

SCIENTIFIC REPORTS



OPEN

The 1257 Samalas eruption (Lombok, Indonesia): the single greatest stratospheric gas release of the Common Era

Received: 10 June 2016
Accepted: 20 September 2016
Published: 10 October 2016

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Large explosive eruptions inject volcanic gases and fine ash to stratospheric altitudes, contributing to global cooling at the Earth's surface and occasionally to ozone depletion. The modelling of the climate response to these strong injections of volatiles commonly relies on ice-core records of volcanic sulphate aerosols. Here we use an independent geochemical approach which demonstrates that the great 1257 eruption of Samalas (Lombok, Indonesia) released enough sulphur and halogen gases into the stratosphere to produce the reported global cooling during the second half of the 13th century, as well as potential substantial ozone destruction. Major, trace and volatile element compositions of eruptive products recording the magmatic differentiation processes leading to the 1257 eruption indicate that Mt Samalas released 158 ± 12 Tg of sulphur dioxide, 227 ± 18 Tg of chlorine and a maximum of 1.3 ± 0.3 Tg of bromine. These emissions stand as the greatest volcanogenic gas injection of the Common Era. Our findings not only provide robust constraints for the modelling of the combined impact of sulphur and halogens on stratosphere chemistry of the largest eruption of the last millennium, but also develop a methodology to better quantify the degassing budgets of explosive eruptions of all magnitudes.

Sulphur (S) gases injected into the stratosphere during plinian eruptions are converted into sulphate aerosols that travel around the globe and backscatter solar radiation, resulting in a net cooling of the troposphere and the Earth's surface¹. Besides the effect of sulphate aerosols, volcanogenic halogens and especially chlorine (Cl) and bromine (Br) may induce the catalytic destruction of the ozone layer². Although these processes have been observed on a very small scale for a number of recent explosive eruptions such as the 1963 Agung, 1982 El Chichón and 1991 Pinatubo events³, the effects of large historic silicic eruptions on climate and atmospheric chemistry remain poorly constrained, with hitherto unexplored feedbacks. The reconstruction of climate forcing associated with explosive eruptions involves several scientific disciplines, as it strongly depends on a variety of parameters including the height of injection, the gas flux and the latitude of the volcanic source. For past plinian eruptions suspected to have triggered or enhanced climate cooling, temperature reconstructions are commonly based on the amount of associated sulphate emissions inferred from glaciological records. It is furthermore commonly admitted that the presence of a pre-eruptive vapour phase renders the quantification of the emissions of sulphate and other volatile species of past eruptions particularly challenging^{4–6}.

The 1257 eruption of Mt Samalas, a part of the Rinjani volcanic complex (Fig. 1) on Lombok Island (Indonesia), has been recognized as the “mystery eruption”⁷ associated with the largest sulphate spike of the last 2.3 ky recorded in cores from both Arctic and Antarctic ice sheets⁸. This continuous four-phase eruption evacuated 40 ± 3 km³ of trachydacitic magma during tens of hours, producing plinian plumes that rose up to 43 km in the stratosphere and tephra fingerprinted up to 660 km from the source, thus standing as the most

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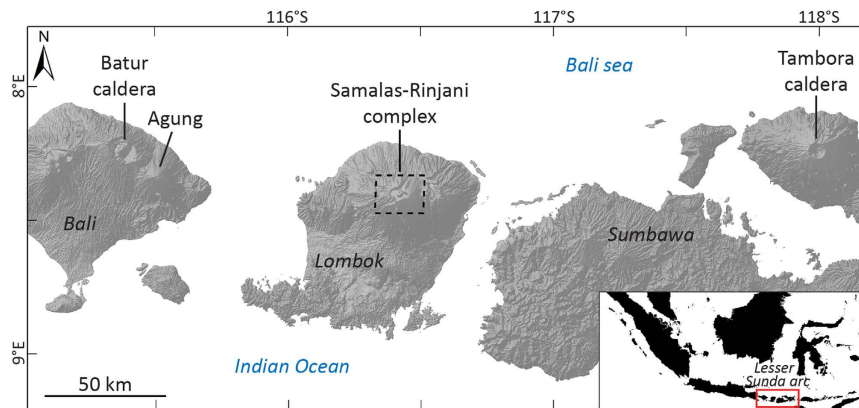


Figure 1. Map of the Lesser Sunda Islands and their active volcanoes. SRTM DEM at 3 arcsecond (~90 m) resolution (<http://srtm.csi.cgiar.org>)⁶³ of Bali, Lombok and Sumbawa. This map was generated using the Esri ArcMap 10.1 software (<http://www.esri.com>).

powerful eruption of the last millennium⁹. Archaeologists recently determined a date of 1258 for mass burial of thousands of medieval skeletons in London¹⁰, that could be linked in some respect to climatic perturbations in the Northern Hemisphere by the 1257 Samalas eruption. Indeed, medieval chronicles in Northern Europe⁷ document the occurrence of initial warming in the early winter of 1258 just following the eruption, that was followed by extensive wet and cold climatic conditions in 1259 that may have impacted crops and contributed to the onset and magnitude of famines at that time for some regions of the Northern Hemisphere. The 1257 Samalas eruption might also have contributed to the onset of the Little Ice Age¹¹.

The climate-forcing associated with the 1258–1259 sulphate anomaly in ice-cores has been the subject of intense research^{12–15}, suggesting a range of sulphate yields until the source of the eruption was found. The recent re-evaluation of the record of volcanic sulphate deposition based on an extensive array of Antarctic ice cores suggested that a total of 170 Mt of SO₂ (85 Tg of S) would have been released to produce the observed sulphate spikes, greater than the 45 Mt of SO₂ estimated for the 1815 Tambora eruption¹⁶. Temperature reconstructions based on tree-ring proxies and ice-core records showed that the emission of 96 to 138 Tg of SO₂, the most probable scenarios, would have induced an extra-tropical summer cooling over land of –0.6 °C to –5.6 °C during a period of 4–5 years¹⁷. The discrepancies between these estimates derived from ice-cores data and modelling reflect the complexity of climate reconstructions based on hypothetical S yields based on distal proxies.

The quantification of volcanogenic volatile emissions at the source commonly relies on the residual amount of volatiles dissolved in silicate melts prior to eruption recorded in small droplets of magma trapped in crystals (melt inclusions) of the erupted products¹⁸. Here we propose an alternative approach based on melt inclusions recording the different steps of magma evolution, including melt inclusions representative of both the magma that fed the eruption (a three-phased mixture of melt, gas and crystals) and of the residual melt.

Pre-eruptive magmatic conditions

The detailed petrology and mineralogy of the Holocene lavas and scoria fallouts of the Rinjani volcanic complex, including the magmatic processes leading to the 1257 Samalas eruption, are presented by Métrich *et al.* (in review), and are briefly summarised hereafter. The 1257 eruption evacuated a chemically homogeneous magma body of trachydacite (64.0 ± 0.4 wt% of SiO₂; 8.1 ± 0.1 wt% of Na₂O + K₂O, volatile free) that derived from a parent high alumina basalt magma through fractional crystallization. The trachydacitic magma crystallised and degassed at temperatures of 900 °C to 980 °C. Its mineral paragenesis typically consists of plagioclase showing a bi-modal distribution with patchy zoned cores (An_{82–75}) surrounded by bands (An_{50–46}), in association with amphibole (magnesian-hastingsite), orthopyroxene (clinopyroxene), rare clinopyroxene (augite), titanomagnetite, iron sulphide and apatite.

