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GEOLOGICAL REPORT NO. 41

An Experiment in Geobotanical Prospecting for Uranium,
Bokan Mountain Area, Southeastern Alaska

By

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AN EXPERIMENT IN GEOBOTANICAL PROSPECTING FOR URANIUM,
BOKAN MOUNTAIN AREA, SOUTHEASTERN ALASKA

By

Gilbert R. Eakins

A B S T R A C T

A project was conducted at the Ross-Adams uranium mine to determine the usefulness of various sample types in uranium exploration in Alaska, where conditions present unusual difficulties. A total of 421 samples were collected, including several varieties of plants, mulch, algae, and stream sediments. The Ross-Adams ore deposit is a uranium-thorium concentration in a small stock of peralkaline granite. The area was selected for testing various types of sampling because the mine has produced high-grade ore and the geology is known.

Highly anomalous uranium values were obtained from the ashed plants and other materials. The highest value of 2396 parts per million uranium was obtained from lodgepole pine. Eleven assay maps illustrate the results of sampling different plants, mulch, and stream sediments. Other maps show a fracture pattern, mine workings, and radiometric surveys. The analyses indicate that the lodgepole pine is the most suitable plant for geobotanical prospecting in the area. Some evidence suggest that mulch and stream sediments may be useful for uranium prospecting.

I N T R O D U C T I O N

OBJECTIVES

This report is the second by the Division of Mines and Geology directed toward assisting those prospecting for uranium in Alaska (Eakins, 1969). Difficulties in radiometric prospecting for uranium over large parts of Alaska, besides the usual ones of remoteness and rugged terrain, are: the presence of tundra cover, dense vegetation, humid climatic conditions, permafrost, complex geologic structure, and widespread metamorphism of both igneous and sedimentary rocks. These conditions prompted the writer to undertake a bio-geochemical prospecting project in an effort to learn if the method would be useful within this state.

The area selected for sampling was the Bokan Mountain uranium-thorium district near the southern end of Prince of Wales Island. High-grade uranium ore has been produced from an alkali granite stock at the Ross-Adams mine, and radioactivity has been found at numerous sites within the surrounding area. The writer spent the month of August, 1969, collecting samples and examining the geology of the mine and nearby prospects.

Evaluation of the properties and general exploration programs by private companies are currently in progress. Competition is very keen, and therefore it would not be appropriate for the writer to attempt a discussion of ore reserves or the economic possibilities of mining properties at this time.

LOCATION AND ACCESS

The Bokan Mountain uranium-thorium area, as defined by MacKevett (1963), includes a northwest trending tract of roughly 70 square miles located on the east side of the southern part of Prince of Wales Island (*fig 1*). Except for a small strip along the eastern shoreline of the island which falls in the Prince Rupert D-6 quadrangle, the area is in the Dixon Entrance D-1 quadrangle. The most important deposit of uranium found so far is at the Ross-Adams mine on the southeast side of Bokan Mountain at an elevation of 950 feet (*fig 2 & 3*). It is about one and three-fourths miles by road from the head of the West Arm of Kendrick Bay, where Newmont Exploration Limited has a camp and dock. The camp is approximately 40 miles southwest of Ketchikan. It is served by chartered boats and float planes, both of which are available in Ketchikan.

Except for the short road between the bay and the Ross-Adams mine, land travel is by foot and generally very difficult. A few trails leading to some claims are not maintained and not easily followed. Helicopters being used by exploration groups can be landed at rather widely-spaced clearings at the higher elevations.

CLIMATE AND VEGETATION

The climate of southern Prince of Wales Island is not severe, but precipitation is heavy, especially during the spring and summer. Strong wind storms are always a possibility. The U. S. Department of Commerce (1968, p 10) recorded an average annual precipitation of 118 inches during the last 10 years at the Annette Island airport, twenty miles northeast of Kendrick Bay. Temperature extremes at the same location were a low of 1° F and a high of 90° F.

Vegetation varies from very dense in rain forests at the lower elevations to nearly absent along rocky slopes, which range up to 2500 feet at Bokan Mountain. Many of the middle and higher areas where soil is thin support only stunted trees and scrub brush. Swamps and muskeg exist where drainage is poor. Forests contain large spruce, cedar, and hemlock, which often have trunks up to four or five feet in diameter. Fallen trees and dense brush consisting of berry bushes, devils club, and other low-growing varieties make some places nearly impenetrable. Thick mosses cover everything in the well-shaded forests (*fig 4*).

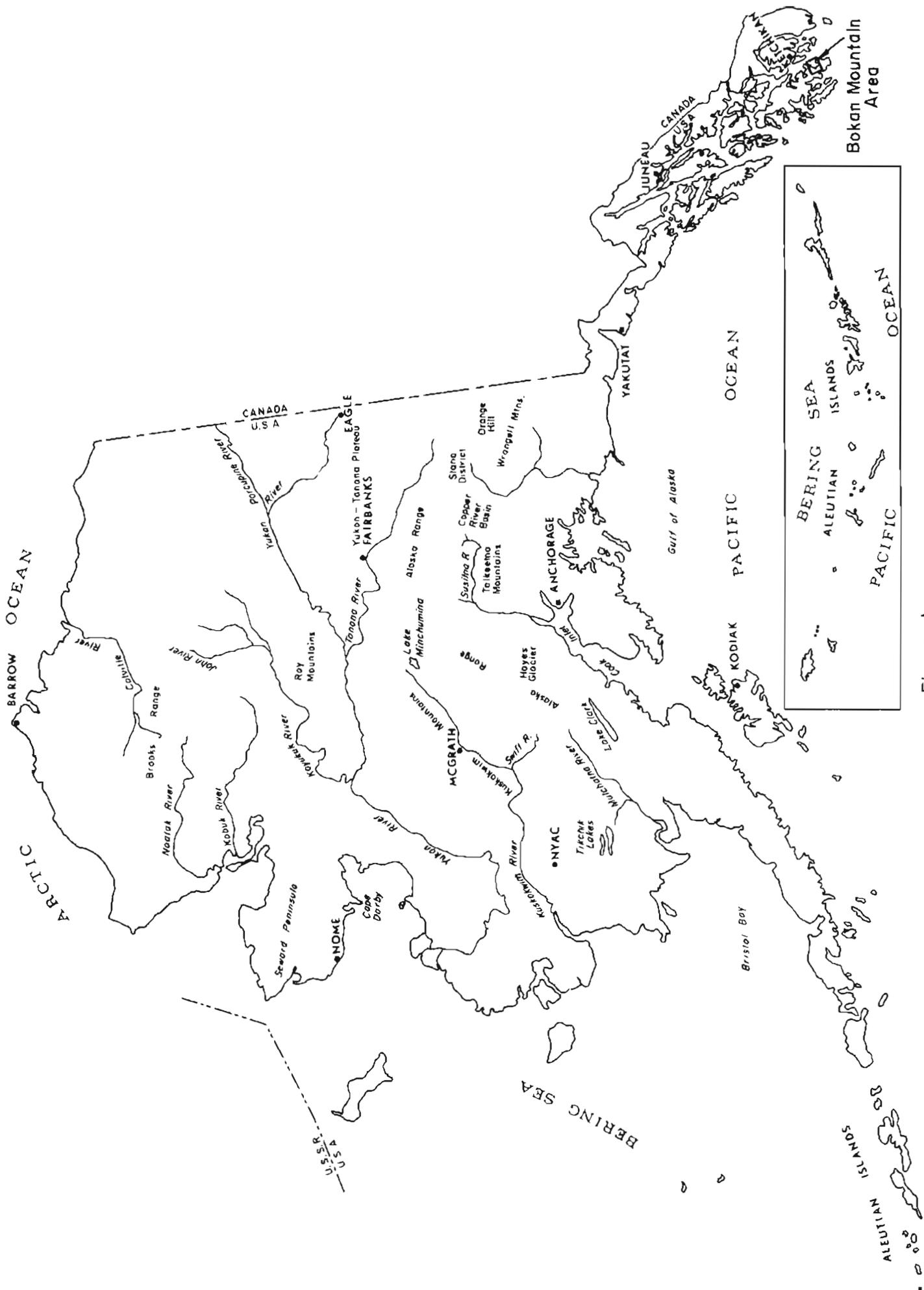


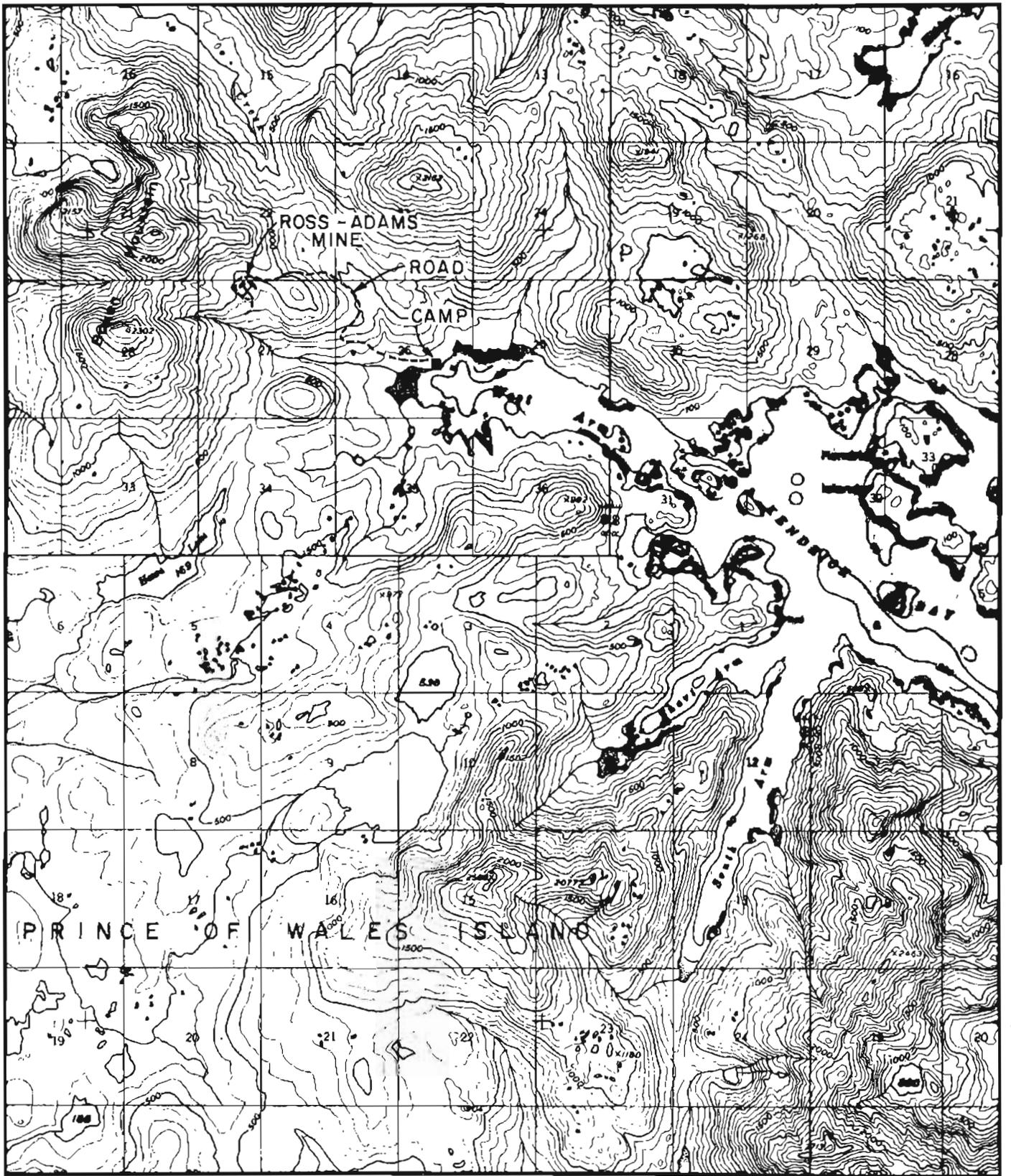
Figure 1

Location Map - Bokan Mountain Area, Alaska

132°10'

132°05'

132°00'



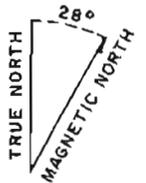
54°55'

54°50'



Figure 2

Bokan Mountain Uranium - Thorium Area



APPROXIMATE MEAN
DECLINATION, 1948



Figure 3-- Bokan Mountain from Kendrick Bay, facing west.
Arrow marks the location of the Ross-Adams mine.

PREVIOUS WORK

Early regional geological reconnaissance was done by Wright and Wright (1908) and Buddington and Chapin (1929). Other publications containing historical information on mining and the geological setting are listed under REFERENCES.

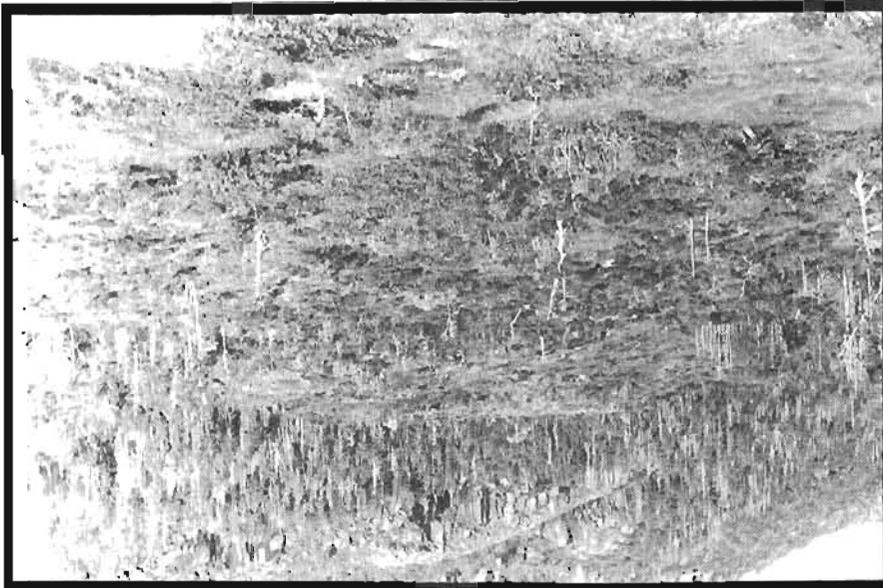
The most complete published report on the Bokan Mountain uranium-thorium area is by MacKevett (1963). It is accompanied by a geological map with a scale of 1:24,000. Much of the following description of geology and mineralogy of the Bokan Mountain area have been drawn from this report. Field work by MacKevett was done during the summers of 1956, 1957, and 1958. Mining at the Ross-Adams mine up through that time had been limited to the shipping of 15,000 tons of ore produced from a trench approximately 400 feet long, 30 to 70 feet wide, and 30 feet deep. Later a 500-foot tunnel was driven at the 700-foot level toward the ore body, and a raise connected the tunnel with the open pit at the 925-foot elevation (*figs 5 & 6*).

Biogeochemical methods used for uranium prospecting in the western states have been developed by the U. S. Geological Survey (Cannon, H. L., 1956, 1957, 1960, 1964, and others). While the writer is not aware of any previous plant analyses for uranium in Alaska, a report on lead and zinc in plants in Southeastern Alaska (Shacklette, 1960) compares metal contents of seven species. Another paper (Chapman and Shacklette, 1960) summarizes the average lead, zinc, copper, nickel, iron, and molybdenum content of several Alaskan plants. Shacklette (1965) has discussed the association of bryophytes with mineral deposits in Alaska. Outside of Alaska, private companies have done extensive work in this field and a number routinely make botanical analyses for uranium in their own laboratories,

Figure 5-- The Ross - Adams mine. View looking south to open pit.



Figure 4-- Typical vegetation of the Ross-Adams mine. View west of mine.



A C K N O W L E D C E M E N T S

The writer wishes to express his gratitude to Newmont Exploration Limited for permission to conduct the biogeochemical test project on their property. He is indebted to Mr. Bill Mounts and Mr. Vivian Vellel of Newmont for their generous assistance and hospitality at the Kendrick Bay camp. Appreciation is due Mr. R. L. Dotson, who holds claims in the area, for his cooperation during the study.

Samples were analyzed by Union Carbide Corporation at their Grand Junction, Colorado laboratory. This service was a great contribution to the project. Thanks are extended to Mr. P. E. Galli, Union Carbide geologist, for his assistance in handling samples and arranging for analyses.

G E N E R A L G E O L O G I C S E T T I N G

Southeastern Alaska is a subprovince of the Western Cordillera tectonic belt. The principal features of the province are the Coast Range Batholith and the outlying subsidiary intrusives. The batholite is complex, consisting of a variety of igneous rocks of various ages.

Marine sediments and volcanics deposited in Paleozoic and Mesozoic euogeosynclines on the western side of the orogenic belt formed thick sequences, totaling possibly 50,000 feet. The oldest dated rocks are Ordovician and Early Silurian (Brew, Loney, and Muffler, 1966).

Deformation and plutonic activity during the Nevadian orogeny have regionally metamorphosed the rocks and created many large-scale faults with lateral movements to more than 50 miles. A conspicuous north-trending linear fault pattern can be seen in many fiords and straits. A number of fault zones contain mineral deposits (Twenhoffel and Sainsbury, 1958). A large vertical fault apparently has controlled the development of the South Arm of Kendrick Bay. Both primary and secondary joints are common. Primary joints, interpreted as being the result of rapid cooling, have localized many of the pegmatite dikes in granite.

Metamorphism on the west side of the Coast Range Batholith has created gneisses which grade westward into the schists, phyllites, and less metamorphosed rocks on the islands of Southeastern Alaska. The strike of most of the bedded rocks and the general grain of the major structures is northeastward.



Figure 6-- Core drilling at the Ross - Adams mine. View southwest of mine.

M I N I N G O F N O N R A D I O A C T I V E M A T E R I A L S

The history of mining in Southeastern Alaska goes back to 1867 when a Russian first mined copper on Prince of Wales Island near new Kasaan. The region contains many old mines and prospects and a wide variety of minerals. The most important deposits are closely related to plutons and structural features of early Mesozoic and early Tertiary age. So far, very few large mineral deposits have been found in the main Coast Range intrusions.

