiled from these different laboratories to produce a more consistent calibration dataset. For example, Yang *et al.* (1996) performed corrections to glass data collected at both Massachusetts Institute of Technology (MIT) and the Smithsonian prior to performing modeling and thermobarometric calculations. Most recently, Gale *et al.* (2013) performed a round-robin using VG2 to obtain correction factors for MORB glasses relative to EPMA analyses from Lamont Doherty Earth Observatory (hereafter, Lamont; Fig. 2). To our knowledge, no such reconciliation has been performed for datasets used to calibrate Cpx-based barometers (or any other mineral barometers), so we quantify the possible effects of interlaboratory offsets in the discussion section.

Additional systematic and random analytical uncertainty during EPMA analyses can result from problems relating to beam damage/charging of materials under the electron beam, poor sample preparation (e.g. bad polishing, sample tilt) and drift of the instrument (e.g. stage Z height, changes in the temperature and pressure in the room; Fig. 1b). It should also be mentioned that other sources of systematic analytical inaccuracy can be introduced from improperly calibrated instruments, for example assumed dead time constants and picoammeter readings from poorly maintained instruments. However, these are almost impossible to quantify retrospectively from published data, so are not discussed further.

Experimental scatter

Analysis of the experimental products used to calibrate thermobarometers are subject to the analytical uncertainty discussed above, along with several other sources of uncertainty (Fig. 1c). Firstly, experimental phases may be heterogeneous because of zoning resulting from changing P-T-fO₂-H₂O conditions during the run, remnant seed crystals, or kinetic effects during growth and sample quench. Capsules may also experience Fe and H⁺ exchange with samples and assemblies (Gaetani & Grove, 1998; Botcharnikov *et al.*, 2005), resulting in poor redox buffering. Relict crystals from the starting material may be present if the starting material was not fully melted (Mutch *et al.*, 2016). The relatively small size of experimental products also enhances EPMA uncertainty relative to natural crystals, because the interaction volume of the electron beam may directly enter other phases, and neighboring phases may influence the measurement through secondary fluorescence (Llovet & Galan, 2003; Fig. 1d).

There are also a number of sources of uncertainty that can arise from the experimental set up. Experiments conducted at <5–7 kbar were mostly performed using TMZ/MHZ vessels (Sisson and Grove, 1993a, 1993b) and internally heated pressure vessels (IHPV, e.g. Cadoux *et al.*, 2014; Di Carlo, 2006; Hamada & Fujii, 2008; Parat *et al.*, 2014). Pressure in IHPV experiments is monitored with strain gauge manometers, Heise Bourdon tube gauges, or digital pressure transducers, which have a precision of ~0.01–0.04 kbar (1–4 MPa; Koepke *et al.*, 2018; Pichavant, 2002). Pressure variations during experiments are mostly within ~0.05 kbar (e.g. Parat *et al.*, 2014). Because pressure is transmitted via a gas to the capsule, excluding calibration offsets in gauges/manometers/transducers, an experimental pressure recorded in one lab is likely to be similar to that in another lab.

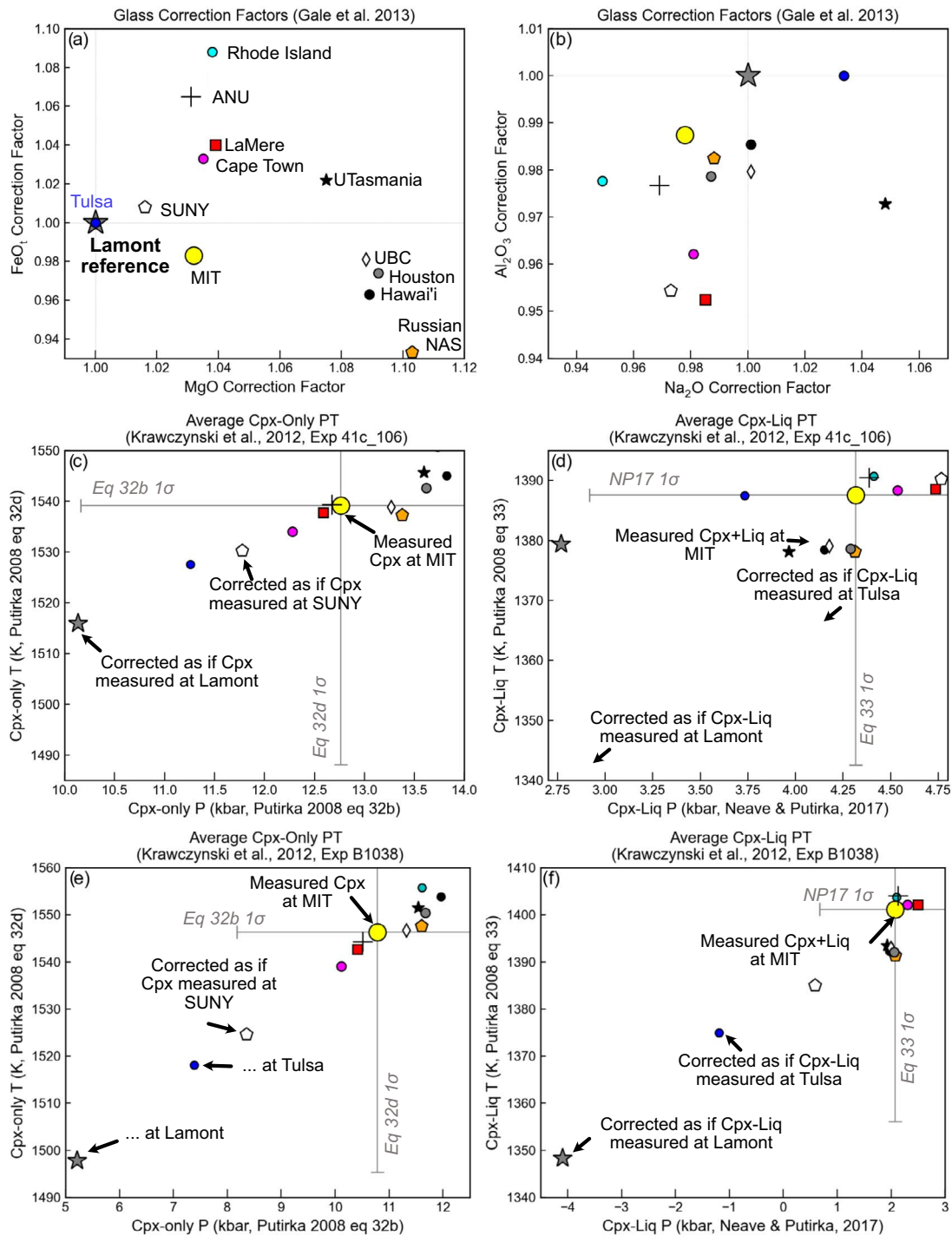


Fig. 2. Assessing the influence of interlaboratory biases on Cpx-only and Cpx-Liq thermobarometry. (a,b) Interlaboratory correction factors from Gale *et al.* (2013) relative to Lamont (plotting at 1, 1). (c,d) Calculated Cpx-only and Cpx-Liq pressures and temperatures for the average reported composition from Experiment 41c-106 of Krawczynski *et al.* (2012, 4.9 kbar, 1248.15 K), corrected as if Cpx and Liq compositions were measured in different laboratories. (e,f) as for (c) and (d), using Experiment B1038 (8 kbar, 1323.15 K). Additional experiments are shown in Supplementary Figs 2–4. Commonly stated SEEs for each thermobarometer are shown for comparison. The SEE for eq32b is the fit to $N = 1173$ data, eq32d is the fit to $N = 910$ anhydrous experiments, and eq33 is the fit to $N = 1174$ data (all from figures in Putirka, 2008). The SEE for Neave & Putirka (2017) is that given in their abstract.

In contrast, most experiments conducted at >5 kbar were performed in piston cylinders (Baker & Eggler, 1987; Gaetani & Grove, 1998; Mercer & Johnston, 2008), where a solid-medium transmits pressure to an enclosed capsule. Pressure in the piston cylinder apparatus is commonly monitored using a Heise gauge, and maintained to within ± 0.5 kbar (Hamada & Fujii, 2008).

In solid-medium experiments the pressure experienced by the experimental capsule can differ from the pressure applied to the hydraulic piston (measured with the Heise gauge), with the correction between the two often referred to as a friction correction (Tamayama & Eyring, 1967; Condamine *et al.*, 2022). This friction/pressure correction can be evaluated for each

experimental set up by running experiments for reactions, which are well constrained in P–T space, and applying a shift to account for any offsets between the observed and predicted transition.

However, different laboratories investigating Cpx equilibria correct for friction in different ways, which could easily introduce bias to an experimental dataset compiled from many different studies. For example, Mercer & Johnston (2008) and Draper & Johnston (1992) do not include a friction correction. Likewise, Bartels *et al.* (1991) do not apply a correction, because their offsets calculated from the melting point of Au are only 0–1 kbar, which is within the uncertainty of the Au calibration (Akella & Kennedy, 1971). Grove *et al.* (2003) corrected for the offset using the reaction between anorthite+gehlenite+corundum \rightarrow Ca-tshermakite (now kushiroite, Hays, 1966), and although they do not give the magnitude of the offset, they stated that after this correction pressures are accurate to ± 0.5 kbar. Gaetani & Grove (1998) used this Ca-tshermakite/kushiroite reaction in addition to the melting point of Au, resulting in a friction correction of 300 MPa. In contrast to this constant offset applied to 12, 16 and 20 kbar experiments, Hamada and Fujii (2008) apply a –10% correction for their 4 kbar experiments, and a –15% correction for their 7 kbar experiments based on a calibration using the water solubility of albite melt (Behrens *et al.*, 2001). Ulmer *et al.* (2018) stated that they follow the experimental approach of Villiger (2004), who applied a correction of -3% calibrated against the quartz-coesite transition at 3.07 GPa and 1000°C (Bose and Ganguly, 1995) and fayalite+qtz \rightarrow orthoferrosilite reaction at 1.41 GPa and 1000°C (Bohlen *et al.*, 1980). Finally, Blatter *et al.* (2013, 2017) calibrated at each pressure by bracketing the CsCl melting curve (Bohlen & Boettcher, 1982).

The wide variation in the reactions chosen for calibration (and the variation in the pressures and temperatures of these calibration reactions) means that it is perfectly plausible that systematic offsets still exist between different laboratories, up to the magnitude of the corrections applied (e.g. $\sim 10\%$ – 15% , or a few kbar). For example, Johannes *et al.* (1971) showed that measurements of the pressure of the albite \rightarrow jadeite+quartz reaction at 600°C using the same starting material in six different laboratories varied from 15.7 to 16.8 kbar. Additionally, Condamine *et al.* (2022) showed that friction can decrease during a run (from 29% in 6 hr runs to 21% in 24 hr runs), which means that the true pressure experienced by the capsule may also be influenced by the length of the experimental run, which varies greatly between studies. Johannes *et al.* (1971) also pointed out that the friction correction can also vary between runs conducted with identical protocols, because of a softer than average batch of the solid-medium material, or more efficient wall lubrication. This means that a piston cylinder experiment reported at 10 kbar in one laboratory may not have experienced the same pressures as an experiment performed at 10 kbar in another laboratory, and variations may exist even within experiments run at 10 kbar in a single laboratory. Uncertainties in the true pressure experienced by the charge explains why experimentalists tend to space out their piston-cylinder pressures within a single study, to ensure that trends are at least coherent. However, when all these data are compiled to calibrate a barometer, offsets on the order of 0.5–1 kbar will increase scatter, and could create systematic uncertainties based on different laboratories investigating different regions of PT space.

Temperature variations in piston cylinders are generally measured with thermocouples (e.g. Pt–Rh, Baker & Egger, 1987, W-Re, Mercer & Johnston, 2008), although the exact geometry varies in different laboratories. Although thermocouples are reasonably

precise and accurate ($\pm 10^\circ\text{C}$), the larger source of error results from thermal gradients within the piston cylinder, requiring a correction to be made to account for the higher temperatures in the hotspot where the experimental capsule is placed versus the position of the thermocouple tip. These corrections depend on the capsule geometry and assembly, but can easily reach 20–40°C (Grove *et al.*, 2003; Brugman *et al.*, 2021). Variable thermal gradients between the capsule and thermocouple could result in systematic differences between runs and laboratories, and thermal gradients within capsules themselves (Harlow, 1997) can cause crystals grown in different parts of the capsule to show different compositions (increasing the compositional scatter seen in experiments). Putirka *et al.* (1996) stated that the most consistent pyroxene analyses are when rim compositions are analyzed along with the neighboring liquid interface, perhaps helping to mitigate the effect of thermal gradients (and thus compositional variations throughout the capsule).