Here we combined the detailed geochemical study of minerals and melt inclusions with matrix glass and whole-rock analyses (major, trace, S, Cl and Br) to characterise the composition of the pre- and post-eruptive magmatic system, and further quantify the volatile budget of the 1257 eruption. Samples from the four phases of the 1257 trachydacitic eruption and of the 712 A.D. basaltic scoria fallout have been examined (see Supplementary Sample description and Figure S1). Our dataset is complemented by melt inclusions from the trachydacitic pumice of the 2550 B.P. eruption¹⁹ (compositions are reported in Supplementary Tables S1, S2 and S3). Major and trace elements denote the bi-modality of the magma compositions and the lack of intermediate andesitic magma (Fig. 2a,b). The overall evolutionary trend is perfectly illustrated by the positive correlation between Rb and Th (Fig. 2b), suggesting that the 1257 trachydacite derived from the basalt through fractional crystallization of ~81% (Methods). Furthermore, compositions of olivine-hosted melt inclusions match that of the whole-rock parent basalt of the 1257 trachydacite (Fig. 2b). Most remarkable are the melt inclusions in plagioclase An_{82–75} that are representative of the whole trachydacitic magma (60.5 ± 1.3 wt% of SiO₂; 9.0 ± 0.9 ppm of Th) whereas more sodic plagioclase (An_{50–46}), amphibole and pyroxenes trapped melts that recorded the evolution towards the residual trachydacitic composition of the matrix glass (68 ± 1 wt% of SiO₂; 16.3 ppm of Th) due to *in-situ* crystallization at shallow depth. Depicting the magma evolution respectively associated with fractional crystallization and *in-situ*

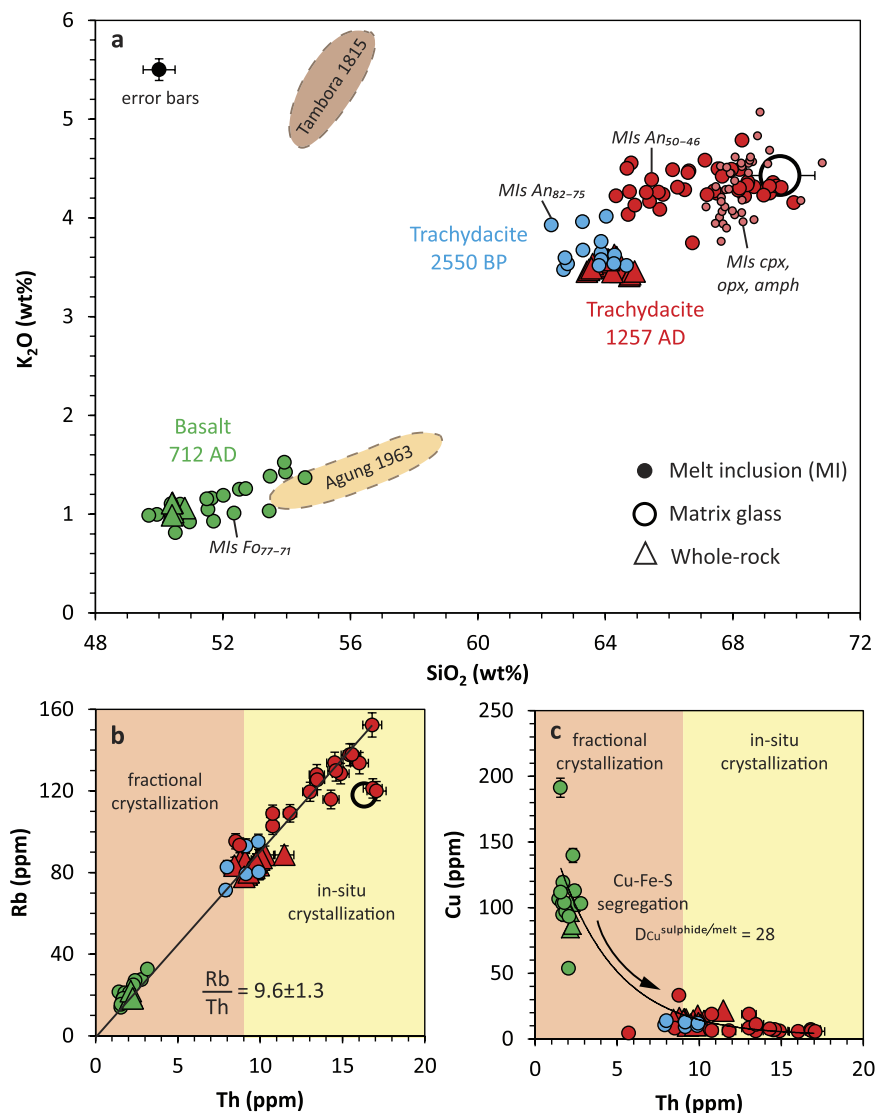


Figure 2. Melt inclusions record the magma evolution leading to the 1257 eruption. (a) K_2O vs SiO_2 variation diagram for the Rinjani-Samalas calc-alkaline suite including whole-rocks, melt inclusions in olivine of the 712 high alumina basalt (Fo is olivine forsterite content, i.e. $100 \times Mg/(Mg + Fe)$), in plagioclase of 2550 B.P. and 1257 pumice clasts (An is anorthite content, i.e. $100 \times Ca/(Ca + Na)$), in clino-(cpx), ortho-(opx) pyroxene, amphibole, and matrix glasses of the 1257 eruptive products. Compositions are normalised to 100 wt%, free of volatiles. The Rinjani-Samalas suite plots in between the whole-rock compositional fields of 1815 Tambora⁵³ and 1963 Agung^{64,65} products, highlighting the enrichment in K_2O of magmas towards the East of the Lesser Sunda arc. (b) Rb-Th positive correlation indicates that the 1257 trachydacite derived from its parent basaltic magma through a dominant process of fractional crystallization. Plagioclase An_{82-75} -hosted melt inclusions are representative of the 1257 whole magma composition, whereas plagioclase An_{50-46} -hosted melt inclusions record its shallow depth *in-situ* crystallization. (c) Cu vs Th variation diagram showing strong Cu fractionation during magma differentiation recording the prevalent Cu-sulphide segregation. See Supplementary Figures S2 and S3 for more major and trace element variation diagrams.

crystallization is an innovative approach that enables to decipher the volatile evolution from the parent basalt to the 1257 trachydacite. We further quantify the distribution of sulphur and halogens between the pre-eruptive silicate melt (melt inclusions), the volatile-bearing mineral phases (sulphide (S), apatite (SO_2 , Cl) and amphibole (Cl)), and the vapour phase (Fig. 3) consisting of gas species exsolved during *in-situ* crystallization of the trachydacite in the shallow crust, with possible incremental influx of deeper-derived vapour phase²⁰. Finally we establish the degassing budget of the 1257 Samalas eruption by comparing the pre-eruption and the post-eruption trachydacitic magma. We stress that volatile partitioning between the vapour phase and such sulphide-saturated calc-alkaline melts at 900–980 °C is still poorly constrained by experimental data.

The evolution of volatile species illustrated in Fig. 4 reflects the differential ability of each species to be fractionated into the vapour phase and minerals. Water contents of melt inclusions remain relatively constant along the path of magma differentiation (Fig. 4a,b). The basalt displays average water contents of 3.5 ± 0.2 wt%, a typical

