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THESIS
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IN THE
UNIVERSITY OF DURHAM

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AN ACCOUNT OF WORK DONE
IN THE SCIENCE DEPARTMENT OF THE
DURHAM COLLEGES
UNDER THE DIRECTION OF
PROFESSOR ARTHUR HOLMES
IN THE YEAR 1935 - 36

ENTITLED
"THE MINOR INTRUSIVES
OF KIRKCUDBRIGHTSHIRE"

BY
BASIL CHARLES KING, B. SC., F.G.S.

May 1934.
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2) Mineral Assemblages
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Plate II: Map showing Localities referred to in text between pages 2 and 3.

Plate III: Diagrammatic Section seen in Tongland cutting between pages 10 and 11.

Plate IV: Diagram illustrating Petrogenetic Sequence between pages 64 and 65.

Plates V to XII: Microscope Fields end papers

Plate XIII: Triangular Composition Diagram end paper.
Introduction:

The One Inch Geological Survey Map of the area (Sheet 5, Scotland) shows an extensive suite of minor intrusives which have been mapped as "porphyrites" and, more rarely as "lamprophyres". These rocks have been intruded as sheets, sills and dykes and are most abundant over an area about five miles square to the north-east of Kirkcudbright.

The country rocks are strongly folded slates and grits of Silurian and Ordovician age, with a prevailing N.E.-S.W. strike and high angle of dip. Over many square miles these have been invaded by the Criffel-Dalbeattie Granodiorite, which forms an irregular oval extending north-eastwards from the neighbourhood of Bengairn and Auchencairn Bay. A metamorphic zone averaging approximately one mile in width surrounds this intrusion.

Along a narrow strip of the Solway shores, Lower Carboniferous conglomerates, grits and sandstones, composed largely of igneous detritus, unconformably overlie the Silurian slates. In the extreme west a patch of Upper Old Red Sandstone underlies the Carboniferous. Nowhere do the intrusives cut the Old Red Sandstone or Carboniferous strata, nor are the latter metamorphosed by the granodiorite. The age of the igneous rocks is therefore known within definite limits.

Not infrequently, too, "porphyrite" dykes and sills are
Map of the area to the North-East of Kirkcudbright, showing the distribution of igneous rocks.
to be found penetrating the granodiorite, hence the relative ages of the plutonic and some of the hypabyssal intrusives can be demonstrated.

The only previous work on the minor intrusives has been carried out by the officers of the Geological Survey, who mapped and briefly described them in the Memoir (Explanation of Sheet 5, 1896). Here, J.J.H. Teall, who was responsible for the petrology, classified the hypabyssal rocks as "porphyrites", "diorites" and "mica-traps". He suggested "that the structural differences between the porphyrites and the hornblende-granites or quartz-diorites are due to differences in the conditions under which the magmas consolidated".

During the present study a general investigation was made of these intrusives. It was found possible in the field to recognise a number of distinct types, which were subsequently confirmed and further discriminated under the microscope. The relative paucity of good exposures inland and the high degree of alteration are severe obstacles in the way of a regional survey of the whole suite. It has been found convenient, therefore, to restrict detailed work to a few localities where good exposures occur of each of the types recognised. (See Plate II).

General Petrological and Field Relations:

The classification of these rocks has been based on purely petrological grounds. At the same time a notable
Map showing Localities referred to in Text.

Type I: Diorite-porphyry  Locality 5.
Type II: Albite-porphyrite  Localities 2 and 8.
Type III: Albitophyre  Locality 1.
Type IV: Albite-porphyry  Localities 3 and 4.
Type V: Plagiophyre  Locality 7.
Type VI: Xenocrystic type  Locality 6.
Type VII: Lamprophyre  Locality 9.
conformance obtains between these divisions and the nature of emplacement of the rocks concerned.

Type I: Quite the most unaltered type, this includes handsome, grey, porphyritic rocks with zoned plagioclase, frequently accompanied by biotite and hornblende. With increase of the latter minerals the rock assumes a lamprophyric appearance. They will be termed "diorite-porphyrite".

All the members of this type encountered are dykes with regular margins and cross-strike trends. They occur principally to the north-west and south-west of the granodiorite area of Bengairn and Forest Hill and are mainly confined within the metamorphic aureole.

Types II and III: These are by far the most abundant rocks of the area. Type II includes red or pink albite-bearing rocks (albite-porphyry) which sometimes internally lose the characteristic porphyritic texture and develop a finely granitic appearance in the hand-specimen. Type III includes reddish- or purplish-green non-porphyritic rocks, also albite-bearing, and possessing an orthophyric felted texture. Greenish, chloritic clots are a common feature of the hand-specimen. They will be termed albitophyre.

Both types are always more or less intensely hydrothermally altered; a process, which, it will be shown, is an essential feature of their petrogenetic evolution.

Between Tongland and Bengairn wide stretches are occupied by these rocks. It is impossible to determine the
exact form of the intrusions owing to the absence of continuous exposures. Their undoubted hypabyssal characters, limited effect on the country rocks and the occurrence within their boundaries of outliers and inliers of overlying and underlying slates, show, however, that their thickness is relatively small.

The discordant relationships of these intrusions to the Ordovician sediments renders the term sill or laccolith inapplicable, nor can the form be regular since it is impossible to recognise any systematic relation between limits of outcrop and contours. It seems most probable, therefore, that the intrusions in question comprise a series of irregular sheets.

Irregular dyke and sill intrusions are common and some of them are connected with the sheets. Their outcrop, modes and trends often vary considerably in different parts of their length, whilst their contacts with the country rocks are always highly irregular. To this type belong the excellently exposed dykes in Tongland road-cutting.

The precise relation between intrusions of type II and those of type III is not quite clear. In Tongland road-cutting the two form distinct dykes. In the area of sheet intrusions to the east the two may form neighbouring exposures, suggesting a closely related origin. Unfortunately, on account of the lack of critical exposures and the complexity in the modes of emplacement of the sheets, no conclusive field-evidence, bearing on the relationship of the two types has been observed.
Type IV: This type occurs as strike dykes, such as that at Balmae Haven, having well-defined vertical walls, and made up of pale cream or pink coloured, compact rocks, with pronounced porphyritic textures. The dyke margins, except for local coincidences, cut across the dip-planes. Albite is the dominant mineral, with rare dark constituents and rather abundant calcite. Quartz, although constituting no less than 26.6 per cent of the normative composition, is quite invisible in the microscope section. These rocks will be termed "leucocratic albite-porphyry".

Type V: Here are included the rocks of a number of intrusions from the Rockcliffe district, characterised by a felted, ortho-phryric texture and having zoned plagioclase felspars. Albite, quartz, chlorite and calcite, all of hydrothermal origin, are important constituents. The rocks form irregular dyke-like intrusions.

Type VI: Another distinct type, so far observed only forming irregular dykes in the Rockcliffe area, contains abundant large felspar aggregates and resorbed phenocrysts, suggesting derivation from xenoliths.

Type VII: A number of lamprophyres have been examined. All occur as narrow, irregular dykes and are typical kersantites or minettes, olive-green in colour and containing abundant flakes of biotite. The felspar can but rarely be identified.

The maps (Figs. I and II) indicate the approximate distribution of the various types and the localities referred to in the detailed descriptions.
### Petrographic Types:

<table>
<thead>
<tr>
<th>Type</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>II:</td>
<td>Albite-porphyry</td>
</tr>
<tr>
<td>III:</td>
<td>Albitophyre</td>
</tr>
<tr>
<td>V:</td>
<td>Plagiophyre</td>
</tr>
<tr>
<td>VI:</td>
<td>Xenocrystic Dykes</td>
</tr>
<tr>
<td>IV:</td>
<td>Leucocratic Albite-porphyry</td>
</tr>
<tr>
<td>I:</td>
<td>Diorite-porphyrite</td>
</tr>
<tr>
<td>VII:</td>
<td>Lamprophyre</td>
</tr>
</tbody>
</table>

### Mode of Occurrence, Trend, Etc.:

<table>
<thead>
<tr>
<th>Type</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Sheets, Irregular Dykes and Sills.</td>
</tr>
<tr>
<td></td>
<td>Regular Strike Dykes.</td>
</tr>
<tr>
<td></td>
<td>Regular Cross-Strike Dykes.</td>
</tr>
<tr>
<td></td>
<td>Small Irregular Dykes.</td>
</tr>
</tbody>
</table>

---

**Type I: Diorite-Porphyrite.**

This, the simplest, least altered type, is conveniently considered first.

A number of dykes of this type occur in the metamorphic aureole of the granodiorite, of which typical examples may be seen on the slopes of Screel Hill, behind Whitehill Farm. Similar dykes are also found south-west of Bengairn and Forest Hill.

Characteristically the dykes cut across the strike of the country rocks at right angles, and lie more or less parallel to the granodiorite margins. The contacts are commonly sharp and straight, while the width of the dykes is rarely above five or six feet.

The dyke-rocks are usually of striking appearance, with large white felspars and dark flakes and prisms of mafic...
minerals in a grey matrix of medium grain. They are characteristically unaltered.

**Petrography:**

The texture is porphyritic, with large tablets and glomero-porphyritic aggregates of plagioclase, together with biotite and hornblende, in a fine-grained groundmass. (Pl. V, 1).

The plagioclase is often quite clear, but may show a patchy clouding due to a "kaolin" type of mineral, while flakes of a "sericitic" mica are common. Occasionally more heavily clouded feldspars are traversed by irregular channels occupied by a clear albitic felspar of low refractive index.

Beautifully zoned crystals are frequently seen; often oscillatory in composition.

**Extinction Angles:** 12° - 16°.

**Optic Sign:** Negative.

**Refractive Indices:**

<table>
<thead>
<tr>
<th></th>
<th>Calcic Zones</th>
<th>Sodic Zones</th>
</tr>
</thead>
<tbody>
<tr>
<td>α</td>
<td>1.546</td>
<td>1.543</td>
</tr>
<tr>
<td>β</td>
<td>1.551</td>
<td>1.547</td>
</tr>
<tr>
<td>γ</td>
<td>1.554</td>
<td>1.551</td>
</tr>
</tbody>
</table>

These characters correspond to oligoclase-andesine ranging in composition from AbAn to AbAn.

The hornblende occasionally possesses euhedral diamond-shaped or rectangular outlines; more commonly, however, the crystal edges are minutely irregular and the centres are riddled with acicular apatites. It is of a brownish-green
colour, and has optical characters normal to common hornblende:

Extinction Angles: \((Z, c)\) 21° - 23°.

Pleochroism: \(X < Y < Z\).

Optic Sign: Negative.

The degree of alteration varies somewhat from dyke to dyke. Clear green flaky chlorite (penninite), granules of epidote and patches of calcite are the secondary minerals.

**Biotite** is frequent as ragged, fibrous crystals. The colour is a deep fawn-brown with strong pleochroism, but bleaching and chloritisation have usually effected considerable alteration marginally and along the cleavage planes. The Chlorite is less flaky than that associated with hornblende and shows abnormal indigo-blue or rusty-brown interference tints. Basal sections with hexagonal outlines occur: percussion figures are sometimes seen.

The **matrix** is fine-grained and generally light-coloured, with a suggestion of flow-structure. Pre-\(\Lambda\) felspar dominates but microlites of biotite and hornblende are also abundant.

**Accessories** are apatite and rare zircon, as well as pyrite and black ores.

As a type these rocks correspond in mineral composition to diorite, of which they appear to represent the hypabyssal equivalent. On these grounds, therefore, the name diorite-porphyrite is fully warranted.
### MINERAL SUMMARY: TYPE I.

#### A. PHENOCRYSTS.

<table>
<thead>
<tr>
<th>Primary Minerals:</th>
<th>Secondary Minerals:</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Plagioclase:</strong> <em>(Ab$_n$An$_m$)</em></td>
<td><strong>Cladding of:</strong> &quot;Kaolin&quot; mineral.</td>
</tr>
<tr>
<td>α 1.546 1.545 1.543</td>
<td></td>
</tr>
<tr>
<td>β 1.551 1.549 1.547</td>
<td></td>
</tr>
<tr>
<td>γ 1.554 1.553 1.551</td>
<td></td>
</tr>
<tr>
<td>Optic Sign: Negative.</td>
<td></td>
</tr>
<tr>
<td><strong>Hornblende:</strong></td>
<td><strong>Channels of:</strong> clear Albite.</td>
</tr>
<tr>
<td></td>
<td>Flakes of Penninite,</td>
</tr>
<tr>
<td></td>
<td>Granules of Epidote.</td>
</tr>
<tr>
<td><strong>Biotite:</strong></td>
<td>Alteration: Bright green Penninite (with indigo-blue and rusty-brown int. tints) Epidote granules. Replacement by Calcite along cleavages.</td>
</tr>
<tr>
<td><strong>Accessories:</strong></td>
<td></td>
</tr>
<tr>
<td>Needles and tiny hexagons of Apatite.</td>
<td></td>
</tr>
<tr>
<td>Pyrite and Black Ores.</td>
<td></td>
</tr>
<tr>
<td>Rare Zircon.</td>
<td></td>
</tr>
</tbody>
</table>
### Primary Minerals:
- Matrix fine-grained, light-coloured and cryptocrystalline with microlites showing tendency towards flow-structure.
- Plagioclase: tiny squares and rectangles, often with indefinite outlines.
- Hornblende and Biotite: tiny dark microlites and shapeless grains.

