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Potassium rich magmatism from a phlogopite-free source

Yu Wang ^{a,*}, Stephen F. Foley ^a and Dejan Prelević ^{b,c}

^a *ARC Centre of Excellence for Core to Crust Fluid Systems/GEMOC; Department of Earth and Planetary Sciences, Macquarie University, NSW 2109, Australia*

^b *Faculty of Mining and Geology, Belgrade University, Đušina 7, 11000 Belgrade, Serbia*

^c *Institute of Geosciences, University of Mainz, Becherweg 21, Mainz 55099, Germany*

SUPPLEMENTARY FILE

This file contains three figures and six tables.

ANALYTICAL METHODS

Major elements: Whole-rock major element compositions of the phyllite and dunite were determined by X-ray fluorescence spectrometry (XRF) using a Philips MagiXPRO spectrometer on fused discs. Major element compositions of melts and minerals were obtained by CAMECA SX-100 electron microprobe (EMP) at the Department of Earth and Planetary Sciences, Macquarie University, with operating conditions of generally 15 kV accelerating voltage and 20 nA beam current. Counting times were 10 s for peaks, and 5 s for each background. The peaks were calibrated by measurements of standards (natural minerals and synthetic oxides). For mineral phases a beam diameter of 1 μm was used, which was increased to 5 μm for the analysis of quenched melt areas.

Trace elements: Trace elements were analysed by laser ablation-inductively coupled plasma-mass spectrometry (LA-ICP-MS) at Macquarie University. For phyllite and dunite, rock powders were melted to form glass beads without any fluxing agent on an iridium strip heater in an argon atmosphere and analyzed by laser sampling of the glasses. All analyses used 266 nm Nd-YAG laser coupled to an Agilent 7500ce ICP-MS. The laser was operated at a repetition rate of 5 Hz, with laser energy at the sample site of 5–6 J/cm², allowing data collection from individual grains in polished thick sections (up to 80 μm thickness) for at least 40 s. Nitrogen was mixed with helium before ablated materials were introduced into the ICP-MS. Data were measured for 60 s plus 30 s background with beam sizes of 30 μm and 50 μm for most samples, 80 μm for some large-grained melt pools, 15 μm and 20 μm for a few small-grained accessory phases and melt samples from experiments. ²⁹Si and ⁴⁴Ca were used as the internal standards applying the Si and Ca concentrations previously determined by electron microprobe. For calibration, NIST SRM 610 was analysed at the beginning and after every 20 measurements on the unknown samples. The time-resolved signal was processed using the program GLITTER 4.4.1 (www.glitter-gemoc.com, Macquarie University), applying the preferred values for NIST SRM 610 reported in the GeoReM database (<http://georem.mpch-mainz.gwdg.de/>) (Jochum et al., 2011). Analytical uncertainty (one sigma) for a spot analysis or line scan was less than 10 %. During each run the basaltic USGS BCR-2G reference glass was analysed as an unknown to monitor accuracy and reproducibility of the analyses.

EXPERIMENTAL METHOD

All experiments were performed in a 0.5-inch piston-cylinder apparatus based on a modified Boyd-England design at Macquarie University (Boyd and England, 1960). Furnace

assemblies (12.7 mm diameter) were made of graphite with cylindrical talc outer sleeve and pyrex® inner sleeve and BN inserts. Temperatures were measured using Pt–Pt₉₀Rh₁₀ thermocouples and automatically controlled by an Electromax V single loop controller. Samples were welded into a 4 by 4 mm Ag₇₀/Pd₃₀ capsule with a wall thickness of 0.2 mm, minimizing iron loss to the capsule. Experimental conditions were applied by first applying pressure, followed by heating until the desired pressure and temperature were reached. Run durations were up to 192 hours. Where present, melt remained free of quench crystallization. Recovered capsules were cut into two parallel to their axis and embedded into epoxy mounts for polishing. Melt proportions in all experimental runs were obtained by program ImageJ using SEM image and mass balance calculation.