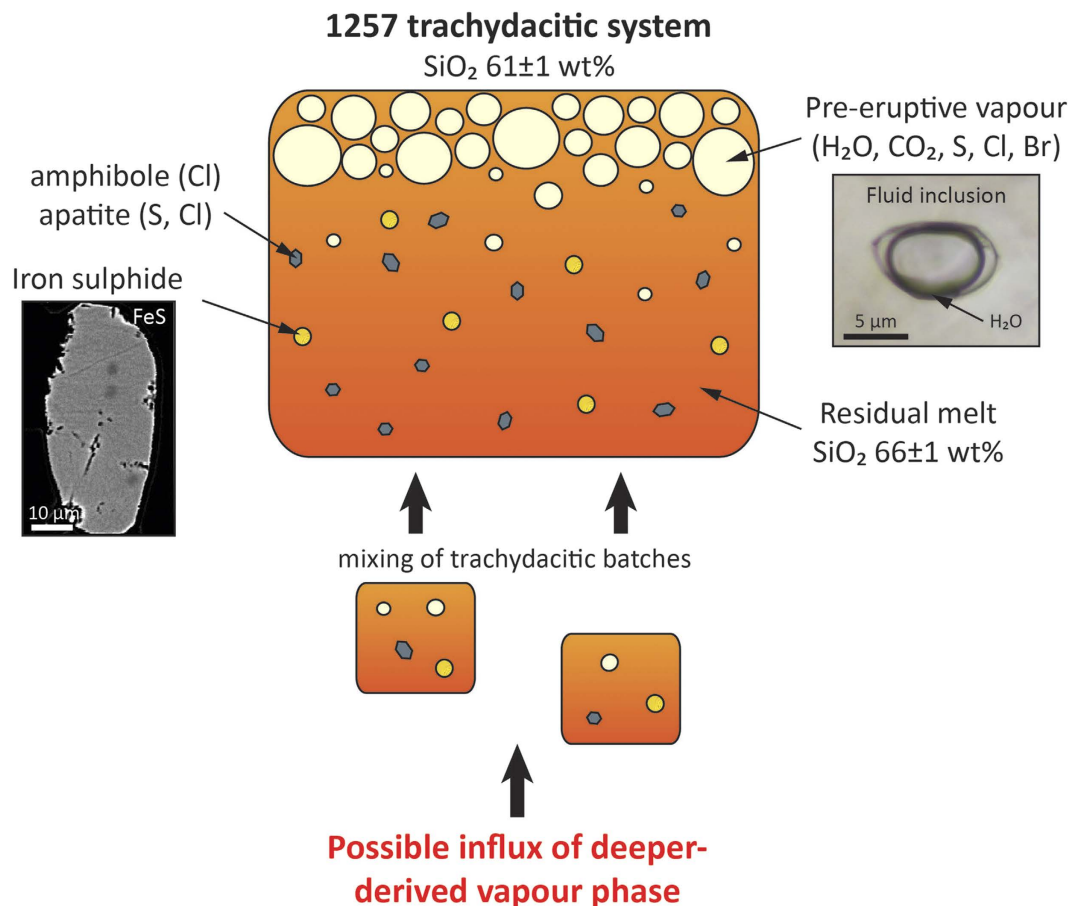


Figure 3. Volatile repositories in the 1257 trachydacitic system. Composition of the residual melt is recorded by melt inclusions trapped in cpx, opx and amphibole whereas that of the whole magma (melt + mineral phases + pre-eruptive vapour) is preserved by melt inclusions in plagioclase An₈₂₋₇₅. The existence of pre-eruptive vapour is illustrated by the occurrence of water-rich fluid inclusions in minerals (Supplementary Figure S5). CO₂ is very likely present in pre-eruptive vapour. The whole system is likely the result of the mixing of trachydacitic magma batches displaying distinct volatile contents (Supplementary Figure S4f).

value for arc basalts²¹. The 1257 trachydacite magma derived from the basalt contains 3.9 ± 0.1 wt% of H₂O, a value measured in melt inclusions in plagioclase An₈₂₋₇₅. Such a dissolved amount of H₂O is well below the maximum water concentrations (8–<10 wt%) of amphibole-bearing dacite crystallizing at 300–400 MPa under water-saturated conditions²². This suggests a drastic loss of water during fractional crystallization and most likely during magma decompression and ascent. The distribution of the H₂O concentrations in melt inclusions in plagioclase An₅₀₋₄₆ indicates that ~70% cluster at 3.7 ± 0.3 wt% H₂O (Supplementary Figure S4a), a value representative of the H₂O concentration of the residual melt (Fig. 4b). A few higher water contents (up to 5.5 wt%) reflect the multi-stage exsolution of water from the melt, and possibly magma mixing during refilling of the 1257 trachydacitic system. This feature is consistent with plagioclase textures and the temperature gradient (900–980 °C) calculated from thermometers (Métrich *et al.*, in review). Fluid inclusions fully support the presence of a free water-rich vapour phase (Supplementary Figure S5a,b). No dissolved or exsolved CO₂ was detected, even though some fluid inclusions show evidence of carbon bearing-complexes (Supplementary Figure S5e). Considering that the trachydacitic system contains an initial amount of 3.9 ± 0.1 wt% of dissolved H₂O, we estimate a proportion of exsolved water of 3.5 wt% during *in-situ* crystallization of the trachydacite (Fig. 4a). The water content of the residual melt corresponds to a fluid pressure ($P_{H_2O} = P_{total}$) of 90–120 MPa under equilibrium conditions, suggesting an upper limit of magma storage of 3.3–4.4 km assuming a crustal density of 2.8 g/cm³²³. Hence, the 1257 Samalas eruption was sustained by a trachydacite reservoir developed in the upper crust similarly to other large plinian eruptions²⁴.

Sulphur concentrations in melt inclusions reflect a more complex behaviour (Fig. 4c,d). The parent basalt displays an average S content of 1940 ± 90 ppm, a value commonly measured in moderately oxidized, hydrous, arc basalts²⁰. Poly-metallic sulphides (Cu-Fe-S) partitioning Cu (Fig. 2c) do not occur in S-rich melt inclusions but are ubiquitous in partly degassed glass embayments. The trachydacitic system displays a total S content of 870 ± 20 ppm distributed between the melt, the S-bearing minerals and the free gas phase. Sulphur concentrations in melt inclusions of clinopyroxene (150 ± 20 ppm), orthopyroxene (180 ± 45 ppm) and amphibole (185 ± 45 ppm) are similar to that of the most evolved plagioclase-hosted melt inclusions (Fig. 4d). Matrix glasses