Southeastern Alaska's lode mining has yielded 6,2 million ounces of gold, 3.3 million ounces of silver, 37 million pounds of copper, 48.3 million pounds of lead, 111 thousand pounds of zinc, and 14 thousand pounds of platinum group metals, mainly palladium (Kaufman, Alvin, 1961). Other ores and potential ores are antimony, barite, garnet, iron, molybdenum, titanium, nickel, and tungsten.

In the Bokan Mountain area, copper and gold deposits were staked during the early 1900's at several localities between McLean Arm and Mallard Bay. The Polson and Ichis copper and gold prospects are just south of McLean Arm, nine miles southeast of the Ross-Adams mine. Ores there are localized in quartz-calcite-barite veins in a series of fault zones. Pyrite, chalcopyrite, hematite, chrysocolla, and traces of bornite and gold have been found. Gold was once produced from auriferous pyrite at the Nelson and Tift mine, a mile west of the mouth of McLean Arm. The ore formed a small lens in calcareous rock that has since been mined out.

Iron claims have been staked in the southern part of Prince of Wales Island where magnetite occurs with hornblende concentrations in diorite and pyroxenite.

G E O L O G Y O F T H E B O K A N M O U N T A I N
U R A N I U M - T H O R I U M A R E A

BEDROCK

Radioactive minerals of the Bokan Mountain district are associated with a small peralkaline granite stock of Mesozoic age which has intruded an older monzonite pluton. According to the Shand classification, peralkaline granite is an igneous rock division in which the molecular proportion of alumina is less than that of soda and potash combined. Potassium-argon and lead-alpha measurements indicate the Bokan Mountain granite to be late Jurassic or early Tertiary in age and the older pluton to be early Paleozoic, probably Ordovician (Lanphere, M. A., MacKevett, E. M., and Stern, I. W., 1964). The peralkaline granite stock is roughly circular and about two miles in diameter. This rock is well exposed on the steep slopes of Bokan Mountain.

Slate of Devonian age extends around three sides of the central stock and along the contact between granite and monzonite. At other places the slate is separated from the granite by a strip of monzonite. Paleozoic gneiss, schist, amphibolite, marble, and calcic hornfels lie north and west of the slate. South and east of Bokan Mountain the rocks are predominantly quartz monzonite and quartz diorite, but other plutonic rocks in the area are pyroxenite, gabbro, and syenite. A variety of dikes, largely aplite, pegmatite, and andesite, are widespread. Radioactivity is frequently encountered in the aplite and pegmatite dikes.

Peralkaline Bokan Mountain granite is the rock of primary interest. It is the host for most of the uranium-thorium deposits, and underlies most of the material sampled for this project.

This granite is unusual and has not been reported from any other locality in Southeastern Alaska. It is characteristically high in quartz and sodium-bearing mafic minerals. Many different textures have been noted and grain size varies from fine to coarse. It is generally light gray to white and speckled with about 10 percent dark minerals. Acmite and riebeckite are characteristic, and each is present in amounts from one to twelve percent. Accessory minerals are chiefly zircon, uranothorite, pyrite, xenotime, fluorite, cordierite, and magnetite. Unusual amounts of the minor elements, uranium, thorium, yttrium, lanthanum, niobium, cerium, and other rare earths are present.

Some of the dikes within the area, especially aplite and pegmatite, genetically related to the Bokan Mountain granite are interesting because they contain uranium, thorium, zirconium, and niobium. These dikes are believed to have crystallized from a volatile-rich fluid during a late intrusive stage. Quartz-rich aplite dikes are up to 1000 feet long and 10 feet wide. Pegmatites are less than 4 feet thick, but are traceable for as much as 3000 feet. They are poorly zoned and contain scattered dark uranium or thorium minerals. Other dikes in the area range in composition from diabase to rhyolite and form a complex assemblage.

URANIUM-THORIUM DEPOSITS

Most of the uranium and thorium minerals in the Bokan Mountain district are primary types and are believed to have formed by hydrothermal solutions during Tertiary time. The principal radioactive minerals there are uranothorite, uranoan thorianite, and uraninite. Phosphates, niobates, and complex silicates are present in minor amounts and carry some uranium or thorium.

Four types of deposits in the Bokan Mountain area have been described by MacKevett:

1. Primary segregation of uranium-thorium minerals in a late stage of the peralkaline granite magma and subsequent hydrothermal deposition. This type occurs at the Ross-Adams mine.
2. Syngenetic deposits in pegmatite and aplite dikes.
3. Epigenetic hydrothermal deposits, chiefly open-space filling, but includes some replacement.
4. A deposit formed in clastic sedimentary rock by filling of the interstices at the Cheri No. 1 prospect.

ROSS-ADAMS MINE

The initial uranium discovery in the Bokan Mountain area was made in 1955 by Don Ross and Kelly Adams using an airborne geiger counter. A total of 60,000 tons of ore averaging almost one-percent of both U_3O_8 and thorium has been produced by various operators. The mine was closed in 1964 after fulfilling a contract with the Atomic Energy Commission. An effort to reopen it is pending results of core drilling and ownership negotiations. Newmont Exploration, under an agreement with Standard Metals, is currently evaluating the property, which covers approximately 500 acres. Drilling has proven the extension of ore beyond previously known limits. The internal structure of granite at the mine has been found to be much more complicated than was suspected before Newmont's work.

The ore body is irregular, but plunges generally southward. The percentage of uranium minerals decreases gradually outward from a high-grade zone, and ore limits are indefinite. Two steep intersecting faults striking N70W and N80E are exposed at the south end of the open pit. These faults cut the ore and displace it to the south. On the largest fault, the writer observed slickensides, which indicate a few feet of recent lateral movement. In addition, many smaller faults and joints are present throughout the mine.

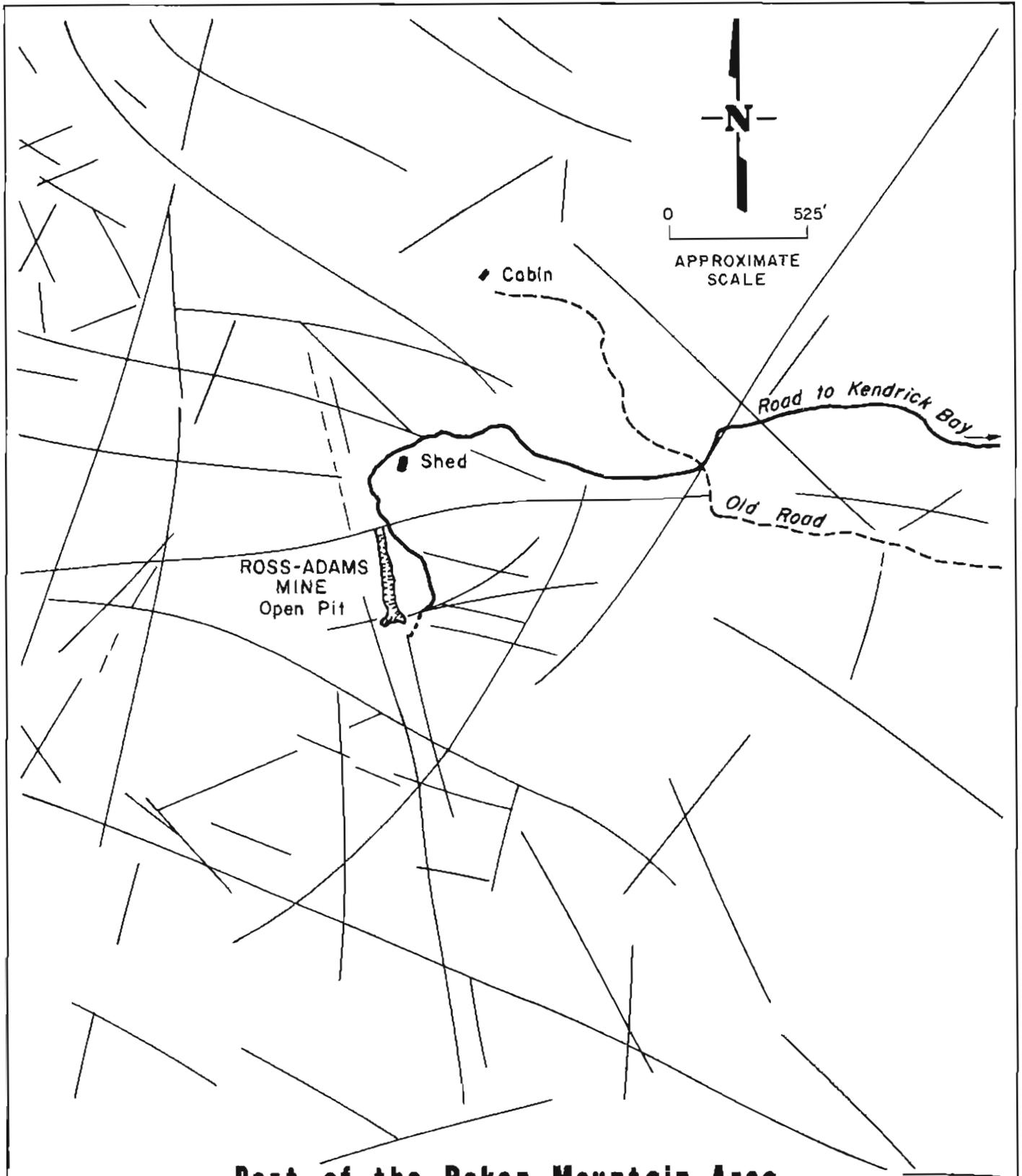
Figure 7 shows lineations evident on aerial photos in the Ross-Adams mine area. These probably represent high-angle joints and faults. Intersections occur that might be loci of crushed zones favorable for mineral deposits. Under the microscope iron-stained fractures can be seen radiating from the ore minerals and from reticulating veinlets along cleavage planes in the feldspars.

The ore body mined in the open pit originally contained a "core" of high-grade ore which averaged over 0.5% U_3O_8 . A large portion contained ore running 1% U_3O_8 , and pods contained up to 3% U_3O_8 . Twelve samples analyzed by the U.S.G.S. yielded from 0.18 to 3.2% chemical uranium. High-grade ore can be distinguished by its dark color due to the presence of associated hematite in the granite. The "core" was enveloped by a zone of lower-grade ore between 2 and 20 feet thick that averaged less than 0.5% U_3O_8 . Information is not available on the ore mined later by underground methods.

Two radiometric surveys by the writer shown in figures 8 & 9 give readings in milliroentgens per hour obtained with a scintillometer in the main open pit and in the short tunnel at the upper level of the mine. Figure 8 shows readings recorded with a geiger counter, both in the main open pit and in the lower tunnel. The geiger counter readings are considerably lower overall than the scintillometer readings, due to differences in instrument sensitivities. The background in both cases averaged about 0.02 milliroentgens per hour at the surface. The highest radioactivity in the entire area, based on geiger counter readings, was at the widest part of the lower tunnel where the short crosscuts connect. This zone is highly broken and sheared.

Mineralogy

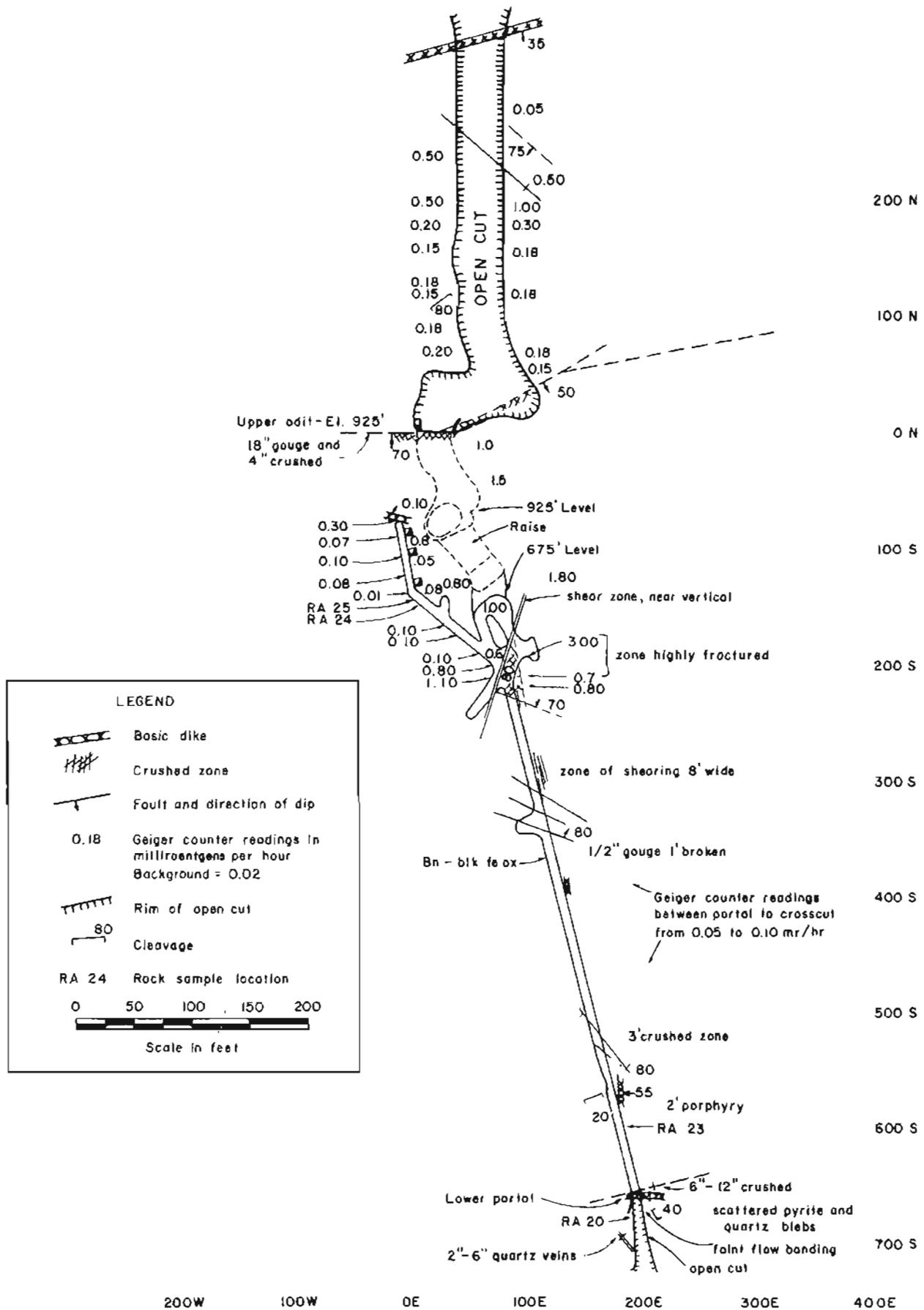
The ore minerals are difficult to identify but almost all are primary. They occur both as scattered grains throughout the peralkaline granite and in numerous thin (0.1 to 0.8 mm) veinlets. Anhedral to euhedral grains up to 2 mm wide are typical. The dominant ore minerals are uranothorite (uranium-bearing thorite) and uranoan thorianite (uranium-bearing thorianite). Coffinite ($U(SiO_4)(OH)_4$) is found in minor amounts. Other vein minerals accompanying the ore minerals are abundant hematite and calcite, and lesser amounts of fluorite, pyrite, hydrous iron sesquioxides, galena, quartz, clay minerals, and chlorite.



**Part of the Bokan Mountain Area
Showing Fracture Pattern**

G. R. Eakins 1969
Data from photo FJB 5-19 (5-10-61)

Figure 7



**SKETCH MAP
ROSS-ADAMS MINE
Geology and Radiometric Survey by G. R. Eakins**

While there is no sharp boundary between the ore and the host granite, the ore contains slightly more iron, lead, aluminum, zirconium, titanium, magnesium, calcium, manganese, and arsenic, but less quartz and potassium, than the surrounding rock. Most of the ore is out of radioactive equilibrium, but the thorium combines with the uranium in such a way to give the effect of apparent equilibrium.

Other uranium-thorium minerals have been found in small amounts at some other prospects in the area. These include uraninite, uranophane, allanite, possibly davidite or brannerite, and ellsworthite. Only minor amounts of secondary uranium minerals have been reported from the Ross-Adams property. These are gummite, sklodowskite, beta-uranophane, bassetite, and novacekite. The scarcity of secondary uranium minerals is undoubtedly due to their solubility and the heavy rainfall in the area.

Ore Genesis

Uranium-thorium mineralization at the Ross-Adams mine is believed to be hydrothermal in origin. It apparently occurred subsequent to the crystallization of the peralkaline granite, and probably was facilitated by a complex fracture system. Tuttle and Bowen (1958) theorized that vapor pressure in crystallizing granite could exceed the overburden limit causing fracturing of the roof with a resulting sudden drop in pressure and rapid crystallization. Such a process might have aided in creating a system of microfractures into which the late stage ore at the Ross-Adams mine was emplaced. Later movements offset the ore body slightly.