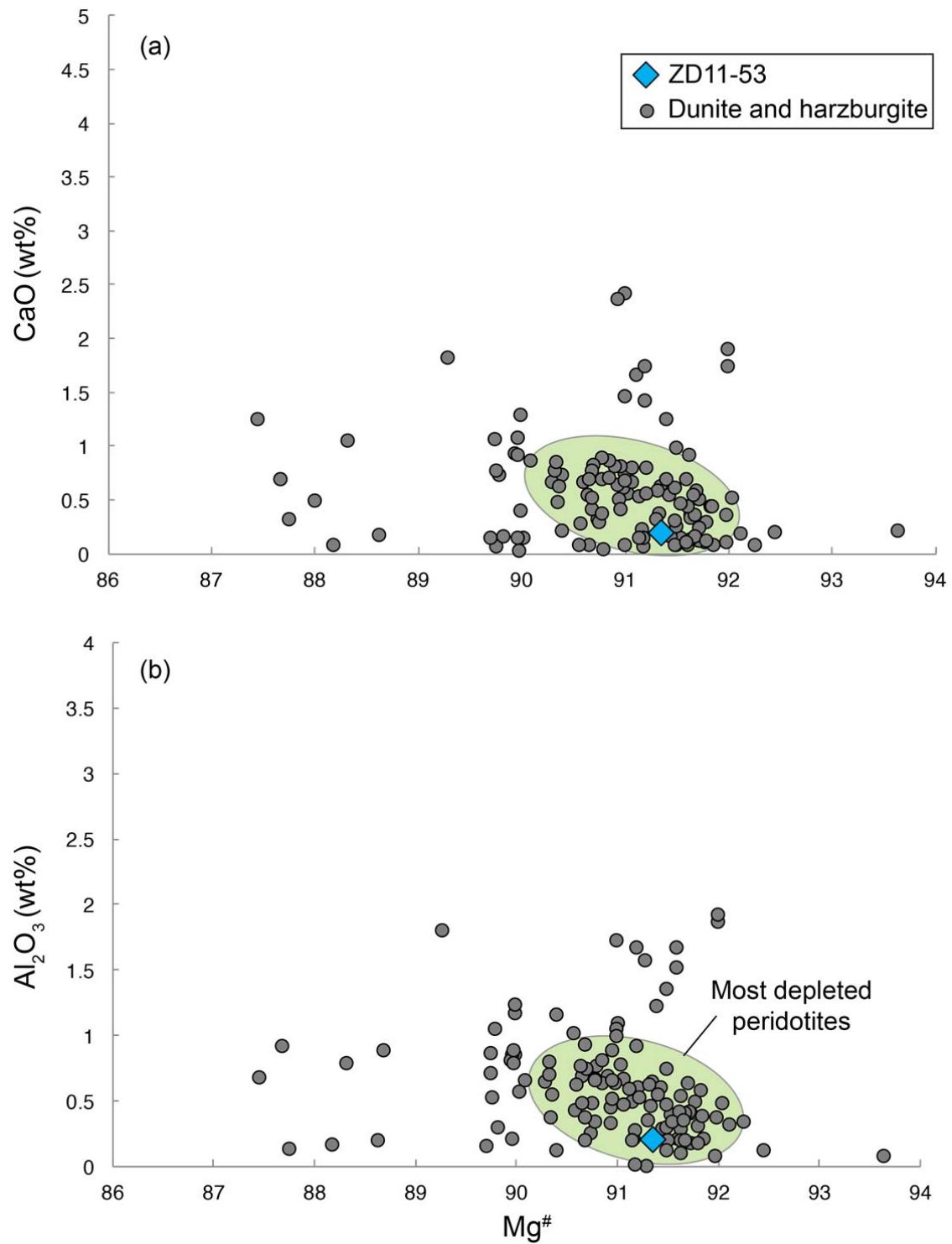


Fig. DR1 Compositions of Mg[#] vs. (a) CaO (wt%) and (b) Al₂O₃ (wt%) of the dunite sample used in this study (ZD11-53) compared to natural depleted peridotites (Ishii et al., 1992; Parkinson and Pearce, 1998; Ohara and Ishii, 1998; Pearce, 1999; Khedr and Arai, 2010; Nozaka, 2014). Mg[#] = 100*MgO/(MgO + FeO) in mole per cent.

