Halogen Enrichment of Siberian Traps Magmas During Interaction With Evaporites

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Volatile emissions to the atmosphere associated with the Siberian Traps eruptions at the Permian-Triassic boundary were sourced from the outgassing of primary magmas and the sedimentary host rocks into which they were intruded. Halogens in volcanic gases may have played an important role in environmental degradation and in stratospheric ozone destruction. Here we investigate how halogens behave during the interaction between salts and basalt magma emplaced as sills and erupted as lava. We present whole-rock, trace, and halogen concentrations for a suite of samples from three locations in the Siberian Traps Large Igneous Province, including basalt lavas erupted, and dolerites intruded into both organic-bearing shales and evaporites. Dolerites are enriched in Cl, Br, and I; their enrichment in Cl is similar to MORB and OIB that have been inferred to have assimilated seawater. The dolerites exhibit halogen compositional systematics, which extend towards both evaporites and crustal brines. Furthermore, all analyzed samples show enrichment in Rb/Nb; with the dolerites also showing enrichment in Cl/K similar to MORB and OIB that have been inferred to have assimilated seawater. We infer that samples from all three locations have assimilated fluids derived from evaporites, which are components of crustal sedimentary rocks. We show that up to 89% of the chlorine in the dolerites may have been assimilated as a consequence of the contact metamorphism of evaporites. We show, by thermal modeling, that halogen transfer may occur via assimilation of a brine phase derived from heating evaporites. Halogen assimilation from subcropping evaporites may be pervasive in the Siberian Traps Large Igneous Province and is expected to have enhanced emissions of Cl and Br into the atmosphere from both intrusive and extrusive magmatism.

Keywords: Siberian Traps, halogens, degassing, contact metamorphism, magma

INTRODUCTION

The Siberian Traps Large Igneous Province, now thought to have been emplaced rapidly within 60 ± 48 ka at the end of the Permian (Reichow et al., 2009; Burgess et al., 2014; Shen et al., 2019), was synchronous with the largest mass extinction in Earth’s history (Renne and Basu, 1991; Renne et al., 1995; Burgess et al., 2017). There remains much controversy about the mass extinction mechanism, but there is a consensus that outgassing of volatiles was a primary environmental perturbation agent
(Campbell et al., 1992; Lomax et al., 2005; Self et al., 2005; Beerring et al., 2007; Black et al., 2012). Carbon-, sulfur- and halogen-bearing gas species all have the potential to affect the environment in ways that could be significant enough to cause environmental degradation and impact life on Earth. Emissions of carbon gases (CO₂ and CH₄) were caused by the devolatilization of carbon-rich shales, coal, and evaporites due to heating from sill intrusions (Svensen et al., 2009), as well as from the outgassing of ascending and erupting magmas (Self et al., 2005; Black et al., 2012). These emissions may have led to significant perturbations to the carbon budget, leading to ocean warming and anoxia, with implications for biological productivity and life (Payne et al., 2004). The long-term warming caused by carbon emissions might have been punctuated by short-term (10–100 years) cooling due to sulfate aerosol formation in the troposphere (Self et al., 2005), which might have persisted to varying degrees throughout the Siberian Traps eruptions (Black et al., 2018). Acid rain affected both terrestrial and ocean environments, promoting weathering and ocean acidification (Black et al., 2014). Methane may have been generated from methanogenic microorganisms in the oceans, activated by nickel (Rothman et al., 2014; Le Vaillant et al., 2017), perhaps transported to the marine environment via sulfate aerosols during Siberian Traps eruptions (Rampino et al., 2017). In addition, it has been suggested that the mantle source feeding the Siberian Traps volcanism was fundamentally rich in carbon and chlorine from recycled oceanic crust that had been inefficiently devolatilised during subduction (Sobolev et al., 2011; Sibik et al., 2015; Broadley et al., 2018). Magma outgassing during emplacement in the crust and during eruption would have transferred these volatiles to the atmosphere.

Halogen outgassing systematics during Siberian Traps Large Igneous Province formation is not well understood. It has been suggested that organohalogens emitted from the Siberian Traps magmas and the country rocks may have been a significant factor in causing mutations and damaging the DNA of life via enhanced UVB radiation fluxes (Beerring et al., 2007; Svensen et al., 2009; Benca et al., 2018). Halogen-bearing gas species, once in the atmosphere, can be converted into the reactive molecules chlorine monoxide (ClO) and bromine monoxide (BrO); and chlorine and bromine atoms (Cl and Br) (Oppenheimer et al., 2006). These species readily enter into self-sustaining ozone destruction reactions (Bobrowski et al., 2003; Gerlach, 2004). Bromine free radicals are 40–70 times more efficient in damaging the ozone layer than chlorine free radicals (Daniel et al., 1999; Sinnhuber et al., 2009); and the destructive potential of bromine is enhanced when accompanied by stratospheric sulfate aerosol loading, which is particularly relevant for an evaluation of the Siberian Traps eruption’s climatic effects.

