

A 10 ka to 1.8 Ma Record of Oxygen-18 and Carbon-13 Variation
in Lake Sediments: Searles Lake, Inyo County, California

by
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DEDICATION

to my parents, Rock and Wheez
for their love, support and humor

and

to the memory of
Dr. Robert Gernant
for introducing me to thinkin' rocks and laughin' places

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ABSTRACT

Searles Valley, California has been the site of continuous, yet variable, lacustrine sediment deposition during the past 3.2 million years. A detailed history of fluctuating lake size, reflecting changing climatic conditions, has been recorded in the ^{18}O and ^{13}C content of primary carbonate sediments at Searles.

The ^{18}O and ^{13}C content of 247 samples from mud units in the upper 400 meters of Searles Lake lacustrine carbonate sediments (core KM-3) were measured using a timed-extraction technique based on the differing rates of reaction during phosphorolysis of co-existing primary carbonates. An average fractionation between calcite and dolomite of +2.8 per mil ^{18}O and +1.0 per mil ^{13}C was determined for the co-existing carbonates.

The samples were assigned ages based on the interpolation of samples between depths having known ages through the ^{14}C , ^{230}Th , ^{36}Cl , and magnetostratigraphic dating of core KM-3 by previous investigators. The resulting stable isotope chronology spans a time period of 10 ka to 1.8 Ma, with a time resolution that ranges from 1,000 to 10,000 years throughout most of the mud units sampled.

An analytical solution of lake volume-isotopic content evolution was used to predict the range of ^{18}O variation

during extremely wet/cold and dry/warm conditions. The measured ^{18}O content of primary carbonates at Searles were within the range predicted by the analytical solution, supporting assumptions of primary origin of the lacustrine carbonates.

Frequencies of lake-size fluctuations on the order of 40 to 50 kyr and 100 kyr are apparent throughout most of the sediments sampled. Searles Lake may have overflowed at approximately 14 ka, 210 ka, 400 ka, 625 ka, 800 ka, and 1.05 Ma. Complete desiccation of the lake is suggested by the stable isotope record at 1.48 Ma, 280 ka, and 27 ka.

Possible origins of the climatic changes producing the lake size fluctuations inferred by the stable isotope record were examined. Lake-size controls may include local or regional controls such as lake topography, lake-atmospheric feedback processes or regional tectonic events such as the uplift of the Sierra Nevada during the Cenozoic. Global controls may include changes in solar insolation due to the Earth's orbital cycles.

Key words: stable isotopes, oxygen-18, carbon-13; paleolake; paleoclimate; California, Searles Lake

I. INTRODUCTION

Problem Addressed

Predicting available water supplies for man's use is a difficult task. Growing populations, accompanied by threatened water supplies due to contamination, add stress to already over-allocated fresh water supplies in some areas. Further, the predicted changes in climate due to the accumulation of "greenhouse gases" in the atmosphere has added to the already complex problems of water-resource planners. However, to begin predict future climates and water supplies, and man's impact upon them, we must first establish an understanding of the frequency, extent and cause(s) of climatic changes in the geologic past, and the effects these changes have had on water supply.

In order to examine past climatic changes, a climate recorder must be found which would have been climatically sensitive, continuously active in the past, and suitable for absolute dating techniques. Stable isotope records from deep sea and ice cores have provided detailed climatic records for Quaternary oceanic and polar regions (Williams and others, 1988; Lorius and others, 1985). Analysis of paleolake levels at low latitudes has revealed systematic lake level fluctuations with latitude through the last 18,000 years (Street-Perrott and Roberts, 1983). However, mid-latitude paleoclimate records are scarce, primarily due to the lack of continental depositional systems operating

continuously in the past. The closed-basin lakes of the western United States Basin and Range provide one of the most continuous and sensitive record of mid-latitude continental Quaternary climate change currently known.

Lake size fluctuates in response to changes in climate. The relationship between lake area/volume and inflow/outflow is termed a lake's "water balance". Outside of tectonic or volcanic events, changes in a lakes's water balance primarily reflect changes in climate.

Lake levels in closed-basin amplifier lakes respond most sensitively to climatic perturbations. "Closed-basin" refers to those lakes which lack a surface outlet, and "amplifier" implies inflow/influx is dominated by surface runoff rather than precipitation on the lake surface, and groundwater transfer is negligible (Szestay, 1974).

Evidence of fluctuating lake levels can be observed in the sequences of abandoned shorelines and spillways, and wave-cut cliffs surrounding closed-basin lakes. Some of these features may be absolutely dated, allowing development of an absolute chronology. Unfortunately, older geomorphic remnants are frequently removed by the rising waters of a younger lake.

A more detailed climatic record can be found in the sediment sequences of closed-basin lakes. Solutes entering the lake are concentrated by evaporation and lack of surface or groundwater outflow. Consequently, most closed-basin

lakes are saline and alkaline. As solute concentrations increase with time or decreasing lake levels, sequences of evaporite minerals are precipitated. With the resumption of large inflows and rising lake levels, non-evaporite minerals are deposited. Thus, lake level fluctuations are recorded in the alternating sequences of muds and evaporites.

In addition to the facies information, the chemical sediment record provides detailed geochemical information. The stable isotope content of lake water is extremely sensitive to inflow/outflow fluxes and their respective isotopic composition. As the lake water balance changes, so will the isotopic composition of the lake water. The temperature, salinity and stable isotope content of the lake water are reflected in the stable isotope content of minerals precipitated in isotopic equilibrium with the lake water. Further, evaporite minerals may be suitable for some absolute dating techniques. Consequently, a continuous record of the stable isotope content of the lake water, together with sediment and geomorphic constraints, can provide a detailed, sensitive record of water balance fluctuations and Quaternary continental climatic change.

Searles Lake, a currently dry, closed-basin amplifier lake in southeastern California (fig. 1), has been the subject of dozens of mineralogic, geochemical and paleohydrologic studies over the past one hundred years. Elevated shorelines and bars in Searles Valley and related

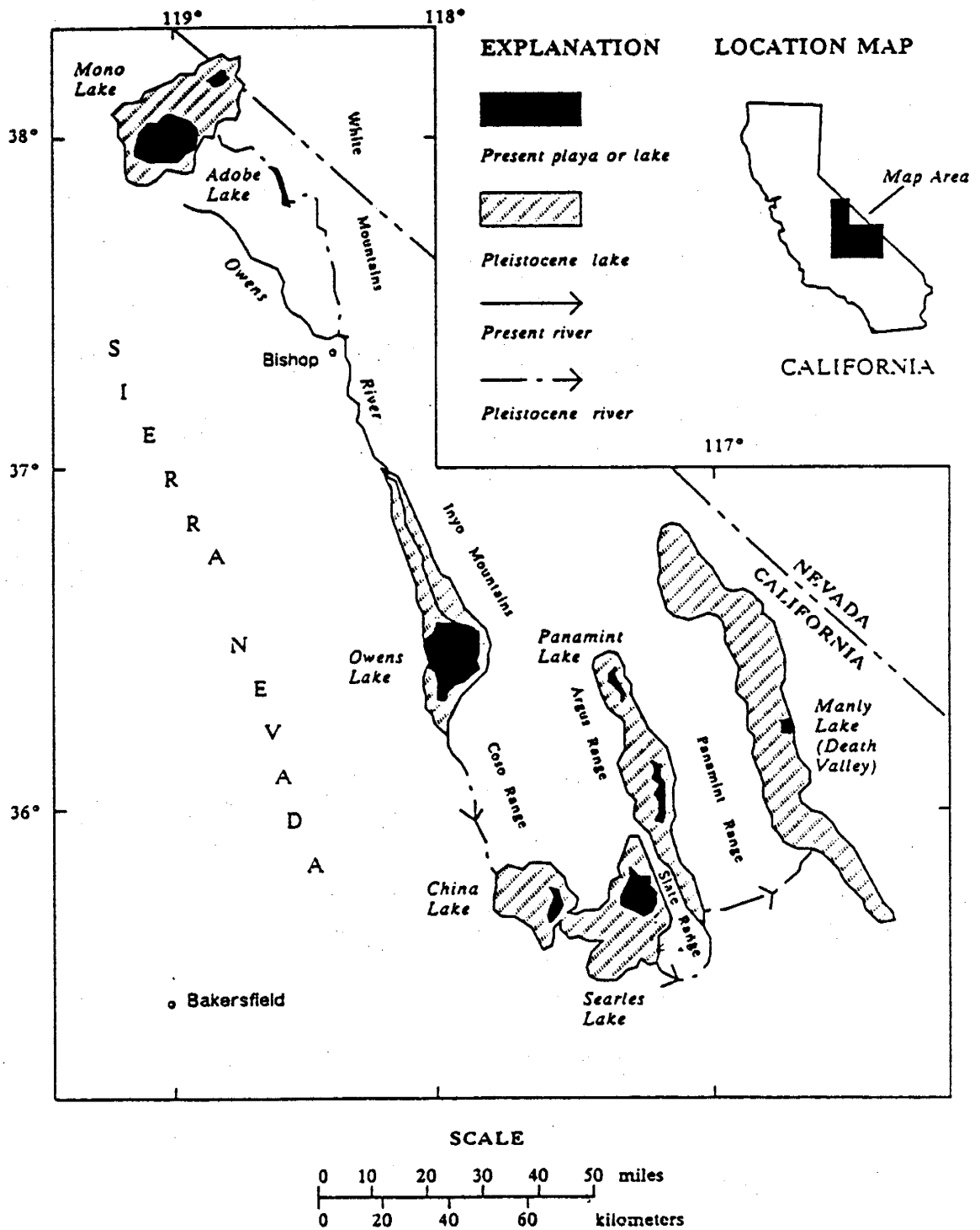


Figure 1 Location of Searles Lake and related Pleistocene lakes. Revised from Smith and others (1983).

basins were recognized by nineteenth-century geologists as evidence of past deep lakes. Early twentieth-century studies revealed the relationship between Searles Lake and four other lakes in the area. Runoff from the eastern Sierra Nevada into the paleo-Owens River created a chain of five paleolakes. Searles Lake was the third and, for most of the Pleistocene, the terminal lake in the chain. Consequently, large amounts of solutes were concentrated in Searles and precipitated as chemical sediments. Exploration cores, drilled by mining companies and the U.S. Geological Survey to define the mineral deposits at Searles, revealed thick sequences of alternating muds and evaporite minerals. The term "muds" has traditionally been used by investigators at Searles to include clays, silts, and marls consisting primarily of alkaline earth carbonates and diagenetic minerals.

