Igneous stratigraphy and internal structure of the Little Minch Sill Complex, Trotternish Peninsula, northern Skye, Scotland

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Abstract – Detailed sampling of the Little Minch Sill Complex reveals that it is composed of both single and multiple sills. These are formed of three main, genetically related units: picrite, picrodolerite and crinanite, which are the result of differentiation of an alkali-olivine basalt magma (approximately 10% MgO) in an upper-crustal magma chamber. Variations in igneous stratigraphy and the presence of internal chills in the Trotternish sills suggest that they were emplaced by multiple intrusion and subsequently differentiated *in situ*. Changes in petrography adjacent to pegmatite veins and textures within picrite units indicate compaction and filter-pressing were important processes after emplacement. Rhythmic layering (1 cm to 1 m thick) is conspicuous in the sills near contacts but does not involve cryptic mineral variation. Such modal layering may be more common than realised in relatively small-scale intrusions and may be modelled in terms of *in situ* differentiation under conditions of significant undercooling in a changing thermal gradient at the synthetic for-sterite-diopside-anorthite eutectic.

1. Introduction

A large Tertiary sill complex, the Little Minch Sill Complex (LMSC; Gibb & Gibson 1989), outcrops on the Trotternish Peninsula of northern Skye and adjacent islands, which include the Shiant Isles and Raasay (Fig. 1). The LMSC covers a large area (4000 km²) and represents a considerable volume of magma. Most of the published work on the LMSC relates to the classic sills on the Shiant Isles (Walker, 1930; Drever & Johnston, 1957, 1965; Gibb, 1973; Gibb & Henderson, 1984, 1989). The sills on the Trotternish Peninsula have formed the basis of relatively few publications (Walker, 1931, 1932; Anderson & Dunham, 1966; T. Simkin, unpub. Ph.D. thesis, Univ. Princeton, 1965; Simkin, 1967) considering their excellent lateral exposure and relatively unaltered nature. This study is primarily concerned with the field relations and petrography of the Trotternish sills, and the geochemistry and petrogenesis of the LMSC are discussed elsewhere (S. A. Gibson, unpub. Ph.D. thesis, Kingston Polytechnic, 1988; Gibson, 1990).

2. Geological setting

The most extensive and laterally persistent outcrops of the LMSC occur on the Trotternish Peninsula where the thickness of individual sills varies from approximately 10 m to 130 m. The LMSC post-dates the overlying Tertiary Skye lavas but is intruded by Tertiary late-stage basic dykes. The sills dip gently westward and transgress upwards through Jurassic sediments in a northerly direction. Lower contacts of the sills with Jurassic sediments are exposed at several localities, e.g. Meall Tuath and Rigg (localities 8 and 18, Fig. 2), but upper contacts are rare. It is unlikely that the whole thickness of the LMSC is completely exposed on the Trotternish Peninsula.

3. Igneous stratigraphy

Detailed field mapping and petrographic evidence suggest that the sills exposed on the Trotternish Peninsula (Fig. 2) are composed of either a single lithology, such as those at Osmigarry, Heribusta and Bornaskitaig (localities 6, 5, 4), or multiple lithologies, such as those at Meall Tuath and Tottrome (localities 8, 19). Three main lithologies are recognizable in the Trotternish sills: picrite, picrodolerite and crinanite (the definitions of Gibb & Henderson [appendix, 1984] that incorporate the traditional names for the Shiant Isles material are used in this study). The multiple sills are generally formed of a lower picrite unit overlain by a crinanite and/or picrodolerite unit. In general the picrite maintains a regular thickness (approximately 50 m) across the Trotternish Peninsula. This is greater than the thickness of picrite in the main Shiant Isles sill (30 m; Gibb & Henderson, 1984). At Osmigarry a lower picrite sill, which also outcrops at Bornaskitaig, is separated from an upper crinanite/picrodolerite (which also outcrops at Heri-

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Figure 1. Outcrop of the Little Minch Sill Complex (after Chesher, Smythe & Bishop, 1983).

busta) by 7 m of sediment (Fig. 3). At Creagan Iar, 2 km north of Osmigarry, the picrite and crinanite/ picrodolerite sills converge and the intervening sediments are absent. This picrite/picrodolerite contact is laterally extensive across the Trotternish Peninsula and can be traced north to Meall Tuath, east to Flodigarry and possibly southeast to Tottrome (Fig. 2). No evidence of chilling is present at this contact which resembles the picrite/picrodolerite 'discontinuity' in the main Shiant Isles sill (Drever, 1953; Gibb & Henderson, 1989).

The picrite sill which outcrops at Rigg may be an offshoot of the sill at Tottrome. The crinanite sill

which outcrops at Holm (NG 519 514; 25 m thick) and Leac Tressirnish (10 m) decreases in thickness northwards and forms the upper unit (5 m) of the multiple sill at Tottrome (Fig. 2). The relationship of the crinanite sills, which outcrop along the east coast between Culnaknock and Staffin, and the rest of the complex is unclear. They lie on the upthrown side of a NW-SE fault, which separates them from sills at Dun Raisburgh and Dun Connavern, and are at a relatively low level in the stratigraphy of the LMSC. A crinanite sill (> 20 m thick) is exposed beneath the main multiple sill at Meall Tuath and may be related to these crinanite sills and also the lower crinanite sill



Figure 2. Variations in igneous stratigraphy of the Trotternish sills. Outcrop of the sills on the Trotternish Peninsula is from Anderson & Dunham (1966). The stratigraphy of the Shiant Isles (Gibb & Henderson, 1984) is shown for comparison. Sample localities are as follows: (1) Skudiburgh; (2) Kilbride Point; (3) Sgeir Lang; (4) Bornaskitaig; (5) Heribusta; (6) Osmigarry; (7) Creagan Iar; (8) Meall Tuath; (9) Balmaquien; (10) Flodigarry; (11) Staffin; (12) Loch Mealt; (13) Dun Raisburgh; (14) Culnaknock; (15) Dunn Connavern; (16) Invertote; (17) Leac Tressirnish; (18) Rigg; (19) Tottrome; (20) Bearreraig.



Figure 3. Transgressive relationships and variations in igneous stratigraphy of the sill in northwest Trotternish.

on Garbh Eilean, Shiant Isles (Gibb & Henderson, 1984).

4. Internal structure

4.a. Internal contacts

Well documented internal discontinuities are present in the main Shiant Isles sill at the picrite/picrodolerite junction (Drever, 1953; Gibb, 1973; Gibb & Henderson, 1984, 1989) and at the crinanite/picrite junction (Gibb & Henderson, 1984, 1989). An upper picrite unit has not been observed in any of the sills exposed on the Trotternish Peninsula but abrupt petrographic and geochemical changes occur at the picrite/picrodolerite junction (sections 5 & 7). Internal chills are present within the Trotternish sills at several localities. Such chills are highly significant in modelling the emplacement of the LMSC and have not been recorded previously in this area. At Skudiburgh and Meall Tuath internal chills occur within 5 m of the lower contact of the picrite sill with sediments. At both localities the upper picrite unit appears to be chilled against the lower one. At Creagan Iar an internal chill is present near the top of the sill. The upper crinanite unit (> 5 m thick) is chilled against the lower crinanite/picrodolerite unit (12 m). An internal chill is also present within a few metres of the crinanite sill/sediment contact on Staffin Island [NG 490 690]. The upper unit appears to be chilled against the lower unit. In a recently exposed quarry section at Invertote (Fig. 4) a crinanite unit (5 m) is clearly chilled against a lower picrite unit (42 m). Further internal contacts are visible in the cliff at Kilt Rock [NG 509 655]. At Kilbride Point a lower picrite unit (20 m) is separated from an upper picrite unit (> 15 m) by 3 m of baked mudstone (Skudiburgh Formation). Both units are chilled against the sediment, which may be a sediment raft incorporated into the centre of the picrite sill or, more likely, a wedge of sediment which separates two sills.

