

TABLE CAPE VENT XENOLITH SUITE, NORTHWEST TASMANIA: MINERALOGY AND IMPLICATIONS FOR CRUST-MANTLE LITHOLOGY AND MIOCENE GEOTHERMS IN TASMANIA

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(with one plate, two text-figures and eight tables)

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The Miocene Table Cape vent erupted a diverse mantle-crust xenolith suite within its fractionated nephelinitic matrix. Assemblages include mantle metaperidotites, garnet-metawebsterites and rarer garnet-metaproxenites, garnet-metawehrlites, metawebsterites and crustal two-pyroxene granulites. Most metaproxenites and granulites represent the Ti-Al-bearing augite suite and their bulk geochemistry indicates transitional olivine basalt magmatic affinities. Metasomatised, hydrous lithologies are only rarely present. Co-existing pyroxenes in the xenoliths provide re-equilibration temperature estimates from 860–1075°C (for the whole suite) and temperature-pressure estimates for the garnet metawebsterites from 1055–1070°C and 1.2–1.4 GPa. This gives a Miocene mantle geotherm gradient at least 80–130°C higher than the Southeast Australian (SEA) western Victorian geotherms. However, considerations of Moho depths from new seismic surveys below Table Cape (~32 km) suggest that the indicated geotherm is more strongly perturbed in its lower levels than at the mantle-crust transition. This localised perturbation is attributed to adjacent magma chamber activity in the mantle (Boat Harbour basalts) just prior to Table Cape vent activity. Tasmanian Miocene geotherms (Table Cape, Bow Hill) achieve relatively high gradients and reinforce suggestions of local variation in East Australian geothermal gradients. They illustrate the potential complexities in comparing xenolith-derived geotherms from different areas in general, both from thermometer/barometer selection and from associated magmatic heat inputs.

Key Words: Table Cape volcano, Cenozoic, xenoliths, mineralogy, mantle, crust, thermobarometry, geotherms, magma chamber.

INTRODUCTION

Table Cape is a prominent bluff of coarse-grained alkali dolerite (although previously mapped as crinanite, the preferred specific nomenclature is now teschenite; see Le Maitre 1989). It forms a massive core of a substantially eroded volcanic centre (fig. 1). The teschenite fronts on to Bass Strait in cliffs up to 179 m a.s.l. (Wynyard 1:25 000 Sheet; 55GCQ930655, AGC66, Everard 2003). This slightly differentiated body only becomes more distinctly alkaline in its chemistry in late-stage pegmatoidal veins (schlieren) and it incorporates ultramafic xenoliths and xenocrysts of mantle origin, particularly in its lower levels (Gee 1971, Piestrzeniewicz 1972, Everard 2001). Lavas from the Table Cape centre form a remnant apron, in parts deeply weathered, that was dated at 13.3 ± 0.2 Ma at Tollymore Road quarry (Sutherland & Wellman 1986). The lavas overlie a range of older lithologies (fig. 1) and extend west of Table Cape to Jacobs Boat Harbour as a mappable unit of normal magnetic polarity (Sutherland *et al.* 2004, fig. 12).

Vent pyroclastic deposits up to 5 m thick outcrop on the south shore of Table Cape (55GCQ930648, AGD66). Here they underlie a north-dipping base to the teschenite and contain a range of mantle and lower crustal fragments within a basaltic and comminuted country rock matrix (Sutherland *et al.* 1989, Everard 2001, Sutherland *et al.* 2004, figs 8d-f, 10f). Rare fragments of Lower Miocene marine sediments in the pyroclastic beds indicate an age younger than the marine beds. The pyroclastics include irregular sill-like bodies of chilled tachylytic basalt that form a more resistant protection to the included fragments. The deeper origin xenoliths are important in reconstructing the underlying Tasmanian lithosphere and its geothermal state at

the time of volcanism, as they not only include garnet-two pyroxene assemblages suitable for geothermobarometry but also diverse mantle peridotites and pyroxenites (Sutherland *et al.* 1994, 2004, Beyer 1998, Beyer *et al.* 2001). The Table Cape suite is one of only two known garnetiferous mantle xenolith sites in Tasmania, the other being at Bow Hill, west of Oatlands in the central midlands (fig. 1).

This study presents a more complete account of the Table Cape vent xenolith mineralogy and uses new thermobarometric methods to discuss their context within recent studies on the Southeast Australian lithosphere in the Cenozoic and its geothermal relationships (Graham *et al.* 2003, Sutherland *et al.* 2004, Roach 2004). The paper integrates the Table Cape vent xenolith data with Boat Harbour (Sutherland *et al.* 1996) and provides a revised lithospheric framework for the Wynyard-Boat Harbour region. The proposed lithology is then correlated with new seismic data on crust-mantle structures under northern Tasmania (Rawlinson *et al.* 2001, 2004).

SAMPLES AND ANALYTICAL METHODS

The Table Cape vent rocks contain numerous xenolithic and xenocrystic fragments. The fragments of mantle-lower crust origin include common spinel-metaperidotites up to 10 cm, sparse garnet-metaproxenites and granulites up to 8 cm, rarer pyroxenites up to 6 cm and sporadic pyroxene and spinel megacrysts up to 3 cm across, as summarised in Everard (2001).

The samples for this mineralogical study were collected by breaking up the dense glassy basalt, which provided more protected, fresher samples than those found in weathered

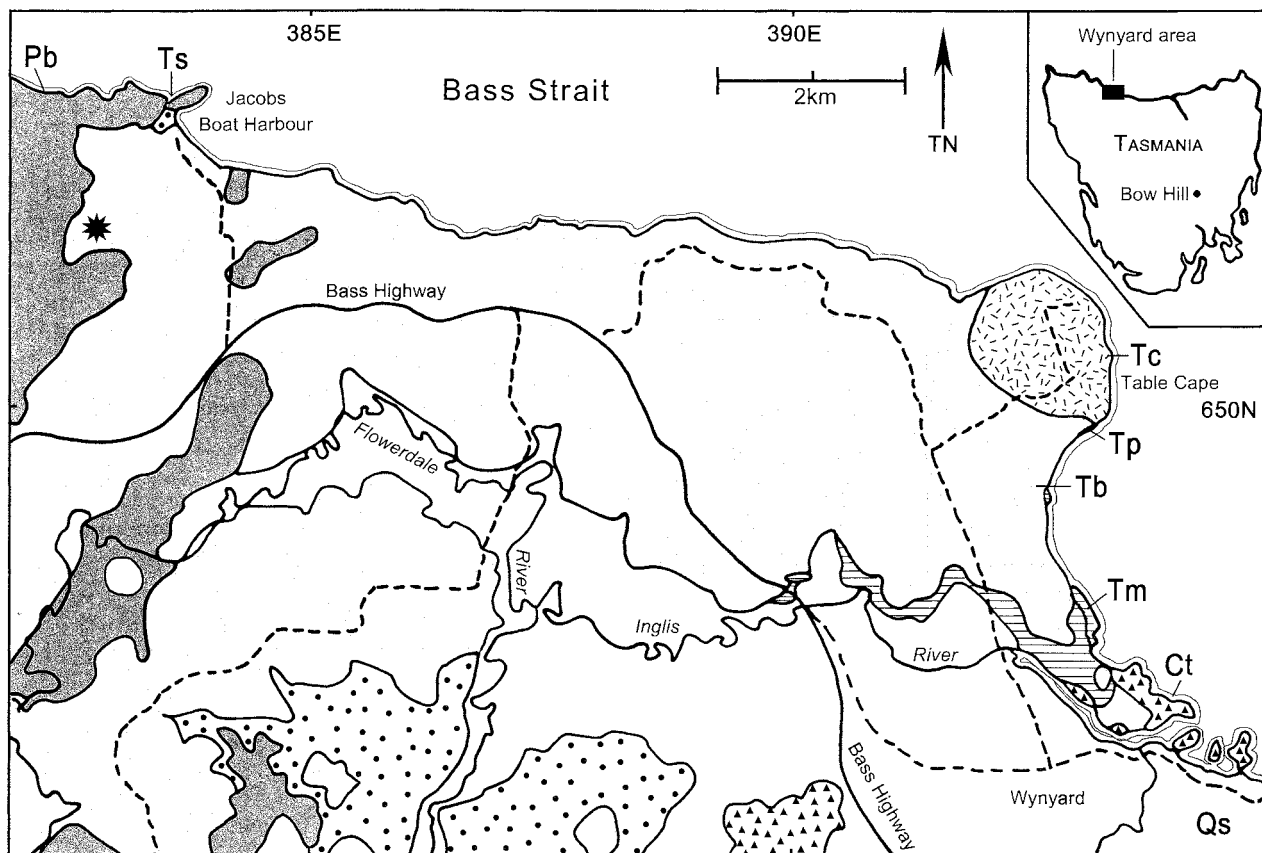


FIG. 1 — Geological map of the Table Cape–Jacobs Boat Harbour area, west of Wynyard, with Table Cape intrusive core (Tc), Table Cape vent pyroclastics (Tp), Table Cape basalt flows (Tb), underlying Miocene marine beds (Tm) and Carboniferous glacial beds (Ct). Basalt (grey areas) include flows from the Cassidy's Creek centre (star) in the western part and overlie Tertiary sands and gravels (Ts) and Precambrian basement beds (Pb). The drainage includes the Flowerdale River (west) which joins the Inglis River with the outlet near Wynyard Creek (east) at Wynyard. The blank land areas include Quaternary alluvium and coastal sands (Qs). Geology after Gee (1971), Sutherland et al. (1996) and Everard (2003).

pyroclastic matrix. Because of the incomplete recovery of xenoliths by this means, a detailed census of xenolith types was not attempted. A range of inclusions was selected for cutting and polishing into discs ready for petrographic microscope examination and analysis by electron microprobe (EMP). The study material is held in the collections of the Australian Museum. The host tachylyte was analysed from hand-picked fragments to avoid included material as much as possible, before crushing for analysis for major, minor and trace elements using conventional X-ray fluorescence (XRF) techniques (table 1).