Despite the importance of halogen outgassing for atmospheric chemistry and the atmosphere’s radiation budget, unequivocal geochemical evidence for their origin and outgassing mechanism during the Siberian Traps eruptions remains elusive. The Cambrian and Devonian age evaporites dominate in the southern and northern parts, respectively, of the Siberian craton sedimentary sequence (Figure 1). Sills comprise a significant volume of the Siberian Traps Large Igneous Province (between 37 and 44%) (Vasil’ev et al., 2000). Most of the lavas and sills on the craton were emplaced in a region of subcropping Devonian evaporites, mainly composed of sulfates (in the thick upper and lower parts) and halite (in the middle part) (Zharkov, 1984). Cambrian evaporites are composed of salts (dominated by halite) and proposed to be volatilized in areas adjacent to sills (Svensen et al., 2009; Svensen et al., 2018). Interactions between magmas and evaporites, including the incorporation of sedimentary pore fluids (e.g., brines), can significantly increase the volatile contents of magmas (Heimdal et al., 2019), as has been suggested to explain some high chlorine contents of melt inclusions in Siberian Traps Large Igneous Province rocks (Sobolev et al., 2009; Black et al., 2012). Previous studies have described the textures and skarn mineralogy produced at the contact between dolerites and evaporites in the Nepa River region (Grishina et al., 1991; Grishina et al., 1992; Istomin et al., 2000; Mazurov et al., 2007; Polozov et al., 2016; Grishina et al., 2018; Grishina et al., 2020). Evidence for interactions with magmas may be retained in the basalt structures. However, the analytical challenges associated with the measurement of bulk rock halogens have, up to now, largely prevented close examination. More than 500 basaltic phreomagmatic and ~250 magnetite-rich breccia pipes are found in the Siberian Traps LIP (Svensen et al., 2009; Polozov et al., 2016), Figure 1. These magmatic breccia pipes likely played a vital role in the liberation of halogens to the atmosphere (Svensen et al., 2009; Aarnes et al., 2010; Polozov et al., 2016).

In this paper, we present whole-rock halogen compositions in a magma-sediment system from the Siberian Traps Large Igneous Province to assess the extent to which halogens have been assimilated into the magmas and outgassed from the surrounding salts; and the mechanisms by which this occurred. We present the results of a detailed geochemical investigation of both dolerite sills and basaltic lava flows emplaced in different basin segments (Dyupkun Lake, Lower Tunguska River and the Nepa River; Figure 1) with the lower parts dominated by evaporites including salt and dolostones (location 3, Nepa River, “NP-43”-series samples; Figure 1), and the shallow parts dominated by carbonates and siliciclastic rocks (locations 1 and 2, Dyupkun Lake and Lower Tunguska River, “S10”-series samples; Figure 1), including thick coal beds. We discuss the influence of these halogen-rich enclosing sedimentary rocks on magma geochemistry and the volatile budget of the Siberian Traps Large Igneous Province eruptions.

**SAMPLES AND METHODS**

The samples of lavas and sills, including sampling coordinates, are provided in the [Supplementary Material](#). The sampling locations are shown in Figure 1. Petrological descriptions of the S10 series at locations 1 and 2 on the map are provided by Sibik et al. (2015), except for the two samples S10–49 and S10–58. The sill and enclosing salt samples from location 3 on the map (“NP”-series, [Supplementary Material](#)) are first described in this
study. Samples from location 1 are basalt flows of the Ayanskaya Formation at Dyupkun Lake (Putorana Plateau), which were emplaced on Cambrian-Silurian carbonate and terrigenous strata, Devonian evaporites with rock salts, and Carboniferous-Permian terrigenous coal-bearing strata and erupted onto basalts of Tutonchana Horizon. Samples of location 2 are dolerite sills in the Lower Tunguska River region which travelled through Cambrian evaporites with anhydrite, Ordovician-Devonian carbonate terrigenous strata, and were emplaced into Carboniferous-Permian organic-rich shales and coal-bearing strata. Samples of location 3 are from a 40-m thick dolerite sill crossed by borehole 194 in the Nepa River region (inset, Figure 1).

**Figure 1** | Schematic geological map of the Siberian craton and the Tunguska Basin (the inset map at the top shows global setting), showing the distribution of outcropping sills, lava flows, and volcanlastic deposits (Malich et al., 1974). Devonian and Cambrian evaporite’s distribution is from (Zharkov, 1984) and (Petrychenko et al., 2005). Sampling areas (1, 2, and 3) indicate the locations at which samples of lava flows and intrusions were collected and then analyzed for halogens. Some basalt pipes and phreatomagmatic vents are shown from (Svensen et al., 2009).
coordinates N59°9′20″ E107°37′2″). This dolerite sill intruded into the salt horizon of the Lower Cambrian evaporites (mostly halite, but also potassium salts) and was sampled from the middle part of intrusion downward to the footwall contact. Dolerite mineralogy comprises clinopyroxene and plagioclase with frequent olivine, as described by Sibik et al. (2015).

Whole-rock major and trace element compositions of lava flows and intrusions from Dyupkun Lake and Lower Tunguska River regions were obtained using combined ICP-MS and ICP-AES analyses (Sibik et al., 2015) (see Supplementary Material for detail of analytical methods). Major and trace elements for the "NP"-series were analyzed at the University of Cambridge (United Kingdom) by ICP-MS. Before halogen analysis, the small amounts (~1.7–2 g) of finely powdered dolerites were mixed with ~20 ml distilled water. The mixtures were then heated to 70°C to dissolve the dolerite-impregnated salts. This procedure was repeated three times to ensure complete removal of salt material, and then the powders were dried in the oven.