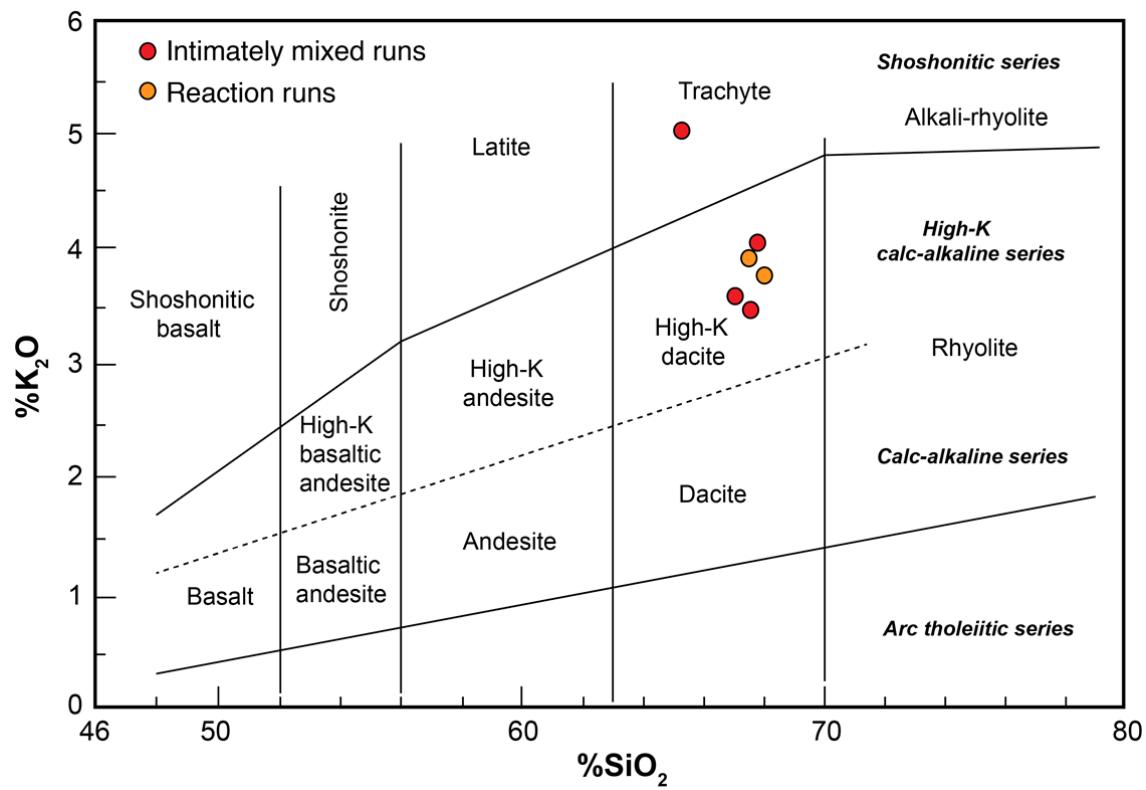


Fig. DR2 SiO₂ (wt %) vs. K₂O (wt %) classification plot for melts produced in this study. Adapted from Peccerillo & Taylor, 1976.

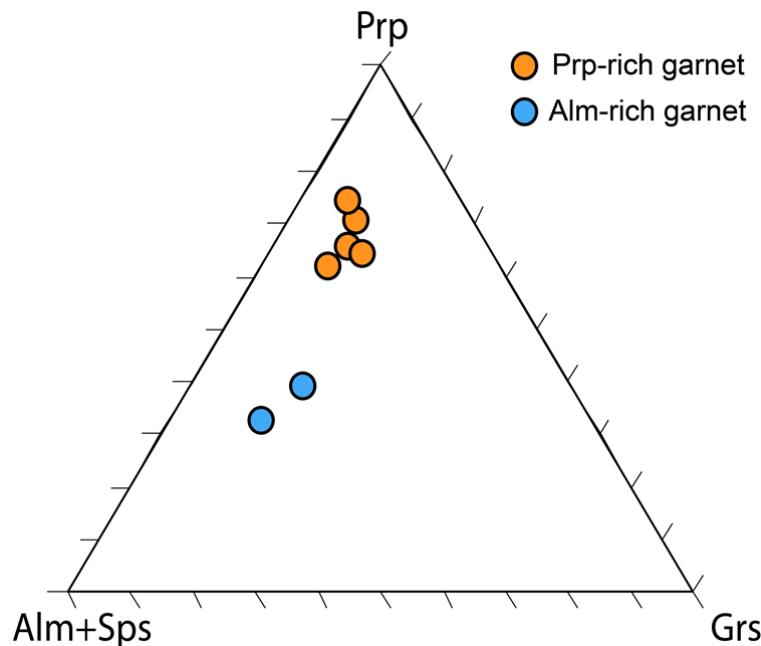


Fig. DR3 Ternary discrimination diagram of garnets produced from the experiments. Note the two compositionally different types. Prp: pyrope; Alm: almandine; Sps: spessartine; Grs: grossular.