The examination of polished samples by a Leica DMLP polarising microscope with DC 300 digital camera attachment identified a number of xenocryst and xenolith assemblages for detailed analysis and photography (pl. 1). These included:

- (a) Pyroxene megacrysts (table 2); clinopyroxene, orthopyroxene.
- (b) Metaperidotites (table 3); spinel-metalherzolite, spinel-metaharzburgite.
- (c) Metapyroxenites (table 4); metawebsterite, mica-metawebsterite.
- (d) Garnet-metapyroxenites (table 5); garnet (\pm spinel)-metawebsterite.
- (e) Garnet-metaclinopyroxenites (table 6); garnet-spinel-metawehrlite, garnet-metaclinopyroxenite.

- (f) Two-pyroxene granulites (table 7); spinel-granulite, amphibole-mica-granulite.

The assemblages also exhibit secondary breakdown mineral products related to decompression and melt fluid interactions during magmatic transport. These include clinopyroxene, orthopyroxene, spinel and amphibole (table 8) among other phases.

The mineral analyses (tables 2–8) were largely carried out using an ETEC Autoprobe, fitted with three wave-length spectrometers and a LINK EDS system in the Electron Microscope Section, University of Sydney (L. R. Raynor and F. L. Sutherland, analysts). However, some analyses were made on an automated ETEC electron microprobe at Macquarie University, Sydney, for comparative purposes; (B. J. Barron, analyst). Analyses were averaged from multiple spot analyses (up to 10 per grain) for several grains of each mineral phase. The analyses were standardised against natural mineral standards and used 15 kV acceleration voltage, a sample current of 20 nA and Bence-Albee corrections. Precisions were better than $\pm 1\%$ for element contents above 10 wt% as oxides, $\pm 5\%$ at 1–10 wt% oxides and $\pm 10\%$ at below 1 wt% oxides. Agreement between analyses for the two instruments was usually within $\pm 2\%$ for oxides over 5 wt%, within $\pm 8\%$ for oxides between 1–4 wt% and within $\pm 30\%$ for oxides below 1 wt%.

TABLE 1
Chemical analysis of tachylytic host to Table Cape xenolith suite

Major elements (wt%)		Trace elements (ppm)		CIPW norm (anhydrous)	
SiO ₂	42.46	Ni	278	Or	13.61
TiO ₂	2.32	Cr	360	Ab	3.00
Al ₂ O ₃	12.39	V	135	An	4.41
Fe ₂ O ₃	2.15	Ba	181	Ne	22.11
FeO	10.74	Sr	1295	Di	25.31
MnO	0.20	Rb	14	Ol	20.49
MgO	10.13	Y	24	Mt	3.18
CaO	8.92	Ga	20	Il	4.49
Na ₂ O	5.08	Zr	318	Ap	3.19
K ₂ O	2.26	Nb	81	Cc	0.28
P ₂ O ₅	1.32	Pb	23	An mol %	58.1
CO ₂	0.12	Zn	156	= 100 An/ (An + 1.061 Ab).	
H ₂ O+	1.59	Cu	61	Differentiation Index (Or+Ab+Ne)	
Total	99.68	Th	7		
Mg/		U	4		38.7
(Mg+Fe ²⁺)%	62.70				

Analyst E.L. Sutherland, Macquarie University, Laboratories, Sydney.

(Major elements from X-ray fluorescence spectrometry, using fused borate buttons, with Na₂O determined by flame photometry, FeO by dissolution with HF and ammonium metavanadate and titration against ceric sulphate, with CO₂ and H₂O determined separately: Trace elements from X-ray fluorescence using pressed powder pellets using boric acid bases).

CIPW norm calculated at a ratio of Fe₂O₃/(FeO + Fe₂O₃) = 0.2

RESULTS

Host Petrology

The host tachylyte consists of a dark, glassy matrix containing small euhedral, partly corroded olivine (ol) crystals (~7%) up to 0.1 mm across (pl. 1A). These phenocrysts lack the anhedral shapes, strain features and reaction margins observed in the incorporated xenocrysts and xenoliths. Analysis of the tachylyte (table 1) corresponds to a mildly fractionated olivine nephelinite (Mg value <63; Ni 278, Cr 360 ppm; Ab < 3%, An mol% 58, D.I. 39) compared to more primitive Tasmanian nephelinites (Sutherland *et al.* 2004). The presence of mantle inclusions, however, indicates a mantle-fractionated melt. Normative ol for the Table Cape nephelinite suggests a fractionated composition for the microphenocrystic olivine, which co-crystallised with minute octahedral inclusions of the ulvospinel-magnetite series.

Megacryst Suite

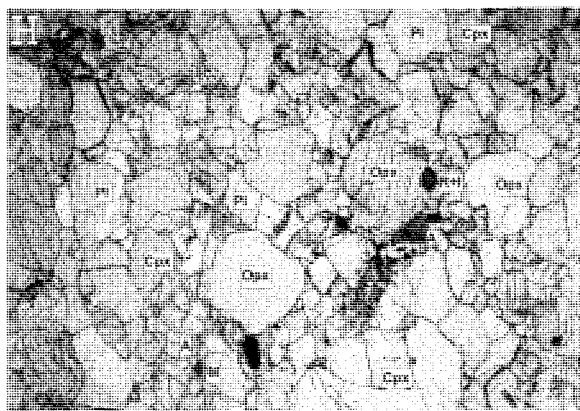
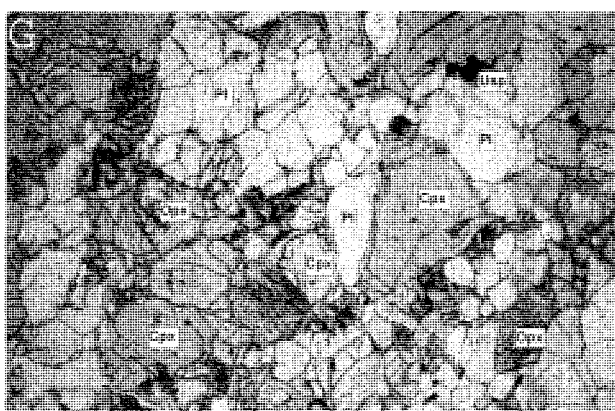
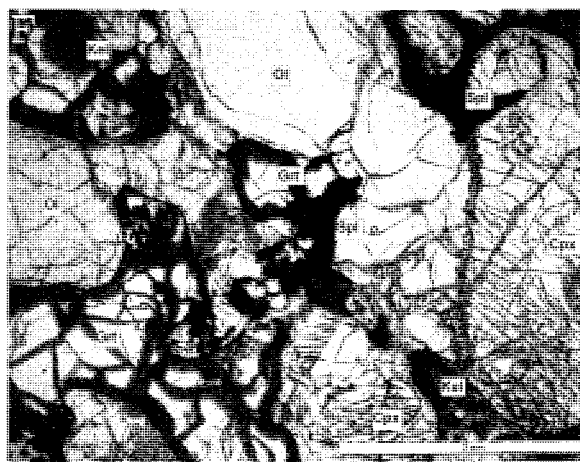
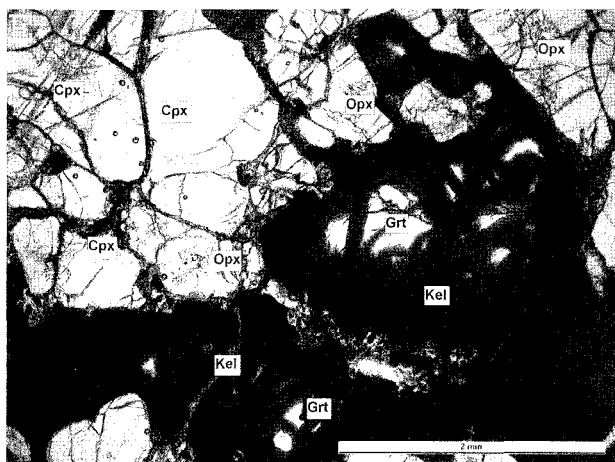
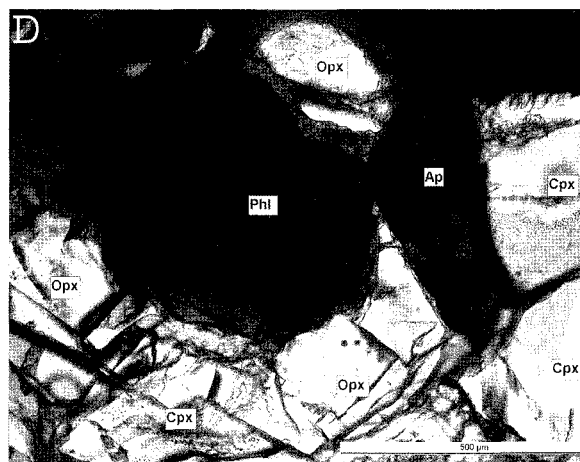
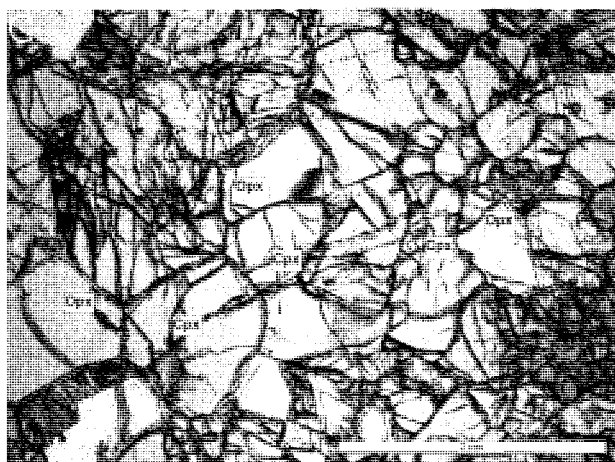
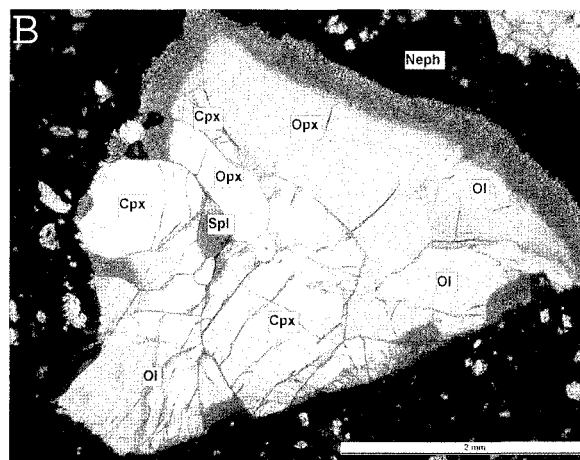
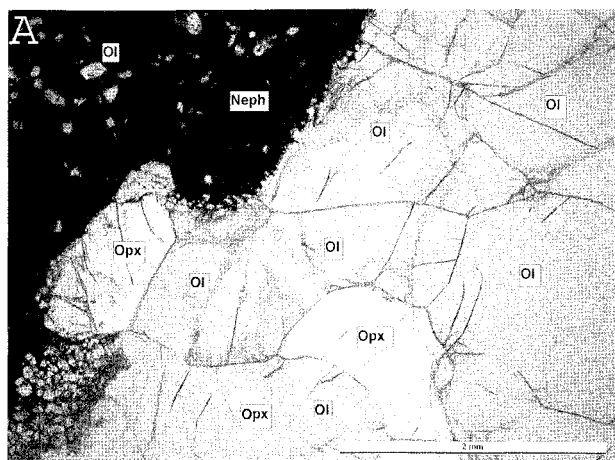
Clinopyroxene (cpx) is the main megacryst within the Table Cape vent (table 2). Unzoned cores are Al-rich augite (En₅₂₋₅₃ Wo₃₅₋₃₆ Fs₁₁₋₁₂). They show sieved, resorbed reaction rims composed of Ti and Na-enriched augite (En₄₄₋₄₅ Wo₄₀ Fs₁₅₋₁₆), which gave low total oxides in EMP analyses due to the inhomogeneous, fluid-bearing melt reactions. A full end-member calculation (Cawthorn & Collerson 1974)