DISCUSSION

Quantifying the effect of interlaboratory EPMA analysis offsets

The dataset of experiments from LEPR used by Putirka (2008) to calibrate their Cpx \pm Liq thermobarometers lists 46 unique experimental laboratories, with a relatively small number of laboratories performing a large proportion of experiments (MIT = 25%, Penn State = 7%, Lamont = 6%, Hannover and University of Oregon = 4–5%). If all laboratories investigated the same P–T–X conditions, and interlaboratory offsets were normally distributed about the nominally correct value, inter-laboratory EPMA offsets would simply add random uncertainty to calibration and test datasets. However, as different experimental laboratories, which mostly use a specific EPMA instrument, tend to target different research questions, the occurrence of these analytical offsets in the experimental dataset is not uniformly distributed, and could thus lead to systematic uncertainty. For example, in the LEPR dataset, at pressures > 10 kbar, experiments performed at Australia National University (ANU) tend to focus on higher SiO₂ liquids than experiments performed at MIT or University of Tasmania (Supporting Fig. 1–2). Thus, interlaboratory offsets mean the calibration could be skewed as a function of liquid composition.

Interlaboratory offsets can also affect the statistics determined from a test dataset if a specific laboratory has an offset relative to the average offset of the LEPR dataset (or even the average value in a specific P–T–X region of calibration dataset). It is noteworthy that the relative contribution of different laboratories at crustal pressures (0–13 kbar) has changed dramatically between our newly compiled ArcPL dataset and LEPR. For example, experiments conducted at Hannover account for 40% of the ArcPL dataset, but just 4% of the LEPR dataset, while MIT comprises 25% of the LEPR dataset and 10% of the ArcPL dataset. If, say the EPMA lab at Hannover (or any other laboratory that has greater output since LEPR was compiled) had an offset from one of the dominant EPMA in the LEPR dataset used for calibration of many published barometers, this could help to account for the fact that new experimental data are predicted poorly by these barometry calibrations.

In general, interlaboratory analytical differences from glass round-robins are <10% (Hunt *et al.*, 1998; Kuehn *et al.*, 2011; Gale *et al.*, 2013), although occasionally it is noted that one or more laboratories produce discrepancies >10% (their identity is kept anonymous in Hunt *et al.*, 1998). Compared to other analytical

techniques, relatively little attention has been paid to interlaboratory EPMA offsets in petrology, with most studies being conducted on silicate glasses by the tephrochronology community (Hunt et al., 1998; Kuehn et al., 2011). To our knowledge, no round-robins have been conducted on pyroxene or other silicate minerals. As many of the EPMA used to measure the pre-2008 experiments in the LEPR dataset have been decommissioned, we conclude that we will never be able to fully determine the exact value of offsets for Cpx. In the absence of other data, we assume that the interlaboratory offsets for VG2 glass reported by Gale et al. (2013; Fig. 2a-b) are a first-order estimate of the interlaboratory differences that may have occurred during Cpx analysis. The round-robin conducted by Gale et al. (2013) is particularly useful because it targeted a number of laboratories, which have analyzed experimental charges used to calibrate and test various thermobarometers.

To assess whether 0–10% offsets between different EPMA laboratories could adversely affect thermobarometric calibrations and assessment using test datasets, we consider the average reported Cpx and glass compositions from experiments from Krawczynski et al. (2012) analyzed on the MIT EPMA. We multiply these measured compositions by the correction factors from Gale et al. (2013) to obtain the compositions that would have been measured on the Lamont EPMA. Using the interlaboratory comparisons to Lamont from Gale et al. (2013), we also estimate the composition that would have been measured at a number of other EPMA labs. We then calculate pressures and temperatures for these corrected compositions by iterating eq32d (T) and eq32b (P) for Cpx-only thermobarometry, and eq33 (T) and eq31 (P) for Cpx-Liq thermobarometry (equations from Putirka, 2008).

The range of calculated pressures resulting from these laboratory offsets is significant (Fig. 2c–f), and varies greatly between different experiments and different thermobarometry expressions (Fig. 2, Supporting Figs 3–5). Cpx-Liq pressures that would have been calculated from mineral compositions measured in different laboratories show offsets from one another of ~0.5 to 6 kbar (Fig. 2d,f, Supporting Figs 3–5), while Cpx-only pressures show offsets of ~3 to 5 kbar (Fig. 2c,e, Supporting Figs 3–5). Interlaboratory offsets also impact calculated temperatures (~10–50 K), but to a lesser extent. These interlaboratory offsets likely contribute to the overall scatter between calculated versus experimental pressures in calibration and test datasets. In fact, it is noteworthy that our calculated offsets are similar in magnitude to quoted SEEs on barometers. These offsets also affect natural samples that barometry equations are applied to; it is highly concerning that calculated pressures may vary by 3–5 kbar based solely on the EPMA laboratory used to perform analyses.

Variability in treatment of chromium

As discussed in the introduction, many experiments in the calibration dataset do not report Cr data, yet the Cpx-only thermobarometers of Wang et al. (2021, eq1, 2, 32dH), Putirka (2008, eq32b, 32d) and Petrelli et al. (2020) include a term for the Cr content of the Cpx. For Cpx-Liq thermobarometry, only the model of Petrelli et al. (2020) is parameterized directly in terms of the Cr content of the Cpx and the liquid. However, Cr in the Cpx is used to calculate the CrCaTs component of the Cpx, which is then used to calculate the diopside-hedenbergite (DiHd) component:

$$\text{CrCaTs} = 0.5 * \text{Cr}_{\text{cat frac}}$$

$$\text{DiHd} = \text{Ca}_{\text{cat frac}} - \text{CaTs} - \text{CaTi} - \text{CrCaTs}$$

The DiHd component is included in the Cpx-Liq barometers of Putirka (2008, eq30 and 31), Neave & Putirka (2017), and the Cpx-only barometers of Putirka (2008, eq32a, 32b). Additionally, even Cpx-Liq or Cpx-only barometers that have no compositional term dependent on Cr in Cpx are normally parameterized in terms of Cpx cation fractions, either directly or in terms of components like jadeite, which are calculated from cation fractions. If Cr is present in the sample, but not measured, the calculated cation fractions of other elements such as Na (and therefore jadeite) are artificially high. Thus, a wide range of barometers are directly or indirectly sensitive to the Cr content of the Cpx.

The proportion of missing Cpx Cr data is high in both the LEPR calibration dataset of Putirka (2008, 42%) and ArcPL (62%). Missing Cr data in the calibration dataset may have resulted in the relationship between components dependent on Cr and P and T being incorrectly parameterized (or correctly parameterized, but with the addition of significant noise). Missing Cr data in our test dataset will also cause noise when comparing predicted versus experimental pressures, because if Cr was present but not reported, calculated pressures and temperatures using true Cr contents may differ from those calculated using the Cr = 0 wt% (and would perhaps lie closer to the 1:1 line). To investigate the possible offsets caused by not reporting Cr when it is actually present, we calculate pressures and temperatures for all Cpx and Cpx-Liq pairs in the ArcPL dataset with reported Cr data, and compare this to calculations using Cr = 0 wt% (Fig. 3).

Different barometers show different sensitivity to Cr. For the Wang et al. (2021) Cpx-only barometer and thermometer, there is a clear correlation between the change in pressure and the Cr content of the Cpx, with the most Cr-rich Cpx showing pressures up to 2.8 kbar too high when Cr is set to 0 wt% (Fig. 3a). The effect on temperature is proportionally smaller (~35 K; Fig. 3b). Strong correlations with true Cr content are also seen for pressures calculated using Putirka (2008) eq32d-32b, with max offsets of 1.9 kbar (and 16 K for temperature). In contrast, the change in pressure using the Petrelli et al. (2020) Cpx-only barometer shows no apparent relationship to the Cr content of the Cpx, and shows significantly larger variations than the other barometers (max $\Delta P = 4.9$ kbar, $\Delta T = 97$ K). This unintuitive response reflects the use of decision trees, where the Cr content affects the route taken through the tree.

For Cpx-Liq thermobarometers, iteration of Putirka (2008) eq33 (T) with eq30 (P), or eq33 (T) with Neave and Putirka (2017, P, red triangles) shows very little sensitivity to Cr content (apart from a single Cpx-Liq pair showing a large change for all equations; Fig. 3c). Iteration of various thermometers with eq32c for pressure shows slightly more sensitivity, with offsets of up to $\Delta P = 1$ kbar and $\Delta T = 10$ K. As for their Cpx-only thermobarometer, the extra-trees regressions for Cpx-Liq of Petrelli et al. (2020) are slightly more sensitive to Cr, with temperatures varying by up to ± 23 K by changing the Cr content, and no clear correlation between the offset and the actual Cr content of the Cpx. Overall, these comparisons show that the presence/absence of Cr data in both calibration and test datasets can clearly introduce noticeable uncertainty in terms of calculated pressures (>1 kbar).

Quantifying the effects of analytical versus experimental variability in test and calibration datasets

Variability in measured phase compositions within a single experiment can also affect calibration and test datasets. Analytical uncertainty associated with the random process of X-ray generation can produce variability in measured phase compositions,

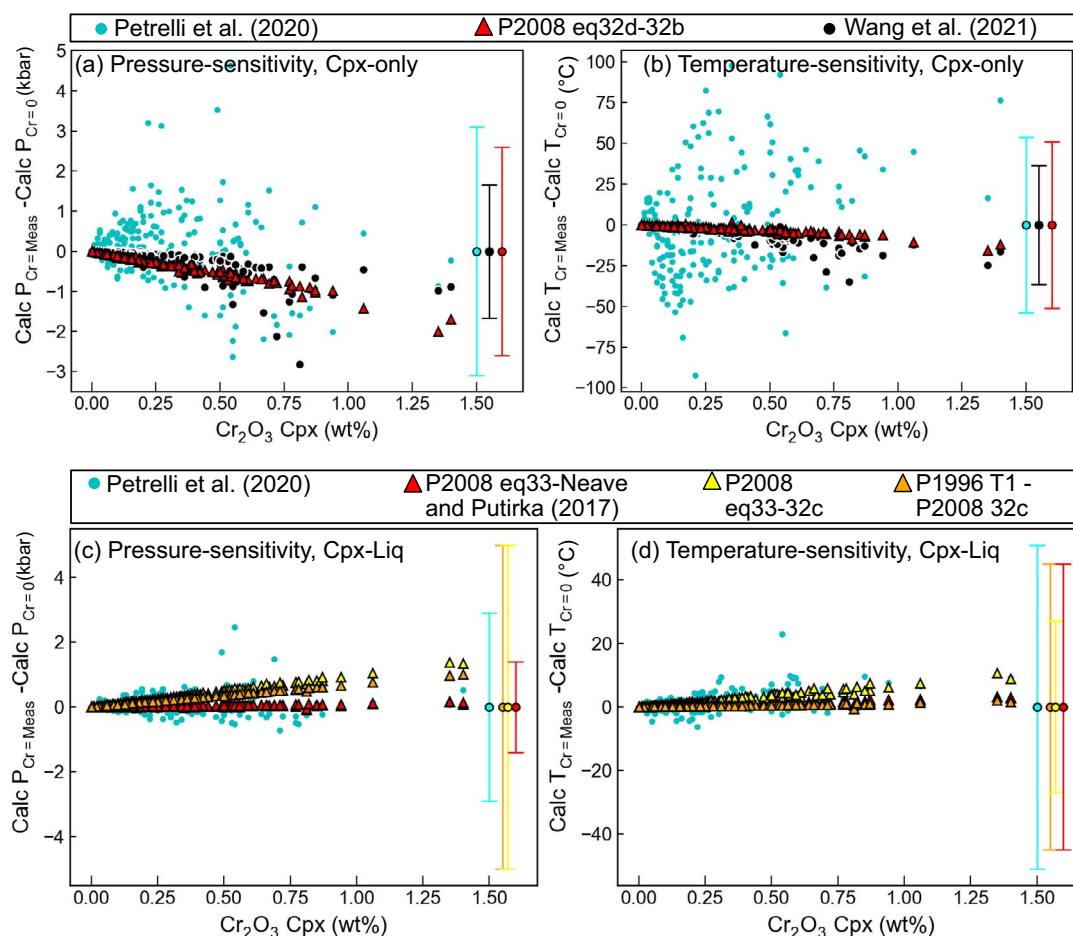


Fig. 3. For each of the $N = 490$ experiments in ArcPL, we calculate pressure (a,b) and temperature (c,d) using the measured Cr content (shown on the x-axis), and using Cr = 0 wt%. The difference between the calculation using measured Cr and Cr = 0 wt% is the y coordinate. Different colors and symbols represent different thermobarometry combinations used to calculate these pressures and temperatures. Commonly stated 1σ errors for these thermobarometers are shown for comparison. In addition to those described in the caption of Fig. 2 from Putirka (2008), eq32c is the fit to a global dataset, the SEE from Petrelli et al. (2020) are the fit to a test dataset, and the error from Putirka et al. (1996) eqT1 is that stated in the conclusion of that paper.

even if the experimental phases themselves are entirely homogeneous. Because analytical uncertainty resulting from counting statistics is normally distributed and random, if infinite numbers of compositionally homogenous Cpx crystals are analyzed from a single experiment, the mean composition will be the same regardless of the precision of each analysis. However, if only a very small number of Cpx are measured, analytical uncertainty can easily yield a reported average composition that is significantly different from the true composition. In the more likely scenario that experimental Cpx are also chemically heterogeneous (e.g. zoned or sector zoned), large numbers of EPMA analyses are required to correctly characterize the average composition of each experiment, even if each EPMA analysis is highly precise. For example, Neave et al. (2019) noted that their Cpx show highly variable Al and Ca contents, which they attributed to sector zoning. They suggested based on their observations of chemical heterogeneity that at least 20–40 analyses are required to meaningfully characterize the composition of Cpx.

