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Petrological and Geochemical Variation during the Soufrière Hills Eruption (1995-2010)

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ABSTRACT
The andesite lava erupted at the Soufrière Hills Volcano is crystal rich with 33-63% phenocrysts, of plagioclase (65%); amphibole (28%), orthopyroxene (7%) and minor Fe-Ti oxide and clinopyroxene microphenocrysts. The andesite hosts mafic enclaves which have similar mineral phases to the andesite. The enclaves are generally crystal poor but can have up to 27% of inherited phenocrysts from the andesite, the majority of which are plagioclase. The eruption is defined by discrete periods of extrusion called phases, separated by pauses. The enclaves exhibit bulk geochemical trends that are consistent with fractionation. We infer that the intruded mafic liquids of Phases I and II interacted and assimilated plutonic residue remaining from the multiple prior mafic intrusions, while the basaltic liquids from Phases III and V assimilated relatively little material. We also infer a change in the basaltic composition coming from depth. The bulk Fe contents of both magma types are coupled and they both show a systematic inter-phase variation in Fe content. We interpret the coupled Fe variation to be due to contamination of the andesite from the intruding basalt via diffusion and advection processes, resulting in the erupted andesite products bearing the geochemical imprint of the syn-eruptive enclaves.

The Soufrière Hills volcano is an andesitic dome complex located on the island of Montserrat in the Lesser Antilles arc. The present eruption began on the evening of July 18th 1995 with ash venting followed by phreatic explosions over the next weeks and months (Young et al., 1998; Robertson et al., 2000). Juvenile material arrived at the surface around November 15th 1995 (Young et al., 1998) building the first lava dome of the eruption. Lava extrusion and dome growth at Soufrière Hills volcano has not been continuous during the eruption; dome growth has been defined by distinct periods of extrusion referred to as ‘phases’ separated by periods of no activity, referred to as ‘pauses’. Each period of extrusion was volcanologically unique, however each of the effusive phases were punctuated by pyroclastic flows generated by dome collapse or Vulcanian explosions.
Phase I of dome growth (mid November 1995 to mid March 1998) was characterized by cyclic growth and frequent collapse of the lava dome, with a mean extrusion rate of 4.5 m$^3$s$^{-1}$ (Wadge et al., 2010) which peaked at about 10 m$^3$s$^{-1}$ (Herd et al., 2005). The volcanic activity of Phase I is described in detail elsewhere (e.g. Aspinall et al., 1998; Young et al., 1998; Miller et al., 1998; Calder et al., 2002; Norton et al., 2002). The cessation of lava extrusion in March 1998 occurred when a dome with a volume of 113 x 10$^6$ m$^3$ was present in the crater (Norton et al., 2002).

Phase II (mid November 1999 to mid July 2003), showed a less variable extrusion rate with a mean of 2.9 m$^3$s$^{-1}$ (Wadge et al., 2010), peaking at about 4 m$^3$s$^{-1}$ (Herd et al., 2005). This resulted in the construction of a larger volume dome than any extruded during Phase I. Dome collapses were less frequent than during Phase I but involved larger volumes, culminating in the largest volume collapse of the eruption to date in July 2003. This removed almost the entire volume of the dome (~200 x 10$^6$ m$^3$ Edmonds et al., 2006). The ensuing period of pause lasted 24 months, four months longer than the first pause and to date is the only prolonged period of the eruption with a relatively dome-free crater.

Phase III (early August 2005 to early April 2007), started slowly and the growth rate for most of August was slow and steady (0.5-0.7 m$^3$/s). Low extrusion rates continued through December and January (3.2-3.9 m$^3$/s), but the extrusion rate exceeded 10 m$^3$s$^{-1}$ in February 2006 and remained high for the rest of the extrusive episode. The dome collapse event on the morning of May 20$^{th}$ 2006 involved ~ 100 x 10$^6$ m$^3$ of dome material and emptied the crater; however an observation flight later that day revealed new extrusion in the vent area. Subsequent growth continued until extrusive activity stopped in April 2007 leaving a dome of ~ 200 x 10$^6$ m$^3$ in the crater. This was the largest dome present during any pause for the current eruption. Phase III was characterized by much less pyroclastic flow activity; to such an extent that the first flow large enough to reach the sea occurred on the morning of May 20$^{th}$ 2006.
**Phases IV and V volcanic activity**

The first three phases of extrusion were characterized by extended periods (on the order of years) of lava production separated by pauses of similar lengths. This was not the case for Phases IV and V. Phase IV was characterized by two separate episodes of extrusion punctuated by explosions, the first episode was in July-August 2008 and the second in December 2008-January 2009. The first episode of Phase IV started with a Vulcanian explosion on July 28th 2008 while the December 2008- January 2009 episode started and ended with Vulcanian explosions on December 3rd 2008 and Jan 3rd 2009 respectively. An approximate volume of $39 \times 10^6$ m$^3$ of andesite was extruded for Phase IV (Wadge et al., 2010). Phase V was also short lived (early October 2009 till mid February 2010), and extruded ~ $74 \times 10^6$ m$^3$. As was the case with Phase IV, Phase V produced a high occurrence of explosive activity (Stinton et al., this volume). The extrusion was also cyclic and the cycles were characterized by periods of intense pyroclastic flow generation which would wax and wane on time scales ranging from 4 to 13 hours (Odbert et al., this volume). The textures present in the Phase IV and Phase V products are similar to that noticed in Phases I, II and III. There however seems to be an increase in the vesicularity of the andesite products with friable hand specimens being more commonplace in Phases IV and V. The presence of banded pumice in the July 28th 2008 explosion products was a new occurrence in the eruption.

**Aim of Current Work**

The initial extrusive episode, Phase I received considerable attention with regards to the petrological and geochemical characteristics of the eruptive products (e.g. Barclay et al., 1998; Devine et al., 1998a; 1998b; Murphy et al., 1998; Murphy et al., 2000; Higgins & Roberge, 2003; Zellmer et al., 2003a; 2003b). The initial research was focused on constraining pre-eruptive storage conditions of the erupted
andesite (e.g. Barclay et al., 1998; Couch et al., 2003a; Rutherford & Devine, 2003) as well as establishing the role of the intruding mafic magma in the onset and fueling of the eruption (e.g. Murphy et al., 1998; Murphy et al., 2000; Devine et al., 2003). Long lived eruptions (such as the eruption at Arenal volcano, Reagan et al., 1987) can produce products with temporal variations in chemistry and mineralogy, which may provide useful data for unlocking processes responsible for driving such eruptions (e.g. Streck et al., 2002; Bolge et al., 2006; Ryder et al., 2006).

The preliminary research has without doubt provided valuable information about the processes responsible for triggering and driving the eruption and hence influencing the eruption style. However the ongoing nature of the eruption now provides an opportunity to address any temporal variations in textures and chemistry of the minerals as well as the bulk rock compositions that may have occurred as the eruption proceeded. The aim of this publication is to present a holistic overview of geochemical and petrological data collected from the enclaves and host andesite throughout the current Soufrière Hills eruption. The Phase IV and Phase V products are compared to the earlier products, highlighting any temporal variations present and their bearing on magma chamber and eruption processes. In particular, we focus on temporal variations in bulk rock FeO, and propose that this is linked to similar compositional variations in some mineral phases. We also make a brief comparison to the Mt Pelée products where the processes responsible for generating the andesitic compositions are well constrained (e.g. Dupuy et al., 1985; Fichaut et al., 1989a; Pichavant et al., 2002).

ANALYTICAL TECHNIQUES

Microprobe Analysis

Samples from Phase IV were analyzed using a Cameca 5-spectrometer SX-100 instrument at the Department of Earth Sciences University of Cambridge. Minerals were analyzed using a 2 µm, 15 kV, 10
nA beam for major elements and a 100 nA beam for minor and trace elements. Glasses were analyzed using a 10 µm, 15 kV, 2 nA beam to avoid alkali migration (Devine et al., 1995; Humphreys et al., 2006), with a 10 nA beam for minor and trace elements (e.g. Cl, Ti, Mg).

The Phase V minerals were probed using a JEOL8600 four spectrometer instrument at the Research Laboratory for Archaeology & the History of Art, University of Oxford. Minerals were analyzed with a 15 keV, 15 nA, 1 micron beam. Peak analysis count times were 30s, except for Na (20s) and Cl (40s).

Glass analysis was carried out with a 15 keV, 6 nA, 10 µm diameter beam. Peak count times for all elements were 30s, except for Na (10s), Cl (40s) and P (60s). Amphibole structural formulae were recalculated on the basis of 23 O atoms following Schumacher (1997) and oxide minerals using Stormer (1983).

**Bulk Rock Analysis**

Andesite and enclave XRF data from Phase II along with some Phase III andesites were obtained by the MVO with the analyses performed at the University of Leicester labs under MVO contract. This is the same laboratory that produced the Phase I data presented in Murphy et al. (2000) and Zellmer et al. (2003a). The remaining andesite bulk rock data from Phase III and all of Phases IV and V were obtained from the University of East Anglia under MVO contract. Enclave bulk rock data from Phase III onwards was also obtained from the University of East Anglia under MVO contract. Bulk enclave data from Phase II were from two sources, the first being McGill University in Canada and the second from the British Geological Survey (BGS) labs in Nottingham England. Detailed information about lab, instrumentation and samples analyzed by each lab is presented in Table 1. Bulk rock data is unavailable for Phase IV enclaves due to the limited volume of observed basaltic material (mm size inclusions) extruded and the lack of extensive deposits available for sampling. Microprobe glass and mineral chemistry data were however obtained from mafic fragments in some thin sections.
Andesite and enclave bulk rock XRF analyses of Phase I major and trace elements were compiled from the literature (Devine et al., 1998a; Murphy et al., 1998; Murphy et al., 2000; Zellmer et al., 2003a).

Previous petrological studies on the eruption products of Phases I, II and III, are present in a variety of sources including Barclay et al. (1998), Devine et al. (1998b), Murphy et al. (2000), Couch et al. (2003a; 2003b), Zellmer et al. (2003a), Humphreys et al. (2009a; 2009b), Barclay et al. (2010) Humphreys et al. (2010). Here we summarize the Phase I bulk rock and petrological characteristics of the Phase I, II and III products from the literature.

**Andesite Textures**

Samples are generally crystal-rich (33-63 vol% phenocrysts), with a phenocryst assemblage dominated by plagioclase (64-78% of phenocrysts) complemented by lesser amphibole (8-28%), orthopyroxene (3-7%) and microphenocrysts of Fe-Ti oxides (1.5-3%). Maximum phenocryst sizes are typically ~7 mm for hornblende, ~3-4 mm for plagioclase and ~5 mm for orthopyroxene. Clinopyroxene microphenocrysts (< 1%) are present, as well as accessory apatite. Quartz is also present as rounded, embayed crystals sometimes jacketed by clinopyroxene. The variations in modal mineral contents are presented in Table 2.