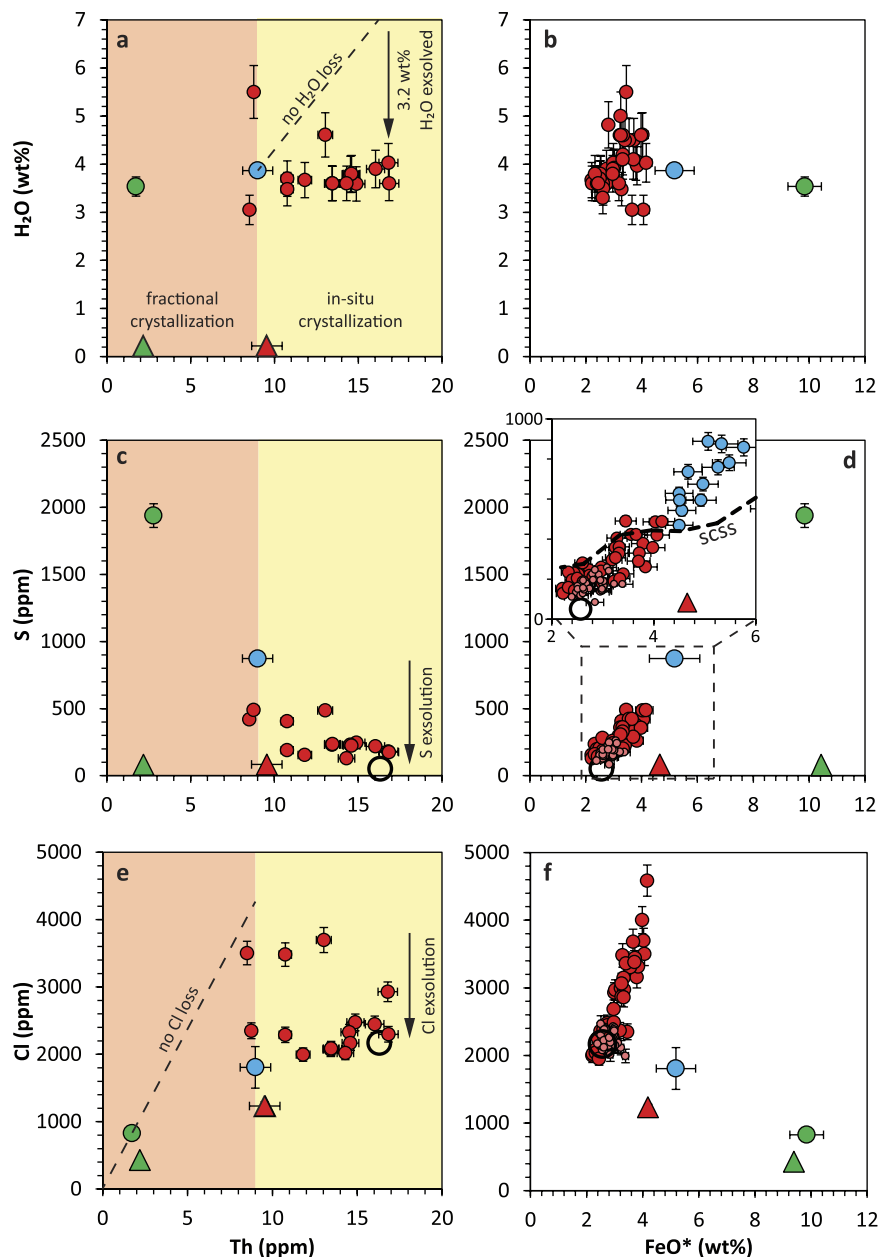


Figure 4. Melt inclusions record the volatile evolution through magma differentiation and *in-situ* crystallization. (a) H_2O vs Th and (b) H_2O vs FeO^* contents in melt inclusions reflect water exsolution through magma differentiation. Reported average H_2O content of basalt are calculated considering melt inclusions unaffected by H^+ diffusion through host mineral (Supplementary Figure S6b). (c) S vs Th variation diagram in melt inclusions. (d) Positive correlation of S and FeO^* in the 1257 melt inclusions is consistent with sulphide saturation (SCSS, dashed curve) was calculated with model B of Fortin *et al.*²⁵. (e) Cl vs Th contents in melt inclusions and whole-rocks suggest that Cl has an incompatible behaviour during basalt differentiation. (f) Cl vs FeO^* contents in melt inclusions, whole-rocks and matrix glasses reflect mixing between trachydacitic melts and Cl exsolution during *in-situ* crystallization of the 1257 trachydacite. The process of mixing is also demonstrated by the positive correlation of S and Cl (Supplementary Figure S4f). Symbols as in Fig. 2. Average initial volatile contents of melt inclusions representative of the basaltic magma and of the whole 1257 system as well as whole-rock compositions are reported with error bars (1σ).

are the most degassed endmember with 50 ± 20 ppm of S (Supplementary Table S2). The positive correlation of S and FeO (Fig. 4d) corresponds to the S concentration at sulfide saturation (SCSS)²⁵, in agreement with the occurrence of iron sulphide globules (up to $60 \mu\text{m}$ in size) entrapped with apatite in phenocrysts (mostly amphibole). However, the extent of FeO content range reflects the possible effect of magma mixing between melts displaying distinct S contents. The composition of iron sulphides (Supplementary Table S4) enabled to estimate a proportion of 0.01% of Fe-sulphides in the system (Methods), suggesting that they removed a negligible amount of S (~ 50 ppm) from the melt. Furthermore, the range of S concentrations in apatite (0.01–0.26 wt% of S,

	Th	S	Cl	Br
Concentrations (ppm)				
Undegassed residual melt $C_{undegassed\ RM}^{RM}$ (MIs cpx, opx, amph)		170 ± 50	2235 ± 155	
Degassed residual melt $C_{degassed\ RM}^{RM}$ (Matrix glass)	16.3	50 ± 20	2170 ± 110	
Pre-eruptive trachydacite $C_{pre-eruptive}^{RM}$ (MIs plagioclase ~An ₇₆)	9.0 ± 0.9	870 ± 20	3500 ± 175	
Trachydacitic whole-rock C^{WR}	9.5 ± 0.9	80 ± 20	1230 ± 45	2.92 ± 0.08
Parent basalt C^{basalt} (MIs olivine ~Fo ₇₆)	1.7 ± 0.2	1940 ± 90	830 ± 80	3.1 ± 0.4
Basaltic whole-rock (712 AD scoria)	2.20 ± 0.07	39 ± 1	428	1.6 ± 0.2
Classic petrologic method				
Volatile <i>i</i> loss $\Delta i = C_{undegassed\ RM}^{RM} - C_{degassed\ RM}^{RM} \times X^{RM}$ (ppm)		140 ± 50	960 ± 190	
Minimum mass released (Tg)		14 ± 5	96 ± 19	
New approach: whole trachydacitic system				
Volatile <i>i</i> loss $\Delta i = C_{pre-eruptive}^{RM} - C^{WR}$ (ppm)		790 ± 30	2270 ± 180	
Mass released (Tg)		79 ± 6	227 ± 18	
Contribution of the parent basalt				
Theoretical amount of volatile available C^{system} (ppm)		10090 ± 1480	4300 ± 720	16 ± 3
$S^{Cu-Fe-S}$ (ppm)		690 ± 225		
S^{fluid} (ppm)		8500 ± 1500		
m_S^{fluid} assuming 1.5 wt% fluid (Tg)		13 ± 6		
Volatile <i>i</i> loss $\Delta i = C^{system} - C^{WR}$ (ppm)				13 ± 3
Maximum mass released (Tg)		92 ± 8		1.3 ± 0.3
Global atmospheric mixing ratios (ppbv of Cl and pptv of Br)			36	90
Stratospheric emissions (Tg)		63–73	23–55	0.13–0.33

Table 1. Volatile degassing budget calculations. MIs: melt inclusions; cpx: clinopyroxene; opx: orthopyroxene; amph: amphibole. Calculations of mass of volatile released and amount of S sequestered in Cu-Fe-S detailed in the method. X^{RM} is the residual melt fraction, i.e. the ratio of Th contents in the 1257 trachydacitic whole-rock and the residual melt (0.59). The concentration C_i^{system} of each volatile species *i* in the 1257 trachydacitic system is derived from the C_i/C_{Th} ratio of the parent basalt assuming the conservation of the ratio of volatile contents respective to Th during basalt differentiation. $m_S^{fluid} = m_{magma} \times X^{fluid} \times S^{fluid}$, where X^{fluid} is the maximum amount of deeper-derived vapour (1.5 wt%). The maximum mass of S released is the sum of the emissions of the trachydacitic system (79 Tg S) and m_S^{fluid} (13 Tg S). See text for Cl and Br. Global atmospheric (troposphere and stratosphere) mixing ratios of halogen X is given by $[X]_{atm} = \frac{n_X}{n_{air}}$, where n_X is the amount of substance (moles) of halogen X released by the eruption, and n_{air} is the amount of substance (moles) of air in the atmosphere, i.e. 1.5×10^{20} mol. Stratospheric emissions calculated as 80% of S and 10–25% of Cl and Br total emissions³⁸. SO₂ emissions (in Tg, i.e. megatons) correspond to twice the reported S emissions.

Supplementary Figure S6) indicates the sporadic influx of SO₂ in the gas phase²⁶, as supported by the occurrence of submicroscopic crystals of anhydrite and gypsum (and other S-bearing complex minerals) on the walls of fluid inclusions (Supplementary Figure S5c–e).

The average Cl concentration in Rinjani-Samalas basalts (830 ± 80 ppm) is typical of a volcanic arc domain²⁷. The trachydacitic system displays a total Cl content of 3500 ± 175 ppm (Fig. 4e) accounting for the Cl distributed between the melt, the Cl-bearing minerals (amphibole and apatite) and the pre-eruptive vapour. Simple mass balance calculation based on apatite composition (42.9 ± 0.2 wt% of P₂O₅ and 0.89 ± 0.02 wt% Cl; Supplementary Table S5) indicates that up to 1.5 ± 0.3% of apatite crystallised (Methods) with a negligible effect on the Cl concentration of the melt (~130 ppm). Furthermore, amphibole from the 1257 trachydacite with an average Cl content of 700 ± 100 ppm (Supplementary Table S6) only removed a minor amount of Cl from the melt. Residual melts trapped in pyroxenes (2250 ± 140 ppm), and amphibole (2170 ± 210 ppm) share similar Cl concentrations with matrix glasses (2170 ± 210 ppm; Supplementary Table S2), indicating low Cl exsolution from the melt during eruption. Cl is therefore dominantly exsolved into the pre-eruptive vapour phase at shallow depth, a feature that explains the significant decrease of the Cl/Th ratio during *in-situ* crystallization of the trachydacitic magma (Fig. 4e). Furthermore, the positive correlation between Cl and FeO (Fig. 4f) corroborates mixing of partly degassed trachydacitic magma batches differing by their volatile contents.