Fine powders of 12 samples of pre-washed dolerites and basalts (Figure 1) were analyzed for halogens at the Alipp6 laboratory, ISTEP, Sorbonne Universitè, Paris, France. Rock powders underwent pyrohydrolysis for halogen extraction: 1 g of the sample powder was mixed with an excess of vanadium pentoxide and then placed in a platinum crucible. It was heated in a combustion tube to 1200°C through water vapor-stream transported by a nitrogen flux. Water vapor containing halogens was condensed by a cooling system and trapped in stainless hydroxide (Michel and Villemant, 2003) followed by Ion Chromatography (IC). The solutions obtained by pyrohydrolysis, the leachates, and dissolved host evaporites from location 3 were analyzed by Ion Chromatography (IC) for chlorine and fluoride and by Inductively-Coupled Plasma Mass Spectrometry (ICP-MS) on a PerkinElmer SCIEX Elan DRC II quadrupole ICP-MS system for bromine and iodine. Fine powders were analyzed for trace elements: 0.1 g of powder was digested in screw-capped PFA (Teflon) containers using the HF-HNO3 technique (Jarvis and Jarvis, 1992) and Evapoclean system for heating and evaporation. The final solution was then diluted by up to 50 ml volume to 2% HNO3. Instrument calibration was performed using international rock standards BCR-2 and BHVO-2 and in-house standards to monitor In, Re, and Rh drift. The measurement’s precision, based on the standard deviation of repeat analysis of standard materials, is 7% for fluorine, 11% for chlorine, 15% for bromine, 57% for iodine, and less than 4% for typical trace elements.

RESULTS

The compositions of lava flows and sills erupted onto or intruded into siliceous continental crust (locations 1 and 2 on Figure 1, samples "S10"-series) and of dolerite sills emplaced in evaporites in the Nepa region (location 3 on Figure 1, samples "NP-43"-series) are reported in Supplementary Material and in Table 1. The major element compositions of the samples plot directly onto the liquid line of descent defined by previous studies of Siberian traps volcanic and plutonic rocks (Hawkesworth et al., 1995; Fedorenko and Czamanske, 1997; Czamanske et al., 2000) and do not show any anomalous enrichment in alkali elements (Supplementary Material Figure S2).

The Rare Earth Element (REE) chondrite-normalized compositions are shown in Figure 2, shown alongside whole rock basaltic compositions from the West Siberian Basin (Reichow et al., 2005) and olivine-hosted melt inclusions from the Gudchikhinsky Formation (Sobolev et al., 2009). Sample S10-49 is more evolved than other samples of the S10-series, with a MgO content of 3.58 wt% compared to 6.93–7.81 wt% for the other S10 series rocks; and higher K2O contents (1.82 against 0.14–1.08 wt%) and significantly higher concentrations of incompatible trace elements (Supplementary Material; Figure 2). Sample S10-49 has a higher La/Yb ratio (10.73) compared to the La/Yb range of 2.77–7.26 in other S10-series magmas. Other S10-series magmas have a variable La/Sm (2.27–4.28) ratio at nearly the same Sm/Yb (1.22–1.70) except the S10-49 sample, which has a Sm/Yb ratio of 2.65.

The basalts and dolerites contain variable concentrations of chlorine (up to 5,331 ppm), bromine (0.44–5.89 ppm), iodine (up to 205 ppb), and fluorine (171–983 ppm) (Table 1; Figure 3, Supplementary Material Figure S3). The basalts erupted onto the siliceous continental crust (location 1) have much lower concentrations of chlorine (< 150 ppm) than the dolerites from location 2, as well as slightly lower fluorine (< 210 ppm compared with 210–990 ppm), bromine (< 1.0 as opposed to 1–1.4 ppm) and similar iodine concentrations (up to 44 and 67 ppb correspondingly). Sample S10-49 (location 2) contains the highest concentrations of F (983 ppm), Cl (5,331 ppm). In contrast, sample S10-44 (location 2) contains the lowest concentration of F (213 ppm) and Cl (325 ppm) within the range of location 2 dolerites (Table 1; Figure 3). When plotted against MgO contents, it can be seen that the Cl contents of the whole rock samples increases with decreasing MgO content (Supplementary Material Figure S3), similar to published melt inclusion compositions (Sobolev et al., 2009; Black et al., 2012). There is no clear relationship between the Cl and Na2O or K2O contents of the rocks, as might be expected for fractionation trends, and the Cl/K2O ratio of 0.33 (derived from a regression through the data in this study) is much larger than the upper limit of 0.06 observed in mid-ocean ridge basalts (Michael and Cornell, 1998). Indeed, there is a very large spread in whole rock and melt inclusion Cl contents over a relatively small range in Na2O and K2O contents, which is not consistent with a fractionation trend (Supplementary Material Figure S3).