TABLE DR1. MAJOR (WT%) AND TRACE (PPM) ELEMENT COMPOSITIONS OF STARTING MATERIALS

Locality	Kopaonik area, Serbia			Zedang area, South Tibet, China
Sample	11KP01	GLOSS	UCC	ZD11-53
Rock	Quartz-phyllite			Dunite
SiO ₂	76.7	58.57	66	42.16
TiO ₂	0.7	0.62	0.5	0.04
Al ₂ O ₃	11.6	11.91	15.2	0.20
FeO _T	3.4	5.21	4.5	8.35
MnO	0.06	0.32	0.08	0.13
MgO	1.1	2.48	2.2	49.53
CaO	1.6	5.95	4.2	0.19
Na ₂ O	1.6	2.43	3.9	0.07
K ₂ O	1.9	2.04	3.4	0.00
Cr ₂ O ₃	0.01	NA	NA	0.36
LOI	1.5			0.25
Sum	100.27			101.28
Mg [#]	37	46	47	91
Li	33	NA	20	2.2
Sc	13	13.1	13.6	4.0
Ti	4269	NA	2458	37
V	58	110	107	16
Cr	1362	78.9	83	1984
Mn	1147	NA	600	1011
Co	11	21.9	17	129
Ga	10	NA	17	0.43
Rb	54	57.2	112	0.003
Sr	288	327	350	0.30
Y	20	29.8	22	0.07
Zr	56	130	190	0.02
Nb	9.7	8.94	12	0.016
Cs	1.6	3.48	4.6	0.003
Ba	355	776	550	0.50
La	30	28.8	30	0.005
Ce	63	57.3	64	0.003
Pr	7.2	NA	7.1	0.001
Nd	28	27	26	0.003
Sm	5.6	5.78	4.5	0.001
Eu	1.1	1.31	0.88	0.001
Gd	4.6	5.26	3.8	0.003
Tb	0.68	NA	0.64	0.000
Dy	3.8	4.99	3.5	0.007
Ho	0.8	NA	0.8	0.002
Er	2.1	2.92	2.3	0.011
Tm	0.31	NA	0.33	NA
Yb	2	2.76	2.2	0.021
Lu	0.26	0.413	0.32	0.005
Hf	1.6	4.06	5.8	0.001
Ta	0.75	0.63	1.0	0.003
Th	8.4	6.91	10.7	0.001
U	1.5	1.68	2.8	0.008

Global Subducted Sediment (GLOSS) from [Plank and Langmuir \(1998\)](#); Upper Continental Crust (UCC) from [Taylor and McLennan \(1995\)](#) and [McLennan \(2001\)](#).

TABLE DR2. □EXPERIMENTAL RUN CONDITIONS AND PRODUCTS.

Run No.	Geometry	P (GPa)	T (°C)	T _C (°C)	Hours	Phases (mode %)
2036	Reaction	2	1000	883	192	Dunite half: Ol (96), Cpx (2), <i>Chr</i> Phyllite half: M (42), Qrz (41), Alm-grt (12), Ky (2), <i>Zir</i> , <i>Apa</i> , <i>Rt</i> Interaction belt: Opx (2), <i>Prp-grt</i>
2037	Mixture	2	1000	933	192	Opx (75), M (23), Ol (5.5), <i>Chr</i> , <i>Prp-grt</i> , <i>Zir</i> , <i>Apa</i> , <i>Rt</i>
2038	Mixture	2	1100	942	168	Opx (71), M (28), <i>Chr</i>
139	Reaction	3	1000	1003	192	Dunite half: Ol (93), Chr (2), <i>Cpx</i> Phyllite half: M (31), Coe (44), Alm-grt (14), Ky (6), <i>Zir</i> , <i>Apa</i> , <i>Rt</i> Interaction belt: Opx (4), <i>Prp-grt</i>
140	Mixture	3	1000	871	192	Opx (71), M (14.5), Prp-grt (6), Coe (6), Cpx (1.5), <i>Zir</i> , <i>Chr</i>
141	Mixture	3	1100	936	168	Opx (66), M (18), Prp-grt (8), Coe (5), Cpx (2), <i>Zir</i> , <i>Chr</i>