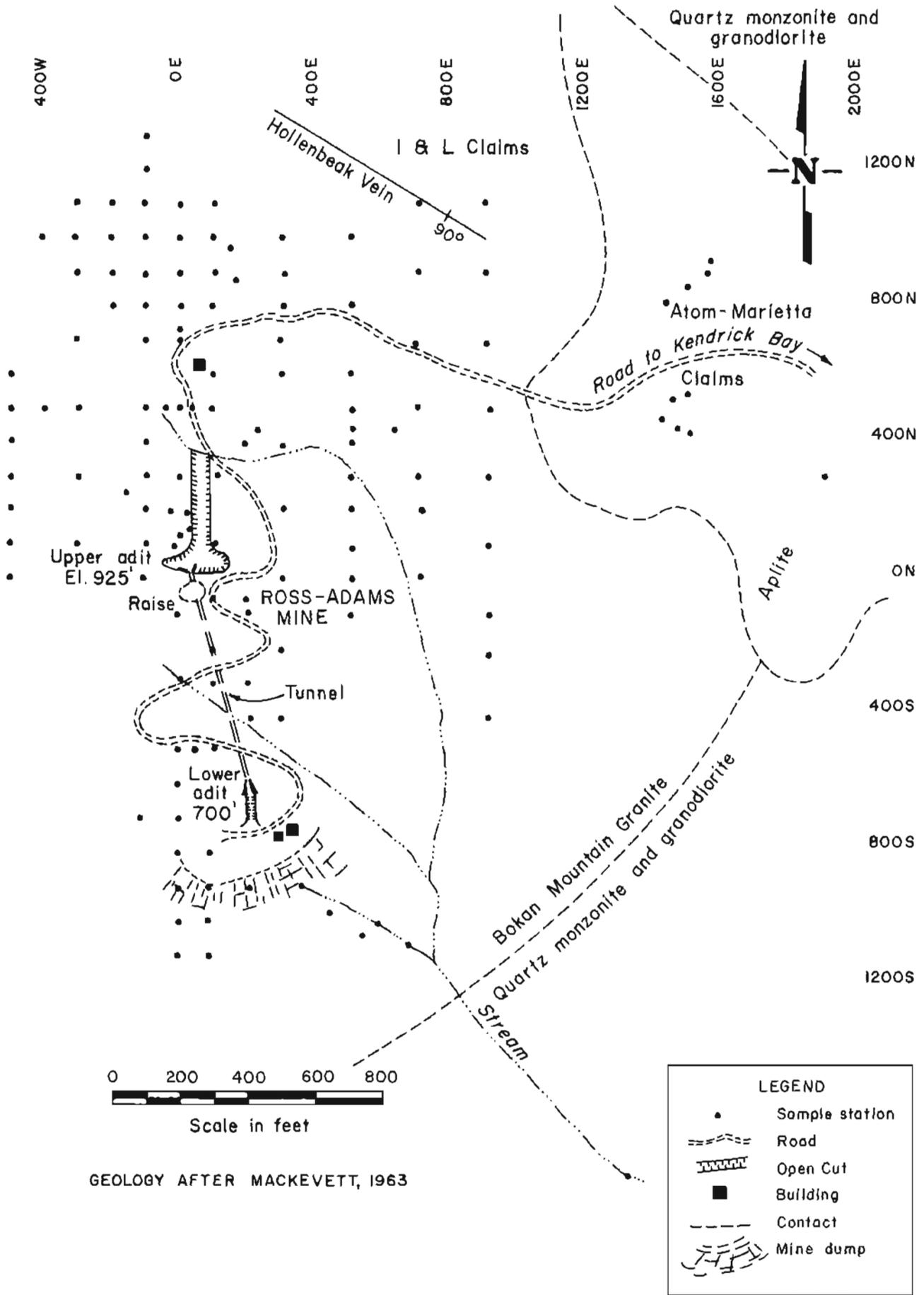
OTHER CLAIMS AND PROSPECTS

Claims and prospects in the area are numerous near altered dacite dikes in albitized zones along the margin of the peralkaline granite. Others are on small pegmatite dikes within the granite. Most claims are only slightly explored. One prospect was located with the intent of mining fluorite.

The Atom-Marietta claims are about 1600 feet east of the Ross-Adams mine (*fig 10*). Shallow pits there have exposed radioactive material in shear zones in the older sedimentary and metamorphic rocks near the contact with the Bokan Mountain Granite. Sample locations 27-30, 59, 60, 61, 65, and 66 are on the Atom-Marietta claims.

Other claims cover weakly radioactive zones scattered in the Bokan Mountain Granite. The "I and L" group of claims adjoin the Ross-Adams mine property. These claims are about 1000 feet north of the Ross-Adams mine at a slightly higher elevation (*fig 10*). Most of the soil and vegetation there has been stripped by bulldozer work and natural erosion. Here radioactive minerals occur in pegmatites and along the northwest trending Hollenbeak vein.

Outside the peralkaline granite stock, weak anomalies occur on prospects near Gardner Bay about eight miles southeast of Bokan Mountain. The anomalies are in pegmatites. Rather weak anomalies have been found in altered andesite dikes cutting syenite near Stone Rock Bay about three miles farther south from Gardner Bay.



GEOLOGY AFTER MACKEVETT, 1963

**Sketch Map of Mine Area
Showing Sample Locations**

B I O G E O C H E M I C A L P R O S P E C T I N G M E T H O D S

Two different botanical prospecting methods have proven effective in locating uranium ores in covered areas where radiometric instruments have failed,

The indicator-plant method of botanical prospecting is based on the principle that certain plant species indicate the presence of abnormal amounts of radioactive elements in soil or ground water associated with weathering mineral deposits. For example, the selenium indicator plants of the Astragalus genus have been used successfully to locate uranium in the Colorado Plateau (Cannon, H. L., 1960). R. L. Dotson, who has been prospecting on Bokan Mountain, has pointed out that the blue flower, lupine, appears to favor the uranium-rich soil in this area. Plant assemblages, or sometimes the lack of them, over known ore deposits compared with assemblages over nonmineralized areas is required for the systematic use of the method.

The biogeochemical method of botanical prospecting involves the chemical analysis of ashed plant material. Deposits of uranium, zinc, tungsten, tin, arsenic, copper, and vanadium have been discovered in this manner. Preliminary plant inventories and analyses are needed to determine which species will best serve to locate particular elements and to establish their normal or background values. The usefulness of plant analyses depends upon finding certain plants of wide distribution which during growth accumulate anomalous amounts of indicator metals. Plant analyses in Grand County, Utah, showed that sulfur, selenium, arsenic, and molybdenum are concentrated with uranium in that area (Cannon, H. L., 1964). A variety of plants, ranging from small bushes and flowers to trees of both conifer and broadleaf types, may be used with varying degrees of success (Cannon, 1960). Lichen and algae also have been found to accumulate uranium in anomalous amounts. A uranium content of 1 part per million or more is frequently considered to be anomalous in the Colorado Plateau region (Froelich, Albert J. and Kleinhampl, 1960, p 59).

Reliable comparison of absolute analytical values can be made only if the samples are collected in a consistent manner and are collected from identical parts of the same plant species. In general, plants with deep roots will serve better than those with shallow roots. Leaves may show greater metal concentrations than the twigs, but both parts can be used. Dead branches of trees in the La Ventana Mesa uranium area, New Mexico, yield more uranium in ash than do live branches (Cannon, H. L. and Starrett, W. H. 1956). Depth of root penetration, amount of moisture in the soil, and depth of the mineral deposit are variables which influence the effectiveness of the method. Fracture patterns, ground water movement, and topography should also be considered when interpreting the results of a botanical sampling program.

Geochemical prospecting for uranium by stream sediment analyses has not been considered very useful because of the high mobility of the U_6 ion (Cohen, Brooks, and Reeves, 1969). However, limited stream sediment sampling by this writer at Bokan Mountain appeared to be effective.

S A M P L I N G P R O C E D U R E

Plants selected for sampling depended largely on their availability. The choice over much of the area was limited to small, stunted evergreens. The species most prevalent were lodgepole pine, western cedar, western hemlock, and common juniper. Trees are larger and denser away from the mine.

Sampling was conducted principally over the Ross-Adams mine property, where the presence of uranium ore is known. Samples were taken mainly on a grid with spacings of 100 or 200 feet (*figs 10 & 12*). Samples of the various plants selected were collected within a radius of 30 feet of the sample station. Spot sampling was done at some of the outlying prospects.

The principal sample area and grid system extend approximately 2500 feet from north to south and 1600 feet from east to west. Bare bedrock is exposed over much of the area and soil is generally only a few inches thick. In the more heavily wooded parts around the margins of the sampling area, the soil was as much as two to four feet thick. The surface around the mine is hummocky and the slopes fairly gentle. Slopes steepen to 20° in marginal areas. Runoff is rapid during periods of rain, and small temporary streams abound (*fig 11*).

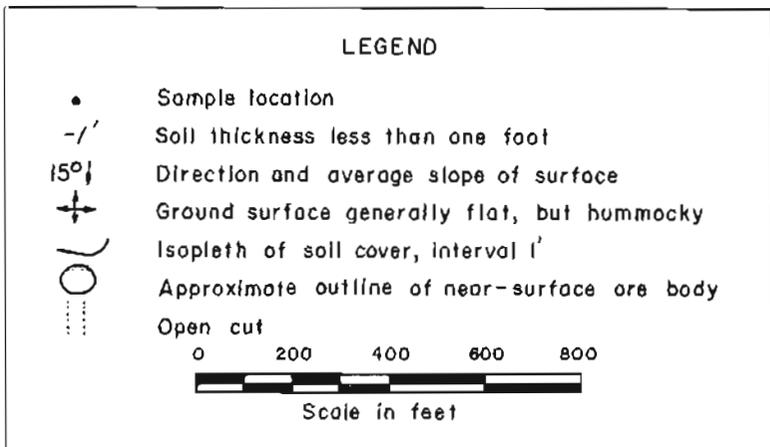
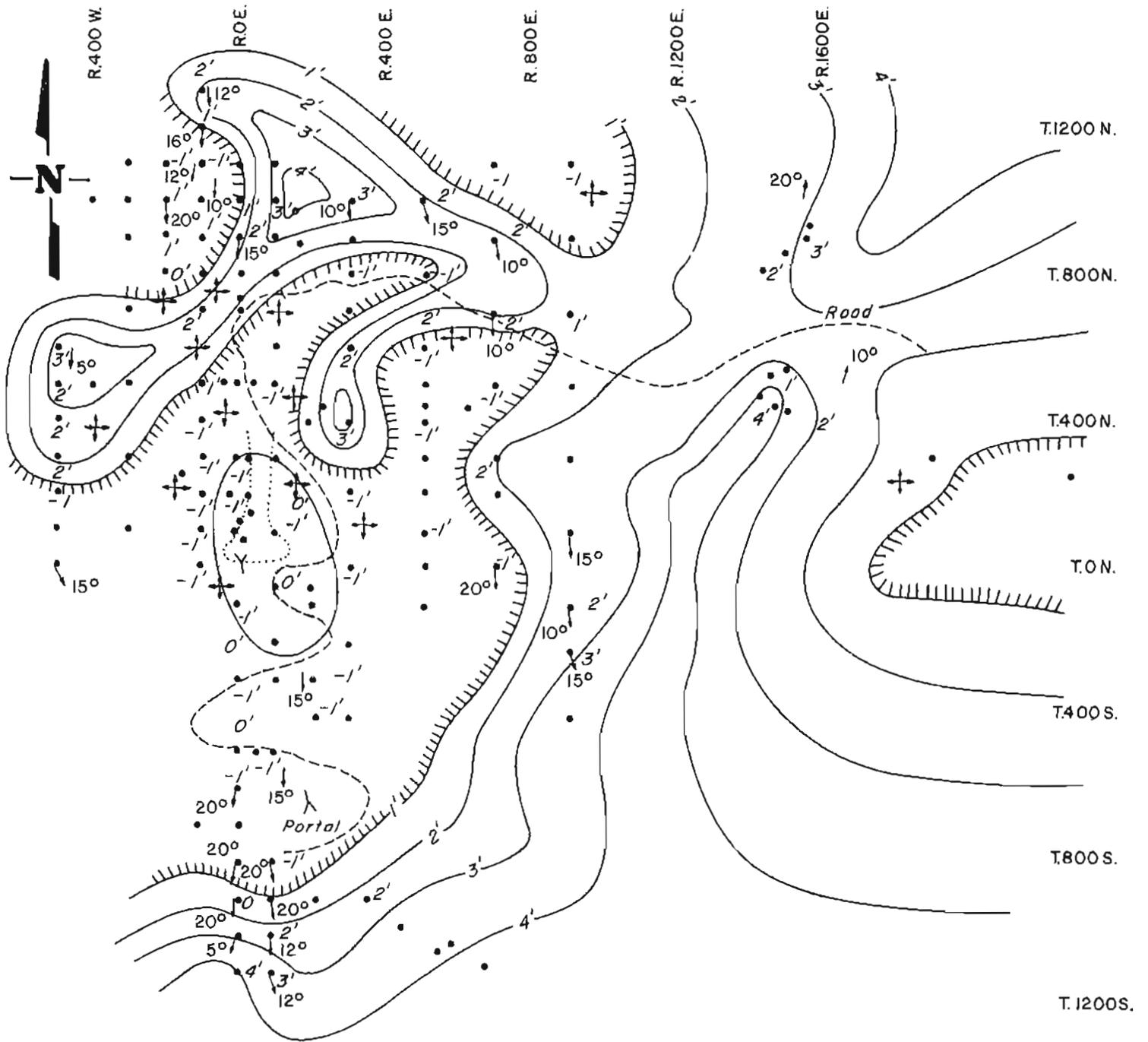
Conditions which make the area less than ideal for test sampling are variable but are generally (1) small thickness of the soil, (2) the shallow root systems of the plants resulting from thin soil and heavy rainfall, (3) the highly variable distribution of the plants, and (4) the possibility of contamination from surface mining operations. However, the choice of areas containing a uranium deposit and known geology was very limited.

The numbers of samples of all types were as follows:

Algae	13
Blueberry	19
Juniper	19
Living lodgepole pine	88
Dead lodgepole pine and burls	22
Miscellaneous plants	13
Mulch and soil	41
Spruce	41
Stream sediment	9
Western cedar	108
Western hemlock	<u>48</u>
TOTAL SAMPLES	421

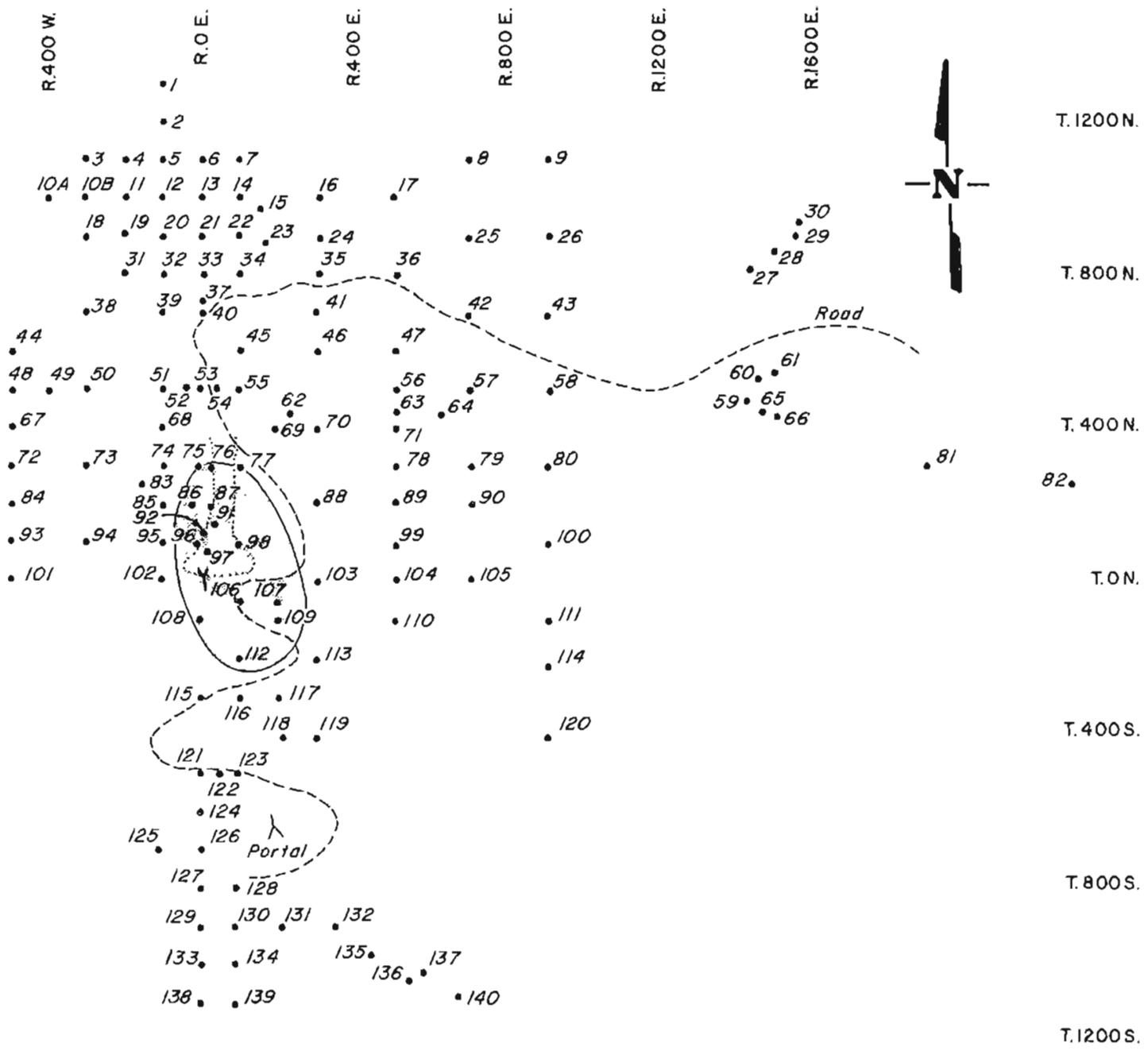
Plant samples were taken by cutting 6 to 10 inches of the branch tips, including leaves, needles and cones, and placing the material in No. 8 paper grocery bags. Because of almost constant rain during the sampling, tearing of bags was a major problem. However, porous bags are required so that samples eventually can dry. Plastic bags are not recommended because plant material will rot if stored in air-tight containers. The weight of the samples ranged from four to eight ounces. Eight ounces or more are preferred in order to have plenty of ash for analyses.

Mulch or humus, which includes accumulated plant material on the ground in various stages of decomposition, has been known to show anomalous metal content over mineralized areas (Ginzburg, 1960, p 233). Therefore, mulch was included in the sampling program. Material was taken from near the surface, just beneath the moss cover. These samples unavoidably contained live moss, grass roots, and soil.



Ross - Adams Mine Area SOIL THICKNESS & SURFACE - SLOPE

Figure 11



LEGEND

- Sample location and number
- Open cut
- Approximate outline of near-surface ore body

0 200 400 600 800

Scale in feet

INDEX MAP

SHOWING ALL SAMPLE LOCATIONS AND NUMBERS

Figure 12

Common blue-green algae can accumulate uranium in flowing streams. Several samples of algae of unidentified types were collected at Bokan Mountain. This plant is difficult to sample, because it is seldom abundant enough to provide an adequate-size sample. It is almost impossible to collect algae without including a certain amount of sediment.