Are we sufficiently averaging experimental and analytical variability?

To address the possible influence of analytical and experimental variability, we compile the number of reported Cpx analyses in the 490 experimental charges from the ArcPL dataset to investigate

how many publications approach the Neave et al. (2019) recommendation of 20–40 analyses. We supplement the ArcPL dataset with a compilation of Cpx measurements in 307 experimental charges in the LEPR dataset conducted at 0–13 kbar in arc-like compositions (ArcLEPR). We note that we do not know for each individual experimental charge whether the number of analyses reported in the associated paper represents measurements of N discrete Cpx crystals, or N analyses, with some Cpx being characterized by more than 1 EPMA spot (e.g. perhaps 10 analyses from 8 Cpx crystals).

When ArcPL and ArcLEPR are combined ($N = 797$), 44 experimental charges (5.5% of the combined dataset) were characterized by a single Cpx analysis. $N = 48$ experiments (6%) performed just 2 Cpx analyses, 64 (8%) performed 3 Cpx analyses, and 78 (10%) performed 4 Cpx analyses (Fig. 4a,b). Overall, 45% of the compiled experimental runs were characterized by ≤ 5 Cpx analyses, 25% by ≥ 10 Cpx analyses, and only 5% by ≥ 20 Cpx analyses (as recommended by Neave et al., 2019).

Concerningly, the experiments with the lowest number of analyses tend to be concentrated at lower pressures, where Na_2O contents are generally lower and more susceptible to large analytical uncertainties (Fig. 4a). It is notable that the discrepancy between the calculated pressure using the reported Cpx composition and the true experimental pressure increases with decreasing number

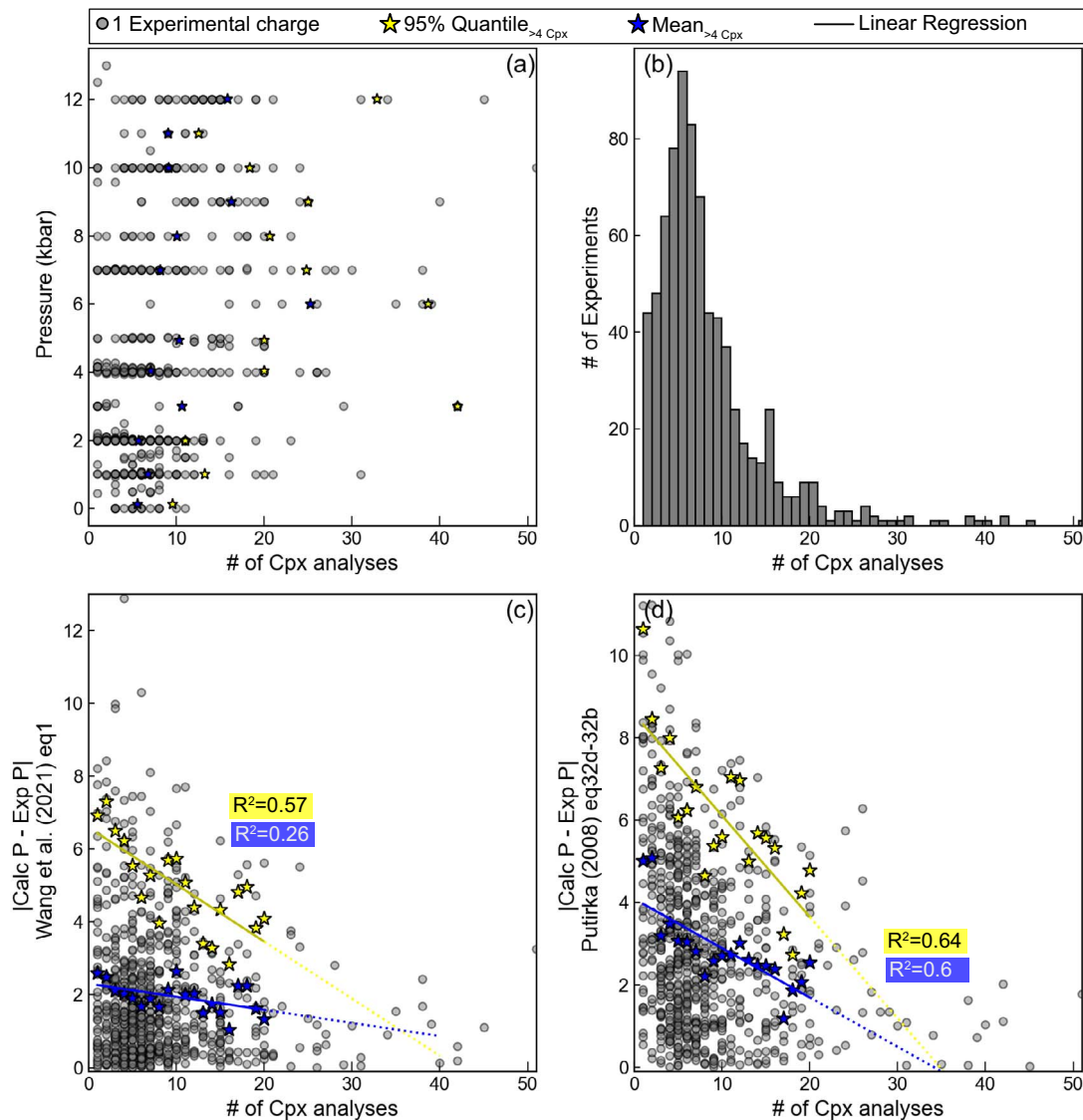


Fig. 4. Number of discrete Cpx analyses performed in each experimental charge plotted against experimental pressure (a) and as a histogram with a bin width of 1 (b). (c,d) For each experimental run, we calculate the absolute difference between the pressures calculated using the Cpx-only barometers of Wang *et al.* (2021) and Putirka (2008) eq32d-32b and the experimental pressure. In general, experimental runs with a smaller number of Cpx analyses show the largest absolute discrepancies. Symbols are semi-transparent, so darker colors represent a tighter clustering of data. The blue stars show the mean value for each discrete number of Cpx in (c) and (d), and for each pressure bin (0 ± 0.5 kbar, 1 ± 0.5 kbar, etc.), and the yellow stars show the 95% quantile.

of Cpx analyses (for the Wang *et al.*, 2021 and Putirka, 2008 Cpx-only barometers; Fig. 4c, d). This indicates that insufficient averaging of analytical variability or true phase variability (e.g. zoning) is affecting the performance of barometers. As the median number of analyses per experiment is 6 for both our new dataset and the subset of LEPR we have compiled the number of analyses for, we assume the statistics given above are representative of the entire LEPR database used for literature thermobarometry calibrations.

Assessing the relative influence of analytical uncertainty and experimental variability in experimental compositions is vital to address how barometry calibrations can be improved. If variability in measured Cpx compositions results from true variation in the composition of that phase (e.g. zoning and other disequilibrium processes), it means we must direct more attention to understanding and identifying disequilibrium in experimental products, measure very large numbers of phases

from each experiment (e.g. Neave *et al.*, 2019), and possibly redo experiments that have not sufficiently approached equilibrium (or remove these experiments from the calibration dataset). In contrast, if analytical variability is the primary culprit, it means we need to direct our attention to improving EPMA analyses of experimental products. Hereafter, we refer to true variation in Cpx compositions in experimental charges as ‘phase variability’ and variability resulting from EPMA analyses as ‘analytical precision.’

Distinguishing phase variability and analytical precision

Unfortunately, the information required to quantify the relative influence of phase variability versus analytical precision is largely absent from the information provided in most legacy experimental studies. In-text pdf tables are normally used to report the mean composition and standard deviation of each phase in each experimental charge. It is very unusual for the individual EPMA

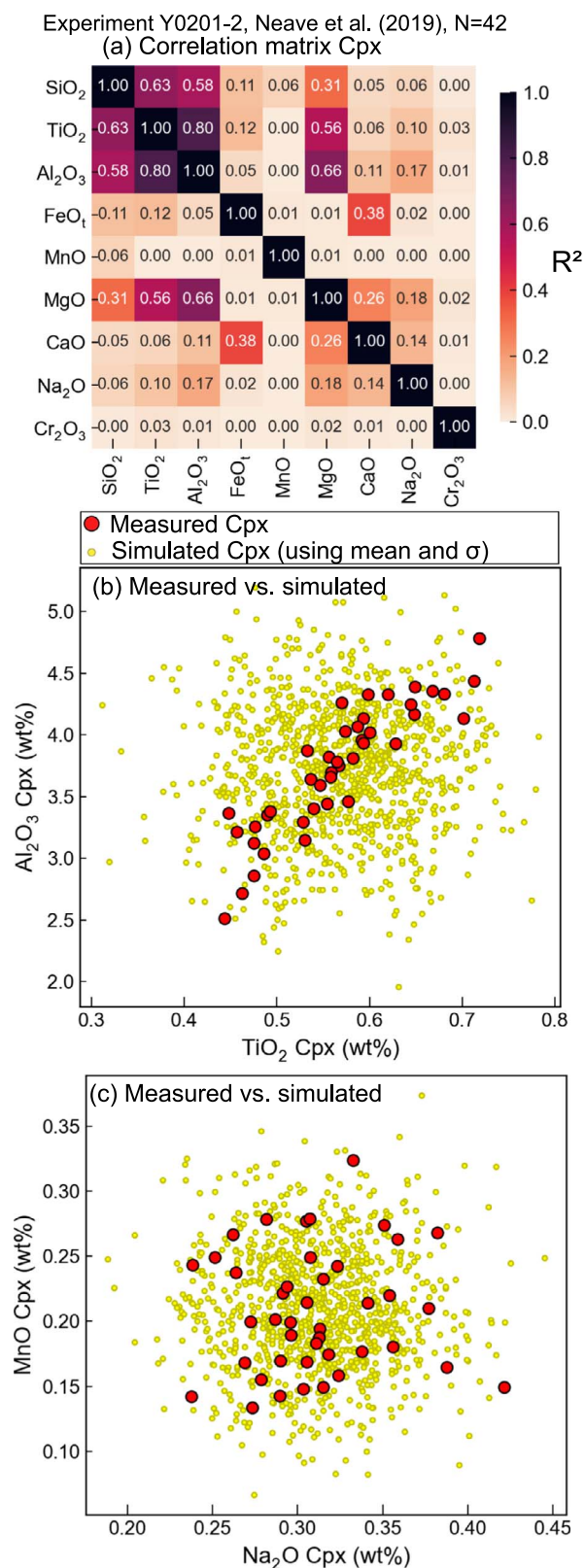


Fig. 5. (a) Correlation matrix showing the correlation coefficient (R^2) between different oxides in the 42 clinopyroxenes measured by Neave et al. (2019) in experiment Y0201-2, with cells colored based on this R^2 value. (b,c) Comparing measured compositions (red dots) to 500 synthetic Cpx analyses with major elements distributed normally using the reported mean and standard deviation of the 42 clinopyroxenes measured in the experiment (yellow dots).

analyses from each experiment to be reported, except in the most recent publications (e.g. Erdmann et al., 2016; Neave et al., 2019; Waters et al., 2021). This makes it difficult to assess the true variability in Cpx compositions, because the quoted mean and standard deviation for each phase in each experiment only provide a good description of the underlying data if it is normally distributed, and lacks covariance between oxides. Yet, almost all of the full experimental datasets we have been able to obtain (Krawczynski et al., 2012; Blatter et al., 2013; Melekhova et al., 2015; Neave et al., 2019) do not show normally distributed elemental variations, and exhibit significant correlations among different oxides (e.g. Fig. 5a, Supporting Fig. 6–15).