The halogen concentrations measured in MORB and OIB are also shown in Figure 3 together with mantle Br/Cl and I/Cl ranges (black, dashed lines) and suggested mantle F/Cl ratios (Kendrick et al., 2020). On a fluorine - chlorine plot (Figure 3A), only basalts from location 1 plot in the MORB field, which is itself not precisely determined (Kendrick et al., 2017). In contrast, all of the dolerites (locations 2 and 3) are chlorine-enriched and plot close to MORB glass inferred to have assimilated seawater Cl (Kendrick et al., 2017). On a bromine—chlorine plot (Figure 3B), all of the basalts and dolerites are outside the mantle range (Br/Cl = 2.8 ± 0.6 × 10^{-3}) (Kendrick et al., 2017). All of the basalts from location 1 and one dolerite (S10-44 of location 2) are slightly Br-enriched (or Cl-depleted); in contrast, all of the other
| Sampling Ht | Location 1, Dyupkun Lake basalts | Location 2, Lower Tunguska River dolerites | Location 3, Nepa River dolerites |
|-------------|----------------------------------|------------------------------------------|--------------------------------|
|             | S10-12                           | S10-44                                   | 864.3 m                       |
|             | S10-15                           | S10-49                                   | 879.2 m                       |
|             | S10-22                           | S10-58                                   | 885.9 m                       |
|             | S10-62                           | NP-9843                                  | NP-9845                       |
|             |                                  | NP-9847                                  | NP-9847                       |
|             |                                  | NP-9848                                  | NP-9848                       |
|             |                                  | NP-9850                                  | NP-9850                       |
| Major elements |                                |                                          |                                |
| MgO         | 7.47                             | 7.81                                     | 4.09                          |
| MnO         | 0.18                             | 0.19                                     | 0.3                           |
| Al₂O₃       | 15.9                             | 16.0                                     | 11.7                          |
| CaO         | 11.5                             | 10.4                                     | 9.20                          |
| TiO₂        | 1.11                             | 1.12                                     | 2.71                          |
| Na₂O        | 2.12                             | 2.22                                     | 2.65                          |
| K₂O         | 0.21                             | 0.43                                     | 0.94                          |
| F₂O₅        | 0.13                             | 0.12                                     | 0.31                          |
| Trace elements |                                |                                          |                                |
| Li           | 21.8                             | 12.7                                     | 19.6                          |
| Br           | 0.54                             | 0.41                                     | 1.1                           |
| Sc           | 44.4                             | 39.8                                     | 53.2                          |
| V            | 321                              | 301                                     | 520                           |
| Cr           | 126                              | 131                                     | 17.2                          |
| Co           | 50.3                             | 53.0                                     | 51.6                          |
| Ni           | 114                              | 136                                     | 25.4                          |
| Cu           | 125                              | 124                                     | 348                           |
| Zn           | 88.7                             | 104                                     | 163                           |
| Ga           | 17.2                             | 17.3                                     | 24.0                          |
| Rb           | 7.68                             | 12.0                                     | 26.6                          |
| Sr           | 210                              | 218                                     | 224                           |
| Y            | 24.5                             | 22.6                                     | 54.7                          |
| Zr           | 81.7                             | 70.4                                     | 198                           |
| Nb           | 4.55                             | 3.81                                     | 9.81                          |
| Sn           | 0.73                             | 0.73                                     | 1.84                          |
| Cs           | 0.30                             | 0.50                                     | 0.52                          |
| Ba           | 111                              | 143                                     | 248                           |
| La           | 7.56                             | 63.9                                     | 14.8                          |
| Ce           | 17.3                             | 15.2                                     | 14.8                          |
| Pr           | 2.33                             | 15.2                                     | 35.8                          |
| Nd           | 10.9                             | 10.1                                     | 24.3                          |
| Sm           | 2.97                             | 2.82                                     | 6.95                          |
| Eu           | 1.02                             | 2.96                                     | 2.21                          |
| Gd           | 3.70                             | 3.41                                     | 8.59                          |
| Tb           | 0.64                             | 1.00                                     | 1.43                          |
| Dy           | 4.15                             | 3.98                                     | 9.58                          |
| Ho           | 0.87                             | 1.66                                     | 1.98                          |
| Er           | 2.50                             | 4.45                                     | 5.66                          |
| Tm           | 0.38                             | 0.35                                     | 0.85                          |
| Yb           | 2.39                             | 3.82                                     | 5.37                          |
| Lu           | 0.35                             | 0.55                                     | 0.79                          |
| Hf           | 2.37                             | 6.70                                     | 5.26                          |
| Ta           | 0.28                             | 0.40                                     | 0.61                          |
| Th           | 1.19                             | 4.99                                     | 2.04                          |
| U            | 0.52                             | 0.83                                     | 0.91                          |
| Halogens |                                |                                          |                                |
| F, ppm       | 204                              | 213                                     | 480                           |
| Cl, ppm      | 92                               | 325                                     | 3022                          |
| Br, ppb      | 400                              | 1400                                    | 3900                          |
| I, ppb       | 19                               | 65                                      | 63                            |
| F/Cl         | 2.2                              | 0.66                                     | 0.16                          |
| Br/Cl (x 10³) | 4.3                            | 4.3                                      | 1.3                           |
| I/Cl (x 10³) | 210                             | 200                                     | 21                            |