Note: Phases in italics are accessory phases present in the run products in the range of 0-1 vol%. T_C denotes temperatures calculated by the two-pyroxene thermometer of [Brey and Köhler, 1990](#), with error of $\pm 16^{\circ}\text{C}$. Mineral abbreviations: M, melt; Ol, olivine; Qrz, quartz; Alm-grt, almandine-rich garnet; Prp-grt, pyrope-rich garnet; Opx, orthopyroxene; Cpx, clinopyroxene; Chr, chromite; Coe, coesite; Ky, kyanite; Zir, zircon; Apa, apatite; Rt, rutile.

TABLE DR3. MAJOR ELEMENT COMPOSITIONS OF MELT (WT%)

Run No.	2036		2037		2038		139		140		141	
P (GPa)	2		2		2		3		3		3	
T (°C)	1000		1000		1100		1000		1000		1100	
Geometry	Reaction		Mixture		Mixture		Reaction		Mixture		Mixture	
N (Number of analyses)	10	1 SD	10	1 SD	11	1 SD	5	1 SD	5	1 SD	3	1 SD
SiO ₂	68.54	0.80	67.16	0.49	67.89	0.33	68.21	0.73	68.46	0.68	65.50	0.87
TiO ₂	0.53	0.02	0.64	0.02	0.79	0.02	0.39	0.01	0.54	0.01	0.62	0.04
Al ₂ O ₃	15.84	0.52	14.88	0.41	14.44	0.09	14.21	0.42	13.15	0.10	13.33	0.16
Cr ₂ O ₃	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
FeO	1.02	0.08	0.98	0.07	1.52	0.04	0.68	0.07	0.61	0.04	1.03	0.18
MnO	0.00	0.00	0.04	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
MgO	0.56	0.16	1.47	0.63	1.91	0.08	0.35	0.03	0.59	0.04	0.84	0.13
CaO	1.62	0.03	1.78	0.07	1.97	0.03	0.86	0.02	0.46	0.02	0.78	0.07
Na ₂ O	2.43	0.35	2.31	0.19	2.20	0.32	1.39	0.49	0.92	0.21	2.26	0.25
K ₂ O	3.84	0.11	3.52	0.11	3.38	0.13	4.08	0.38	4.22	0.46	5.09	0.49
Total	94.37		92.79		94.1		90.18		88.94		89.45	
Mg#	50		73		69		48		64		59	
A/(CNK)	1.43		1.37		1.33		1.72		1.90		1.25	
Degree of melting (%)	42		23		28		31		14.5		18	

Mg# = 100*MgO/(MgO + FeO) in mole per cent;

A/(CNK) = Aluminum saturation index, defined as the molecular ratio Al₂O₃ / (CaO + Na₂O + K₂O);

1 SD = one standard deviation, N refers to number of replicate analyses.

TABLE DR4. MAJOR ELEMENT COMPOSITIONS OF GARNET (WT%)