The covariance structure of oxides in each phase gives important clues into the source of variability in experimental products. For example, strong correlations between SiO₂-TiO₂, SiO₂-Al₂O₃, SiO₂-MgO, Al₂O₃-MgO in Cpx are indicative of sector zoning (e.g. Fig. 3b, Neave et al., 2019; Ubide et al., 2019, Supporting Fig. 6–14), whereas elements with variability arising from EPMA counting statistics will be uncorrelated (e.g. MnO-Na₂O; Fig. 5c). Importantly, when the full data are not reported, simulations using the reported mean and standard deviation provide a poor match to the real experimental data when a strong covariance structure is present (e.g. Fig. 5b, simulated Cpx as yellow dots, measured Cpx as red dots from Neave et al., 2019). The poor match of data simulated using just the mean and standard deviation means that it is nearly impossible to simulate the true variability in the LEPR dataset, where individual analyses could not be recovered.

Similarly, the information required to reconstruct the analytical precision for each element is seldom (if ever) reported. Ideally, authors would report the software-calculated sigma values for each analysis, which uses the number of counts on the peak and background of a specific EPMA spot to estimate the uncertainty related to counting statistics (Equation 4). Alternatively, more approximate estimates of error can be calculated from Equations 2–3, but this requires knowledge of the counts on the primary calibration standards, as well as the beam current and count times used for each element. We were unable to find any relevant experimental papers reporting software-calculated errors or sufficient information to use Equations 2–4. However, we were able to obtain software-calculated precisions upon request from the authors for a subset of experimental Cpx analyses from Krawczynski et al. (2012) conducted on the old MIT JEOL 733 and the newer MIT JEOL8200 (installed ~2007), and from Neave et al. (2019) on the Hannover Cameca SX100 (installed ~2001, decommissioned ~2021). We supplement these estimates with precision from natural pyroxene analyses on the University of Cambridge Cameca SX100 (Gleeson et al., 2021) and secondary standard analyses for the Kakanui Augite from the Oregon State University Cameca SX100. We also obtained peak and background counts from primary standard analyses on the MIT JEOL 733 (installed in the 80s) and the Bristol JEOL JXA8530F (installed ~2012) to model precision as a function of count time and beam current.

For the following discussion, we focus on Na to highlight an element affected by analytical error that is pertinent to the majority of Cpx barometry parameterizations, because of its relationship to jadeite content. As expected, compiled analytical precisions from different EPMA systems increase in magnitude as Na₂O content decreases below 1 wt% (Fig. 6a,c), closely following the trajectory of modeled precision curves. The exact trajectory of these precision versus concentration curves depends on spectrometer efficiency (ratio of peak: background counts in a primary standard), as well as the energy of the beam (voltage, current) and the count time. As the majority of Cpx analyses are performed at

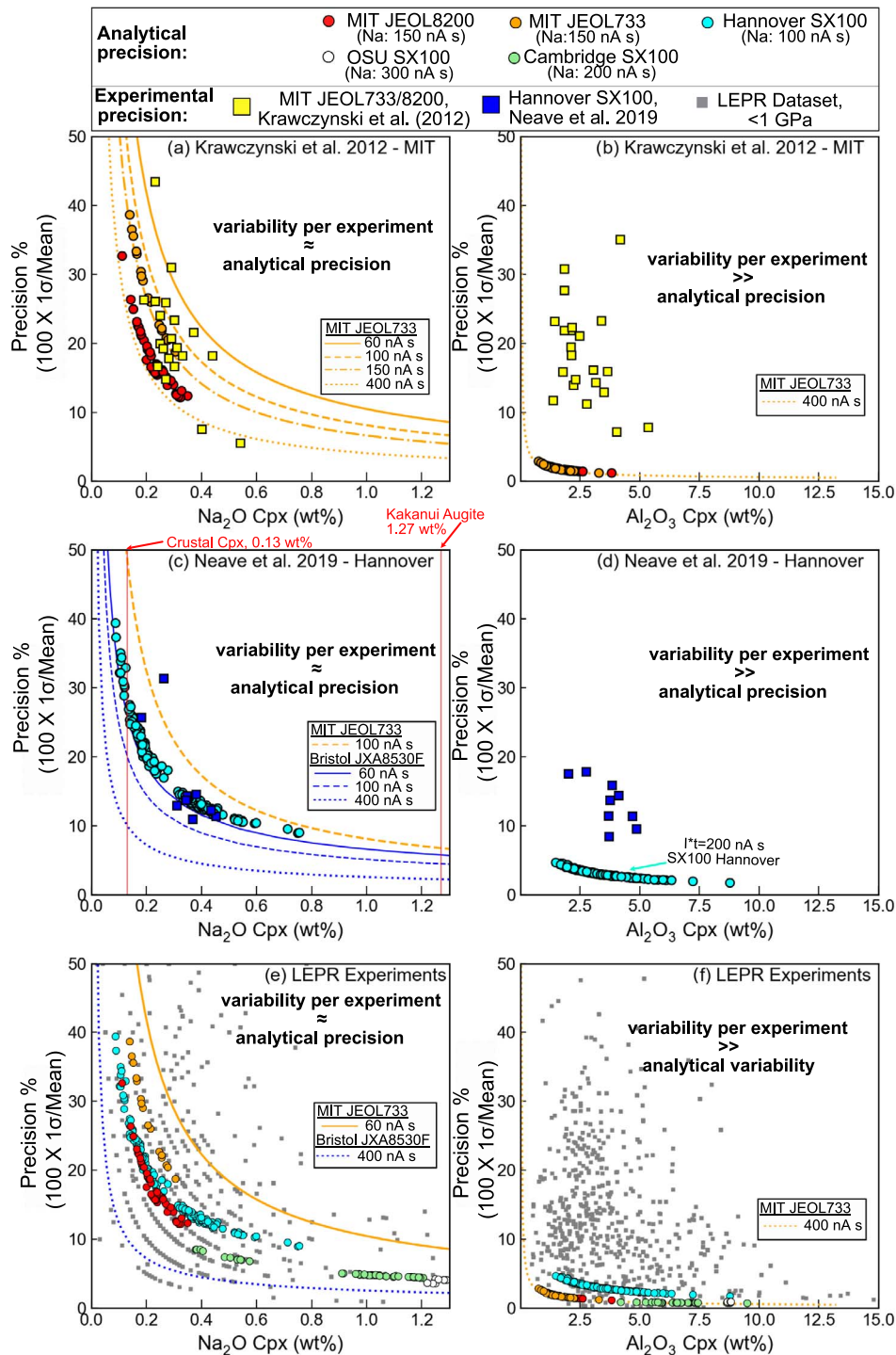


Fig. 6. Analytical precision as a function of Na₂O content and analytical conditions. (a) Theoretical precision versus concentration curves calculated for four different beam conditions on the MIT JEOL733 are shown in orange. EPMA software-calculated precisions for individual Cpx analyses are shown in red circles from Krawczynski *et al.* (2012) on the MIT JEOL8200 and orange circles from the older JEOL733. Phase variation/precision calculated from each experiment of Krawczynski *et al.* (2012; combining analyses from both probes) are shown as yellow squares. Analyses were performed with $I^*t = 150$ nA s. (b) Same for Al₂O₃. (c) Software-calculated precision for Cpx measurements from Neave *et al.* (2019) on the University of Hannover SX100 (cyan dots) with phase variation overlain (blue squares). The necessary count rates to use equation 2 could not be obtained from this instrument, so we overlay curves calculated using various analytical conditions for the old MIT JEOL733 and newer Bristol JEOL JXA8530F. (d) Same for Al₂O₃. Theoretical lines are not shown as there is substantially more variation in instrument efficiency and analytical conditions between different labs for Al compared to Na (See Supporting Table 1). (e, f) Experimental variability for Cpx in different experimental conditions reported in the LEPR dataset are shown as grey dots. We also overlay software-calculated precisions from MIT and Hannover, as well as measurements on the University of Cambridge Cameca SX100 from Gleeson *et al.* (2021) and measurements of Kakanui augite on the Oregon State Cameca SX100. We overlay theoretical precision lines representing best-case (newer Bristol JEOL JXA8530F, $I^*t = 400$ nA s) and worst-case scenarios (older MIT JEOL733, $I^*t = 60$ nA s).

15 kV, variations in beam current and count time can be combined into a single variable, $I \cdot t$ (10s and 20 nA gives identical counting statistics to 20 s and 10 nA).

Using count data from the Bristol JEOL JXA8530F TAP crystal for Na on an Albite primary standard, we calculate that measurement of a Cpx with $\text{Na}_2\text{O} = 0.13 \text{ wt\%}$ if $I \cdot t = 60 \text{ nA s}$ has a precision of 25% (e.g. 10 s, 6 nA), while measurement if $I \cdot t = 400 \text{ nA s}$ has a precision of 9.7% (e.g. 20s, 20 nA; Fig. 6c). For the MIT JEOL 733 with a TAP crystal calibrated for Na on Albite, the precision for $\text{Na}_2\text{O} = 0.13 \text{ wt\%}$ is 60% for $I \cdot t = 60 \text{ nA s}$, and 23% for $I \cdot t = 400 \text{ nA s}$ (Fig. 6a). The lower precision on the Bristol versus MIT JEOL for a given $I \cdot t$ value reflects the increase in detector efficiency on the newer Bristol instrument. This can also be seen by comparing the software-reported precision for the old MIT JEOL 733 (orange dots; Fig. 6a) to the newer MIT JEOL 8200 (red dots; Fig. 6a), where all other variables are kept constant (same calibration routine, same current and count times).

When information is available on both the software-reported analytical precision for each individual EPMA spot and the true phase variability observed in measured Cpx compositions within a single experimental charge, these can be compared. If variability for a specific element is dominated by analytical precision, the percentage variation calculated from multiple measurements of Cpx in that experimental charge ($100 \cdot \sigma / \text{Mean concentration}$) will be the same as the % precision value predicted from counting statistics. In contrast, if true phase variability is dominant (e.g. zoning, disequilibrium), the observed variability in measured Cpx compositions will greatly exceed the analytical precision. We were only able to obtain estimates of analytical precision for the experiments of Krawczynski *et al.* (2012) and Neave *et al.* (2019). In both sets of experiments, the analytical precision for Na_2O is very similar to the observed variability for measurements of Na_2O in Cpx (Fig. 6a, c). In contrast, the variability of Al_2O_3 in experimental Cpx greatly exceeds the analytical precision, indicating true phase variability (Fig. 6b, d). In general, variability in elements with $>1 \text{ wt\%}$ concentration in the experiments of Neave *et al.* (2019) and Krawczynski *et al.* (2012) significantly exceed the analytical precision, while lower abundance elements ($<1 \text{ wt\%}$) show variability similar to that expected from counting statistics (Fig. 6, Supporting Figs 16–17). While the strong correlations between Al_2O_3 versus TiO_2 and Al_2O_3 versus SiO_2 in the experiments of Neave *et al.* (2019) are strongly indicative of sector zoning (compare Supporting Figs 6, 11), the weaker correlations for the Krawczynski *et al.* (2012) data make it hard to know the exact origin of true phase heterogeneity without detailed elemental mapping of crystals (Supporting Fig. 8).

We wish to determine whether elemental variability for the other experiments in the LEPR database are similarly controlled by analytical precision for Na_2O , and true phase variability for higher concentration elements. While we cannot reconstruct the exact analytical precisions for these studies, we can use the curves calculated for the new Bristol JXA8530F and old MIT JEOL733 probes as end-member examples of detector efficiency to estimate precision for different $I \cdot t$ values. This approach is supported by the fact that the software-calculated Cameca SX100 errors from Hannover lie within the modeled lines for these two probes (Fig. 6c). To get an idea of common $I \cdot t$ values, we compile count times and beam currents from 39 randomly selected experiments in LEPR used to calibrate the Putirka (2008) and Petrelli *et al.* (2020) Cpx-only and Cpx-Liq expressions, as well as a number of Cpx containing experiments conducted since 2008 (see Supporting Table 1). This compilation is also hampered by insufficient reporting of analytical conditions. Of the 39 LEPR experimental

studies, 31 stated the beam current while only 16 (~40%) gave the count time for Na_2O . The 22 experiments published since 2008 are somewhat better, with 21 providing beam current, and 14 (~60%) providing count times for Na_2O . Although incomplete, our compilation shows that $I \cdot t$ mostly varies between 60 and 400 nA s (Supporting Table 1). Interestingly, many papers include statements to the effect that well known or established analytical conditions were used for analysis without actually reporting what these were. The range of compiled $I \cdot t$ values shows that this is clearly insufficient, as there is no such thing as 'normal' analytical conditions (Supporting Table 1).