Of particular interest is Kerr-McGee core KM-3, a 930 meter core drilled to bedrock in 1968 (Smith and others, 1983). Of the 930 meters of core, 693 meters are believed to be lacustrine deposits which date to the early Pliocene. An absolute chronology for KM-3 has been established by means of numerous carbon-14 (Flint and Gale, 1958) (Stuiver, 1964) (Stuiver and Smith, 1979) , uranium-thorium (Peng and others, 1978) (Bischoff and others, 1985), chlorine-36 (Phillips and others, 1986) (Jannik, 1989) and paleomagnetic (Liddicoat and others, 1980) dates. Therefore, with the

benefit of a detailed and lengthy chronology already in place, in addition to geochemical and stratigraphic evidence assembled, core KM-3 can provide a long record of mid-latitude continental climatic change.

Objective and Approach

The primary objective of this study was to construct a record of stable isotope variation in Searles Lake sediments for the last 2.0 Myr [in this paper I follow the convention of designating ages by 'Ma' (million years before present) or 'ka' (thousand years before present) and periods of time by 'Myr' (million years) or 'kyr' (thousand years)]. Preliminary interpretation of the record, in addition to comparison with other isotope records, was made. The approach taken to accomplish the objective is as follows:

- 1) Sample Searles Lake Core KM-3 with enough frequency to determine degree and duration of climatic cycles.

It has been recognized from deep-sea and ice core isotopic records that global climate has fluctuated in a cyclic fashion throughout the Pleistocene (Hays and others, 1986; Dansgaard, 1969). A sampling interval in time comparable to those in deep-sea cores would allow a similar definition of climate frequencies at Searles. Further, detailed sampling would allow time-series analysis of the Searles Lake isotope chronology so that frequencies and possible causes of the climatic fluctuations may be ascertained.

- 2) Examine composite chronology proposed by Jannik(1989). Evaluate sources of error for

each dating technique. Assign ages to all KM-3 samples using composite chronology.

The chronology proposed by Jannik is a composite of selected ^{14}C , U-Th, ^{36}Cl , and paleomagnetic dates, in addition to dates based on accumulation of acid-insoluble residue. An examination of the dating methods, dates selected, and relative error is warranted before time-series analysis, comparison with other paleoclimate records, or numerical modeling of the lake begins.

3) Develop methodologies to separate primary carbonate minerals from diagenetic carbonates, and to separate primary carbonates from each other.

Searles lake sediments contain a variety of primary and diagenetic carbonate minerals. Carbonates termed "primary" include dolomite, calcite, aragonite and northupite. Some gaylussite may also be primary. These minerals are presumed to have precipitated in isotopic equilibrium with the lake water. Diagenetic carbonates are formed through the reaction of infiltrating concentrated brines with primary carbonates to form new minerals, such as gaylussite and pirssonite. Therefore, diagenetic carbonates may not reflect the isotopic composition of the original lake water.

4) Perform mass spectrometric analysis of primary carbonates to determine oxygen-18 (^{18}O) and carbon-13 (^{13}C) content.

Interpreting the ^{13}C and ^{18}O content of CO_2 extracted from primary lake carbonates assumes the CO_2 incorporated into the mineral at the time of precipitation was in isotopic equilibrium with the dissolved lake CO_2 . A second assumption is that the dissolved lake CO_2 was in isotopic equilibrium with lake water. If these assumptions are born out, analysis of ^{13}C and ^{18}O content of carbonates would allow determination of the isotopic content of the paleolake water.

5) Estimate paleometeorological parameters such as relative humidity, temperature, net evaporation. Use analytical model to predict range of expected ^{18}O and ^{13}C during infill, overflow and desiccation episodes. Compare range of values to mass spectrometer results.

Comparison of the isotopic content of the Searles Lake carbonates with the isotopic content predicted by an analytical solution allows evaluation of the methodologies used to separate and extract the carbonates from the lake mud, and the validity of the assumptions of primary origin of the carbonates. Second, the analytical results will allow validation of the numerical model under construction. Finally, the analytical model will allow preliminary interpretation of the Searles Lake isotope chronology.

6) Interpret Searles Lake isotope chronology. Compare Searles Lake isotope chronology to other Great Basin and deep-sea isotope records.

The record of the stable isotope content of the lake water through time will permit interpretation of changes in lake water balance (Phillips and others, in progress). From the water balance, changes in climate may be inferred, and the response of the Searles Lake water balance due to changes in climate may be determined.

Comparison of the Searles Lake isotope record to other paleoclimate records will allow examination of the relationship between oceanic and continental paleoclimates. The geographic extent of climate change affecting Searles may be determined: local or regional changes affecting only Searles, the Great Basin, or global climatic changes. The correlation of climatic changes to other events will be examined: whether the climate changes relate to volcanic or tectonic activity, or to the waxing and waning of global ice volumes, presumably due to changes in solar insolation.

II. RELATED STUDIES

Closed Basin Lakes

One of the earliest, most insightful, and possibly the most enjoyable reports on a closed-basin lake of the Great Basin can be found in Russell (1885). Russell recognized that the Quaternary closed-basin lake levels were climate-dependent and hypothesized the changes in humidity and temperature that would have been required to produce the large saline lakes. Meinzer (1922) took a broader approach to paleoclimate reconstructions of closed-basin lakes. All closed-basin lakes in the western U.S. were included in a discussion of the relationships between the water balance of these lakes and their respective locations, geology, bathymetry, and climate.

One of the most extensive quantitative reconstructions of paleoclimates using closed-basin lake records can be found in Mifflin and Wheat (1979). Paleotemperatures were assumed in order to calculate the changes in evaporation, precipitation and runoff necessary to produce the paleolake levels indicated by geomorphic evidence in Lake Lahontan, Nevada.

Smith and Street-Perrott (1983) calculated the required inflow from the Owens River necessary to produce the Owens River system closed-basin paleolakes, including Searles Lake. Assumed paleotemperatures were used to calculate

paleo-evaporation and precipitation. Street-Perrott and Harrison (1985) summarized the factors affecting the sensitivity of closed-basin lakes to climatic change, the various approaches to reconstructing lake levels, and evidence used by to investigators accomplish the reconstruction.

This paper is the first part of a study to reconstruct paleo-water balances through the construction of stable isotope records from the sediments of two Pleistocene lakes: Searles Lake in southeastern California (this paper) and Lake San Agustin in west central New Mexico (Johnson, in progress). Some of the previous investigations that used stable isotopes to determine the water balance of lakes can be found summarized in Pearson and Coplen (1978).

A numerical approach to isotopic evolution in closed-basin lakes was taken by Phillips and others (1986). A lumped-parameter numerical model was created to simulate the isotopic evolution of a lake through time, given specified fluxes, meteorological and chemical parameters. The model was validated by simulating the isotopic evolution of Owens Lake (California) during a desiccation episode. The numerical model in development to reconstruct the paleo-water balance of Searles and San Agustin is based on the Phillips and others (1986) model.

The causes of lake level fluctuations have been discussed by several investigators. Kutzbach and Street-

Perrott (1985) correlated fluctuations in tropical lake levels with fluctuations in solar insolation calculated from the Milankovitch theory. Changes in sedimentation in Searles Lake due to orbital forcing have also been suggested by Smith (1984); Bischoff and others (1985); and Jannik (1989). Lake level fluctuations caused by changing atmospheric circulation patterns have been suggested by Street-Perrott and Roberts (1983).

Many records for Quaternary climates have been published. Among the most important are the stable isotope records from ocean cores. Most recently, Martinson and others (1987) presented a 1.8 Myr "stacked" and "orbitally-tuned" ^{18}O record: the record was "stacked" in that it was a normalized composite of several ocean cores, and "orbitally-tuned" in that orbital forcing was presumed to be the driving force of sea-level fluctuations, and the chronology was adjusted to be "in sync" with orbital frequencies. Williams and others (1988) presented a 1.8-Myr ^{18}O record that was also a normalized composite of several cores, yet was not "orbitally tuned".