A few moss samples were included to determine to what extent they accumulate metals. While having very shallow roots, mosses are widespread throughout Alaska. Huckleberry, one of the most common bushes in the wooded areas on Bokan Mountain, also was sampled wherever it occurred at a sample station.

A stream crossing the Ross-Adams mine property and flowing to Kendrick Bay was sampled at several points between its mouth and the mine to see how well stream sediment fines would serve for geochemical prospecting for uranium. Stream sediment analysis is not a biogeochemical method.

A N A L Y T I C A L M E T H O D S

Analyses of all the samples collected for this project were made by Union Carbide Exploration Corporation at their laboratory in Grand Junction, Colorado. A summary of the methods used follows:

Uranium:	assayed by the fluorimetric procedure using a flux of sodium and lithium fluoride. Accuracy to 1 ppm* uranium.
Vanadium:	assayed by the hydrogen-peroxide colorimetric procedure. Sensitivity to 1 ppm vanadium.
Molybdenum:	assayed by atomic absorption. Sensitivity to 2 ppm molybdenum.
Arsenic:	assayed by the silver diethyldithiocarbonate colorimetric procedure. Sensitivity to 1 ppm arsenic.
Copper:	assayed by atomic absorption. Sensitivity to 2 ppm.
Manganese and zinc:	assayed by atomic absorption. Because of the high concentrations both were reported in percentages.

* ppm means parts per million

S A M P L I N G R E S U L T S

Analyses of the various plants, mulch, and stream sediment samples are listed in tables 1 through 12. Maps showing the sample locations and analyses for each type of plant and material are included. Extremely high uranium values were obtained from many of the samples collected near the ore body. Average background values cannot be calculated because the laboratory reported that the sample size in many cases was too small for exact determination of uranium below a certain level, generally 3 to 20 ppm. The range of uranium assays, including all samples, is from less than 1 to 2396 ppm. The higher values correspond well with the known near-surface distribution of uranium mineralization. The few scattered samples collected from the Atom-Marietta claims gave rather poor and inconsistent results.

Background or normal uranium content for most of the materials and plants sampled, based on the lower values reported, apparently is less than 3 ppm. The small number of stream sediment samples prevents estimating a background value for this material, but the lowest was 7 ppm.

There is little or no correlation between uranium content and vanadium, molybdenum, arsenic, copper, manganese, and zinc content. These accessory metals are more characteristic of sedimentary-type uranium deposits rather than vein-type or those associated with granitic rocks.

It was not possible to standardize the sampling, because of the variable surface conditions and unequal distribution of plant types. Therefore, it is difficult to accurately compare the relative merits of using one or another plant for geobotanical prospecting in the area. However, lodgepole pine is the most useful of the plants sampled in the area.

All the plant types sampled, with the possible exception of blueberry and alder, evidently can accumulate anomalous amounts of uranium. Also, uranium in the test area is concentrated in humus. Radiometric measurements showed that mulch or humus generally contains more radioactive matter than underlying granite.

Comments on each of the types of materials sampled follow:

Living lodgepole pine (table 1 and fig 13) Lodgepole pine was sampled at 88 locations and produced uranium assays ranging from less than 1 to 2396 ppm. This plant is one of the most abundant and widely distributed in the area, and appears to be the most useful in geobotanical prospecting for uranium there. The better root system no doubt helps account for its relatively high sensitivity to uranium in soil. The pattern of analyses reflects the near-surface mineralization very well.

Dead lodgepole pine (table 2 and fig 14) The 22 samples of twigs and burls from dead lodgepole pine had assays ranging from 7 to 1230 ppm uranium. The average uranium concentration appears to be higher in the dead material than in the living plants, especially those from locations north and up slope from the Ross-Adams mine.

Spruce (table 3 and fig 15) Spruce trees were sampled at 41 locations, and yielded values from less than 2 to 315 ppm. Spruce is less abundant than lodgepole pine and cedar, and does not appear to accumulate uranium as well. However, relatively high values were obtained from the vicinity of the old mine workings.

Western cedar (table 4 and fig 16) Western cedar grows abundantly in the area and was sampled at 108 locations. Uranium assays ranged from less than 1 to 2127 ppm. Background appears to be less than 3 ppm. Cedar assays generally are lower than for lodgepole pine at the same locations and do not define the mineralized areas as well as the pine assays.

Western hemlock (table 5 and fig 17) Western hemlock is not as abundant in the area as cedar and lodgepole pine and was found at only 48 of the sample sites. The assays ranged from less than 2 to 901 ppm. The distribution of significantly high assays corresponds well with the known mineralization, but the number of assays is too small to compare the merits of hemlock with other plants.

Juniper (table 6 and fig 18) Nineteen samples of juniper had uranium values ranging from less than 2 to 159 ppm. While the higher assays from over the ore zone seem to indicate that juniper can serve as an accumulator plant, its shallower root system probably makes it less efficient than larger trees.

Blueberry (table 7 and fig 19) Nineteen samples of blueberry bushes had uranium values ranging from less than 20 to 30 ppm. Compared with other plants, blueberry showed only low uranium values. Though the samples collected were the same size as those of other plants, they proved to be insufficient for accurate ash analyses. All but two samples were simply reported as less than 20 ppm or less than 30 ppm.

Algae (table 8 and fig 20) Thirteen algae samples have uranium values ranging from 2 to 1833 ppm. The highest values were from small streams and pools a few hundred feet north of the open cut, suggesting that this material can accumulate large amounts of uranium even in areas with relatively low uranium in the bedrock. It appears that the algae north of the mine are accumulating uranium derived from the granite upslope rather than an ore body. However, sampling of algae in routine prospecting is not practical because they usually are not available in adequate quantities and carbonaceous or silty material cannot be separated completely from such samples.

Miscellaneous plants (table 9 and fig 21) This group of samples includes the following: 5 alder, 2 club moss, 1 luketkea pectinata, 3 crowberry, and 2 unidentified shrubs. The range in uranium assays was from less than 3 to 923 ppm. These plants were too sparsely distributed to be useful. The uranium content of such low-growing, shallow-root plants seems unusually high, but the high values were from immediately over mineralized ground.

Mulch and soil (table 10 and fig 22) The 41 mulch and soil samples yielded uranium assays from 1 to 441 ppm and show that uranium is concentrated in the humus here. The material sampled was the humic layer just below the moss cover. While it is called mulch, the material unavoidably contained some silt and live moss roots. The ease and convenience of sampling mulch rather than plants seems to make it preferable where conditions are similar to those at Bokan Mountain. Mulch sample coverage of the test area is poor, partly because of the lack of sufficient mulch around the mine. However, analyses show high values which indicate that there has been a buildup of uranium at the surface by the accumulation of decomposed plant material.

Stream sediments (table 11 and fig 23) Nine stream sediment samples were collected from a stream beginning above the Ross-Adams open cut to a point near its entrance into Kendrick Bay, a distance of slightly more than one mile. Uranium values ranged between 7 and 554 ppm and produce a nearly ideal assay profile. Three samples near the upper end of the open cut had values of 7, 8, and 33 ppm. Starting at a point just below the lower tunnel the values in sequence downstream were 554, 253, 291, 198, and 77 ppm. Surface mining and the mine dump have probably caused a considerable increase in the uranium in the sediments above what it would be if the area had not been disturbed. Many more stream samples should be collected and analyzed for uranium to determine their value for uranium prospecting in the region. While stream sediment sampling has generally been considered unsuitable for uranium prospecting because of its solubility, it is much more convenient and rapid than geobotanical prospecting.

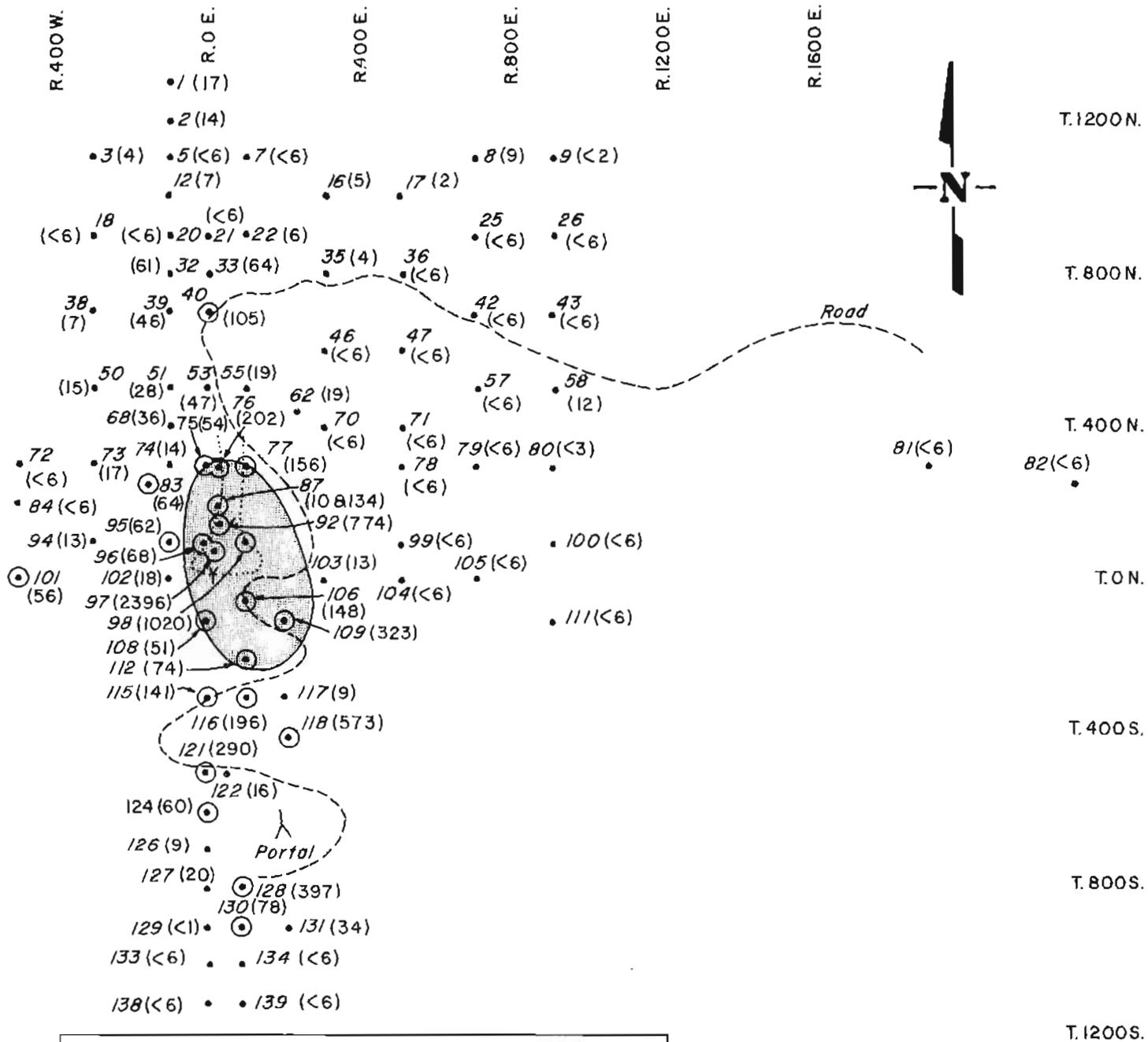
Lichen On the west side of the saddle on the south side of Bokan Mountain at an elevation of about 1600 feet, the writer noted a bright orange lichen growing on granite outcrops. This lichen evidently has acted as a uranium accumulator. Repeated checking with a scintillometer showed the lichen-coated portions of the granite to have two to four times as much radioactivity as the other nearby rock.

The lichen, identified by Dr. Leslie A. Viereck, U. S. Forest Service, as a species of Lecidea (possibly Lecidea Lapidica) may serve as a good indicator plant for uranium. This lichen is known to have the ability to extract iron from rock, and the concentration of uranium by it at Bokan Mountain may illustrate an ability to extract certain ions. The single area where this plant was noted was perhaps 100 feet wide. This limited observation does not permit one to say whether lichen grows preferentially upon high uranium-bearing rock or not, but only that it at least has a tolerance for uranium and can concentrate the metal under proper conditions.

C O N C L U S I O N S

Sampling at the Ross-Adams mine shows that a variety of plants and materials may produce highly anomalous uranium in a uranium mining district, but in varying degrees. Of the plants sampled, the lodgepole pine proved the most useful because of its wide distribution, the relative ease of obtaining an adequate sample, and the plant's sensitivity to uranium in the soil. Mulch overlying the Bokan Mountain granite showed definite uranium concentrations. Measurements with a scintillometer frequently failed to indicate radioactivity. Geobotanical sampling may be especially helpful in defining the outlines and limits of uranium deposits once they have been detected.

While stream sediment sampling produced an almost ideal assay pattern along a stream draining from the mineralized area, the number of samples collected and the area sampled were too limited to determine their value for prospecting. The effects that mining has had on the sample results are not known. The Division plans to continue testing stream sediment sampling and other prospecting techniques to aid in locating the most favorable geologic settings for possible vein-type and sedimentary-type uranium deposits in Alaska.



LEGEND

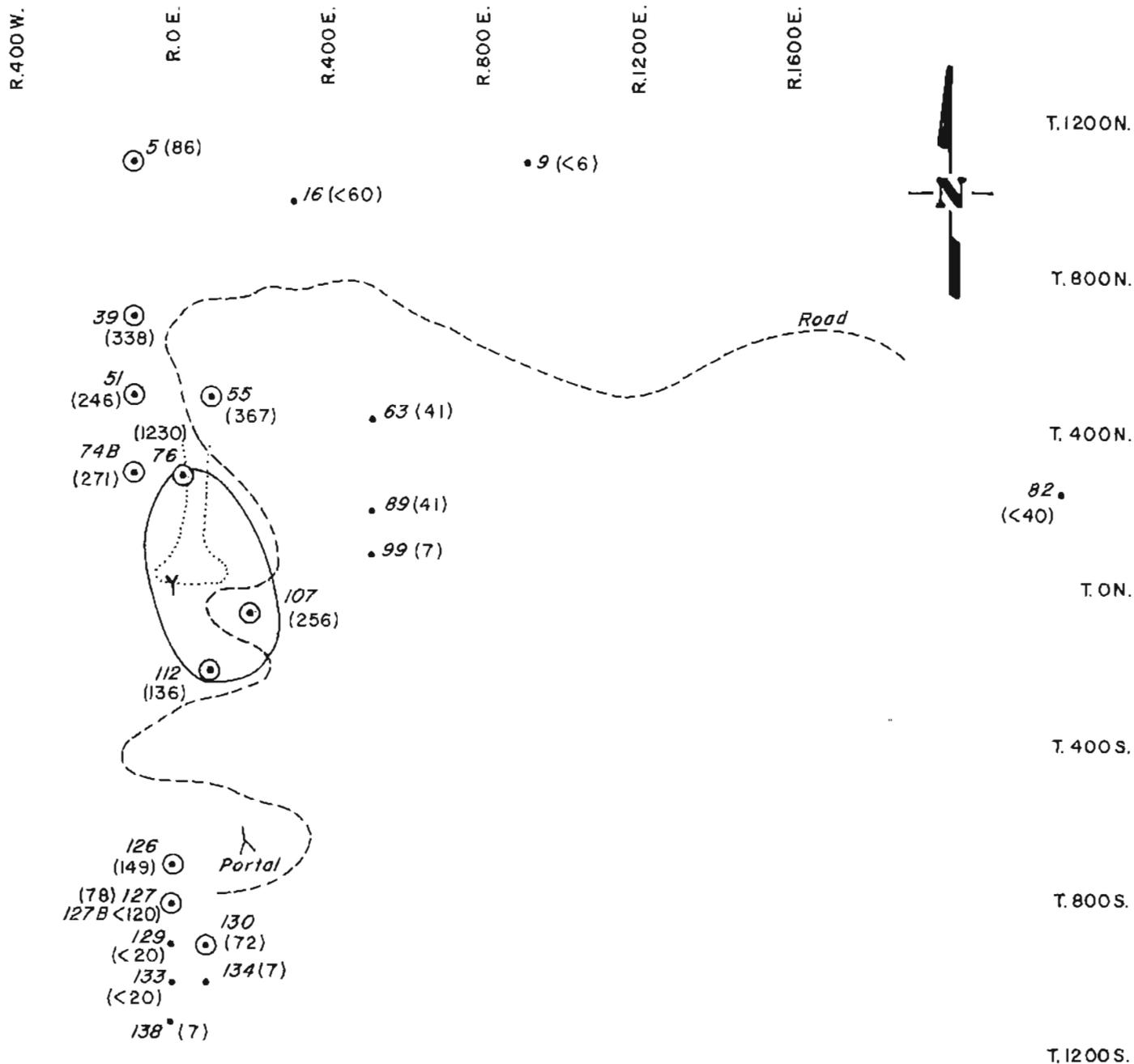
127 • Living Lodgepole Pine sample location and number
 (7) Uranium assay, parts per million
 ⊙ Uranium content over 50 parts per million
 < Signifies less than
 ⋮ Open cut
 ⊙ Approximate outline of near-surface ore body

Number of samples: 88
 Range of uranium values: < 1- 2396 parts per million (Table 3)

0 200 400 600 800
 ───────────────────────────────────
 Scale in feet

Sample Locations LIVING LODGEPOLE PINE

Figure 13



LEGEND

- 127 • Dead Lodgepole Pine sample location and number
- 127B • Burl sample location and number
- (7) Uranium assay, parts per million
- ⊙ Uranium content over 50 parts per million
- < Signifies less than
- ⋯ Open cut
- ⊖ Approximate outline of near-surface ore body

Number of samples: 22

Range of uranium values: 7- 1230 parts per million (Table I)