When the worst-case scenario (60 nA s on the MIT JEOL733, orange line) and best-case scenario (400 nA s on the Bristol JX8530F, blue dashed line) are overlain on the reported 1 sigma values for element variability within a single experimental charge from LEPR (Fig. 6e-f), it is apparent that the majority of Cpx Na_2O variability reported within single experiments results from analytical precision. The various measures of software-reported precision we compiled for different EPMA facilities (e.g. Cambridge, OSU) lie in the middle of the cloud defined by LEPR experiments, further supporting this assertion. The small number of points that lie above the MIT JEOL733 line could be influenced by experimental scatter, or even lower spectrometer efficiency than the MIT JEOL733 (or $I \cdot t < 60 \text{ nA s}$). In contrast to Na_2O , Al_2O_3 in LEPR experiments show significantly more variability than software-estimated precision, indicating that features such as sector zoning may be nearly ubiquitous in experimental Cpx (Fig. 6f).

Our modeling and compilations of analytical precision highlight a major issue with the way EPMA precision is being assessed and reported within the petrology community. If mentioned at all, precision is generally discussed with reference to repeated measurements of secondary standard materials (e.g. Neave *et al.*, 2019; Waters *et al.*, 2021). Although secondary standards are very helpful to assess accuracy and drift during an analytical session, they only provide useful insights into precision if secondary standards and samples have similar concentrations for a given element. This is a particular problem for Cpx, because the commonly used Kakanui augite secondary standard has 1.2–1.3 wt% Na_2O (Fig. 6b), whereas experimental Cpx produced at $<10 \text{ kbar}$ have median Na_2O contents of only 0.55 wt% (and most crustal Cpx have similarly low concentrations). The theoretical lines calculated for the MIT JEOL 733 show that analysis of Kakanui augite would indicate 5–10% error, but the lower Na_2O contents of crustal Cpx yield errors of up to ~40% (Fig. 6c). Thus, secondary standards must only be used to estimate precision for elements where the standard and sample have very similar concentrations.

Effect of major element variability on calculated pressures

For experiments where individual Cpx measurements within a single experimental charge could be obtained, we calculate Cpx-only pressures for each reported measurement (Krawczynski *et al.*, 2012; Blatter *et al.*, 2013; Melekhova *et al.*, 2015; Erdmann *et al.*, 2016; Neave *et al.*, 2019). Despite the fact that Cpx within a single experimental charge experienced a narrow range of pressures within their capsule, calculated pressures span up to ~4 kbar (Fig. 7a–e). The Cpx-only barometer of Putirka (2008) shows the widest range in most cases (red diamonds), with the barometers of Wang *et al.* (2021, black dots) and Petrelli *et al.* (2020, cyan squares) showing a slightly smaller range for a given experiment. The range of calculated pressures in a single experiment is in many cases comparable to the reported SEE on the barometer (colored error bar). Visually, it is easy to see how this variation can lead to large reported errors on barometers if only a small number of Cpx are

analyzed. For example, if Neave *et al.* (2019) had measured just one Cpx, the pressure calculated using the Cpx-only barometer and thermometer of Putirka (2008, eq32a-32d) could range from ~4 to 10 kbar. In turn, when plotted against experimental pressure (3 kbar), this pressure calculation could lie 1 to 7 kbar off the 1:1 line. When all experiments in a test dataset are considered, these offsets cause very large SEE estimates on the barometer.

However, it is still unclear from these discussions whether major elements (predominantly controlled by true phase variability) or minor elements (controlled by analytical precision) are producing this spread of calculated pressures in a single experimental charge. Qualitatively, when we plot the calculated pressures from five experiments against Na₂O for two barometers, the strong correlation between calculated pressures and Na indicates that analytical precision is a major contributor to the spread in calculated pressures (Fig. 7f).

To investigate the relative effect of analytical precision versus true phase variability more quantitatively, we use Monte Carlo simulations implemented in the Python3 thermobarometry tool Thermobar (Wieser *et al.*, 2022b). For specific experimental charges from Krawczynski *et al.* (2012) and Neave *et al.* (2019), we create 5000 synthetic Cpx compositions with the concentration of each element distributed normally about the mean composition of the Cpx from each experiment, and the standard deviation equal to the average reported 1 sigma error from EPMA software for analyses from that experimental charge. Each major element in each of the 5000 Cpx is sampled randomly from a normal distribution. The uncoupled variations of these different oxides means that cation sums in these simulated Cpx do not lie substantially outside the distribution of cation sums in the measured Cpx (Supporting Fig. 18–19). We then calculate pressures and temperatures for these synthetic Cpx using a variety of Cpx-only barometers (Fig. 8–9). To aid visualization of the spread of simulated pressures and temperatures, we contour the results of the Monte Carlo simulations and overlay contours incorporating 67% and 95% of simulated pressures and temperatures using the Python3 tool Pyrolite (Williams *et al.*, 2020). If analytical variability is the dominant cause of pressure–temperature variability, the Monte Carlo simulations will encompass the variability of calculated P and T for each measured Cpx composition. In contrast, if true phase variability dominates, calculated P and T for measured Cpx compositions (yellow and red dots) will plot well outside the PT region defined by the simulations (Figs 8 and 9).

For experiment Y0272–1 from Neave *et al.* (2019), the 95% contour around the Monte Carlo simulation almost completely incorporates the spread of calculated pressures and temperatures in individual Cpx measured in that experiment for all three Cpx-only thermobarometers (yellow dots; Fig. 8a–c, see also Supporting Fig. 20). Thus, while experimental products show major element variability resulting from sector zoning (Neave *et al.*, 2019), the dominant control on calculated pressures and temperatures results from analytical precision.

For Cpx-Liq calculations, no EPMA-estimates of glass analytical uncertainties could be obtained for Neave *et al.* (2019), so calculations were performed using the average glass composition paired with simulated Cpx compositions (so this calculation underestimated the true variability caused by analytical uncertainty). Despite neglecting analytical variability in the glass phase, the span of P–T in measured Cpx-Liq pairs is very similar to our Monte Carlo simulations for Putirka's (2008) eq 33 for temperature, iterated with either eq 30 (Fig. 8e) or Neave & Putirka (2017) for pressure (Fig. 8e and Fig. 8d, respectively). It is noticeable on these

figures that the machine learning-based Cpx-only and Cpx-Liq thermobarometers from Petrelli *et al.* (2020) show significantly less spread in calculated pressures and temperatures both for measured experimental products, and our Monte Carlo simulations (Fig. 8c,f). However, using a different experiment, this barometer shows a much larger spread (Supporting Fig. 20), illustrating the unintuitive behavior of regression tree machine-learning algorithms versus empirical expressions.

In the experiments of Krawczynski *et al.* (2012), Cpx in a single experiment have compositions such that sometimes jadeite is calculated using Na and sometimes using Al. This complicates the relationship between the spread of pressures and the analytical precision because Al is mostly affected by true phase variability (e.g. sector zoning) while Na is mostly affected by analytical precision. Our simulations do not show the same P–T extent as measured values. However, if one imagines shifting the simulation away from the mean, the spread shown by Cpx where jadeite is calculated from Na (yellow dots) is reasonably similar to the simulation, while Cpx with jadeite calculated from Al (red dots) lie well outside the Monte Carlo simulation (Fig. 9).

Overall, these comparisons indicate that apparent phase variability resulting from analytical precision alone can yield a wide range of pressures. Although analytical precision seems to dominate the spread of calculated pressures in experiments where jadeite is calculated using Na, measuring sufficient Cpx to average true phase variability is vital where jadeite is calculated from Al. When either source of elemental variability is insufficiently averaged by measuring a large number of Cpx, this can lead to large calculated errors for barometers. For x–y plots of calculated versus experimental pressure, the effect of measuring only a single Cpx on the y axis value is easy to visualize from Fig. 8–9.

We investigate whether insufficient averaging of analytical and/or true phase variation can explain the notable decline in the discrepancy between predicted and experimental pressure with increasing number of Cpx analyses per experiment (Fig. 4c–d). We use software-reported estimates of analytical precision from an experimental charge of Neave *et al.* (2019) to produce 5000 synthetic Cpx compositions. This simulates what would happen if 5000 discrete analyses were made of entirely homogenous experimental products, with variability in measured oxides resulting from counting statistics. We then calculate the composition obtained from sampling discrete numbers of Cpx (N = 1 to N = 100), and feed these averaged compositions into various Cpx-only barometers. We calculate the discrepancy between the calculated pressure for this averaged composition, and the pressure calculated from the mean composition of all 5000 Cpx. For each discrete number of averaged Cpx analyses, we overlay the mean discrepancy (blue star), and the 95% quantile (yellow star) on the data (Fig. 10a–b). We repeat this process 700 times using a for loop in Python, meaning we have simulated randomly sampling one Cpx 700 different times, two Cpx 700 times, etc. A schematic showing the loop and subsampling routine is provided in Supporting Fig. 21.

Figure 10a,b demonstrates that when only one EPMA analysis is taken from a homogenous Cpx population, the calculated pressure can differ from the pressure calculated from the true composition by up to ±3–4 kbar for Putirka (2008; Fig. 10a) and ±2.5–3.5 kbar for Wang *et al.* (2021; Fig. 10b). The mean absolute discrepancy for N = 1 Cpx is ±0.89 kbar for Putirka (2008) and ±0.55 kbar for Wang *et al.* (2021, blue stars), while the 95% quantile is ±2.09 kbar for Putirka (2008) and ±1.39 kbar for Wang *et al.* (2021, yellow stars; Fig. 10a,b). The absolute discrepancy from each experiment, the mean, and 95% quantiles decline very

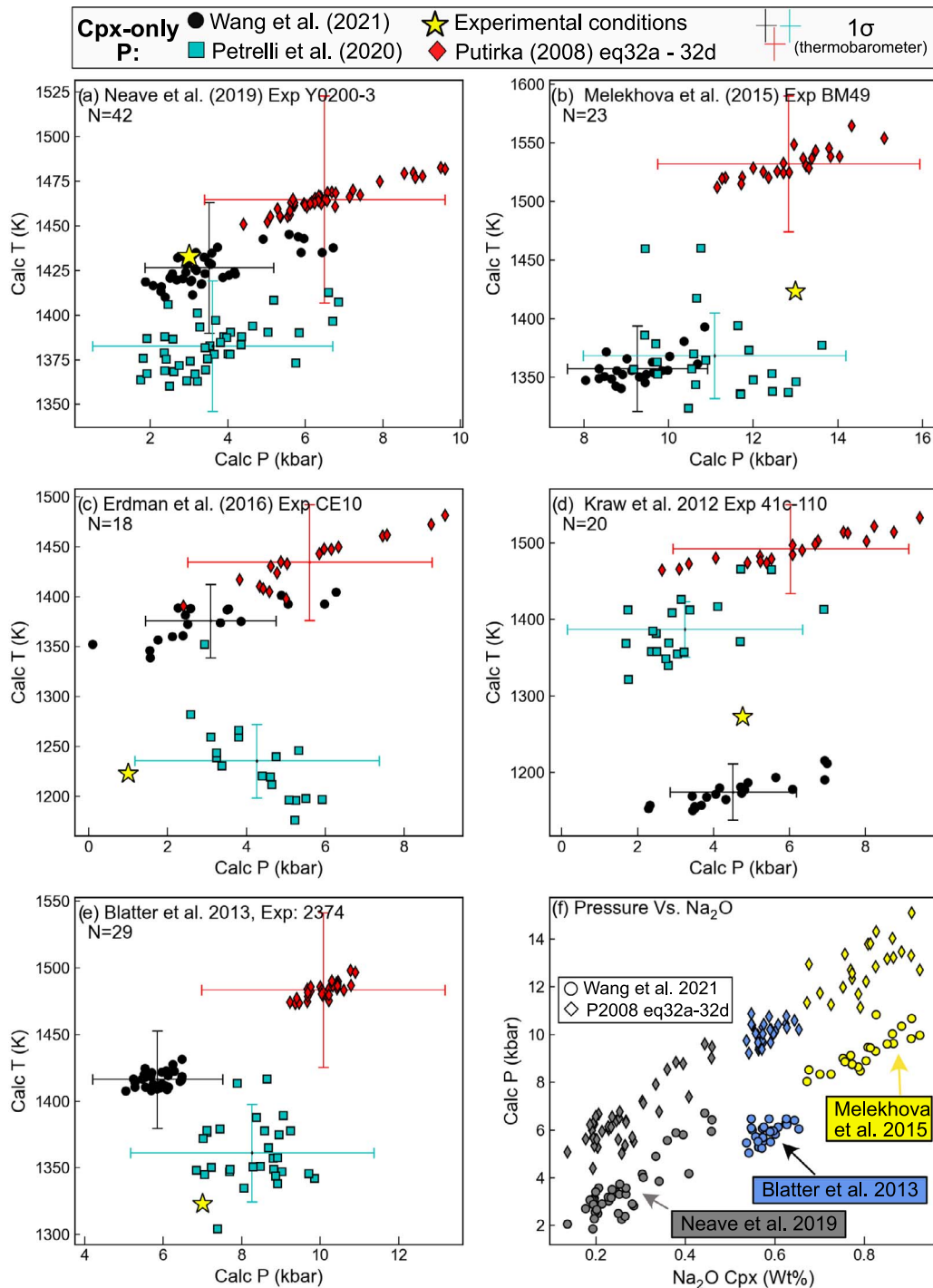


Fig. 7. (a–e) Pressures and temperatures calculated from individual Cpx compositions from specific experiments using Cpx-only thermobarometers from Petrelli et al. (2020), Putirka (2008), and Wang et al. (2021). Error bars showing published/commonly quoted SEE for each thermobarometer (see caption for Figs 2 and 3). (a) The pressures and temperatures calculated from the 42 Cpx from experiment Y0200–3 of Neave et al. (2019). The yellow star shows the P–T conditions at which the experiment was conducted. (b) Experiment BM49 from Melekhova et al. (2015). (c) Exp CE10 from Erdmann et al. (2016). (d) Exp 41c–110 from Krawczynski et al. (2012). (e) Exp 2374 from Blatter et al., 2013. In (f), for three different experiments, we show the strong correlation between Na₂O in the Cpx and calculated pressure using the Putirka (2008) barometer (diamonds) and the Wang et al. (2021) barometer (circles). Experiments showing a larger spread of Na₂O values (Melekhova et al., 2015; Neave et al., 2019) show a much wider spread of pressures than the Blatter et al. (2013) experiment.

rapidly between $N = 1$ and $N = 7$ Cpx. When >7 Cpx are averaged, there are very few experiments >1.5 kbar from the mean value calculated from all 5000 Cpx, and very few individual experiments >1 kbar ($\sim 5\%$). However, it must be remembered that we are only simulating analytical uncertainty; analyses will also be affected by true heterogeneity in experimental products.