Table 1: Whole rock compositions, including major, trace and halogens, of the rocks analyzed in this study. Methods are described in the text and in the Supplementary Material. Locations from which the samples are derived are at the top of the table, as well as sampling height (metres) for the dolerites from location 3, borehole1 9 5.
dolerites (locations 2 and 3; Figure 1) are Cl-enriched (or Br-depleted). On an iodine—chlorine plot (Figure 3C), again almost all of the studied basalts and dolerites plot outside of the mantle range (I/Cl = 60 ± 30 × 10³) (Kendrick et al., 2017). Two basalts (from location 1) and one dolerite (S10-44 of location 2) are I-enriched (or Cl-depleted), whereas all other dolerites (from locations 2 and 3) are Cl-enriched (or I-depleted). Three dolerites from location 3 (the Nega River region) and one dolerite from location 2 (S10-62, the Lower Tunguska River) have an I/Cl ratio that almost coincides with the mantle but lies far from mantle Br/Cl values. However, these samples coincide with the field for MORB and OIB glasses inferred to have assimilated seawater Cl (Kendrick et al., 2017). As a whole, there is enrichment in Cl, and to some extent, Br and I over MORB values observed for both dolerites intruded into evaporites (location 3) and into siliceous continental crust (location 2). On a I/Cl vs Br/Cl plot (Figure 3D), all dolerites and basalts plot significantly outside the mantle field and are dispersed between the endmember compositions of evaporites, seawater, and crustal brines (Kendrick et al., 2017). The borehole 194 samples display a lower I/Cl ratio than “typical” evaporites (Zherebtsova and Volkova, 1966; McCaffrey et al., 1987; Kendrick et al., 2017); the I content of the Nega evaporites is below detection (Table 2). Sedimentary evaporites of the Nega River area have the lowest values of I/Cl, with iodine contents close to the limit of detection.

The stratigraphy of the sill intruded into salts (inset in Figure 1) is illustrated in Figure 4 with the halogen and sulfur concentrations measured in this study. The halogen analysis of leachates from the purification of sill samples intruded into salts (NP-43), and the chlorine, bromine, iodine, and sulfur concentrations in both dolerite (Table 1) and evaporite (Table 2) lithologies are also shown. There are considerable fluctuations in chlorine concentrations in the evaporites on the bottom contact with the sill; above the sill, the lowest chlorine concentrations are found near the contact. Bromine concentrations in the evaporites increase towards the contact with the sill on both sides of the intrusion, but particularly towards the lower contact, from 122 ppm 61 m from the contact to 869 ppm 1.5 m from the contact, which results in an increase in Br/Cl from 0.22 × 10³ to 1.38 × 10³ (Figure 4; Table 2). Iodine is present at higher concentrations in the sill than in the adjacent sedimentary rocks (Figure 4). The evaporite samples are depleted in I (relative to Cl) compared to “typical” evaporites (Figures 3D; Figure 4).

Trace element ratios and chlorine contents for the studied migmatic rocks are plotted in Figure 5. All the dolerite samples have a La/Yb ratio similar to MORB and OIB basalt glasses, but they are Cl-enriched compared to MORB and OIB (except one; S10-44; Figure 5A). On a plot of CI/K vs Rb/Nb (Asafov et al., 2020) all dolerite and basalt samples plot outside the field of MORB and OIB basalt glasses (Kendrick et al., 2017) and are shifted to higher Rb/Nb ratios except for one basalt sample (S10-15) that has the lowest Rb (2.2 ppm) content and coincides with the field of reference samples inferred to have assimilated seawater Cl. All studied samples form a generally positive array on the CI/K vs Rb/Nb plot (Figure 5B).

**DISCUSSION**

**Origin of Halogens in Basalts and Dolerites of the Siberian Traps Large Igneous Province**

Halogens are expected to behave similarly to incompatible elements during mantle melting, and crystallization. The fluorine and chlorine concentrations in the basalts increase with La/Yb for MORB and OIB melts uncontaminated with seawater chlorine (Kendrick et al., 2012; Figure 5A); thus mantle melts with high La/Yb, are expected to be Cl-rich (Figure 5A). As they are all similarly incompatible, the ratios between the halogens do not change significantly during melting or fractionation. These ratios reflect the abundances of the halogens in the MORB-OIB mantle source uncontaminated by seawater (Kendrick et al., 2012). Some OIB glasses proposed to be contaminated with seawater do not demonstrate Cl-enrichment beyond what is expected for their La/Yb ratio (Figure 5A). In contrast, all seawater-contaminated MORB glasses, some contaminated OIB glasses, and almost all studied dolerites of the Siberian Traps are demonstrably Cl-enriched (Figure 5A). Chlorine enrichment for the seawater-contaminated MORB-OIB glasses can also be seen on the CI/K vs Rb/Nb plot (Figure 5B). However, unlike seawater-contaminated MORB and OIB, all of the Siberian basalts and dolerites, except one, are shifted from the cloud of MORB-OIB mantle melts towards high Rb/Nb, with less significant CI/K enrichment. We note that olivine-hosted melt inclusions from the Gudchikhinsky Formation (Sobolev et al., 2009) also show a correlation between CI/K and Rb/Nb and extend to the same region of the plot in Figure 5B. We propose that the trends towards high Cl and LILE exhibited by the Siberian Traps Large Igneous Province magmas from all three locations studied here are...
### TABLE 2 | Whole rock compositions of evaporite samples from the Nepa River borehole 194 measured in this study (methods described in main text and in the Supplementary Material). Sample names are given in the first column. The distance from the contact with the dolerite, in meters, is given for evaporites above the upper contact with the dolerite, and below the lower contact, along with sampling height in the borehole. The halogen compositions are given in parts per million (ppm). Note that the iodine composition of the evaporite samples is at or below detection, which is 5 ppb (parts per billion).