shows the cores are notably more Mg-rich (~En₄₅ Wo₂₄₋₄₃ Fs₉₋₁₀ Tsc₁₂₋₁₃ Jad₈₋₉, Cta₁₋₂ Cfs₁) than in most Tasmanian cpx megacrysts (Sutherland *et al.* 2004). The distinctive megacryst chemistry compared to that of clinopyroxenes in the accompanying xenoliths suggests the former represent a high-pressure liquidus phase crystallised from an Mg-rich parental melt. Experimental work on highly undersaturated nephelinitic or basaltic melts suggests such clinopyroxenes can crystallise as liquidus phases from volatile-bearing melts between 1250–1300°C and 2.4–2.6 GPa (Adam 1990).

Orthopyroxene (opx) megacrysts with anhedral, embayed rims and high Mg (~En₈₈ Fs₁₁ Wo₁) also occur at Table Cape (table 2). Their compositions, however, closely match those of enstatites in the spinel-metaperidotite xenoliths. Thus, they may derive from these sources rather than from liquidus crystallisation, particularly as opx is commonly reacted out among high pressure liquidus products (Adam 1990). Similarly, the spinel (spl) megacrysts are close to spinel compositions within the xenoliths.

Metaperidotites

Two distinct types were identified. The more common (pl. 1B) is a coarse polygonal-textured assemblage (~75% ol, 15% opx, 5% cpx and 5% spl from visual grid estimates) with strain/extinction in the silicates. The ol (Fo₈₉) and opx (En₈₈) are Mg-rich, the cpx is Al-Na-rich diopside (Wo₄₈ En₄₆ Fs₆) and the spl is Cr-bearing pleonaste (e.g., Sp₇₈ Hc₁₂ Cm₈ Mt₁,



Usp₁) (table 3). This type dominates mantle xenoliths in the Table Cape region and elsewhere in Tasmania (Piesterzeniewicz 1972, Varne 1977, Sutherland *et al.* 1996).

Rare spinel-metaharzburgite (pl. 1A) exhibits a medium to coarse granular polygonal texture (grains to 5 mm across). It is richer in ol (78.5%) and opx (20%) and impoverished in spl (1%) and cpx (0.5%) compared to the spinel-metaherzolitites. The ol (Fo₉₁), opx (En₈₉), cpx (En₅₂ Wo₄₃ Fs₅) and spl are all Mg-rich phases, with the spl being a Cr-rich magnesiochromite (Mc₄₇ Sp₂₅ Hc₂₂ Mf₅ Usp₁) (table 3). Such highly depleted metaperidotites are rarely described from Tasmanian mantle xenolith suites.

Metawebsterites

Two types of metapyroxenites were found. One lacked obvious accessory phases and consisted of subequal polygonal grains of cpx and opx up to 4 mm across (pl. 1C). Pleochroic cpx (α pale green, β pale brown, γ pale pink) is Al-rich, Na-Ti-bearing augite (En₄₈ Wo₄₀₋₄₁ Fs₁₁₋₁₂) and pleochroic opx (α pale grey, β pale yellow, γ pale green) is moderately Al- and Fe-enriched (En₇₉₋₈₁ Fs₁₇₋₁₉ Wo₂) (table 4). Exsolution lamellae and blebs of one pyroxene within the other occur in some grains.

The other type is cpx-rich, with large pleochroic cpx grains up to 15 mm across. These enclose and intergrow with opx, mica and cloudy apatite grains up to 1 mm across (pl. 1D). The cpx is Al-Mg-rich, Cr-bearing diopside (Wo₄₇ En₄₇ Fs₆), the opx is Al-bearing enstatite (En₈₈), and the mica is Ti-rich magnesian phlogopite (table 4). Such pyroxenites with Cr-diopside suite characteristics (Wilshire & Shervais 1975) are rare in Tasmania (Everard 2001) and also within eastern Australia (O'Reilly & Griffin 2000).

Garnet-metawebsterites

These consist of polygonal, amoeboid-like and blebby intergrowths of cpx, opx, garnet (grt) and in some cases spl, in grains up to 6 mm across (pl. 1E). Smaller areas of finer grained recrystallised mosaics can occur within coarse-grained matrix. Assemblages carry up to 25% grt, commonly in relicts within reaction rims and patches but rarely in completely fresh grains. Fresh cores are unzoned pyrope of restricted end-member range (Py₆₃₋₆₄ Am₂₂₋₂₃ Gr₉₋₁₀ An₃₋₄ Uvo₀₋₁ Sp₀₋₁; after Knowles 1987); the pleochroic cpx and opx range from sub-equal to cpx-dominant amounts with only 5% opx and include Al-Ti-Na-rich augite (En₄₅₋₄₇ Wo₄₀₋₄₁ Fs₁₃₋₁₄) and Al-rich opx (En₇₇₋₇₈) (table 5).

Two of the five studied garnet-metawebsterites contain up to 5% spinel as blocky interstitial grains or as cores up to 1.5 mm across within reacted garnet enclosures. The spl is Cr-bearing pleonaste (Sp₆₅₋₇₂ Hc₂₄₋₂₅ Mt₂₋₆ Cm₀₋₃ Usp₀₋₂) (table 5). Some grains resemble Fe-hydroxide altered olivine (5%) in one assemblage, which has pervasive fluid inclusion trails and alterations along grain boundaries. These garnet (\pm spinel)-two pyroxene assemblages allow TP estimates for the Table Cape vent suite, but are notably more Fe-enriched cpx (FeO total >5wt%) than in most garnet-websterite suites (Sutherland *et al.* 1994).