The andesite groundmass is generally microcrystalline but can contain up to 25% rhyolitic glass in rapidly erupted samples. Cristobalite is present in slower erupted dense dome samples and can be up to 15 wt% in the groundmass, or more commonly as a vapor phase precipitate in the vesicles. The glass is commonly observed to have undergone phase separation and or devitrification in dome samples. Hornblende is replaced by orthopyroxene and clinopyroxene, thus yielding a groundmass normally comprising plagioclase, two pyroxenes and Fe-Ti oxides, with wide variations in microlite textures. No systematic differences in texture modal proportion or mineral composition have been observed within the andesites of Phases I, II and III (Table 2).
Enclave Textures

Mafic enclaves have been present in the eruptive products of the Soufrière Hills volcano as far back as 24ka (Wadge & Isaacs, 1988). The enclaves from Phases I, II and III are typically small (< 30cm) with a maximum reported length of 80 cm (Barclay et al., 2010; Plail et al., this volume); their shapes vary from ellipsoidal to angular. The ellipsoidal enclaves have smooth or crenulate, chilled contacts with the andesite. Small mm-scale fragments of enclaves are also observed in the andesite. The enclaves contain similar mineral phases to that of the andesite; they are however recognized by the higher abundance of mafic minerals, and a higher abundance of clinopyroxenes relative to orthopyroxene. There are no true phenocrysts present, but based on chemistry and texture there are andesite-derived xenocrysts of plagioclase, amphibole and orthopyroxene present in the enclaves.

The xenocryst contents can be up to 5% by volume for Phase I enclaves (Murphy et al., 1998; 2000), and up to 27% for Phase III; two thirds of which are normally plagioclase (Plail et al., this volume). The enclave groundmass is typically fine grained, consisting of randomly oriented interlocking elongate or acicular crystals of plagioclase, orthopyroxene, clinopyroxene and amphibole forming a diktytaxitic framework. About ~10% of residual rhyolitic glass with variable amounts of devitrification is observed in most thin sections, banded textures are also present in some scoriaceous samples.

Mineral Textures and Composition – (Andesite and Enclave)

Due to their similarities in crystal phases, the initial findings on the mineral compositions of the andesite and enclaves are both summarized together in this section. A through overview of the mineral chemistry from the early erupted products is presented in Barclay et al. (1998), Devine et al. (1998a; 1998b) and
Murphy et al. (1998; 2000) for Phase I and Humphreys et al. (2009a; 2009b) for the products of Phases II and III.

Plagioclase

Plagioclase is present as phenocrysts, microphenocrysts and microlites in the andesite with the phenocrysts showing a variety of zoning textures (Table 3; see also Murphy et al., 2000; Humphreys et al., 2009a; 2009b). Oscillatory and patchy zoning are common in the phenocryst cores, including the cores of sieved crystals, which are interpreted to have experienced direct contact with melt of a more mafic nature. The modal proportion of each plagioclase textural type varies between samples. Plagioclase is present in the enclaves mostly as microlites, with minor inherited phenocryst from the andesite also present.

Plagioclase phenocrysts in the andesite have sodic cores (typically An$_{48-58}$, but with calcic zones up to An$_{80}$) with either normal or reversed zoned rims, while the microlites have more calcic cores An$_{60-75}$ and can be normally, or to a lesser extent, reversed zoned. The plagioclase microlite cores in the enclaves are calcic up to An$_{93}$ and normally zoned. Apart from the differences in anorthite content, the andesite phenocrysts generally have lower Fe contents than the enclave crystals. Most andesite microlites, some microphenocrysts and the sieved rims are generally richer in FeO content relative to the phenocryst rims, their elevated FeO contents are similar to that observed in the enclave plagioclase crystals (Figure 1e & f, Table 4).

Amphibole

Amphibole phenocrysts in the andesite show green-brown pleochroism and can show oscillatory zoning (Rutherford & Devine, 2003; Humphreys et al., 2009b). Amphiboles can have irregular rims (indicating slight resorption) or pristine rims, but commonly show a variety of breakdown textures, of which three
end-members can be identified (summarized from Devine et al., 1998a; Murphy et al., 2000; Rutherford & Devine, 2003; Buckley et al., 2006; Humphreys et al., 2009b; Plechov et al., 2008):

(i) Fine-grained intergrowths of clinopyroxene, orthopyroxene, Fe-Ti oxides and plagioclase, growing where the amphibole is in contact with melt. This breakdown texture can pseudomorph the original crystal and is thought to form as a result of decompression during ascent due to changes in $\text{pH}_2\text{O}$. The development of these rims can be used as an indicator of ascent conditions, with the thicker rims developing during slower ascent. The rim thicknesses observed in the samples represent ascent rates of $> 0.019 \text{ m/s}$ (Devine et al., 1998a; Rutherford & Devine, 2003).

(ii) Coarser-grained reaction rims dominated by clinopyroxene (preferentially aligned parallel to the c-axis of the amphibole, and commonly in optical continuity). This rim type can also pseudomorph the entire crystal and is interpreted as a thermal breakdown texture.

(iii) Partial to complete opacitisation, particularly along cleavage and cracks or adjacent to vesicles.

Decompression breakdown rims, thermal breakdown rims and resorbed rims are present in the majority of analyzed samples; typically a given sample will also show a range of decompression rim thicknesses. In the mafic enclaves, some amphibole microphenocrysts have undergone partial breakdown or opacitisation while microlites are normally euhedral and yellowish in appearance. The inherited amphibole phenocrysts in the enclaves most commonly show thermal breakdown textures. Amphibole compositions in the andesite range from Mg-hornblende to Mg-hastingsite, with some pargasite present. Amphibole compositions in the enclaves range from pargasite to Mg-hastingsite. In general, andesite amphibole phenocrysts have lower $\text{Al}_2\text{O}_3$ (6-8 wt%) relative to the enclave amphibole contents of (12-14.5 wt%).

Pyroxenes

Orthopyroxene phenocrysts are typically euhedral, although some show slight rounding, which may be followed by reversely zoned euhedral outer rims. The reversed zoned phenocrysts represent a minor
fraction of the total orthopyroxene population and are not observed in every thin section. Some crystals have rounded cores with clinopyroxene overgrowths. The reverse zoning is interpreted to occur as a result of heating, with the clinopyroxene overgrowth rims indicating direct contact with mafic melt.

Clinopyroxene normally occurs in the andesite as microphenocrysts and microlites as well as overgrowths on orthopyroxene, quartz and sometimes in the thermal breakdown rims of amphibole (Table 3); it can be very common in the groundmass of some thin sections. Clinopyroxene is also common in mafic enclaves, as microlites and microphenocrysts.

There are three orthopyroxene phenocryst types, identified on the basis of having unzoned rims, reversely zoned rims or clinopyroxene overgrowths, all have similar core compositions ($\text{En}_{56-70}$, $\text{Fs}_{26-41}$, $\text{Mg#}_63-68$) where ($\text{Mg#} = \text{atomic units Mg}/(\text{Mg+Fe}^{2+})$). Type 1 consists of unzoned or weak normally zoned crystals with rim compositions $\text{En}_{57-61}$ and $\text{Wo}_{1.8-2.3}$. Type 2 consists of reversed zoned crystals with rim compositions $\text{En}_{62-73}$ and $\text{Wo}_{2.2-3.7}$. Type 3 consists of crystals with clinopyroxene overgrowths.

Orthopyroxene microlites have Mg#s in the range 58-74 and display the second most magnesium rich compositions of the orthopyroxenes present in the andesite after the reversed zoned phenocryst rims (Table 6a & 6b). Orthopyroxene in the mafic enclaves have similar Mg#s to the andesite microlites. The clinopyroxenes have Mg#s in the range 66-74, and they are not in equilibrium with the orthopyroxene phenocrysts of the andesite.

**Fe-Ti Oxides**

In the andesite, phenocrysts are rare while microphenocrysts (>100 µm) are common and can be zoned (Table 3) with rounded irregular cores, resorbed rims and exsolved textures. Fe-Ti oxides also occur as inclusions in the phenocrysts and microphenocrysts of amphibole, pyroxene and plagioclase, where they tend to be subhedral. Enclave microphenocrysts can be euhedral homogeneous crystals or heterogeneous crystals with exsolution textures (Table 3).
Titanomagnetite is the most common oxide in the eruptive products, with minor ilmenite present.

Individual hand specimen samples can show significant inter-crystal compositional variation; the extent of the variation can be such that Devine et al. (1998a) reported observing reversed zoning in half of the titanomagnetite crystals analyzed. The oxide compositions have shown little temporal variation for the first three eruption phases. The andesite titanomagnetite microlites and enclave titanomagnetites generally have higher titanium contents than the andesite microphenocrysts with the microphenocryst rims being slightly more titanium rich than the cores.

**Bulk Rock Composition**

To date only the Phase I bulk chemistry has been published, the products are predominantly andesite displaying SiO\(_2\) contents in the range (58-62 wt%), (Murphy et al., 2000; Zellmer et al., 2003a; Barclay et al., 2010), with an average composition of 59.7 wt% SiO\(_2\) (Murphy et al., 2000). The enclaves are reported to have SiO\(_2\) contents of 51-56 wt% (Murphy et al., 2000) and 49-55.5 wt% (Zellmer et al., 2003a). Both compositions are low-K with the andesites having a calc-alkaline affinity while the enclaves are of tholeiitic affinity (Murphy et al., 2000). Bulk geochemical trends are similar in both lava types. With increasing SiO\(_2\); TiO\(_2\), Al\(_2\)O\(_3\), FeO, MgO and CaO all show decreasing trends, while Na\(_2\)O and K\(_2\)O show increasing trends. The Phase I andesites have relatively restricted major element compositions and the differences in the modal proportions of mineral phases were proposed as a likely source of the andesite heterogeneity within Phase I (Zellmer et al., 2003a).

**Initial Conclusions**

The experiments of Barclay et al. (1998) showed that the andesites were stored at a minimum pressure of 115-130 MPa at temperatures (820-840) °C before being erupted, with measured water contents in the range (4.27 ±0.54) wt% H\(_2\)O (Barclay et al., 1998), 4.7 wt% (Devine et al., 1998a) or up to 6.2 wt% H\(_2\)O (Humphreys et al., 2009b). Textural characteristics of the amphibole phenocrysts such as coarser grain
reaction rims dominated by clinopyroxene are thought to represent thermal breakdown during reheating
events, while partial to complete opacitisation, particularly along cleavage and cracks or adjacent to
vesicles is thought to develop during shallow dome storage with passive gas fluxing (e.g. Garcia &
Jacobson, 1979; Devine et al., 1998b; Humphreys et al., 2009b).