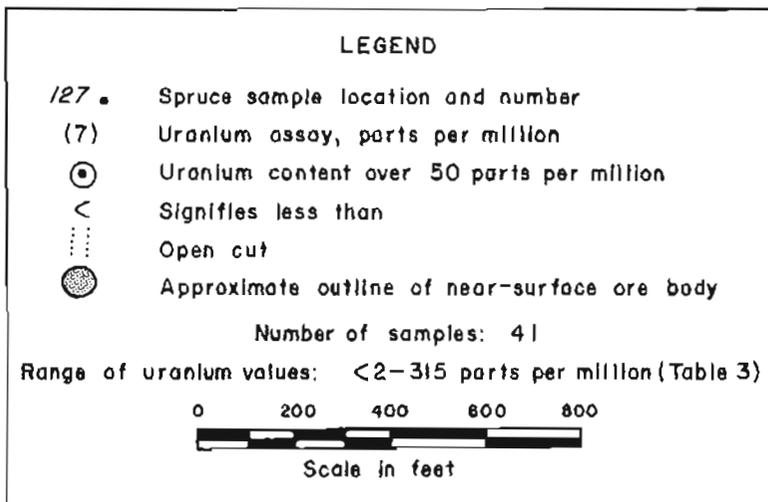
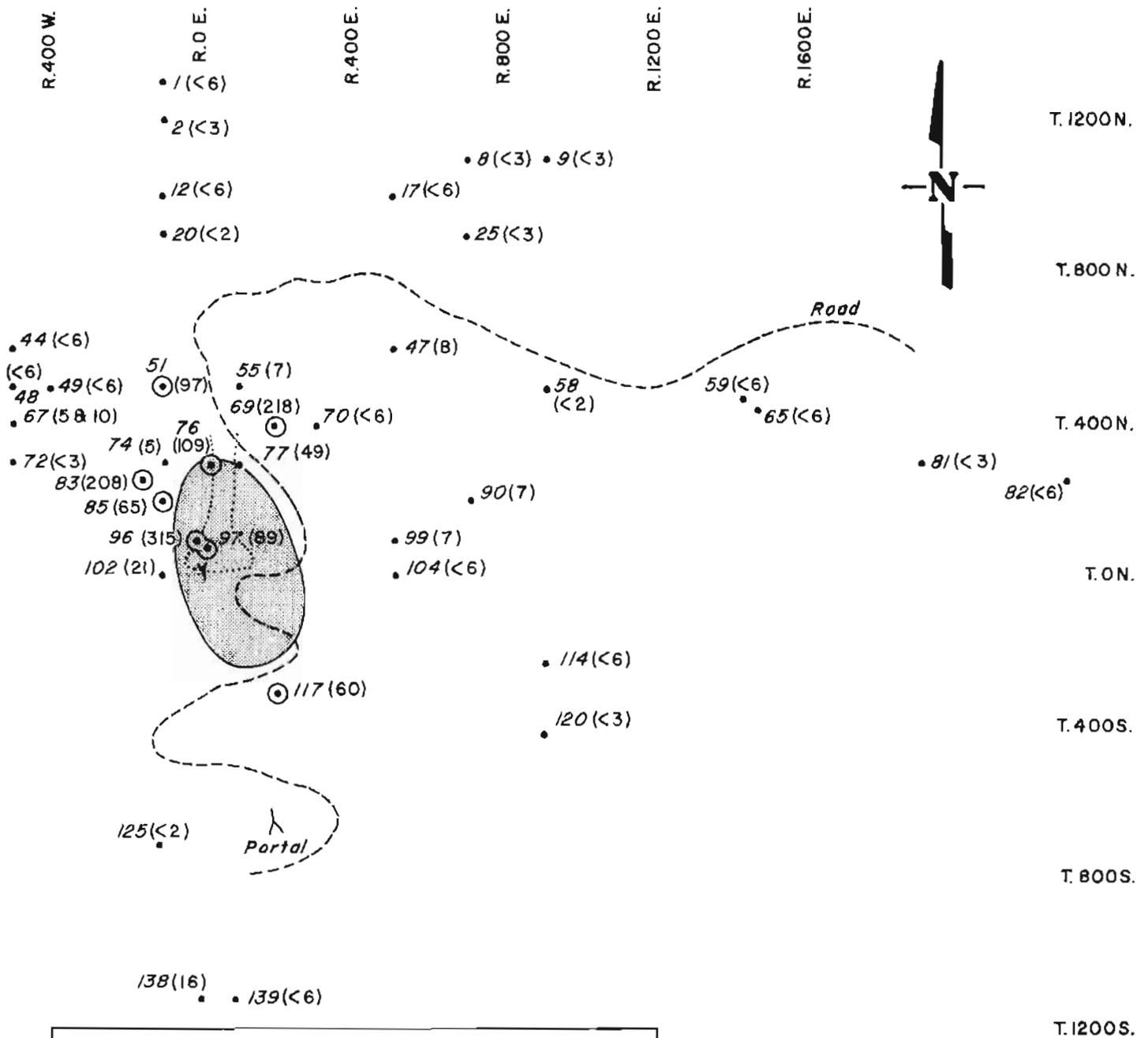
0 200 400 600 800



Scale in feet

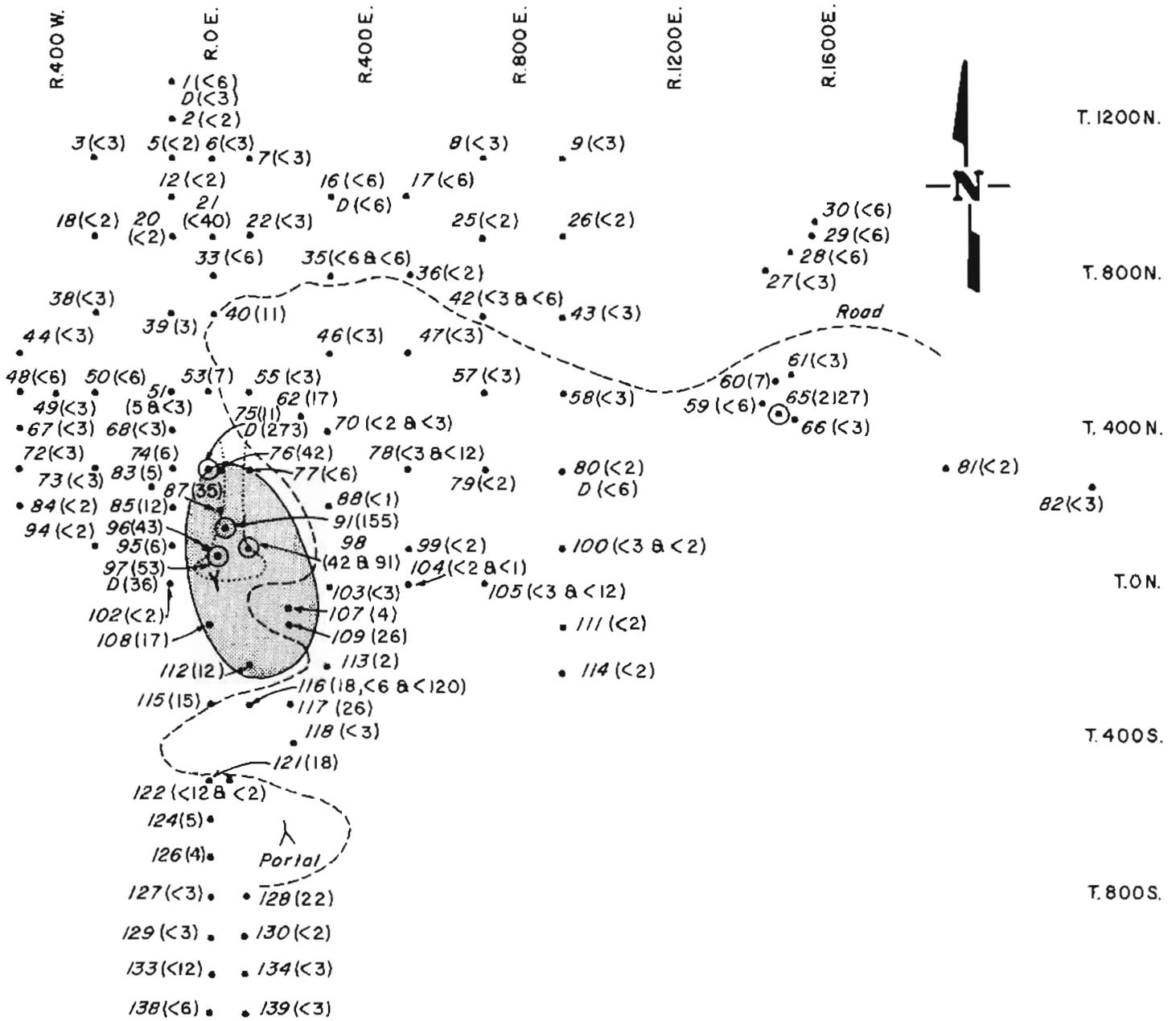
Sample Locations DEAD LODGEPOLE PINE & BURLS

Figure 14



Sample Locations SPRUCE

Figure 15



LEGEND

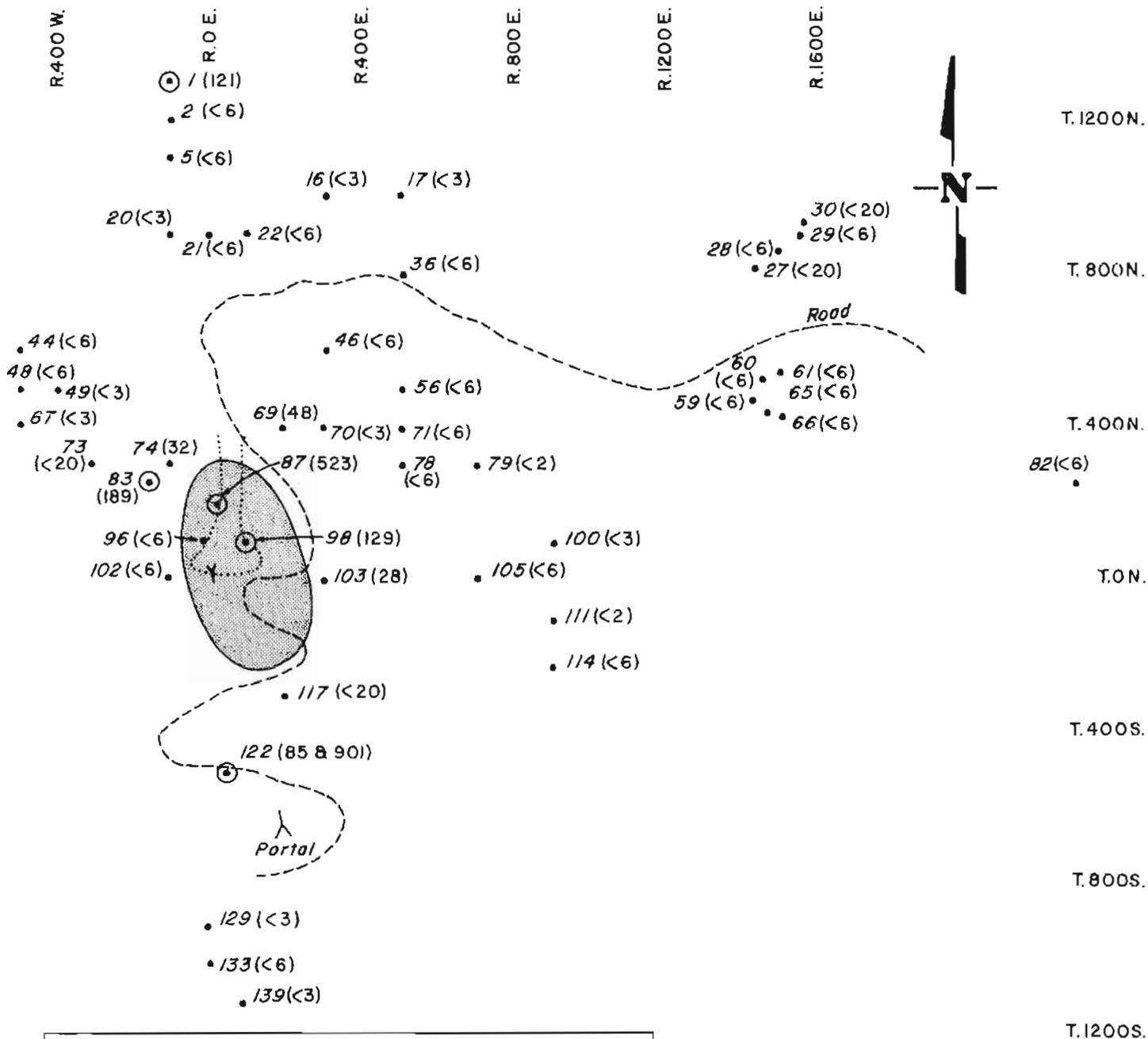
- 127 • Live Western Cedar sample location and number
- 127D • Dead Western Cedar sample location and number
- (7) Uranium assay, parts per million
- ⊙ Uranium content over 50 parts per million
- < Signifies less than
- ⋯ Open cut
- Approximate outline of near-surface ore body

Number of samples: 108
Range of uranium values: <1-2127 parts per million (Table 4)

0 200 400 600 800
Scale in feet

Sample Locations WESTERN CEDAR

Figure 16



LEGEND

- 127 • Western Hemlock sample location and number
- (7) Uranium assay, parts per million
- ⊙ Uranium content over 50 parts per million
- < Signifies less than
- ⋯ Open cut
- ⊙ (shaded) Approximate outline of near-surface ore body

Number of samples: 48

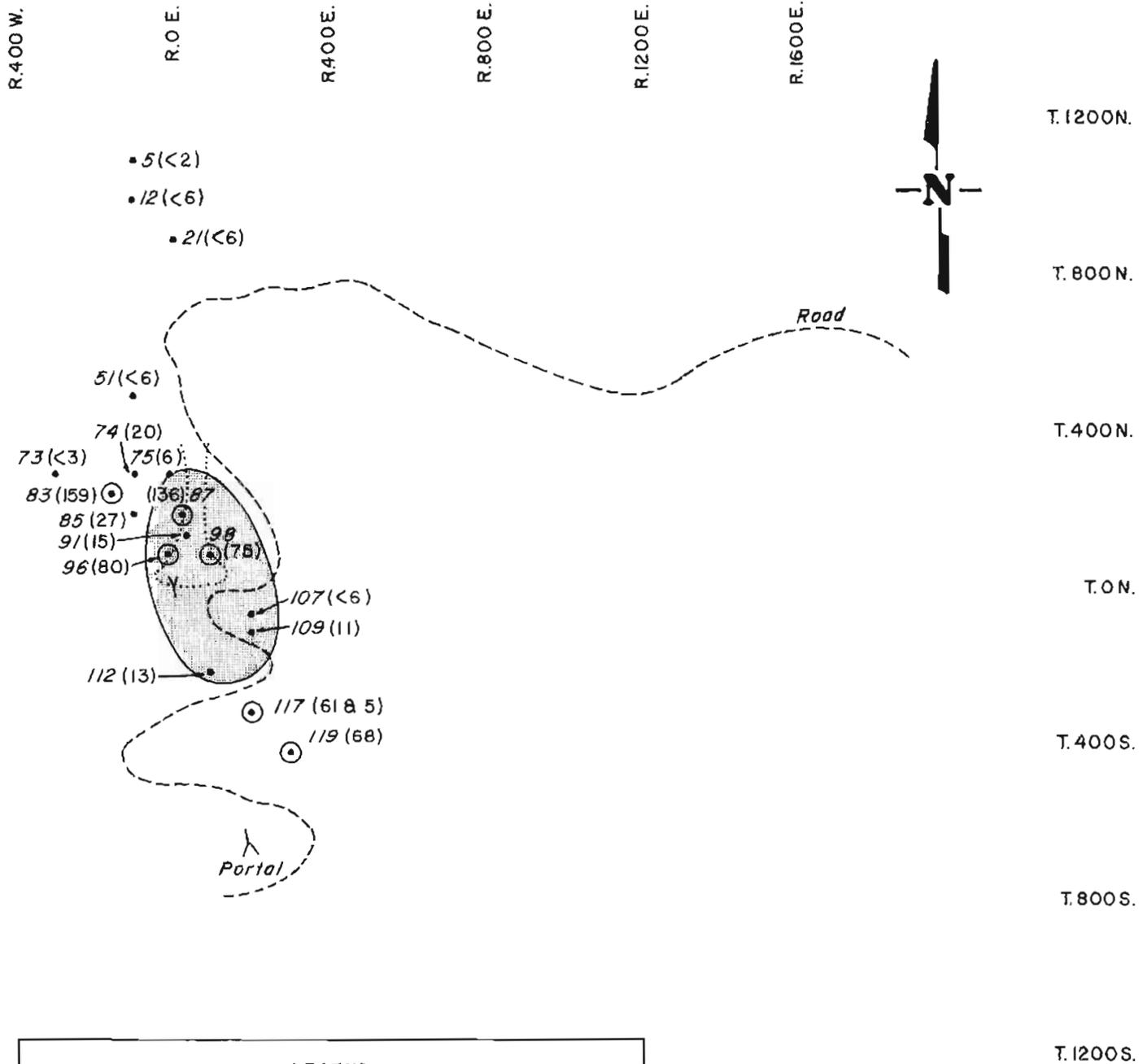
Range of uranium values: <2-901 parts per million (Table 5)