Garnet-metaclinopyroxenite and Garnet-metawehrlite

A rare garnet-metaclinopyroxenite contains abundant reacted garnet (35%). This includes and is intergrown with rounded, embayed and polygonal cpx (55%) grains up to 4.5 mm across. The reacted areas enclose many secondary spl grains, rimmed by pod-like growths of secondary opx, while alteration prevails along cpx grain boundaries. The grt (Py₆₂ An₂₃ Gr₁₁ An₃ Sp₁) and Al-Na-Ti-rich augite (En₄₅ Wo₄₁ Fs₁₄) compositions are similar to those in the garnet-metawebsterites and one cpx is mantled by a gedritic amphibole (table 6).

A garnet-metawehrlite contains grt (42.5%), cpx (37.5%), spl (10%), and ol (10%). Poikilitic areas of garnet (to over 12 mm across) enclose grains and polygonal clusters of the other minerals (pl. 1F). The garnet is pyrope (Py₆₃₋₆₄ Am₂₂₋₂₃ Gr₁₀₋₁₁ An₃ Sp₁) in composition. The cpx (up to 4.5 mm across) is Ti-Na-rich augite (En₄₆ Wo₄₂ Fs₁₂), while blocky to rounded spl grains (up to 1 mm across) with narrow secondary opx fringes are pleonaste (Sp₆₉₋₇₀ Hc₂₂₋₂₃ Mt₅₋₆ Usp₂ Cm₀₋₁) and ol (Fo₈₁) forms subhedral grains up to 2.5 mm across. Other than olivine, the mineralogy (table 6) closely compares with phases in the garnet (spinel)-metapyroxenites.

Two-pyroxene Granulites

These feldspathic rocks form two types. One is a medium-grained, banded assemblage with conspicuous zones of polygonal plagioclase (55%) interspersed with subequal mosaics of pleochroic opx and cpx, mostly 0.5–2.5 mm across, and sparse interstitial irregular spl grains up to 0.5 mm across (pl. 1G). The zoned plagioclase shows lamellar and pericline twinning and is calcic (An₅₈₋₆₀); the cpx is Al-bearing diopside (Wo₄₆₋₄₇ En₃₆ Fs₁₇₋₁₈), the opx is Fe-enriched

PLATE 1

Photomicrographs of main Table Cape vent xenoliths. (A) Metaharzburgite, in host olivine nephelinite (top left). (B) Spinel-metaherzolitite, in host olivine nephelinite. Note reaction rim on xenolith. (C) Metawebsterite, with mosaic-textured pyroxenes. (D) Mica-apatite-bearing metawebsterite. Note basal phlogopite plate (left), cloudy apatite crystal (right) and fluid inclusion trails in the pyroxenes. (E) Garnet-metawebsterite. Note 'kelyphite' alteration rims around inner cores of garnet. (F) Garnet-spinel-metawehrlite. Note large fresh olivine grains, 'kelyphite' rims and veins in garnet grains and interstitial spinel grains. (G) Two-pyroxene granulite. Note polygonal texture in plagioclase and pyroxenes and interstitial accessory ulvospinel. (H) Metasomatised two-pyroxene granulite. Note accessory mica, amphibole and rutile grains mantled with ilmenite. All photomicrographs have equal magnification, except for (D) which is enlarged 4X. Symbol for host nephelinite is Neph and for minerals are Ol (olivine), Opx (orthopyroxene), Cpx (clinopyroxene), Spl (spinel), Usp (ulvospinel), Grt (garnet), Kel ('kelyphite'), Pl (plagioclase), Phl (phlogopite), Ap (apatite), A (amphibole), M (mica), R (rutile), I (ilmenite). Microphotographs were taken in transmitted light, I. T. Graham photographer.

TABLE 2
Representative analyses of pyroxene megacrysts,
Table Cape pyroclastics

	Clinopyroxene		Orthopyroxene
	core	resorbed rim	
SiO ₂ wt%	50.70	48.76	54.54
TiO ₂	0.61	1.36	0.12
Al ₂ O ₃	8.03	8.49	4.70
Cr ₂ O ₃	0.25	-	0.27
FeO total	6.52	8.26	6.87
MgO	16.61	13.25	32.00
CaO	15.54	16.46	0.64
Na ₂ O	1.18	1.76	-
Total	99.44	98.34	99.14
Cation	Mg _{52.8}	Mg _{44.7}	Mg _{88.1}
ratio%	Ca _{35.5}	Ca _{39.9}	Fe _{10.6}
	Fe _{11.6}	Fe _{15.5}	Ca _{1.3}

Analyst F. L. Sutherland

TABLE 3
Comparative representative analyses, peridotitic xenoliths (DR 10491)

Mineral/ Rock Type	Olivine		Orthopyroxene		Clinopyroxene		Spinel	
	SML	SMH	SML	SMH	SML	SMH	SML	SMH
SiO ₂ wt%	40.38	41.05	53.90	55.46	50.99	52.71	0.09	0.09
TiO ₂			0.15	0.08	0.74	0.11	0.18	0.12
Al ₂ O ₃		0.09	4.62	2.65	6.94	3.04	60.33	26.80
Cr ₂ O ₃		0.03	0.29	0.77	0.67	1.16	7.69	40.25
FeO total	11.09	8.95	6.89	5.70	3.01	2.89	11.14	15.44
MnO	0.14		0.13	0.14	0.08	0.09	0.15	0.37
MgO	49.29	49.31	32.28	33.36	14.40	18.12	20.61	16.29
CaO	0.06	0.09	0.68	1.11	20.78	21.01		
Na ₂ O			0.11	0.08	1.95	0.37		
Total	100.96	99.52	99.05	99.35	99.56	99.50	100.19	99.36
Cation	Mg ₈₉	Mg ₉₁	Mg ₈₈	Mg ₈₉	Mg ₄₆	Mg ₅₂	Mg ₆₇	Cr ₄₆
ratio %	Fe ₁₁	Fe ₉	Fe ₁₁	Fe ₉	Ca ₄₈	Ca ₄₃	Fe ₂₀	Mg ₃₅
			Ca ₁	Ca ₂	Fe ₆	Fe ₅	Cr ₁₃	Fe ₁₉

SML Spinel-metalherzolite SMH Spinel-metaharzburgite. Analyst L. R. Raynor.

Calculated bulk rock compositions, based on modal proportions of mineral phases

Rock	SML	SMH	SML (ol 75%, opx 15%, cpx 5%, spl 5%)	SMH (ol 78.5%, opx 20%, spl 1%, cpx 0.5%)
SiO ₂	40.93	43.57		
TiO ₂	0.16	0.02		
Al ₂ O ₃	4.06	0.89		
Cr ₂ O ₃	0.41	0.58		
FeO total	10.07	8.46		
MnO	0.14	0.03		
MgO	43.56	45.65		
CaO	1.14	0.40		
Na ₂ O	0.12	0.02		
Total	100.59	99.62		

TABLE 4
Comparative representative analyses, metabasites and mica-metabasite

Mineral/ rock type	Clinopyroxene			Orthopyroxene			Mica
	MW(WW14)	MW(WW16)	MW(WW6)	MW(WW14)	MW(WW16)	MW(WW6)	MW(WW6)
SiO ₂ wt%	49.85	49.87	52.08	52.71	52.83	55.01	36.69
TiO ₂	1.17	1.08	0.23	0.35	0.34		4.38
Al ₂ O ₃	7.86	7.50	7.12	5.40	5.53	4.90	16.85
Cr ₂ O ₃	0.29	0.40	0.75	0.17	0.23	0.24	0.49
FeO total	7.27	6.22	3.10	12.52	10.82	6.90	4.77
MnO	0.16	0.15		0.19	0.18		
MgO	14.21	14.41	14.28	27.70	28.55	32.11	19.87
CaO	16.94	17.36	19.60	1.11	1.21	0.71	0.15
Na ₂ O	2.27	2.03	1.87	0.25	0.16		0.35
K ₂ O							8.50
Total	100.02	99.02	99.03	100.40	99.85	99.87	92.05
Cation	Mg ₄₈	Mg ₄₈	Mg ₄₇	Mg ₇₉	Mg ₈₁	Mg ₈₈	Mg ₈₀
ratio %	Ca ₄₀	Ca ₄₁	Ca ₄₇	Fe ₁₉	Fe ₁₇	Fe ₁₁	Fe ₁₁
	Fe ₁₂	Fe ₁₁	Fe ₆	Ca ₂	Ca ₂	Ca ₁	Ti ₉

MW Metabasite (WW14, WW16). WW14 cpx av. 4 analyses, opx av. 4 analyses; WW16 cpx av. 8 analyses, opx av. 5 analyses. Analyst L. R. Raynor.

MW Mica + apatite-bearing metabasite (WW6). Analyst F. L. Sutherland.