Couch et al. (2003b) proposed that the textural variations observed in the andesitic groundmass can be
attributed to variations in ascent rates in a system undergoing open system degassing. The presence of
enclaves in the erupted lavas (Murphy et al., 1998), the reverse zoning in the orthopyroxene and
plagioclase phenocrysts (Murphy et al., 1998; 2000) along with the higher temperature groundmass
assemblage (Humphreys et al., 2009a) were used as evidence that the eruption was triggered by an
intruding mafic magma, heating and remobilizing the andesite. It was also demonstrated that amphibole
fractionation is important in driving the enclave compositions (Zellmer et al., 2003a).

Reheated textures are fairly localized in the andesite (Murphy et al., 1998; 2000), which is evident in
the low modal abundance of reversed zoned plagioclase and orthopyroxene phenocrysts. Based on the
absence of amphibole microlites in the Phase I andesite, Murphy et al. (1998) had initially suggested that
enclave disaggregation was minor if at all present. More recently, Humphreys et al. (2009a) have shown
that the physical destruction and dispersion of some enclaves also contributes to the mafic nature of the
microlite assemblage in the Phase II and Phase III andesite groundmass, while Genareau & Clarke (2010)
showed that basaltic melt microscopically intrudes the margins of the andesite. Based on the thickness of
amphibole thermal breakdown rims, extrusion occurs after reheating on time scales of hours to months
(Devine et al., 2003; Rutherford & Devine, 2003; Devine & Rutherford, this volume).

Zellmer et al. (2003a) showed that the currently erupting Soufrière Hills andesite can be generated by
fractional crystallization of plagioclase and amphibole from the South Soufrière Hills (SSH) basalts. In
contrast, while the SSH lavas are compositionally similar to the mafic enclaves of the current Soufrière
Hills eruption (Figures 9 & 10), incompatible trace element ratios demonstrate that neither the current andesite nor the SSH lavas can be derived from the mafic enclaves (Zellmer et al., 2003a).

RESULTS

Phases IV and V

The textures of andesite samples erupted in Phases IV and V vary from the banded pumice produced during the July 28th 2008 explosion to the dome rock produced throughout both phases. We have already discussed how the volcanic activity of Phases IV and V differed to the first three extrusion phases; we now present the results of petrological and geochemical analysis from Phases IV and V.

Textures

The mineral phases present in the respective andesite and enclave lavas have remained consistent throughout the five eruption phases. Mineral textures associated with disequilibrium in Phases I, II and III andesites e.g. reversely zoned orthopyroxene phenocrysts, sieve textured plagioclase and embayed quartz are all observed in the andesites of Phases IV and V. Similarly to Phases I, II and III, amphibole is absent from the andesite microlite assemblage. Dome samples from Phases IV and V are generally crystal rich (~30-46% porphyritic) with phenocrysts and microphenocrysts of plagioclase, amphibole and orthopyroxene present along with microphenocrysts of clinopyroxene and Fe-Ti oxides (Figure 2). The groundmass in dome samples can be coarsely crystalline and vesicular (up to 35%) with microlites and glass present.

The enclave textures observed in Phases IV and V are also similar to those observed in the previous three phases. Plail et al. (this volume) categorizes the enclaves using parameters such as the volume of crystals inherited from the host andesite. They have identified three distinct enclave textures: a diktytaxitic framework dominated by plagioclase and pargasitic amphibole (Figure 3a); a diktytaxitic framework dominated by plagioclase and clinopyroxene (Figure 3b) and a composite (mixed enclave) of the above
two textures (Figure 3c). Xenocrysts (plagioclase, hornblende and orthopyroxene) are present in all enclave types (Plail et al., this volume).

The inter-phase differences between andesites of the eruption may be subtle. For example, some of the dome rock samples from Phases IV and V tend to be less dense and friable; their groundmass tends to be finer grained than the denser dome blocks. The feature seems more common in Phases IV and V relative to the earlier phases. Secondly, while the mafic enclaves are ubiquitous throughout the eruption, there is an apparent increase in enclave abundance over time (see discussion in Plail et al, this volume). The average macroscopic modal abundance of enclaves in the andesite products ranges from 1-2% in the Phase I products (Murphy et al., 1998; 2000) to up to 8% in the products of Phase III (Barclay et al., 2010) and 8.5 % in Phase V (Plail et al., this volume), while Komorowski et al. (2010) report a mafic content of 12% in the tephra generated from explosions during Phase IV activity (although these three sets of values were determined using different methods).

Banded Pumice

The banded sample from the July 28th 2008 explosion (sample number MVO1532d) is atypical of previously erupted samples. The sample shows clear textural heterogeneity in the form of whitish and dark grey streaks dissimilar to the chilled margins displayed by the mafic inclusions and resembles banded pumice. The sample comprises of a silicic part with pale brown to colourless glass, very few microlites and relatively low vesicularity, as well as a more mafic part which is dark brown with a crystalline groundmass and no obvious clear glass. Clearly mixed patches containing brownish glass are evident. The ‘mixed’ and mafic parts occur together spatially in patches hundreds of µm across, while the silicic parts form larger patches.

The mixed and mafic parts contain the typical assemblage present in the andesite such as phenocrysts of plagioclase with oscillatory and sieve textures, hornblendes, orthopyroxenes with clear rims or overgrown by clinopyroxene, and microlites of plagioclase + 2 pyroxenes + magnetite. In contrast, the silicic part
contains hornblende, no sieved plagioclase, only pristine orthopyroxene and few small microlites (~5-10 µm) of plagioclase, orthopyroxene and titanomagnetite; clinopyroxene is absent. Hornblende phenocrysts are commonly either pristine or have a very thin incipient reaction rims (5-8 µm). The silicic matrix also contains numerous larger (50-200 µm) broken fragments of plagioclase phenocrysts. Euhedral, prismatic quartz microlites are common; these are normally ~50 µm (±20) in size and are not typical of the Soufrière Hills andesite. They do not replace the typical rounded, embayed quartz phenocrysts, which are present as usual.

Plagioclase Composition

The Phase IV and Phase V andesite plagioclase phenocryst cores and rims exhibit a wide range of compositions (An$_{46}$ to >An$_{80}$, Table 4) which overlaps with phenocryst compositions from the previous phases (Figure 1a). The oscillatory plagioclase cores are generally more calcic than the cores of phenocrysts with other textures (Figure 1d). The mean An content for both Phase IV and Phase V phenocryst rims is ~ An$_{56}$, within error of the mean An contents from phenocryst rims of the previous phases (Table 4, Figure 1a). As with the first three extrusion phases, the microlite rims of Phases IV and V are more calcic than their respective phenocryst rims (Figure 1a & b, Table 4), and the enclave microlites are generally more calcic than the andesite microlites (Figure 1b). There is inter-phase variation of Fe contents in the andesite hosted microlite crystals, it is however non-systematic.

Amphibole Composition

Phase IV and Phase V andesite amphiboles are predominantly magnesio-hornblende, the same as the andesite amphiboles from Phases I to III (Figure 4a) with minor amounts of tschermakite. Mg-numbers are in the range 0.44-0.55. There is a systematic change in enclave amphibole FeO content (Figure 4d) from Phases I to III (unfortunately, no data from Phase IV enclave amphibole is available since no enclave products from Phase IV were collected). The enclave amphiboles from Phases I to III display a
steady increase in magnesium numbers (0.5 to 0.59), and the SiO₂ content also decreases from Phases I to III (Table 5b, Figure 4c &d). The absence of Phase IV enclave amphibole data produces an undesired gap in the dataset however, it can be seen that the Phase V enclave amphibole have higher mean contents of wt% Al₂O₃, MgO, CaO, TiO₂ and Na₂O relative to the amphibole in the enclaves of the first three eruption phases (Table 5b). Apart from having the highest FeO and SiO₂ contents, the Phase I enclave amphiboles are lowest in wt% Al₂O₃, MgO and TiO₂.

Orthopyroxene Composition

Andesite orthopyroxene phenocryst core compositions have been relatively homogeneous throughout the eruption (En₅₆-₇₀, Fs₂₆-₄₁) with a mean magnesium number of 0.6 (Table 6a). The normally zoned rims (Mg# 0.58-0.65 mean ~ 0.61) and reversed zoned rims (Mg# 0.68-0.75 mean ~ 0.7) have been fairly unchanged during the eruption (Figure 5a). The andesite microlites generally have higher magnesium numbers, as well as higher CaO and Al₂O₃ contents, than the phenocryst cores throughout the eruption (Figure 5b). Microlites from Phases I, II, III and IV have the same mean magnesium number of 0.67 (Figure 5b). However, the microlites of Phase V have a slightly lower mean magnesium number of 0.63 (Table 6a, Figure 5b).

Clinopyroxene Composition

The clinopyroxene composition has varied little throughout the eruption; they are mostly augites, calcic-augites and diopsides with cores and rims in the compositional range En₃₉-₄₉, Wo₃₁-₄₇ and mean compositions of En₄₂-₄₄, Wo₄₀-₄₃. Kushiro (1960) has demonstrated that it is the SiO₂ wt% content of a magma and not temperature that is the major control of the SiO₂/Al₂O₃ ratio of the crystallizing clinopyroxenes and that the ratio increases with increasing magma SiO₂ content. The andesite clinopyroxenes show distinct SiO₂/Al₂O₃ ratios by eruption phase (Table 6c). The enclave clinopyroxenes also show a similar inter-phase trend in FeO from Phase I to III like the enclave amphiboles.
Oxide Composition

As was the case in the first three eruptive phases, titanomagnetite is the most common oxide in the products of Phases IV and V with only minor ilmenite present. The andesite microphenocryst titanomagnetite cores and rims from the five extrusion phases have overlapping compositions (21-30 mol% Usp), with no systematic variation between phases. The Phase IV microlites have higher TiO$_2$ contents (up to 16 wt%) than the microphenocrysts (7-10 wt%), as is the case with the titanomagnetites from Phases I, II and III. However, this is not the case for Phase V where the microlites have similar TiO$_2$ wt% contents to the microphenocrysts from Phase V (Table 7a).

Glass composition

The andesite and enclave glass compositions for the products of Phases IV and V have shown little variation relative to the previous phases, glass compositions have been consistently rhyolitic in both magma types (70-82 wt% SiO$_2$) throughout the eruption (Table 8). Some authors have shown that the glass present in both magma types are essentially indistinguishable (e.g. Murphy et al., 1998; 2000), while others (e.g. Humphreys et al., 2010) have shown evidence indicating subtle differences between enclave and andesite glass. The K$_2$O content shows a bimodal distribution of high and low K$_2$O glass samples with some intermediate compositions observed (Figure 6).

Humphreys et al. (2010) interpreted the glass compositions enriched only in K$_2$O, as an indication of diffusive contamination by high-K mafic inclusion glass; some samples are also enriched in TiO$_2$, suggesting physical mixing of remnant glass. In general with increasing SiO$_2$, there is a decreasing trend of Al$_2$O$_3$, while K$_2$O increases (Figure 6). There are a few samples with SiO$_2$ contents of 80-82 wt% that have high Na$_2$O, high CaO, low FeO and low K$_2$O, which are likely due to post extrusion phase
separation of two types of glasses of differing chemistry as described by Cashman (1992) for the Mt St Helens lava dome and indicated by Humphreys et al. (2009a) for the products of Phases II and III.