### Secondary Minerals:
- Alteration slight: "Kaolin" clouding.
- Chloritisation general.

---

**TYPE II: ALBITE-PORPHYRY.**

The northern dyke at Tongland road-cutting has been examined in greatest detail and is, therefore, taken as the type example. Similar dykes or sills occur near Billies Farm and near Little Sypland, a few miles north-east and east, respectively, of Tongland.

The southern margin of the Tongland dyke makes a sharp contact with the country rocks. These are here greenish-grey, highly cleaved slates, dipping steeply away from the dyke. The latter cuts obliquely across the cleavage, which appears in no way to affect the trend of the igneous contact. Near the dyke margin the slates contain "splashes" or patches of porphyry of varying shape and size, up to an inch or more in diameter. In none of the cases examined could these
Diagrammatic Section of Dykes seen in Tongland Road-cutting.

(Not drawn to scale).
PLATE III.

SOUTH DYKE

NORTH DYKE

Not accurately to scale.
Total length of section approximately 180 feet.

Figure refers to specimen numbers.

Width of South Dyke about 30 feet.
be traced to the main dyke, but they may have been originally connected via a mass now removed.

Inwards from the contact, the igneous rock, at first markedly porphyritic, with phenocrysts up to about 3 - 4 mms. in length, gradually becomes more even-grained owing to increasing coarseness in the texture of the matrix. Even in the field this change is clearly evident, since from being porphyritic the rock develops a fine-grained granitic texture.

Further north in the cutting, the porphyry becomes bleached, crumbly and very susceptible to weathering by surface waters. When, further on, the fresh rock reappears, it is again of the porphyritic variety and, although the northern contact cannot be seen owing to the termination of the cutting, the approximate position of the junction can be located. Evidently the porphyritic variety is the marginal facies to the granitic variety of the interior. (See Plate III).

**Petrography:** (See Plate V, Fig. 2).

The phenocrysts are:

(a) felspar, easily dominant;

(b) hornblende, often partially or completely pseudomorphed by chlorite; and

(c) a pale green chlorite, thought to represent original biotite.

The hornblende, in particular, frequently shows nearly euhedral outlines.

(a). Occasionally the felspar phenocrysts are euhedral;
more commonly, however, they are aggregates of several crystals, irregularly disposed and intergrown. The crystal edges often show re-entrants and acute angles, but either resorption (if they are xenoliths), or incomplete growth (if they are segregations) is suggested where the boundaries are rounded or indistinct. The fact that both phenocrysts and aggregates occur right up to the contact indicates that they were in existence before the emplacement of the magma, an inference which is supported by the frequent occurrence at the dyke margin of cracked crystals, due, presumably, to the stresses set up during intrusion. The occurrence of cracked crystals also suggests a probable explanation for the angular form of certain phenocrysts, which thus appear to be fragments of formerly larger felspars.

Further evidence regarding the mode of development of the felspar phenocrysts is provided by a dyke of the same general type occurring near a small quarry about one mile N.N.W. of Argreennan Mains (two miles north of Tongland). Here all the felspars, although often nearly euhedral in form, are riddled with patches of matrix. Commonly a narrow marginal zone is free from inclusions, which presumably precludes the possibility of corrosion of pre-existing crystals by the matrix magma. (Cf. MacGregor, 1931). The inference is, therefore, that portions of the matrix were enclosed while the felspars were crystallising. Frequently such a "sieve" texture is encountered in metamorphic rocks, where a mineral has developed in a solid medium, e.g. "sieved" pyroxenes, whilst chiastolite presents a similar,
though not quite comparable, phenomenon. (Plate IX, Fig. 2).

The riddled phenocrysts also raise the question of whether the "shadowy-looking" felspars of type II as a whole have ever been more completely developed than they now are. A comparison with the "incipient" felspars of the matrix, (q.v. pp. 17 - 18) which have only begun to crystallise, suggests that the phenocrysts themselves may be similarly imperfectly formed. Such an assumption would materially help in reconciling the two apparently incompatible ideas:

(i) that some of the felspars were already crystallised before the emplacement of the magma, and

(ii) that in other cases the felspars have developed during the consolidation of the matrix.

It is only a question of the circumstances controlling the clearing strength of growing crystals, as to which size alone is demonstrably no gauge.

The felspar phenocrysts are usually more or less clouded. Most sections show lamellar twinning, but zoning is never seen.


Optic Angle: Large - approaching 90°.

Optic Sign: Positive.

Refractive Indices:

\[
\begin{align*}
\alpha & = 1.528 \\
\gamma & = 1.537
\end{align*}
\]

These characters correspond to those of albite.

Apart from the presence of patches and channels of clear
secondary albite, clouding is ubiquitous; it resolves under high power to diffuse, brownish specks, suggesting dispersion of a kaolin-like substance.

Tiny flakes of a white "sericitic" mica, recognised by its bright interference tints, are usually present. Generally they show a parallelism to the traces of the dominant cleavage direction or twinning planes in the section.

Little clear, pale green flakes and fibres of chlorite are occur similarly, though less abundantly than the mica.

Occasionally, too, quartz occurs as flakes, nests and fine-veinlets and calcite as greyish patches, resolved into numerous tiny crystals under high-power. These minerals have effected replacement of a somewhat different nature from that of the other products of alteration of the felspar. Both quartz and calcite occur in the matrix and phenocrysts indiscriminately.

Some difficulty is encountered in attempting to establish the sequence of formation of these minerals. Clouding and sericitisation appear to be the earliest, since they are almost invariable and are absent where channelled by clear albite. Similarly chlorite, quartz and calcite would appear to represent the probable order among the later minerals.

(b). The hornblende in some specimens (e.g. 95) is entirely or largely unaltered. Commonly it occurs as medium-sized, approximately euhedral phenocrysts, up to a millimetre or so in length and having characteristically rectangular or diamond-
shaped outlines. The cleavages, where developed at all, are generally fine.

The colour is a brownish olive-green, most sections showing a moderately strong pleochroism.

Simple twins are not uncommon. The twinning plane is (100), since it is parallel to the length in longitudinal sections which show moderately low birefringence, and bisects the acute cleavage-angle in basal plates.

**Extinction Angles:** $(Z_c) \quad 22^\circ - 23^\circ$.

**Pleochroism:**

<table>
<thead>
<tr>
<th></th>
<th>X</th>
<th>Y</th>
<th>Z</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pale</td>
<td>Yellowish-</td>
<td>Brownish</td>
<td>Brownish</td>
</tr>
<tr>
<td>green.</td>
<td>green.</td>
<td>green.</td>
<td>green.</td>
</tr>
</tbody>
</table>

$X < Y < Z$.

**Birefringence:** 0.023.

**Optic Angle:** Moderate.

**Optic Sign:** Negative.

A suggestion of marginal zoning to a darker variety has been noted. These properties indicate a hornblende of about normal composition; the brown colour may be due to a relatively high soda content.

Although unaltered hornblende is found, obvious pseudo-morphs in chlorite and calcite are more common. All gradations exist between the two extremes, and intermediate forms, with residual patches of optically continuous hornblende, are abundant. (See Plate VI, Fig. 2, D and E).

Alteration to chlorite proceeded first; it accurately
conforms to the original outlines and encloses residual hornblende. Chloritisation commenced marginally and tended to follow the cleavages.

Granules and aggregates of epidote are commonly associated with the chlorite. Occasionally, too, black ores are much in evidence, occurring as euhedral or rounded grains, marginally distributed.

Patchy calcite, with "lobate" margins, replaces both hornblende and chlorite. Often it follows the original outlines as closely as the chlorite, elsewhere its distribution is less regular. Epidote tends, however, to remain enclosed and unreplaced. Some examples of otherwise unaltered hornblende veined with calcite occur.

Rarely albite can be seen irregularly replacing hornblende.

(c). Unaltered biotite never occurs, but chlorite suspected of pseudomorphing biotite is common. The evidence that the original mineral was not hornblende, but probably biotite, is as follows:

(i) Chlorite is sometimes found in rectangular sections with the characteristic ragged ends and strong basal cleavage of biotite. Basal flakes with no cleavage and nearly hexagonal outlines may be seen.

(ii) The general outline and cleavages of these pseudomorphs are quite unlike those of chloritised hornblende. Traces of cleavage after hornblende, if seen at all, are
always fine, straight and infrequent. In these pseudomorphs, however, the cleavages may be waved or bent; a feature which is common in biotite.

(iii) Complete "biotite" pseudomorphs are found in one specimen (95) together with nearly unaltered hornblende. Their derivations from hornblende is therefore extremely unlikely.

(iv) The epidote (see pp. 22 - 23) is partly of a fibrous type, following the cleavages, not found in pseudomorphs after hornblende.

The matrix, is conveniently considered under three main headings, according to the relative position in the dyke, which each are in turn divided into two sub-headings, describing primary and secondary constituents.

(A). Marginal Portion of Dyke (e.g. specimens 95 and 96).

Here the matrix is fine-grained and definite crystal outlines cannot be distinguished.

(a) Primary Constituents:

Between crossed-nicols a mottled effect is seen which is due to the incipient development of "shadowy" felspars. These have been "frozen" before assuming more distinct outlines or clearing themselves of minute inclusions. Red and black specks of ore and laths of chlorite pseudomorphs are abundant. The general reddish colour is due, presumably, to a dispersion of hydrated iron oxides.

(b) Secondary Constituents:

Flakes, streaks and tiny fibres of chlorite have been
extensively introduced. Their distribution is highly irregular, but tends to be concentrated around phenocrysts of felspar and pseudomorphs after hornblende and biotite. Elsewhere, too, in the matrix occur chlorite aggregates and patches.

Quartz is not uncommon, being usually closely associated with felted fibres of chlorite and forming colourless patches with shadowy extinction. Occasionally, too, grains of clear secondary albite, showing twin-lamellae occur with, and are eat-by partly replaced by quartz. The latter appears to replace chlorite also, since it cuts across the fibres.

Tiny rounded calcite patches are distributed fairly uniformly throughout the matrix, together with occasional irregular replacements of greater extent. Calcite appears to be always the latest replacing mineral, penetrating matrix, chlorite and quartz alike.

The degree of alteration of the matrix bears no apparent relation to the proportion of secondary minerals in the matrix. Thus, although the matrix is identical in specimens 95 and 96, yet in the former the hornblende is fresh, while in the latter it is entirely chloritised.

(B). Intermediate Varieties (e.g. specimens 99 and 100).

(a) Primary Constituents:

Here the matrix is less obscurely developed. The felspars have assumed more or less rectangular outlines with definite crystal edges. Even so, under the microscope, the
rock still retains a porphyritic texture.
(b) Secondary Constituents:

A more advanced type of hydrothermal alteration can here be recognised. The most notable feature is the occurrence of "nests" of quartz and albite, sometimes with other minerals (e.g. epidote, chlorite and calcite) in addition. The albite is frequently untwinned. Where associated with chlorite, the latter is the replacing mineral, penetrating the albite in fibrous growths and encroaching on its crystal margins. It is clear, as evidenced by the fact that the chlorite fibres are always cut by quartz, but never cut it, that quartz is a still later mineral. Replacement phenomena between quartz and albite are well exhibited:

(i) Smooth rounded encroachments of quartz into albite crystals;
(ii) Fingering penetrations of quartz into albite;
(iii) Partial replacements of quartz after albite, in which only a fine network of optically continuous albite remains;
(iv) Fine intergrowths of quartz and albite. These frequently occur beyond the edges of unaffected albite laths or tablets. They may also form a peripheral rim to a felspar which happens to be in the vicinity of a quartz-albite nest.

Albite, chlorite and quartz are also dispersed throughout the matrix, always showing evident replacement of earlier
minerals. Quartz, although tending to occur in the interstice of the matrix felspars, commonly makes lobate penetrations into their margins.

A clear, bright yellow, pleochroic epidote of aggregate-columnar or radiating-fibrous habit, often occupies the middle of quartz-albite nests. (Plate VI, Fig. 2, A & C).

Calcite occasionally replaces the matrix, primary and secondary minerals alike, forming clear optically-continuous patches with irregular, rounded or angular edges.

