

THE DETAILED CHEMISTRY OF THE REACTION  $SP + PX = GT + OL$  IN PYROXENITE  
INCLUSIONS FROM SALT LAKE CRATER, HAWAII

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In most of the garnet pyroxenite inclusions at Salt Lake Crater, Hawaii, garnet is not a primary mineral but rather has formed during the cooling of an assemblage of aluminous pyroxene(s) and spinel (Green, 1966). The garnet occurs mainly as reaction rims between spinel and clinopyroxene; it may also occur as directly exsolved blebs in certain occasional grains of clinopyroxene. The goal of this study has been to reconstruct the modal proportions and chemical compositions of the pre-garnet mineralogy in these rocks, and thereby identify the components of the initial pyroxenes and spinels which have contributed to the formation of the garnet and olivine in the rock. For each major element determined by microprobe analysis in the phases of a given rock, a mass-balance equation can be written:

$$a [e_i]_{SPI} + b [e_i]_{CPXI} + c [e_i]_{OPXI} = d [e_i]_{GTII} + e [e_i]_{OLVII} + f [e_i]_{OPXII} + g [e_i]_{CPXII} + h [e_i]_{SPII}$$

(where I refers to the initial, pre-garnet rock and II refers to the present analysed rock. a through h are modal proportions (wt.%) and  $e_i$  refers to one of the eight analysed elements. All of the terms on the right side of the equation are known, leaving a system of eight equations and at the outset, 27 unknowns.)

A number of additional compositional and textural constraints exist. Even in a single rock, spinels are surrounded by garnet coronas of differing relative size, and show a systematic linear variation in composition as a function of the degree to which the garnet-forming reaction has proceeded. (As the reaction progresses the remnant spinel becomes richer in Cr, Fe, Ti, and Mn and poorer in Al and Mg.) Although the reasons for this variation are unclear, one can extrapolate these trends to  $GT/SP = 0$ , and closely fix the composition of the pre-garnet spinel (SPI). The modal proportion of SPI is determined by knowing the compositions of SPI and SPII, the modal proportion of SPII, and by the observation that to a first order approximation the initial spinel's Cr budget simply becomes concentrated in the eventual, smaller, garnet-rimmed spinel. (In such rock studied in detail, SLC8,  $Cr_2O_3SP/Cr_2O_3GT$  ranges from 10 for low Cr spinels (2.3% $Cr_2O_3$ ) to 22 for spinels with about 9%  $Cr_2O_3$ ;  $Cr_2O_3SP/Cr_2O_3CPX$  ranges from 18 to 47 over the same range).

The combined composition and modal proportion of CPXI plus OPXI are thus determined, allowing a balanced relation, for SLC8, to be written:

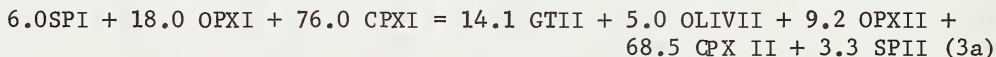


or equivalently



where SPR and PXR refer to the components contributed to the reaction by the initial spinel and combined pyroxene. If just prior to garnet formation, the rock had simply consisted of SPI + a single subcalcic pyroxene (combination of OPXI + CPXI), PXR would be the actual contribution from that pyroxene. Textures indicate however that such a single pyroxene, if it ever existed, had exsolved to discrete grains of opx and cpx prior to garnet formation.

An approximate composition for CPXI is obtained by the recombination of CPXII and GTII, in those isolated clinopyroxenes showing exsolved garnet. This composition (particularly its  $\text{Al}_2\text{O}_3$ ) is very sensitive to the proportions of CPXII and GTII used in the vicinity of the modally determined values. The best values for the compositions and modal abundances for both CPXI and hence for OPXI are obtained by choosing a CPXII/GTII ratio which allows the ratio  $\text{Al}_2\text{O}_3\text{CPXI}/\text{Al}_2\text{O}_3\text{OPXI}$  to be 1.5, a value consistently found in aluminous spinel lherzolites at Salt Lake Crater and Kilbourne Hole. (See table I for compositional data.) Equations (2a) and (2b) can now be rewritten:



and



Two interesting geophysical results follow from these data. (1) As predicted by MacGregor's experimental study of the effects of Cr content on the P-T position of the spinel-garnet boundary (MacGregor, 1970), the increasing Cr content of the garnet-rimmed spinels as the reaction proceeds indicates that spinel and garnet should coexist over a certain range of pressures for any given bulk composition, and that even if the upper mantle were regionally homogeneous with respect to  $\text{Cr}/(\text{Cr} + \text{Al})$ , one would not expect a sharp boundary or density contrast to exist at any depth. (2) It should be noted that the dominant contribution of CPXI to the garnet forming process is high in Mg (23.20%) and low in Ca (9.78%). This suggests that caution is needed when comparing apparent equilibration temperatures for garnet-bearing and garnet-free rocks using the simple En-Di solvus for clinopyroxene temperatures.

Table I

	SPI	SPII	SPR	CPXI	CPXII	CPXR
SiO <sub>2</sub>	.00	.00	.00	50.71	51.97	42.86
TiO <sub>2</sub>	.30	.23	.38	.74	.88	.00
Al <sub>2</sub> O <sub>3</sub>	64.00	62.90	65.31	7.36	6.80	13.04
Cr <sub>2</sub> O <sub>3</sub>	1.50	2.75	.00	.17	.16	.27
FeO	12.10	12.50	11.62	4.39	3.74	10.68
MnO	.10	.09	.11	.14	.10	.51
MgO	21.70	21.32	22.15	15.54	14.83	23.20
CaO	.00	.00	.00	18.60	19.71	9.78
Na <sub>2</sub> O	<u>.00</u> 99.70	<u>.00</u> 99.84	<u>.00</u> 99.57	<u>1.59</u> 99.24	<u>1.74</u> 99.95	<u>.34</u> 100.68
	OPXI	OPXII	OPXR	GTII	OLIVII	
SiO <sub>2</sub>	54.38	55.30	53.33	41.87	40.78	
TiO <sub>2</sub>	.34	.17	.51	.14	.00	
Al <sub>2</sub> O <sub>3</sub>	5.42	4.30	6.59	23.70	.00	
Cr <sub>2</sub> O <sub>3</sub>	.09	.10	.08	.16	.00	
FeO	7.24	6.80	7.70	8.93	11.18	
MnO	.12	.11	.14	.33	.14	
MgO	31.52	31.99	30.99	19.81	47.72	
CaO	.67	.66	.66	4.91	.08	
Na <sub>2</sub> O	<u>.06</u> 99.84	<u>.11</u> 99.55	<u>.12</u> 100.12	<u>.00</u> 99.91	<u>.00</u> 99.97	

## References:

- (1) Green, D.H. (1966) The Origin of the "eclogites" from Salt Lake Crater, Hawaii, EPSL, 1, 414-420.
- (2) MacGregor, I. (1970) The effect of CaO, Cr<sub>2</sub>O<sub>3</sub>, Fe<sub>2</sub>O<sub>3</sub> and Al<sub>2</sub>O<sub>3</sub> on the stability of spinel and garnet periodites, Phys. Earth Planet. Interiors, 3, 372-377.