To investigate the combined effect of analytical and experimental variability (i.e. the measured variability in a given experiment), we repeatedly resample the 18 reported Cpx from experiment B1038 of Krawczynski et al. (2012), averaging different numbers of measured Cpx compositions to create a theoretical ‘average’ composition (see supporting Fig. 22 for a schematic of

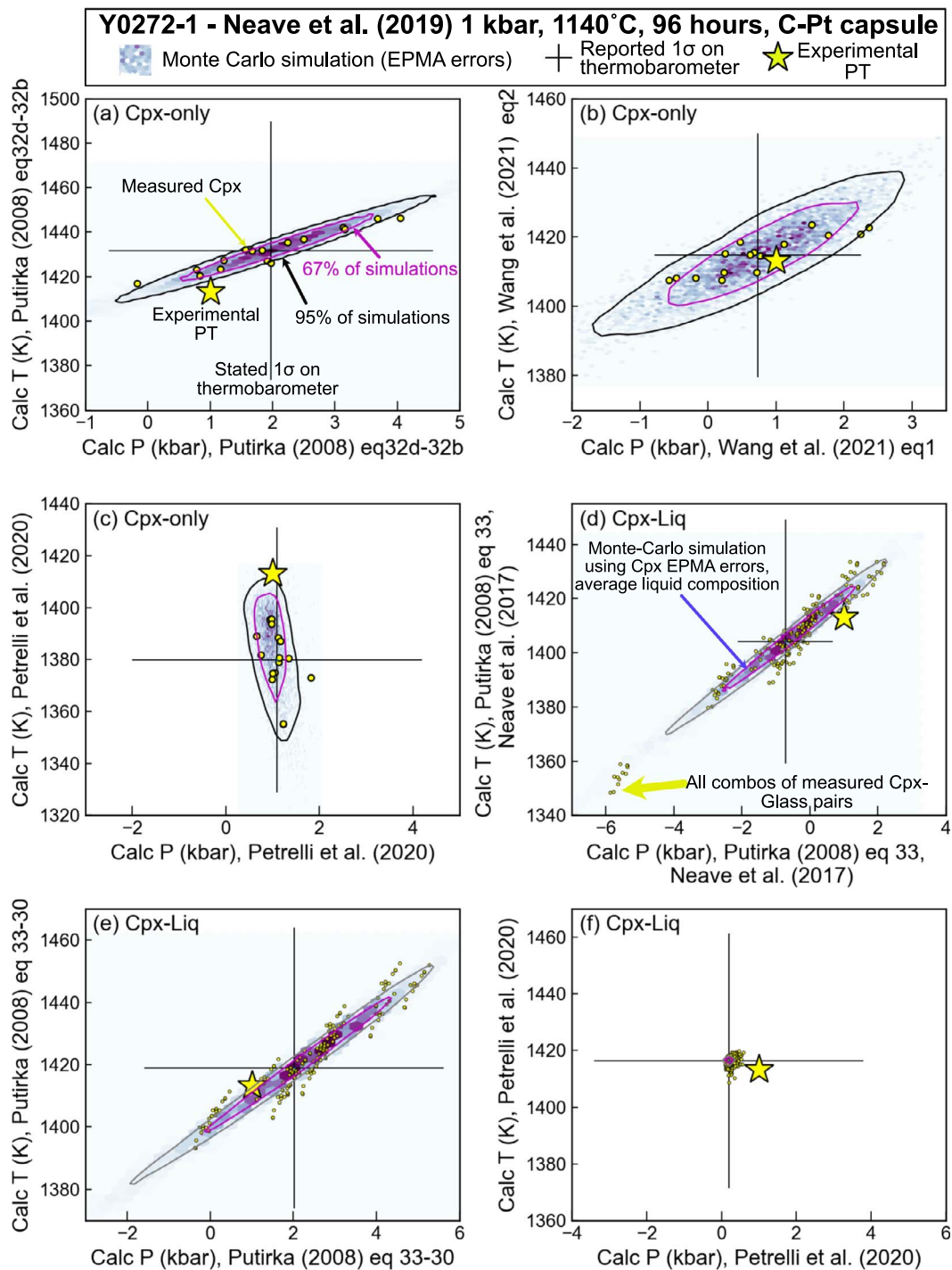


Fig. 8. Monte Carlo simulations showing the spread of Cpx-only and Cpx-Liq pressures resulting from electron microprobe-calculated analytical precision of experimental Cpx from Exp Y0272-1 of Neave et al. (2019). Measured Cpx compositions are overlain in (a–c), and all possible combinations of measured Cpx and Liquids in (d) and (e). The error bar on each plot shows the reported error on each thermobarometer, and the yellow star shows the analytical conditions. In (d–f), Monte Carlo simulations show errors for Cpx and the average glass composition. Monte Carlo simulations and thermobarometry calculations performed in Thermobar, contours showing 67% and 95% of simulations calculated using Pyrolite. Supporting Fig. 20 shows the same plots for experiment B0184–10.

this loop). For $N = 1$ Cpx, the mean offsets reach ~ 2 kbar, with 95% quantiles of ~ 4 –5 kbar (Fig. 10c,d).

We also consider the effect of sampling on Cpx-Liq barometry, using individual measurements of experimental Cpx and glass from experiment B1038 from Krawczynski et al. (2012). First, we match up all possible pairs of measured experimental Cpx and Liq ($N = 360$ pairs). For each pass through the loop, we randomly

select 10 Liq that have been paired with a given Cpx, and calculate an average Liq composition. We then consider averaging different numbers of Cpx paired with these average Liq compositions (see Supporting Fig. 23 for schematic of this loop). This approach of using average liquid compositions means we are only investigating the effect of how many Cpx analyses are averaged, because there is no clear correlation in between number of Cpx and

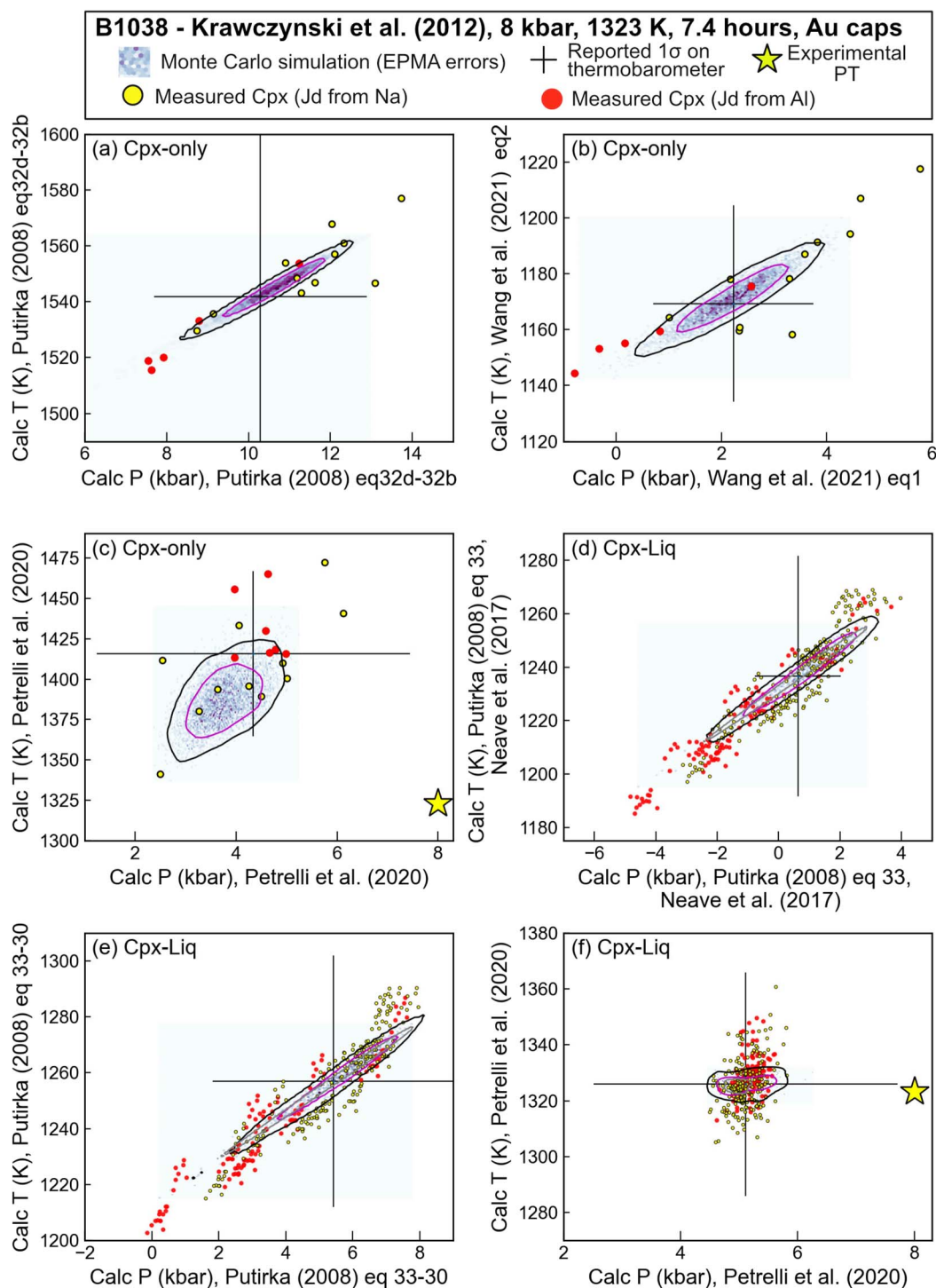


Fig. 9. As in Fig. 8 but for Exp B1038 from Krawczynski et al. (2012). In (d–f), Monte Carlo simulations show errors for both Cpx and Glass. The 95% contour using only analytical noise for Cpx (and not glass) is shown in grey for comparison with Fig. 8 (where the analytical error on glass analyses from Neave et al., 2019 was not available).

Liq analyses per experiment in our compiled dataset. Depending on the choice of barometer, mean offsets decline rapidly with increasing N from ± 2 kbar for $N = 1$ Cpx, with a 95% quantile of ± 4 –5 kbar (Fig. 10e–f).

Importantly, the large offsets for $N = 2$ –4 Cpx highlight a particular problem that can arise when measuring only a small number of Cpx. Specifically, when a Cpx has Na–Al contents such that the jadeite content is zero, averaging the composition of that Cpx with a second Cpx with Jd > 0 can result in a very low (but non-zero)

jadeite content. While Jd = 0 will return a NaN for pressure (not a number), because most Cpx–Liq thermobarometers contain a log term involving the jadeite content (and $\log(0) = \text{NaN}$), the log of a very low but non-zero jadeite content produces a very large negative number, yielding an anomalous pressure (see Supporting Information Fig. 24).