4.b. Layering

The occurrence of rhythmic layering in the LMSC has been implied by Bell & Harris (1986) and Brown (1969), and recorded in the syenites of Eilean Mhuire, Shiant Isles (Drever & Johnston, 1957). Such layering is rarely described in relatively small-scale intrusions (e.g. Dunham, 1965) but is present at several localities in the LMSC. It usually occurs adjacent to sill contacts and is most noticeable on weathered surfaces. The layering is typically developed on a centimetre scale within a metre of the contact but is sometimes more extensive. Good examples occur in the cliffs beneath Duntulm Castle [NG 409 744], at Rubha nam Brathairean [NG 527 628], Flodigarry, Leac Tressirnish, Meall Tuath (Fig. 2), the Ascrib Islands [NG 304 632] and Flodigarry Island [NG 483 716], where

layering has developed on a 1 m scale up to 40 m away from the lower contact between the sill and sediment (Fig. 5). At localities where sills make major transgressions, such as at Duntulm [NG 410 730], large rafts (up to 10 m) of Jurassic sediment have been incorporated into the sills and layering has developed parallel to the surfaces of these rafts. Here, layer thickness increases but layer characteristics become less pronounced away from the contact. In general each layer has a mafic base which gradationally becomes more felsic and resistant to weathering upwards. The junctions between layers are abrupt. In thin section the layering at Rubha nam Brathairean is well defined by the variation in modal proportions of olivine, clinopyroxene and plagioclase (Fig. 6). No marked variation in mineral chemistry (cryptic layering) has been detected. The mafic bases are olivinerich (14%) with large (up to 6 mm), ophitic, zoned clinopyroxene (29%) which enclose laths (< 2.5 mm) of plagioclase feldspar (51%). Titaniferous magnetite occurs interstitially, sometimes partially enclosing plagioclase, and forms approximately 6% of the less resistant layers. With increasing height above the mafic base, there is a corresponding increase in plagioclase (59%) and a decrease in olivine (7%) and clinopyroxene (22%). The increase in clinopyroxene is accompanied by a change in habit, from large ophitic plates to small granules (< 2 mm). Once more, the top of some layers is marked by a distinctive finegrained feldspathic horizon. The isolation of individual large laths of plagioclase by poikilitic plates of clinopyroxene suggests in situ crystallization. The distance between the plagioclase and olivine crystals is too great for them to have been supported by grainto-grain contact. The absence of cross-cutting layers and the relatively small thickness of the sills also support an origin by in situ crystallization. In the Trotternish sills layering commonly occurs close to contacts with sediments. At the time of emplacement, large differences in temperature between the magma and Jurassic sediments would produce large thermal gradients and a large ΔT at the contacts, where a boundary layer may have formed within the magma. In the Trotternish sills this appears to have been on a centimetre scale and the changing thermal gradient was a significant factor in the formation of layering (S. A. Gibson, unpub. Ph.D. thesis, Kingston Polytechnic, 1988). In larger intrusions such as Rhum and Skaergaard, ΔT would be small (perhaps < 5 °C) and of only minor significance at the margins.

By analogy with phase relationships in experimental Fe-free basaltic systems, undercooling at the forsterite-diopside-anorthite eutectic may account for the alternating phase relationships (S. A. Gibson, unpub. Ph.D. thesis, Kingston Polytechnic, 1988). After crystallization of spinel, the liquid would be driven towards the forsterite-diopside-anorthite eutectic. If undercooling occurred at this point olivine,



Figure 4. Quarry section at Invertote shows the contact between a lower picrite (P) unit (42 m) and an upper crinanite (C) unit (5 m). A pegmatite vein (Peg) occurs near the top of the picrite unit.

being the easiest phase to nucleate, would crystallize and be accompanied by clinopyroxene. Clinopyroxene would have a relatively high growth/nucleation ratio and would become ophitic towards plagioclase. Olivine crystallization would drive the liquid towards the diopside-anorthite cotectic. At relatively small degrees of undercooling, clinopyroxene would have a high nucleation rate and form granular crystals (Carmichael, Turner & Verhoogen, 1974). As clinopyroxene crystallized the liquid would become enriched with respect to plagioclase and would produce the feldspathic unit which is sometimes present at the top of layers. The relative rates of nucleation and growth appear to have been more important than the sequence of phases at the liquidus in the formation of layering at the margins of the Trotternish sills.

5. Petrography

Petrographic variations in the LMSC relate primarily to modal differences in olivine, clinopyroxene and plagioclase. The main lithologies picrite, picrodolerite



Figure 5. Rhythmic layering in southeast Flodigarry Island. Height of cliffs is approximately 60 m. The base of the sill is at sea level.

and crinanite are also distinguished by textural variations and mineral chemistry (Fig. 7). Representative analyses of mineral compositions are presented in Table 1 and clinopyroxene crystallization trends in Fig. 8.

5.a. Picrites

The most distinctive feature displayed by the picrites is their very high content of modal olivine (generally 40-60%). The olivine occurs as either large (up to 5 mm in length) euhedral phenocrysts or as slightly smaller subhedral phenocrysts. Occasionally the olivines have re-entrants. The olivine core compositions are almost constant (Fo₈₂-Fo₇₉) with slight Feenrichment towards rims (Fig. 7). The olivine is poikilitically enclosed by clinopyroxene (100(Mg/ Mg + Fe = 84.1-74.6, which forms 9-24% of the rock and only rarely occurs as isolated crystals. Plagioclase forms 27-46% of this lithology; small laths (< 0.5 mm) often occur in aggregates enclosed by the ophitic clinopyroxene, and slightly larger, unenclosed plagioclase phenocrysts (up to 1 mm) show normal zonation (An₆₄-An₃₁). In some picrite samples, plagioclase is tangentially arranged around olivine crystals (Fig. 9). Chrome spinel (which is always enclosed by olivine), ilmenite and analcime are present as minor phases (< 2%).

A local variation of the picrites shows large poikilitic laths of plagioclase (< 6 mm) which sub-enclose olivine. The olivine may have a skeletal appearance and a maximum length of 6 mm. Clinopyroxene is no longer ophitic, but is euhedral/subhedral and sector zoned. Some of the clinopyroxene has a poikilitic relationship with olivine and is itself partially enclosed by plagioclase.

In the picrite sills a distinct textural variation occurs within 5 m and 10 m of the lower and upper contacts respectively. In these 'marginal picrites' the olivine content decreases towards the contacts (from 40% to 7%) and is accompanied by a gradual decrease in grain size of the groundmass. At the sill contacts the olivine phenocrysts are euhedral to subhedral, 2 mm in length with re-entrants and often pseudomorphed by iddingsite (Fig. 10). A few metres away from contacts with sediments, fresh olivine phenocrysts are strikingly zoned (Fo₇₆-Fo₂₂). Plagioclase is also present as a phenocryst phase (0.5 mm in length) and shows normal zonation (An₆₇-An₆₄). The plagioclase phenocrysts commonly form open spherulites which are sometimes intergrown with olivine.