Pumice clasts from the four phases of the eruption contain on average 1230 ± 45 ppm of Cl (Table 1). This value is in agreement with the residual Cl content of matrix glasses after correction for crystallization. Br contents average 1.6 ± 0.2 ppm in basaltic scoria and 2.92 ± 0.08 ppm in 1257 pumice clasts (Table 1). The average Br/Cl ratio of the basaltic whole-rock ($3.7 \pm 0.4 \times 10^{-3}$) is in agreement with the upper range of published values for arc/back-arc basin basalts ($1-3.2 \times 10^{-3}$)²⁸. This probably suggests a minimum effect of syn-eruptive degassing on the Br/Cl ratio²⁹. Considering that the whole-rock Br/Cl ratio ($3.7 \pm 0.4 \times 10^{-3}$) is characteristic of the basaltic magma ratio, we calculated indirectly a Br concentration of 3.1 ± 0.4 ppm in the basalt containing 830 ± 80 ppm of Cl (Table 1). Such a Br content is comparable to the concentrations measured in basaltic melt inclusions of

Masaya, Nicaragua³⁰. We stress that from the onset to the end of the 1257 eruption, whole-rocks record similar volatile concentrations (S, Cl, Br), suggesting a relatively constant extent of the syn-eruptive degassing.

Atmospheric volatile emissions of the 1257 Samalas eruption

The atmospheric volatile yields from volcanic eruptions are usually estimated using the *petrologic method*, which involves subtracting the volatile concentrations in the residual melt corrected for crystallization from those of undegassed melt inclusions³¹. The pre-eruptive volatile contents of the residual melt (170 ± 50 ppm of S and 2235 ± 155 ppm of Cl) and that of the degassed matrix glass (50 ± 20 ppm of S and 2170 ± 110 ppm of Cl) indicate volatile losses of 140 ± 50 ppm of S and 960 ± 190 ppm of Cl (Table 1). When scaled to the mass of erupted magma (calculated using a volume of 40 km^3 and a DRE density of 2500 kg/m^3 , see Methods), the syn-eruptive devolatilisation of the melt would have released $14 \pm 5 \text{ Tg}$ of S (28 Tg of SO_2) and $96 \pm 19 \text{ Tg}$ of Cl. This calculation neglects, however, the amounts of S and Cl stored in the pre-eruptive vapour phase, and thus provides minimum estimates.

In the following, we quantify the atmospheric yields using the representative volatile contents of the whole 1257 trachydacitic system. Its S content is recorded by melt inclusions trapped in plagioclase An_{82-75} (Table 1) and accounts for the S allocated in the trachydacitic melt, the pre-eruptive vapour phase, and the iron sulphides (Fig. 3). We further assume that sulphide breakdown prior to eruption through interaction with a pre-existing fluid phase³² did not occur, as they likely remained stable prior to eruption. Given the negligible effect of iron sulphide on the S dissolved in the melt, the S degassing (melt + vapour) is provided by the difference between the initial S concentration of the system (870 ± 20 ppm) and the S content of the 1257 whole-rock (80 ± 20 ppm), which yields a loss of 790 ± 30 ppm (Table 1). Such a loss corresponds to the emission of $79 \pm 6 \text{ Tg}$ of S ($158 \pm 12 \text{ Tg}$ of SO_2).

The highest Cl content (3500 ± 175 ppm) recorded by the trachydacitic melt inclusions representative of the 1257 whole-rocks (Fig. 4e) accounts for the Cl allocated in the melt and in the pre-eruptive vapour phase. Using the average Cl concentration in the whole-rock (1230 ± 45 ppm), we calculate a Cl loss of 2270 ± 180 ppm, indicative of an atmospheric discharge of $227 \pm 18 \text{ Tg}$ of Cl (Table 1).

Underplating basalt may contribute to volatile emissions during plinian eruptions of silicic magmas⁶. Despite the absence of any obvious petrologic evidence of the contribution of the parent basalt into the degassing budget, we further explore how the potential contribution of a deeper-derived vapour phase would have increased the atmospheric emissions of the 1257 eruption. Using the S concentration of basaltic melt inclusions, we calculate that the trachydacitic system derived through 81% of fractional crystallization of the parent basalt would have contained 1 wt% of S (S^{system} ; Table 1), distributed between the trachydacitic melt (S^{melt}), the pre-eruptive vapour phase (S^{fluid}), and the poly-metallic sulphides ($S^{\text{Cu-Fe-S}}$). The Cu concentration decreases from basalt to trachydacite (Fig. 2c) indicating that segregated Cu-bearing sulphides (Cu-Fe-S) sequestered a maximum of 690 ppm of S (Methods; Table 1). Considering such a sink of S in Cu-sulphides and the proportion of S dissolved in the trachydacitic melt (870 ± 20 ppm), the pre-eruptive vapour phase would have stored a maximum of 8500 ppm of S. This suggests a partition coefficient $D_S^{\text{fluid/melt}}$ of 13, which fills a gap of experimental constraints on S solubility for these magma conditions. In order to calculate the mass of S that could have been emitted by additional pre-eruptive vapour, it is necessary to assess the amount of water lost during the magma differentiation. An amount of 5–6 wt% fluid is thought to be an upper limit beyond which percolation would occur and the fluid would be lost from the system³³. Furthermore, an average value of 5 wt% fluid is consistent with the discrepancy between the water content of the trachydacitic melt (3.7 ± 0.3 wt%) and the maximum H_2O concentrations that an amphibole-bearing dacite could display ($8 < 10$ wt%)²². Assuming a maximum of 5 wt% of fluid in the system including 3.5 wt% exsolved during *in-situ* crystallization of the trachydacite, the parent basalt would have contributed to 1.5 wt% of supplementary vapour. This corresponds to an addition of $13 \pm 6 \text{ Tg}$ of S onto the S budget (Table 1). The large uncertainty on this estimate reflects the complexity of quantifying the proportion of water, the major component of the vapour phase. The 1257 eruption would thus have released a maximum of $92 \pm 8 \text{ Tg}$ of S ($184 \pm 16 \text{ Tg}$ of SO_2).

We estimate similarly a total theoretical Cl available in the trachydacitic system of 4300 ± 720 ppm (Table 1). Within the error, such concentration is relatively in agreement with the initial Cl content of the melt inclusions representative of the 1257 whole-rocks (3500 ± 175 ppm), suggesting minor Cl fractionation by Cl-bearing phases during basalt differentiation. The potential contribution of the parent basalt onto the Cl budget is thus likely negligible.

Partial exsolution of magmatic Br into pre-eruptive vapour may occur, and its fluid/melt partition coefficient $D_{\text{Br}}^{\text{fluid/melt}}$ is typically much larger than $D_{\text{Cl}}^{\text{fluid/melt}}$ ³⁴. Following the previous reasoning, the initial Br concentration calculated for the parent basalt indicates a total of 16 ± 3 ppm Br available in the trachydacitic system (Table 1) allocated in both the melt and the pre-eruptive vapour phase. Using the average Br content of the 1257 whole-rocks (2.92 ± 0.08 ppm), we calculate a Br loss of 13 ± 3 ppm that corresponds to a maximum emission of $1.3 \pm 0.3 \text{ Tg}$ of Br (Table 1).