Detailed isotope records from polar ice cores have also been published. Dansgaard (1969) presented a deuterium chronology from a Greenland ice core. Isotope records from Antarctica ice cores were published by Barkov and others (1979) and Lorius and others (1985).

Several long paleoclimate records from the Great Basin have been developed in the past few years. Winograd and others (1985) measured the deuterium content in calcite fluid inclusions from eastern Nevada calcite veins. The calcite was precipitated from calcite-saturated groundwater, and therefore, the deuterium content of the fluid inclusions should reflect the deuterium content of the groundwater. A systematic depletion in the deuterium values during the past 2.0 Myr was theorized to be due to the uplift of the Coastal Ranges and the Sierra Nevada during the Quaternary. A 50 ka to 310 ka ^{18}O record from a southwestern Nevada calcite vein was reported by Winograd and others (1988). The stable isotope record mimicked ocean and ice core ^{18}O records presented by Martinson and others (1987) and Lorius and others (1985) respectively. However, the timing of the ^{18}O fluctuations in the Nevada calcite did not correspond to that of the ocean and ice core records. Rather, the isotopic events identified in the Nevada ^{18}O record occurred decreasingly earlier through time than the other two records suggest. A mechanism for a systematically decreasing time-lag of climatic events among the three study areas is not apparent. It would appear that some systematic error occurred in the development of the absolutely-dated chronology for at least one of the ^{18}O records. The discrepancy in the timing of climatic events suggest by the ocean ^{18}O record and the Nevada ^{18}O record is of great

interest in regard to previously inferred dependence of climatic fluctuations upon orbited forcing (Hays and others, 1976). If the Nevada ^{18}O chronology is assumed to be correct, the differences in the timing of events between the Nevada ^{18}O chronology and the Martinson and others chronology, which was "orbitally tuned", suggest that solar insolation fluctuations predicted by the Milankovitch (1941) theory are not the primary forces contributing to climatic change (Winograd and others, 1988, p.1279).

A detailed discussion and comparison of the lake level histories in four Great Basin closed-basin lakes is provided in Benson and others (in press). Geomorphic, sedimentological and geochemical evidence of lake level fluctuations for Lake Bonneville, Lake Lahontan, Lake Russell and Searles Lake were discussed, and the lake level fluctuations in the four lake systems were concluded to be penecontemporaneous.

Searles Lake

An impressive volume of work has been published on the geology, chemistry, and chronology of tectonic and climatic events in Searles Valley. Russell (1885) included the paleolake occupying Searles Valley on Plate 1 of his section on Great Basin paleolakes. Bailey (1902) recognized the relationship between Searles Lake and other Owens River paleolakes. The discovery of large deposits of potash in

Searles Lake drew investigators to the area to define and exploit the mineral deposits in the early twentieth century. Gale (1914) reviewed the known available mineral assemblages at Searles and drew remarkably insightful conclusions as to the history of sedimentation in Searles and related paleolakes.

Investigations at Searles were resumed during the late 1950's with the advent of suitable radiometric dating techniques and a drive by the United States Geological Survey (USGS) to better define the lacustrine deposits beneath Searles Valley. Hundreds of sediment cores were drilled by mining companies and the USGS. Stratigraphic studies based on several of these cores can be found in Smith and Pratt (1957), Flint and Gale (1958), Haines (1959), Smith (1962; 1979) and Smith and others (1983).

Detailed discussions of Searles Lake mineralogy include Droste (1961); Hay and Moiola (1964); Smith and Haines (1964), and Smith (1976; 1979). The geochemistry of Searles Lake deposits and brines include reports by Eugster and Smith (1965), Eugster and Hardie (1978), Smith (1979), Guldman (1984), and Hay and Guldman (1987). The hydrology of Searles Lake is reported in Hardt and others (1972).

The recognition by investigators of the usually favorable opportunity to obtain a detailed record of continental climatic change prompted numerous studies employing isotopic, and paleomagnetic techniques to date

Searles Lake sediments. Investigators took advantage of new developments in radiocarbon techniques to establish a chronology for the shallow sediments beneath Searles Lake. These include Flint and Gale (1957), Stuiver (1964) and Stuiver and Smith (1979). Reasonable agreement of ^{230}Th ages of upper stratigraphic units with radiocarbon dates (Peng and others, 1978) allowed a continuation of the ^{230}Th chronology to units at the top of the Mixed Layer (Bischoff and others, 1985). Identification of paleomagnetic boundaries in Searles Lake core KM-3 extended the absolute chronology to 3.15 Ma (Liddicoat and others, 1980). Several halite units were dated by the ^{36}Cl method (Phillips and others, 1983; Jannik, 1989). A tephra layer in core KM-3 was tentatively correlated with the Lava Creek B ash (Hay and Guldman, 1987). Radiocarbon dates on rock varnish from high-stand shorelines are presented by Dorn (in Benson and others, in press).

Paleoclimatic interpretations of Searles Lake based on stable isotope information have been presented in Oana and Deevey (1960), Stuiver (1964), and Horita (1987). Paleoclimatic interpretations based on sedimentological and outcrop evidence have been presented in Smith (1968; 1976; 1979; 1984; 1987), and Smith and others (1983).

III. STUDY AREA

Location

Searles Lake is located in the southwestern corner of the Basin and Range physiographic province along the boundaries of Inyo and San Bernadino counties, California (35 45'N, 117 20'W). Currently, Searles Lake is a dry (playa) lake, with standing water only after intense rainfall events (Smith, 1979).

During pluvial periods, Searles Lake was the third in a chain of five paleolakes created by large runoff from the eastern Sierra Nevada into the ancient Owens River. However, Owens Lake was the frequent terminus of the Owens River during interpluvial periods, as it is today. According to Smith and others, (1983) when flow in the Owens River increased, Owens Lake filled until at approximately 64 m in lake depth, Owens Lake overflowed into Indian Wells Valley to form China Lake. Upon reaching a depth of 8 m, China Lake, in turn, overflowed into Searles Valley to form Searles Lake. When Searles Lake deepened to ~180 m, China Lake and Searles Lake coalesced to form one large lake (Searles Lake), which at a depth of 200 m overflowed into Panamint Valley, creating Panamint Lake. During extreme pluvial periods, Panamint Lake would overflow into Death Valley. Lake Manly (Death Valley) was also the ultimate sink for the Armagosa and Mojave River paleolake systems.

In addition, lakes in the Mono Basin, such as Mono Lake and Adobe Lake may have overflowed during extreme pluvial periods, contributing to the flow in the ancestral Owens River system. Figure 1 shows the relationships between all of the paleolakes. Figure 2 is a schematic cross-sectional representation of the Owens River paleolakes.

Hydrology

Currently, Searles Lake playa, located in the bottom of Searles Valley, has a surface area of approximately 100 km². According to Smith (1979), Searles Valley floor is 60 km long and up to 15 km wide. The playa surface itself is 15 km long and 11 km wide; approximately 66 percent of the playa is silt covered, and the remaining third is hard salt pan.

Searles Valley has a drainage area of approximately 1600 km² (Smith, 1979). However, during intense pluvial periods, most of the water in Searles Lake would have originated from eastern Sierra Nevada runoff via the Owens River. Lee (1912) reports that the Owens River has a drainage area of approximately 3300 mi² (~8550 km²) of which 536 mi² (1390 km²), in the high altitude area of the eastern Sierra Nevada, contributes most of the runoff reaching Owens River. The remaining drainage area is predominantly either desert or transition slopes. From Lee's work, Jannik (1989) postulated that even during intense pluvial periods, the