0 200 400 800 800

Scale in feet

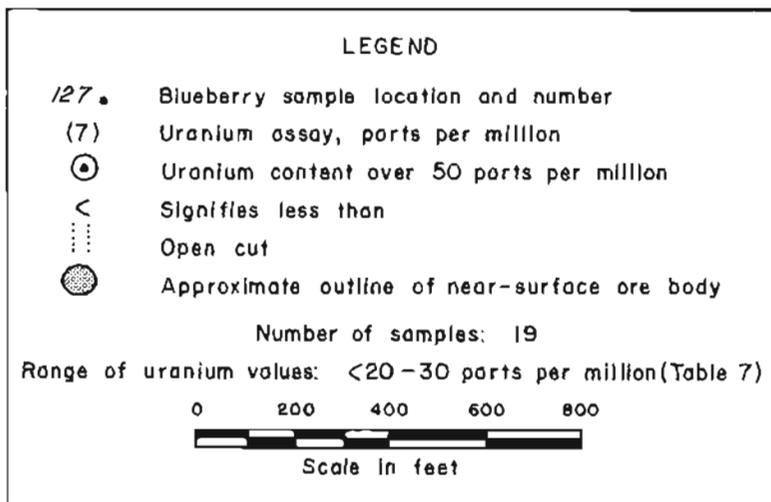
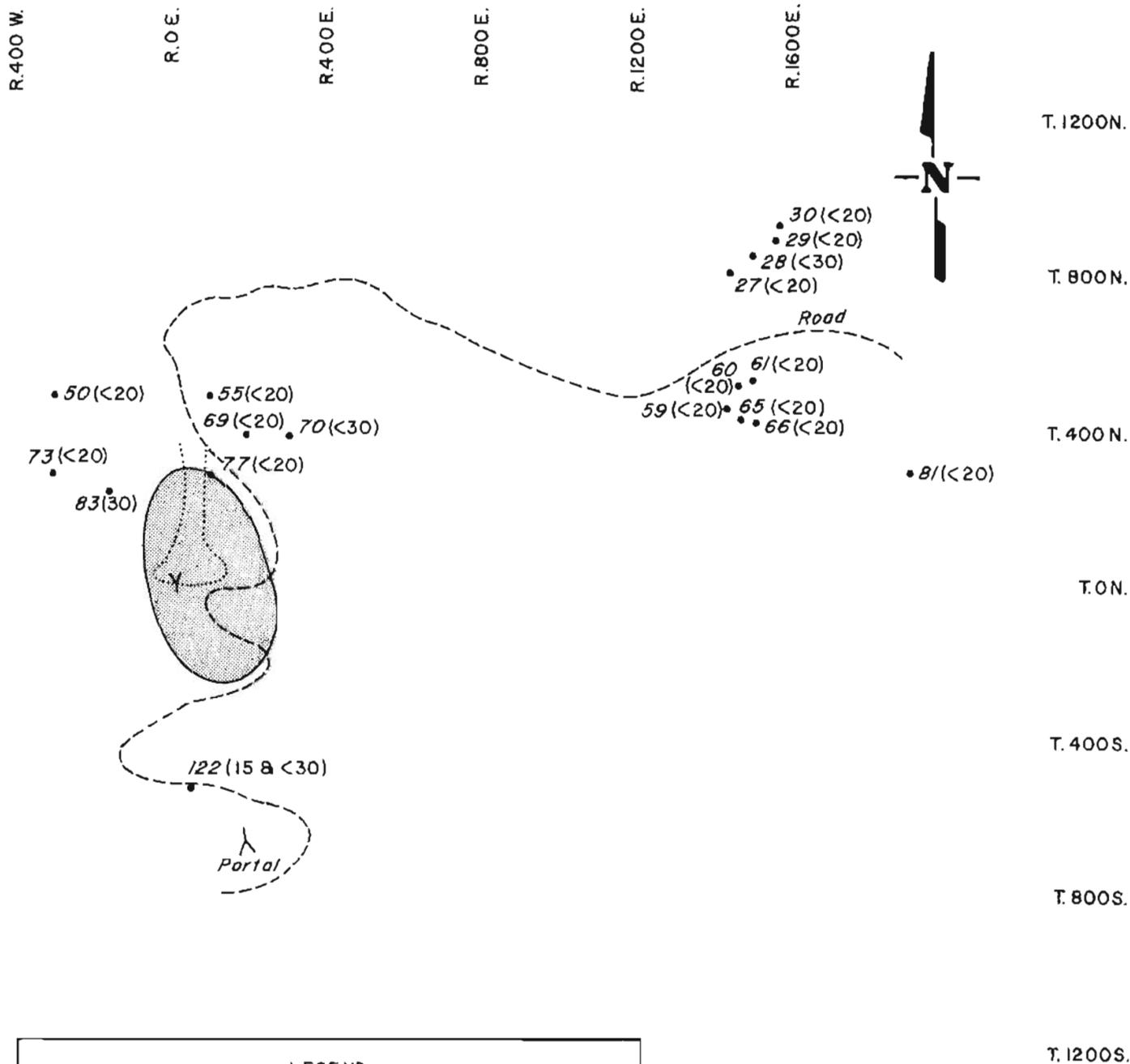
Sample Locations WESTERN HEMLOCK

Figure 17



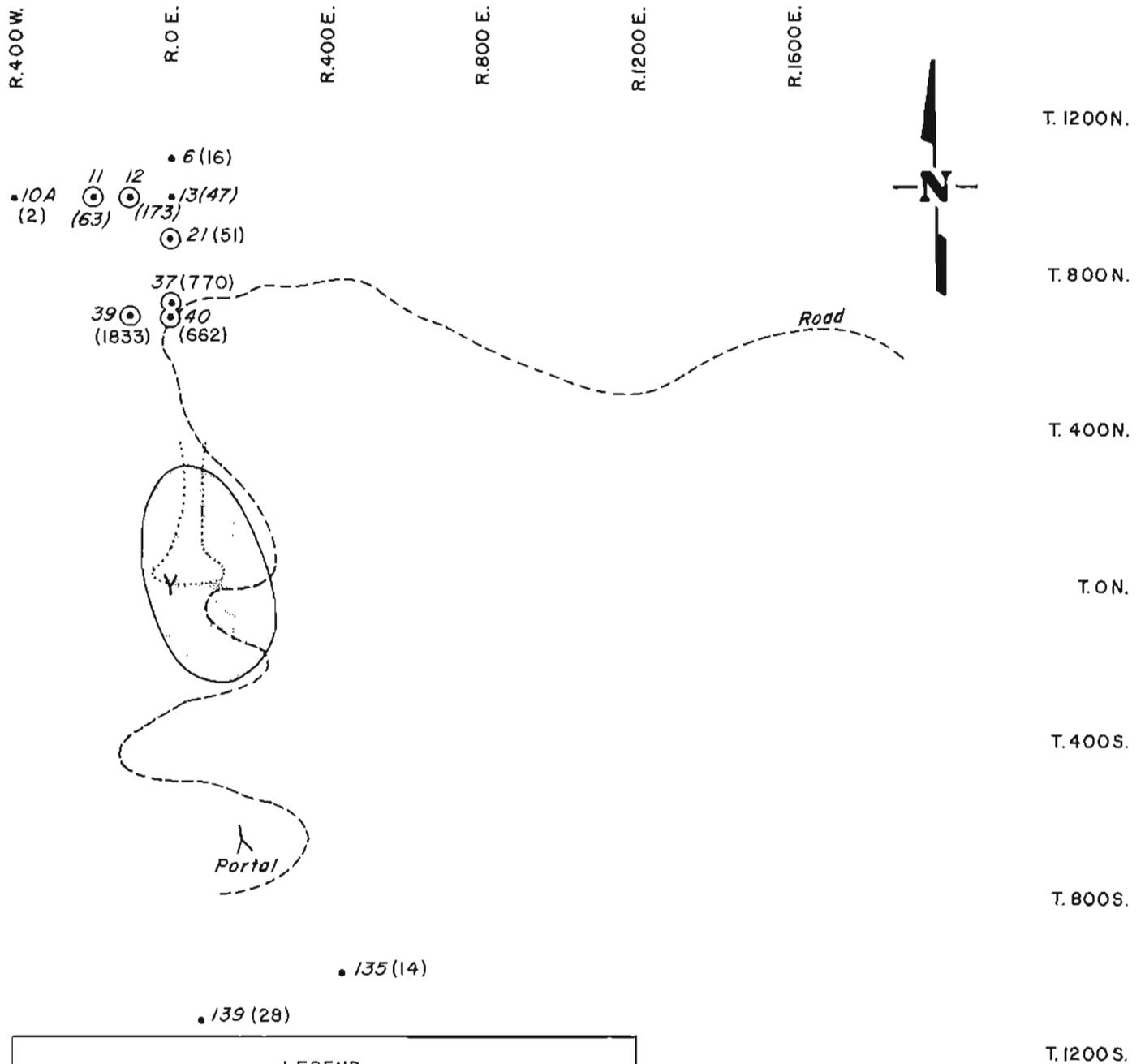
Sample Locations JUNIPER

Figure 18



Sample Locations BLUEBERRY

Figure 19



LEGEND

- 127 • Algae sample location and number
- (7) Uranium assay, parts per million
- ⊙ Uranium content over 50 parts per million
- < Signifies less than
- ⋯ Open cut
- ⊙ (shaded) Approximate outline of near-surface ore body

Number of samples: 13
Range of uranium values: 2-1883 parts per million (Table 8)

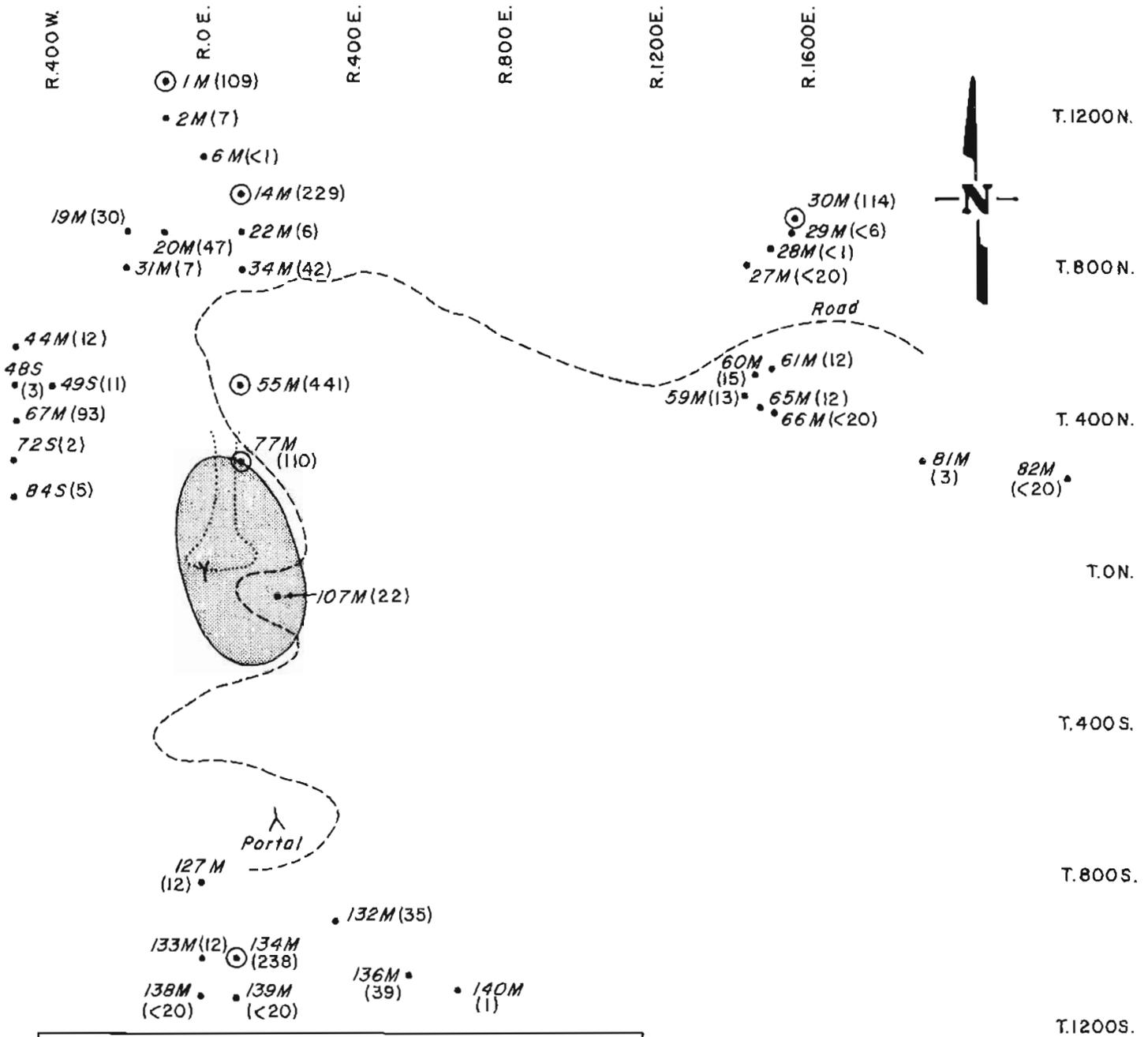
0 200 400 600 800

Scale in feet

NOTE: Algae samples from locations off map:
1. Near camp on Kendrick Bay (5)
2. West end of Kendrick Bay at shoreline (<20)

Sample Locations ALGAE

Figure 20



LEGEND

127M. Mulch sample location and number
 127S. Soil sample location and number
 (7) Uranium assay, parts per million
 ⊙ Uranium content over 50 parts per million
 < Signifies less than
 ⋮ Open cut
 ⊙ Approximate outline of near-surface ore body

Number of samples: 41
 Range of uranium values: 1-441 parts per million (Table 10)

0 200 400 600 800
 Scale in feet

• 141M(<6)

NOTE: Mulch samples from locations off map:

1. (1)
2. (8)
3. (<3)

Sample Locations MULCH AND SOIL

Figure 22

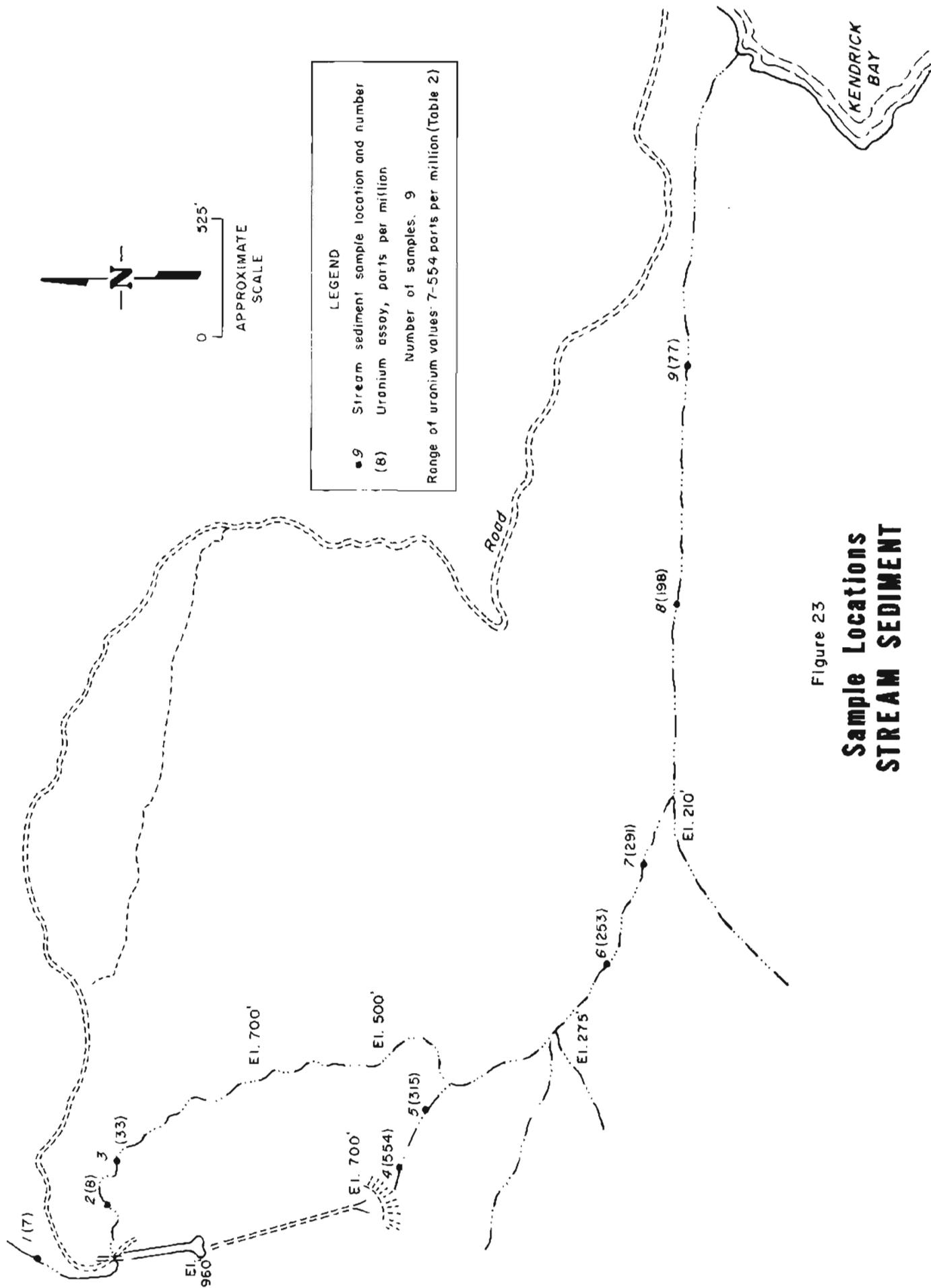


Figure 23

**Sample Locations
STREAM SEDIMENT**

EXPLANATION OF THE TABLES

- indicates "less than"

I.S. indicates insufficient sample for analysis

U = Uranium
 V = Vanadium
 Mo = Molybdenum
 As = Arsenic

Cu = Copper
 Mn = Manganese
 Zn = Zinc

Table 1
 Living Lodgepole Pine
 Ash Analyses

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
1	100W-1300N	43323	17	-150	20	49	-10	5.00	.28
2	100W-1200N	43322	14	-150	10	3	20	4.10	.22
3	U300W-1100N	43330	4	- 75	-10	5	35	3.40	.27
5	100W-1100N	43378	-6	-150	60	4	20	3.10	.44
7	100E-1100N	43394	-6	-150	10	1	10	5.30	.22
8	700E-1100N	43337	9	-150	-10	32	10	1.20	.16
9	900E-1100N	43334	-2	- 50	-10	1	10	0.73	.09
12	100W-1000N	43321	7	-150	40	7	-10	2.80	.260
16	300E-1000N	43325	5	-150	10	34	-10	4.10	.19
17	U500E-1000N	43324	2	- 75	15	3	20	1.50	.19
18	300W-900N	43348	-6	-150	-10	10	2	1.30	.25
20	100W-900N	43376	-6	-150	-10	4	-10	4.20	.14
21	U00W-900N	43361	-6	-150	40	20	15	2.10	.22
22	U100E-900N	43342	6	-150	20	-10	11	2.70	.30
25	700E-900N	43360	-6	-150	-10	-10	19	2.20	.20
26	900E-900N	43347	-6	-150	-10	-10	3	1.30	.35
32	100W-800N	43370	61	-150	-10	4	-10	3.20	.23
33	0W-800N	43379	64	-150	50	1	30	4.70	.54
35	U300E-800N	43326	4	- 75	10	6	30	1.55	.15
36	U500E-800N	43372	-6	-150	-10	4	-10	2.60	.19
38	U300W-700N	43332	7	-150	-10	30	10	1.60	.24
39	100W-700N	43381	46	-150	60	9	20	5.00	.35
40	00W-700N	43335	105	- 75	-10	13	30	1.75	.21
42	700E-700N	43345	-6	-150	10	-10	5	1.00	.35
43	900E-700N	43404	-6	-150	-10	3	10	1.40	.23
46	U300E-600N	43385	-6	-150	-10	2	20	2.60	.25
47	500E-600N	43386	-6	-150	-10	8	30	4.20	.22
50	U300W-500N	43338	15	-150	60	23	10	2.70	.23
51	100W-500N	43390	28	- 75	-10	3	85	5.01	.21
53	0W-500N	43685	47	I.S.	-20	3	100	2.8	.099