Run No.	2036	2037				139				140				141	
P (GPa)	2	2				3				3				3	
T (°C)	1000	1000				1000				1000				1100	
Geometry	Reaction	Mixture				Reaction				Mixture				Mixture	
N (Number of analyses)	3	1 SD	2	1 SD	6	1 SD	3	1 SD	2	1 SD	5	1 SD	11	1 SD	
Type	Almandine-rich	Pyrope-rich			Pyrope-rich	Almandine-rich			Pyrope-rich	Pyrope-rich			Pyrope-rich	Pyrope-rich	
SiO ₂	39.41	1.06	41.31	0.34	41.61	0.58	38.49	0.26	42.33	0.64	42.32	0.78	42.60	0.99	
TiO ₂	1.11	0.24	1.05	0.26	0.91	0.37	1.27	0.42	0.73	0.42	0.81	0.05	0.60	0.32	
Al ₂ O ₃	22.33	0.16	22.95	0.04	22.17	0.72	21.33	0.56	22.26	0.01	22.50	0.56	22.95	0.89	
Cr ₂ O ₃	0.00	0.00	0.13	0.00	0.36	0.40	0.00	0.00	0.08	0.00	0.06	0.00	0.19	0.09	
FeO	19.83	0.92	11.74	0.77	13.82	1.11	24.47	1.18	11.18	0.91	10.64	0.12	9.31	0.66	
MnO	0.52	0.08	0.24	0.01	0.35	0.04	0.46	0.06	0.31	0.01	0.30	0.03	0.26	0.07	
MgO	10.91	0.82	18.05	0.39	17.33	0.86	8.53	0.33	18.23	0.54	19.58	0.40	22.08	0.58	
CaO	6.60	0.32	4.90	0.45	4.35	0.38	5.44	0.51	5.77	0.48	4.49	0.33	2.71	0.70	
Na ₂ O	0.09	0.02	0.13	0.01	0.06	0.02	0.22	0.07	0.09	0.01	0.24	0.04	0.11	0.04	
K ₂ O	0.05	0.02	0.00	0.00	0.06	0.00	0.04	0.01	0.03	0.00	0.08	0.01	0.05	0.01	
NiO	0.00	0.00	0.00	0.00	0.05	0.00	0.00	0.00	0.00	0.00	0.06	0.00	0.06	0.00	
P ₂ O ₅	0.19	0.06	0.00	0.00	0.14	0.00	0.15	0.06	0.00	0.00	0.20	0.08	0.16	0.09	
Total	101.1	100.5			101.2	100.4			101.0	101.3			101.1		
Pyrope	40%	64%			61%	32%			63%	68%			75%		
Almandine	41%	23%			27%	52%			22%	21%			18%		
Grossular	18%	12%			11%	15%			14%	11%			7%		
Spessartine	1%	0%			1%	1%			1%	1%			1%		

TABLE DR5. MAJOR ELEMENT COMPOSITIONS OF PYROXENES (WT%)

Run No.	2036	2037	2038	139	140	141	2036	139	140	141
P (GPa)	2	2	2	3	3	3	2	3	3	3
T (°C)	1000	1000	1100	1000	1000	1100	1000	1000	1000	1100
Geometry	R	M	M	R	M	M	R	R	M	M
N	5	1 SD	9	1 SD	10	1 SD	6	1 SD	11	1 SD
Phase	opx	opx	opx	opx	opx	opx	cpx	cpx	cpx	cpx
SiO ₂	55.68	0.75	56.66	0.53	56.08	0.50	56.78	0.54	57.75	0.64
TiO ₂	0.23	0.05	0.20	0.03	0.21	0.04	0.07	0.04	0.13	0.03
Al ₂ O ₃	3.81	0.43	2.76	0.61	3.20	0.35	1.88	0.52	1.95	0.64
Cr ₂ O ₃	0.10	0.04	0.15	0.07	0.21	0.09	0.39	0.28	0.00	0.00
FeO	6.70	0.72	7.32	0.47	7.11	0.46	6.63	0.81	6.32	0.86
MnO	0.10	0.01	0.12	0.01	0.12	0.01	0.14	0.03	0.08	0.02
MgO	32.82	0.90	32.07	0.79	31.92	0.61	33.50	0.72	33.12	0.53
CaO	0.42	0.12	0.53	0.18	0.56	0.10	0.60	0.14	0.21	0.07
Na ₂ O	0.05	0.01	0.09	0.03	0.11	0.03	0.11	0.10	0.21	0.12
K ₂ O	0.00	0.00	0.03	0.01	0.11	0.12	0.00	0.00	0.11	0.09
Total	99.91		99.93		99.62		100.1		99.88	
Wollastonite	1%		1%		1%		0%		1%	
Enstatite	89%		87%		87%		88%		89%	
Ferrosilite	10%		11%		11%		10%		10%	
Aegirine/Acmite	0%		0%		0%		1%		1%	

Note: R: Reaction; M: Mixture.