Calculated bulk rock compositions, based on modal proportions of mineral phases

Rock	WW14	WW16	WW6	WW14 (cpx 50%, opx 50%) WW16 (cpx 50%, opx 50%) WW6 (cpx 95%, opx 5%, phl < 0.5%, ap < 0.5%)
SiO ₂ wt%	51.28	51.36	52.33	
TiO ₂	0.76	0.71	0.22	
Al ₂ O ₃	6.63	6.52	7.01	
Cr ₂ O ₃	0.23	0.31	0.72	
FeO total	9.90	8.52	3.30	
MnO	0.18	0.17		
MgO	20.96	21.49	15.18	
CaO	9.03	9.29	18.66	
Na ₂ O	1.26	1.10	1.78	
K ₂ O			<0.05	
P ₂ O ₅			<0.05	
Total	100.23	99.47	99.20	

TABLE 5
Comparative representative analyses, garnet-two pyroxene assemblages

Mineral/rock type	Clinopyroxene	Orthopyroxene	Garnet	Spinel
	GW(WW8-WW13)	GW(WW8-WW13)	GW(WW8-WW13)	GW(WW12-13)
SiO ₂	49.47-50.24	51.53-52.41	40.49-41.28	
TiO ₂	1.28-1.40	0.40-0.50	0.45-0.59	0.76-0.80
Al ₂ O ₃	8.05-8.36	5.58-5.98	22.54-22.99	58.26-59.31
Cr ₂ O ₃	0.00-0.25	0.03-0.15	0.06-0.40	0.41-2.36
FeO total	7.21-7.71	12.46-12.95	12.35-13.66	21.47-22.76
MnO	0.00-0.16	0.15-0.19	0.33-0.41	0.08-0.13
MgO	13.77-14.10	27.26-27.94	16.98-17.45	16.68-16.81
CaO	16.62-16.93	1.12-1.28	5.09-5.26	0.00-0.02
Na ₂ O	2.05-2.28	0.16-0.24		
K ₂ O				
Total	99.48-100.34	99.68-100.88	99.82-100.53	99.58-100.17
Cation	Mg ₄₅₋₄₇	Mg ₇₇₋₇₈	Mg ₆₀₋₆₁	Mg ₅₆₋₆₀
ratio %	Ca ₄₀₋₄₁	Fe ₁₉₋₂₀	Fe ₂₆₋₂₇	Fe ₄₀₋₄₃
	Fe ₁₃₋₁₄	Ca ₂₋₃	Ca ₁₃₋₁₄	Cr ₁₋₄

GW Garnet-websterite (\pm spinel). WW8-WW10; cpx av. 5-13 analyses, opx av. 4-8 analyses, grt av. 5-6 analyses. GW (WW12-13) cpx av. 8 analyses, opx av. 4 analyses, grt av. 8 analyses, spinel av. 2-3 analyses. Analyst L. R. Raynor.

(En₅₇₋₅₈) and the spinel is ulvospinel (Usp₉₂₋₉₃ Mf₅₋₆ Sp₁₋₂ Mc₀₋₁) (table 7).

The other granulite is a fine-grained, slightly foliated assemblage composed of plagioclase (35%), cpx (30%) and opx (25%) forming strongly polygonal recrystallisations in grains mostly 0.5-1 mm across (pl. 1H). The large pyroxenes show strain lamellae and undulose extinction and the cpx is pleochroic. The plagioclase is distinctly calcic (An₇₃₋₇₅); the cpx is Al-rich diopside (Wo₄₈₋₄₉ En₄₀₋₄₁ Fs₁₁₋₁₂), the opx is moderately Fe-enriched (En₆₈₋₇₀) and small interstitial accessory minerals (10%) include orange-brown rutile (5%) altered on rims and cleavages to opaque Mg-rich ilmenite, pargasitic amphibole, Ti-rich phlogopite and rare corundum (table 7).

Secondary Minerals

The garnet-bearing assemblages show varied degrees of reaction of the garnet, which alters to cloudy, fine-grained admixtures of spinel and feldspar containing secondary spl, opx and rarely cpx grains. The spl is pleonaste (Sp₆₇₋₇₄ Hc₂₃₋₃₀ Mt₂₋₃ Usp₀₋₁ Cm₀₋₁), the opx is moderately Fe-enriched (En₇₅₋₇₉) and cpx is Al-rich augite (En₄₉₋₅₀ Wo₃₇₋₃₈ Fs₁₃) (table 8). The Ca-contents in the opx (Wo₃₋₄) suggest relatively high temperature reactions were involved in their formation (Brey & Köhler 1990) while up to 0.7% Na₂O in the cpx suggests a metasomatic component helped in the garnet breakdown. The steps involved in such breakdown reactions were detailed in Griffin *et al.* (1984).

Secondary amphibole forms brown infillings between spl and cpx grains in a garnet-spinel-metacclinopyroxenite. It is kaersutite in composition, distinctly more Ti-rich than the gedritic amphibole that mantles cpx in the same assemblage, and may derive from fluid infiltrations from the host melt.

Thermobarometry

Several thermometers and barometers were used to investigate temperature (T) and pressure (P) conditions for metamorphic re-equilibration in the xenolith assemblages. Previously reported couplings (Ellis & Green 1974 T and Wood 1974 P; Brey & Köhler 1990 TP) for Table Cape (Sutherland *et al.* 1994) are not further considered because of apparent compositional and experimental incompatibilities, discussed later. The Wells (1977) two-pyroxene thermometry, with appropriate allocations of Fe²⁺ and Fe³⁺, provides good results at T < 1300°C, with uncertainties in the precision within \pm 50°C (Tribaudino & Bruno 1993, Taylor 1998). This gives a thermal ordering of re-equilibration temperatures (quoted to the nearest 5°C) for the Table Cape vent meta-assemblages. The method gives a range of 1055-1070°C for garnet (\pm spinel)-metawebsterites and metawebsterites, 810-1080°C for spinel-metaperidotites, 950-960°C for mica (\pm apatite)-metawebsterites and 860-870°C for granulites. The relatively wide spinel-metaperidotite temperatures overlap those for the granulites and some metawebsterites and can be refined further using the Sachtleben & Seck (1981) and Witt-Eickschen & Seck (1991) thermometric methods. This gave likely T ranges between 915-1010°C for the spinel-lherzolite and 965-1075°C for the spinel-bearing harzburgite, which preserves the relative ordering for these assemblages established with the Wells thermometer. The modified metaperidotite T results also delineate the crustal granulite range (< 900°C) from the mantle T range (> 900°C) and suggest the crust-mantle boundary (MOHO) intersected an ambient geotherm at around 900°C (fig. 2).

The garnet-metawebsterites permit three-phase thermobarometry. This study used the Wells (Fe³⁺) two-pyroxene thermometric results with the Wood (1974) garnet-orthopyroxene barometer. This coupling provides a narrow TP range for these assemblages (1050-1070°C; 1.22-1.36 GPa), compatible with the tight compositional range for the assemblage analyses and the experimental spinel-pyroxenite/

TABLE 6
Comparative representative analyses, garnet-metawehrlite and metaclinopyroxenite

Mineral/ Rock type	Clinopyroxene		Garnet		Olivine	Spinel		Amphibole
	GW	GC	GW	GC	GW	GW	GC	GC
SiO ₂	49.41	49.66	41.52	41.00	39.18	0.29	0.29	43.89
TiO ₂	1.47	1.43	0.43	0.49		0.62	0.24	0.74
Al ₂ O ₃	9.14	8.35	23.41	23.11		61.01	65.17	19.01
Cr ₂ O ₃	0.02		0.03		0.03	0.47	0.18	
FeO total	6.65	7.68	12.08	13.61	17.49	19.62	15.66	12.67
MnO	0.07		0.30	0.32	0.16	0.13	0.15	0.26
MgO	13.66	13.84	17.70	17.08	42.96	18.20	19.02	16.63
CaO	17.60	17.21	5.30	5.31	0.20			4.00
Na ₂ O	2.07	1.98						0.57
K ₂ O								0.58
Total	100.09	100.15	100.77	100.92	100.02	100.34	100.71	98.35
Cation	Mg ₄₆	Mg ₄₅	Mg ₆₃	Mg ₆₀	Mg ₈₁	Mg ₆₂	Mg ₆₈	Mg ₆₂
ratio %	Ca ₄₂	Ca ₄₁	Fe ₂₄	Fe ₂₇	Fe ₁₉	Fe ₃₇	Fe ₃₂	Fe ₂₇
	Fe ₁₂	Fe ₁₄	Ca ₁₃	Ca ₁₃		Ti ₁	Ca ₁₁	

GW Garnet-spinel metawehrlite (WW4). Analyst L. R. Raynor. GC Garnet-amphibole-metaclinopyroxenite (WW15). Analyst B. J. Barron.