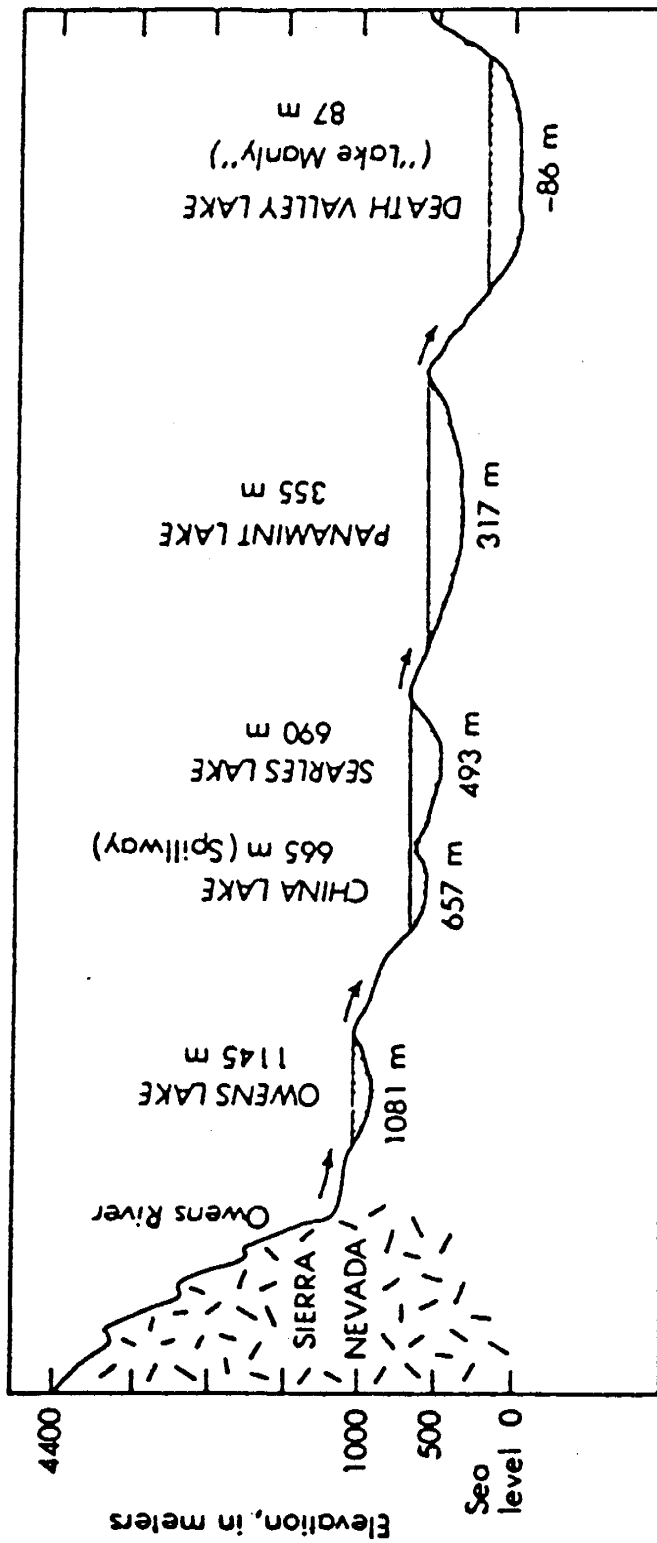


Figure 2 Schematic cross-section of lake basins in the paleo-Owens River Chain. Smith and Street-Perrott (1983).

Sierra Nevada created such a strong rain shadow that most of the water in Searles Lake still originated from the high-altitude section of the eastern Sierra Nevada. Any precipitation falling within Searles Lake's drainage area would have been negligible compared to the inflow into Searles via Owens and China Lakes.

Hydrothermal springs in the Owens Valley area also contributed to the flow in the Owens River paleolake systems (Jannik, 1989). Of equal interest, the springs may have contributed some of the unusual solutes found in Searles Lake evaporite deposits.

Climate

Most of the following climate information was found in Smith (1979). The climate at Searles Lake is hot and arid. Temperature extremes range from -12.2 C to 47.8 C, with an average annual temperature of 19.1 C (for a 36-year period ending 1973). Annual precipitation during this period averaged 96.3 mm, with extremes ranging from 23 mm to 291 mm per year. Most of the precipitation occurs during the winter. I was not able to acquire relative humidity data, although it is reported (Horita, 1987) that Kerr-McGee Corporation in Trona, California maintains a weather station, and humidity values are recorded. A linear regression of relative humidity data in the surrounding area gave an approximate annual average relative humidity for

Searles Valley of 22 percent (Appendix G).

Searles Valley playa supports no vegetation. Smith (1979) reported vegetation on the surrounding mountains and alluvial fans as typical of the lower Sonoran zone: desert holly (Atriplex hymenelytra), creosote bush (Larrea tridentata), burro-bush (Franseria dumosa), and hop sage (Grayia spinosa). Piñon pine and joshua trees are reported in the surrounding mountain ranges.

Geology

Geologic Environment According to Smith (1979), Searles Valley is a closed tectonic depression of Cenozoic age. The valley floor is surrounded by several mountain ranges: the Argus Range and Spangler Hills to the west, the Slate Range on the north and east, and the Lava Mountains to the south. The Argus Range consists primarily of Mesozoic granites. The Slate Range contains Precambrian metamorphic units, small amounts of Paleozoic limestone, Mesozoic granites and metavolcanics and Cenozoic volcanics (Jennings and others, 1962 in Jannik, 1989). The Lava mountains consist primarily of late Tertiary volcanics and sandstone (Smith, 1968).

Core KM-3 Numerous sediment cores have been recovered from the lake bed in Searles Valley. A detailed examination of the chemistry and mineralogy of sediments from several cores in Searles Lake is provided in Smith (1979). In 1968, a 930

m exploration core designated KM-3 was drilled to bedrock by Kerr-McGee Corporation. In 1976, the United States Geological Survey (USGS) was granted permission to examine company data on the core, and the core itself, and to publish the results of their investigation, including internal Kerr-McGee data. The findings of the USGS investigation, including detailed core description, chemical analyses, and interpretation can be found in Smith and others (1983).

Detailed geochemical analyses were performed on the KM-3 sediments by Kerr-McGee Corporation. The core was divided up into 254 units based on lithology. Analysis for percent acid-insoluble residue, and ten elements in the acid-soluble fraction. In addition, the percent by weight of important evaporite minerals were calculated, including halite and several carbonate, sulfate, and borate minerals. The results of these analyses (Smith and others, 1983, Plates 2-4) were used as a general guide in selecting samples from core KM-3 for stable isotope analysis.

The following core descriptions, stratigraphic and mineralogic information were obtained from Smith (1979) and Smith and others (1983). Core KM-3 was drilled near the center of Searles Valley (fig. 1). The ~8 cm diameter core consisted of three distinct sections: 1) 15 m of quartz monzonite bedrock of which only ~3 m of core was actually recovered; 2) 212 m of alluvial arkosic sand and gravel, and

3) 693 m of lacustrine sediments. The lacustrine sediments were divided into fourteen informal stratigraphic units based on lithology and mineralogy. The informal stratigraphic units are those that have been used by the mining companies at Searles Lake, and have been adopted by investigators. A copy of the core log summary presented in Smith and others (1983) is shown in table 1, and is shown graphically in figure 3.

In the broadest sense, Searles Lake sediments have been divided into two categories: "muds" and "salts". These are not sedimentologically exact terms, but they are useful in delineating sediments deposited in different environments and have a long history of application to Searles sediments. As mentioned earlier, the "muds" can consist of marl, silt or clay, and are plastic when wet. The muds vary in origin from deep lake marls and oozes to playa sediments, tephra, and possibly paleosols. The marls contain clays as well as alkaline earth carbonates (calcite, dolomite, aragonite), clastics, and diagenetic carbonate minerals such as gaylussite and pirssonite. The saline units can be thin seams of single or multiple evaporite minerals, such as hanksite, borax, tincalconite, thenardite, northupite, nahcolite, and halite, to thick coarsely crystalline units of halite or trona. Many of the saline minerals can also be found disseminated within the mud units.

Table 1. Generalized Searles Lake core KM-3 log (Smith and others, 1983).

Depth to base of unit (m)	Thickness of unit (m)	Description
5.8	5.8	Not cored. Overburden Mud is included in this zone.
19.9	14.1	<i>Upper Salt.</i> —Salines, mostly trona and halite; abundant hanksite near top, borax at base; light- to medium-gray (N6-8) and yellowish-gray (5Y8/1), with some thin interbeds of olive-gray (5Y4/1) mud; mostly indistinctly bedded to massive, locally vuggy.
25.0	5.1	<i>Parting Mud.</i> —Megascopic crystals of gaylussite and pirssonite in soft mud composed of microscopic crystals of dolomite, halite, aragonite, other evaporite minerals, and elastic silicates; light- to moderate-olive gray (5Y3-6/1-4); upper part finely laminated, lower part massive.
37.9	12.9	<i>Lower Salt.</i> —Seven saline layers interbedded with six mud layers; salines are mostly halite and trona in upper two layers, trona, halite, and burkeite in underlying two layers, and trona in lower three layers; interbedded mud layers contain megascopic crystals of gaylussite and pirssonite; salines range in color from white through dark gray to yellowish orange (N5-8, 10YR6/6), mud from dark olive gray to brown (5Y4-6/1-4); salts poorly bedded to massive; some mud layers have thin laminar bedding.
69.0	31.1	<i>Bottom Mud.</i> —Mud containing megascopic gaylussite crystals; mud is composed of microscopic crystals of dolomite, aragonite, calcite, and other carbonate minerals, and about 30

Depth to base of unit (m)	Thickness of unit (m)	Description
		percent acid-insoluble silicates and organic residues; thin-bedded to massive, with some laminar bedding; medium- to dark-brown, brownish-gray, and olive (5YR4/4 to 5Y3-4/1-2). Discontinuous saline layers at 41.4 m (0.5 m thick), 48.5 m (0.4 m thick), and 54.4 m (0.8 m thick).
90.8	21.8	Interval of poor core recovery; recovered core (3.4 m) is composed of mud containing megascopic gaylussite crystals, massive, medium- to dark-brown and olive (5YR5/2 to 5Y3/2). Top of interval probably represents top of Mixed Layer. This and following three units probably represent Units A and B of Mixed Layer, of which most of the saline layers were lost during drilling (see text).
95.1	4.6	Salines, mostly trona, containing mud; faintly bedded to massive; moderate-brown to olive (5YR4/4 to 5Y5/1).
99.8	4.4	Mud, mostly acid-insoluble material, some dolomite; light-olive-gray (5Y5-6/1-2), thin-bedded.
114.0	14.2	Salines, mostly trona, with small amounts of other minerals and extensive mud impurities; light- to dark-green and brown (5GY5/1, 5Y4-6/1, 5YR3/4); faint bedding in lighter colored salines, with interbeds of mud common near base. Contact between Units B and C of Mixed Layer is at base of this interval.
124.0	10.0	Salines, with some interbedded mud; saline minerals are mostly halite, with smaller amounts of trona and other evaporite minerals; indistinct bedding, beds mostly 1 to 2 cm thick; salines light- to dark-olive-gray (5Y4-7/1-2) and moderate-brown (10YR4-6/2-4).
130.4	6.4	Salines and some mud; salines are about two-thirds halite and one-third trona, with some thenardite; yellowish-gray (5Y5-7/2); upper part of unit contains largest percentage of mud impurities.
135.6	5.2	Salines, mostly halite, with minor trona and thenardite; dark- to medium-gray (N3-5); upper part of unit contains mud impurities.
151.2	15.6	Interbedded mud and salines; salines are mostly halite, with some trona; salines olive-gray (5Y4/1 to 5Y6/1), mud brownish-black (5YR2/1); saline layers, 0.3 to 0.6 m thick, constitute about one-third of zone.
166.4	15.2	Salines, mostly halite, with smaller amounts of trona and thenardite, nearly pure in lower part; mostly gray to yellowish-gray (N4-7 to 5Y4-6/1); bedding 1 to 2 cm thick, some zones porous but most nonporous. Contact between Units C and D+E of Mixed Layer is at base of this interval.
178.6	12.2	Mud containing megascopic crystals of gaylussite and pirssonite, microscopic crystals of