**Bulk Rock Composition**

The andesites of Phases IV and V have a similar range in bulk rock SiO$_2$ (56-62 wt%) to the products of the previous phases (Table 9). The trends of decreasing TiO$_2$, Al$_2$O$_3$, FeO, MgO and CaO and increasing Na$_2$O and K$_2$O with evolution reported for the Phase I products are present in the products of the later eruptive phases (Figure 7). There is considerable inter-phase compositional overlap for most elements at similar SiO$_2$ contents. As is the case with the andesite, the enclave products of the later phases show compositional overlap and similar trends to the Phase I enclave products.

**Whole Eruption**

**Trends and Variations in Bulk Rock Composition**

There are some clear differences between the eruption phases in terms of bulk rock composition. Firstly, the FeO contents show systematic inter-phase variation with the mean FeO contents showing a clear decrease during the first three phases. The andesite products of Phase I have a mean FeO content of (7.32 wt%), while the Phase II and Phase III products have mean values of 7.14 and 6.6 wt% FeO respectively (Table 9). There is a relative increase in mean FeO to 7.4 wt%, for samples erupted during Phase IV followed by a subsequent decrease in Phase V to 7.08 wt% FeO (Figure 8a, Table 9). Although not as systematic as FeO, MgO contents also show some variation in the andesite (Table 9).

The enclave bulk rock FeO trend behaves in a similar way to the andesite bulk FeO trend (Figure 8a). The inter-phase contrast in FeO content is however more pronounced in the enclaves than in the andesites, thus at approximately 50 wt% SiO$_2$ there is a bulk rock compositional difference of up to 2 wt% FeO between the enclave products of Phases I and III (Figure 7). One observation of note is that the difference between the mean bulk enclave FeO content and their syn-eruptive host andesites shows a
steady decrease with a change of ~ 1% over the eruption, from 2.3 wt% in Phase I to 1.4 wt% in Phase V, this is outside the error of the FeO contents. In other words, the difference between enclave and host FeO contents has been narrowing with time. There are also subtle differences in MgO and CaO that are outside analytical error.

Also of note is that the Phase I and Phase II andesite compositions are more scattered than those of Phases III, IV and V (Figures 7 & 10). This results in better defined linear trends of (for example) Al₂O₃, CaO, TiO₂ and Sc with increasing FeO/MgO for the andesite products from Phases III, IV and V; linear trends are also present in the enclave products of Phases III and V (Figure 10). There is a bit of scatter at higher ratios (>2.2 FeO/MgO) in the Phase III and Phase V enclaves in all of the plots. The enclave products of Phases III and V define a different slope to the Phase I and Phase II products on the SiO₂ vs. FeO plot (Figure 7) and thus together seem distinct from the products of Phases I and II. Phase IV enclave data are not available, but given the behaviour of the Phase IV andesite products and the apparently coupled FeO variation of both magmas, we anticipate that the bulk chemistry of the Phase IV enclaves would quite likely show well defined trends with little scatter similar to that observed in the enclaves of Phases III and V. It must be noted that the inter-phase variation in plagioclase microlite FeO content (Figure 8d) does not mirror bulk rock inter-phase FeO variation (Figure 8a).

Generally, the Phase V andesites have the highest SiO₂ contents while the Phase II and Phase IV andesites have the lowest SiO₂ contents. The Phase V enclaves are also the most SiO₂-rich (Plail et al., this volume); regardless to this the SiO₂ contents of the enclaves and host andesites are not coupled. This is highlighted by the Phase II enclaves being more SiO₂-rich than the Phase I and Phase III enclaves (Table 9). The Phase V andesites also show elevated total alkali contents, relative to Phases I, III and IV andesites (Figure 5c) as well as lower wt% TiO₂, Al₂O₃, MgO and CaO (Table 9). Phase IV andesites have the highest wt% FeO, MgO and Al₂O₃ while the Phase II andesites are richer in CaO wt%. The Phase V enclaves are richer in total alkalis.
Due to its amalgamated nature, the integrity of the bulk rock data set needs be examined in order to verify whether the observed variations in FeO are real or an artifact of using different laboratories over several years. This is addressed in Appendix I where we show that the observed variations are independent of laboratory bias and are indeed real.

Estimates of temperature, pressure, water content and log fO2

A number of different methods have been used to estimate the pre eruption temperatures of the andesites and mafic enclaves, including estimates obtained from orthopyroxene (using QUILF, Anderson et al., 1993) in single pyroxene mode, following Murphy et al. (2000), use of magnetite-ilmenite pairs (e.g. Devine et al., 2003), the hornblende-plagioclase geothermometer of (Blundy & Holland, 1990), the 2-pyroxene geothermometer of Lindsley (1983), the plagioclase-liquid thermometer of (Putirka, 2005), and clinopyroxene-melt equilibria of Armienti et al. (2007), after Putirka et al. (1996; 2003) and Putirka (1999) which also produces pressure estimates. Pressures were also estimated with VolatileCalc (Newman & Lowenstern, 2002) by measuring glass volatile contents and assuming H2O saturation.

For Phase I, these approaches generated values of 785-980 °C for the andesite orthopyroxene phenocryst cores with an average of 851 ±20 °C (Murphy et al., 1998; 2000), a similar temperature range of 834-850 °C was obtained by Devine et al. (1998a), and 820-840 °C by Devine et al. (2003) both using magnetite-ilmenite phenocryst pairs. This is consistent with the experiments of Barclay et al. (1998) which showed that the andesites were stored at temperatures 820-840 °C before being erupted. The reversed zoned orthopyroxene rims give higher temperatures up to1100 °C (Murphy et al., 1998; 2000). Enclave hosted orthopyroxene cores give temperatures in the range 1020-1050 °C (Murphy et al., 1998). New data obtained by MVO for Phases II,III,IV and V (some Phase II and Phase III data reported in Humphreys et al., 2009a) are reported below and presented in Table 11.
Pyroxene: Orthopyroxene phenocryst cores in the andesite give temperatures of 824-965 °C (mean 866 °C), with unzoned rims at 845-895 °C (mean 869 °C) and reversely zoned rims at 1018-1032 °C (mean 1025 °C). Orthopyroxene inclusions in other minerals or in crystal clots also gave low temperatures 809 °C. Orthopyroxene microlites in the andesite gave temperatures of 958-1017 °C, while orthopyroxene in mafic enclaves gave 903-1305 °C (mean 1070 °C), with 1074-1196 °C for the two-pyroxene geothermometer. These two-pyroxene temperatures may not reflect equilibrium conditions.

Plagioclase-liquid: The plagioclase-liquid thermometer produced a wide range of temperatures from 821 °C (for melt inclusions trapped in plagioclase phenocrysts) to 1100 °C (for matrix glasses paired with microlite rims); with increasing temperatures correlating with decreasing melt H₂O content. This is similar to the style of variation reported in Blundy et al. (2006) but given the huge range in temperature we suggest that many of the higher groundmass temperatures may reflect disequilibrium.

Hornblende-plagioclase: Inclusions of plagioclase in andesite hornblende phenocrysts give temperatures in the range 804-890 °C (mean 833 °C). Paired analyses from crystal clusters, interpreted as fragments of disaggregated mafic enclaves, give temperatures of 849-947 °C (mean 894 °C).

Clinopyroxene-melt: The clinopyroxene-melt thermometer gave temperatures in the range 1097-1145 °C (mean 1110 °C) for microlites/matrix glass and microphenocrysts/matrix glass in the mafic enclaves. As with the two-pyroxene and plagioclase-melt temperatures, these may not reflect equilibrium conditions.

Pressure and water estimates: Pressure estimates obtained from the Cpx-melt equilibrium for crystals from both enclaves and andesite range from 10-280 MPa with a mode of 50-100 MPa. Previous measurements of melt H₂O contents from melt inclusions are in the range 4.27 ± 0.54 wt% H₂O (Barclay et al., 1998), ~4.7 wt% (Devine et al., 1998a) and 1.1-6.2 wt% (Humphreys et al., 2009b). Assuming water saturation in the absence of CO₂ (Newman & Lowenstern, 2002), these estimates equate to pressures up to 214 MPa which is consistent with the estimates from the Cpx-melt equilibrium.
Temperature variation

Different methods tend to produce distinct temperature ranges and this is probably because they are recording temperatures from different parts of the reservoir or localized equilibrium at differing times. Taken together, the hb-plag temperatures, single-orthopyroxene core temperatures, 2-oxide temperatures and plagioclase-liquid temperatures from melt inclusions are probably representative of phenocryst crystallization and hence the ambient temperature within the andesite storage region. These temperatures are typically in the range 785-900 °C with an average at the lower end of that range and are consistent with the experiments of Barclay et al. (1998) on Phase I products.

Higher temperature estimates are obtained from the disequilibrium features outlined above, e.g. 826-1101 °C from reverse zoned orthopyroxene rims (Table 11). This is consistent with the very An-rich compositions observed in sieved plagioclase crystals, which requires very high temperatures and a more mafic melt (Couch et al., 2003a). These temperatures are inferred to be recording the reheating of the andesite crystals by the basalt intruding into the base of the reservoir. The estimates from the enclaves are expectedly higher, 903-1305 °C for the single orthopyroxene method, 1074-1196 °C using the 2 pyroxene method and 1097-1145 °C using the clinopyroxene-melt equilibrium, although some of these temperatures may also reflect disequilibrium mineral compositions.

The variability within these estimates may be due to heterogeneities within the andesite storage region, and/or due to the localized nature of the basaltic intrusion and thus equilibrium at higher temperatures (e.g. Pichavant et al., 2007). The temperature estimates obtained using the various methods all show a range of values with the smallest difference between maximum and minimum temperatures being 14°C for any given method. There is no evidence in our data to suggest systematic changes in temperature as the eruption proceeds. However Devine & Rutherford (this volume) used oxide thermometry to show that the mean ambient temperature of the andesite before being reheated shows a 10 °C increase from Phase I (825 °C) to Phase V (835 °C).
Estimates of log \( f_{O_2} \)

Log \( f_{O_2} \) values calculated from andesite hosted ilmenite-magnetite pairs are presented in Table 8. The inter-phase values show some overlap with no clear trend. The Mg-Fe exchange coefficient \( K_d \), (Putirka et al., 2003) is used to establish mineral melt equilibrium of clinopyroxenes where \( K_d \text{ cpx-liq} = [(X_{MgO \text{ liq}}/X_{FeO \text{ Cpx}}) / (X_{MgO \text{ Cpx}}/X_{FeO \text{ liq}})] \). Suitable mineral-melt equilibrium is thought to be attained when a \( K_d \) value of 0.27±0.03 is obtained for clinopyroxene. Bédard (2010) showed that clinopyroxene \( K_d \) values will vary as a function of temperature, pressure, log \( f_{O_2} \) as well as melt SiO\(_2\) and, total alkalis.