(C). **Innermost Varieties** (e.g. specimens 101 and 102).

(a) Primary Constituents:

These rocks correspond with the "granitic" variety of the field description (see p.11). The matrix is even coarser than in 99 and 100 and individual felspars have reached the proportions of small phenocrysts. The even-grained appearance of the groundmass is lost, but the distinctive feature of the rock lies in its secondary characters.

(b) Secondary Constituents:

Intergrowth-textures are here dominant. The larger felspars are often surrounded by "rosettes" of albite and quartz. The secondary albite appears to have grown in continuity with the original felspars and to have been partially replaced by quartz. Where the intergrowths are very fine discrimination is often possible only if the component minerals can be traced to discrete crystals. The various
fine replacement textures are:

(i) Striation type, in which the two minerals are evenly streaked together. The striations are straight and parallel.

(ii) Vermicular type. The component minerals can here be readily distinguished and traced to larger areas. This closely resembles myrmekitic textures.

(iii) "Herring-bone" type. This usually forms the rosettes surrounding felspar crystals. Each "spoke" of a rosette has the appearance of a herring-bone of which quartz forms the backbone and ribs and albite the intervening spaces.

(iv) Mesh type, in which the albite remains as a fine lattice-like framework.

Frequently these intergrowths become perceptibly resolved only under high-power. (Plate VII, Figs. 1 and 2).

Chlorite may assume a spherulitic habit in quartz-albite nests; with it are occasionally associated radiating fibres of what is probably a pale actinolite (see pp. 23-24)

Chlorite of several varieties is abundant in all the dyke-rocks. The mineral occurs as:

(a) Pseudomorphs replacing hornblende and "biotite".

(b) Fibrous aggregates and scattered flakes in the matrix and through the felspars.

(c) Constituents of albite-quartz nests, often with spherulitic radiating habit.
Penninite (negative): This, the most abundant variety, pseudomorphs hornblende and occurs through the matrix and felspars. It is pale green in colour, slightly pleochroic and fibrous or flaky in habit. Pseudomorphs consist of aggregates of separate flakes. It has a negative optical sign and shows abnormal prussian-blue interference tints.

Penninite (positive): Another pale green chlorite occurs in association with a brownish epidote in the "biotite" pseudomorphs. The interference tints are deep purplish-indigo, sometimes changing to a rusty-brown on rotation. A further distinction is its positive optical sign.

Colourless Chlorite: Occasionally a colourless radiating fibrous mineral is found associated with positive penninite in biotite pseudomorphs. It tends to form sheafs of acicular crystals along the cleavage planes of the pseudomorphs, appearing to push aside the epidote granules and penninite. The crystals are commonly length-slow; otherwise optical properties are not determinable.

Epidote is a very widely distributed mineral. Optical properties are usually difficult or impossible to determine, but it is convenient to recognise the following varieties, according to differences in form and mode of occurrence:

(a) Epidote I: Greenish or brownish-yellowish granules occurring in the hornblende pseudomorphs. The grains have high relief and characteristic birefringence (about 0.030), but their crystal orientation is usually difficult to
determine. Elongated crystals appear to show typical straight extinction and may be either length-slow or fast. The sign is negative and the greatest absorption is always with the length.

(b) Epidote II: A second variety, brownish, cloudy and often fibrous, is almost exclusively associated with positive penninite in biotite pseudomorphs. The fibres are more or less parallel to the cleavage of the chlorite, but may occur in lenticular aggregates of sheaf-like habit. The extinction is usually but not invariably straight. Under high-power the real colour is seen to be pale yellow, the brown colour being due to the cloudiness, which may obscure the interference tints. The optical properties are the same as those of the first variety, which also occurs in the biotite pseudomorphs.

(c) Epidote III: A bright yellow, aggregate-columnar or radiating-fibrous variety is found in the middle of albite-quartz nests. Here the epidote is very clear and has strong pleochroism. The sign is negative, the acute bisectrix at right angles to the length and the elongation usually nearly with the optic normal. In the aggregates, transverse sections showing acute and obtuse angles are common.

A radiating, fibrous mineral is occasionally associated with chlorite in hornblende pseudomorphs or in the chlorite of quartz-albite nests. It has the following properties, suggesting actinolite:

(a) Fibrous, pale green in colour, faintly pleochroic and
with brownish clouding arranged in concentric bands in the sheafs;

(b) Moderately high relief, greater than that shown by quartz, felspar and chlorite;

(c) Oblique extinction, up to about 20°;

(d) Slow with length;

(e) Negative sign.

Since its needles penetrate the associated chlorite, its position in the paragenetic sequence appears to be later than that mineral. Relations with quartz have not been observed.

Distinction is often difficult between quartz and albite in the nests and matrix replacements, since:

(a) The albite is frequently clear and free from visible signs of twinning;

(b) The albite-quartz replacement textures may be so fine that discrimination is impossible.

Distinction, where possible, has been based on the following features:

(a) Lamalār twinning, where found, is indicative of albite;

(b) Where albite and quartz occur together without overlapping the relative refractive indices may be determined;

(c) If the orientation is suitable a clear uniaxial or biaxial figure can sometimes be found. An albite section, in which the twinning is not seen, is often nearly perpendicular to the acute bisectrix;

(d) Where the intergrowths textures are very fine the component
minerals may be sometimes traced to larger crystals, the identification of which are possible.

Evidence of Veins.

A few thin veins occur in the central portions of the dyke. The vein material varies from place to place, containing sometimes quartz, elsewhere albite, and occasionally chlorite or calcite. Quartz and albite, both of which are colourless, are most readily distinguished when passing through albite phenocrysts. The clear albite of the vein is sharply contrasted to the cloudy albite of the phenocrysts, but both have the same refractive index. Quartz, however, has a relief perceptibly higher than either.

Accessories:

Accessories are apatite and zircon. Apatite is a relatively abundant mineral, occurring as euhedral inclusions in the mafic or felspathic constituents or scattered through the matrix. The crystals are not infrequently acicular in form, suggesting a direct magmatic source. This variety contrasts with the squat, rounded crystals from the dyke margin and the slates, the forms and restricted occurrence of which suggest a sedimentary origin (see below, p. 26).

At least one or two isolated crystals of zircon have been recorded from every dyke section. In a section from the margin no fewer than five were found within a few square millimetres. The outlines are always much rounded and the
similarity of their form to those found in the slates suggests a sedimentary origin.

Relations with Slates.

At its southern end the dyke can be seen making a sharp, though irregular contact with the slates. Under the microscope igneous material is observed to have penetrated the sediments, irregular offshoots containing phenocrysts of hornblende, felspar and pseudomorphous chlorite in a chilled matrix.

The slates consist of irregular quartz grains of varying size, together with much finely disseminated green chloritic material and black ore. Occasional albite porphyroblasts occur, and, here and there, nearly colourless flakes of mica, showing high interference tints.

Rounded crystals of zircon and apatite were observed in the slates, apatite being by far the more abundant.

In the margins of the igneous rock, too, apatites occur, often rounded, larger and in greater abundance than in the rest of the dyke. Rounded zircons were also found in the marginal rock and, as already mentioned (p. 25), in most of the slides of the dyke similar rounded zircons can be found. The detrital appearance of the apatite and zircon and their general similarity to those in the slates suggest their derivation from the latter.
### MINERAL SUMMARY: TYPE II.

#### A. PHENOCRYSTS.

<table>
<thead>
<tr>
<th>Primary Minerals:</th>
<th>Secondary Minerals:</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Albite:</strong></td>
<td></td>
</tr>
<tr>
<td>Form: euhedral tablets; &quot;fragmentary&quot; crystals or aggregates; re-entrants common; angles may be rounded; sometimes matrix inclusions. Lamellar Twinning. No Zoning. Extinction Angles: 17° - 18°. Refractive Indices: ( \alpha = 1.528 ) ( \gamma = 1.537 ) Optic Sign: Positive.</td>
<td>Alteration and replacement: &quot;Kaolin&quot; clouding. Extensive saussuritisation: &quot;Sericite&quot;, rarer Chlorite, clear secondary Albite - as flakes; Quartz and Calcite - irregular patches.</td>
</tr>
<tr>
<td><strong>Hornblende:</strong></td>
<td></td>
</tr>
<tr>
<td>Form: small euhedral crystals. Colour: brownish-green. Simple Twinning. Extinction Angle: ( Z_a - c 23° ). Birefringence: 0.023. Optic Sign: Negative. Optic Angle: Moderate. Pleochroism: ( X &lt; Y &lt; Z ) (mod. to strong).</td>
<td>Often entirely pseudomorphed: Negative Penninite and Epidote granules; Black Ores; Rare Actinolite; Calcite; Rare Albite.</td>
</tr>
<tr>
<td><strong>Biotite:</strong></td>
<td></td>
</tr>
<tr>
<td>Entirely pseudomorphed. Form: rectangles with ragged terminations; hexagonal flakes.</td>
<td>Positive Penninite and much fibrous Epidote.</td>
</tr>
<tr>
<td><strong>Accessories:</strong></td>
<td></td>
</tr>
<tr>
<td>Rounded Apatite near dyke margins; Acicular Apatite internally; Rounded Zircons - especially near margins.</td>
<td></td>
</tr>
</tbody>
</table>
B. MATRIX:

<table>
<thead>
<tr>
<th>Primary Minerals:</th>
<th>Secondary Minerals:</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Albite:</strong></td>
<td><strong>Chlorite:</strong></td>
</tr>
<tr>
<td>In finest types; light-coloured patches - incipient crystals.</td>
<td>Flakes dispersed through groundmass - concentrations round pseudomorphs.</td>
</tr>
<tr>
<td><strong>Hornblende:</strong></td>
<td></td>
</tr>
<tr>
<td>Needles and laths; more or less chloritised.</td>
<td>Nests of Chlorite, clear Albite Quartz and Calcite, together with sheafs of clear yellow Epidote.</td>
</tr>
<tr>
<td><strong>Ores:</strong></td>
<td></td>
</tr>
<tr>
<td>Dispersion of hydrated ores gives general red coloration Black specks and granules.</td>
<td>Quartz-Albite intergrowths; intricate and finely textured; often margining felspar phenocrysts; most important where non-porphyritic.</td>
</tr>
</tbody>
</table>

**TYPE III: ALBITOPHRE (ALBITISED PORPHYRITE).**

This is the most widespread of all the "porphyrite" types. Owing to its wide variations in texture and degree of alteration it cannot be briefly defined. Most of the dykes between Kirkcudbright and Castle Douglas and the extensive sheet-like expanses round Culdoach Hill and from Bentudor to Black Stockarton Moor are of this type. At intervals intrusions of the more massive red albite-porphyry are associated with those of Type III, but good contacts between the two types have not been found.

In the field it is evident that the rocks of Type III are always more or less intensely altered. They are grey,
purplish-brown, dull-green or reddish-brown in colour and, though not markedly porphyritic, they all contain numerous green chloritic clots up to several millimetres in diameter.

Most of the varieties are excellently exposed in the southern dyke of the Tongland road-cutting (see Pl.III), which is taken for convenience as the type example. This is the smaller of the two dykes exposed in the cutting and has a width of approximately thirty yards.

The slates may be seen to the south of the dyke and again between the two dykes. Close to the contact they are somewhat baked but otherwise they are but little affected by the intrusion. The cleavage dips at about 65° and strikes in the characteristic N.E. - S.W. direction.

The southern contact of the dyke is sharp, with irregular injections into the slates. The contact is interrupted by a V-shaped shatter-system of jumbled slates and dyke-rock and is charged with calcite veins.

The "porphyrite" near the contact has a dark greenish-grey or purplish colour, with a flaky surface and well-jointed structure.

Beyond the shattered zone is a jointed rock of a reddish colour with chlorite clots. This is succeeded by a more massive "porphyrite" of similar colour, having strong rectangular joints, the planes of which are coated with calcite and chlorite. In turn, this variety passes sharply into well-jointed red rock, which, for the next twenty yards,
becomes progressively more massive.

The dyke at its northern end makes a sharp penetrating contact with the slates, irregularly truncating the cleavage planes.

**Petrography:**

The microscope shows that the field variations are purely mechanical, and that, conversely, the wide textural variations disclosed in thin section are in no way reflected in the appearance of the rocks in the hand-specimen.

The specimens of which slides have been made can be classified into the following groups:

(a) **Porphyritic Variety:** fine-grained matrix containing felspar microlites.

(b) **Variety with matrix intermediate between (a) and (c).**

(c) **Coarse-grained Variety:** "cracked" orthophyric felspars; characterised by biotite.

(d) **Variety intermediate between (c) and (e).**

(e) **Variety with saussuritised, brown felspars;** characterised by chloritisation of all mafics.