Depth to base of unit (m)	Thickness of unit (m)	Description	Depth to base of unit (m)	Thickness of unit (m)	Description
		dolomite, halite, and probably other acid-soluble minerals; brownish-black (5YR2/1) in upper and lower part, moderate-brown (5YR3/4) in middle.	291.1	14.2	Mud, similar to that at 249.5 m. Contact between Units F and G (new) of Mixed Layer is at base of this interval.
186.5	7.9	Mud, with interbedded salts at 179 and 181 m; salts, in beds 0.1 and 0.6 m thick, are mostly halite and trona, with some thenardite and northupite; mud is composed largely of microscopic crystals of dolomite and other carbonates; olive to brownish-black (5Y2:1 to 5YR2/1).	294.4	3.3	Impure salines with mud impurities similar to interval above; salines, mostly halite and thenardite, are mottled aggregates surrounded by mud.
192.0	5.5	Salines interbedded with mud containing scattered saline crystals; salines are mostly halite, with subordinate trona, thenardite, and other minerals; salines olive-gray and medium- to dark-gray (5Y6/1 to N4-6), mud dark olive-black (5Y1-2/1).	299.3	4.9	Mud, dusky yellow-green (5GY5/2); upper part mottled, lower part has buff laminae and thin beds.
196.1	4.1	Mud, dark olive-black (5Y1-2/1).	306.3	7.0	Salines and some mud; salines are chiefly halite and thenardite, white to light-gray (N5-8); mud pale-green (10G6/2); upper part faintly bedded, lower part mottled.
196.6	0.5	Salines, mostly halite; olive-gray (5Y4/1).	324.3	18.0	Two saltbeds separated by mudbeds (see pl. 1); salts, largely halite, massive, light-greenish-gray (5GY6-8/1); muds massive, greenish-gray (5G6/1).
204.5	7.9	Mostly mud, with some disseminated saline crystals; brown (5YR3/4) in upper part, olive-gray (5Y4/1) in lower.	333.1	8.8	Three mud and two impure salt layers (pl. 1); mud pale-green (10G6/2), massive, with dispersed salts; saline layers, consisting of mottled zones of lighter colored secondary crystals oriented randomly in mud matrix, are halite, with some thenardite, glauberite, and anhydrite.
207.1	2.9	Salines, with interbedded mud; salines are mostly halite, distinctly bedded, averaging 1 cm in thickness; salines yellowish-gray (5Y7/2), mud olive-gray (5Y4/1).	334.7	1.6	Impure salts, halite; dark-greenish-gray (5GY4/1).
210.9	3.5	Mud, moderate-brown (5YR4/1) in upper half, olive-black (5Y2/1) in lower part.	337.3	2.6	Mud containing some halite; brownish-gray (5YR4/1).
213.6	2.7	Salines, mostly halite, with mud impurities; olive-black (5Y2/1) to light-olive-gray (5Y6/1); upper part faintly bedded, lower part massive to mottled.	341.1	3.8	Muddy salt grading downward into impure mud; salts largely halite and anhydrite; mud dark-greenish-gray (5GY4/1).
218.1	4.7	Mud, massive, olive-black (5Y3/1).	345.3	4.2	Mud with some dispersed salts; massive, faint mottled coloring; olive-gray (5Y4/1).
218.5	0.1	Salines, trona and halite.	405.7	60.4	Nine alternating salt and mud layers in nearly equal volumes, with individual layers generally 4 to 6 m thick (see pl. 1); salts are halite and other saline minerals, mostly light- through medium-gray (N5-7) to light-olive-gray (5Y6/1) and grayish-orange-pink (5YR7/2); mud greenish-gray (5G4-6/1) to dark-greenish-gray (5GY4/1); mud is massive except in 0.5-m-thick zone below saltbeds, where it is thin bedded; salts are faintly bedded to massive.
227.4	8.9	Mud, massive, olive-black (5Y2/1).	413.3	7.6	Mud; inadvertently not photographed, but reported by field log as green to brown.
227.7	0.3	Salines, trona and halite. Contact between Units D+E and F of Mixed Layer is at base of this interval.	422.5	19.3	Core not recovered.
248.1	20.4	Mud, mostly grayish-olive (5GY4/1), with a grayish-brown (5YR3/2) zone at 236-238 m and a greenish-gray (5GY6/2) zone at 244-245 m.	425.5	22.3	Mud, soft and plastic, olive-black (5Y2/1). Contact between Units G and H (new) of Mixed Layer is at base of this interval.
249.5	1.4	Mud, dark-greenish-gray (5GY4/1), mottled to faintly bedded; lower half extremely hard (limestone).	437.7	12.2	Mud, with small crystals of thenardite dispersed randomly; more coherent than interval above; average moderate-brown (5YR3/4).
271.4	21.9	Mud, with irregular concentrations of a few mottled areas caused by light-colored dolomite or salts; mostly pale- to grayish-green (10G4-6/2), upper 3 m grayish-olive-green (5GY3/2); massive except near 260 and 268 m, where thin to laminar bedding is defined by pale-orange (10YR6-8/2-4) layers (dolomite?).	444.6	6.9	Mud, soft and plastic, olive-black (5Y2/1).
276.9	6.5	Mud, similar to interval above but containing searlesite.			

Depth to base of unit (m)	Thickness of unit (m)	Description
449.6	5.0	Mud, more coherent than interval above; moderate-brown (5YR3/4).
451.4	1.8	Salts and mud; salts in thin beds and mottled areas, yellowish-gray (5Y8/1), chiefly glauberite and anhydrite, with some halite; mud massive, olive-gray (5Y4/1).
482.5	31.1	Mud, moderate-brown (5YR3/4), with some zones of light-olive-gray (5Y5-6/1-2); salts largely halite and anhydrite, both dispersed and concentrated in mottled zones.
483.4	0.9	Mud and some salts; mud olive-gray (5Y4/1), thin bedded; salts are chiefly anhydrite and halite.
494.4	11.0	Mud, moderate-yellowish- to pale-brown (10YR5/2-4), massive.
507.5	13.1	Mud, light-olive-gray (5Y5/2) to pale-yellowish-brown (10YR/2), 2-m-thick zone at 502 m is pale brown (5YR5/2).
507.8	0.3	Mud and salt; mud light-olive-gray (5Y5/2); salts are mostly glauberite.
516.6	8.8	Mud and disseminated salts; light-olive gray (5Y6/1) to pale-yellowish-brown (10YR6/2).
524.0	7.4	Mud; moderate-brown (5YR4/1) in upper part, yellowish-brown (10YR6/2) to pale-brown (5YR5/2) in lower part.
530.1	6.1	Mud; upper part greenish-gray (5GY6/1), mottled, thin-bedded to massive; lower part yellowish-brown (10YR6/2), thin-bedded.
541.6	11.5	Mud, mottled, pale-brown (5YR5/2) to pale-yellowish-brown (10YR6/2). Contact between Units II and I (new) of Mixed Layer is at base of this interval.
582.2	40.6	Mud, olive-gray (5Y4/1) down to 558 m, light-olive-gray (5Y6/1) below that depth, with 2-m-thick pale-brown (5YR5/2) zone at base.
631.0	51.8	Mud, light-olive-gray (5Y5-6/1-2).
610.1	6.1	Mud, brownish-black (5YR2/1) to olive-black (5Y2/1).
619.2	9.1	Mud, light-olive-gray (5Y5-6/1-2) and pale-olive (10Y6/2).
658.7	9.5	Mud; grayish-olive (10Y4/2) in upper part, yellowish-gray (5Y6/2) in lower part.
681.5	22.8	Mud, mostly pale-olive (10Y6/2) to yellowish-gray (5Y7/2), with zones near-olive-gray (5Y4/1) at 662, 665, and 669 m.
684.0	2.4	Tuff mixed with mud, grading down into pure tuff; impure tuff is olive-gray (5Y4/1), pure tuff yellowish-gray (5Y7/2); well indurated in basal 40 cm.
690.4	6.4	Mud, silt- to sand-size; mottled, ranging in color from olive gray (5Y4/1) to dark yellowish brown (10YR4/2)
690.9	0.5	Tuff, yellowish-gray (5Y6-8/1), well-indurated, crossbedded.