Pegmatite veins (up to 1 m thick) are common within the picrites and are essentially composed of clinopyroxene (~ 30%), plagioclase (~ 65%), ilmenite and zeolites. Olivine is absent. Clinopyroxene is present as either large (1.5–14.0 mm in diameter), subhedral phenocrysts or small (< 0.5 mm) acicular crystals in the groundmass. Sector zonation is conspicuous in both generations of clinopyroxene. Some clinopyroxenes, adjacent to alkali-rich mesostases, show extreme Fe-enrichment (Fig. 8) and green aegirine rims. These contain up to 13 wt.% Na₂O (Table 1) and approximately 6 wt.% ZrO₂ (qualitative analysis). Plagioclase laths are generally less than



Figure 6. Photomicrograph of rhythmic layering in the Trotternish sills. Most of the olivine is altered to iddingsite and the strongly birefringent mineral is clinopyroxene. (1) is the less resistant olivine-rich horizon which occurs at the base of layers; (2) is a horizon of granular clinopyroxene which overlies (1); and (3) is the resistant feldspathic horizon which is present at the top of layers.

		Picrite SG114		Marginal picrite SG223		Picrodolerite SG19		Crinanite SG40		
		OL(C)	OL(R)	OL(C)	OL(R)	OL(C)	OL(R)	OL(C)	OL(R)	
••••••••••••••••••••••••••••••••••••••	Wt. %							Ji	- 1996 A	
	SiO,	39.05	38.73	39.10	32.29	38.66	38.43	35.80	33.68	
	Al,Õ,	0.29	0.17	1.07	1.19	n.d.	n.d.	0.69	1.00	
	FeO	16.89	20.62	20.96	52.64	24.32	25.12	34.88	45.90	
	MnO	0.34	0.18	0.29	1.09	0.26	0.40	0.31	0.96	
	MgO	42.39	39.45	38.11	10.29	35.15	34.35	25.77	15.85	
	CaO	0.52	0.45	0.64	0.99	0.55	0.57	0.58	0.64	
	NiO	0.26	0.17	0.11	n.d.	0.10	0.21	0.19	0.32	
	Total	99.74	99.77	100.28	98.50	99.04	99.08	98.22	98.35	
	Atoms per	4 oxygens								
	Si	0.984	1.002	0.993	1.003	1.019	1.019	1.009	1.009	
	Al	0.006	0.003	0.021	0.044	0.000	0.000	0.023	0.035	
	Fe	0.984	1.002	0.445	1.368	0.553	0.557	0.821	1.149	
	Mn	0.005	0.003	0.004	0.029	0.003	0.009	0.007	0.024	
	Ma	1 502	1 554	1 443	0.025	1 379	1 350	1 083	0.707	
	Co	0.000	0.008	0.011	0.477	0.015	0.016	0.017	0.707	
	Ni	0.003	0.008	0.011	0.000	0.013	0.010	0.017	0.021	
	-	0.005	0.005	0.002	0.000	0.002	0.002	0.005	0.007	
	Fo	82	77 	76	26	72	71	38	57	
	Picrite SG21		Marginal picrite SG128		Pirodolerite SG251		Crinanite SG358		Pegmatite SG31	
	CPX(C)	CPX(R)	CPX(C)	CPX(R)	CPX(C)	CPX(R)	CPX(C)	CPX(R)	CPX(C)	CPX(R)
Wt %			·····							
SiO ₂	50.11	50.45	48.21	52.36	50.51	51.35	52.85	50.34	50.67	51.82
Al,Ō,	4.09	3.44	6.53	2.05	3.47	3.45	3.74	2.06	3.61	1.63
TiÔ	0.55	1.05	2.09	0.53	1.15	1 49	1.00	1.00	1.68	4.24
r 0 ⁴	0.55	1.95				1.72			1.00	
Feu	5.71	^{1.95} 7.81	8.99	14.45	7.27	10.16	6.94	15.46	7.58	
FeO Fe _n O _n	5.71	· 7.81	8.99	14.45	7.27	10.16	6.94	15.46	7.58	23.61
FeO Fe ₂ O ₃ MnO	5.71 0.13	7.81 0.16	8.99 0.26	14.45 0.48	7.27 0.18	0.32	6.94 0.16	0.29	7.58	23.61 0.27
Fe ₂ O ₃ MnO MgO	0.13 0.13 15.89	1.95 7.81 0.16	8.99 0.26	14.45 0.48 8.88	7.27 0.18 14.40	0.32 12.39	6.94 0.16 13.91	0.29 9.45	7.58 0.28 14.96	23.61 0.27
FeO Fe ₂ O ₃ MnO MgO C2O	0.13 5.71 0.13 15.89 20.29	0.16 14.65	8.99 0.26 11.86 21.11	14.45 0.48 8.88 21 18	7.27 0.18 14.40 20.29	0.32 12.39 20.28	6.94 0.16 13.91 20.54	0.29 9.45 18.97	0.28 14.96	23.61 0.27 1.14 3.53
FeO Fe ₂ O ₃ MnO MgO CaO Na O	0.13 5.71 0.13 15.89 20.29	1.95 7.81 0.16 14.65 19.94	8.99 0.26 11.86 21.11 1.03	14.45 0.48 8.88 21.18	7.27 0.18 14.40 20.29 0.90	0.32 12.39 20.28	6.94 0.16 13.91 20.54 0.83	15.46 0.29 9.45 18.97 0.73	7.58 0.28 14.96 19.05	23.61 0.27 1.14 3.53
FeO Fe ₂ O ₃ MnO MgO CaO Na ₂ O Cr O	0.13 5.71 0.13 15.89 20.29 1.95	1.95 7.81 0.16 14.65 19.94 1.04	8.99 0.26 11.86 21.11 1.03 0.09	14.45 0.48 8.88 21.18 n.d. 0.07	7.27 0.18 14.40 20.29 0.90 0.37	1.49 10.16 0.32 12.39 20.28 n.d.	6.94 0.16 13.91 20.54 0.83 0.57	15.46 0.29 9.45 18.97 0.73	7.58 0.28 14.96 19.05 1.91	23.61 0.27 1.14 3.53 12.01
FeO Fe ₂ O ₃ MnO MgO CaO Na ₂ O Cr ₂ O ₃ Total	0.13 5.71 0.13 15.89 20.29 1.95 1.12	1.95 7.81 0.16 14.65 19.94 1.04 n.d. 99.44	8.99 0.26 11.86 21.11 1.03 0.09	14.45 0.48 8.88 21.18 n.d. 0.07	7.27 0.18 14.40 20.29 0.90 0.37 08 54	10.16 0.32 12.39 20.28 n.d. 0.05	6.94 0.16 13.91 20.54 0.83 0.57	15.46 0.29 9.45 18.97 0.73 0.08	7.58 0.28 14.96 19.05 1.91 0.09	23.61 0.27 1.14 3.53 12.01 n.d. 08.25
FeO Fe ₂ O ₃ MnO MgO CaO Na ₂ O Cr ₂ O ₃ Total	0.13 5.71 0.13 15.89 20.29 1.95 1.12 99.84	1.95 7.81 0.16 14.65 19.94 1.04 n.d. 99.44	8.99 0.26 11.86 21.11 1.03 0.09 100.17	14.45 0.48 8.88 21.18 n.d. 0.07 100.00	7.27 0.18 14.40 20.29 0.90 0.37 98.54	10.16 0.32 12.39 20.28 n.d. 0.05 99.49	6.94 0.16 13.91 20.54 0.83 0.57 100.54	15.46 0.29 9.45 18.97 0.73 0.08 98.38	7.58 0.28 14.96 19.05 1.91 0.09 99.83	23.61 0.27 1.14 3.53 12.01 n.d. 98.25
FeO Fe_2O_3 MnO MgO CaO Na_2O Cr_2O_3 Total Atoms pe	0.13 5.71 0.13 15.89 20.29 1.95 1.12 99.84 er 6 oxygens	1.95 7.81 0.16 14.65 19.94 1.04 n.d. 99.44	8.99 0.26 11.86 21.11 1.03 0.09 100.17	14.45 0.48 8.88 21.18 n.d. 0.07 100.00	7.27 0.18 14.40 20.29 0.90 0.37 98.54	10.16 0.32 12.39 20.28 n.d. 0.05 99.49	6.94 0.16 13.91 20.54 0.83 0.57 100.54	15.46 0.29 9.45 18.97 0.73 0.08 98.38	7.58 0.28 14.96 19.05 1.91 0.09 99.83	23.61 0.27 1.14 3.53 12.01 n.d. 98.25
FeO Fe ₂ O ₃ MnO MgO CaO Na ₂ O Cr ₂ O ₃ Total Atoms pe Si	0.13 5.71 0.13 15.89 20.29 1.95 1.12 99.84 er 6 oxygens 1.840	1.95 7.81 0.16 14.65 19.94 1.04 n.d. 99.44 1.843	8.99 0.26 11.86 21.11 1.03 0.09 100.17 1.803	14.45 0.48 8.88 21.18 n.d. 0.07 100.00 1.989	7.27 0.18 14.40 20.29 0.90 0.37 98.54 1.885	10.16 0.32 12.39 20.28 n.d. 0.05 99.49 1.978	6.94 0.16 13.91 20.54 0.83 0.57 100.54 1.931	15.46 0.29 9.45 18.97 0.73 0.08 98.38 1.962	7.58 0.28 14.96 19.05 1.91 0.09 99.83 1.878	23.61 0.27 1.14 3.53 12.01 n.d. 98.25 2.100
FeO Fe ₂ O ₃ MnO MgO CaO Na ₂ O Cr ₂ O ₃ Total Atoms pe Si Al	0.53 5.71 0.13 15.89 20.29 1.95 1.12 99.84 er 6 oxygens 1.840 0.670	1.95 7.81 0.16 14.65 19.94 1.04 n.d. 99.44 1.843 0.148	8.99 0.26 11.86 21.11 1.03 0.09 100.17 1.803 0.288	14.45 0.48 8.88 21.18 n.d. 0.07 100.00 1.989 0.092	7.27 0.18 14.40 20.29 0.90 0.37 98.54 1.885 0.152	1.45 10.16 0.32 12.39 20.28 n.d. 0.05 99.49 1.978 0.152	6.94 0.16 13.91 20.54 0.83 0.57 100.54 1.931 0.161	15.46 0.29 9.45 18.97 0.73 0.08 98.38 1.962 0.095	1.38 7.58 0.28 14.96 19.05 1.91 0.09 99.83 1.878 0.158	23.61 0.27 1.14 3.53 12.01 n.d. 98.25 2.100 0.078