Calculated bulk rock compositions, based on modal proportions of mineral phases

Rock	WW4	WW15	WW4 (grt 42.5%, cpx 37.5%, spl 10%, ol 10%)
SiO ₂ wt%	40.13	41.66	WW4 (cpx 55%, grt 35%, spl 10%)
TiO ₂	0.79	1.00	
Al ₂ O ₃	19.48	19.18	
Cr ₂ O ₃	0.07	0.01	
FeO total	11.33	10.58	
MnO	0.19	0.11	
MgO	18.76	15.59	
CaO	8.87	11.33	
Na ₂ O	0.78	1.09	
Total	100.40	100.55	

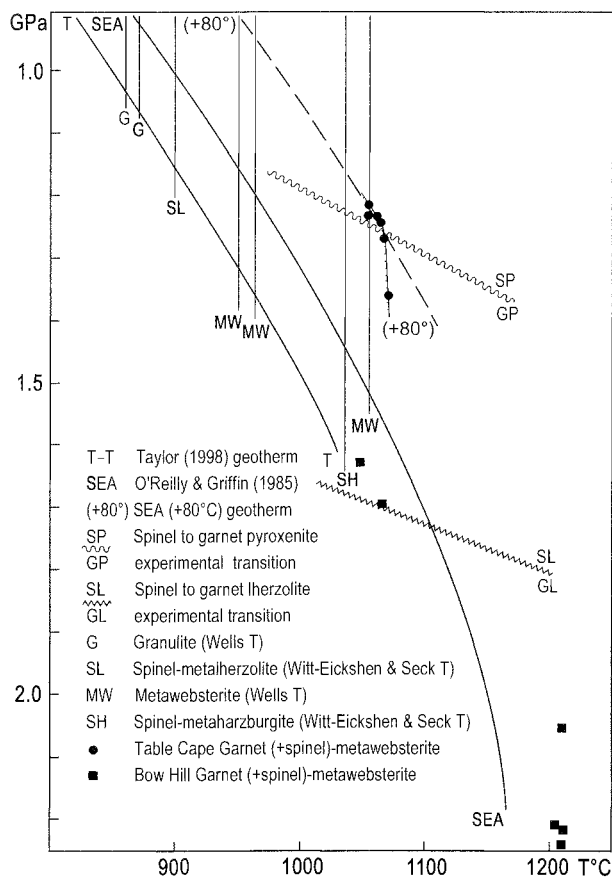


FIG. 2—Temperature (T °C)-Pressure (GPa) diagram, relating Table Cape (Wells T, Wood P) and Bow Hill (Taylor TP) plots to experimental spinel to garnet pyroxenite and spinel to garnet lherzolite phase transitions. Re-equilibration temperature lines for two-pyroxene granulites and metawebsterites (Wells T) and metaperidotites (Witt-Eickshen & Seck T) are indicated to show intersections on different south-eastern Australian geotherms. The TP points for Table Cape garnet (\pm spinel)-websterites are 1060°C, 1.23 GPa (WW8); 1065°C, 1.24 GPa (WW9); 1055°C, 1.23 GPa (WW10); 1070°C, 1.36 GPa (WW12), 1065°C, 1.27 GPa (WW13). The Wood barometry uses the Cr-Na-Al modification. The TP points for Bow Hill garnet (\pm spinel)-lherzolites and garnet-websterite are 1048°C, 1.63 GPa (DR12508), 1065°C, 1.69 GPa (DR 12501), 1205°C, 2.21 GPa (EBH-E), 1213°C, 2.22 GPa (EBH-D); 1210°C, 2.24 GPa (BHS); 1212°C, 2.05 GPa (QQ20a).

garnet-pyroxenite transition (fig. 2). The Table Cape TP results are 60–85°C higher at equivalent pressures than for the Southeast Australian (SEA) geotherm of O'Reilly & Griffin (1985), based on Ellis & Green (1979) T with Wood (1974) P, and 110–135°C higher than for the revised SEA geotherm, using Taylor (1998) TP (fig. 2). This reinforces conclusions from previous thermobarometry for the Table Cape vent suite that suggest a relatively hotter Cenozoic geotherm for the Tasmanian lithosphere (Sutherland *et al.* 1994), but the new thermobarometry provides a more controlled result.

DISCUSSION

The Table Cape vent xenoliths allow partial insights into the underlying lower crust-upper mantle lithology and its thermal state when sampled by a Mid-Miocene nephelinitic melt. The host nephelinite was fractionated prior to the eruptive sampling and this evolution took place at TP conditions exceeding those estimated for the deepest contained mantle xenoliths (i.e., > 1070°C and 1.4 GPa).

Host Evolution

The host nephelinite has generally low Zr/Nb (4.0), Rb/Sr (0.017), Ba/Nb (2.2) and Ba/Th (26) ratios, which link it to nephelinites of the low Rb and Ba group in Tasmania (e.g., West Scottsdale, Northeast Tasmania; see Sutherland *et al.* 2004). These particular nephelinites are attributed to low degrees of partial mantle melting in which some phlogopite is retained in the mantle source. In this respect, the Table Cape vent nephelinite differs from the nearby, but older, nephelinite at Boat Harbour, which relates to the high Ba and Rb group (Sutherland *et al.* 1996). Similar nephelinites to the Table Cape vent melt represent initial melts with liquidus temperatures around 1250–1300°C at 2.4–3.5 GPa based on experimental work on Tasmanian examples from Lughrata and Scottsdale (Tilley & Thompson 1972, Frey *et al.* 1978, Adam 1990). The experiments on the Scottsdale nephelinite suggest such melts with only slightly (1–2 wt%) higher SiO₂ can be generated from garnet lherzolite mantle that contains volatile (H₂O + CO₂) concentrations of 6.5 wt% at 3.4 GPa (Adam 1990).

The precise high pressure mineral phases that crystallised during evolution of the erupted Table Cape vent melt are difficult to completely designate. They probably include clinopyroxene similar to the Mg-rich augite megacrysts (see Adam 1990, fig. 7). Other potential phases such as olivine, spinel and garnet may be difficult to distinguish in composition from those released from disaggregated mantle xenoliths, although some cases may be distinguished texturally and based on inclusions.

Assemblage Affinities

The xenoliths include mantle metaperidotites and metapyroxenites and crustal granulites, but their chemical characterisations are handicapped by lack of sufficiently large and fresh samples for analysis within the altered pyroclastic deposits. Only garnet-metapyroxenite and granulites were found suitable for conventional analyses (Sutherland *et al.* 2004). Approximate bulk compositions, however, were calculated from small, fresh samples by using estimated proportions of mineral phases analysed by EMP, using a visual grid.

The metaperidotites show significant olivine alteration in larger xenoliths, but compositions were estimated from mineral analyses in fresh spinel-metaharzburgite and metaharzburgite (table 3). These suggest the metaharzburgite is enriched in Al, Fe, Ca and Na and depleted in Si relative to the harzburgite. 'Mantle' CIPW norms based on the analyses, after Wilkinson & Stolz (1997), gave reasonable matches with the modal mineralogy, showing higher Ca-tschermakite in the metaharzburgite (Cats 4.4 *cf* 1.5 wt%) and higher Hy in the metaharzburgite (22.8 *cf* 13.5 wt%). The higher

TABLE 7
Comparative representative analyses, two-pyroxene granulites

Mineral/ rock type	Clinopyroxene		Orthopyroxene		Plagioclase		Amphibole	Mica
	BG(WW1)	MG(WW3)	BG(WW1)	MG(WW3)	BG(WW1)	MG(WW3)	MG(WW3)	MG(WW3)
SiO ₂ %	50.99	50.71	50.80	52.67	53.58	49.69	42.34	37.52
TiO ₂	0.22	0.56	0.14				2.92	5.45
Al ₂ O ₃	3.70	4.94	1.79	2.49	29.78	32.84	14.27	15.82
Cr ₂ O ₃		0.23					0.27	
FeO total	10.53	6.57	26.25	19.49			10.32	10.52
MnO	0.15		0.51	0.23				
MgO	12.15	13.35	20.09	24.24			13.35	14.78
CaO	21.79	22.12	0.43	0.54	12.08	15.11	11.69	3.23
Na ₂ O	0.47	0.48			4.64	2.86	1.57	0.65
K ₂ O					0.11	0.21	2.11	6.00
Total	100.00	98.96	100.01	99.66	100.19	100.71	98.84	93.97
Cation	Ca ₄₆₋₄₇	Ca ₄₈₋₄₉	Mg ₅₇₋₅₈	Mg ₆₈₋₆₉	Ca ₅₈₋₅₉	Ca ₇₃₋₇₄	Mg ₄₈₋₄₉	Mg ₆₃
ratio%	Mg ₃₆	Mg ₄₀₋₄₁	Fe ₄₁₋₄₂	Fe ₃₀₋₃₁	Na ₄₀₋₄₁	Na ₂₅₋₂₆	Ca ₃₀₋₃₁	Fe ₂₅
	Fe ₁₇₋₁₈	Fe ₁₁₋₁₂	Ca ₁	Ca ₁	K ₀₋₁	K ₁₋₂	Fe ₂₁	Ti ₁₂

BG Banded granulite (WW1), includes ulvospinel (Fe₆₆Ti₂₉Mg₅). MG Micro-granulite (WW3), includes rutile with Mg-rich ilmenite rims (Ti₅₀Fe₂₉Mg₂₁). Analyst F. L. Sutherland.