TABLE DR6. TRACE ELEMENT COMPOSITIONS OF MELT (PPM)

Run No.	2036	2037		2038		139		140		141	
P (GPa)	2	2		2		3		3		3	
T (°C)	1000	1000		1100		1000		1000		1100	
Geometry	R	M		M		R		M		M	
N	7	1 SD	6	1 SD	9	1 SD	6	1 SD	9	1 SD	8
Li	86	6.3	33	1.9	42	2.9	82	4.7	26	3.1	23
Be	4.5	0.32	2.1	0.43	2.8	0.37	4.9	0.62	4.1	0.52	3.0
Sc	4.2	0.49	6.9	0.84	7.1	0.86	3.6	0.55	4.2	0.56	5.0
Ti	2950	138	2653	185	3654	277	2764	253	2574	304	3089
V	9.1	1.5	29	2.3	17	3.1	19	2.1	17	1.5	18
Cr	70	16	640	79	378	61	12	1.8	398	43	221
Mn	109	21	508	87	381	45	107	27	272	25	304
Co	3.6	0.61	43	8.6	27	2.8	2.7	0.38	25	3.1	21
Ni	7.9	1.2	708	82	287	39	155	24	479	45	243
Cu	2.8	0.44	1.2	0.31	2.1	0.27	3.0	0.39	5.7	0.57	2.0
Zn	21	3.8	25	3.2	17	1.9	31	4.0	50	4.4	10
Ga	23	2.0	9.3	1.5	9.2	1.2	27	2.6	23	2.7	14
Rb	167	9.1	99	8.9	119	23	206	14	219	29	191
Sr	294	16	167	13	209	24	346	25	336	37	341
Y	2.6	0.52	14	2.9	19	2.7	3.9	0.22	6.7	1.1	5.1
Zr	212	7.3	168	11	327	45	198	18	235	30	354
Nb	11	1.6	11	2.3	17	2.4	11	1.7	14	1.3	22
Cs	5.4	0.54	3.5	0.46	3.9	0.50	6.8	0.77	7.2	1.1	5.6
Ba	698	39	402	27	495	46	841	62	871	98	792
La	49	4.9	26	3.0	33	3.8	46	3.6	46	5.6	62
Ce	105	10	62	6.4	76	8.2	99	7.9	100	13	96
Pr	11	1.1	6.8	0.89	8.5	1.7	10	0.89	11	1.9	10
Nd	34	3.2	24	2.8	32	4.6	29	3.3	34	3.8	37
Sm	4.2	0.63	4.9	0.76	6.2	1.1	3.3	0.63	3.6	0.60	5.5
Eu	0.85	0.12	1.1	0.21	1.3	0.37	0.68	0.17	0.63	0.20	1.2
Gd	1.9	0.32	3.6	0.44	4.7	0.58	1.2	0.26	2.3	0.33	2.5
Tb	0.17	0.05	0.47	0.17	0.62	0.16	0.16	0.03	0.33	0.16	0.28
Dy	0.7	0.13	3.0	0.15	4.0	0.59	0.93	0.13	1.4	0.48	1.3
Ho	0.09	0.04	0.53	0.09	0.73	0.15	0.13	0.05	0.23	0.07	0.19
Er	0.17	0.09	1.6	0.32	2.0	0.47	0.47	0.04	0.58	0.12	0.41
Tm	0.03	0.01	0.18	0.05	0.25	0.03	0.10	0.03	0.10	0.05	0.06
Yb	0.17	0.06	1.2	0.24	1.6	0.34	0.55	0.08	0.56	0.09	0.31
Lu	0.02	0.01	0.12	0.03	0.27	0.04	0.10	0.02	0.1	0.03	0.07
Hf	6.3	0.78	4.0	0.66	8.4	1.1	4.8	0.69	6.0	0.77	8.8
Ta	1.1	0.22	0.93	0.23	1.4	0.29	1.0	0.09	1.4	0.17	1.9
Th	23	3.3	12	1.9	15	1.0	20	1.7	22	2.5	18
U	5.0	0.45	3.5	0.26	3.9	0.34	4.6	0.36	5.5	0.53	4.4
Th/La	0.47	0.08	0.46	0.09	0.45	0.06	0.43	0.05	0.48	0.08	0.29
Rb/Cs	31	3.5	28	4.5	31	7.1	30	4.0	30	6.1	34
Sm/La	0.09	0.02	0.19	0.04	0.19	0.04	0.07	0.01	0.08	0.02	0.09
Sr/Nd	8.6	0.94	7.0	0.98	6.5	1.2	12	1.6	9.9	1.6	9.2
Nb/Ta	10	2.5	12	3.8	12	3.0	11	2.0	10	1.5	12
Zr/Hf	34	4.3	42	7.5	39	7.4	41	7.1	39	7.1	40

Note: N: number of analyses; R: Reaction; M: Mixture.

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