|                       | Distance from contact, m | Sampling height in the borehole, m | F, ppm | Cl (×10³) ppm | Br, ppm | I, ppb | S, ppm | Br/Cl (×10³) | I/Cl (×10⁶) |
|-----------------------|--------------------------|-----------------------------------|--------|---------------|---------|--------|--------|-------------|-------------|
| Above upper contact   |                          |                                    |        |               |         |        |        |             |             |
| NP 9835               | 79.5                     | 765.5                             | b.d.   | 930           | 302     | 5      | 3596   | 0.32        | 5.38        |
| NP 9828               | 4                        | 841                               | b.d.   | 547           | 329     | 5      | 313    | 0.60        | 9.14        |
| NP 9859               | 0.4                      | 844.6                             | 229    | 606           | 477     | 5      | 229    | 0.79        | 8.25        |
| Below lower contact   |                          |                                    |        |               |         |        |        |             |             |
| NP 9852               | 1.5                      | 886.2                             | 252    | 629           | 869     | 5      | 2323   | 1.38        | 7.95        |
| NP 9838               | 9                        | 889.8                             | b.d.   | 536           | 702     | 5      | 4033   | 1.31        | 9.33        |
| NP 9861               | 27.4                     | 904.2                             | b.d.   | 795           | 515     | 5      | 697    | 0.65        | 6.29        |
| NP 9854               | 61                       | 915.7                             | 269    | 550           | 122     | 5      | 406    | 0.22        | 9.09        |

FIGURE 3 | Fluorine (A), bromine (B), iodine (C) versus Cl contents and I/Cl vs Br/Cl ratios (D) for the studied samples. Reference basalt glasses from the Pacific, Atlantic, and Indian MORB (grey unfilled circles) and OIB (grey unfilled triangles) combined together with suggested having assimilated seawater Cl (Kendrick et al., 2017). Error bars corresponded to the analytical uncertainty of measurement omitted for clarity. F/Cl mantle value is not well constrained and varies from 1 to 7 (Kendrick et al., 2017). Legend as in Figure 2.
caused by varying degrees of assimilation of crustal sedimentary rocks (including evaporites) by the Siberian Traps magmas during their emplacement, as has been suggested by others (Lightfoot et al., 1990; Wooden et al., 1993; Callegaro et al., 2021).

In some basalts/dolerites (those from locations 1 and 2) (Figure 3) bromine and, to a lesser extent, iodine are at higher values than MORB-OIB basalts with their F/Cl ratio close to the mantle range 1–7 (Kendrick et al., 2017). The Br/Cl ratio for all basalts and one dolerite (S10-44) exceeds those of mantle melts (determined as $2.8 \pm 0.6 \times 10^{-3}$ by Kendrick et al., 2017) (Figure 3B), and plot close to seawater–crustal brines–sedimentary rocks compositions (Figure 3D). All other dolerites (from locations 2 and 3) intruded into the terrigenous crustal rocks and evaporites are Cl-enriched and plot close to seawater–evaporite compositions (Figure 3D). These deviations in halogen compositions from MORB-OIB basalts are not consistent with degassing from evolved, uncontaminated magmas. Instead, we suggest that the enrichment in Cl and mobile elements (as represented by Rb) is due to the magmas passing through sedimentary strata containing evaporites, which are present at all of the studied localities (Figure 1). Here we note that the halogen systematics of these samples (largely extending to lower Br/Cl than the mantle and towards evaporite compositions; Figure 3D); are not consistent with their having interacted with a postulated high Br/Cl sub-continental lithosphere, as has been previously proposed as a potential halogen source (Broadley et al., 2018). It is very likely that in our sample set, and perhaps other published sample sets, we have no halogen-uncontaminated Siberian Traps rocks; thus, it is not possible to estimate the Siberian Trap’s primary halogen contents based on the current samples. It is likely that halogens are held in the form of halogen-bearing minerals (apatite, mica, amphibole) in the dolerites, although this requires further detailed microanalysis and imaging. The pipes in the Siberian Traps, for instance at Nepa, contain abundant and halogen-rich apatite (Neumann et al., 2017).

The basalts from location 1 (Lake Dyupkun) are subaerial lavas, and are relatively depleted in Cl relative to the dolerite samples from locations 2 (Lower Tunguska) and 3 (Nepa River)
The basalts from location 1 are more primitive than those from locations 2 and 3, although there is some overlap with one sample from location 2 (Figure 2). The basalts from location 1, which are lavas erupted onto the surface, may have undergone a larger degree of degassing relative to the intruded dolerites (which were emplaced at depths of >1 km), which may explain some of the observed depletion in chlorine. Chlorine partitions into a hydrous magmatic vapor during decompression, which may then be outgassed (Webster et al., 1999). The magnitude of chlorine degassing, however, is difficult to estimate from these data. The proportion of Cl degassing from basalt lava flows is estimated to lie in the

(Figure 3; Supplementary Material Figure S3). The basalts from location 1 are more primitive than those from locations 2 and 3, although there is some overlap with one sample from location 2 (Figure 2). The basalts from location 1, which are lavas erupted onto the surface, may have undergone a larger degree of degassing relative to the intruded dolerites (which were emplaced at depths of >1 km), which may explain some of the observed depletion in chlorine. Chlorine partitions into a hydrous magmatic vapor during decompression, which may then be outgassed (Webster et al., 1999). The magnitude of chlorine degassing, however, is difficult to estimate from these data. The proportion of Cl degassing from basalt lava flows is estimated to lie in the

![Figure 6](image-url)
range 27–72% for other flood basalt eruptions (Thordarson and Self, 1993; Thordarson et al., 1996; Self et al., 2008) consistent with experimental studies on basalts and andesites which have yielded aqueous fluid-silicate melt partition coefficients for Cl of between 1 and ∼20 (Bureau et al., 1998; Webster et al., 1999; Villemant et al., 2008).