TABLE 8
Comparative representative analyses, secondary phases, garnet-pyroxene and pyroxene assemblages

Mineral/ rock type	Cpx	Opx	Opx	Opx	Spl	Spl	Spl	Spl	Amp
	WW4	WW4	WW9	WW15	WW9	WW12	WW14	WW15	WW15
SiO ₂ wt%	48.36	52.17	48.46	50.06	0.25			0.34	36.12
TiO ₂	0.68	0.23	0.39	0.54	0.21	0.14	0.22	0.19	5.45
Al ₂ O ₃	11.00	8.17	11.00	10.81	63.98	65.08	64.53	65.00	6.84
Cr ₂ O ₃	0.04	0.04	0.22	0.02	1.18	0.39	0.35	0.13	0.02
FeO total	7.17	11.03	12.27	10.50	16.48	15.02	17.27	14.85	19.70
MnO	0.28	0.27	0.36	0.27	0.23	0.13	0.17	0.16	0.28
MgO	15.34	26.82	24.95	26.66	18.30	19.17	17.85	19.55	13.28
CaO	16.11	1.86	1.68	1.75					6.80
Na ₂ O	0.68	0.18	0.09	0.11					1.22
K ₂ O									1.50
Total	99.66	100.77	99.42	100.72	100.63	99.93	100.39	100.22	91.21
Cation	Mg ₄₉₋₅₀	Mg ₇₈	Mg ₇₅₋₇₆	Mg ₇₈₋₇₉	Mg ₆₄₋₆₅	Mg ₆₉₋₇₀	Mg ₆₄₋₆₅	Mg ₇₀	Mg ₄₅₋₄₆
ratio %	Ca ₃₇₋₃₈	Fe ₁₈	Fe ₂₀₋₂₁	Fe ₁₇₋₁₈	Fe ₃₁₋₃₂	Fe ₃₀₋₃₁	Fe ₃₅	Fe ₃₀	Fe ₃₇₋₃₈
	Fe ₁₃	Ca ₄	Ca ₃₋₄	Ca ₃₋₄	Cr ₄₋₅	Ti ₀₋₁	Cr ₀₋₁	Ti ₀₋₁	Ca ₁₆₋₁₇

Cpx (clinopyroxene). Opx (orthopyroxene). Spl (spinel). Amp (amphibole)
WW4 Garnet-metawehrlite. WW9 Garnet-metawebsterite. WW12 Garnet-spinel-metawebsterite. WW14. Metaclinopyroxenite. WW15
Garnet-amphibole-metawebsterite. Analyst L. R. Raynor.

Mg/(Mg+Fe²⁺) for the metaharzburgite (0.923) compared to metalherzolite (0.906) indicates its more depleted nature. Two potential sources exist for the metaharzburgite xenolith. One is a highly-depleted mantle component, while the other is a higher level fragment derived from an underlying, unexposed Cambrian ultramafic complex as seen outcropping elsewhere in northwestern Tasmania (Brown 1988). Cambrian harzburgites, however, contain Cr-rich spinels with higher 100xCr/(Cr+Al) and 100xMg/(Mg+Fe²⁺) values (Brown 1988, fig. 8) than the Table Cape vent metaharzburgite (Cr value 0.50, Mg value 0.65). This favours a separate origin for the metaharzburgite, which although a scarce lithology in Southeast Australian mantle peridotite suites, can be significant at rare sites (e.g., Mt Leura, Victoria; Chen *et al.* 1989). The calculated bulk analysis and mineralogy for the Table Cape vent metaharzburgite match those of the high Mg value (0.92–0.93) Mt Leura harzburgites, rather than the lower Mg value (0.87–0.90) harzburgites that Chen *et al.* (1989) interpreted as cumulates. The Table Cape vent metaharzburgite pyroxenes have higher Ca in the opx and lower Na and Al in the cpx than the Mt Leura examples. They seem a deeper, higher T variant than at Mt Leura where they dominate in the uppermost mantle zone.

Garnet-metawebsterites dominate the Table Cape vent garnet-metapyroxenites. A bulk analysis gives a relatively low Si, high Mg composition (Sutherland *et al.* 2004) but with significantly lower Mg/(Mg+Fe²⁺) than for the metaperidotites (0.79 *cf.* 0.90). The 'mantle' norm is dominated by Hy (37%) and Cats (33%), with subordinate Ol (12%). This suggests a recrystallised mafic parental cumulate, either from transitional olivine basalt or olivine tholeiite composition. The high Ni (419 ppm) and Cr (387 ppm) and low Zr (47 ppm), Ba (7 ppm) and Rb (3 ppm) are compatible with a relatively mafic cumulate or little fractionated basalt melt. Although high Ni and Cr can be introduced by mantle wall rock-melt interactions (O'Reilly *et al.* 1989), this is considered unlikely here as no composite xenoliths are evident. The present neoblastic assemblage of cpx, opx, spl and grt formed from progressive recrystallisation of an original magmatic parent along reactions similar to those outlined from western Victorian suites (O'Reilly *et al.* 1989). The rarer wehrlites and olivine-bearing clinopyroxenite at Table Cape vent may represent less re-constituted lithologies or possibly assemblages derived from more alkaline parental melts. Their calculated bulk compositions (table 6) and 'mantle' norms (Cats 34–37, Hy 35–37, Ol 9–16 wt%) overlap those of garnet-metawebsterites. The Table Cape garnet-metapyroxenites are essentially 'dry' assemblages and differ from the hydrous amphibole-mica-bearing suites in some SE Australian xenolith suites (e.g., Bullenmerri, Gnotuk, Anakie East; Graham *et al.* 2003). They also differ in mineral and bulk compositions from 'dry' garnet-metapyroxenite suites that typify many eastern Australian suites (Irving 1974, Stolz 1984, Griffin *et al.* 1987), in being more Fe-rich. Relatively little experimental work is available on Australian garnet-metawebsterites, except for an Fe-poor Delegate example (Irving 1974). This work gave a TP point for the co-existing grt, cpx and opx at 1200°C and 1.5 GPa, which lies on the high T side (fig. 2) of the Table Cape vent Fe-rich metawebsterite re-equilibration conditions (1055–1100°C, 1.2–1.4 GPa).

Metawebsterites, lacking garnet, form a minor component at Table Cape vent and include two distinct types. The more common type consists of a polygonal-textured pyroxene

mosaic with opx compositions that overlap those in the garnet (\pm spinel)-metawebsterites (see tables 4 and 5). This suggests links between these two assemblages. Calculated bulk analyses, however, are notably higher in Si and Mg and lower in Al and Fe for the metawebsterites (table 4) and give 'mantle' norms with high Hy (51 wt%), moderate Di (26 wt%), minor Cats (10 wt%) and negligible or no Ol. This suggests a tholeiitic parental affinity, unless the metawebsterites are only pyroxene-rich parts from inhomogeneous garnet (\pm spinel)-metawebsterite bodies. Alternatively they may be recrystallisations of coarse subcalcic clinopyroxenites, originally crystallised as liquidus precipitates, as proposed for some metawebsterite xenoliths near Glen Innes, New South Wales (Wilkinson & Stolz 1997). This origin is favoured by mutual unmixing in the pyroxenes and a bulk recalculated composition close to that of a subcalcic clinopyroxene. The Table Cape vent metawebsterites, however, have lower Ca and higher Ti than in the Glen Innes subcalcic clinopyroxene (table 4, *cf.* with Wilkinson & Stolz, tables 5 and 10). In contrast, the other type of Table Cape vent metawebsterites approach Cr-bearing diopside in composition (table 4) and are higher in Ca and lower in Mg and Fe than the Ti-Al-rich websterites. The accessory Ti-bearing phlogopite and apatite suggest an introduced metasomatic component enriched in Ti, K and P. The cloudy apatite resembles that in diopsidic vein material within metasomatised metalherzolites in south-eastern Australia (O'Reilly & Griffin 2000). Such mantle apatites in southeastern Australian xenoliths come from a relatively young metasomatic event, originally assigned to a deep (plume-related) fluid. Re-examination of the apatite neon isotope data, however, now suggests it derives from Mid-Ocean Ridge Basalt (MORB) fluids rather than a primitive plume source (Gautheron & Moreira 2003).

The crustal granulites are two-pyroxene, Ca-rich feldspar assemblages and include both anhydrous and hydrous metasomatised types. An anhydrous banded granulite (Sutherland *et al.* 2004) corresponds chemically to a transitional olivine tholeiite. It equates to a basalt protolith (SiO₂ 51, Di 25, Hy 22, Ol 4 wt%) of relatively unfractionated nature, with Mg/(Mg+Fe²⁺) = 0.67 and An/(An+Ab) = 0.59. The hydrous micro-granulite exhibits a diverse accessory mineralogy that includes Ti-bearing amphibole and mica, rutile-ilmenite and rare corundum. It is even more Mg-rich and primitive in its features than the anhydrous granulite, with Mg/(Mg+Fe²⁺) = 0.74 and An/(An+Ab) = 0.68, high Ni+Cr (522 ppm) and low Zr (53 ppm) contents. It also corresponds to transitional olivine basalt in its SiO₂ (50 wt%) and other normative contents (Hy 32, Di 9, Ol 2 wt%). The accessory minerals and textures suggest modification of a basaltic parent by volatile, Al, Ti, Fe and K-bearing fluids prior to metamorphic re-equilibration.

The garnet-metawebsterites and two-pyroxene granulites in the Table Cape vent suite suggest transitional olivine basalt protoliths were emplaced at both mantle and crustal levels. Normative wt% ratios for these rocks (Hy 44–75, Di 20–48, Ol 8–23) plot across relatively unoccupied positions in the Ol-Di-Hy normative tetrahedron for eastern Australian assemblages (O'Reilly *et al.* 1989). This supports an unusual lithospheric section below Table Cape, in keeping with the moderately Fe-rich pyroxene mineralogy in these assemblages. The Table Cape vent calcic plagioclase granulites differ from granulite in the Boat Harbour section, which contains Na-rich plagioclase (Sutherland *et al.* 1996).

Although more silica-saturated protoliths dominate the Table Cape vent Al-Ti-bearing augite suites, rare metawehrlite and garnet-metaclinopyroxenites suggest more olivine-rich and perhaps even alkaline progenitors may have been present.