$Fe_{2}O_{3}$ $Fe_{2}O_{3}$ MnO MgO CaO $Na_{2}O$ $Cr_{2}O_{3}$ $Total$ $Atoms pe$ Si Al Ti	0.53 5.71 0.13 15.89 20.29 1.95 1.12 99.84 er 6 oxygens 1.840 0.670 0.025	1.95 7.81 0.16 14.65 19.94 1.04 n.d. 99.44 1.843 0.148 0.054	8.99 0.26 11.86 21.11 1.03 0.09 100.17 1.803 0.288 0.059	14.45 0.48 8.88 21.18 n.d. 0.07 100.00 1.989 0.092 0.015	7.27 0.18 14.40 20.29 0.90 0.37 98.54 1.885 0.152 0.032	1.47 10.16 0.32 12.39 20.28 n.d. 0.05 99.49 1.978 0.152 0.042	6.94 0.16 13.91 20.54 0.83 0.57 100.54 1.931 0.161 0.028	15.46 0.29 9.45 18.97 0.73 0.08 98.38 1.962 0.095 0.793	1.88 7.58 0.28 14.96 19.05 1.91 0.09 99.83 1.878 0.158 0.047	23.61 0.27 1.14 3.53 12.01 n.d. 98.25 2.100 0.078 0.129
FeO Fe ₂ O ₃ MnO MgO CaO Na ₂ O Cr ₂ O ₃ Total Atoms pe Si Al Ti Fe ²⁺	0.53 5.71 0.13 15.89 20.29 1.95 1.12 99.84 er 6 oxygens 1.840 0.670 0.025 0.169	1.95 7.81 0.16 14.65 19.94 1.04 n.d. 99.44 1.843 0.148 0.054 0.215	8.99 0.26 11.86 21.11 1.03 0.09 100.17 1.803 0.288 0.059 0.281	14.45 0.48 8.88 21.18 n.d. 0.07 100.00 1.989 0.092 0.015 0.459	7.27 0.18 14.40 20.29 0.90 0.37 98.54 1.885 0.152 0.032 0.227	1.47 10.16 0.32 12.39 20.28 n.d. 0.05 99.49 1.978 0.152 0.042 0.317	6.94 0.16 13.91 20.54 0.83 0.57 100.54 1.931 0.161 0.028 0.212	15.46 0.29 9.45 18.97 0.73 0.08 98.38 1.962 0.095 0.793 0.504	1.33 7.58 0.28 14.96 19.05 1.91 0.09 99.83 1.878 0.158 0.047 0.235	23.61 0.27 1.14 3.53 12.01 n.d. 98.25 2.100 0.078 0.129
FeO Fe ₂ O ₃ MnO MgO CaO Na ₂ O Cr ₂ O ₃ Total Atoms pe Si Al Ti Fe ²⁺ Fe ³⁺	0.53 5.71 0.13 15.89 20.29 1.95 1.12 99.84 er 6 oxygens 1.840 0.670 0.025 0.169	1.95 7.81 0.16 14.65 19.94 1.04 n.d. 99.44 1.843 0.148 0.054 0.215	8.99 0.26 11.86 21.11 1.03 0.09 100.17 1.803 0.288 0.059 0.281	14.45 0.48 8.88 21.18 n.d. 0.07 100.00 1.989 0.092 0.015 0.459	7.27 0.18 14.40 20.29 0.90 0.37 98.54 1.885 0.152 0.032 0.227	1.40 10.16 0.32 12.39 20.28 n.d. 0.05 99.49 1.978 0.152 0.042 0.317	6.94 0.16 13.91 20.54 0.83 0.57 100.54 1.931 0.161 0.028 0.212	15.46 0.29 9.45 18.97 0.73 0.08 98.38 1.962 0.095 0.793 0.504	7.58 0.28 14.96 19.05 1.91 0.09 99.83 1.878 0.158 0.047 0.235	23.61 0.27 1.14 3.53 12.01 n.d. 98.25 2.100 0.078 0.129 0.953
$Fe_{2}O_{3}$ $Fe_{2}O_{3}$ MgO CaO CaO $Cr_{2}O_{3}$ $Total$ $Atoms pe$ Si Al Ti Fe^{2+} Fe^{3+} Mn	0.53 5.71 0.13 15.89 20.29 1.95 1.12 99.84 er 6 oxygens 1.840 0.670 0.025 0.169 0.004	1.95 7.81 0.16 14.65 19.94 1.04 n.d. 99.44 1.843 0.148 0.054 0.215 0.004	8.99 0.26 11.86 21.11 1.03 0.09 100.17 1.803 0.288 0.059 0.281 0.008	14.45 0.48 8.88 21.18 n.d. 0.07 100.00 1.989 0.092 0.015 0.459 0.015	7.27 0.18 14.40 20.29 0.90 0.37 98.54 1.885 0.152 0.032 0.227 0.005	1.40 10.16 0.32 12.39 20.28 n.d. 0.05 99.49 1.978 0.152 0.042 0.317 0.010	6.94 0.16 13.91 20.54 0.83 0.57 100.54 1.931 0.161 0.028 0.212 0.005	15.46 0.29 9.45 18.97 0.73 0.08 98.38 1.962 0.095 0.793 0.504 0.010	1.00 7.58 0.28 14.96 19.05 1.91 0.09 99.83 1.878 0.158 0.047 0.235 0.009	23.61 0.27 1.14 3.53 12.01 n.d. 98.25 2.100 0.078 0.129 0.953 0.009
$Fe_{2}O_{3}$ $Fe_{2}O_{3}$ MgO CaO CaO $Cr_{2}O_{3}$ $Total$ $Atoms pe$ Si Al Ti Fe^{2+} Fe^{3+} Mn Mg	0.53 5.71 0.13 15.89 20.29 1.95 1.12 99.84 er 6 oxygens 1.840 0.670 0.025 0.169 0.004 0.854	1.95 7.81 0.16 14.65 19.94 1.04 n.d. 99.44 1.843 0.148 0.054 0.215 0.004 0.798	8.99 0.26 11.86 21.11 1.03 0.09 100.17 1.803 0.288 0.059 0.281 0.008 0.661	14.45 0.48 8.88 21.18 n.d. 0.07 100.00 1.989 0.092 0.015 0.459 0.015 0.503	7.27 0.18 14.40 20.29 0.90 0.37 98.54 1.885 0.152 0.032 0.227 0.005 0.801	1.45 10.16 0.32 12.39 20.28 n.d. 0.05 99.49 1.978 0.152 0.042 0.317 0.010 0.690	6.94 0.16 13.91 20.54 0.83 0.57 100.54 1.931 0.161 0.028 0.212 0.005 0.758	15.46 0.29 9.45 18.97 0.73 0.08 98.38 1.962 0.095 0.793 0.504 0.010 0.549	1.88 7.58 0.28 14.96 19.05 1.91 0.09 99.83 1.878 0.158 0.047 0.235 0.009 0.827	23.61 0.27 1.14 3.53 12.01 n.d. 98.25 2.100 0.078 0.129 0.953 0.009 0.070
$Fe_{2}O_{3}$ $Fe_{2}O_{3}$ MgO CaO $Na_{2}O$ $Cr_{2}O_{3}$ $Total$ $Atoms pe$ Si Al Ti Fe^{2+} Fe^{3+} Mn Mg Ca	0.53 5.71 0.13 15.89 20.29 1.95 1.12 99.84 er 6 oxygens 1.840 0.670 0.025 0.169 0.004 0.854 0.800	1.95 7.81 0.16 14.65 19.94 1.04 n.d. 99.44 1.843 0.148 0.054 0.215 0.004 0.798 0.780	8.99 0.26 11.86 21.11 1.03 0.09 100.17 1.803 0.288 0.059 0.281 0.008 0.661 0.846	14.45 0.48 8.88 21.18 n.d. 0.07 100.00 1.989 0.092 0.015 0.459 0.015 0.503 0.862	7.27 0.18 14.40 20.29 0.90 0.37 98.54 1.885 0.152 0.032 0.227 0.005 0.801 0.811	1.49 10.16 0.32 12.39 20.28 n.d. 0.05 99.49 1.978 0.152 0.042 0.317 0.010 0.690 0.811	6.94 0.16 13.91 20.54 0.83 0.57 100.54 1.931 0.161 0.028 0.212 0.005 0.758 0.805	15.46 0.29 9.45 18.97 0.73 0.08 98.38 1.962 0.095 0.793 0.504 0.010 0.549 0.793	1.88 7.58 0.28 14.96 19.05 1.91 0.09 99.83 1.878 0.158 0.047 0.235 0.009 0.827 0.757	23.61 0.27 1.14 3.53 12.01 n.d. 98.25 2.100 0.078 0.129 0.953 0.009 0.070 0.155
$Fe_{2}O_{3}$ $Fe_{2}O_{3}$ MgO CaO $Na_{2}O$ $Cr_{2}O_{3}$ $Total$ $Atoms pe$ Si Al Ti Fe^{2+} Fe^{3+} Mn Mg Ca Na	0.53 5.71 0.13 15.89 20.29 1.95 1.12 99.84 er 6 oxygens 1.840 0.670 0.025 0.169 0.004 0.854 0.800 0.058	1.95 7.81 0.16 14.65 19.94 1.04 n.d. 99.44 1.843 0.148 0.054 0.215 0.004 0.798 0.780 0.074	8.99 0.26 11.86 21.11 1.03 0.09 100.17 1.803 0.288 0.059 0.281 0.008 0.661 0.846 0.074	14.45 0.48 8.88 21.18 n.d. 0.07 100.00 1.989 0.092 0.015 0.459 0.015 0.503 0.862 0.000	7.27 0.18 14.40 20.29 0.90 0.37 98.54 1.885 0.152 0.032 0.227 0.005 0.801 0.811 0.090	1.49 10.16 0.32 12.39 20.28 n.d. 0.05 99.49 1.978 0.152 0.042 0.317 0.010 0.690 0.811 0.000	6.94 0.16 13.91 20.54 0.83 0.57 100.54 1.931 0.161 0.028 0.212 0.005 0.758 0.805 0.059	15.46 0.29 9.45 18.97 0.73 0.08 98.38 1.962 0.095 0.793 0.504 0.010 0.549 0.793 0.080	7.58 0.28 14.96 19.05 1.91 0.09 99.83 1.878 0.158 0.047 0.235 0.009 0.827 0.757 0.137	23.61 0.27 1.14 3.53 12.01 n.d. 98.25 2.100 0.078 0.129 0.953 0.009 0.070 0.155 0.936