**Thermal and Geochemical Modeling of Sill Emplacement**

Magmas intruded into volatile-rich sediments, such as evaporites, can cause extensive outgassing as well as partial melting of the sediment (Aarnes et al., 2011a; Iacono-Marziano et al., 2012). To understand how the evaporites are affected by sill intrusions, a one-dimensional thermal model for sill emplacement into country rocks is developed, incorporating the thermal properties of pure halite (NaCl) and the thermal convection of magma (details of the model and the code may be found in the Supplementary Material). We note that most evaporites contain additional chemical components, which will affect the solidus temperature, as well as H2O (Grishina et al., 2018). We choose a host rock starting temperature of 50°C, but acknowledge that the host rocks may have been hotter, which would affect the results. The thermal modeling results using the parameters listed in Supplementary Material for thick (40 m) and thin (2 m) intrusions are shown in Figures 6A,B. Br and Cl loss was modeled based on the heating experiments carried out by Svensen et al. (2009) that show Br and Cl halocarbon formation in evaporites heated to 275°C. In reality, devolatilization might begin at lower temperatures, although very little published data exist for evaporite’s phase stability. Importantly, convection in magma has the consequence that heat transfer occurs faster. Thus, a 40 m thick sill emplaced at 1,200°C produces melting of a 7 m thick layer on the upper contact. In

**TABLE 3** Estimates of the proportion of chlorine (Cl) assimilated by the magmas that formed the whole rocks analysed in this study, using the method of Kendrick et al. (2013 (described in main text)). Sample Br/Cl and Cl contents (ppm) are shown (also shown in Table 1). Assimilated Cl (%) is estimated based on a mass balance, using values of Br/Cl of 2.8 ± 0.8 × 10⁻³ for the mantle and 0.3 × 10⁻³ for brines derived from evaporites (Kendrick et al., 2017).

| Location                  | Br/Cl | Cl, ppm | Assimilated Cl, % | Primary Cl, ppm |
|---------------------------|-------|---------|-------------------|-----------------|
| 1: Dyupkun Lake           | 4.3   | 92      | —                 | 92              |
|                           | 6.6   | 151     | —                 | 151             |
|                           | 7.9   | 63      | —                 | 63              |
| 2: Lower Tunguska River   | 4.3   | 325     | —                 | 325             |
|                           | 0.24  | 5331    | —                 | 172–281         |
|                           | 0.64  | 1589    | 82–89             | —               |
|                           | 0.68  | 2068    | 80–88             | 253–414         |
| 3: Nepa River borehole    | 1.3   | 3022    | 47–68             | 975–1591        |
|                           | 2.2   | 1852    | 0–39              | 1135–1852       |
|                           | 1.1   | 4143    | 58–74             | 1069–1744       |
|                           | 1.2   | 2307    | 53–71             | 670–1093        |
|                           | 1.6   | 3675    | 32–58             | 1541–2514       |
contrast, the aureole affected by Cl and Br devolatilization is 29 m thick on the upper contact and 23.5 m on the lower contact (Figure 6A). If the magma temperature is initially 1,100°C (i.e., 100°C lower), the molten salt layer on the top of the sill is 1.5 m thick (Figure 6C), and for a temperature of 1,090°C, no melting of the salt strata occurs (Figure 6D). There is no melting for a thin intrusion (2 m) even when the surrounding salt reaches the highest temperature of 1,200°C, and the thickness affected by devolatilization is 4.24 m on the top margin and 4.09 m on the bottom margin of the intrusion (Figure 6B). These models are consistent with previous estimates reporting the aureole thickness to be typically within 30–250% of the intrusion’s thickness (Aarnes et al., 2010). Moreover, it has been suggested that hydrocarbon generation by devolatilization may increase by nearly 35% in the case of two separate sills emplaced in organic-rich sediments with a vertical spacing of 5–10 times the intrusion thickness (Aarnes et al., 2011a), over the sum of the effects of two entirely separate sills.