There is a general relationship between these varieties and their position in the dyke. Variety (c) occurs on either side of (e), which is more or less central in position and the most extensive. Variety (d) is intermediate between these two, both (d) and (a) are, however, practically suppressed on the northern limb of the dyke. (Plate III)
Variety (a):

This has a characteristic porphyritic texture, with phenocrysts of plagioclase and more or less chloritised biotite (Pl. VIII fig. 4).

The felspar phenocrysts, though occasionally showing nearly euhedral outlines, are generally grouped as irregular aggregates. Angles and edges may have rounded outlines, suggesting resorption. Crystal margins frequently grade into the matrix, with no sign of a definite edge.

Irregular patchy inclusions in the phenocrysts, having crenulate or smoothly rounded margins are common; a general tendency towards concentration in the crystal centres being observed (cf. p. 12). The margins are always clear. In composition the patches resemble the matrix, with which, it is presumed, they are to be identified. This interpretation suggests that some of the phenocrysts were unable to clear themselves of matrix inclusions; perhaps because they separated from a highly viscous magma.

The felspars always have an appearance of being highly altered. All the larger and many of the smaller crystals are nearly opaque, apart, as a rule, from a narrow marginal zone. This is only in part due to the development of a kaolin type of mineral. Under high magnification a confused mass of "sericite", chlorite, clear albite and epidote is revealed. The clear margins are occupied by clear albite, which sends penetrating processes into the opaque products of
the interior of the crystals. Those felspars which are clear throughout have refractive indices lower than that of balsam and are entirely made up of unzoned albite. The alteration processes appear to be, therefore, saussuritisation (responsible for the opaque products), succeeded by marginal or complete replacement by albite.

No original felspar can be found; hence the original composition can be inferred only from that of the felspars of the internal varieties of the dyke.

A greenish biotite, more or less pseudomorphed by chlorite, is relatively abundant. The replacing chlorite has the same bright green colour and can be distinguished only by its low interference colours. Granules of epidote are associated with the chlorite. Frequently the chlorite develops beyond the original outlines of the biotite. Replacement of the biotite by another colourless mica, has, in some cases, also occurred. The later mica is probably more closely related to sericite as it forms tiny flakes making an angle with the original cleavage.

Certain dirty, brownish, flaky aggregates showing variable birefringence may be partly micaceous, partly chloritic. Similar patches, having approximately hexagonal outlines, may represent former basal sections of mica. They show an undulose extinction, resembling that often seen in biotite flakes. The extreme alteration of the rock, however, makes identification impossible.
There is no evidence suggesting the original occurrence of hornblende in the rock.

In addition to the chlorite which partially pseudomorphs biotite, a pale green penninite occurs. Occasionally it has sharp polygonal margins; more generally it sends off penetrating fibres and irregular processes into the matrix. Enclosed mesh-works of black ore are common, suggestive of derivation from an earlier mineral.

The characters of certain albite-chlorite "clots" are of considerable interest. Plate X, fig. 1 illustrates a typical example. Here there are areas of pale green chlorite partially or completely enclosing crystals of clear albite. The albite is penetrated by chlorite flakes and fibres; the former is evidently the earlier mineral.

The matrix is fine-grained with an abundance of tiny felspar laths and needles, forming a characteristic criss-cross pattern. The background of the matrix is partly felspathic, but contains, in addition, much disseminated chloritic and black ore material.

Little "nests" of quartz aggregates are scattered through the matrix. The latter is evidently replaced, since the felspar microlites are abruptly truncated.

Variety (c): (Plate VIII, Fig. 2).

This is a coarser-grained rock than the last described; all the constituent minerals are visibly crystalline and the
porphyritic texture is correspondingly less conspicuous.

The dominant mineral is a relatively clear, often nearly euhedral felspar. All gradations in size occur. The felspars, especially the smaller ones, tend to occur in squat-rectangular or lath-like forms. The grain-size and dominant form of the felspars controls the appearance of the rock, giving it an orthophyric texture.

The sharp edges, emphasised cleavages and cracks, and streaky grey clouding are features which are characteristic of the felspars of this variety.

Lamellar twinning is usual and slight marginal zoning not uncommon in larger crystals.

<table>
<thead>
<tr>
<th>Extinction Angles:</th>
<th>Centres of zoned crystals:</th>
<th>Smaller crystals and marginal zones:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>14°</td>
<td>7°</td>
</tr>
<tr>
<td>Reffactive Indices:</td>
<td>α 1.545</td>
<td>1.540</td>
</tr>
<tr>
<td></td>
<td>γ 1.553</td>
<td>1.548</td>
</tr>
<tr>
<td>Optic Angle:</td>
<td>Large - approaching 90°.</td>
<td>Negative.</td>
</tr>
</tbody>
</table>

The felspar therefore varies in composition from Ab\textsuperscript{70}\textsubscript{30} to Ab\textsuperscript{70}\textsubscript{30}.

The slight clouding, presumably due to a kaolin type of mineral, is usually concentrated along streaks isolating clear areas of felspar.

Flakes of a colourless mica, showing high birefringence, are abundant. It appears to be an early replacement, since with "kaolin" it occurs in otherwise unaltered felspars.
Clear albite, usually untwinned, develops along the margins and twinning planes of the oligoclase crystals. Smaller laths may become completely albitised. The albite is recognisable by its low refractive index and positive sign.

Chloritisation apparently occurs in two stages. The earlier chlorite forms tiny flakes in the more altered-looking felspars. Where the felspar is clear albite little or none of this chlorite occurs. Later chlorite forms fibrous streaks and veinlets which penetrate the albite along cleavages and otherwise.

Granules of epidote, smaller than those in pseudomorphous chlorite are relatively common. Frequently they are associated with streaks of albite, thereby suggesting derivation from the anorthite part of the felspar.

Quartz effects an indiscriminate type of replacement. Commonly it occupies interstices, from which it replaces neighbouring crystals. Lobate encroachments into felspar margins and penetrations along cleavage planes are frequent. With albite, however, quartz forms more complex intergrowths. Occasionally quartz replaces and partly isolates portions of albite crystals, leaving rounded margins. Elsewhere quartz penetrates albite in fine striae or vermicules, which can often be traced back to recognisable discrete crystals. Sometimes albite tablets have been almost entirely pseudomorphed, leaving only a fine albite mesh.

The biotite is identical with that occurring in the
marginal variety (a). It is green or brownish-green in colour and has a fibrous structure. Chloritisation has proceeded along the margins and cleavages, associated with the deposition of granules of epidote and black ore.

Aggregates of chloritised biotite often enclose mesh-works of black ore and relatively large epidote crystals. The chlorite is paler in colour than the biotite and shows grey interference tints. From larger masses fibrous processes pass into adjacent felspars. Quartz, however, always cuts the chlorite fibres and is therefore later.

The biotite and chlorite are often nearly opaque, owing to a brownish-grey clouding, probably due to a fine dispersion of hydrated ores. Definite characters are therefore impossible to determine.

Epidote is always of a cloudy, brown, granular variety. The true yellow colour and pleochroism can only be recognised under high-power. Optical properties are difficult to determine.

Apart from the flaky chlorite pseudomorphing biotite, chlorite occurs as fibrous patches, having a replacing habit towards the surrounding matrix minerals. Sometimes these form marginally on biotite-chlorite clots. Occasionally plumose, fibrous aggregates of a pale green penninite are found.

Calcite is uncommon in this variety. Its importance increases through the intermediate variety (d) to a maximum
in (e).

Tiny needles and basal sections of apatite are abundant. They are especially frequent in biotite or chlorite-chlorite clots, but are unusual in the felspars.

**Variety (e):** (Plate IX, Fig. 1).

This is characterised by brown felspars, without sharp edges, cleavages or cracks, as in variety (c). The texture is typically orthophyric. No original biotite remains; the chlorites are clear and pale green in colour and calcite is abundant.

The felspar is all either saussuritised or nearly entirely albiteised. The final albition has developed along twinning planes or slightly winding channels, evidently replacing the confused "saussurite".

The chlorite is a clear, pale-green penninite with abnormal blue interference tints and negative optical sign. Often it has definite rectangular outlines, but if it be a pseudomorph there is no evidence as to the identity of the original mineral. It is noteworthy that chloritisation is not complete in the intermediate variety (d).

Epidote granules are abundant, both as an alteration product of the felspars and as an associate of chlorite.

Calcite has a characteristically grey, cloudy appearance; it is generally pseudomorphous after felspar, biotite or chlorite. In the intermediate variety (d) there are calcite pseudomorphs in which residual fibres of biotite
Irregular grains and aggregates of quartz are abundant. Often they appear to have a purely interstitial disposition, but closer examination reveals lobate extensions into the felspars or chlorite, transgressions across felspar laths and penetrations along the cleavage planes in the felspars. Calcite, however, is never affected by the quartz and is therefore presumably of contemporaneous or later development.

Colourless needles of apatite are common throughout the rock, enclosed by both felspars and chlorite.

The "Red Spot".

This is a patch of reddish material occurring in variety (e) with a calcite-quartz infilling towards one side. It was in no way connected with vein material, since, when found, the spot was entirely surrounded by normal dyke rock.

The spot shows striking replacement phenomena by albite and quartz. All the original felspar has been albitised and radial sheafs of albite are common. The spot is rimmed by these sheafs, which often enclose lath-shaped pseudomorphs of original felspar. Quartz has also been an active replacing mineral, occurring both interstitially and as penetrations into the felspar laths. Frequently, too, zig-zag fingerings of quartz cut across the albite sheafs.

The infilling itself (see Pl. XI, fig. 1) consists of chlorite, quartz and calcite, developed in that order from the margins to the centre. These minerals are surrounded by a
series of albite sheafs. Each of these minerals in turn shows replacement relations towards the earlier mineral on the outer side and normal growth features towards the interior of the infilling.

The evidence for the order of replacement is as follows:

(i) Calcite is latest, since it replaces both chlorite and quartz with development in each case of similar marginal textures. The quartz appears to have straight edges inwards; actually, under high magnification, they are found to be minutely ragged and indented by calcite. Towards the margins of the infilling (that is, where development was possible only by replacement), the calcite ceases to be optically continuous and is formed of patchy granules.

(ii) Quartz is later than the chlorite and generally occurs between chlorite and calcite. Inwards it has developed in tooth-shaped crystals; towards chlorite, however, it forms irregular patchy replacements which truncate the chlorite fibres.

(iii) Albite is replaced by chlorite and quartz, both of which cut across the albite sheafs.

Thus the order of mineral replacement is established:

1) Albite, 2) Chlorite, 3) Quartz, 4) Calcite.

The Significance of the Internal Modifications of Type III.

As regards the differences between varieties (a), (b) and (c) the controlling factor appears to have been distance from the margin, the effect of which on the course of crystal-
lisation has been a corresponding change from a porphyritic to a coarse-grained equigranular type of rock.

The relation between (c) and (e) is of quite a different nature, despite the fact that there is every gradation from one to the other. The main features distinguishing the extremes are:

(i) Marked differences in the felspars as regards degree of "definition", colour and intensity of alteration. The colour-change — from grey in (c) to reddish-brown in (e) — is probably due to hydration of disseminated iron-ores in (e). Moreover, original oligoclase remains only in (c), whereas in (e) a more advanced stage of alteration and replacement has been reached.

(ii) No original biotite remains in variety (e).

(iii) Variety (e) shows a much greater development of calcite.

It is therefore clear that variety (e) has suffered more severe hydrothermal alteration than (c). Nor can its position in the dyke be without significance. Just as (c) approaches nearer to the contact on the northern than on the southern limb of the dyke, so does the more altered variety (e). The latter thus occupies what may be termed the "genetic centre" of the dyke.

The process of hydrothermal alteration in this example is considered to be autolytic (see pp. 65-71). It is convenient at this juncture to summarise the implications of this hypothesis.

As the magma crystallised, water and carbon dioxide,
together presumably with some excess of soda, would become concentrated in the still fluid portions, particularly in the "genetic" centre of the dyke. Eventually aqueous solutions, containing sodium and carbonate ions and silica would remain. First the silica and sodium assisted in the breakdown of the plagioclase into albite and epidote, (Na Si) driving out (Ca Al). At the same time hydration led to chloritisation of biotite with release of iron-oxides, together with potassium. The latter would facilitate production of "sericite". Later, residual silica was deposited as quartz, initially in the interstices and afterwards as replacements of such neighbouring minerals as obstructed its crystal development. Finally, "carbonate" with released calcium formed the latest replacing mineral, calcite.

The absence of any fine network of veins supports this hypothesis of auto-alteration by residual solutions, since the latter would already be widely dispersed throughout the interior of the dyke.