Table 1. Representative olivine and clinopyroxene compositions

Analyses of mineral chemistry were measured on a JEOL T300 SEM and Link system 860 probe. Abbreviations: OL, olivine; CPX, clinopyroxene; R, rim; C, core; n.d., not detected.

4 mm in length and are strongly zoned. Zeolites (usually thomsonite) occur as spectacular radiating fibrous aggregates (up to 1.5 mm in diameter) and form a maximum of 5% of this lithology. The pegmatite veins have a gradational lower contact and a sharp upper contact with the surrounding picrite. The picrite shows an upward increase in modal plagioclase and clinopyroxene towards the base of the pegmatite veins but no evidence of chilling.

5.b. Picrodolerites

Picrodolerites are distinguished from picrites by their lower content of modal olivine (10–20%). The greatest amount of olivine occurs adjacent to the picrite/ picrodolerite junction and decreases gradually into the crinanites. This decrease is compensated by an increase in clinopyroxene which becomes increasingly ophitic with plagioclase. Olivine in the picrodolerites is subhedral/anhedral and shows a greater Feenrichment (Fo₇₃-Fo₇₆) than has been observed in the



Figure 7. Variations in mineral composition with height and lithology in the northwest Trotternish sill. Symbols are as follows: \bigcirc olivine and \blacksquare clinopyroxene.

picrites. Glomerophyric aggregates of olivine and clinopyroxene (100(Mg/Mg+Fe) = 83.2-43.9), 3-4 mm in diameter, are common in the picrodolerites and typically consist of a large olivine crystal (2 mm) surrounded by smaller (0.5 mm) granular clinopyroxene crystals. A few larger clinopyroxene crystals (1.5 mm) subophitically enclose plagioclase. Plagio-clase laths (< 1.5 mm) exhibit strong normal zonation (An₆₅-An₁₈) and form 50-60% of this lithology. They have a subophitic relationship with ilmenite (< 1 mm).