Box plots of clinopyroxene microlite \( K_d \) values by eruption phase are presented in Figure 5d, the \( K_d \) value for the clinopyroxene microlites varies by eruption phase with Phase III having the highest median values while Phases V and II have the lowest median values. The data obtained from the Cpx-melt equilibria showed no systematic variation in pressure and temperature with eruption phase. The total alkali contents display a negative correlation with the clinopyroxene \( K_d \) values, hence it seems the Cpx \( K_d \) values are responding to changes in total alkali content and not log \( f_{O_2} \).

DISCUSSION

Although the andesitic rocks erupted since 1995 at Soufrière Hills are fundamentally similar in terms of textures and mineral compositions, there are some key systematic variations over time that may reveal details about the nature of the plumbing system at depth. In particular, the key observation is that the bulk rock FeO content of both andesite and enclave appear to be coupled. Both andesites and enclaves from Phases I and II show higher major element oxide contents apart from SiO\(_2\), and more scatter at similar FeO/MgO ratios compared with the later phases (Figure 10), they also have higher CaO/Al\(_2\)O\(_3\) ratios at similar FeO/MgO ratios. Here we review the overall relationships between the andesite and enclave magmas and hence interpret the possible causes of these compositional variations.
Mt Pelée Martinique has produced a spectrum of products ranging from basaltic-andesites to dacites; some compositions were only erupted at certain periods in the volcano’s history due to the changing nature of volcano’s plumbing system with time (Fichaut et al., 1989a). Andesites are however ubiquitous in the Mt Pelée deposits and have been of a similar composition for the past 100,000 years (Dupuy et al., 1985). The Mt Pelée andesites also contain microlites inherited from intruding basalts (Martel et al., 2006) as do the Soufrière Hills andesites (Humphreys et al., 2009a).

The volcanic stratigraphy and petrology of Mt Pelée, is well constrained and the physical evolution of the plumbing system through time has been modeled (Fichaut et al., 1989a). It has been established that the current shallow reservoir is zoned (Dupuy et al., 1985, Pichavant et al., 2002) and is relatively small (< 1km³) at about 10km bsl (Fichaut et al., 1989a). It has also been established that the basaltic magmas are parental to the andesites via magma mixing and fractional crystallization (Fichaut et al., 1989b).

Magma mixing occurs in relation to basaltic intrusions which generate eruptions with heterogeneous magma compositions (e.g. Fichaut et al., 1989a; Gourgaud et al., 1989), they are normally larger in volume than the homogeneous eruptions since lower parts of the reservoir gets tapped (Dupuy et al., 1985). Fractional crystallization of the hybrid magmas created by the mixing events produces fairly homogeneous magmas which tend to evolve towards more acid compositions; these are the magmas that feed the smaller volume homogeneous eruptions (Fichaut et al., 1989a; Pichavant et al., 2002). Thus andesite generation at Mt Pelée is fairly well understood.

Andesite phenocryst-groundmass relationship

In the current Soufrière Hills eruption, microlite rims and some cores are generally more mafic than phenocryst rims (in the case of plagioclase) or both cores and rims (in the case of orthopyroxene). This is contrary to what is expected from closed-system crystallization and cooling, and demonstrates the effects
of mafic magma input and the short time scale between heating and eruption. The nature and extent of physical interaction between the andesites and basalts is also evidenced in the ubiquitous presence of mafic enclaves within the andesite, as well as rarer macroscopic features such as the banded pumice. The contrasting temperatures of the intruding mafic magma and the silicic host also leads to heat transfer (Snyder, 2000), resulting in remobilization of the andesite (Murphy et al., 2000).

The andesite contains various disequilibrium textures which provide evidence of reheating driven by mafic magma, including reverse zoned orthopyroxene (Barclay et al., 1998; Murphy et al., 1998; 2000); sieve textures and calcic overgrowth rims on plagioclase phenocrysts (Couch et al., 2003a); two oxide disequilibrium (Devine et al., 1998a; 2003); resorbed quartz with clinopyroxene mantles; thermal breakdown of amphiboles (Humphreys et al., 2009b); and clinopyroxene overgrowths on orthopyroxene (Murphy et al., 1998; Humphreys et al., 2009a). These observations are consistent with the higher range of temperatures obtained from the andesite by some of the thermometry methods employed.

Other Disequilibrium Features
The andesite microlite assemblage generally consists of normally zoned plagioclase with euhedral to hopper or skeletal textures (suggesting rapid growth, Hammer & Rutherford, 2002), normally zoned orthopyroxenes, and normally zoned clinopyroxenes with a skeletal appearance. Although all orthopyroxene microlites are typically more mafic than phenocryst rims, they are not always compositionally identical to the enclave microlites, but can sometimes exhibit intermediate or even more mafic compositions relative to some enclave microlites (Figure 5b).

The absence of groundmass amphibole in the Phase I andesite and the proposed compositional differences between the plagioclase groundmass in the Phase I andesite and the enclave plagioclase were used as evidence against enclave disaggregation being a significant process (Murphy et al., 2000), the absence of groundmass amphibole in the andesite was also used by Couch et al. (2003a) as evidence against magma
mixing. More recently (Humphreys et al., 2009a; Genareau & Clarke, 2010; Humphreys et al., 2010) proposed enclave disaggregation and mixing as significant processes to support chemical and textural observations in the products of Phases II, III and IV.

We have also shown that the Phase I andesite plagioclase microlite population has elevated Fe contents similar to the enclave plagioclase and the andesite microlites from subsequent phases (Table 4, Figure 1e), which suggests that disaggregation probably did occur during Phase I. We have also highlighted the dominance of normally zoned plagioclase microlites in the andesite. These additional observations suggest that the mafic signature of the Soufrière Hills andesite groundmass may not be due to a simple case of crystal transfer due to enclave disaggregation, and that reheating and recrystallization of resident crystals might also be an important process in generating the hotter groundmass assemblage. The lack of change in modal abundance of reverse zoned plagioclase and orthopyroxene phenocryst indicates that the heating is still ongoing on a scale that has remained relatively unchanged. Finally, we suggest that the dominance of normally zoned plagioclase microlites reflects decompression-driven crystallization continuing after enclave disaggregation.

The microprobe SiO$_2$/Al$_2$O$_3$ ratio of crystallizing clinopyroxene is heavily dependent on the bulk wt% SiO$_2$ content of the host magma (Kushiro, 1960); with increasing bulk rock SiO$_2$ favouring increased SiO$_2$/Al$_2$O$_3$ ratios in the cpx crystals. Thus the higher SiO$_2$/Al$_2$O$_3$ values of the Phase V microphenocrysts (Table 6c) suggests that the clinopyroxene microphenocrysts found in Phase V andesites crystallized in a more SiO$_2$-rich melt relative to the microphenocrysts from the first three phases. There are a number of other observations for the Phase V andesites such as the elevated total bulk rock alkali contents relative to Phases I, III and IV andesites (Figure 5c); lower TiO$_2$ wt% contents in the oxide microlites (Table 7a) and the slightly lower magnesium numbers of the orthopyroxene microlites (0.63) (Table 6a, Figure 5b) that are all consistent with the Phase V andesite products being more evolved.
Bulk Rock Variation

Enclave Compositions

The fundamental trends in the enclaves are related to fractional crystallization. The enclave products of Phases I and II are much more scattered compositionally (Figure 10). We interpret the scatter in the Phase I and Phase II enclaves as an indication of assimilation of plutonic residues remaining from previous mafic injections. There is a bit of scatter in the more evolved samples (>2.2 FeO/MgO) of the Phase III and Phase V enclaves in all of the plots; the linear trends observed in the enclave products of Phases III and V could be consistent with fractional crystallization, less affected by assimilation processes. Hence the minimal scatter in the Phase III and Phase V enclaves indicates they may have erupted through a relatively residue-free pathway.

Irrespective of the scatter present in the products of Phases I and II there are still discrepancies in compositions of the enclave products that need to be accounted for. The ratios of incompatible trace elements should not be affected by shallow crustal process therefore; incompatible trace element ratios should be representative of the source region. The near constant ratio of Zr/Rb in the mafic enclaves (Table 10) indicates a fairly homogeneous source composition and suggests that the observed major element differences are likely due to differentiation rather than heterogeneities in the source region. There are a number of possible processes that might cause the enclave compositions to change and to better constrain why, we address the following relevant questions.

i. Are the discrepancies purely due to the effects of contamination, thus are Phase I and Phase II enclaves more hybridized products of the same magma, parental to the products of Phases III and V?

ii. Are the enclave compositions representing localized compositions in a chemically heterogeneous basaltic system?
iii. Do the enclaves of each phase represent discrete chemically distinct batches of basalt magma from depth?

Assimilation of plutonic residue

Earthquake swarms in the century leading up to the current eruption were considered to represent the intrusion of basalt into the andesite storage area (Shepherd et al., 1971). The mafic magma interacting with the andesite will over time cool and crystallize, resulting in crystals accumulating at the base of the reservoir. Subsequent intrusions of basalt may therefore interact with any crystal mush or plutonic residue present, assimilating crystals and creating hybridized liquid compositions. Thus it is possible that the primary magma composition intruding into the andesite reservoir has not changed with time, but the compositions of the early erupted products of Phases I and II are driven by contamination from residue left from previous basaltic intrusions into the andesite reservoir. If this were the case, then the bulk chemistry for enclave products from Phases III and V should be generally more basic than the products of Phases I and II, and the bulk enclave compositions of the Phases I and II should be reproducible by addition of crystals to the bulk enclave compositions of Phases III and V.

There are a number of observations inconsistent with this model. First is that most recent enclave products of Phase V are more evolved than the earlier enclave products (Plail et al this volume). Secondly, crystal accumulation would increase the SiO$_2$ contents of the Phase I and Phase II enclave products and drive them towards the compositions of the Phase III and Phase V enclave products on the SiO$_2$ vs. FeO/MgO plot (Figure 10). Thus there is therefore no evidence to suggest that the enclave compositions of Phases III and V represent less hybridized liquids of the same magma batch that produced the enclaves of Phases I and II and we therefore reject this theory. This however indicates that there are other processes influencing the enclave compositions other than simple crystal assimilation and fractional crystallization.
Chemical heterogeneity within a magma reservoir could be characterized by vertical compositional stratification due to the relationships between magma density and chemistry, with the less evolved, denser magmas residing in the lower regions of the reservoir (e.g. Trail and Spera, 1990), as is the case in the present day reservoir of Mt Pelée (e.g. Dupuy et al., 1985, Pichavant et al., 2002). Continuous or periodic tapping of a vertically heterogeneous reservoir is thus expected to produce a compositional variation towards lesser evolved lavas as the deeper reservoir is tapped, but this is inconsistent with the more evolved nature of the Phase V enclaves (Plail et al this volume); therefore we do not accept this model for explaining the inter-phase enclave and andesite compositional variations.

