PARENT MAGMAS OF SNC HARZBURGITES: PHASE EQUILIBRIA MODELING

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The clan of SNC meteorites includes two igneous harzburgites - ALHA77005 and LEW88516, both having similar textures and mineralogy [1,2]. A distinctive feature of these rocks is the presence of large (up to few mm) low-Ca pyroxene oikocrysts (hereafter Opx) enclosing olivine (OI) and chromite (Chr). Mineral assemblage in non-poikilitic areas is represented by maskelynite, Ol, Opx, high-Ca pyroxenes (Aug), Chr, ilmenite, and minor phosphates (±sulfides). OI in ALH is almost uniform in composition (Fo₇₀₋₇₃) [2-4], whereas olivines in LEW are more iron-rich ranging from Fo₆₄ to Fo₇₀ [1,4]. Pyroxene compositions display similar relations. Based on the textural and mineralogical data, the SNC harzburgites have been interpreted to be igneous cumulates solidified from primary magma mixtures of crystals (Ol+Chr±Opx) cumulus with intercumulus liquid [2.4]. Despite some doubts that the intercumulus minerals were crystallized as a closed system [5], few attempts have been estimate the parent magma done to compositions (PMC) for ALH using its bulk rock and mineral compositions [2,5], as well for LEW the components from magmatic usina inclusions [1]. These attempts, however, did not result in an unambiguous solution, mainly because of the lack of constraints following from phase equilibria. Knowledge of the phase equilibria for the natural cumulates could allow one to estimate more correctly the range of trapped liquid compositions, and finally to make it narrower the range of parent magmas to be searched.

Calculating phase equilibria for SNC harzburgites. To accomplish the goal, we used the METEOMOD program designed for the calculations of melting-crystallization relationships in iron-rich meteoritic igneous systems [6]. The basic block of METEOMOD is a set of empirically calibrated equations, which describe equilibria between silicate melt and minerals such as OI, Opx, Aug, and PI in terms of temperature, $f_{\Omega 2}$, and liquid compositions. crystallization The program calculates sequences step by step, as the total amount of crystals is increased. The accuracy of the calculations for temperature is of ±10-15°C, with the contents of major end-members in the minerals calculated within ±1-2 mol% [6].



Using METEOMOD we calculated the course equilibrium crystalliof zation for 2 melts corresponding to the bulk compositions of ALH [2] and LEW [4]. These calculations were carried out at 1 atm, with the crystal increment of 1%, at WM buffer, and in the range of meltina 100<*F*<20 wt%. Results the modeling of are shown in Fig.1. Compositions for some of the residual melts are listed in the Table.

Fig.1. Modeled crystallization sequences

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These compositions can be used to estimate the **PMC** for ALH and LEW, if average compositions of their cumulus minerals were similar to those which should be in equilibrium with the trapped melt. The main problem of accurate defining the **PMC** along the calculated liquid lines of descent is the search for additional constraints following from primary cumulus mineral-melt equilibria. These constraints may include proportions or/and compositions of initial cumulus minerals.

The first method, based on the modal mineralogy analysis, results in too large differences between the estimated *OI* proportion for ALH (45.8 wt% [2,5] and 60.2% [4]). The problem of mafic mineral reequilibration during a post-cumulus process was also widely discussed in the literature. An alternative approach includes

Table. Calculated mineral-melt compositions and phase proportions in terms of the temperature

Component,	LEW88516 (bulk)		ALHA77005 (bulk)		
wt%	1135.6°C	1226.2°C	1140.3°C	1229.6°C	1336.5 °C
SiO ₂	50.45	51.03	50.16	51.53	47.66
TiO ₂	1.07	0.67	1.94	1.30	0.99
Al ₂ O ₃	12.68	7.35	12.88	9.33	6.98
FeO	14.17	17.99	13.10	16.32	21.10
MnO	0.36	0.47	0.34	0.43	0.53
MgO	5.93	9.33	6.32	9.19	13.51
CaO	10.58	10.60	10.97	9.14	7.16
Na ₂ O	2.28	1.23	2.15	1.43	1.07
K ₂ O	0.14	0.08	0.19	0.12	0.09
P_2O_5	2.33	1.25	1.96	1.22	0.91
Total	100.0	100.0	100.0	100.0	100.0
Mg/(Mg+Fe)	0.427	0.480	0.462	0.501	0.533
Ca/(Ca+Al)	0.431	0.567	0.436	0.471	0.483
Calculated proportions of melts and minerals, wt%					
Melt	21.0	39.1	21.2	34.1	45.5
01	54.4	53.9	64.6	65.6	54.5
Орх	15.4	7.0	12.1	0.3	-
Aug	9.2	-	0.9	-	-
PI	-	-	1.1	-	-
Calculated mineral compositions, mol%					
Fo (Ol)	69.1	73.0	72.2	74.8	78.5
En-Wo (Opx)	69.3-6.6	74.8-3.5	71.4-6.7	76.3-3.1	-
En-Wo (Aug)	50.4-33.5	-	51.7-34.3	-	-
An (Pl)	-	-	74	-	-

Longhi&Pan,1989 (T=1240 °C) Bartels&Grove,1991 Al2O3 in melt, wt% 12 1.0.01872 1140 1180 8 Al2O3 in Chromite cores in ALHA77005 Chr, wt% 0 Т Т Т Т Т Т Т Т 0 4 8 12 16 20 24

Fig. 2. Experimental Chromite and Glass compositions

the use of chromite core compositions.

Constraints from spinel compositions.

The chromite cores in SNC harzburgites were found to contain small amounts of Al₂O₃, mostly in the range of 6-9 wt.% [3,4,8]. Experimental data obtained igneous eucrite/SNC in systems indicate that such spinels could be crystallized from the only melts containing no more than 8 wt% Al₂O₃ [7,8]. In fact, there is a strong dependence between the observed spinel and experimental glass compositions (Fig. 2), which could be used to predict alumina content in the trapped melt (PMC?), if an composition average of chromite cores is known. For example, if the Chr cores in ALH contain in average of about 7.5% Al₂O₃, one can estimate the alumina content

in its trapped melt to be of 7 wt.%. Such a melt corresponds to the temperature of 1340° C (see Table). More detail studies of Chr chemistry are necessary to estimate the *PMC* for LEW.

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