Table 1 (Contd)

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
53	OW-500N	43333	47	- 75	60	11	55	0.95	.25
55	100E-500N	43393	19	-150	-10	3	-10	6.40	.27
57	700E-500N	43373	-6	-150	10	4	-10	2.30	.11
58	900E-500N	43340	12	-150	10	26	60	1.60	.38
62	200E-450N	43403	19	-150	10	3	60	1.60	.180
68	100W-400N	43357	36	-150	-10	-10	17	1.70	.19
70	U300E-400N	43344	-6	-150	-10	-10	4	4.00	.16
71	500E-400N	43377	-6	-150	40	1	-10	2.60	.26
72	500W-300N	43354	-6	-150	-10	10	14	2.80	.21
73	U300W-300N	43329	17	-150	-10	4	-10	3.10	.20
74	100W-300N	43405	14	-150	-10	2	30	3.70	.15
75	OW-300N	43355	54	-150	-10	-10	1	2.00	.20
76	25E-300N	43387	202	-150	60	7	10	3.50	.20
77	100E-300N	43362	156	-150	-10	20	21	2.80	.22
78	500E-300N	43365	-6	-150	-10	19	-10	1.50	.18
79	U700E-300N	43375	-6	-150	10	3	-10	4.30	.17
80	900E-300N	43358	-3	- 75	35	25	6	2.80	.12
81	AT-9	43396	-6	-150	-10	3	10	0.40	.15
82	AT-10	43395	-6	-150	20	2	30	2.70	.24
83	150W-250N	43380	64	-150	50	6	50	2.10	.30
84	500W-200N	43397	-6	-150	-10	5	-10	2.00	.16
87	35E-200N	43383	10	-150	40	14	30	3.20	.24
87	OW-200N	43369	134	-150	10	1	20	1.50	.19
92	50E-150N	43389	774	-150	-10	17	40	3.20	.26
94	U300W-100N	43339	13	-150	60	34	10	4.00	.18
95	100W-100N	43320	62	- 75	20	6	40	3.57	.215
96	OE-100N	43363	68	-150	-10	15	20	2.30	.170
97	OW-100N	43428	2396	-100	15	7	15	1.90	.33
98	U100E-100N	43341	1020	-150	-10	30	41	2.10	.19
99	500E-100N	43366	-6	-150	50	15	-10	2.20	.20
100	900E-100N	43374	-6	-150	-10	2	-10	4.20	.16
101	500W-ON	43356	56	-150	30	-10	15	1.80	.15
102	100W-ON	43319	18	- 75	10	6	15	3.20	.185
103	U300E-ON	43331	13	-150	20	11	-10	2.40	.28
104	500E-ON	43359	-6	-150	-10	-10	15	1.20	.11
105	700E-ON	43343	-6	-150	-10	-10	9	3.60	.26
106	100E-50S	43353	148	-150	10	-10	2	3.90	.23
108	OE-100S	43346	51	-150	-10	-10	4	2.20	.23
109	200E-100S	43400	323	- 75	5	2	30	2.95	.20
111	900E-100S	43350	-6	-150	-10	-10	11	3.60	.17

Table 1 (Contd)

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
112	100E-200S	43401	74	- 75	-10	1	45	1.40	.115
115	OE-300S	43368	141	-150	40	21	10	2.70	.25
116	ID100E-300S	43336	196	- 50	-10	11	33	1.63	.13
117	200E-300S	43398	9	- 50	-10	1	20	1.43	.10
118	200E-400S	43382	573	-150	70	15	50	2.30	.29
121	OE-500S	43399	290	- 75	-10	2	30	4.00	.26
122	50E-500S	43327	16	- 75	-10	4	-10	2.65	.15
122	ID50-500S	43328	I N S U F F I C I E N T			S A M P L E			
124	OE-600S	43391	60	-150	-10	5	40	2.60	.23
126	OE-700S	43367	9	-150	20	6	10	1.80	.21
127	OE-800S	43392	20	-150	-10	4	50	1.80	.24
128	100E-800S	43388	397	-150	10	11	40	4.10	.35
129	OE-900S	43384	-1	-150	50	6	30	5.10	.28
130	100E-900S	43364	78	-150	-10	14	10	2.60	.18
131	300E-900S	43406	34	-150	-10	5	10	1.50	.28
133	OE-1000S	43352	-6	-150	-10	-10	2	5.40	.21
134	100E-1000S	43349	-6	-150	-10	-10	4	4.30	.30
138	OE-1100S	43371	-6	-150	-10	3	-10	3.10	.19
139	100E-1100S	43351	-6	-150	-10	-10	6	5.40	.18

Table 2
Dead Lodgepole Pine and Buris

Ash Analyses

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
5	100W-1100N	43421	86	I.S.	-20	2	-20	-.20	.04
9	900E-1100N	43427	-6	I.S.	-20	2	-20	.20	.07
16	300E-1000N	43420	-60	I.S.	-50	5	-50	-.50	.16
39	100W-700N	43412	338	I.S.	-50	7	-50	-.30	.10
51	100W-500N	43416	246	-150	-10	6	-10	.10	.04
55	100E-500N	43413	367	I.S.	-20	9	-20	.40	.07
63	500E-450N	43426	41	I.S.	-10	2	80	.60	.10
74	100W-300N	43408	271	I.S.	-50	4	30	.40	.17
76	25E-300N	43418	1230	-150	-10	2	-10	-.10	.02
82	AT10	43423	-40	I.S.	-30	3	-30	.30	.11
89	500W-300N	43411	41	I.S.	-20	7	-20	.80	.09
99	500E-100N	43410	7	-150	-10	2	-10	.10	.04
107	200E-50S	43414	256	-150	-10	8	80	.40	.06
112	100E-200S	43422	136	-50	-10	1	10	-.10	.01
126	0E-700S	43419	149	-150	-10	1	-10	.10	.03
127	0E-800S	43407	-120	I.S.	-100	14	-100	-1.0	.21
127	0E-800S	43415	78	I.S.	-20	13	-20	-.20	.06
129	0E-900S	43686	-20	-150	-15	11	60	2.33	.077
130	100E-900S	43424	72	I.S.	25	2	10	.30	.06
133	0E-1000S	43409	-20	-275	-20	2	-20	.13	.06
134	100E-1000S	43425	7	-100	-10	1	-10	-.10	.03
138	0E-1100S	43417	7	I.S.	-20	11	-20	.266	.07

Table 3
Spruce
Ash Analyses

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
1	100W-1300N	43537	-6	-150	-10	14	-10	9.7	.02
2	100W-1200N	43536	-3	- 75	- 5	7	- 5	7.8	.01
3	700E-1100N	43545	-3	- 75	20	9	10	3.55	.40
9	900E-1100N	43569	-3	- 75	5	1	- 5	1.20	.01
12	100W-1000N	43544	-6	-150	10	24	-10	7.00	.04
17	U500E-1000N	43538	-6	-150	-10	10	-10	5.8	.03
20	100W-900N	43553	-2	- 50	7	-1	- 3	1.77	.01
25	700E-900N	43566	-3	- 75	20	3	- 5	3.00	.02
44	500W-600N	43562	-6	-150	-10	-1	-10	4.30	.04
47	500E-600N	43539	8	- 75	30	8	- 5	7.65	.03
48	500W-500N	43559	-6	-150	-10	-1	-10	4.30	.03
49	400W-500N	43546	-6	-150	-10	2	-10	3.60	.04
51	100W-500N	43568	97	I.S.	-12	9	-12	4.22	.04
55	100E-500N	43564	7	- 50	17	4	- 3	1.50	.008
58	900E-500N	43552	-2	- 50	10	1	10	2.63	.30
59	AT-4	43570	-6	-150	-10	6	-10	4.80	.03
65	AT-1	43560	-6	-150	-10	7	-10	2.40	.03
67	500W-400N	43561	5	- 75	15	8	- 5	6.10	.03
67	500W-400N	43573	10	I.S.	-20	16	-20	1.35	.11
69	200E-400N	43557	218	I.S.	-12	6	-12	4.78	.04
70	U300E-400N	43549	-6	-150	-10	7	-10	4.90	.04
72	500W-300N	43548	-3	- 75	- 5	12	10	4.90	.35
74	100W-300N	43551	5	- 75	25	5	5	6.85	.02
76	25E-300N	43563	109	I.S.	-15	42	-15	2.25	.10
77	100E-300N	43535	49	-150	-10	13	-10	4.9	.03
81	AT-9	43541	-3	- 75	40	7	15	1.65	.02
82	AT-10	43571	-6	-150	10	13	-10	6.00	.04
83	150W-250N	43543	208	-150	-10	12	-10	4.10	.04
85	100W-200N	43567	65	-150	-10	5	-10	5.30	.05
90	700E-200N	43534	7	-150	-10	17	-10	2.5	.03
96	OW-100N	43572	315	-150	-10	2	-10	7.30	.60
97	OW-100N	43558	89	- 50	16	10	3	1.57	.30
99	U500E-100N	43542	7	- 75	15	7	10	4.65	.06
102	100W-ON	43533	21	-150	-10	14	-10	7.9	.04
104	500E-ON	43547	-6	-150	10	18	-10	4.50	.05
114	900E-200S	43554	-6	-150	-10	60	-10	10.30	.02
117	200E-300S	43550	60	I.S.	-15	8	-15	8.25	.07
120	900E-400S	43565	-3	-150	-10	-1	-10	7.40	.02
125	100W-700S	43556	-2	- 50	- 3	-1	- 3	3.23	.27
138	OE-1100S	43555	16	-150	10	7	-10	4.40	.04
139	100E-1100S	43540	-6	-150	-10	12	-10	7.50	.04

Table 4
Western Cedar
Ash Analyses

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
1	100W-1300N	43479	-6	-150	-10	4	-10	1.4	.03
1	100W-1300N	43530	-3	- 75	- 5	5	- 5	-.1	.01
2	100W-1200N	43463	-2	- 50	- 3	2	- 3	-.1	.01
3	300W-1100N	43467	-3	- 75	5	7	- 5	-.1	.01
5	100W-1100N	43480	-2	- 50	3	1	- 3	-.1	.008
6	U0W-1100N	43474	-3	- 75	- 5	1	- 5	-.1	.01
7	100E-1100N	43518	-3	- 75	5	2	- 5	-.1	.02
8	700E-1100N	43477	-3	- 75	5	- 1	- 5	-.1	.01
9	900E-1100N	43486	-3	- 75	5	5	- 5	-.1	.01
12	100W-1000N	43461	-2	- 37	- 3	2	5	-.1	.008
16	U300E-1000N	43529	-6	-150	-10	13	-10	-.1	.02
16	U300E-1000N	43437	-6	-150	-10	5	-10	.223	.0400
17	U500E-1000N	43491	-6	-150	-10	10	-10	-.1	.02
18	U300W-900N	43441	-2	- 50	20	- 1	- 3	-.1	.01
20	100W-900N	43465	-2	- 37	- 3	1	- 3	-.1	.01
21	U00W-900N	43501	-40	I.S.	-50	9	-50	-.4	.03
22	100E-900N	43502	-3	- 75	- 5	9	- 5	-.1	.01
25	700E-900N	43513	-2	- 50	- 3	2	- 3	-.1	.01
26	900E-900N	43466	-2	- 50	- 3	1	- 3	-.1	.009
27	AT-8	43498	-3	- 75	5	10	- 5	0.3	.01
28	AT-7	43520	-6	-150	-10	3	-10	0.5	.04
29	AT-6	43499	-6	-150	10	17	-10	0.2	.02
30	AT-6	43526	-6	-150	-10	13	-10	-.1	.03
33	OW-800N	43500	-6	-150	-10	32	-10	-.1	.02
35	U300E-800N	43528	-6	-150	-10	14	-10	-.1	.02
35	300E-800N	43483	-3	- 75	- 5	6	- 5	-.1	.02
36	U500E-800N	43489	-2	- 50	10	3	- 3	-.1	.009
38	300W-700N	43485	-3	- 75	20	8	- 5	-.1	.01
39	100W-700N	43492	3	- 75	- 5	4	- 5	-.1	.01
40	OW-700N	43682	11	- 75	75	3	- 5	-.1	.02
42	700E-700N	43472	-3	- 75	5	5	- 3	0.2	.02
42	700E-700N	43673	-6	-150	80	7	-10	0.50	.04
43	900E-700N	43454	-3	- 75	5	5	- 5	-.1	0.12
44	500W-600N	43456	-3	- 75	10	8	5	0.3	.02
46	U300E-600N	43523	-3	- 75	20	3	- 5	-.1	.02
47	U500E-600N	43490	-3	- 75	5	5	- 5	-.1	.01
48	500W-500N	43475	-6	-150	- 5	3	- 5	-.1	.03
49	400W-500N	43444	-3	- 75	20	1	20	.45	.02
50	U300W-500N	43484	-6	I.S.	-10	1	-10	0.6	.03
51	100W-500N	43430	5	- 75	5	1	- 5	-.1	.02

Table 4 (Contd)

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
51	100W-500N	43508	-3	- 75	5	1	- 5	-.1	.02
53	0W-500N	43450	7	- 75	20	1	10	-.1	.02
55	100E-500N	43674	-3	- 75	25	5	5	-.1	.02
57	700E-500N	43482	-3	- 75	5	7	- 5	-.1	.02
57	700E-500N	43476	-3	- 75	-10	2	-10	-.1	.01
58	900E-500N	43469	-3	- 75	- 5	5	- 5	-.1	.01
59	AT-4	43515	-6	-150	-10	5	-10	0.2	.03
60	AT-5	43521	7	-150	-10	16	-10	0.4	.02
61	AT-2	43435	-3	- 75	- 5	3	- 5	-.1	.02
62	200E-450N	43519	17	-150	10	3	-10	0.9	.03
65	AT-1	43683	2127	I.S.	-15	18	13	.005	.06
66	AT-3	43448	-3	- 75	15	2	- 5	.80	.01
67	500W-400N	43429	-3	- 75	- 5	2	- 5	-.1	.02
68	100W-400N	43507	-3	- 75	5	2	- 5	-.1	.01
70	U300E-400N	43436	-2	- 50	4	1	- 3	-.1	.01
70	300E-400N	43497	-3	- 75	- 5	10	- 5	-.1	.01
72	500W-300N	43431	-3	- 75	5	2	- 5	-.1	.01
73	300W-300N	43522	-3	- 75	5	8	- 5	-.1	.01
74	100W-300N	43488	6	- 75	10	4	- 5	-.1	.02
75	0W-300N	43438	11	- 50	10	1	- 3	.0160	.0233
75	0W-300N	43532	273	I.S.	-15	5	-15	-.2	.03
76	25E-300N	43459	42	- 75	20	5	5	0.1	.02
77	100E-300N	43666	-6	I.S.	80	9	-10	-.1	.02
78	U500E-300N	43681	-3	- 75	25	4	10	-.1	.02
78	500E-ON	43667	-12	I.S.	70	8	-10	0.3	.02
79	700E-300N	43670	-3	- 75	35	6	- 5	0.3	.007
80	900E-300N	43503	-2	- 50	- 3	8	- 3	-.1	.009
80	900E-300N	43531	-6	-150	-10	10	-10	-.1	.03
81	AT-9	43517	-2	- 50	13	1	7	-.1	.01
82	AT-10	43495	-3	- 75	20	5	- 5	0.3	.02
83	150W-250N	43512	5	- 75	10	3	- 5	-.1	.02
84	500W-200N	43453	-2	- 37	- 3	1	- 3	-.1	.006
85	100W-200N	43493	12	- 75	10	5	- 5	-.1	.02
87	35E-200N	43457	35	- 50	3	3	- 3	-.1	.01
88	U300E-200N	43433	-1	- 30	13	- 1	- 3	.0140	.0266
91	50E-150N	43458	155	- 50	16	3	3	-.1	.01
94	U300W-100N	43470	-2	- 37	- 3	1	- 3	-.1	.005
95	100W-100N	43487	6	-150	50	13	-10	-.1	.04
96	0W-100N	43527	43	- 75	20	6	5	-.1	.06
97	0E-100N	43442	53	- 50	16	3	7	.17	.02

Table 4 (Contd)