Distinct chemical pulses

The nickel contents of the enclaves are all <20 ppm and they are hence not representative of primary compositions, which would be expected to have 200-300 ppm nickel (Rhodes & Dungan, 1977). The absence of modal olivine and the presence of modal plagioclase are also consistent with the parental compositions of the enclaves not being primary. Therefore the magma injected into the base of the andesite reservoir must have previously undergone differentiation, most recently in the deeper reservoir of the Soufrière Hills magma system at 10-13km (e.g. Elsworth et al., 2008; Foroozan et al., 2010; 2011). This is consistent with the fractional crystallization trends present in the Phase III and Phase V enclaves. However, the nature of basalt delivery into the base of the Soufrière Hills andesite reservoir is still a matter of discussion. A scenario of continuous recharge is proposed by Wadge et al. (2010) and Foroozan et al. (2011) due to the patterns observed in the GPS deformation signal. Contrary to this, Humphreys et al. (2009b) showed that the amphibole phenocrysts in the andesites have variations in their molar (Cl/OH) ratios and temperatures attributed to non-continuous volatile release which likely coincide with basalt recharge episodes.
If the basalt input into the base of the andesite reservoir is pulsed, then the enclaves from each eruption phase may represent chemically distinct magma pulses entering the andesite reservoir. The liquids likely represent melts from the same reservoir whose compositions are being driven by differentiation processes occurring on timescales shorter than that of the eruption. Edwards & Russell (1998) have demonstrated that AFC processes in basalts can occur on the time scale of weeks to years, a timescale which is consistent with the timescale of the first three extrusion phases and intermediate pauses. This model of periodic basalt delivery is also consistent with the observations of periodic heating Zellmer et al. (2003b), periodic volatile release Humphreys et al. (2009b) and the short time scales of hours to months between reheating, mixing and eruption of the hybrid material (e.g. Snyder, 2000; Devine et al., 2003; Humphreys et al., 2010; Devine & Rutherford, this volume). This model is thus consistent with the petrological and geochemical data and is therefore accepted as a possible cause for the basalt compositional variation.

*Enclave Amphibole*

We have demonstrated that the inter-phase variation of FeO content in enclave amphibole mirrors FeO variation in the bulk rock for the first three extrusion phases (Figure 4d, Figure 8a), and that there is a systematic inter-phase increase in the enclave amphibole Mg# (Figure 4c, Table 5b), and an inter-phase trend of increasing Mg and Al with falling Si from Phases I to III (Table 5b).

The composition of amphibole is heavily influenced by the composition of the liquid from which they crystallize. In particular, the Al content is related to crystallization pressure (Hammarstrom & Zen, 1986; Hollister et al., 1987; Johnson & Rutherford, 1989; Rutter et al., 1989; Blundy & Holland, 1990; Schmidt, 1992; Devine et al., 1998a; Ernst & Liu, 1998; Ridolfi et al., 2010). This is consistent with the experimental data of Barclay et al. (1998) which showed that higher pressure crystallization runs on the Soufrière Hills andesite products produced amphiboles with higher Al contents than the amphiboles which occurred naturally in the lavas. Variations in amphibole Al content can also be influenced by temperature and \( fO_2 \) (Bachmann & Dungan, 2002; Anderson & Smith, 1995). In general for calcic
amphiboles, increasing P and T leads to increases in MgO, Mg# and Al, while decreases in Si along with total (Fe + Mn + Ca) can also expected.

There are no independent crystallization temperature estimates for the enclave amphiboles and hence no way to assess how much influence the temperature or pressure has on the enclave amphibole Al contents. The increasing Mg#s in the enclave amphiboles from Phases I to III are consistent with the parental magmas becoming increasingly mafic from Phase I to Phase III. This would also explain the decreasing Si and increasing Al, and would be consistent with increasing crystallization temperatures. We do note however that the high Al and Ca contents coupled with high Mg#s of the Phase V enclave amphiboles (Table 5b) is inconsistent with the more evolved nature of the host magmas as indicated by Plail et al. (this volume). We also note that the lack of systematic inter-phase variation in the andesite-derived amphiboles is consistent with fairly stable crystallization conditions.

To summarize, the pattern of varying bulk rock enclave compositions is consistent with temporal variations in the composition of the parental mafic magma entering the volcanic system at depth. This can also explain the coupled changes in enclave amphibole composition, which is consistent with crystallization from an increasingly more mafic melt for the first three extrusion phases.

**Andesite Compositions**

The observed chemical heterogeneities in the Phase I andesitic products are attributed by Murphy et al. (2000) to the proposed mode of formation, which is mainly the disaggregation and remobilization of multiple previously intruded andesitic bodies. The scatter present in the Phase I andesite is similar to that observed in the Phase I enclaves; and lessening of the scatter as the eruption proceeds is also present in the andesite as well as the enclaves. The behaviour of FeO in both lava types suggests that the composition of both is coupled to some extent, but the reason for this is not clear. We suggest that the bulk andesite compositions may be dependent on the composition of the syn-erupted basalt with which it
interacts, inheriting the geochemical imprint of the mafic magma. Below we consider the possible mechanisms for transferring such a geochemical signature from the mafic magma to the andesite.

Andesite Hybridization

Hybridization requires mass transfer which has been shown to occur at the Soufrière Hills with the proposed processes ranging from simple crystal transfer, (Murphy et al., 1998; 2000) to the physical destruction of the enclaves by varying mechanisms (e.g. Humphreys et al., 2009a; Edmonds et al., this volume; Plail et al, this volume), to localized mixing, percolation and incorporation of melt into the host andesite margins (Genareau & Clarke, 2010). It must be noted that although not systematic and coupled like FeO, inter-phase variation is still observed in all of the major elements of both lava types. The elevated Ca, Al and Mg contents of the groundmass assemblage makes it apparent that Fe is being systematically transferred from the enclaves to the host andesite while the other major elements are transferred on a more irregular basis. Therefore the process/processes responsible for transferring the Fe signature from enclave to andesite must be examined.

Enclave disaggregation

Variations in the FeO contents of enclave clinopyroxene and amphibole are similar to the observed bulk rock variations, while no systematic variation in FeO is observed for any of the other enclave mineral phases. We therefore suggest that a possible explanation for the bulk rock compositional variation is the physical destruction of mafic enclaves, and incorporation of clinopyroxene and amphibole crystal fragments into the andesite. The presence of high-Al amphibole fragments in the groundmass of Phase II and Phase III andesite supports such a process.

This model was tested by performing mass balance calculations on the andesite compositions using FeO content as the index for choosing end members, which are the low FeO and high FeO samples within each eruption phase. We use the enclave mineral compositions present in each respective phase for the
calculations. If enclave destruction is responsible for the Fe signature in the andesite then the mass balance results should be consistent with the low modal occurrence of high Al amphibole fragments in the groundmass. However, the goodness of fit (reflected by $R^2$ values) for our modeled results is rather low, and the results show that the required mass of crystals to be added is inconsistent with the observed modes (Table 12). Therefore this model for Fe transfer is rejected.

**Diffusion/advection from enclaves**

Grasset & Albarede (1994), propose a mechanism whereby diffusion and buoyancy-driven convection are responsible for the chemical exchange between mafic enclaves and their hosts. They propose that the density difference between the enclave and host will induce relative motion which entrains the enclave-host interface thus generating a flow pattern inside the enclave. Thermal equilibrium occurs on much shorter time scales than chemical equilibrium via diffusion (Sparks & Marshall, 1996) thus hybridization by chemical diffusion requires both magmas be liquid and the presence of mobile fluids for transporting the chemical species.

The chilled margins and abundant interstitial voids in the enclaves (Murphy et al., 1998), suggest that the enclaves were intruded into the andesite as liquid blobs (e.g. Bacon, 1986; Clynne, 1999; Saito et al., 2003). These textures however also indicate fairly rapid heat transfer and quenching, their high vesicularity suggest that they may be permeable to gas flow and melt percolation. The small size of the enclaves and short timescales (days to months) between reheating, mixing and eruption (Snyder, 2000; Devine et al., 2003; Humphreys et al., 2010) will greatly restrict the volume of andesite that could be contaminated by diffusion or advection from enclaves, and we reject this model based on time and surface area constraints.
Diffusion/advection from basal flow

Due to the difference in density between the two magmas, basaltic injections into silicic chambers may tend to occupy the base of the reservoir. Here the intricacies of the mixing could result in significant hybridization of small volumes of silicic magma, trapped beneath the intruding mafic magma (e.g. Snyder & Tait, 1995; 1998a; 1998b). Snyder & Tait (1998b) propose that isotopic and trace element signatures can be transferred from basalt to andesite without significant effect on the major element chemistry apart from FeO. This is achieved through a combination of diffusion and advection via the large surface area of the fingered morphology created by the trapped andesite rising up through the overriding basalt due to differences in density (Snyder & Tait, 1995; 1998a; Perugini & Poli, 2005).

Contamination of the andesite will increase its density due to water loss to the basalt and the migration of FeO from the basalt to the andesite. The contaminated liquid thus forms a layer between the basalt and uncontaminated andesite, in the same region where enclave formation is thought to occur (e.g. Eichelberger, 1980; Thomas et al., 1993). The relatively higher temperature of the contaminated andesite layer should delay quenching, thus giving more time for diffusion from the upper layer of the basalt. The relatively higher density of the contaminated andesite liquid should make it easier for the basaltic liquids to physically interact with and generate vesiculated blobs as enclaves into the contaminated andesite (e.g. Thomas & Tait, 1997). Both compositions are subsequently erupted together as enclaves and host andesite. Such a process would be consistent with the FeO content of the enclaves and their host andesite lavas being coupled.

For a series of injections closely spaced in time, the process is repeated and any remnant contaminated andesite liquids are further contaminated by the new influx. Subsequent basalt intrusions would create a step like dispersal of the contaminants upwards via double diffusive convection. Rather than a rapid dispersal via convection, trace element and isotopic gradients will be set up in the andesite as a function of the number of injections encountered.
However if each injection leads to total extrusion of the contaminated liquid, the extruded andesite should have an imprint that is only controlled by the relative chemistry and volume of the basalt it encounters.

There is no robust trace element and isotopic dataset for products erupted after Phase I, so we cannot test this model conclusively. However the migration of FeO from the basalt to the andesite as predicted by the model is consistent with our observations of the coupled FeO behaviour in both magmas and this therefore seems a good way to explain the behaviour of Fe in both magmas.

We previously highlighted a steady decrease in the gap between the Fe content of the andesite and the syn-erupted enclaves as the eruption progresses. This could be due to Fe-diffusion becoming more efficient with time. A second more plausible possibility is that residual liquids are being further contaminated over time by subsequent basalt influxes, and hence more efficient mixing may be occurring as the eruption proceeds due to the converging chemistries of the contaminated residue and the intruding basalt. We interpret the closing gap in relative bulk Fe contents as an indicator of further contamination of un-erupted previously contaminated andesite. This observation is consistent with a periodic basalt delivery into the andesite reservoir rather than a continuous trickle.