Depth to base of unit (m)	Thickness of unit (m)	Description
693.4	2.5	Mud and sand; faint to conspicuous thin beds, light-olive-gray (5Y5/2), with streaks of pale-brown (5YR5/2). Contact between Unit I of Mixed Layer and alluvial sand and gravel is at base of this interval.
915.3	211.9	Pebbly arkosic sand and gravel, most commonly moderate brown (5YR3-4/4), with zones that average light brown (5YR6/4) in color between 726-740 and 748-798 m; mostly coarse to very coarse sand, poorly sorted, containing quartz monzonite and volcanic-rock fragments, as large as 15 cm in diameter; faintly bedded to massive; not cored between 748.3-793.4, 804.7-826.3, and 839.1-903.1 m.
929.6	14.3	Quartz monzonite, light- to medium-gray (N5-7), with pale-brown (5YR5/2) stains along fractures extending through cored interval; rock bit was used from 915.3 to 926.3 m, and so no core was recovered.

DEPTH (ft) vs LITHOSTRATIGRAPHY

AGE (Ma) vs LITHOSTRATIGRAPHY

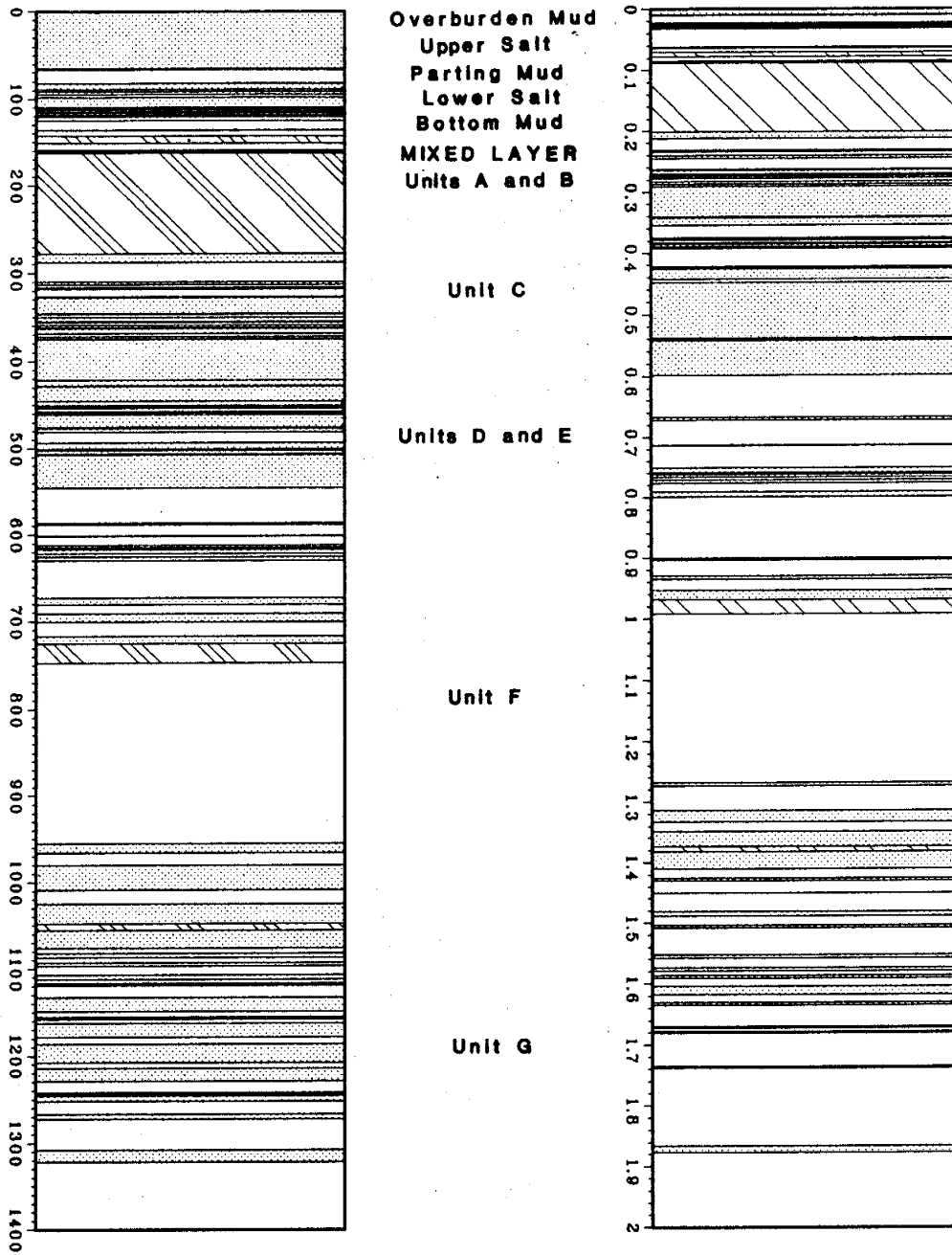


Figure 3 A comparison of core KM-3 stratigraphy with depth and with age. Cross-hatching indicates poor core recovery; stippling, intervals of predominantly evaporite minerals; and absence of pattern, predominantly lacustrine mud.

Mineralogy The buildup of solutes in Searles Lake water provided an unusual environment for the accumulation of thick sequences of evaporite minerals. These minerals, and the accompanying brines, have been the target of mining companies at Searles Lake for the last 100 years. A list of evaporite minerals and the diagenetic products of some of the minerals (Smith, 1979) is shown in table 2. Of greatest interest, for the purposes of this paper, are primary and diagenetic carbonates. Those carbonates assumed to be of primary origin are dolomite, calcite, aragonite, and northupite. Gaylussite and pirssonite are assumed to be of diagenetic origin, although fine-grained gaylussite may be primary.

Assumptions of diagenetic or primary origin of these carbonates are crucial in attempting to reconstruct lake history by stable isotope records. If isotopic exchange had occurred with interstitial brines, or by any other mechanism, paleohydrologic information from the isotopic record may be lost. Therefore, a brief discussion of the evidence leading to conclusions of primary or diagenetic origin is warranted.

The mineral of greatest interest in this paper is dolomite. Smith and others (1983) reported dolomite to exist throughout most of core KM-3, and its abundance and frequent occurrence makes the mineral very attractive in constructing a detailed and consistent isotope record at

Table 2. Names and composition of the non-clastic minerals found in the KM-3 core. (Smith and others, 1983).

<i>Mineral</i>	<i>Composition</i>
Analcime	$\text{NaAlSi}_2\text{O}_6 \cdot \text{H}_2\text{O}$
Anhydrite	CaSO_4
Aragonite	CaCO_3
Borax	$\text{Na}_2\text{B}_4\text{O}_7 \cdot 10\text{H}_2\text{O}$
Burkeite	$2\text{Na}_2\text{SO}_4 \cdot \text{Na}_2\text{CO}_3$
Calcite	CaCO_3
Celestite	$(\text{Sr}, \text{Ba}) \text{SO}_4$
Dolomite	$\text{CaMg}(\text{CO}_3)_2$
Gaylussite	$\text{CaCO}_3 \cdot \text{Na}_2\text{CO}_3 \cdot 5\text{H}_2\text{O}$
Glauberite	$\text{Na}_2\text{SO}_4 \cdot \text{CaSO}_4$
Gypsum	$\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$
Halite	NaCl
Hanksite	$9\text{Na}_2\text{SO}_4 \cdot 2\text{Na}_2\text{CO}_3 \cdot \text{KCl}$
Heulandite	$\text{CaO} \cdot \text{Al}_2\text{O}_3 \cdot 6\text{SiO}_2 \cdot 5\text{H}_2\text{O}$
Magnesite	MgCO_3
Nahcolite	NaHCO_3
Northupite	$\text{Na}_2\text{CO}_3 \cdot \text{MgCO}_3 \cdot \text{NaCl}$
Pirssonite	$\text{CaCO}_3 \cdot \text{Na}_2\text{CO}_3 \cdot 2\text{H}_2\text{O}$
Searlesite	$\text{NaBSi}_2\text{O}_6 \cdot \text{H}_2\text{O}$
Thenardite	Na_2SO_4
Tincalconite	$\text{Na}_2\text{B}_4\text{O}_7 \cdot 5\text{H}_2\text{O}$
Trona	$\text{Na}_2\text{CO}_3 \cdot \text{NaHCO}_3 \cdot 2\text{H}_2\text{O}$

Searles. Smith (1979) suggested a primary origin for dolomite based on several factors: 1) the mineral occurs only in microscopic form, 2) it is found in lake bed outcrops surrounding the valley floor, where other minerals of diagenetic origin are missing, 3) dolomite occurs in distinct zones, suggesting it occurred only when the lake chemistry was conducive to dolomite formation, 4) the magnesium content of interstitial brines is very low, and other minerals occurring with dolomite contain very small amounts of magnesium; the source of magnesium during the possible alteration of calcium carbonate to dolomite is not readily apparent, and 5) dolomite does not occur in areas where minerals present could react diagenetically to form dolomite.