Our simulations (Fig. 10) demonstrate that insufficient EPMA analyses to characterize the true composition of experimental phases can account for the observation that experiments char-

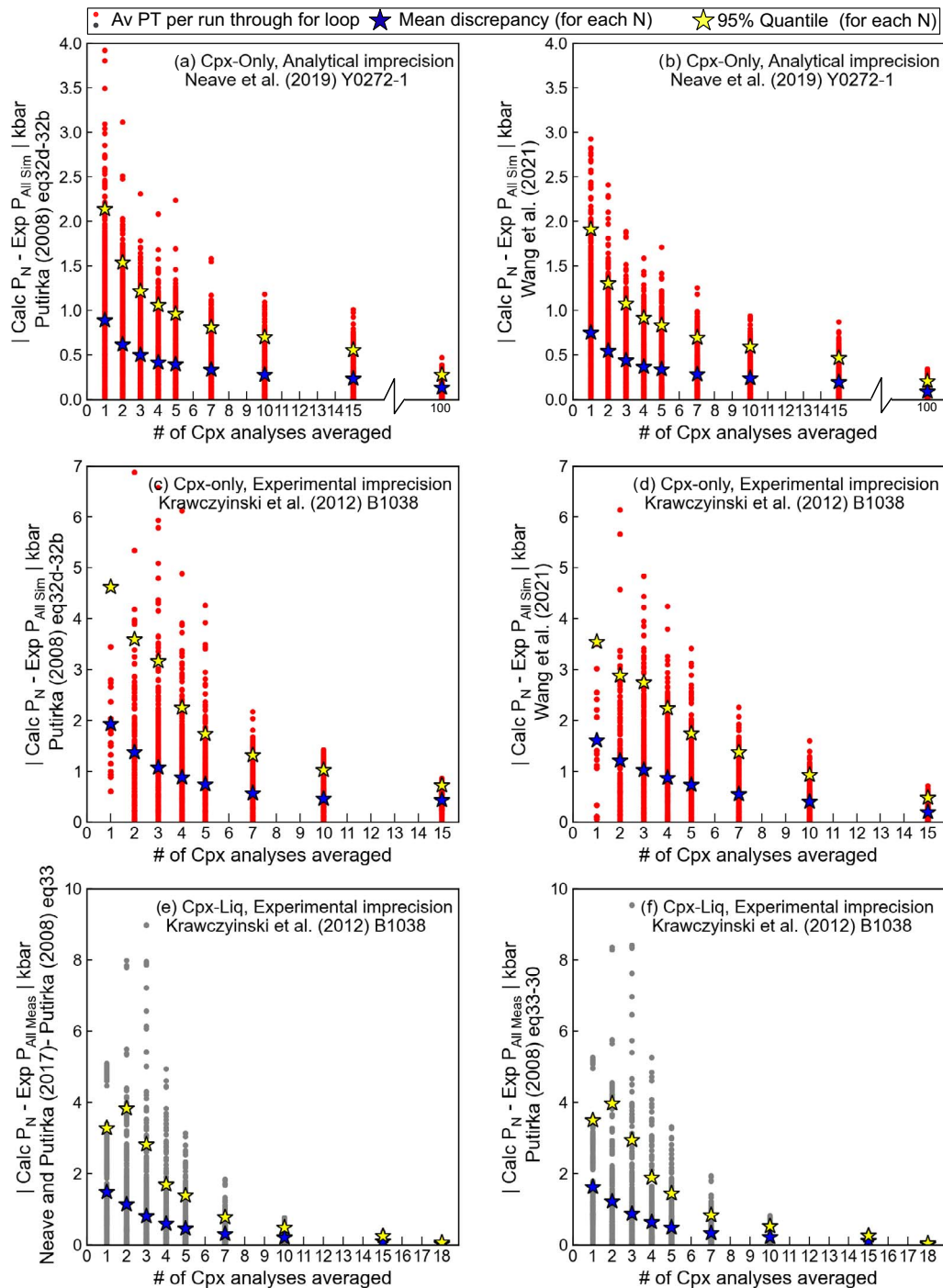


Fig. 10. Demonstration of the absolute pressure offset that can be introduced by measuring insufficient Cpx from any given experimental study. (a,b) Groups of 1–100 Cpx were randomly sampled from an underlying distribution of 5000 Cpx simulated using reported EPMA errors for experiment Y0272-1 of Neave et al. (2019). The difference in pressure calculated for each group is compared to the average pressure calculated for all 5000 Cpx, and plotted as a single dot (see Supporting Fig. 21). (c,d) To simulate the combined influence of true phase variability and analytical errors, we repeatedly resample and average the 18 measured Cpx from Krawczyński et al. (2012, see Supporting Fig. 22). (e,f) Random sampling of measured Cpx-Liq pairs from Krawczyński et al. (2012, see Supporting Fig. 23). On all plots, the y-axis shows the absolute discrepancy between the pressure calculated during each run through the loop and the average pressure. In (a–d), this is the average PT for the entire simulated cloud of Cpx, in (e,f) it is the average for the reported mean Cpx and glass composition from the experiment. For each discrete number of Cpx, we calculate the mean (blue star) and 95% quantile (yellow star).

acterized by a small number of analyses can show large discrepancies between calculated and experimental pressures (Fig 4c,d). Most simply, imagine that barometers perfectly predict the relationship between composition and pressure. Even in this perfect scenario, measuring a small number of Cpx (\pm Liq) in each exper-

iment can result in the measured composition not being representative of the true composition, resulting in offsets between calculated and experimental pressures. Of course, in some cases, a single measurement will obtain the correct value (i.e. sampling the mean of the normal distribution by fluke), accounting for

the fact that some experiments analyzing a small number of Cpx show small offsets on Figs 4 and 10. However, the smaller the number of Cpx analyses performed, the more likely it is that the composition of the Cpx obtained differs from the true composition. In reality, barometers are not perfectly accurate, which explains why there are still offsets between calculated and experimental pressure even when larger numbers of Cpx are analyzed.

The y-axis in these subsampling simulations represents the distance a single experiment may lie from the 1:1 line in a plot of experimental pressure versus calculated pressure, simply because of insufficient averaging of analytical and experimental variability. Given that ~43% of experiments we have compiled performed ≤ 5 Cpx analyses per experiment, even if barometers perfectly recreate the relationship between pressure and temperature for the true composition of Cpx and Cpx-Liq pairs from that experiment, we would expect scatter about a 1:1 line of ± 2 –3 kbar based on insufficient averaging. In reality, the scatter off the 1:1 line will be larger than this, because insufficient averaging of experimental products affects the calibration of barometers, as well as the assessment of these barometers using test datasets. Thus, we suggest that insufficient averaging of analytical precision (with a contribution from true phase variability) is the ultimate reason barometers have SEE of ~2–4 kbar when applied to global test datasets, regardless of the exact calibration strategy.

Implications for thermobarometry on natural systems

The Monte Carlo simulations of calculated pressures and temperatures resulting from analytical uncertainty shown in Figs 7 and 8 for Cpx and Cpx-Liq are equally applicable to natural systems. Namely, even if erupted Cpx are entirely homogenous and come from a single magma storage region at a single pressure and temperature, calculated pressures may span ~4 kbar using Cpx-only thermobarometry, and ~6 kbar using Cpx-Liq thermobarometry because of low analytical precision (assuming analytical conditions similar to those used by Neave *et al.*, 2019 and Krawczynski *et al.*, 2012, i.e. 100–150 nA s). When plotted in P-T space, completely random, normally distributed analytical error produces a strongly correlated pressure–temperature array (Figs 8 and 9). It is crucial to recognize that a wide spread of pressures calculated for a natural system using popular Cpx-only and Cpx-Liq barometers should not automatically be interpreted as representing magma storage across a broad region of the crust (i.e. transcrustal magma storage). Rather it should be assumed that this is an artifact of analytical uncertainty until proven otherwise. Specifically, Monte Carlo simulations with errors determined from software-calculated analytical uncertainty (rather than secondary standards) should be used to calculate the spread in pressures and temperatures that result from analytical precision. If the spread of pressures and temperatures from natural samples exceeds the simulated spread (and even if it does not), the role of P-T spread resulting from sector zoning and other disequilibrium features should also be investigated (e.g. Hammer *et al.*, 2016). True transcrustal storage should only be invoked after ruling out these null hypotheses.

Although the overall structure of pressure and temperature estimates should certainly be considered, averages of many individual measurements help to eliminate analytical and true phase variability. For example, Putirka *et al.* (1996) showed that the fit between calculated and experimental pressures are substantially improved when all experiments conducted at a given

pressure are averaged, rather than considered individually (compare their Fig. 3a and Fig. 4a). To avoid averaging out true variations in magma storage conditions, it may be best to perform and average multiple analyses within the core and rim of any given Cpx (e.g. Klügel *et al.*, 2005).

SUMMARY AND FUTURE DIRECTIONS

There is broad consensus that improved methods for estimating the pressures of igneous processes and magma storage is vital to advance the field of igneous petrology (Hilley *et al.*, 2022; McGuire *et al.*, 2017). In the preceding sections we have highlighted a number of sources of uncertainty affecting Cpx-only and Cpx-Liq barometry, which are two of the most popular barometry tools. We show that insufficient averaging of measurements made with low analytical precision, combined with heterogeneity in experimental products, and interlaboratory offsets fundamentally limit the precision of Cpx-based barometric estimates to ± 2 –4 kbar for crustal Cpx. Below, we highlight a number of ways in which we can improve how experimental products are analyzed and reported to improve the future calibration and application of Cpx-based barometers. These recommendations can be summarized as: 1) collect more counts, 2) measure more phases in each experimental charge, 3) address interlaboratory biases and 4) better data reporting.

Can we simply collect more counts for Na in Cpx?

Our Monte Carlo simulations demonstrate that analytical precision associated with the measurement of Na₂O resulting from insufficient count times and/or beam currents is a major source of uncertainty affecting Cpx-only and Cpx-Liq barometry. Poor precision affects the experimental data used for barometer calibration and testing, as well as calculations of pressure in natural sample suites. Many papers suggest that they are using such short analysis times and low beam currents because of fears of Na migration (e.g. Neave *et al.*, 2019). Although beam migration is a justifiable concern in hydrous glasses and Na-rich feldspars, to our knowledge there are no reports of Na migration within crustal pyroxenes.

To investigate the beam sensitivity of pyroxene, we perform tests of Na mobility during analysis at 15 kV, and 20, 40 and 100 nA using a 1 μ m spot on the Oregon State Cameca SX100. We track changes in peak intensity, and calculated Na concentrations using typical analytical routines for a Na-rich end-member jadeite (~7.1 wt% Na₂O) and a Kakanui Augite with lower Na contents (~1.2 wt% Na₂O), as well as a hydrous rhyolite glass for comparison.

Peak intensities in the hydrous rhyolite glass show a rapid decrease with increasing exposure, even at 20 nA, as is widely documented (e.g. Morgan & London, 2005; Fig. 11a, black triangles). The jadeite is significantly less beam-sensitive, with peak counts only decreasing by 15%–20% after ~5 minutes (and only at 40 and 100 nA; Fig. 11a,b). In contrast, we see no evidence for a decrease in Na peak intensities collected on the Kakanui augite even at 100 nA over ~6 minutes (Fig. 11b).