Gas fluxing

It has been demonstrated that the intruding basalt is the primary source of the SO$_2$ that is emitted from the Soufrière Hills volcano (Edmonds et al., 2001). The volcano has exhibited a type of behaviour termed as persistent degassing (Shinohara, 2008). Thus there has been a constant influx of volatiles into the andesite reservoir from the intruding basalt. The SO$_2$ flux through the andesite reservoir varies on timescales ranging from weeks to years and in most cases is independent of magma extrusion (Christopher et al., 2010). There is no evidence to indicate that the fluxing of volatiles through the andesitic system has a major influence on the andesitic groundmass composition.
**Expected Future Trends and Behaviour**

The changes in magma composition from Phase III onwards, coincide with an increase in the violence of activity during the extrusion phases. To date there is no strong evidence linking eruption style to magma composition during the eruption. However we expect subsequent extrusion phases during the present eruption to produce andesite and enclaves showing coupled Fe contents with both compositions showing similar inter-phase Fe trends. The Fe content of both magma types is further expected to converge due to the continued Fe enrichment from the repeated contamination of previously contaminated un-erupted andesite residue. We also expect the products of subsequent phases to show linear trends with little scatter.

**Future Research**

The variation in FeO content observed in the enclaves indicates significant changes in the basalt composition. It would be therefore useful to constrain the nature of basalt delivery into the andesite reservoir. Several petrogenetic processes are occurring contemporaneously during this eruption such as magma mixing, fractional crystallization, mafic recharge and possibly crustal contamination; all of which influence the magma compositions. The effects of each are poorly constrained and need to be deciphered by robust analysis of the trace element and isotopic geochemistry. The current trace element dataset is sparse and thus needs supplementing. Trace element and isotope data of the crystal cargo would also help better constrain the interaction between the basalt and andesite as well as any recharge episodes that would have occurred (e.g. Tonarini et al., 1995; Browne et al., 2006; Davi et al., 2009).
Conclusions

i. Basaltic intrusion and the subsequent formation of mafic enclaves have played an integral role in eruptions at the Soufrière Hills volcano for millennia. The intruding basalt has been providing heat to the surrounding host andesite magma and hence energy for remobilizing the andesite throughout the current eruption.

ii. The early intruded basalt erupted in Phases I and II assimilated plutonic residue left over from previous basalt intrusions, while the basalt products of Phases III, IV and V encountered a relatively residue free reservoir floor.

iii. There are genuine inter-phase changes in the composition of the basalt arriving from depth, especially in the FeO content.

iv. Crystal fractionation of the basalt and subsequent mixing/mingling and diffusion produces a whole suite of hybrid magmas which are erupted as enclaves and host andesites; hence we are unable to sample the true composition of the intruded basalt or the pre intrusion andesite.

v. The andesites are likely hybridized by the basalt due to trapping of ambient andesite liquid beneath the basal intrusion which allows for diffusion and transfer of the basalt Fe signature to the andesite while it percolates upward through then resides on the basalt sheet. Transfer of elements such as Ca, Al and Mg are controlled predominantly by mixing and enclave destruction.

Acknowledgements

MCSH is supported by a Royal Society University Research Fellowship. We acknowledge assistance from Victoria Smith and Chiara Petrone during electron microprobe analysis as well as Crystal Mann for her helpful discussion. We also acknowledge the support of the British Geological Survey.
Appendix I - Possible inter-laboratory variability

The majority of bulk rock geochemical data collected and presented here for the first three phases of extrusion are from two main labs, Brown University (Joe Devine) and University of Leicester (Murphy and Sparks, 1999; Murphy et al., 2000; Zellmer et al., 2003a; MVO contract), with additional data from the labs at Bristol University, McGill University, British Geological Survey and University of East Anglia. To rule out laboratory bias as a source of the inter-phase FeO variation, we generated plots of FeO vs. SiO$_2$ for products from Phases I, II and III (Figure 11) demarcated by data source and eruption phase.

Despite some scatter, andesite analyses from a given eruptive phase overlap within error regardless of the laboratory used. For example, Phase II andesites clearly show reduced FeO contents relative to Phase I andesites while the Phase III andesites generally have lower FeO contents than Phase II andesites at similar SiO$_2$ (Figure 11a). The same pattern is true for the mafic enclaves where the Phase III products from two different labs have distinctly lower FeO contents (Figure 11b). To further investigate this, andesite and enclave splits from Phase I products were recently analyzed at the UEA lab and are included on the plot (Figure 11a & b). They show similar FeO contents to Phase I analyses from other labs (Figure 11a & b). We are therefore confident that the variations in Fe content are real and not a function of source laboratory. In addition, the correlation of FeO contents with other geochemical variables (e.g. trace elements such as Sc, Ba and V) indicates that these distinct FeO compositions are related to real differences in bulk rock compositions and not related to laboratory analysis.


Buckley V., Sparks R. & Wood B. 2006. Hornblende dehydration reactions during magma ascent at Soufrière Hills Volcano, Montserrat. Contributions to Mineralogy and Petrology 151, 121-140.


ZELLMER G. F., HAWKESWORTH C. J., SPARKS R. S. J., THOMAS L. E., HARFORD C. L., BREWER T. S. &

ZELLMER G. F., SPARKS R.S.J., HAWKESWORTH C. J. & WIENDEMECK M. 2003b. Magma Emplacement and
Remobilization Timescales Beneath Montserrat: Insights from Sr and Ba Zonation in Plagioclase
Figure captions

Figure 1: Box plots showing relative plagioclase An contents by eruption phase for andesite phenocryst rims A, andesite microlites B, sieved rims, oscillatory rims and enclave crystals C, phenocryst cores by texture D. Microlite Fe content vs An content by phase E, Fe content vs An content for sieve and enclave crystals F.

Figure 2: Typical andesite whole rock texture found in Phases IV and V. Pl – plagioclase, Px – orthopyroxene, Cpx – clinopyroxene, Ox – Fe-Ti oxide.

Figure 3: Examples of enclave textures described by Plail et al (this volume). Pl – plagioclase, Px – clinopyroxene, Hbl – hornblende, Ox – Fe-Ti oxides, V – vesicles.

Figure 4: Amphibole compositions in andesite and enclaves by phase A, box plots for mg#'s of amphibole phenocryst rims in the andesite B, enclaves by phase and mg# C, enclaves by phase and Fe content D. an- andesite, en- enclaves. The number of analyses used is given in each box, horizontal line represents median values.

Figure 5: Box plots of mg# for andesite orthopyroxene unzoned rims (by phase) and rev-zoned rims A, andesite orthopyroxene microlites (by phase) and enclave microlites B, bulk rock total alkali by eruption phase C, andesite hosted clinopyroxene microlite Kd values by phase D. The number of analyses used is given in each box, horizontal line represents median values. an – andesites, en – enclaves.

Figure 6: Abundances of major elements in andesite and enclave glass from microprobe analyses, andesites- An, enclaves- En. High K2O > 3wt%, low K2O < 3wt%.

Figure 7: Bulk andesite and enclave major element abundances by phase, vertical dashed line demarcates boundary between silica content of the andesite- An, enclave- En.

Figure 8: Box plots showing inter-phase bulk rock contents of FeO, V, Ba and plagioclase microlite Fe content throughout the eruption. The number of analyses used is given in each box, horizontal line represents median values, mic- microlite, an – andesites, en- enclaves.
Figure 9 Selected trace element plots for bulk rock andesite and enclave products. Symbols and labels are the same as Figure 7, vertical line demarcates boundary between andesite and enclave silica content.

Figure 10 Major element behaviour with changing FeO/MgO showing changes as function of crystal control. Arrows indicate removal of particular mineral phase. An- andesite, En – enclaves. Fractionation and assimilation vectors for the other oxides are same as in $\text{Al}_2\text{O}_3$.

Figure 11 Harker plots of $\text{SiO}_2$ vs FeO for phases 1, II and III by lab showing an absence of any systematic inter-lab variation in the data for andesites- A and enclaves-B. L-leicester, JD- Joe Devine, SS- (Murphy et al., 1998), BGS- British geological survey, UEA- University of East Anglia, MGU- Mc Gill University.

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Table 1 Laboratory and sample information for the XRF data used in this study; An-andesite, En- enclave.

Table 2 Published modal mineral contents for the first three extrusion phases. Arbitrary distinction between microlites and microphenocryst defined by 100$\mu$m, *Recalculated from point counts in Humphreys et al., 2009a, range shown.

Table 3 Mineral phases and associated textures in the andesite and enclave lavas of the current eruption.

Table 4 Averaged andesite plagioclase phenocryst and microlite microprobe data, comparing compositions from Phases IV and V with the published compositions of Phases I, II and III. pc-phenocryst core, pr - phenocryst rim, mc- microlite core, mr- microlite rim, mpc- microphenocryst core, mpr – microphenocryst rim, An- andesite, En – enclave.

Table 5a Averaged andesite amphibole compositions by texture and eruption phase.

Table 5b Averaged enclave amphibole compositions from eruption Phases I, II, III and V along side high aluminum andesite hosted microphenocrysts and groundmass fragments. Mphx – microphenocrysts, Frag – groundmass fragments. Phase V data from Plail et al. (this volume)
Table 6a Averaged Orthopyroxene phenocryst and microlite compositions comparing the andesite crystals of phase IV and V with crystals from phases I, II and III, errors are ± 1σ, mic – microlites, phx-phenocrysts.

Table 6b Averaged compositions of andesite hosted reverse zoned orthopyroxene phenocryst rims along with andesite and enclave microphenocrysts, errors are ± 1σ, Mphx- microphenocrysts, Phx- phenocrysts

Table 6c Averaged compositions of clinopyroxene microphenocrysts, microlites and overgrowths, errors are ± 1σ, mic – microlites, mphx- microphenocrysts.

Table 7a Averaged andesite titanomagnetite compositions, sorted by eruptive phase and textural type. Usp/Ilm mol% after Stormer (1983), mic – microlites.

Table 7b Averaged compositions of enclave oxides along with andesite derived ilmenite.

Table 8 Averaged microprobe glass compositions for andesites and enclaves, along with log fO2 values from touching ilmenite–magnetite pairs.

Table 9 Averaged bulk rock major element compositions for the andesite and enclaves of the different eruption phases.

Table 10 Averaged bulk rock trace element for andesites and enclaves of the different eruption phases.