That dolomite occurs as a primary evaporite mineral in some lakes has been proven by several investigators (Clayton and others, 1968; De Decker and Last, 1988). Of great interest is the evidence presented by Clayton and others (1968), following Jones (1965), of modern dolomite formation in Deep Springs Lake, California. Smith (1979) suggested that current dolomite formation in Deep Springs may be a modern analog of dolomite formation in Searles Lake, as the two lakes are presumed to be chemically very similar. Calcite in core KM-3 is also assumed to be of primary origin. Calcite occurs less frequently than dolomite in core KM-3, yet is predominant in a few sections of the core,

most notably in the Parting Mud and very deep in the core (Units G through I). Calcite is well known as a primary precipitate from lake waters, and no compelling evidence suggests a diagenetic origin. However, Smith (1979) strongly suggested that some of the calcite may be recrystallized metastable aragonite.

Aragonite occurs only in the upper stratigraphic units, mainly the Parting Mud and the Bottom Mud. As aragonite is thermodynamically metastable near the earth's surface, any aragonite that may have been precipitated earlier in the lake's history has probably been altered to calcite, or reacted with the interstitial brines to form gaylussite or pirssonite. No evidence suggests that the aragonite present in KM-3 is of any but primary origin.

Megascopic and microscopic northupite occurs in discrete intervals throughout most units of the Mixed Layer. Primary northupite is thought to have originated by the mixing of magnesium-bearing waters with concentrated sodium carbonate and chloride brines (Smith and Haines, 1964). However, Smith (1979) suggested that microcrystalline northupite may have a diagenetic origin in some sections of the core. As no northupite/water fractionation factor has been published, northupite was not used in the stable isotope analysis of core KM-3.

Several carbonate minerals are considered to be of at least partially diagenetic origin in Searles Lake, including

gaylussite and pirssonite. The alteration of calcium carbonate by percolating sodium and carbonate-rich brines is suggested by the inverse relative proportions of the diagenetic carbonates to the primary calcium carbonates (Smith, 1979). Textural evidence, such as growth of macrocrystalline gaylussite across mud laminae without deformation of the laminae suggests that the diagenetic minerals formed after compaction of the lake sediments. Gaylussite is known to occur as a primary mineral in Owens Lake (Smith and others, 1987), and disseminated crystals in the sediment may be of primary origin.

Brines The interstitial brines have been classified by Eugster and Hardie (1978) as being of the Na-CO₃-SO₄-Cl type. The origin of the brines has been discussed at length in Eugster and Hardie (1978). Figure 4 graphically displays the chemical evolution of the brines at Searles.

Hardt and others (1973) described the relationship of the relatively fresh water in the alluvial fans to the brines. The authors showed that the hydrostatic head of the brines is, and perhaps generally has been, higher than the head in the surrounding freshwater aquifers. Brine movement is therefore down, through the lake sediments, and possibly eventually moving out laterally from the basin (Friedman and others, 1982).

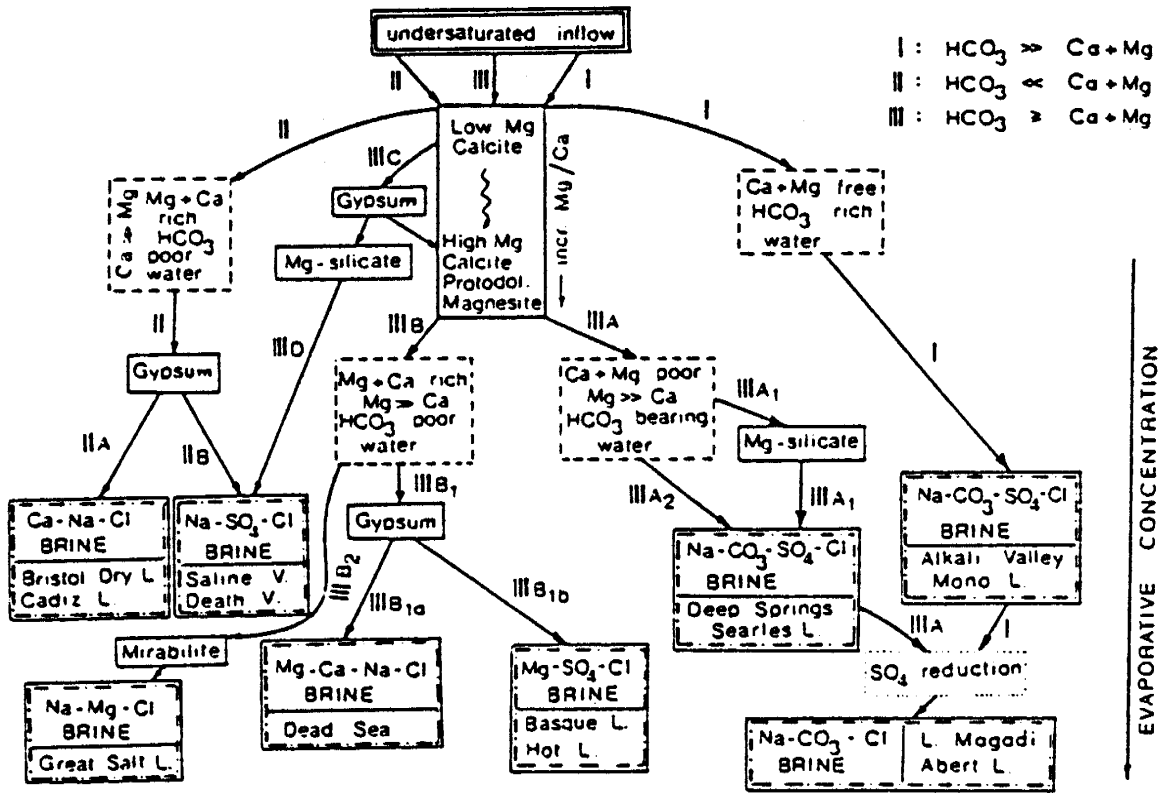


Figure 4 Water quality path of Searles Lake brines during evaporative concentration (Eugster and Hardie, 1978).

IV. CHRONOLOGY DEVELOPMENT

Carbon-14

During the late 1950's and 1960's, numerous investigators used radiocarbon techniques to obtain numerical age-estimates of lake sediments at Searles Lake (Flint and Gale, 1958; Stuiver, 1964; Stuiver and Smith, 1979). Eighty-eight radiocarbon dates for sediment in the Overburden Mud, Upper Salt, Parting Mud, Lower Salt and the Bottom Mud have been published; a summary of these dates can be found in Benson and others (1989). Materials used for radiocarbon age-estimates include wood, disseminated organic carbon, and primary and diagenetic carbonate minerals.

The reliability of the materials for radiocarbon dating at Searles has been debated at length (Benson and others, in press; Stuiver and Smith, 1979). Potentially diagenetic carbonate minerals are not considered to be useful except as a supplement to more suitable materials. Briefly, the likely contamination of diagenetic carbonates with carbon from interstitial brines casts considerable doubt on the reliability of these minerals for radiocarbon dating.

Debate over the reliability of radiocarbon dates from primary carbonates is due to several factors. First, non-equilibrium of the $^{14}\text{C}/^{12}\text{C}$ ratio between the lake water and atmosphere, the "reservoir effect", can cause the lake water to have an apparent age of hundreds to thousands of years

(Broecker and Walton, 1965; Peng and others, 1978; Stuiver and Smith 1979). Therefore, radiocarbon age-estimates of carbonates precipitated in isotopic equilibrium with lake water will be too "old". Second, re-crystallization or the isotopic re-equilibration of carbonate minerals with interstitial brines may have occurred. Depending upon the source of the brine, either from above or below the sediment, the radiocarbon ages could be too "old" or too "young". However, the possibility that isotopic re-equilibrium or exchange with interstitial brines has occurred cannot be ruled out. A third source of error in the radiocarbon age-estimates of inorganic carbonates could be the assumption that the carbonate minerals were precipitated in isotopic equilibrium with the lake water. This problem is most likely to occur at low lake levels where kinetic effects due to supersaturation of carbonates can produce non-equilibrium precipitates (Stiller and others, 1985). However, resulting errors in radiocarbon ages are likely to be small in comparison with other sources of error. Finally, Stuiver and Smith (1979) list substantial evidence that possible reworking of older sediments has occurred, especially in the Overburden Mud and the Upper Salt. Re-deposition of older sediments could lead to ^{14}C dates that are too "old". Further, the dissolution of dead carbon into the lake would also cause precipitating carbonates to appear too "old".

Disseminated organic radiocarbon dates would be affected by many of the same processes as inorganic carbonates. The $^{14}\text{C}/^{12}\text{C}$ ratio of organisms utilizing dissolved carbon in the lake water will reflect the "apparent age" of the water. Similarly, isotopic exchange with interstitial fluids will produce errors in the radiocarbon age-estimates of organic sediments. Finally, bacterial contamination of wet-sediment cores may produce radiocarbon age-estimates of organic sediments that are too "young" (Geyh and others, 1974 in Benson and others, in press).

Radiocarbon age-estimates on wood are probably the most reliable (Stuiver and Smith, 1979). Only two wood dates are available at Searles Lake: one in the Overburden Mud and one in a lower mud unit in the Lower Salt. No evidence to cast doubt the reliability of these dates has been put forward.