The evidence of the "Red Spot" is also of significance, since it appears to have been a centre of intense hydrothermal alteration. The red igneous material is probably only the normal dyke-rock which has suffered extreme alteration and replacement, the encircling rim of albite sheafs representing the "fore-front" of this localised activity. The calcite-quartz-chlorite area is partly an infilling, towards the centre; partly a replacement, towards the margins. The order of replacement in the "spot" and the order of infilling of the
cavity, both agree with the observed order of replacement throughout the dyke.

The "spot" therefore represents a centre of unusual concentration of volatile material, which, after the rock had crystallised, reacted with the primary minerals in the manner already outlined for the dyke as a whole.

Von Eckermann (1928) describes similar quartz or calcite infillings in a lamprophyre ("hamrongite") from near Gavle in northern Sweden. Both these minerals also occur interstitially.

Miss Reynolds (1931) figures a drusy cavity, closely analogous to the "red spot", in a kersantite from the Ards Peninsula, Co. Down. Here, calcite, the final infilling material, replaces both quartz and felspar.

It is significant, too, that it is the interior of the northern dyke (Type II), that shows the greatest alteration features. Here again, veins along which hydrothermal solutions could have travelled from an external source, are extremely rare. Quartz-albite nests, however, which appear to have the same significance as the "red spot", are common, and in the interior of the dyke are the centres of intense hydrothermal activity.

The analyses of varieties (c) and (e) shown on p. 44 serve to emphasise the soda-rich character of these rocks. The close similarity between the two analyses suggests that a molecular rearrangement would largely account for the mineralogical differences observed under the microscope.
### MINERAL SUMMARY: TYPE III.

<table>
<thead>
<tr>
<th>Primary Minerals:</th>
<th>Secondary Minerals:</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Plagioclase</strong>: (Ab\textsubscript{ab} - Ab\textsubscript{an})</td>
<td>Alteration and replacement:</td>
</tr>
<tr>
<td>Form: euhedral with sharp edges; smaller crystals rectangular. Lamellar Twinning. Marginal Zoning occasionally. Extinction Angles: \begin{align*} \text{margins} &amp;:\phantom{1.540} 7^\circ \ \text{centres} &amp;:\phantom{1.540} 14^\circ \end{align*} Refractive Indices: \begin{align*} \alpha &amp;:\phantom{1.540} 1.540 \ \gamma &amp;:\phantom{1.540} 1.548 \end{align*} Optic Sign: Negative.</td>
<td>Frequently original felspar remains unreplaced. &quot;Kaolin&quot; not abundant; cloudy patches. &quot;Sericite&quot;; abundant flakes. Chlorite; greenish flakes and fibres. Albite; clear, often untwinned; frequently effects complete replacement. Epidote; tiny granules. Quartz; irregular areas of replacement.</td>
</tr>
<tr>
<td><strong>Biotite</strong>:</td>
<td>Extensive chloritisation together with granules of epidote and calcite.</td>
</tr>
<tr>
<td>Green or brown colour. Ragged and fibrous; sometimes in clotted aggregates with ores. Often nearly opaque owing to clouding.</td>
<td>Quartz and calcite - indiscriminate replacement.</td>
</tr>
<tr>
<td><strong>Accessories</strong>:</td>
<td></td>
</tr>
<tr>
<td>Black Ores; Apatite needles.</td>
<td></td>
</tr>
<tr>
<td><strong>Matrix</strong>: (where porphyritic)</td>
<td>- contains abundant tiny felspar laths.</td>
</tr>
<tr>
<td>NO.: 80.</td>
<td>CHEMICAL ANALYSIS:</td>
</tr>
<tr>
<td>----------</td>
<td>--------------------</td>
</tr>
<tr>
<td>SI0₂</td>
<td>58.20</td>
</tr>
<tr>
<td>Al₂O₃</td>
<td>17.08</td>
</tr>
<tr>
<td>Fe₂O₃</td>
<td>3.07</td>
</tr>
<tr>
<td>FeO</td>
<td>1.94</td>
</tr>
<tr>
<td>MgO</td>
<td>4.82</td>
</tr>
<tr>
<td>CaO</td>
<td>1.96</td>
</tr>
<tr>
<td>Na₂O</td>
<td>6.18</td>
</tr>
<tr>
<td>K₂O</td>
<td>1.65</td>
</tr>
<tr>
<td>H₂O</td>
<td>0.40</td>
</tr>
<tr>
<td>H₂O⁺</td>
<td>2.36</td>
</tr>
<tr>
<td>CO₂</td>
<td>1.04</td>
</tr>
<tr>
<td>P₂O₅</td>
<td>0.27</td>
</tr>
<tr>
<td>TiO₂</td>
<td>0.92</td>
</tr>
<tr>
<td>MnO</td>
<td>0.17</td>
</tr>
<tr>
<td></td>
<td>100.54%</td>
</tr>
</tbody>
</table>

**Analyst:**
A. Huddleston.

<table>
<thead>
<tr>
<th>NO.: 89.</th>
<th>CHEMICAL ANALYSIS:</th>
<th>NORMATIVE COMPOSITION:</th>
<th>CALCULATED MINERAL COMPOSITION:</th>
<th>1</th>
</tr>
</thead>
<tbody>
<tr>
<td>SI0₂</td>
<td>58.10</td>
<td>Quartz 7.28</td>
<td>Quartz 9.39</td>
<td>Qu</td>
</tr>
<tr>
<td>Al₂O₃</td>
<td>17.38</td>
<td>Albite 51.02</td>
<td>Albite 57.70, 68.74</td>
<td>Al</td>
</tr>
<tr>
<td>Fe₂O₃</td>
<td>2.49</td>
<td>Anorthite 6.68</td>
<td>Anorthite 5.26, 27.96</td>
<td>Fe</td>
</tr>
<tr>
<td>FeO</td>
<td>2.76</td>
<td>Corundum 3.14</td>
<td>Corundum 1.54</td>
<td>Ca</td>
</tr>
<tr>
<td>MgO</td>
<td>4.10</td>
<td>Orthoclase 10.24</td>
<td>Orthoclase 10.24, 12.34</td>
<td>Ca</td>
</tr>
<tr>
<td>CaO</td>
<td>2.26</td>
<td>Hypersthene 10.84</td>
<td>Chlorite 13.42</td>
<td>Ca</td>
</tr>
<tr>
<td>Na₂O</td>
<td>6.03</td>
<td>Dolomite 1.90</td>
<td>Calcite 1.90</td>
<td>Ca</td>
</tr>
<tr>
<td>K₂O</td>
<td>1.73</td>
<td>Magnetite 3.61</td>
<td>Ores 4.84</td>
<td>Or</td>
</tr>
<tr>
<td>H₂O⁺</td>
<td>0.40</td>
<td>Ilmenite 1.96</td>
<td>Apatite 0.60</td>
<td>Or</td>
</tr>
<tr>
<td>H₂O</td>
<td>2.61</td>
<td>Apatite 0.60</td>
<td>Water 1.43</td>
<td>Or</td>
</tr>
<tr>
<td>CO₂</td>
<td>0.90</td>
<td>Water 3.21</td>
<td>Water 1.43</td>
<td>Or</td>
</tr>
<tr>
<td>P₂O₅</td>
<td>0.25</td>
<td>Water 3.21</td>
<td>Water 1.43</td>
<td>Or</td>
</tr>
<tr>
<td>TiO₂</td>
<td>1.03</td>
<td>Water 3.21</td>
<td>Water 1.43</td>
<td>Or</td>
</tr>
<tr>
<td>MnO</td>
<td>0.19</td>
<td>Water 3.21</td>
<td>Water 1.43</td>
<td>Or</td>
</tr>
<tr>
<td></td>
<td>100.43%</td>
<td>SYMBOL: II.4524</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Analyst:**
A. Huddleston.

*Chlorite calculated as Penn formula: Fe₀.₄Mg₀.₃Si₀₂·Al
<table>
<thead>
<tr>
<th>NO.</th>
<th>CHEMICAL ANALYSIS</th>
<th>NORMATIVE COMPOSITION</th>
<th>CALCULATED MINERAL COMPOSITION</th>
<th>MODAL(VOL.) COMPOSITION</th>
</tr>
</thead>
<tbody>
<tr>
<td>89</td>
<td>SiO₂ 58.10</td>
<td>Quartz 8.39</td>
<td>Quartz 9.93</td>
<td>*Quartz 3.3</td>
</tr>
<tr>
<td></td>
<td>Al₂O₃ 17.86</td>
<td>Albite 52.27</td>
<td>Albite 55.53</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Fe₂O₃ 3.07</td>
<td>Anorthite 3.26</td>
<td>Anorthite 56.87</td>
<td></td>
</tr>
<tr>
<td></td>
<td>CaO 1.96</td>
<td>Corundum 3.94</td>
<td>Corundum 1.34</td>
<td></td>
</tr>
<tr>
<td></td>
<td>MgO 4.82</td>
<td>Orthoclase 9.73</td>
<td>(Orthoclase) 9.73</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Na₂O 6.18</td>
<td>Hypersthene 10.29</td>
<td>Chlorite* 15.00</td>
<td></td>
</tr>
<tr>
<td></td>
<td>K₂O 1.65</td>
<td>Dolomite 3.13</td>
<td>Calcite 3.13</td>
<td></td>
</tr>
<tr>
<td></td>
<td>H₂O 0.40</td>
<td>Magnetite 4.34</td>
<td>Ores 5.4</td>
<td></td>
</tr>
<tr>
<td></td>
<td>CO₂ 1.54</td>
<td>Hasmatite 2.76</td>
<td>Apatite 0.64</td>
<td>etc. 100%</td>
</tr>
<tr>
<td></td>
<td>P₂O₅ 0.27</td>
<td>Ilmenite 1.74</td>
<td>Water 0.91</td>
<td></td>
</tr>
<tr>
<td></td>
<td>TiO₂ 0.92</td>
<td>Water 2.76</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>MnO 0.18</td>
<td>100.49%</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Symbol:** II.(4)5.2.4. *Quartz largely intergrown with Felspars, hence low modal value.

<table>
<thead>
<tr>
<th>NO.</th>
<th>RED ROCK DYKE</th>
<th>CHEMICAL ANALYSIS</th>
</tr>
</thead>
<tbody>
<tr>
<td>34</td>
<td>SiO₂ 67.32</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Al₂O₃ 16.04</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Fe₂O₃ 0.41</td>
<td></td>
</tr>
<tr>
<td></td>
<td>CaO 0.98</td>
<td></td>
</tr>
<tr>
<td></td>
<td>MgO 1.48</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Na₂O 2.02</td>
<td></td>
</tr>
<tr>
<td></td>
<td>K₂O 5.06</td>
<td></td>
</tr>
<tr>
<td></td>
<td>H₂O 0.70</td>
<td></td>
</tr>
<tr>
<td></td>
<td>CO₂ 2.20</td>
<td></td>
</tr>
<tr>
<td></td>
<td>P₂O₅ 0.38</td>
<td></td>
</tr>
<tr>
<td></td>
<td>TiO₂ 0.52</td>
<td></td>
</tr>
<tr>
<td></td>
<td>MnO 0.18</td>
<td>Erase 100.07%</td>
</tr>
</tbody>
</table>

**Symbol:** I.4.2.4. *Quartz 26.60, Orthoclase 11.72, Albite 42.76, Anorthite 4.75, Corundum 3.81.

| Analyst: A. Huddleston. *Calcite largely as partial pseudomorphs, hence high modal value. *Chlorite calculated as Penninite, having formula:** FeO₄MgO₃SiO₆Al₂O₃₄H₂O. |
TYPE IV:  (LEUCOCRATIC) ALBITE-PORPHYRY.

This type forms large, wall-like dykes, sometimes over 10 feet in width, always with well-defined straight margins. They are intruded parallel to the strike of the country rocks and their relations to the cleavage-planes are always discordant. The large dykes at Red Rock near Balmae (south of Kirkcudbright) and on the eastern foreshore of Brighouse Bay, and several other strike dykes near Kelton Hill (three miles south-west of Castle Douglas) are typical examples.

The dyke at Red Rock has a width of approximately 30 feet and is well-exposed on the foreshore. Inland, within 60 yards it is displaced south-eastwards by two small faults. The displacement in each case is about the same as the width of the dyke.

The rock is homogeneous in appearance, of a pink or cream colour and has a flaky fracture. Fine jointing, parallel to the dyke margins, is especially marked towards the contacts. Weathered surfaces show numerous small felspar phenocrysts. Calcite veins are common, sometimes penetrating the slates.

Petrography:

The rock is an interesting type inasmuch as almost all its recognisable constituents are albite.

The felspar phenocrysts occur as rectangular tablets, irregular or indented crystals, or glomero-porphyritic aggregates. Generally the degree of alteration is not
considerable; slight clouding and scattered flakes of a sericitic mica being usual. Occasionally calcite has replaced patches of the felspar.