We also perform six repeated analyses at 100 nA at the same stage position on the jadeite and Kakanui augite, with a total beam-on time of ~60 s per analysis. Peak count times were Na (3.3 s), Ca (10 s), Al (10 s), Si (10 s) and Ti (10 s). We plot the concentration of each element against the average beam-on time for each analysis. For the jadeite, the concentration of Na₂O declines reasonably coherently by ~10% after ~6 minutes of beam exposure (Fig. 11c). For the Kakanui Augite there is no coherent

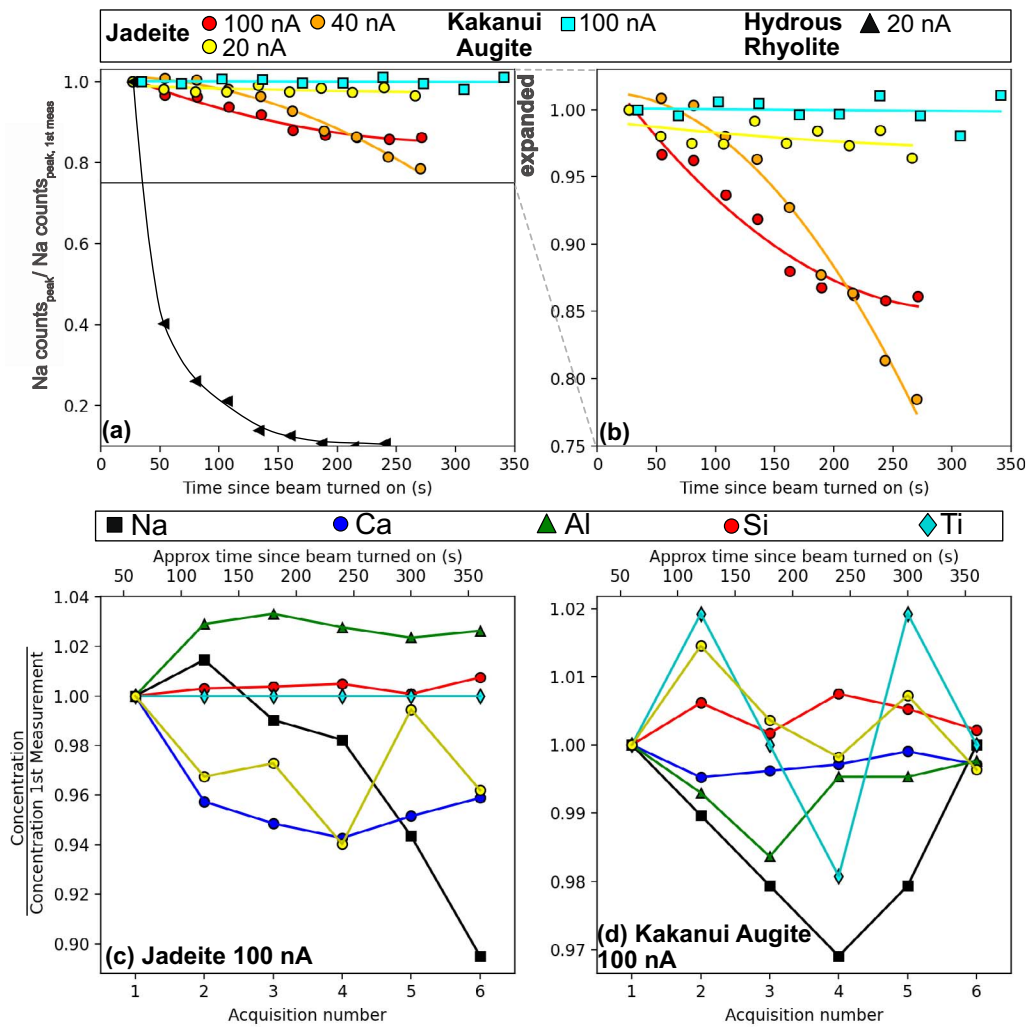


Fig. 11. Investigation of Na migration. (a,b) Tracking the number of counts on the Na peak with increasing beam exposure on a hydrus rhyolite, a jadeite and a Kakanui augite. Polynomial fits are included through the data to help with visualization. Note the compressed y-axis scale on b. (c,d) Change in elemental concentration for six repeated quantitative analyses on the same location in jadeite versus Kakanui augite. The approximate beam exposure time is shown as a secondary axis (although different elements were acquired at slightly different points during each acquisition).

decrease in Na_2O , with concentrations varying well within the expected threshold given the low count time (Fig. 11d). Further details of additional tests are provided in the supporting information (Supporting Figs S25–30). Overall, we conclude that in pyroxenes with relatively low Na_2O contents, there is no need to restrict count times or beam currents, as Na migration seems negligible.

Given the lack of measurable Na migration in natural Cpx, we suggest that the simplest way to increase precision on reported Na in future experiments and natural samples is a combination of longer count times and/or a higher beam current during Cpx analyses. Increasing the count time at a specific beam current is the best strategy if the interaction volume needs to be minimized (e.g. for analyses of small experimental products), while increasing the beam time and current together is most efficient if a tiny interaction volume is not critical.

In many analytical routines, increasing the count time on Na will not increase the total analytical time if other spectrometers are already counting minor elements for longer (e.g. Cr and Mn on the LIF). Additionally, in a number of studies, Na and Al are both being measured on the TAP or LTAP (e.g. Krawczynski *et al.*, 2012; Hammer *et al.*, 2016; McCane, 2022; Wieser *et al.*, 2022a), with

Al being counted significantly longer than Na, despite its higher concentration in Cpx. If total analytical time is an issue, count times can be adjusted on the TAP/LTAP to count for longer on Na instead.

Although secondary standards have great utility for assessing accuracy and drift, they should only be used to assess precision if the standard has a very similar elemental concentration to the analyte. If elements are present in lower concentrations in natural samples, software-calculated precisions should be used to assess uncertainties resulting from counting statistics. Overall, we suggest that users optimize their EPMA acquisition parameters to achieve <5% precision for the range of Na_2O contents found in their samples.

Perform more measurements in each experimental charge.

Analytical noise is particularly problematic for calibration datasets if only a small number of Cpx in each experiment are measured. The large number of experiments performing <5 individual Cpx measurements, combined with low precision on Na_2O measurements, explains why no Cpx-based barometers can predict pressure in a global dataset of experiments with errors

smaller than $\pm 2\text{--}5$ kbar (despite many different calibration strategies, Wieser *et al.*, 2023). Even if analytical precision is improved by increasing the number of counts, performing multiple Cpx analyses remains vital to minimize the effects of experimental heterogeneity (e.g. sector zoning, Neave *et al.*, 2019). This is particularly true if jadeite is calculated from Al, or when using Cpx-Liq barometers relying on Al content (e.g. eq32c, Putirka, 2008) as Al and other major elements in Cpx and glass are influenced more by true experimental phase variability than analytical uncertainty.

Quantify and resolve interlaboratory offsets for glasses and silicate minerals

Even if analytical conditions are optimized to make measurements more precise, and a larger number of products are measured in each experiment, interlaboratory offsets may still introduce random, or even systematic offsets into barometry calibrations. To properly resolve the extent of interlaboratory biases, we suggest that mineral round-robins are conducted on the current generation of EPMA at institutions which are contributing a significant amount of experimental data (and ideally between all laboratories). Such round-robins will be vital to determine whether our assumption that mineral offsets may be as large as offsets measured for glasses is correct. It may be that by applying such corrections, more precise barometry calibrations can be obtained.

Longer-term, it is not practical to rely on large round-robins among all possible EPMA labs that may wish to perform mineral-melt barometry on natural samples, or conduct experiments used to calibrate these expressions. These interlaboratory biases highlight the need for the development of a new generation of homogenous reference materials to replace the heterogeneous NMNH standards. One recent suggestion, put forward in an open letter to the microanalysis community with over 90 co-signers (<https://probesoftware.com/smf/index.php?topic=1415.0>), is to create a global reference set of high purity, stoichiometric end-member synthetic compositional standards in approximately 500 to 1000 gram quantities. This would ensure that every e-beam microanalysis laboratory in the world could readily obtain sufficient material to last decades of polishing, use, repolishing and reuse. Even better, such synthetic standard materials can always be reproduced in the future as necessary, since they would be selected such that their synthesis would be well-constrained by both purity and thermodynamics. These efforts are being formalized within the focused interest group microanalytical standards committee of the Microbeam Analysis Society (<https://the-mas.org/about-us/focused-interest-group-figmas/>).

Better reporting of compositional, analytical and metadata

It may be possible to improve barometry calibrations by excluding experiments with numbers of analyses below a certain threshold, or experiments where phases were only briefly characterized to identify phase occurrence, rather than to provide reliable phase compositions. At a minimum, filtering based on numbers of analyses will require this information to be copied over from pdf tables in papers and entered into the LEPR dataset. However, to evaluate analytical noise within a single experiment rigorously, we also need an estimate of the analytical precision. For example, it may be more rigorous to include an experiment where only two Cpx were measured but at high precision (e.g. Na = 10%) versus an experiment with five Cpx with low analytical precision (e.g. Na = 50%; Fig. 6b). However, this information cannot be obtained

for the majority of the LEPR database, particularly as many of the EPMA instruments used for these measurements have been decommissioned.

Thus, for future experimental work, we suggest that authors report the following information:

- 1) The beam current, voltage, crystal, primary calibration material, and peak and background count times for every element;
- 2) The software-calculated 1 sigma value for each analysis;
- 3) The elemental data for every spot analysis of every phase in each experimental charge and every natural mineral analysis, rather than providing a mean composition and standard deviation;
- 4) Detailed information on how thermal gradients were assessed, and any friction corrections.

Providing this information will allow future attempts at calibrating barometers to better filter the underlying data. We also encourage authors to think carefully about the influence of minor elements such as Cr on Cpx phase stability (Voigt *et al.*, 2017) and to carefully report whether an attempt was made to measure an element but it was below the detection limit, or whether no measurement attempt was made. If an element was below detection limit, and an estimate of the detection limit is provided, the number can be more reliably imputed than when no information is given.

Remeasure existing experimental products

We recognize that experimental studies are seldom performed with the sole aim of calibrating thermobarometers. Instead, the authors may have simply wanted to constrain a phase diagram, so only a small number of Cpx were analyzed in each experiment to confirm phase occurrence, and there was no reason for analytical conditions to be optimized for low concentration components like Na₂O, or to measure all elements (e.g. Cr₂O₃ in Cpx and glass). Although it will require a significant community effort to find and share the experimental charges, this is likely much less effort than redoing experiments from scratch, and it would be worthwhile to reanalyze a large proportion of the experimental charges compiled in LEPR. This would take advantage of the higher precision of modern EPMA instruments (compare the precision of the old and new MIT JEOL on Fig. 6c) and help to fill in the large amount of missing data in LEPR (Cr₂O₃, H₂O, P₂O₅, etc). Ideally, reanalysis would take place on a single EPMA instrument or on a set of instruments where secondary standards are exchanged to correct for interlaboratory biases.

CONCLUSIONS

In their current state, Cpx-based barometers struggle to precisely and accurately constrain the location of magma storage in the crust (with errors of $\pm 2\text{--}5$ kbar, $\pm 8\text{--}19$ km). In addition to fundamental thermodynamic limitations, we suggest that the poor performance of barometers results from the fact that the true composition of Cpx in experiments has not been precisely determined, because insufficient analyses were conducted to average out low analytical precision and true phase variability. Calibrating and testing Cpx-based barometers has been further hindered by interlaboratory offsets during EPMA analyses, and large amounts of missing elemental data reported by experimental studies. We suggest that pressures calculated from individual Cpx analyses in both experiments and natural samples must be evaluated in

the context of the expected range of pressures obtained from the propagation of analytical errors for that specific study. To invoke true transcrustal storage, it must be demonstrated that the range of calculated pressures greatly exceeds that expected from analytical precision alone.

We believe a new generation of more precise barometers could be calibrated through a community effort to obtain an experimental dataset, which properly averages analytical imprecision (for low concentration elements such as Na₂O) and true phase variability (e.g. Al₂O₃ concentrations affected by sector zoning). There is growing recognition in the Machine Learning community that big data are not as important as good data. In a recent interview (2022), Machine Learning expert Andrew Ng states 'In many industries where giant datasets simply do not exist, I think the focus has to shift from big data to good data.' We suggest that the same reasoning should be applied to petrological thermobarometers. Ideally, an independent high quality test dataset would be isolated during model calibration and tuning to allow robust estimates of the precision that can be expected when these methods are applied to 'unknown' samples. Improving Cpx-based barometers is vital for reliable interpretation of a volcanic plumbing system geometry (e.g. distinguishing between a single reservoir, discrete reservoirs, and true transcrustal magmatic systems).

DATA AND CODE AVAILABILITY

Jupyter Notebooks used to produce Figs 2–11 in this paper have been uploaded to Penny Wieser's GitHub (https://github.com/PennyWieser/BarometersBehavingBadly_Wieser2022), archived on Zenodo: <https://zenodo.org/badge/latest/doi/571745416>. These Jupyter Notebook read from the following supporting Excel files, also available on GitHub and zenodo:

- Supporting_Data_1.xlsx – Contains Interlaboratory offsets from Gale *et al.* (2013), the Cpx calibration dataset used by Keith Putirka which we use to examine the prevalence of missing data, and the compilation of Cpx-Liq experiments used in this study (i.e. ArcPL).
- Supporting_Data_2.xlsx – Analysis of individual phases from the experiments of Krawczynski *et al.* (2012).
- Supporting_Data_3.xlsx – Estimates of analytical precision from a subset of experimental analyses by Krawczynski *et al.* (2012),
- Supporting_Data_4.xlsx – All Cpx-bearing experiments from the LEPR (downloaded in 2021).
- Supporting_Data_5.xlsx – Analysis of individual phases from the experiments of Neave *et al.* (2019).
- Supporting_Data_6.xlsx – Analysis of individual phases from the experiments of Erdmann *et al.* (2016), from their supporting information
- Supporting_Data_7.xlsx – Analysis of individual phases from the experiments of Melekhova *et al.* (2015).
- Supporting_Data_8.xlsx – Analysis of individual phases from the experiments of Waters *et al.* (2021).
- Supporting_Data_9.xlsx – Analysis of individual phases from the experiments of Blatter *et al.* (2013)
- Supporting_Data_9.xlsx – Analysis of individual phases from the experiments of Blatter *et al.* (2013)
- Supporting_Data_10.xlsx – Investigation of Na migration on the OSU SX100

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