The largest volatile release of the Common Era

The 1257 Samalas eruption produced a S injection of $79 \pm 6 \text{ Tg}$, equivalent to $158 \pm 12 \text{ Tg}$ of SO_2 (Table 1), and a maximum of $184 \pm 16 \text{ Tg}$ of SO_2 if the parent basalt contributed to the degassing budget. Hence, the influx of volatiles from the basalt would not exceed 15% of the total budget. Furthermore, our new approach increases by a factor of 5 the minimum S emission derived from the classic petrologic method. We stress that such emission probably occurred within a day⁹, and is much higher than the total annual global volcanic SO_2 flux of $15\text{--}21 \text{ Tg/yr}$ associated with atmospheric emissions from both quiescent and explosive degassing volcanoes³⁵. In addition to the prodigious S yield, the 1257 Samalas eruption emitted $227 \pm 18 \text{ Tg}$ of Cl, and a maximum of $1.3 \pm 0.3 \text{ Tg}$ of Br taking into account the potential contribution of the basalt. These yields stand as the largest volatile emissions

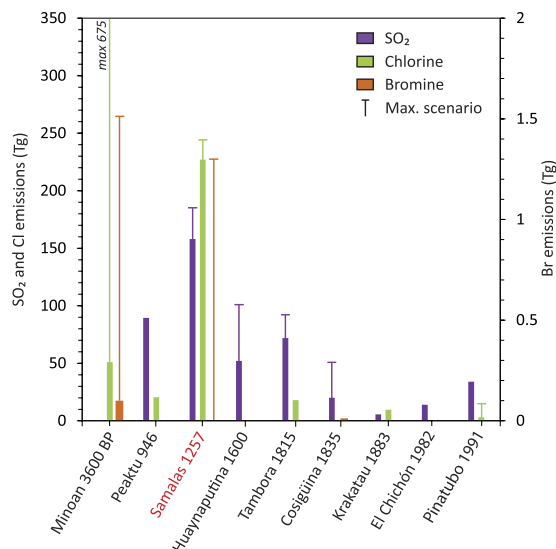


Figure 5. The 1257 Samalás eruption produced the largest volatile emissions of the Common Era. Plots of SO₂, Cl and Br emissions (in Tg, i.e. megatons) for climate-impacting plinian eruptions. Yields of eruptions before 1980 are from petrological studies. Fine lines represent the maximum estimates (associated with the contribution of the parent basaltic magma in the case of the 1257 eruption). References: Minoan 3,600 y B.P. (Greece)³⁷; Peaktu 946 A.D. (DPRK/China)³⁶; Huaynaputina 1600 (Peru)⁶⁶; Tambora 1815 (Indonesia), emissions re-calculated at 36–45 Tg S (73–91 Tg SO₂) and 18–23 Tg Cl based on the syn-eruptive losses (400 ppm S; 200 ppm Cl)^{53,67} and new volume estimates of 41 ± 4 km³ DRE⁶⁸ and 51 km³ DRE⁶⁹; Cosigüina 1835 (Nicaragua)^{30,70}; Krakatau 1883 (Indonesia)⁷¹; El Chichón 1982 (Mexico)⁷²; Pinatubo 1991 (Philippines)⁴.

of the Common Era (Fig. 5), exceeding by a factor of two the SO₂ emissions of the 946 Peaktu eruption (90 Tg of SO₂)³⁶, and of the 1815 Tambora eruption (73–91 Tg of SO₂) that had a devastating impact on climate on a global scale. Furthermore, the Cl yield of the Samalás eruption constitutes the largest emission since the Minoan eruption of Santorini, 3,600 y B.P.³⁷. Although the Br emissions have been estimated for only a few Holocene eruptions, the Br release of the 1257 Samalás eruption (1.3 ± 0.3 Tg) is similar to the maximum scenario of Br yield of the Minoan eruption (1.5 Tg of Br)³⁷, and one order of magnitude higher than the Br emission of the Chiltepe eruption, 1.8 ka (0.125 Tg)³⁰.

Stratospheric injection and potential impact on climate and atmospheric chemistry

Sulphur gases are scavenged by incorporation into ice particles in volcanic plumes before reaching the tropopause, resulting in stratospheric emissions which probably represents 80% of the initial injection³⁸. Such a process would suggest that the 1257 Samalás eruption produced a stratospheric injection of ~126 Tg of SO₂ (Table 1), which is within the range of the two SO₂ stratospheric scenarios of 96 and 138 Tg of SO₂ estimated based on climate modelling and tree-ring based hemispheric temperature reconstructions¹⁷. If the parent magma contributed to the degassing budget, the stratospheric injection of SO₂ could reach up to 150 Tg, which should be considered as a maximum value. Hence, our estimates corroborate the upper range of sulphate yields derived from ice-core data and temperature reconstructions. We also stress that the low discrepancies between our stratospheric estimates and ice-cores based scenarios may reflect the efficient gas injection from the onset to the end of the eruption. Indeed, the stratospheric loading of volatiles is correlated to the initial dynamics of an eruption that needs to be sufficiently powerful to inject a pre-existing vapour phase accumulated at the top of the magmatic reservoir^{33,39} into the stratosphere via a high-flux gas jet and convective column before wholesale column collapse.

The input of both volcanic Br and Cl enhances the catalytic destruction of ozone because the resulting BrO is a reaction partner for ClO⁴⁰. Such reactions can occur either when halogens are released at tropospheric altitudes or when they are directly injected into the stratosphere. The Cl and Br emissions of the 1257 Samalás eruption represent initial atmospheric concentrations of 36 ppbv of Cl and 90 pptv of Br (Table 1). For comparison, the pre-industrial atmospheric Cl mixing ratio was 0.55 ppbv, corresponding to methyl chloride emissions from the oceans, i.e. before the onset of anthropogenically dominated emissions, and increased up to 3.8 ppbv in the late 1990s, during the Antarctic ozone hole climax^{41,42}. The pre-industrial Br mixing ratio was 5 pptv, and increased up to 20 pptv in the late 1990s⁴³. Hence, the 1257 Samalás eruption produced increments in globally averaged Cl and Br concentrations that exceed the pre-1980 levels by a factor of 65 for Cl and 18 for Br, and the ozone-hole climax concentrations by factors of 9 for Cl and 5 for Br. These results likely suggest that the 1257 eruption discharged enough halogen gases into the atmosphere to trigger ozone destruction cycles. We stress that such comparisons should be made carefully, however, given that the mixing ratios calculated for the 1257 Samalás eruption correspond to initial and local increases of Cl and Br concentrations, whereas background mixing ratios are global annual values.

A significant proportion of Cl and Br released by the 1257 Samalás eruption was likely scavenged by hydrometeors during ascent in the plume and may not have reached the stratosphere. Although satellite-based

measurements of present-day volcanism enhanced our understanding of the fate of volcanic halogens in the stratosphere, much remains to be investigated, in particular concerning Br emissions and potential effects of large tropical explosive eruptions on stratospheric ozone⁴⁴. Sophisticated plume experiments and modelling³⁸ showed that 10% to 25% of halogen gases can reach the stratosphere, although the scavenging efficiency strongly depends on several parameters such as the latitude, the salinity of the fluid, the gas phase composition and the ability of ash particles to capture halogens⁴⁵. Comparison of tropical versus high latitude eruptions shows that tropical humid atmospheric conditions, including the occurrence of a typhoon, may cause efficient scavenging of halogens as observed for the 1991 Pinatubo eruption^{46,47}. It has been observed in the case of the 1982 El Chichón eruption, however, that a high mass flux rate within a plinian plume can partly preserve halogens from being scavenged, thus enhancing the stratospheric injection⁴⁸. Considering that: (i) the 1257 Samalas eruption more likely occurred between May and July 1257, i.e. during the Indonesian dry season^{7,17}; and (ii) the strong mass flux rates of the two plinian phases (2.8×10^8 kg/s during phase 1 and 4.6×10^8 kg/s during phase 3)⁹ are one order of magnitude higher than the maximum flux rate of the 1982 El Chichón eruption (6.8×10^7 kg/s)^{49,50}, we assume that the stratospheric injection of halogens could likely have reached 10 to 25% of the initial load. This would correspond to stratospheric injections of 23–57 Tg of Cl and 0.1–0.3 Tg of Br (Table 1) for the 1257 eruption. Volcanic-induced ozone depletion has been observed as a consequence of the 1991 Pinatubo eruption that caused the destruction of 2–6% of global average total ozone⁴⁶ due to the release of 3–16 Tg of Cl⁴ most of which was scavenged by typhoon Yunya during the eruption. The combined Cl (51–675 Tg) and Br (0.1–1.5 Tg) emissions of the Minoan eruption of Santorini (Greece) would have provoked reductions of 20 to >90% of ozone at northern high latitudes³⁷. This strongly suggests that the 1257 Samalas eruption likely provoked strong ozone destruction, even for the lower-bound 10%-injection scenario. This process is enhanced by the simultaneous emission of S, as sulphate aerosols provide surface area for heterogeneous chemical reactions that activate Cl and Br species, thus enabling and enhancing the catalytic destruction of ozone^{51,52}.