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
97	OE-100N	43684	36	I.S.	45	15	95	1.07	.11
98	100E-100N	43671	42	- 75	40	7	- 5	-.10	.01
98	100E-100N	43473	91	- 75	5	5	- 5	-.1	.02
99	U500E-100N	43524	-2	- 50	- 3	5	- 3	-.1	.006
100	900E-100N	43672	-3	- 75	40	3	- 5	0.35	.006
100	900E-100N	43464	-2	- 50	4	1	- 3	-.1	.01
102	100W-ON	43449	-2	- 50	10	- 1	- 3	-.1	-.01
103	U300E-ON	43455	-3	- 75	- 5	7	- 5	-.1	.01
104	500E-ON	43504	-2	- 37	- 3	5	- 3	-.1	.007
104	500E-ON	43516	-1	- 30	6	10	- 2	-.1	.006
105	U700E-ON	43525	-3	- 75	10	7	- 5	-.1	.02
105	700E-ON	43669	-12	-150	20	12	-10	0.5	.02
107	200E-50S	43514	4	- 75	20	- 1	- 5	-.1	.01
108	OE-100S	43439	17	- 75	- 5	2	- 5	.021	.0250
109	200E-100S	43511	26	- 75	10	- 1	- 5	-.1	.01
111	900E-100S	43446	-2	- 50	10	- 1	- 3	-.1	.01
112	100E-200S	43468	12	- 50	7	1	3	-.1	.01
113	300E-200S	43494	2	- 50	3	3	- 3	-.1	.01
114	900E-200S	43471	-2	- 50	7	1	- 3	-.1	.01
115	OE-300S	43462	15	- 50	7	1	- 3	-.1	.01
116	100E-300S	43434	18	- 75	15	3	- 5	.0240	.0500
116	100E-300S	43665	-6	I.S.	40	17	-10	0.40	.04
116	ID100E-300S	43668	-120	I.S.	95	40	-15	-.7	.02
117	200E-300S	43496	26	- 75	5	9	- 5	-.1	.02
118	200E-400S	43509	-3	- 75	- 5	2	- 5	-.1	.02
121	OE-500S	43478	18	- 75	10	2	- 5	-.1	.02
122	50E-500S	43481	-12	I.S.	-20	2	-20	-.1	.05
122	U50E-500S	43505	-2	- 50	13	2	- 3	-.1	.008
124	OE-600S	43443	5	- 50	10	2	7	-.10	.01
126	OE-700S	43506	4	- 75	5	2	- 5	-.1	.01
127	OE-800S	43432	-3	- 50	10	- 1	- 3	.0103	.0233
128	100E-800S	43452	22	- 50	7	- 1	- 3	-.1	.007
129	OE-900S	43440	-3	- 75	5	6	- 5	-.1	.01
130	100E-900S	43451	-2	- 37	5	1	- 3	-.1	.006
133	OE-1000S	43510	-12	I.S.	- 5	1	- 5	-.1	.03
134	100E-1000S	43447	-3	- 75	35	1	- 5	-.1	.01
138	OE-1100S	43445	-6	-150	-10	2	-10	.50	.01
139	100E-1100S	43460	-3	- 75	10	5	- 5	0.7	.02

Table 5
Western Hemlock
Ash Analyses

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
1	100W-1300N	43605	121	-150	-10	- 1	-10	8.20	.04
2	100W-1200N	43583	-6	-150	-10	4	-10	10.90	.03
5	100W-1100N	43585	-6	-150	-10	8	-10	9.10	.03
16	U300E-1000N	43587	-3	- 75	- 5	7	- 5	7.15	.01
17	U500E-1000N	43581	-3	- 75	20	5	- 5	4.10	.01
20	100W-900N	43593	-3	- 75	35	4	- 5	6.45	.007
21	0W-900N	43575	-6	-150	-10	7	-10	1.70	.02
22	100E-900N	43602	-6	-150	-10	- 1	10	5.70	.04
27	AT-8	43582	-20	I.S.	-13	14	-13	7.67	.03
28	AT-7	43578	-6	-150	20	117	-10	3.90	.02
29	AT-6	43601	-6	-150	10	- 1	40	3.20	.02
30	AT-6	43621	-20	I.S.	13	9	35	4.00	.03
36	U500E-800N	43600	-6	-150	10	- 1	30	3.20	.06
44	500W-600N	43577	-6	-150	10	- 1	-10	5.40	.02
46	U300E-600N	43615	-6	-150	10	3	-10	6.40	.05
48	500W-500N	43616	-6	-150	-10	- 1	10	8.60	.03
49	400W-500N	43592	-3	- 75	- 5	1	- 5	6.40	.01
56	500E-600N	43607	-6	-150	-10	- 1	-10	4.10	.02
59	AT-4	43597	-6	- 15	40	- 1	-10	3.50	.02
60	AT-5	43589	-6	-150	10	- 1	-10	5.20	.02
61	AT-2	43590	-6	-150	-10	- 1	-10	6.00	.02
65	At-1	43599	-6	- 50	40	- 1	10	5.60	.02
66	AT-3	43611	-6	-150	50	10	-10	6.80	.02
67	500W-400N	43613	-3	- 75	- 5	3	- 5	3.85	.01
69	200E-400N	43594	48	-150	90	4	35	9.00	.04
70	U300E-400N	43595	-3	- 75	- 5	- 1	5	8.35	.006
71	500E-400N	43574	-6	-150	-10	113	-10	3.90	.02
73	U300W-300N	43619	-20	I.S.	25	11	45	8.25	.06
74	100W-300N	43603	32	-150	-10	- 1	10	3.70	.05
78	U500E-300N	43580	-6	-150	40	5	-10	5.00	.02
79	U700E-300N	43606	-2	- 50	- 3	- 1	13	5.00	.02
82	AT-10	43579	-6	-150	-10	7	-10	6.90	.02
83	150W-250N	43620	189	-150	23	8	55	4.56	.05
87	35E-200N	43614	523	-150	10	125	40	8.40	.04
96	0E-100N	43604	-6	-150	-10	- 1	-10	6.80	.02
98	100E-100N	43576	129	-150	-10	7	-10	7.90	.03
100	900E-100N	43586	-3	- 75	20	9	10	7.05	.02
102	100W-0N	43588	-6	-150	-10	- 1	-10	6.70	.60
103	U300E-0N	43617	28	-150	-10	2	10	4.20	.08
105	U700E-0N	43610	-6	-150	30	10	-10	8.10	.02

Table 5 (Contd)

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
111	900E-100S	43598	-2	- 50	10	- 1	3	5.97	.008
114	900E-200S	43596	-6	- 15	40	- 1	10	9.80	.02
117	200E-300S	43591	-20	I.S.	-20	- 1	-20	7.83	.06
122	U50E-500S	43618	85	I.S.	-50	30	-50	6.00	.08
122	ID50E-500S	43612	901	-150	-10	9	20	4.70	.04
129	OE-900S	43609	-3	75	15	- 1	20	6.25	.02
133	OE-1000S	43584	-6	-150	30	5	-10	13.80	.04
139	100E-1100S	43608	-3	75	- 5	- 1	5	6.10	.01

Table 6

Juniper

Ash Analyses

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
5	100W-1100N	43637	-2	- 50	7	1	10	0.27	.01
12	100W-1000N	43636	-6	-150	10	14	-10	0.30	.02
21	0W-900N	43627	-6	-150	-100	9	-100	1.3	.03
51	100W-500N	43635	-6	-150	30	27	-10	1.60	.02
73	100W-300N	43633	-3	- 75	15	6	15	0.50	.02
74	100W-100N	43626	20	- 75	5	3	15	0.05	.02
75	25E-300N	43622	6	- 75	10	- 1	- 5	0.80	.01
83	150W-250N	43634	159	-150	10	17	-10	1.20	.02
85	100W-200N	43624	27	-150	-10	18	-10	0.60	.02
87	35E-200N	43625	136	-150	-10	11	10	1.80	.02
91	50E-150N	43628	15	- 75	10	4	10	0.65	.03
96	0W-100N	43631	80	- 75	35	9	25	0.95	.03
98	100E-100N	43640	75	-150	20	13	-10	1.70	.02
107	200E-50S	43632	-6	-150	-10	12	-10	1.10	.02
109	200E-100S	43639	11	- 75	35	8	5	0.70	.02
112	100E-200S	43630	13	- 75	10	3	10	1.00	.02
117	200E-300S	43638	61	- 75	15	8	15	1.00	.04
117	300E-200S	43629	5	- 75	- 5	2	5	0.45	.02
119	300E-400S	43623	68	- 75	5	7	25	1.25	.04

Table 7
Blueberry
Ash Analyses

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
27	AT8	43655	-20	I.S.	-20	4	60	8.20	.05
28	AT7	43657	-30	I.S.	-20	13	20	6.75	.06
29	AT6	43656	-20	I.S.	80	6	40	5.00	.03
30	AT6-B2	43659	-20	I.S.	-20	6	40	2.40	.05
50	U300W-500N	43647	-20	I.S.	14	3	40	4.14	.05
55	100E-500N	43658	-20	I.S.	25	3	-15	9.00	.04
59	AT4	43649	-20	I.S.	-20	2	80	3.60	.06
60	AT5	43645	-20	-150	-10	6	140	5.89	.06
61	AT2	43648	-20	-150	24	2	125	5.78	.05
65	AT-1	43642	-20	I.S.	33	6	80	1.13	.04
66	AT3	43652	-20	I.S.	35	8	55	7.50	.04
69	200E-400N	43651	-20	I.S.	-20	21	60	6.17	.04
70	U300E-400N	43654	-30	I.S.	98	19	-25	7.25	.05
73	U300W-300N	43646	-20	-150	55	5	35	5.44	.04
77	U100E-300N	43644	-20	I.S.	-12	4	12	5.38	.04
81	AT9	43643	-20	I.S.	14	7	90	3.33	.06
83	150W-250N	43641	30	I.S.	-12	11	85	8.67	.05
122	50E-500S	43653	15	I.S.	95	9	75	5.75	.06
122	U50E-500S	43650	-30	I.S.	-30	12	-30	1.50	.03

Table 8
Algae
Ash Analyses

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
6	AOW-11N	43746	16	- 75	-50	3	-50	0.50	-.001
10	A4W-10N	43752	2	- 30	- 2	- 1	- 2	-.1	-.001
11	A2W-10N	43751	63	60	- 2	3	2	-.1	-.001
12	A1W-10N	43748	173	I.S.	-30	3	-30	1.00	.02
13	AOW-10N	43745	47	- 30	14	- 1	4	0.84	.003
21	AOW-9N	43744	51	- 75	10	- 1	- 5	3.95	.24
37	AOW-7N-2	43743	770	- 30	8	4	- 2	-.1	.001
39	A1W-7N	43747	1833	- 30	14	2	- 2	1.74	.09
40	AOW-7N-1	43742	662	- 50	13	7	3	.2	.02

Table 8 (Contd)

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
135	A1	43740	14	I.S.	-20	37	-20	-.2	.006
139	A1E-11S	43750	28	-150	-10	1	-10	0.50	.009
*	A2	43741	5	- 50	- 3	24	- 3	-.1	.01
**	AKB-2	43749	-20	I.S.	-15	1	-15	-.2	.007

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* Sample location -- near Kendrick Bay Camp -- not shown on map

** Sample location -- west end of Kendrick Bay -- not shown on map

Table 9

Miscellaneous Plants

Ash Analyses

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
24	300E-900N	<u>A</u> 43660	-20	-150	75	8	45	0.33	.03
32	100W-800N	<u>U</u> 43687	-20	-150	45	7	35	2.44	.057
32	100W-800N	<u>A</u> 43663	- 6	-150	70	15	20	1.70	.09
51	100W-500N	<u>U</u> 43688	- 3	- 75	35	4	5	-.1	.02
53	0W-500N	<u>A</u> 43664	- 6	-150	120	9	40	0.70	.06
57	700E-500N	<u>A</u> 43662	- 6	-150	130	5	90	0.80	.09
76	25E-300N	<u>T</u> 43679	374	-150	80	5	10	0.80	.04
81	AT9	<u>M2</u> 43676	-20	I.S.	-15	5	65	2.01	.04
81	AT9	<u>T</u> 43680	-20	I.S.	-20	25	13	1.67	.05
87	35E-200N	<u>M</u> 43675	923	-150	70	19	60	1.60	.08
95	100W-100N	<u>M2</u> 43677	832	I.S.	-20	2	140	4.00	.05
107	200E-50S	<u>M2</u> 43678	119	I.S.	-15	4	130	4.25	.06
117	300E-300S	<u>A</u> 43661	- 6	-150	80	5	30	0.70	.07

A = Alder U = Unidentified Shrub T = Club Moss M2 = Crowberry M = Luketkea Pectinata

Table 10
Mulch and Soil Analyses

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
1	M1W-13N	<u>M</u> 43693	109	I.S.	23	27	35	0.57	
2	M1W-12N	<u>M</u> 43694	7	- 37	20	5	15	-.1	
6	COE-11N	<u>M</u> 43716	- 1	- 30	10	4	- 2	-.1	.004
14	M1E-10N	<u>M</u> 43692	229	- 75	40	11	20	2.05	.03
19	M2W-9N	<u>M</u> 43695	30	- 30	60	6	160	.1	
20	M1W-9N	<u>M</u> 43725	47	- 75	- 5	6	20	-.1	.004
22	M1E-9N	<u>M</u> 43713	6	- 50	3	7	- 3	-.1	
27	AT-8M	<u>M</u> 43718	-20	I.S.	-15	9	-15	.2	.009
28	AT-7M	<u>M</u> 43717	- 1	- 30	- 2	12	- 2	-.1	.005
29	AT-6M	<u>M</u> 43690	- 6	-150	-10	12	-10	0.1	.02
30	AT6-M2	<u>M</u> 43710	114	- 30	16	- 1	12	.2	
31	2W-8N	<u>M</u> 43699	7	- 30	- 2	4	- 2	-.1	
34	M1E-8N	<u>M</u> 43712	42	-150	-10	16	50	.1	
44	S5W-6N	<u>M</u> 43715	12	- 30	14	5	- 2	-.1	.003
48	S5W-5N	<u>S</u> 43739	3	- 30	- 3	1	- 3	-.1	.003
49	S4W-5N	<u>S</u> 43738	11	- 30	- 2	2	- 2	-.1	.003
55	M1E-5N	<u>M</u> 43723	441	-150	-10	23	70	-.1	.01
59	AT4-M	<u>M</u> 43701	13	- 50	7	- 1	20	.2	
60	AT5-M	<u>M</u> 43709	15	-150	-10	25	40	.1	
61	AT2-M	<u>M</u> 43707	12	- 37	7	- 1	5	-.1	
65	AT-1M	<u>M</u> 43697	19	- 50	- 3	11	15	.2	
66	AT-3M	<u>M</u> 43698	-20	I.S.	-15	15	25	1.45	
67	S5W-4N	<u>M</u> 43706	93	-150	-10	323	40	.3	
72	S5W-3N	<u>S</u> 43737	2	- 30	- 3	2	- 3	-.1	.007
77	M1E-3N	<u>M</u> 43691	110	- 30	12	4	4	-.1	.001
81	AT-9M	<u>M</u> 43719	3	- 30	18	4	10	-.1	.004
82	AT-10M	<u>M</u> 43720	-20	I.S.	-15	6	35	.2	.009
84	S5W-2N	<u>S</u> 43736	5		2	16	- 2	-.1	.003
107	S200E-50S	<u>M</u> 43705	22	- 30	- 2	2	- 2	-.1	
127	MOE-800S	<u>M</u> 43722	12	- 30	20	6	- 2	-.1	.006
132	M-1	<u>M</u> 43701	35	-150	-10	25	50	.1	
133	MOE-10S	<u>M</u> 43714	12	- 75	15	80	25	.2	
134	M1E-10S	<u>M</u> 43724	238	- 75	5	12	30	-.1	.004
136	M-2	<u>M</u> 43726	39	- 75	-15	9	45	.2	.02
138	MOE-11S	<u>M</u> 43721	-20	I.S.	-20	16	-20	.3	.01
139	M1E-11S	<u>M</u> 43700	-20	I.S.	-15	19	45	.4	
140	M-3	<u>M</u> 43702	1	- 30	10	5	6	-.1	
141	M-4	<u>M</u> 43696	- 6	-150	-10	4	20	.4	
	M-7	<u>M</u> 43703	- 3	- 75	- 5	7	05	-.1	
	M-6	<u>M</u> 43704	8	- 30	34	1	14	.1	
	M-5	<u>M</u> 43711	1	- 30	- 2	- 1	2	.2	

M = Mulch S = Soil

Table 11
Stream Sediment Analyses

Map No.	Field No.	Analytical Code No.	U ppm	V ppm	Mo ppm	As ppm	Cu ppm	Mn %	Zn %
1	RAC-10 OW-900N	43733	7	-30	-2	2	-2	-.1	.003
2	RAC-11 300E-500N	43734	8	-30	-2	2	-2	.1	.001
3	RAC-12 650E-450N	43735	33	-30	-2	2	-2	-.1	.003
4	C-1	43727	554	-30	-2	5	-2	-.1	.002
5	C-2	43728	315	-30	-2	3	-2	-.1	.003
6	C-3	43729	253	-30	-2	2	-2	-.1	.002
7	C-4	43730	291	-30	-2	3	-2	.1	.004
8	C-5	43731	198	-30	-2	4	-2	-.1	.003
9	C-6	43732	77	100	-2	6	4	-.1	.005

Table 12
Semi-Quantitative Spectrographic Analyses of Six Geobotanical Samples

Map No.	98	118	97	91	83	4
Type of Sample:	Lodgepole Pine	Lodgepole Pine	Lodgepole Pine	Western Cedar	Spruce	Stream Sediment
Lab. No.	43341	43382	43428	43458	43543	43727

The following are reported as oxides of the elements indicated.

Element:	%	%	%	%	%	%
Si	6.	4.	5.	3.	8.	N.D.
Ca	12.	12.	12.	40.	15.	0.25
Mg	8.	15.	6.	5.	20.	.35
K	18.	30.	15.	15.	18.	5.
Na	3.5	3.	3.5	.6	3.5	6.
Al	2.5	2.	2.5	.4	2.5	18.
P	3.5	4.	5.	3.	6.	--
Fe	1.	.8	1.2	.5	2.	7.
Mn	1.2	2.	1.	.08	4.	.3
B	.2	.3	.25	.3	1.	--
Pb	.03	.02	.04	.015	.06	.02
Ga	.001	--	.001	--	.001	.01
Mo	--	-.002	--	--	--	--
Cu	.02	.015	.02	.003	.02	.001
Yb	.002	-.001	.002	-.001	.001	.004
Zn	.35	.15	.4	--	.1	--
Ti	.025	.03	.03	.03	.06	.3
Ag	.0005	.001	-.0005	--	--	--
Zr	.025	.004	.04	.05	.02	.2
Ni	.003	.004	.003	.001	.005	.001
Sr	.04	.03	.02	.05	.05	.01
Cr	.0005	.0007	-.0005	-.0005	.0008	.0005
V	.002	.002	-.001	--	.002	--
Y	.02	.015	.01	.003	.03	.06
Ba	.01	.01	.01	.03	.01	.015

Balance in first five samples: Not found. N.D: Si + nondetectables: Balance in #43727.

Uranium in ppm by Fluorimetric method:	1020	573	2396	155	208	554
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