Thermal modeling demonstrates that if the basalt temperature is < 1,100°C, melting of the evaporites is negligible. Therefore, congruent melting and assimilation of melted evaporite is not the mechanism leading to halogen enrichment in the intruded magma. Further, if the sill had assimilated evaporites by congruent melting, the sills would be enriched in all halogen species that are found in the evaporites (Cl, Br, and I), as well as Na and K, which is not observed (see whole rock compositions plotted in Supplementary Material Figure S2 and S3). Instead, we suggest that the geochemical trends are consistent with halogens being assimilated into the magma in the form of brine. Thermal models suggest that the metamorphic aureole within which devolatilization of halogens is possible may extend to several tens of meters for the case of discrete emplacement of a 40-m thick sill. In reality, the aureole’s devolatilized region will be larger than modeled as the evaporite is likely to contain salts with lower melting and devolatilization temperatures (i.e., sylvite, carnallite) (Grishina et al., 1992). Water is driven off the heated sediments, and halogens (and other elements) will partition into the hydrothermal fluids produced. The increasing Br content of the evaporites with decreasing distance from the dolerite contact, particularly at the lower contact (Figure 4; Table 2) may reflect the accumulation of heating-derived fluids in pore spaces in the salt, with the evaporites beneath the dolerite trapping more fluid as a consequence of the low permeability of the overlying dolerite, a pattern also observed in previous studies of total organic carbon content of shales intruded by dolerite sills (Aarnes et al., 2011b). At low pressures and high temperatures, the hydrothermal fluids produced during heating of evaporites will comprise a coexisting brine and vapor (Liebscher et al., 2006; Counou et al., 2009; Kendrick and Burnard, 2013). It is not well understood whether fractionation of Br and Cl occurs during the phase separation stage (Kendrick, 2018). Data from conjugate vapor and brine fluids from hydrothermal vents, for example, have similar Br/Cl ratios (Von Damm et al., 1997; Wu et al., 2012) which is consistent with experimental data (Berndt and Seyfried, 1997). We propose that the separated, low-density vapor phase migrated up towards the surface, perhaps through permeable country rocks or explosively through gas pipes, as was described and demonstrated experimentally by Svensen et al. (2009). A circulating hydrothermal system may have allowed the higher density brine to follow activity gradients to become assimilated into the sill, as was demonstrated for intrusions in Duluth, Minnesota, where the flow of hydrothermal brines into sills has been identified based on osmium isotopic variations (Ripley et al., 2001). Sill solidification may also be accompanied by the formation of pressure gradients from the aureole into the sill (Galerne et al., 2010; Svensen et al., 2010; Iyer et al., 2018). The brine’s ability to flow into the sill is supported by observations of veins and fissures in the sill filled in with salt precipitates 5–8 m from the contacts. In addition, high chlorine concentration on the rims of biotites in sills in the Lower Tunguska area have been interpreted as being due to chlorine transfer via a hydrous fluid (Heimdal et al., 2019; Callegaro et al., 2021) as observed for aureoles surrounding mafic sills at Norilsk (Pang et al., 2013).

The thermal modelling, combined with the geochemical data, suggests that many, if not all, samples in the set analysed here have been affected by assimilation of brines derived from heating evaporites in the subsurface prior to or during emplacement. Given these findings, we may use the geochemical data shown in Figure 3D to estimate the proportion of the Cl in the samples that was assimilated from a brine derived from heating evaporites. We use the method developed for estimating the amount of brine assimilation undergone by submarine magmas (Kendrick et al., 2013) whereby:

\[
\% Cl_{\text{assimilated}} = 100 \times \left( 1 - \frac{\left( \frac{Br}{Cl} \right)_{\text{bulk}} - \left( \frac{Br}{Cl} \right)_{\text{sample}}}{\left( \frac{Br}{Cl} \right)_{\text{bulk}} - \left( \frac{Br}{Cl} \right)_{\text{source}}} \right)
\]

The results of the analysis are shown in Table 3. The basalts from location 1 (Dyupkun Lake) may have been affected by degassing, so we do not discuss these further. The dolerites from the Lower Tunguska River (location 2) and the Nepa River borehole (location 3), likely assimilated between 0 and 89% of their chlorine contents during interaction with evaporite-derived brines. These upper values are similar to those estimated for the most contaminated submarine glasses from the Galápagos Spreading Centre and Lau Basin (~95%) (Kendrick et al., 2013). Using the observed Cl concentrations (Table 1), we may therefore estimate the “primary” Cl contents of the melts that formed these rocks (prior to Cl assimilation), which likely ranged from 172 to 2,514 ppm (Table 3).

CONCLUSION

We present whole-rock trace and halogen concentrations for a suite of samples from three locations in the Siberian Traps Large Igneous Province. The samples comprise basalt lava flows from Dyupkun Lake, dolerite sills from the Lower Tunguska River (intruded into organic-rich shales), and a dolerite sill intruded into evaporites from a borehole in the Nepa River region. The data show that the dolerites are enriched in Cl, similar to MORB.
and OIB that have been inferred to have assimilated seawater, and show halogen systematics which extend towards both evaporites and crustal brines. Furthermore, all analyzed samples show enrichment in Rb/Nb; with the dolerites also showing enrichment in CI/K similar to MORB and OIB that have been inferred to have assimilated seawater. We interpret the data to show that samples from all three locations have assimilated components of crustal sediments (which include evaporites) (Figure 7). We estimate that dolerites in our sample set have assimilated up to 89% of their Cl from fluids derived from evaporites. Primary Cl contents of the magmas that formed the whole rocks analysed here (prior to contamination) may have ranged from 172 to 2,514 ppm. Halogen assimilation from subcropping evaporites may be pervasive throughout the region, regardless of whether or not the magmas are emplaced within evaporites (Figure 7). The basalt lava flows from Dyupkun Lake have undergone degassing at atmospheric pressure, which has depleted their CI contents.

Thermal modeling shows that the intrusion of a sill of a few tens of meters thickness, active over timescales of a few years, causes the evaporites surrounding the sill to become depleted in halogens through devolatilization, whereby heating drives off water and halogens partition into a hot aqueous fluid. At low pressures in the crust, where we envisage much of the sill emplacement to have occurred, phase separation into a low-density vapor and a higher density brine would occur. Basaltic magmas in the sill may assimilate such brines, becoming enriched in halogens. Vapor-transported halogens may have been outgassed to the atmosphere (via pipes or a permeable fracture network). These mechanisms of magmatic halogen assimilation from evaporites have implications for the amounts of Br and Cl reaching the atmosphere during both intrusive (whereby magmatism is dominated by sill emplacement) and extrusive activity (dominated by surface lavas flows).

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author.

AUTHOR CONTRIBUTIONS

SS analysed the samples. All authors interpreted the data and co-wrote the manuscript.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/feart.2021.741447/full#supplementary-material.

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