**Extinction Angles:** 17° - 19°.

**Optic Sign:** Positive.

**Refractive Indices:**

\[ \alpha = 1.526 \]
\[ \gamma = 1.536 \]

Evidently the composition approximates closely to that of albite.

**Mafic minerals are rare.** Original biotite has not been observed, but chlorite-"sericite"-calcite aggregates, having a form suggestive of pseudomorphs after biotite, are present.

The **matrix** is as pale as the felspar phenocrysts and its dominant constituent has a low refractive index. It can largely be resolved even under the low power and consists principally of angular albite crystals, together with calcite, "sericite", and some chlorite.

Though there is nothing in the mode which can be identified as **Quartz**, the latter mineral forms no less than 26.6 per cent of the normative composition of the rock.

The **accessories** are a lustrous, pale yellow metallic sulphide, sometimes as relatively large crystals, and tiny needles of apatite.

See analysis on p. 44.
**MINERAL SUMMARY: TYPE IV.**

<table>
<thead>
<tr>
<th>Phenocrysts:</th>
<th>Matrix:</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Albite:</strong></td>
<td>Dominantly Pelspathic.</td>
</tr>
<tr>
<td>Form: euhedral tablets; angular fragments and aggregates.</td>
<td>Pale colour.</td>
</tr>
<tr>
<td>Refractive Indices: α 1.526, γ 1.536.</td>
<td></td>
</tr>
<tr>
<td>Optic Sign: Positive.</td>
<td></td>
</tr>
<tr>
<td>Alteration: slight - &quot;Kaolin&quot; mineral; &quot;Sericite&quot;.</td>
<td></td>
</tr>
</tbody>
</table>

**Mafic Minerals: (rare)**

| Chlorite-"Sericite"-Calcite pseudomorphs; (? after Biotite) Rare Epidote granules with chlorite. |

**Accessories:**

| Opaque yellow Sulphide Ores; Tiny Apatite needles. |

---

**TYPE V: PLAGIOPHYRES.**

These were encountered at Portling Bay, in the Rockcliffe district. Except for a relatively narrow belt, widening eastwards, in the neighbourhood of the high-water mark, the foreshore at Portling Bay consists of Carboniferous basement beds. A number of irregular dykes occur in the Silurian. In the central part of the bay, just below the road, there are outcrops of two plagiophyre dykes and one of xenocrystic type,
the latter being east of the others.

The plagiophyres have a dark, green or grey colour, with scattered pinkish felspars and numerous fine joints. Specks of pyritic ore are freely disseminated. Felspathic veins and streaks are common, as well as irregular veins and impregnations of carbonates.

**Petrography:**

The dyke and veins will be described separately:

(a) **Main-Dyke Plagiophyre:**

The rock consists of about equal proportions of light and dark coloured minerals, with an evenly medium-grained texture and no groundmass.

The felspars are usually euhedral in form, ranging from rectangular tablets 1.0 to 1.5 mm. in length, down to small laths. All crystals tend to be arranged in a decussate pattern, modified somewhat by the mafic minerals, whilst sometimes the "hornblende" pseudomorphs interrupt the regular form of the felspars, elsewhere laths of felspar penetrate the pseudomorphs or "control" their outlines.

Where not obscured by alteration, the felspars show excellent zoning and lamellar twinning.

The central zones have extinction angles of $18^\circ$ - $19^\circ$ and a refractive index higher than that of balsam, corresponding to sodic andesine ($Ab_xAn_y$). The margins have a lower refractive index than balsam and are evidently near albite in composition.
The cores of the crystals are often nearly opaque as a result of clouding by alteration products; marginally they may be nearly or completely fresh. The alteration is of the saussuritic type, with abundant flakes of sericitic mica, rarer chlorite, clear albite and a dispersed brownish material, presumably a clay mineral. Often, too, the central zones contain disseminated flakes of calcite, resulting in a grey clouding which "twinkles" on rotation in polarised light.

The mafic minerals have been entirely replaced by chlorite. According to their form, pseudomorphs after hornblende and biotite can be recognised. These diagnoses are supported by the fact that the chlorite replacing presumed hornblende differs from that after biotite. The former always occurs as small green flakes and fibres, with an approach to parallel orientation; has a prussian-blue interference tint; and is associated with numerous tiny granules and granular aggregates of epidote and sparse ores. The flakes are too small and felted for determination of the optic sign.

The chlorite after biotite has a clear emerald green colour, with rarer epidote and ores, and preserves the biotite cleavage. It shows a rusty-brown or indigo interference tint, gives a positive optic sign and has an absorption formula $X > Y > Z$. These properties agree with those for positive penninite. Frequently it forms irregular patches with winding processes cutting indiscriminately across the felspars. Here replacement has occurred at a late stage.

Accessories are apatite, inacicular needles, and sulphide
ores in numerous grains; both are included by all the other minerals.

Besides the minerals already mentioned, others occur which are characteristic of a hydrothermal assemblage; their development is probably related to the alterations which have affected the primary constituents.

**Clear albite**, distinguished by its refractive index, often free from twin-lamellae, is frequently moulded irregularly on the outer plagioclase zones; it may also enter into dentate intergrowths with quartz. In one slide a thin albite vein (up to about 0.02mm. in width) replaces all the primary minerals in its course. Occasionally calcite fills the vein for short distances. When it passes through the hornblende pseudomorphs only some of the chlorite fibres disappear and those left have lost their ubiquitous epidote granules. Significant, too, is the fact that the clear patches of penninite cut across the vein and were therefore relatively later. This is concordant with the relative order established between penninite and albite in Types II and III.

**Quartz**, though in part occupying the interstices of the felspar mesh-work also embays and sends fingering processes into the felspar laths. Irregular crystals of quartz may lie right across the felspars or entirely within them. As calcite alone is unaffected by the quartz it is presumably of later development.

Apart from the diffuse flakes in the felspars, calcite
forms clear, discrete crystals exhibiting replacement relations towards the earlier-formed minerals. Association with quartz and chlorite is common.

A micrometric measurement of one section (No. 27) gave the following mode (proportions by volume):

<table>
<thead>
<tr>
<th>Mineral Type</th>
<th>Proportion by Volume</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hornblende pseudomorphs</td>
<td>29.1</td>
</tr>
<tr>
<td>Penninite, including pseudomorphs after biotite</td>
<td>18.3</td>
</tr>
<tr>
<td>Plagioclase</td>
<td>47.1</td>
</tr>
<tr>
<td>Quartz</td>
<td>2.8</td>
</tr>
<tr>
<td>Calcite</td>
<td>4.0</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td><strong>101.3%</strong></td>
</tr>
</tbody>
</table>

(b) **Leucocratic Vein: Quartz-Plagiophyre**: (Pl. XII, Fig. 1).

The felspathic veins show a rather coarser and more even grain than the main rock. Felspars range in size up to about 2.5 mm in length and the smaller crystals often have nearly square outlines. They exhibit characteristic euhehedral outlines and show marked zoning. The central zones have extinction angles (about 15°) corresponding with calcic oligoclase.

The chlorite is all of the emerald-green, positive penninite variety, with associated epidote granules and black ore. Much of it is evidently pseudomorphous after biotite, the cleavage and general form of which are still preserved. Occasionally the chlorite occurs as elongated, straight or twisted crystals up to a millimetre or so in length and a small fraction of a millimetre in width. Penninite also occurs as irregular flakes and aggregates, with frequent fibrous processes
replacing surrounding minerals.

Albite forms irregular outgrowths from the plagioclases. Interstitial quartz embays the felspars, just as in the main dyke rock. Calcite is frequent, usually in association with quartz.

The mineral composition and textures of this variety are thus very similar to those of the main dyke. The relative proportions of the constituents are, however, widely different. Hornblende pseudomorphs are absent and penninite constitutes only about 9 per cent. Quartz has increased considerably, to approximately 10 per cent, whilst plagioclase, constituting most of the remainder, is also much higher.

Conclusions:

(a) Genesis of the Magmas.

The close similarity between the main dyke and its more leucocratic veins, in texture, and mineral composition and type of hydrothermal alteration, indicates an intimate relation between their respective magmas. In each case the magma was endowed with the same tendencies during crystallisation, both as to the production of minerals and the development of textures. Crystal-sinking of the mafic constituents in situ to form the residual magma of the felspathic veins can scarcely be invoked, but the possibility must not be overlooked that the veining magma was the squeezed or drained out residual liquid from the main dyke after the latter had practically consolidated, the process taking place more or less in situ. The complete absence of former hornblende, reduction in the proportion of
"biotite" and increase of quartz seem favourable to this view. Against this mechanism may be cited the following points:

(i) A residual liquid produced after most of the main magma had consolidated would be unlikely to deposit zoned plagioclase with central zones nearly as calcic as the crystals of the parent.

(ii) Although there is considerably more quartz in the leucocratic veins, albite has not increased in corresponding proportion. A late magmatic liquid would theoretically contain a higher ratio of albite to calcic felspar than that found. Hence it is extremely unlikely that any process of crystallisation-differentiation in situ could account for the proved facts.

It is presumed, therefore, that the leucocratic vein magma was differentiated by some mechanism operating in depth or by some syntectic-metasomatic process operating nearly in situ. Of neither process is there any evidence and it would serve no useful purpose to speculate further.

(b) Hydrothermal Activity.

Both the dykes and the vein systems show identical hydrothermal textures, which are characteristic of the type, and in marked contrast to those of Types II and III.

It is quite evident that the group of hydrothermal minerals — albite, chlorite, quartz and calcite — were most actively developed after the magma had practically consolidated. For even albite, the earliest hydrothermal, forms veins with
sharp margins, pointing definitely to development in a solid medium.

Such activity is frequently regarded as having been effected by low temperature aqueous solutions, arising from some source in depth, after an indefinite pause succeeding complete consolidation.

If such were here the case it seems difficult to explain why the development of hydrothermal quartz, for example, is much greater in the veins than in the main dyke.

The facts suggest that the hydrothermal agencies were inherent in each initial magma and that early crystallisation gradually increased the concentration of "volatiles". (See further pp. 65-71).

(c) Nomenclature.

Tyrrell (1912 - 13) suggested the term "plagiophyre" for rocks with the texture and appearance of orthophyres. Most of the features of the examples he described accord closely with those of the type here under discussion. "The rock consists essentially of euhedral plagioclase in short laths, with diverse, non-fluidal arrangement". The plagioclases are similarly "highly zonal" and have the same composition, whilst "chloritic decomposition products" and "iron-ores" occupy the interstices.

He also records a quartz-plagiophyre with an abundance of "interstitial euhedral quartz" and notes that the chlorite "has probably overspread its original euhedral boundaries" and "sends processes into the felspars".
The terms "plagiophyre" and "quartz-plagiophyre" appear, therefore, to be quite appropriate for the dyke-rock and the leucocratic veins, respectively.

**MINERAL SUMMARY: TYPE V.**

<table>
<thead>
<tr>
<th>Primary Minerals:</th>
<th>Secondary Minerals:</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Plagioclase:</strong>  ((\text{Ab}_x\text{An}_y))</td>
<td>Brownish clouding of &quot;Kaolin&quot; mineral.</td>
</tr>
<tr>
<td><strong>Hornblende:</strong></td>
<td>Flakes and fibres of Chlorite (prussian-blue int. tints) Much granular Epidote, Rarer Ores.</td>
</tr>
<tr>
<td>Pseudomorphs with hornblende form. (Absent from felspathic veins).</td>
<td></td>
</tr>
<tr>
<td><strong>Biotite:</strong></td>
<td>Clear green Penninite (indigo-blue int. tints) (positive sign). Rarer Epidote granules.</td>
</tr>
<tr>
<td>Pseudomorphs with biotite form.</td>
<td>Positive Penninite also oversteps &quot;biotite&quot; margins - and as irregular patches.</td>
</tr>
<tr>
<td><strong>Accessories:</strong></td>
<td>Quartz: interstitial and encroaching on felspars. Calcite: similar distribution.</td>
</tr>
<tr>
<td>Apatite needles; Black Ores and Pyrite.</td>
<td></td>
</tr>
</tbody>
</table>
TYPE VI: XENOCRYSTIC VARIETIES.

A number of dykes occur in which the phenocrysts have an appearance and composition quite incompatible with their having crystallised directly and completely from the same magma as that forming the matrix. The most striking occurrence is a dyke from the eastern end of Portling Bay, nine miles south-south-east of Dalbeattie.

The rock is markedly porphyritic in texture, with large phenocrysts of quartz, felspar and biotite in a fine-grained matrix.

Most common are large rounded tablets of orthoclase, with a lower refractive index than that of balsam; many of these show simple twinning and somewhat wavy extinction. Perlitic replacement by albite is frequent, and in some cases albition has proceeded until little orthoclase remains. Large plagioclases, showing strong oscillatory zoning, occur in aggregates; the most calcic component ranges up to oligoclase-andesine in composition.