Table 11 Temperature and Pressure estimates for the eruptive products obtained using a number of different techniques.
Table 12 Intra-phase andesite mass balance calculations for addition of enclave crystals, weight % values are relative to initial magma compositions.
Figure 1 Box plots showing relative plagioclase An contents by eruption phase for andesite phenocryst rims A, andesite microlites B, sieved rims, oscillatory rims and enclave crystals C, phenocryst cores by texture D. Microlite Fe content vs An content by phase E, Fe content vs An content for sieve and enclave crystals F.
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a: diktytaxitic plag-pargasitic amphibolite
b: diktytaxitic plag-cpx
c: composite (hand specimen)
Figure 4 Amphibole compositions in andesite and enclaves by phase A, box plots for mg#s of amphibole phenocryst rims in the andesite B, enclaves by phase and mg# C, enclaves by phase and Fe content D. The number of analyses used is given in each box, horizontal line represents median values.
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Figure 7 Bulk andesite and enclave major element abundances by phase, vertical dashed line demarcates boundary between silica content of the andesite- An, enclave- En.
<table>
<thead>
<tr>
<th>Extrusion Phase/lava</th>
<th>PI an</th>
<th>PII an</th>
<th>PIII an</th>
<th>PIV an</th>
<th>PV an</th>
<th>PI en</th>
<th>PII en</th>
<th>PIII en</th>
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<td>8</td>
<td>9</td>
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<td>V ppm</td>
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<td>Ba ppm</td>
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<td>100</td>
<td>150</td>
<td>200</td>
<td>250</td>
<td>300</td>
<td>350</td>
<td>400</td>
<td></td>
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Figure 8 Box plots showing inter-phase bulk rock contents of FeO, V, Ba and plagioclase microlite Fe content throughout the eruption. The number of analyses used is given in each box, horizontal line represents median values, mic- microlite, an – andesites, en- enclaves.
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<table>
<thead>
<tr>
<th>Lab/Location</th>
<th>Instrument</th>
<th>Samples analyzed by Lab</th>
<th>Phase</th>
<th>Total Number of samples</th>
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<tr>
<td>University of East Anglia</td>
<td>Brucker-AXS S4 Pioneer FluorescenceS spectrometer</td>
<td>An: 1523,1524, M003, M010 En: M001&amp; M002, M004 to M009, M011 to M019</td>
<td>III</td>
<td>An - 4 En - 17</td>
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<td></td>
<td></td>
<td>An: 1532,1535</td>
<td>IV</td>
<td>An - 7</td>
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<td></td>
<td></td>
<td>An: 1537 to 1544</td>
<td>V</td>
<td>An - 12</td>
</tr>
<tr>
<td>McGill University, Canada</td>
<td>Philips PW2440 4 kW automated XRF</td>
<td>An: CM 36 to 38</td>
<td>I</td>
<td>En - 3</td>
</tr>
<tr>
<td>BGS labs</td>
<td>Spectron X-LAB 2000 ED-XRF</td>
<td>An: 1247/AND (1 to 3)</td>
<td>II</td>
<td>An – 3 En - 9</td>
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<tr>
<td>University of Leicester</td>
<td>PANalytical Axios Advanced PW4400 XRF</td>
<td>An: 40,45 to 57,174 to 242,244 to 332,665 to 725,1003 to 1091</td>
<td>I</td>
<td>An – 46 En - 11</td>
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<td>An: 1325 (a &amp; b), 1326 (a &amp; b)</td>
<td>III</td>
<td>An – 9 En 4</td>
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Table 1 Laboratory and sample information for the XRF data used in this study; An-andesite, En-enclave.
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<th>Opx</th>
<th>Ti-mag</th>
<th>Qtz</th>
<th>cpx</th>
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<tr>
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<td>30-35</td>
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<td>2-5</td>
<td>2-4</td>
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<tr>
<td>Devine et al, 1998 Phase I</td>
<td>~40</td>
<td>~30</td>
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<td>Humphreys et al., 2009a*, Phase III</td>
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<td>20-32</td>
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<td>0.05-0.4</td>
<td>0.5-0.66</td>
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Table 2 Published modal mineral contents for the first three extrusion phases. Arbitrary distinction between microlites and microphenocryst defined by 100µm, *Recalculated from point counts in Humphreys et al., 2009a, range shown.
<table>
<thead>
<tr>
<th>Mineral Phase</th>
<th>Crystal Textures Present</th>
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<td><strong>Andesite</strong></td>
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<tr>
<td>Plagioclase</td>
<td>subhedral – euhedral phenocrysts, zoning (sieved, normal, reverse, patchy and oscillatory) normally zoned microlites euhedral, hopper shaped or skeletal</td>
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<td>Amphibole</td>
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Table 3 Mineral phases and associated textures in the andesite and enclave lavas of the current eruption.
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Table 4 Averaged andesite plagioclase phenocryst and microlite microprobe data, comparing compositions from Phases IV and V with the published compositions of Phases I, II and III. pc- phenocryst core, pr - phenocryst rim, mc- microlite core, mr- microlite rim, mpc- microphenocryst core, mpr – microphenocryst rim, An- andesite, En – enclave.
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Table 5a Averaged andesite amphibole compositions by texture and eruption phase.
Table 5b Averaged enclave amphibole compositions from eruption Phases I, II, III and V along side high aluminum andesite hosted microphenocrysts and groundmass fragments. Mphx – microphenocrysts, Frag – groundmass fragments. Phase V data from Plail et al. (this volume)

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Table 6a: Averaged Orthopyroxene phenocryst and microlite compositions comparing the andesite crystals of phase IV and V with crystals from phases I, II and III, errors are ± 1σ, mic – microlites, phx – phenocrysts.
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<td>1.08</td>
</tr>
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<td>1.5</td>
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<td>0.7 ± 0.02</td>
<td>0.64 ± 0.05</td>
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Table 6b: Averaged compositions of andesite hosted reverse zoned orthopyroxene phenocryst rims along with andesite and enclave microphenocrysts, errors are ± 1σ. Mphx-microphenocrysts, Phx-phenocrysts.
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<th>P5 Andesite mphx</th>
<th>P1-P3 Andesite mic</th>
<th>P4 Andesite mic</th>
<th>P5 Andesite mic</th>
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<td>0.27</td>
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<td>98.97</td>
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<td>99.17</td>
<td>98.84</td>
<td>99.21</td>
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<tr>
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<td>32.43</td>
<td>17.78</td>
<td>16.68</td>
<td>17.47</td>
<td>13.92</td>
<td>15.49</td>
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<tr>
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<td>42.8</td>
<td>40.75</td>
<td>42.56</td>
<td>42.73</td>
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Table 6c Averaged compositions of clinopyroxene microphenocrysts, microlites and overgrowths, errors are ± 1σ, mic – microlites, mphx- microphenocrysts.
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<tr>
<th></th>
<th>P1-P3 mph cores</th>
<th>P4 mph cores</th>
<th>P5 mph cores</th>
<th>P1-P3 mph rims</th>
<th>P4 mph rims</th>
<th>P5 mph rims</th>
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<th>P4 mic</th>
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<td>0.11</td>
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<td>1.97</td>
<td>2.19</td>
<td>1.76</td>
<td>1.72</td>
<td>2.02</td>
<td>1.82</td>
<td>1.83</td>
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<td>81.17</td>
<td>82.27</td>
<td>80.94</td>
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<td>80.6</td>
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<td>0.6</td>
<td>0.62</td>
<td>0.59</td>
<td>0.62</td>
<td>0.67</td>
<td>0.63</td>
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<tr>
<td>MgO</td>
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<td>1.32</td>
<td>1.18</td>
<td>1.16</td>
<td>1.39</td>
<td>1.26</td>
<td>1.15</td>
<td>1.38</td>
<td>1.08</td>
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<tr>
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<td>0.03</td>
<td>0.04</td>
<td>0.04</td>
<td>0.16</td>
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<td>0.09</td>
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<td>94.46</td>
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<td><strong>25</strong></td>
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<td>± 1σ</td>
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<td>6</td>
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<td>4</td>
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<td>9</td>
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Table 7a Averaged andesite titanomagnetite compositions, sorted by eruptive phase and textural type. Usp/IIm mol% after Stormer (1983), mic – microlites.
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<td>1.26</td>
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<td>2.04</td>
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<td>0.05</td>
<td>0.05</td>
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<td>96.68</td>
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<td>83</td>
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<td>8</td>
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<td>6</td>
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Table 7b Averaged compositions of enclave oxides along with andesite hosted ilmenite.
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<th>SiO2</th>
<th>TiO2</th>
<th>Al2O3</th>
<th>FeO</th>
<th>MgO</th>
<th>CaO</th>
<th>Na2O</th>
<th>K2O</th>
<th>Total</th>
<th>Mg#</th>
<th>Fe/Mg</th>
<th>Ca/Al</th>
<th>Tot alkali</th>
<th>fO2 log units</th>
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<td>1.3</td>
<td>0.5</td>
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<td>1</td>
<td>0.6</td>
<td>1.4</td>
<td>1.1</td>
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<td>7.56</td>
<td>NNO + 0.5- +1.3</td>
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<tr>
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<td>0.45</td>
<td>0.5</td>
<td>0.09</td>
<td>0.6</td>
<td>0.7</td>
<td>1.5</td>
<td>1.2</td>
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<td>1.1</td>
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<td>3.85</td>
<td>98.9</td>
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<td>7.7</td>
<td>NNO+ 0.5- +1.4</td>
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<tr>
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<td>1.45</td>
<td>0.6</td>
<td>0.14</td>
<td>0.9</td>
<td>0.7</td>
<td>1.3</td>
<td>1.3</td>
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<td>99.8</td>
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<td>7</td>
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<td>0.08</td>
<td>1.11</td>
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<td>3.1</td>
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<td>0.08</td>
<td>0.03</td>
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Table 8 Averaged microprobe glass compositions for andesites and enclaves, along with log fO2 values from touching ilmenite–magnetite pairs.
<table>
<thead>
<tr>
<th>Andesite</th>
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<th>TiO₂</th>
<th>Al₂O₃</th>
<th>FeO</th>
<th>MnO</th>
<th>MgO</th>
<th>CaO</th>
<th>Na₂O</th>
<th>K₂O</th>
<th>P₂O₅</th>
<th>Tot</th>
<th>Fe/Mg</th>
<th>Ca/Al</th>
<th>Tot alk</th>
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<td>0.61</td>
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<td>± σ</td>
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<td>0.36</td>
<td>0.39</td>
<td>0.02</td>
<td>0.19</td>
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<td>0.18</td>
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<td>0.1</td>
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<td>0.63</td>
<td>17.86</td>
<td>7.14</td>
<td>0.18</td>
<td>2.81</td>
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Table 9 Averaged bulk rock major element compositions for the andesite and enclaves of the different eruption phases.
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Table 10 Averaged bulk rock trace element for andesites and enclaves of the different eruption phases.
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Table 11 Temperature and Pressure estimates for the eruptive products obtained using a number of different techniques.
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Table 12 Intra-phase andesite mass balance calculations for addition of enclave crystals, weight % values are relative to initial magma compositions.