5.c. Crinanites

The crinanites have an essential mineralogy of clinopyroxene (25-40%), plagioclase (55-65%) and analcime (< 3%) and, in comparison with the picrites and picrodolerites, are olivine-poor (< 5%). The olivine (Fo₈₁-Fo₃₈) is anhedral, up to 3.5 mm in diameter and sub-encloses plagioclase. Phenocrysts in



Figure 8. (a) Variations in clinopyroxene compositions in the Trotternish sills. Symbols are as follows \bigcirc picrite; \bigcirc marginal picrite; \triangle picrodolerite; \blacksquare crinanite; \blacktriangle pegmatite. (b) Comparison between the observed trend (shaded area) with other well-known intrusions (1) Black Jack sill (Wilkinson, 1957); (2) Main Shiant Isles sill (Gibb, 1973); (3) Main Shiant Isles sill (Murray, 1954); (4) Eilean Mhuire sill, Shiant Isles (A. R. O. Mohammed, unpub. Ph.D. thesis, Univ. Oxford, 1982); (5) Skaergaard Intrusion (Wager & Brown, 1967); (6) Shonkin Sag laccolith (Nash & Wilkinson, 1970).



Figure 9. Plagioclase laths tangentially arranged around olivine in a picrite sample.



Figure 10. Photomicrograph showing euhedral olivine phenocrysts (altered to iddingsite) in the chilled margin at the lower sill/sediment contact, Osmigarry.

the chilled margins are euhedral and comparable in composition with picritic olivines ($Fo_{81}-Fo_{79}$). They become more Fe-rich away from the chilled margins. Large clinopyroxene crystals (up to 5 mm in length) are strongly zoned ($Mg_{72.7}-Mg_{51.0}$) and ophitically enclose small plagioclase laths (< 1.5 mm). Unenclosed plagioclase laths (< 2.5 mm) sometimes occur as open spherulites. Large titaniferous magnetite crystals (up to 2 mm) are common (up to 5 modal %) and are intergrown with plagioclase. Analcime occurs interstitially in cavities with thomsonite.

A few pegmatite veins occur in the crinanites. A thick (~ 1 m) pegmatite vein is present two-thirds above the base of the crinanite sill (36 m thick) at Staffin [NG 494 680]. These pegmatites generally consist of plagioclase (65%), granular clinopyroxene (15%), anhedral olivine (15%) and opaque oxides (5%).

6. Interpretations of petrography

Poikilitic textures in the picrites resemble those of other well-known cumulate intrusions such as Skaergaard. Textures relating to adcumulus growth do not occur in the picrites and indicate that the rate of solidification exceeded the rate of convection (Sparks *et al.* 1985). The elongate crystals of plagioclase and olivine in the 'big feldspar picrite' may imply rapid crystallization and be analogous with undercooling along the synthetic forsterite-plagioclase cotectic. Donaldson (1976) crystallized olivines resembling those in the marginal picrites and suggested that they

are the result of either rapid cooling (50 °C/hour) and undercooling $(T \circ C = 40)$ or rapid growth due to a large olivine content in the melt. Previously, Drever & Johnston (1957) suggested that re-entrants in the shapes of these olivines were evidence of resorption. The proximity of such olivines to the margins of the sills suggests that the re-entrants are the result of rapid cooling. The significant Fe-enrichment at the rims of olivines in the marginal picrites may be interpreted in terms of changing thermal and chemical gradients surrounding the olivine phenocrysts. Rapid cooling close to the contacts would inhibit total re-equilibration and homogenization of the olivine. The presence of spinel, olivine and plagioclase phenocrysts in the chilled margins of the picrites implies that they were all near-liquidus phases in the magma at the time of emplacement.

Interlocking relationships between the pegmatite veins and surrounding picrite suggest that the formation of the veins is contemporaneous with crystallization of the picrite. The increasing amount of modal plagioclase and decreasing modal olivine within the host picrite towards the base of the veins suggests that the veins may have been derived by filter-press action. This mechanism involves the post-cumulus movement of interstitial liquid, as a result of compaction in the cumulus pile due to density differences between olivine and plagioclase. Interstitial liquid expelled upwards from the cumulus pile during compaction may have accumulated locally as veins. The relatively thin veins may have coalesced to form much thicker ones, such as the 75 cm thick pegmatite vein at Ru Bornaskitaig [NG 370 729]. This pegmatite occurs close to the upper contact of the picrite sill, possibly at the boundary of the upper and lower crystallization fronts. In this zone the expelled liquid may have ponded beneath a relatively impervious layer, but in an area favouring freedom of nucleation and crystal growth. The presence of plagioclase laths tangentially arranged around olivines in the picrites (Fig. 9) may reflect this compaction. Similar compaction textures have been observed in the Eastern Layered Series of Rhum (Young & Donaldson, 1985). The decrease in modal olivine towards the top of some picrite sills, such as the one at Skudiburgh [NG 374 674], may also be due to compaction. Swallow-tail terminations in clinopyroxenes imply undercooling, rapid crystallization and supersaturation, whilst biotite and zeolites infilling drusy cavities suggest an increase in water saturation during the later crystallization stages of the pegmatite veins. The absence of amphibole may reflect a deficiency of K_2O or low temperatures.

The glomerophyric clusters of clinopyroxene surrounding olivine that occur in the picrodolerites resemble reaction textures. Similar textures occur in Hawaiian tholeiites, where olivine is mantled by a rim of orthopyroxene, formed by reaction with surrounding silica-rich liquid (BVSP, 1981). In the multiple sills of northwest Trotternish the olivine crystals in the picrodolerites may be cognate xenocrysts from the underlying picrite which have been incorporated into, and reacted with, the more silica-rich magma of the picrodolerites to form clinopyroxene. Such 'xenocrysts' are present up to 5 m above the picrite/ picrodolerite contact and may suggest significant mixing and relatively rapid cooling in order to prevent the olivine from complete reaction with the surrounding 'silica-rich' magma. Clinopyroxene supersaturation and the cooling rate appear to have been lower in the picrodolerites than in the crinanites so that the growth/nucleation rate was relatively high and zoned ophitic clinopyroxene crystallized.

7. Geochemical variations in igneous stratigraphy

Stratigraphical variations in major and trace elements are shown in a composite profile from the sills in northwest Trotternish (Fig. 11). Representative wholerock analyses are presented in Table 2. All elements have relatively constant concentrations within the picrite unit, with marked variations occurring at the marginal picrite/picrite transition and at the picrite/ picrodolerite junction. Both changes reflect, primarily, a sharp decrease in modal olivine. Interaction at the sill margins with Jurassic sediments produces increases in CaO, MnO and Sr contents and the scatter of element concentrations. The gradual petrographic transition from picrodolerites to crinanites shows corresponding changes in geochemistry. The highest concentrations of MgO (30 wt.%) occur in the centre of the picrite unit, gradually decreasing outwards, towards the picrite/picrodolerite and marginal picrite/picrite transition, comparable to variations in modal olivine. The lowest concentrations of MgO occur in the crinanites (< 7.5 wt.%) which are structurally highest in the stratigraphy. FeO^T $(= FeO + Fe_2O_3)$ and MnO exhibit similar variations to MgO. This correlation between MgO and FeO^T relates to the modal distribution of chrome spinels and olivine. The cumulative concentration of FeO^T in both of these minerals is responsible for the high FeO^T (13%) in picrites. Other major elements show an antipathetic relationship to this trend.