The quartz shows very striking resorption phenomena. Optically continuous, rounded relics of formerly continuous quartz crystals are now surrounded and penetrated by the matrix. The matrix itself becomes modified against the resorbed margins, resulting in "spectral" zoning following the outlines of each individual. (See Pl. XI, fig. 3).

Biotite also exhibits rounded outlines. It is of a fawn-brown colour with a woody, fibrous cleavage and strong
pleochroism. Partial or complete alteration to chlorite is frequent. Some chlorite-calcite pseudomorphs are, however, more probably after hornblende.

The matrix has an indefinite, microfelsitic texture. The poorly-formed felspars occasionally show lamellae and always have refractive indices lower than that of balsam. The presence of albite is therefore indicated.

Patchy calcite, occasionally occupying vein-like channels, has been a late replacing mineral, affecting biotite, felspar and matrix alike.

Accessories are apatite and ores; both of which are frequently included in the biotite.

The following features suggest that the rock is a mixture of partly resorbed xenoliths and xenocrysts in a magma which broke through a pre-existing plutonic or pegmatitic rock: (a) The occurrence of xeno-porphyritic aggregates; (b) The strong resorption which shows that marked instability existed between phenocrysts and liquid; (c) The occurrence of perthite, the crystallisation of which from a magma under hypabyssal conditions, such as doubtless controlled the consolidation of the matrix, would be unlikely; (d) The occurrence of orthoclase, a mineral not found in any of the other types and therefore suspected to be foreign to the type in question.

In the Dalbeattie "granite" the following minerals have been described, (M. MacGregor, 1936): (i) Plagioclase, with oscillatory zoning and composition
of oligoclase-andesine (average);

(ii) Microperthite;

(iii) Biotite: ragged crystals, often in clots with hornblende and containing inclusions of apatite and iron-ores;

(iv) Hornblende: common variety.

The similarity between these and the xenocrysts suggests that the Dalbeattie granite or a similar rock might be looked to as a source for at least some of the xenocrysts.*

The following order of degree of resorption can be clearly made out in the rock under discussion:

1) Quartz - most;
2) Biotite;
3) Orthoclase and Perthite;
4) Plagioclase - least.

The matrix corresponds closely to a normal albitophyre. Since it would naturally deposit albite, the above order can be readily explained. The outer zones of plagioclase would be little affected, but orthoclase would tend to be replaced. It seems not unlikely that at least some of the perthite was formed in this manner. To an even greater degree would quartz depart from equilibrium, whilst biotite and hornblende naturally became chloritised. There would also be a quantity of calcite replacing these minerals indiscriminately.

Original Magma + Xenolithic Material => Matrix + Corroded and Altered Xenoliths and Xenocrysts

*(Cf. Fenner 1933).*
TYPE VII: LAMPROPHYRES.

A number of lamprophyres were briefly examined. All those encountered form narrow, irregular dykes, a few feet in width, usually with intricate indented margins, and accompanied by numerous short apophyses and embayments into the slates.

In appearance they are all dark grey or greenish, fine-grained, compact rocks with abundant glistening flakes of mica. Calcite veins are always common.

All are mica-bearing, but under the microscope many show an extreme type of carbonate alteration, whereby their original characters are largely obscured.

Petrography:

A brief description will be given of one relatively fresh dyke from the right bank of the Dee estuary near Nun Mill. This is the southernmost of three lamprophyres intruded into Silurian grits and volcanic agglomerate.

In texture the rock is strikingly porphyritic with abundant phenocrysts of biotite and carbonate pseudomorphs after olivine. The biotite is of a yellowish colour, but with a strong absorption parallel to X, of a deep fawn-brown. Hexagonal basal flakes are common, usually showing a number of concentric zones of deeper colour and giving a wavy extinction. Calcite filled veins and channels often penetrate the crystals. Silicified patches along the margins of the flakes are common.

Euhedral pseudomorphs, having the form of olivine, are
frequent (see Pl. XII, fig. 2). The edges are generally delineated by black ores, with carbonates occupying the interior. A staining test indicated that the carbonate is not calcite, but an iron- or magnesium-rich variety. Most of the carbonate has developed in minute flakes, having a general brownish coloration, but in places larger, colourless crystals are found.

The matrix is dark and cryptocrystalline, but shows numerous tiny shadowy laths of felspar. The latter are always too small and obscure to be determined optically.

Relations:

H. H. Read (1926) has shown, in Wigtownshire, that the mica-bearing dykes are restricted to the country south-east of a line from the Mull of Galloway to the neighbourhood of Creetown. A continuation of this line eastwards would leave the similar dykes of the Kirkcudbright area also to the south-east; they evidently belong to the same swarm.

CONCLUSIONS: AFFINITIES AND HYDROTHERMAL AGENCIES.

A characteristic and significant feature of Types II, III and V, especially as compared with Type I, is the striking evidence they exhibit of intense hydrothermal alteration and replacement. Although the altered types are distinguished among themselves by certain significant differences of texture, which, to some extent, correspond with differences of composition, they have, nevertheless, been affected by later agencies
which have produced a similar assemblage of minerals.

1). Replacement Textures.

Gilluly (1932) adduces strong textural, mineralogical and chemical evidence in favour of a replacement origin for the albite in granites near Sparta, Oregon. He shows that much of the present albite has replaced original andesine and that the rock as a whole is an albitised diorite. Quartz has in turn replaced albite, with development of textures which are identical with those of the same two minerals in Types II and III. The following comparable observations by Gilluly may be noted:

(i) General saussuritic alteration of the felspar is characteristic.

(ii) Clear albite replaces andesine as marginal coatings, penetrative veins and scattered patches.

(iii) Quartz replacements of albite vary from simple embayments to intricate pseudomorphs, in which only a mesh of albite remains.

(iv) Rosettes and myrmekitic intergrowths of quartz and albite frequently surround albite crystals.

Gilluly cites a wealth of literature in which other examples are described of replacement of plagioclase by albite and of the origin of myrmekitic textures by replacement.

Gilson (1929) has shown replacement of plagioclase by orthoclase and microcline in the Plioche District of Nevada. Bailey and Grabham (1909) described albite replacement
of plagioclase in basaltic lavas of Carboniferous and Permocarboniferous age.

Penner (1936) has recently described cases where replacement of plagioclase by orthoclase is actually being effected at the present day. Dacites are being transformed into rhyolites by geyser emanations in Yellowstone Park. Bore-hole evidence shows it to be a dynamic process in the course of which soda, displaced by potash in depth, is redeposited at higher levels, whilst steam alone escapes at the surface.

2). Mineral Assemblages.

The production of the final assemblage of minerals (albite, chlorite, epidote, quartz and calcite) is of such widespread occurrence as to suggest that it represents a stable, low-temperature mineral-facies. It is highly significant that this assemblage is frequently encountered in schists as well as in igneous rocks.

Eskola (1925) has listed characteristic series of assemblages in certain Fennoscandian schists. The final changes under low-temperature metamorphism involves the transformations:

Albite - Epidote - Actinolite,

↓

Albite - Epidote - Chlorite - Calcite.

In the schists of central and western Otago, New Zealand, Turner (1935) has observed the following change:

Augite - Hornblende - Plagioclase,

↓

Actinolite - Albite - Epidote - Chlorite.
He concludes that such transformations are only possible at low grades in the presence of sufficient water and, when calcite occurs, upon the presence of an adequate concentration of carbon dioxide to combine with the lime set free during the chloritisation of amphiboles. The occurrence in the hornblende pseudomorphs of Type II of actinolite (see pp. 23-24) is of interest in view of the assemblages cited above.

3). Mineral Transformations.

Some idea of the nature of the hydrothermal agencies may be gained from an analysis of the mineral transformations which have taken place in the breakdown of original constituents.

Brammall (1936) has demonstrated that such changes are more realistically shown by an atomic, rather than a molecular scheme.

One of the most characteristic changes of the types is the replacement of a plagioclase (oligoclase-andesine) by albite. This involves the replacement of (Ca Al) by (Na Si). Since only about one-fourth of the original felspar was anorthite and, since sericite, kaolin (?) and, to a smaller extent, chlorite, epidote and calcite result from the mineral transformations, removal of Ca by the incoming of Na is not so complete as might appear at first sight.

The change hornblende→chlorite involves the release of Ca, Si and some Na and requires the addition of (H0) and, possibly some Al. Chloritisation of biotite releases Si, K and possibly Al, and requires (H0). The frequent association
of iron-ores with chlorite suggests that the Fe/Mg ratio may also be altered during the destruction of the original minerals.

Of the other new minerals formed, epidote takes up Ca and Al; sericite K, Al and (HO); and calcite Ca and (CO₃).

\[
\begin{align*}
\text{Chlorite} & \rightarrow \text{Albite} \\
\text{Biotite} & \rightarrow \text{Plagioclase} \\
\text{Hornblende} & \rightarrow \text{Chlorite}
\end{align*}
\]

\[
\begin{align*}
\text{Si} & \rightarrow \text{K} \\
\text{K} & \rightarrow \text{Al} \\
\text{Al} & \rightarrow \text{Ca} \\
\text{Ca} & \rightarrow \text{Si} \\
\text{Si} & \rightarrow \text{Na}
\end{align*}
\]

It appears probable, therefore, that to effect the breakdown of the original constituents and the production of many of the later minerals only HO and CO₃ were required. The compositions of the rocks are, however, all more than normally rich in Na and it is quite evident that for the considerable development of hydrothermal albite and quartz extra Na and Si must have been derived from a source other than that provided by the destruction of early-formed minerals. This aspect will be considered later in relation to the magmatic history of the intrusions (see pp. 65 - 72).

4. Order of Replacement.

From a detailed examination and comparison of the mode of occurrence of the replacing minerals a general order has been established (q.v. under separate descriptions). The following order has been found to hold, more or less consist-
Diagram illustrating Petrogenetic Sequence, with particular reference to Types III and V.
<table>
<thead>
<tr>
<th>Procession</th>
<th>Hydothermal</th>
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<tbody>
<tr>
<td>Nodules</td>
<td></td>
</tr>
<tr>
<td>Sulfides</td>
<td></td>
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<tr>
<td>Anhydrite</td>
<td></td>
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<tr>
<td>Chert</td>
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</tbody>
</table>

| Epidote   |            |
| Sulfate   |            |
| Quartz    |            |
| Calcite   |            |

| Apaite    |            |
| Epidote   | Ores       |
ently, throughout (a) individual dykes; (b) all dykes of one type; and (c) all the different types which show hydrothermal alteration:

(i) Chloritisation of hornblende and biotite (epidotisation);
(ii) Albitisation (epidote);
(iii) Second period of chloritisation (probably continuous with the first);
(iv) Development of quartz; nests with albite;
(v) Calcite replacements.

Considerable overlapping of the successive stages has undoubtedly occurred. Thus albite entering into felspar replacements is earlier than that occurring in nests with quartz. Similarly chloritisation commenced by pseudomorphing the original mafic minerals and then proceeded to develop more irregularly. The hydrothermal sequence is represented diagramatically in Plate IV.

5). The Spilite-Keratophyre Problem.

A close similarity exists between these rocks and those of the more acid members of the spilite suite. The following characters are common to both:

(i) A high content of soda;
(ii) The occurrence of albite as the dominant felspar, often showing clear signs of a late replacement origin;
(iii) Chlorite as dominant mafic mineral;
(iv) Abundance of epidote, associated with both chlorite and albite;
(v) Abundance of calcite.
There has been, and is still, much controversy as to the source and nature of the albitising agents.

Eskola (1925) and more recently Gilluly (1932) have summarised the conflicting opinions relating to this problem, the main features of which will be indicated here.

Bailey and Grabham (1909) considered that albitisation was the work of residual solutions, that is, a process of auto-metasomatism. They contended that the original magma was abnormally rich in soda, carbon dioxide and water.

Dewey and Flett (1911), working on pillow-lavas, suggested that spilites were a distinct suite, associated with periods of subsidence without considerable folding. They conceived the albite to be often primary in the keratophyric types, but more generally due to "pneumatolitic emanations", rich in water, soda, silica and probably carbon dioxide.

Sundius (1915), from his work on the Kiruna greenstones, contended that alteration was due to "regional metasomatosis"; more recently (1930), he suggested that the spilites should be "distinguished as a special rock type", and maintained that the albite is of primary origin.

Wells (1922 and 1923) discussed the spilites and their associates and considered the "suite" to be established as a petrogenetic concept, particularly as these rocks are not commonly associated with basalts.