Tasmanian/East Australian Geotherms

The SEA geotherm was established using Ellis & Green (1979) grt-cpx exchange thermometry, which, while accommodating Fe²⁺ and Fe³⁺ assignments in the minerals, generally leads to wider temperature differences between these allocations than does the Wells thermometry (Griffin *et al.* 1984, O'Reilly & Griffin 1985). The Ellis & Green thermometer, however, has proved unreliable in its application. It generally overestimates T relative to experimental results for the TP range for such xenolith suites (see Sutherland *et al.* 1994, 1998, Ai 1994, Taylor 1998) or is erratic where Ca exceeds a concentration of 0.15 atoms in the garnet (Alaoui *et al.* 1997). The Table Cape vent garnets all have Ca <0.15, but application of Ai (1994) thermometry nevertheless gives highly erratic results. For these reasons, grt-cpx exchange thermometry is abandoned for the Table Cape vent grt-bearing assemblages, in favour of Wells thermometry. The comparison in T range between Ellis & Green (Fe³⁺) and Wells (Fe³⁺) for Table Cape vent grt-websterites is 1005–1220°C and 1055–1075°C respectively, which leads to corresponding differences in P estimates when inserted into the Wood barometer (e.g., 0.74–1.86 GPa, Sutherland *et al.* 1994, cf 1.23–1.36 GPa, Sutherland *et al.* 2004).

Wood (1974) barometry, however, is retained for the Table Cape vent study as a suitable barometer for Fe-rich grt-metawebsterites, particularly as some other potential barometers were experimentally calibrated against more Mg-rich assemblages. This includes Taylor (1998) thermobarometry, which provides a lower intersection than the original SEA gradient. The new Table Cape vent thermobarometry indicates a coherent grouping for grt-websterites near the base of the sampled mantle section at temperatures well above SEA (fig. 2). The Table Cape vent TP section extends to 45 km depth, so represents a higher level section than at Bow Hill, Tasmania, which extends into the spinel-garnet-metalherzolite transition.

The Mg-rich Bow Hill grt-assemblages (Sutherland *et al.* 1984) need different thermobarometry than for Table Cape assemblages. With Wells (1977) thermometry (with only Fe²⁺ allocated) and Wood (1974) barometry, the wide range of results (1000–1180°C, 1.0–2.5 GPa) extend well beyond the experimental garnet-spinel lherzolite transition zone, suggesting an unsuitable coupling. The Taylor (1998) TP method, which gives T estimates closest to Wells T, was chosen as a more appropriate coupling. In calculating TP, Fe²⁺ and Fe³⁺ was allocated in the garnets (Ca concentration 0.12 to 0.14 atoms) by charge balance and in the pyroxene by the cation allocation method of Cawthorn & Collerson (1974). The results (I. Roach, Australian National University, pers. comm. 2004) range from 1048–1065°C, 1.63–1.69 GPa (grt-spl-metalherzolite) to 1205–1213°C, 2.21–2.24 GPa (grt-metalherzolite) and 1213°C and 2.05 GPa (grt-metawebsterite). These results lie on the high T side of the Taylor (1998) modified SEA geotherm (T-T, fig. 2) and the higher P assemblages even lie above the original SEA geotherm. Thus, both Table Cape and Bow Hill suites suggest that extreme geothermal gradients existed in the Tasmanian Miocene lithosphere.

The precise profiles and extent of variations within Mesozoic-Cainozoic East Australian geotherms have engendered debate as to whether the TP profiles overestimate the data or mark more conservative profiles, depending on the choice of thermometer/barometer couplings and other effects (McDonough *et al.* 1991, Sutherland *et al.* 1994, 1998, O'Reilly *et al.* 1997, Sutherland 1999, Gaul *et al.* 2003, Roach 2004). Various protocols for thermometer and barometer usage in general, that incorporate differences in mineralogical compositions of assemblages, have been proposed (O'Reilly *et al.* 1997, Xu *et al.* 1998, Smith 1999). These suggested couplings do not include some well-based newer couplings (Taylor 1998 TP). Thus, uncertainties exist in geotherm comparisons for the East Australian/East Asian xenolith suites. In a study of xenoliths in Miocene basalts from northern China, Chen *et al.* (2001) preferred Brey & Köhler (1990) TP for the Mg-rich garnet-websterites, rather than the Ellis & Green (1979) and Wood (1974) TP, and the results clearly plot well above the SEA geotherm which was based on the latter coupling. However, they retained Ellis & Green T and Wood P for an Fe-rich garnet-metawebsterite, which placed this plagioclase-bearing assemblage at over 70 km depth, well-divorced from the other TP plots.

Despite the complications in selecting thermobarometric methods, the two revised Tasmanian Miocene geotherms both suggest elevated gradients that exceed both SEA and the modified SEA (T-T, fig. 2) gradients. This supports a range of gradients for eastern Australian Mesozoic-Cenozoic geotherms, going from cooler gradients (e.g., Mesozoic Ruby Hill and Jugiong suites; Sutherland 1999, Gaul *et al.* 2003) into higher gradients (e.g., Cenozoic western Victoria SEA; O'Reilly *et al.* 1997, Taylor 1998) and some more extreme gradients (e.g., Cenozoic Tasmania, this study). The exact geotherm gradient becomes critical when correlating other lithological temperatures, uncontrolled by pressure, as for spinel-metalherzolites, garnet-metaclinopyroxenites and non-garnetiferous granulites, leading to potential depth uncertainties in crust-mantle sections (Sutherland *et al.* 1998).

Some temperature intervals for Table Cape vent assemblages are shown in relation to the TP controlled segments of Tasmanian Miocene geotherms (fig. 2). The spinel-metalherzolite T ranges at Table Cape intersect at up to 1.15 GPa on the modified SEA geotherm, up to 1.0 GPa on the SEA geotherm and extrapolate to 0.8 GPa on a geotherm 80°C higher than SEA. Similarly, granulites at Table Cape intersect at up to 1.05 GPa on modified SEA, up to 0.9 GPa on SEA and extrapolate to 0.75 GPa on SEA + 80°C geotherms. These potential differences in lithological assignments (0.3 GPa) graphically illustrate their dependence on the precise geotherm, and are further complicated by pressure-dependant variation in metaperidotite T estimates using Witt-Eickschen & Seck (1991) thermometry.

The seismic data for the Table Cape region suggests a Moho depth at 32 km (Rawlinson *et al.* 2001), i.e., 1.05–1.1 GPa, which would give T intersections of 870°C on modified SEA, 920°C on SEA and 1000°C on SEA + 80°C geotherms. The first two T intersections are compatible (within error) with the Table Cape granulite-metalherzolite interval (870–915°C), but less so for the SEA (+80°C) geotherm. Thus, an apparent discrepancy exists between more elevated gradients in the lower (grt-metawebsterite) part of the section and reduced gradients at the mantle-crust transition zone. Unless all the thermobarometry (past and present) on Table Cape garnet-metapyroxenites is seriously

in question, this implies a localised thermal anomaly.

One possibility for such a thermal anomaly is local heating of mantle adjacent to magma chambers. Such mantle magma chambers exist nearby at Boat Harbour, where basanite magmas were fractionated to erupt as lherzolite-bearing nepheline mugearites (Sutherland *et al.* 1996). The Boat Harbour fractionated lavas are dated at 14.2 ± 0.1 Ma, just prior to eruption of the Table Cape centre (minimum 13.3 ± 0.2 for late stage lavas). Thus, additionally heated mantle may have already been present when fractionated olivine nephelinite brought up the Table Cape xenolith suite. This scenario would represent small-scale local perturbation of a prevailing geotherm, within a limited segment of the lithosphere. It would contrast with the western Victorian scenario where extensive basaltic intrusions around the mantle-crust boundary were identified on seismic sections and were considered to generate the SEA geotherm (Cull *et al.* 1991). The Bow Hill TP plots also suggest a declining geothermal gradient, so that thermal contribution from deep mantle magma chambers that fractionated the host nepheline hawaiite there also needs consideration.

SYNTHESIS

The Table Cape vent xenolith suite (erupted with evolved nephelinite) represents a unique lithospheric section, both within Tasmania and elsewhere among eastern Australian xenolith suites. It is one of two known garnet-bearing (two-pyroxene) xenolith suites in Tasmania, but represents a higher level mantle section with more Fe-rich minerals in the assemblages than occurs at Bow Hill.

Revised thermobarometry for Tasmanian garnet-bearing xenoliths confirmed that the associated Miocene geotherms in part exceeded the western Victorian (SEA) gradient.

Considerations of xenolith temperatures for the crust-mantle boundary lithology at Table Cape, in relation to the seismic Moho (32 km depth) and proposed SEA and Table Cape geotherms, suggest the Table Cape geotherm may mark a locally highly perturbed gradient.

The Table Cape geotherm may reflect thermal input from adjacent magma chamber activity that gave 14 Ma Boat Harbour fractionated basalts, prior to Table Cape activity.

Xenolith-derived geotherms related to Mesozoic-Cenozoic basaltic activity have been constructed from many eastern Australian and Asian sites. However, not only do they involve complications in thermobarometry in defining such geotherms, but also incorporate diverse magmatic thermal inputs that may include localised effects.

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