Our study underscores the fundamental importance of considering magmatic systems in their totality as well as the evolution of the behaviour of volatiles during magma differentiation in order to significantly improve the quantification of the degassing budget of large climate-impacting explosive eruptions. The application of this methodology to other magmatic systems that have produced volatile-rich intermediate magmas would enhance the reconstruction of the climate impact of past volatile emissions associated to eruptions of large to moderate volumes of S-rich magmas (e.g. 1815 Tambora⁵³, 1835 Cosigüina⁵⁴). Given the prodigious amounts of volatiles released by the 1257 Samalas eruption, interactions between processes involving S, Cl and Br should be considered in future global scale climate-modelling. Although the probability of occurrence of such a large eruption in the next decades is statistically low, the ozone-destruction power of volcanic eruptions in general should be systematically assessed, particularly given that the Antarctic ozone-hole is decreasing⁴³ and that stratospheric halogen concentrations are expected to reach their pre-industrial level during the second half of the century.

Methods

Calculation details. Fraction of solid removal. We apply Rayleigh's law to Th concentrations (Th is highly incompatible) in basaltic and trachydacitic melts as $C_{Th}^{pre-eruptive} = C_{Th}^{basalt} \times f^{D_{Th}-1}$, where $C_{Th}^{pre-eruptive}$ is the Th content of the 1257 system (9.0 ± 0.9 ppm), C_{Th}^{basalt} is the Th content of the basalt (1.7 ± 0.2 ppm), f is the proportion of remaining melt, and the global partition coefficient of Th D_{Th} is $\ll 1$. Equation is then simplified as $\frac{C_{Th}^{pre-eruptive}}{C_{Th}^{basalt}} = \frac{1}{f}$. We calculate a proportion of trachydacitic melt f of 0.19 ± 0.03 , i.e. a solid removal $(1-f)$ of 0.81.

Water loss during in-situ crystallization of the 1257 trachydacite. The conservation of the ratio of water respective to Th during *in-situ* crystallization of the trachydacite is expressed as $\frac{C_{H_2O}^{pre-eruptive}}{C_{Th}^{pre-eruptive}} = \frac{C_{H_2O}^{RM}}{C_{Th}^{RM}}$. *In-situ* crystallization of the 1257 trachydacite with 3.9 ± 0.1 wt% H₂O and 9.0 ± 0.9 ppm Th indicates a theoretical water content of 7.2 wt% in the residual melt (RM). The latter actually contains 3.7 ± 0.3 wt% H₂O, suggesting the exsolution of 3.5 wt% of H₂O.

Proportion of apatite. Using P₂O₅ concentrations of the 1257 trachydacite (0.42 ± 0.05 wt%; Supplementary Table S3) and of the 1257 matrix glass (0.13 ± 0.06 wt%; Supplementary Table S3), we estimate a P₂O₅ loss ΔP_2O_5 of 0.6 ± 0.1 wt% due to apatite crystallization during *in-situ* crystallization. The proportion of P₂O₅ in apatite ($X_{P_2O_5}^{apat}$) is given by $X_{P_2O_5}^{apat} = \frac{\Delta P_2O_5}{C_{P_2O_5}^{apat}}$, where $C_{P_2O_5}^{apat}$ is the P₂O₅ content of apatite (42.9 ± 0.2 wt%; Supplementary Table S5). We thus calculate a proportion of 0.015 ± 0.003 of apatite that would account for the removal of a maximum of 130 ppm of Cl from the melt.

Proportion of Fe-sulphides in the 1257 trachydacitic system. The proportion of iron-sulphide in the 1257 trachydacite is derived from $C_S^{WR} = C_S^{degassed RM} \times X^{RM} + C_S^{sulphide} \times X^{sulphide}$, where X^{RM} is the residual melt fraction (0.59), and $C_S^{sulphide}$ is the S content of Fe-sulphides (38.9 ± 0.5 wt%; Supplementary Table S4). This equation yields a proportion of iron-sulphides $X^{sulphide}$ of 0.01% in the 1257 trachydacite.

Mass of volatile released. The mass of volatile i released is calculated as $m_i = m_{magma} \times \Delta i \times 10^{-15}$, where 10^{-15} is the conversion factor of ppm into kg, and $m_{magma} = \rho_m \times V$ is $1.00 \pm 0.08 \times 10^{14}$ kg (DRE magma density ρ_m is 2500 kg/m³, DRE erupted volume V is 40 ± 3 km³).

Sulphur uptake by poly-metallic sulphides. Using Cu concentrations of the basalt (114 ± 32 ppm; Supplementary Table S3) and of the 1257 trachydacite (11.4 ± 0.8 ppm; Supplementary Table S3) derived through 81% of crystallization, we calculate that 580 ppm of Cu (ΔCu) were fractionated by Cu-bearing sulphides segregated during the basalt differentiation. The amount of associated S is given by $S^{Cu-Fe-S} = \Delta Cu \times C_S^{sulphide} / C_{Cu}^{sulphide}$ where $C_S^{sulphide}$ (32.1 ± 0.6 wt%) and $C_{Cu}^{sulphide}$ (27.0 ± 1.1 wt%) are S and Cu contents in Cu-Fe sulphides, respectively (Supplementary Table S4). This calculation yields an amount of 690 ± 225 ppm of S removed from the melt by poly-metallic sulphides. This amount is considered as a maximum because we neglected the possible fractionation of Cu into the vapour phase.

Analysis of melt inclusions, matrix glasses and fluid inclusions. *Water and CO₂ analysis.* Melt inclusions were analysed using a Thermo-Nicolet 6700 FTIR spectrometer coupled with an optical/IR Spectra-Tech microscope (IPGP, France). Water and carbon concentrations were calculated using the Beer-Lambert law ($C(\text{wt}\%) = 100 \times A \times M / (\varepsilon \times \rho \times e)$, where A is the absorbance, M the molar mass (g/mol), ε the molar absorptivity (L/mol·cm), e the thickness (cm), and ρ the glass density in g/L). Melt inclusion densities were calculated with the model of Lange and Carmichael (1987), using a partial molar volume of water of 12.0 ± 0.5 cm³/mol⁵⁵. Dissolved total water concentrations ($H_2O_{molec} + OH^-$) were calculated using the broad absorption band at 3550 cm⁻¹ and absorption coefficients of 62.8 L/mol·cm⁵⁶ for basalt, and of 68.7 ± 1.0 L/mol·cm⁵⁷ for trachydacite. Carbon concentrations were determined using the 1515 cm⁻¹ CO₃²⁻ absorption band after subtraction of the background, previously acquired on a volatile-free basalt. Deconvolution of the carbonate peaks taking into account the contribution of the 1630 cm⁻¹ H₂O_{molecular} band was performed using PEAK FIT software. The absorption coefficient was calculated as $\varepsilon_{1515} = 451 - 342 \times Na / (Na + Ca)$ ⁵⁸. The analytical uncertainties on CO₂ and water concentrations were 17% and $\geq 10\%$, respectively. CO₂ or carbonate ions were not observed in trachydacitic melt inclusions.