Eskola (1925) considered the albitisation as due to late residual fluids, rich in soda and silica, permeating
and replacing the consolidated magma. More recently he has cooperated in an experimental investigation of the "spilite reaction" (Eskola, Vuoristo and Rankana 1935). A generalised equation competent to represent the transformation is:

\[ \text{Na}_x \text{CO}_3 + \text{CaAl}_2\text{Si}_3\text{O}_8 + 4 \text{SiO}_2 \rightarrow \text{CaCO}_3 + 2 \text{NaAlSi}_2\text{O}_6. \]

The problems are, therefore; (a) whether the albite and other characters are primary or due to later replacement; (b) whether the parental magma is distinct from the normal calc-alkali (basaltic) types or not; and (c) whether the chemical and mineral assemblages are essential features determined by the initial composition of the magma. The possibilities may be summarised thus:

1. Albite may be of primary origin;
2. Albite may be due to later replacement, by the action of material from:
   1. an internal source, i.e. from the immediate magma of the rock concerned, with or without extensive internal migration;
   2. an external source, i.e. from a regional deep-seated source.

In the case of Types II and IV there is no trace of an original, more calcic felspar, and all the available evidence suggests that the albite of the phenocrysts and the incipient crystals of the matrix were of primary origin. At the same time clear albite, quartz and chlorite, all of obvious hydrothermal origin, are much in evidence.

The phenomena of Types III and V, on the other hand,
raise the question of the source of the albitising agencies, whether 2 (i) or 2 (ii).

On the triangular diagram (Pl.XIII) points have been plotted to represent the relative proportions of mafic oxides, alkalies and lime for representative spilites, albite-diabases and keratophyres. Points representing various basalts, porphyrites and andesites are added for comparison. Significant fields have been enclosed by lines. A continuous gradation is demonstrable from basalts to spilites and albite-diabases, and from andesites and porphyrites to keratophyres. Similarly spilites and albite-diabases clearly grade into keratophyres. It should be noted, however, that a marked overlap occurs between the fields of the latter groups; this was also disclosed when alkalies and lime were plotted against either silica or the ferromagnesian ratio. Clearly then some confusion has occurred in the discrimination between spilites (and albite-diabases) and keratophyres, and the ranges of the two appear to require more exact definition. From the diagram it is also evident that the sodic character of the spilite-keratophyre rock group is both marked and fundamental.

Gilluly (1932) has criticised the method adopted by Sundius (1930) of selecting analyses to establish a hiatus between spilites and more normal basalts. It is difficult to understand why so much emphasis should be laid on the presence or absence of such a hiatus as evidence for or against the indepence of the spilites as a group. All gradations exist between boys and men, yet their individuality remains
unquestioned. Whether spilites were produced from normal basalts by albitisation or consolidated from distinct magmas, intermediate types would be expected. In the one case Na-H₂O-CO₂ emanations are added to magmatic material after its consolidation; in the other they were present in the magmatic material before its consolidation. The real problem is the source of these "emanations".

Gilluly (1935) has advanced a number of arguments with which an autolytic explanation has to contend. He casts doubt upon the contention that "although the aggregate magma was more than usually sodic, it did not make the primary felspar as sodic as possible, but retained the alkalies in solution to low temperatures". Surely if this were impossible, zoned plagioclases could never be produced during the course of a crystallising magma.

A more important, but purely theoretical criticism lies in what Gilluly calls "the quantitative difficulty". He demonstrates that the volume of "residual self-contained solutions" required to alter an original more calcic rock, would be out of all proportion to the volume of the original magma, and that, "if entirely indigenous", such solutions "would be expected to be revealed by vugs, comby cavities and other evidences of the volume they occupied while the rock was consolidating". To support this view, he compares the composition of a "pre-spilite" (basalt) with an albite-diabase, and shows that, assuming normal solubilities in an aqueous medium, that the solutions required would occupy 50 per cent
of the volume of the rock. By employing the analysis of a basalt to represent the "pre-spilite" of his calculations, he makes the implicit assumption that crystallisation of the rock would have led to the formation of basaltic minerals before the residual solutions had affected it. Moreover, there is no justification for the explicit assumption that the active agents were aqueous solutions of laboratory type. The existence of residual solutions commences with the beginning of crystallisation and their reaction with the early-formed crystals is not restricted merely to one definite stage towards the end of consolidation of the rock.

As regards the albitic rocks of Kirkcudbrightshire, it has been shown (pp. 63 - 64) that no drastic amounts of introductions in solution would be necessary to alter the original constituents. These, in fact, could readily be altered by the continued solvent and subsequent precipitating action of relatively small quantities of water and carbon dioxide, together with some silica and soda. It is the quartz, albite and calcite of the "nests" and interstices that exhibit the most striking display of hydrothermal crystallisation, and where, moreover, there is every reason to regard the development of these minerals from residual solutions as almost certain.

It has perhaps been too readily assumed by many petrologists that the nature of hydrothermal solutions is entirely different from that of the original magma. A sudden and drastic change, from magmatic to post-magmatic
conditions, is often inherently implied. It is, therefore, especially noteworthy that particularly in the case of the plagiophyres (Type ¥), it is extremely difficult to decide where a distinction is to be drawn between late-magmatic and "post-magmatic" (hydrothermal) activity. (Cf. Krokstrom, 1936).

Against the application to the present rocks of the hypothesis of wide-spread hydrothermal solutions permeating the rocks after consolidation, and derived from some hypothetical source in depth, are ranged the following points:

(i) Apart from occasional calcite veins, which extend for short distances from the dykes, the country rocks are unaffected by hydrothermal agencies, beyond the usual modifications for a few inches outside the dyke-margins.

(ii) Examples of both relatively fresh and intensely altered dykes may occur within a few feet of each other.

(iii) The unaltered dykes invariably have textures quite distinct from those showing hydrothermal alteration. This suggests that the latter are not merely modified varieties of a normal type.

(iv) The larger dykes which have been examined in detail disclose evidence of a definite relationship between distance from the dyke-margin and degree of hydrothermal alteration. This strongly suggests that hydrothermal agencies were inherent in the magma of each individual dyke.

The evidence thus consistently suggests that a process of "autolysis" (auto-metasomatism) was responsible for the characteristic hydrothermal features of the altered types of
intrusives in the Kirkcudbright area.

ACKNOWLEDGMENTS.

It is with the greatest of pleasure that I express my gratitude to Professor Arthur Holmes, by whom it was my privilege to be directed during the course of this investigation, for his kindly and constructive criticism at every stage of the work.

I am also greatly indebted to Miss D.L. Reynolds for her constant encouragement and assistance.

My thanks are also due to Mr. A. Huddleston for his kindness in making two analyses.

I also wish to thank Mr. G. O'Neill who was responsible for photographing my drawings and making a number of thin-sections and Mr. S. H. Morris (of Bedford College) who took the two micro-photographs (Plate VII) and made a number of photographic enlargements.

The rock specimens and thin-sections made in connection with this work are preserved at the Science Department, South Road, Durham.
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## APPENDIX.

### ANALYSES EMPLOYED IN PLATE XIII.

<table>
<thead>
<tr>
<th>Andesites:</th>
<th>Analyst:</th>
<th>Reference:</th>
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<td>2. Andesite, J.S. Grant Wilson</td>
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<tr>
<td>O. R. S. lava, Cheviots.</td>
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| Porphyrites: | | |
| 4. Q-diorite-porphyrite, E.G. Radley | | |
| Pre-Cambrian(?) intr. Leicestershire. | | |

| Basalts: | | Johannsen, A., "The average chemical composition of various rock types". Neuen Jahr-buch f. Mineralogie, etc. |
| 5. Average Basalt | | |
6. Basalt,  
   Tertiary lava,  
   Mull.  

Keratophyres, etc.  
7. Average Keratophyre  
   Pre-Devonian (?) sill,  
   Cornwall.  

8. Keratophyre,  
   W. Pollard Guppy, Eileen M., idem.  
   Pre-Devonian sill,  
   Cornwall.  

9. Keratophyre,  
   E.G. Radley Guppy, Eileen M., idem.  
   Pre-Devonian sill,  
   Cornwall.  

10. Albite-trachyte,  
    E.G. Radley Guppy, Eileen M., idem.  
    Ordovician lava,  
    Skomer Is., Pembroke.  

    Carrock Fell.  
    Vol. IV. 1917*,  

12 Albitophyre,  
   A. Huddleston (Present Work, p. 44).  

13. Tongland,  
    Kirkcudbrightshire.  

Spilites:  
14. Spilite,  
    E.G. Radley Guppy, Eileen M., idem.  
    Pre-Camb. lava,  
    Argyllshire.  

B.B. 64, Abt. A. 1931,  
   p. 511.  

Johannsen, A., idem.  
   p. 511.  

No. 244.  

No. 326.  

No. 327.  

No. 328.  

No. 315.  

No. 315.
15. Spilite, Pre-Camb. (?) lava, Argyllshire.


17. Spilite, Ord. lava, Ayrshire.

18. Spilite, Lava, Aunus.


Albite-Diabases:


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26. Albite-porphyry W. Herdsman (present work p. 44).
Fig. 1. Diorite-porphyrite (Type I) illustrating Plagioclase, Hornblende and Biotite phenocrysts: matrix conventional. (Diameter of field - 2.25 mms.)

Fig. 2. Albite-porphyrite (Type II) illustrating texture of porphyritic variety. (Diameter of field - 2.25 mms.)
Fig. 1. Albite-porphyrite showing outlines of Albites in marginal variety. (Diameter of field - 2.25 mms.)

Fig. 2. A, B and C are Quartz-Albite "nests": A and C show radiating-fibrous Epidote. D is an Epidote (fibrous) and Chlorite pseudomorph after Hornblende. E is a partial pseudomorph in Chlorite showing residual, optically-continuous Hornblende.
Fig. 1. Typical "Myrmekite" in Albite-porphyrite (slide 102). Micro-photograph. (Diameter of field - 0.70 mm.)

Fig. 2. As above: (diameter of field - 0.35 mm.)
Fig. 1. Albitophyre (Type III) showing texture.
(Diameter of field - 2.25 mms.)
Variety (a).

Fig. 2. Albitophyre (Type III) showing texture:
(Diameter of field - 2.25 mms.)
Variety (c).
Fig. 1. Albitophyre; Variety (e) showing texture: colourless areas are of pale green penninite. (Diameter of field - 2.25 mms.)

Fig. 2. Albite-porphyrite (Type 11) from near Argrennan Mains, showing "riddled" felspars. (Diameter of field - 2.25 mms.)
Fig. 1. Chlorite-Albite "Clot":
A: Diameter of field - 2.25 mms.)
B: Diameter of field - 0.40 mms.

Fig. 2. Albite (unshaded) and Epidote replacing Oligoclase-Andesine. Much "sericite" and Chlorite (both as minute flakes) omitted.
(1/10th. inch objective employed).
Fig. 1. A: Minerals of Infilling: (diameter of field - 2.25 mms.)
(a) Albite (near margins of field), with "sheaf" habit;
(b) Chlorite (dark and flaky);
(c) Quartz (unshaded or even grey), with "toothed" habit;
(d) Calcite - final infilling.

B: Albite (with sheaf habit) penetrated by Quartz (unshaded). (Diameter of field - 0.40 mm.)

C: Calcite (unshaded) replacing Quartz (light shading) and Chlorite (fibrous). (Diameter of field - 0.40 mm.)

Fig. 2. A: Albite (unshaded) replacing Oligoclase-andesine: chlorite and epidote also indicated.

B: Quartz (light stippling) replacing Albite and Chlorite (darker shading).

C: Quartz (unshaded) replacing Albite with "myrmekite" texture. Partial pseudomorph of Albite by Quartz shown. Epidote, Chlorite and "Sericite" stippled.

Fig. 3. A: Strongly resorbed crystal of Quartz.

B: Rounded Biotite crystal; margins are chloritised and flakes of chlorite have developed beyond rounded margins of Biotite: Calcite has effected replacement along cleavage-traces.

C: Outline drawing of portion of a xenoporphyriftic aggregate of Plagioclase crystals. (Diameters of fields - 2.25 mms.)
PLATE XI.

Fig. 1.

A.  B.  C.

Fig. 2.

A.  B.  C.

Fig. 3.

A.  B.  C.
Fig. 1. Quartz-plagiophyre (Type V):
Quartz (Qu) and Albite (Ab) are indicated.
(Diameter of field - 2.25 mms.)

Fig. 2. Large basal flake of Biotite, showing slight marginal zoning, and carbonate pseudomorph after olivine. From Lamprophyre (near Nun Mill).
(Diameter of field - 2.25 mms.)
Triangular Composition Diagram illustrating relations of Spilite-Keratophyre suite to corresponding calcic rocks (Basalts, Andesites, etc.,).

12 and 13 are from the analyses of Albitophyres shown on p. 44 (Type III).

26 is from the analysis (p. 44) of the Albite-porphyry (Type IV) at Red Rock.

References to the other analyses employed are given in the Appendix, p. 76.