The inverse relationship between MgO and CaO suggests that clinopyroxene has had little influence on MgO concentrations. Increases in TiO₂ (from 0.5% to 2.7%) with height in the sill reflect the increasing modal amounts of ilmenite in picrodolerites and titaniferous magnetite in crinanites. Trace element concentrations parallel the major element variations and highlight the contrasting trends. Concentrations of Cr and Ni vary uniformly with MgO. A two-fold jump in Sc (from 20 ppm to 42 ppm, at Creagan Iar; Fig. 2) at the picrite/picrodolerite junction reflects a similar two-fold increase in modal clinopyroxene.



Figure 11. Stratigraphic variations in major and trace elements in the northwest Trotternish sill. Symbols are as follows: • picrite; \triangle marginal picrite; \triangle picrodolerite; \blacksquare crinanite.

	C SG14	PD SG19	P SG20	P SG24	PEG SG31	P SG90	P SG114	MP SG123	MP SG125
Wt. %									
SiO,	48.39	48.81	45.58	43.65	45.19	42.1	40.83	45.76	45.55
TiO,	1.93	1.08	0.73	0.67	2.61	0.67	0.75	1.84	1.56
Al ₂ Õ ₃	15.54	12.26	9.55	8.41	13.07	6.9	6.51	13.3	13.54
FeOT	10.74	9.32	11.13	11.43	8.00	13.97	15.66	12.87	11.71
MnO	0.17	0.16	0.18	0.18	0.13	0.22	0.22	0.21	0.2
MgO	4.76	12.11	19.85	25.07	6.57	30.46	29.94	12.07	14.28
CaO	10.18	12.32	6.51	5.24	12.93	4.25	4.63	8.89	9.08
Na ₂ O	3.65	2.05	1.65	1.59	4.05	0.99	1.15	3.49	2.56
K,Ō	0.67	0.34	0.25	0.19	0.15	0.11	0.12	0.18	0.24
P ₂ O ₅	0.17	0.1	0.09	0.09	0.28	0.07	0.09	0.19	0.19
Total	96.2	98.55	95.52	96.52	92.98	99.74	99.89	98.79	98.91
ppm									
Ba	182	87	93	72	81	26	32	232	108
Cr	29	939	1263	2862	142	2491	2447	755	889
Cu	110	64	53	48	83	74	79	105	113
Nb	5.5	2.5	4.6	na	8.2	1.9	na	na	na
Ni	53	227	638	1052	107	1272	1268	465	617
Rb	20	13	9	13	na	13	11	12	12
Sc	30	42	20	17	27	15	16	31	30
Sr	402	275	202	178	639	108	121	504	527
Th	1.2	0.4	0.6	na	1.3	0.4	na	na	na
U	0.4	0.1	0.2	na	0.4	0.1	na	па	na
v	432	296	192	177	330	175	192	394	380
Y	20	12	7	9	26	7	8	20	19
Zn	111	80	106	92	59	102	104	108	116
Zr	134	68	58	47	196	42	48	112	101
La	12.4	7.4	8.9	7.3	12.6	6.3	6.9	8.1	8.3
Sm	4.4	0.3	na	na	4.3	0.8	na	na	na
Eu	1.9	0.1	na	na	1.9	0.5	na	na	na
Gd	5.4	3.6	na	na	5.7	1.4	na	na	na
Dy	4.4	2.9	па	na	4.1	0.8	na	na	na
Ho	0.8	0.6	na	na	0.7	0.2	na	na	na
Er	2.3	1.3	na	na	2.1	0.5	na	na	na
Yb	2.2	1.4	na	na	2.2	0.6	na	na	na
Lu	0.4	0.3	na	na	0.4	0.1	na	na	na
(⁸⁷ Sr/ ⁸⁶ Sr) ₆₀	0.704933±8	0.704968 <u>+</u> 9	0.704785 ± 10		0.704852±17	0.704922 ± 10	0.703680±8		

All major and trace elements except Nb were determined by ICP AES. Nb was analysed by ICP MS. Isotope ratios were determined on a VG 354 thermal ionisation mass spectrometer for which SRM gave 0.710241 ± 8 . Abbreviations: C, crinanite; PD, picrodolerite; P, picrite; PEG, pegmatite; MP, marginal picrite; na, not analysed.

87Sr/86Sr ratios have been obtained for 9 representative samples, from the Trotternish sills and a svenite from Eilean Mhuire, Shiant Isles. The results of these determinations are as follows: ⁸⁷Sr/⁸⁶Sr ratios range from 0.70360 to 0.70479 in picrites, 0.70492 to 0.70497 in picrodolerites and 0.70493 to 0.70531 in crinanites. They have a negative correlation with MgO (Fig. 11). The variation in Sr isotopes with petrography is consistent with major and trace element variations and suggests the picrites, picrodolerites and crinanites are genetically related. The syenite from Eilean Mhuire, Shiant Isles and a pegmatite vein from Bornaskitaig (SG31) have very low ⁸⁷Sr/⁸⁶Sr ratios (0.70432 and 0.70485 respectively) considering their highly evolved petrography. The similar low ⁸⁷Sr/⁸⁶Sr ratio of the Eilean Mhuire syenite and the pegmatite vein are compatible with a direct relationship with the picrites, probably by filter pressing (S. A. Gibson, unpub. Ph.D. thesis, Kingston Polytechnic, 1988). Previously, Walker (1930) suggested that the syenites formed by auto-intrusion and A. R. O. Mohammed (unpub. Ph.D. thesis, Univ. Oxford, 1982) suggested that the syenites may be the result of an upward migration of volatiles.

8. Parental magma

The whole-rock chemistry of chilled margins is often used to estimate parental magma compositions of plutons. Phenocrysts of olivine and plagioclase in the chilled margins of the Trotternish sills suggest that the magma had started fractionating prior to sill emplacement and these chilled margins may only approximate to liquid compositions. However, samples from picritic chilled margins (SG61, SG88, SG254) have whole-rock compositions which are in equilibrium with olivine Fo_{82} - Fo_{84} (calculated from Roeder & Emslie, 1970). Frey, Green & Roy (1978) suggested that primary basic magmas derived by partial melting of upper mantle peridotite should have Mg/(Mg+Fe) ratios between 68 and 75. All of the chilled margins have Mg values which are lower than this and are too evolved to be a partial melt of normal upper mantle peridotite. This may be explained by one or all of the following processes:

(i) The parental magma of the Trotternish sills may have undergone crystal fractionation at the base of the crust, prior to emplacement in the magma chamber. Fractionation of olivine and/or clinopyroxene would decrease the Mg/(Mg + Fe) ratio of the magma.

(ii) Highly forsteritic olivine may have fractionated and accumulated at the base of the magma chamber where it remains hidden.

(iii) The olivines (Fo_{83}) in the sills may represent sub-solidus re-equilibrated compositions.

Overall variations in whole-rock geochemistry suggest that crystal fractionation and accumulation were important processes during magma evolution. This fractionation and accumulation have been modelled using a least-squares computer program based on that of Bryan, Finger & Chayes (1969) and a parental magma with the same composition as SG254 (Table 3). Analysed minerals were used to estimate the compositions of the fractionating phases. Although the calculations assume that mineral compositions are constant, which may be petrologically unrealistic, the results (Table 3) show similarities with petrographic interpretations. The most MgO-rich picrites such as SG90 (30.46 wt.%) are calculated to have 62% olvine (Fo₈₃), 2.9% plagioclase and 1.2% chrome spinel. Clinopyroxene is an insignificant cumulus phase in the calculations.