Major element, S and Cl analysis. Major element concentrations in glasses and host minerals were measured by electron probe micro-analysis (EPMA) using a Cameca SXFive electron probe (Camparis, Paris, France). In melt inclusions they were measured with an accelerating voltage of 15 kV, a 4–10 nA defocused beam and peak counting times of 10–30 s depending on the element. Sodium was measured first with a 5 s peak count time in order to minimise alkali loss. S, Cl and P concentrations were determined with a 30 nA defocused beam and counting times of 120–200 s on peak. S speciation was investigated in these analytical conditions in a large olivine-hosted melt inclusion of the dataset, by scanning the peak position for the wavelength of S K α radiation ($\lambda SK\alpha$)⁵⁹. Accuracy of S and Cl measurements were ensured by analysis of KE12 (3332 ± 42 ppm Cl; 183 ± 6 ppm S; $N = 57$), of the alkali trachyte CFA47 (5399 ± 175 ppm Cl; 67 ± 17 ppm S; $N = 61$), and of the international standard Vg2 (291 ± 14 ppm Cl; 1435 ± 40 ppm S; $N = 43$). The analytical errors are $\leq 6\%$ and 3% for S concentrations < 200 ppm and above 1400 ppm, respectively; 4% and $\leq 2\%$ for Cl concentrations < 300 ppm and > 3300 ppm, respectively. The detection limits were 55 ppm for S and 60 ppm for Cl.

Trace element analysis. Abundances of 30 traces elements (Ba, Ce, Co, Cu, Ni, Sc, V, Dy, Er, Eu, Gd, Hf, Ho, La, Lu, Nb, Nd, Pb, Pr, Rb, Sm, Sr, Ta, Tb, Th, Tm, U, Y, Yb and Zr) were determined in melt inclusions by laser ablation inductively coupled plasma mass spectrometry (LA-ICP-MS) at the Laboratoire Magmas et Volcans (Université Blaise Pascal, Clermont-Ferrand, France), using a 193 nm ArF excimer laser ablation system (Resonetics M50) coupled to a 7500 cs Agilent ICP-MS system, with helium as the ablation gas. Samples were analysed using a laser repetition rate of 2 Hz, laser spot diameter range of 20–40 μ m and pulse energy of 6 mJ (14 J/cm²). The background was measured for 30 s before ablation, and each analysis lasted 100 s. Measurements were calibrated against NIST 612 glass standard⁶⁰, using CaO as the internal element reference. Repeated analyses of BCR2-G glass international standard for each spot size were reproducible within 1 to $\leq 5\%$ RSD for most elements and $\leq 7\%$ RSD for Gd and some elements with low abundances such as Yb and Hf (< 5 ppm) and Tb, Tm, Lu and Ta (< 1 ppm). Our measurements on BCR-2G standard were compared to published values (Supplementary Table S7). Melt inclusions were analysed in single spot. Careful examination of counting statistics of each element and data reduction was performed using the Glitter software⁶¹.

Fluid inclusions characterisation. Plagioclase-hosted primary fluid inclusions of the 1257 A.D. rocks were analysed by Raman microspectrometry at the Ecole Normale Supérieure, Paris (France). Spectra were acquired in ambient conditions using a Renishaw INVIA spectrometer. This device is equipped with an Ar laser source giving an incident beam with a 514.5 nm wavelength, focused through a Leica microscope. The Rayleigh scattering component was removed by a Notch filter, and the Raman-scattered light was dispersed by a holographic grating with 1800 lines/mm and detected by a CCD camera. A power of 2 mW was used to avoid heating effects and sample damage. Raman spectra were acquired between 200 and 3900 cm⁻¹ in order to identify S-, CO₂- and H₂O-bearing components of the fluid phase.

Whole-rock analysis. Major and trace element analyses of whole-rock samples are reported in Métrich *et al.* (in review). In this work, S and halogens (Cl, F and Br) were extracted by the pyrohydrolysis method⁶². About 500 mg of powdered whole-rock mixed with ~ 500 mg of V₂O₅ in a platinum crucible was heated at 1200 °C in a quartz combustion tube through a H₂O-vapour stream transported by a nitrogen flux. The extracted S and halogen species are converted into acids by hydrolysis in the H₂O-vapour which was further condensed in a cooling system. The condensate was collected in a vial containing 10 mL of a NaOH solution (25 mmol/L). The vapour flux was adjusted in order to complete the extraction in 45–60 min and collect 80–100 ml of solution. The dilution factors ($mass_{solution} / mass_{sample}$) ranged from 180 to 250. After the complete extraction, the solutions were immediately analysed (for Cl and F) by liquid chromatography using a Dionex DX120 ion chromatograph

(IC) with an Ion Pac AS9-HC (Dionex) anionic column, performed in suppression mode (ASRS-UltraII). The detection limit was 100 $\mu\text{g/L}$ for both F and Cl in the sample solution. Solution concentrations were calculated using a calibration curve (0–50 ppm), and were further converted into rock concentrations using the dilution factor. Taking into account the mean dilution factor, detection limits in rock samples were 15 to 20 mg/kg for both elements. Br contents were determined using a 7900 Agilent ICP-MS device, in low resolution with a Scott spray chamber and a micro nebulizer (0.2 ml/min). Br was analysed in the ‘no gas’ mode, with an acquisition time of 0.3 sec, measuring 5 points as peak pattern, 3 replicates and 100 sweeps by replicates, 70 sec for the uptake, 40 sec as stabilisation time and 60 sec for rinsing time. Br solution concentrations were further calculated using a calibration curve (0–50 ppb), and were further converted into rock concentrations using the dilution factor. To ensure a complete extraction and the accuracy of the analyses, pyrohydrolysis/IC/ICP-MS was performed on international standards covering a wide compositional range and S, Cl, Br and F concentrations. We stress that our results were reproducible within 1–10% RSD for Cl, within 6–16% RSD for Br, and within 2–16% RSD for S except for AGV-1 (35% RSD). The efficient extraction of S from sulphide-bearing samples was guaranteed by the analysis of a sulfide-bearing syenite standard SY-2. Our results compared to published reference values are reported in Supplementary Table S8.

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Acknowledgements

We thank the Indonesian Ministry of Research and Technology RISTEK and the Nusa Tenggara Barat Governor's Office, M. Hendrasto (CVGHM), Surono (Badan Geologi) and J.-P. Toutain (IRD) for their permission to conduct fieldwork and administrative support. We also thank H. Rachmat, Muntaharlin and J.-P. Degeai for providing additional samples. We thank M. Fialin, N. Rividi, J.-L. Devidal, D. Deldicque and V. Zanon for dedicated assistance during EPMA, LA-ICP-MS and Raman analyses. ^{14}C dates were obtained by C. Moreau and J.-P. Dumoulin (LMC14, CNRS UMS2572). SRTM data was provided by CIAT-CSI. This study was partly funded by INSU-CNRS CT3-ALEA projects ECRin I and II and INSU-CNRS Artemis 2014 for ^{14}C dating in the framework of C.M. Vidal PhD's thesis. This is IGP contribution 3754.

Author Contributions

C.M.V., N.M. and J.-C.K. conducted fieldwork, sampling, sample preparation and analysis, interpreted the results, performed degassing budget calculations and co-wrote the manuscript and co-drafted the figures. A.M. conceived and conducted the pyrohydrolysis/IC/ICP-MS analyses. I.P., N.K., V.R. and F.L. conducted fieldwork and sampling. All authors reviewed the manuscript.

Additional Information

Supplementary information accompanies this paper at <http://www.nature.com/srep>

Competing financial interests: The authors declare no competing financial interests.

How to cite this article: Vidal, C. M. *et al.* The 1257 Samalas eruption (Lombok, Indonesia): the single greatest stratospheric gas release of the Common Era. *Sci. Rep.* **6**, 34868; doi: 10.1038/srep34868 (2016).



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