The differentiation of a parental magma with the same composition as SG254 to MgO-poor crinanites such as SG14 (4.92 wt.%) has also been modelled (Table 3). Plagioclase (An₈₀) may have been an important fractionating phase (32%) in addition to olivine (Fo₈₃; 14.6%), clinopyroxene (9.4%) and titaniferous magnetite (6.4%). The picrodolerites may

Table 3. Least squares computer modelling of crystal fractionation and accumulation in the Trotternish sills

	M SG254 + -	lodel 1 OL+SP+PL → SG90	Mod SG254 -	el 2 → SG14	
Parent liquid or accumulate Mixture (wt. %) Crystals:	SG90		SG2	54	
Olivine Clinopyroxene	Fo ₈₃	62.0 % 	Fo ₈₃ C ₃₉ M ₅₀ F ₁₁	14.6% 9.4%	
Plagioclase Fe-oxide	An ₈₀ Sp	2.9 % 1.2 %	An ₈₀ Mgt	32.8 % 6.4 %	
Residual liquid or liquid fraction of cumulate	SG254	33.9%	SG14	36.8%	
ΣR^2		0.997		0.883	

Model 1 demonstrates the crystal accumulation of a picrite (SG90) from the proposed parental magma (SG254). Model 2 shows the degree of crystal fractionation required to form a crinanite (SG14) from SG254. ΣR^2 is the sum of squares for oxide percentage differences between observed and calculated compositions ($\Sigma R^2 < 1$ are considered to be significant). Abbreviations: OL, olivine; SP, Sp, spinel; PL, plagioclase; Fo, forsterite; An, anorthite; Mgt, magnetite.

represent the accumulation of these fractionated phases.

9. Emplacement models

9.a. Previous emplacement models

In the first major petrographic study of the LMSC, Walker (1930) accounted for the basal concentration of olivine in the main Shiant Isles sill, by in situ gravitational differentiation of crinanitic magma, due to olivine sinking and filter-pressing. However, Walker (1932) suggested that large-scale differentiation in inter-crustal reservoirs, prior to the intrusion of at least two separate pulses of magma, is necessary to explain the considerable volume of picrite in the Trotternish sills. Drever & Johnston (1959) suggested, with the discovery of the picrite/picrodolerite discontinuity in the Main Shiant Isles sill, that the picrite was emplaced as a separate intrusion to the rest of the sill. Simkin (unpub. Ph.D. thesis, Univ. Princeton, 1965; Simkin, 1967) examined variations in modal mineralogy within the sills in northwest Trotternish and concluded that they were emplaced by flow differentiation. Gibb (1973) proposed a mechanistic emplacement model for the Shiant Isles, expanding on the ideas of Drever (1953) and Drever & Johnston (1959, 1965, 1967). He envisaged the continuous emplacement of the picrite unit as a 'dense plug of olivine-rich sludge', after emplacement of the crinanite. Following the discovery of an upper picrite unit in the main Shiant Isles sill, Gibb & Henderson (1989) suggested that the main Shiant Isles sill was emplaced by multiple intrusion of picrite followed by crinanite.

9.b. Proposed emplacement model

The following petrographic and geochemical evidence suggests that the Trotternish sills were emplaced as a series of multiple intrusions with subsequent *in situ* differentiation:

(i) the presence of internal chills;

(ii) sills composed of single and multiple lithologies;

(iii) abrupt changes in petrographic textures and mineral compositions at the picrite/picrodolerite junction.

Huppert & Sparks (1989) have described how chilled margins in sills may be partially or totally remelted and how internal contacts may result from a single intrusive episode. They suggested that the amount of remelting should be highly asymmetric due to convection resulting in high heat flux at the roof. However, Gibb & Henderson (1989) dismissed a model involving a single flux of magma and remelting to explain the picrite/picrodolerite contact in the main Shiant Isles sill due to the distance from the lower and upper sill/sediment contacts.

Multiple intrusion may have resulted from withdrawal of magma from either interconnecting discrete magma chambers or from different levels in a single compositionally zoned magma chamber. Several factors suggest that the LMSC was emplaced by multiple intrusion from a single compositionally zoned magma chamber:

(i) The similar chemical compositions of olivine phenocrysts (Fo₈₂) in both the crinanite and picrite chilled margins indicate that a common source of olivine was available, prior to emplacement.

(ii) Continuous and systematic variations in mineral chemistry, whole-rock composition, Sr isotope ratios and petrographic textures with stratigraphic height (Gibson, 1990) suggest the tapping and subsequent emplacement from a single, compositionally zoned magma chamber.

Field relationships are not so simple, and a more complex emplacement mechanism is required for the LMSC than sequential withdrawal of less-fractionated magmas from progressively deeper levels in a zoned reservoir. At Invertote, east Trotternish, the picrite appears to have been emplaced before the overlying crinanite. The presence of strongly zoned olivine in the picrite directly below the contact suggests that the crinanite unit was emplaced close to the upper contact of the picrite unit. Nevertheless, the chill at the base of the crinanite unit suggests that the picrite had sufficient time to cool to a lower temperature than the crinanite magma. At Invertote, columnar jointing is continuous across the picrite/crinanite chill, which indicates that the sill eventually cooled as a single unit. At the picrite/picrodolerite junction in northwest Trotternish there is no evidence of chilling but the glomerophyric clusters of olivine and clinopyroxene in the picrodolerites resemble xenocrysts.

The emplacement of the Trotternish sills appears to have taken place discontinuously and may relate to intervals during withdrawal of magma from the chamber. Murata & Richter (1976) observed a correlation between discharge rate and olivine content in the 1959 Kilauean eruptions; at high discharge rates olivine-rich magmas (containing up to 50%) olivine phenocrysts) were erupted. The olivine distribution in the Trotternish sills suggests that the discharge rates were relatively high during the initial stages of magma withdrawal and olivine-rich magma was tapped from low levels in the chamber. A decrease in discharge rates and/or the outlet width (Blake & Ivey, 1986), would result in the withdrawal of crinanitic magma from the upper regions of the chamber. This olivine-poor magma was emplaced as homogenous sills, such as Staffin, or formed the upper parts of multiple intrusions, such as in northwest Trotternish. Sr isotope ratios show a continuous correlation with whole-rock geochemistry and provide no evidence of successive magma replenishment. The increase in modal olivine away from the contacts of picrite sills suggests that flowage differentiation may have been a significant process at the margins. The

effects of this process away from the contacts in thick intrusions (> 10 m) is limited by grain dispersive pressure (Barrière, 1976).

10. Conclusions

Variations in petrography, major and trace elements, and Sr isotopes with igneous stratigraphy suggest that the picrites, picrodolerites and crinanites in the Trotternish sills are genetically related and result from the multiple intrusion of magma from a compositionally zoned chamber in the upper crust. Picrites (18–30 wt.% MgO) may be interpreted as olivine cumulates, and crinanites (3–8 wt.% MgO) as olivinepoor fractionates of an alkali olivine basalt (10 wt.% MgO) parental magma. Pegmatite veins and the olivine distribution in picrites are evidence of postemplacement compaction. Rhythmic layering at sill contacts may be the result of variable growth/ nucleation ratios at the forsterite-diopside-anorthite eutectic in response to undercooling.

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