Inorganic carbonate minerals are least reliable for radiocarbon dating due to "reservoir effects", possible isotopic exchange with interstitial fluids, and re-deposition of older sediments. Stuiver and Smith (1979) estimate errors associated with the ^{14}C dates of carbonate minerals range from ± 500 to ± 2500 years, in addition to reported counting errors. Estimated errors for disseminated organic dates range from 500-2500 years too "old", in addition to counting errors. As there are sufficient disseminated organic carbon and wood dates available, the

inorganic dates will not be considered further in this paper.

Uranium/Thorium

To expand the absolute-age chronology of Searles Lake sediments, Peng and others (1978) compared radiocarbon dates of Lower Salt muds (Stuiver and Smith, 1979) with ^{230}Th dates on the surrounding salt beds. Nineteen ^{230}Th dates were published on halite and trona samples from two cores. The uranium-thorium dates were in general agreement with the radiocarbon dates. This fact encouraged further uranium-series dating of the Bottom Mud and Mixed Layer Unit A+B by Bischoff and others (1985). An additional nineteen ^{230}Th dates on trona, gaylussite and dolomite were published, extending the absolute-age chronology to approximately 230 ka.

The reliability of the ^{230}Th age-estimates depends upon how well the technique satisfies two basic assumptions: 1) all ^{230}Th present in the sample resulted from post depositional in situ decay of uranium (it is assumed that there is no non-radiogenic ^{230}Th present), and 2) the sediment has remain a geochemically closed system; there has been no gain/loss or exchange of uranium or thorium (Ku and Liang, 1984). In Peng and others (1978), several additional assumptions were made in calculating the ^{230}Th age. First, an estimated correction factor for non-radiogenic ^{230}Th was

arrived at by assuming that the initial $^{230}\text{Th}/^{232}\text{Th}$ ratio was the same for all salt units in the Lower Salt. Second, it was assumed that the salt units had been precipitated in an extremely short amount of time.

Varying degrees of error are introduced by the above assumptions. Fluctuations in initial $^{230}\text{Th}/^{232}\text{Th}$ ratios are expected. Isotopic exchange with interstitial brines is possible, though difficult to determine. Age-estimates of the Lower Salt based on the accumulation of acid-insoluble residue (Jannik, 1989) support the assumption that the salt units were deposited in less than 1000 years and possibly less than 100 years.

A potential problem of post-depositional contamination also exists with the age-estimates presented by Bischoff and others (1985). Finally, because intervals of core loss were not always noted during drilling, there may be uncertainty concerning the sample depth. With all possible errors in mind, the counting errors associated with the ^{230}Th dates for Searles Lake salts represent only a minimum error estimate.

Magnetostratigraphy

Interpretation of paleomagnetic polarities from Searles Lake core KM-3 allowed development of a magnetostratigraphy for the paleolake sediments (Liddicoat and others, 1980). Using the revised time scale of Mankinen and Dalrymple (1979), the deepest lacustrine sediments were correlated

with the base of the Mammoth subchron of the Gauss chron (3.15 Ma). The youngest boundary defined was the Brunhes-Matuyama (~730 ka). A total of twenty-four polarity boundaries were described, including five zones that were not included in the Mankinen and Dalrymple time scale.

Possible interpretation errors due to core sections reversed in the core boxes were discussed by the investigators. Only deep in the core, beyond the maximum depth of stable isotope analysis, was this problem suspected to exist. Reported errors associated with zone definition ranged from ± 0.4 m to ± 10.6 m. Using an average sedimentation rate for the core of 45 yr/cm, a minimum age error of ± 1.8 kyr to ± 45.5 kyr may be assumed for these paleomagnetic age-estimates.

Tephrochronology

Series of ash units were deposited in Searles Lake sediments as a result of widespread Cenozoic volcanic activity. Fourteen tephra layers were identified in core KM-3 (Hay and Guldman, 1987). One ash unit, at ~168.6 m, was tentatively identified, and was correlated with the Lava Creek B (formerly 'Pearlette Type O') ash. The most recent age-estimate for the Lava Creek B ash is 610 ka (J.D. Obradovich, written communication in Izett, 1981). The age is based on a mean of several K-Ar dates. No error associated with this date has been published. Fission-track

ages of zircons within the Lava Creek B ash from several localities are 0.6 ± 0.07 Ma (Naeser and others, 1973) and 0.6 ± 0.15 Ma (Christiansen and Blank, 1972 in Naeser and others, 1973). These ages are in close agreement with the K-Ar date of 0.610 Ma.

The correlation of the tephra unit in Core KM-3 with the Lava Creek B ash was made partially due to its stratigraphic position relative to the Brunhes-Matuyama paleomagnetic boundary. There appears to be some circular reasoning with this correlation, as the paleomagnetic boundary was originally identified partially based on its proximity to this tephra unit, then assumed to be the Bishop Tuff (Liddicoat and others, 1980). However, additional mineralogic information supports the identification of this ash unit as the Lava Creek B.

Chlorine-36

Absolute age-estimates based on the decay of ^{36}Cl in halite have extended the chronology of sediments at Searles Lake. Six ^{36}Cl dates on halite units from 14.2 m to 401.3 m in depth ranged from 10 ka to 1.95 Ma (Phillips and others, 1983). The ^{36}Cl dates were in general agreement with radiocarbon, uranium-series, and paleomagnetic age-estimates of the same units. Based on the success of Phillips and others (1983), Jannik (1989), determined an additional fourteen ^{36}Cl age-estimates on halite units in core KM-3

from 3.0 m to 401.3 m in depth. ^{36}Cl dates on these new samples ranges from 6.7 ka to 1.27 Ma. The deepest sample at 401.3 m (1.95 Ma) had previously been suspected of contamination by Phillips and others (1983), and the new date (1.25 Ma) of this depth confirmed earlier suspicions. A summary of all ^{36}Cl age-estimates can be found in Jannik (1989).

As the ^{36}Cl method is somewhat new, a brief discussion of ^{36}Cl theory as it applies to the Searles Lake sediments is warranted. The hydrophilic nature of chlorine allows the chloride ion to travel through a hydrologic system with minimal chemical interaction. As Searles Lake was the frequent terminus of the Owens River paleolake system, chloride accumulated in the lake water and eventually precipitated out in evaporite minerals during arid periods. Upon deposition, the decay of ^{36}Cl began.

The various sources of ^{36}Cl in the evaporite minerals are discussed in great detail in Jannik (1989). Briefly, there are three main sources of ^{36}Cl . First, the initial ^{36}Cl input to a hydrologic system is through the inclusion of atmospheric-produced ^{36}Cl in precipitation. Second, surface production of ^{36}Cl in rocks is caused by cosmic-ray spallation of calcium and potassium and neutron activation of ^{35}Cl . The weathering of these rocks will add ^{36}Cl to surface or groundwater systems. Third, post-deposition production of ^{36}Cl due to activation of ^{35}Cl by cosmic-ray

neutrons, in addition to neutron activation by the decay of uranium and thorium in the sediments, can occur, increasing the $^{36}\text{Cl}/\text{Cl}$ ratio in the evaporite minerals. Phillips and others (1983) estimated the $^{36}\text{Cl}/\text{Cl}$ input from the three different sources for the halite units in the upper Salt. Approximately thirty percent of the ^{36}Cl in the halite is derived from atmospheric sources, about seventeen percent from subsurface production, and apparently more than fifty percent from surface production.

In addition to the production of ^{36}Cl by the three sources mentioned above, several other site-specific sources or sinks of chloride bear mentioning. The mixing ratios of the various source waters, especially thermal springs in the Owens Valley, can alter the final $^{36}\text{Cl}/\text{Cl}$ ratio in the Owens River water and its related paleolakes. Second, the inflow/overflow/desiccation history of each lake will affect the $^{36}\text{Cl}/\text{Cl}$ ratio of evaporite minerals precipitating out of a given lake.

In light of all the possible contributions of ^{36}Cl to the eventual $^{36}\text{Cl}/\text{Cl}$ ratio of the Searles Lake halite units, the determination of age-estimates based on the ^{36}Cl method depends upon several conditions. First, the $^{36}\text{Cl}/\text{Cl}$ ratio of inflow waters at Searles Lake was constant or known. Second, subsurface production of ^{36}Cl is assumed to be negligible. And finally, no post-deposition transfer of chloride to or from the halite crystal has occurred. The

general agreement of the ^{36}Cl age-estimates with other radiocarbon, uranium-series, and paleomagnetic age-estimates support these assumptions. However, a few ^{36}Cl measurements from Searles Lake showed abnormally high $^{36}\text{Cl}/\text{Cl}$ ratios. Jannik (1989) suggested that some sort of laboratory contamination had taken place, yet she did not rule out other sources of error.

Analytical errors on the Searles Lake ^{36}Cl dates ranged from ± 1.0 kyr in shallow samples to ± 176 kyr for deep samples. As with all other methods described previously, these analytical errors represent only a minimum error.

AIR-Interpolated Ages

With a detailed absolute chronology established, assigning ages to samples between absolutely-dated points usually involves a linear interpolation of ages based on an assumption of constant sedimentation rates. However, large variations in sedimentation rates at Searles Lake are expected due to the extreme fluctuations between deep-water muds and shallow-water evaporite sequences. Therefore, Jannik (1989) developed a technique to minimize the errors associated with interpolated ages. This technique is based on evidence for a relatively constant accumulation of acid-insoluble residue (AIR) in lake sediments.

The variability of sedimentation rates in Searles is largely due to the accumulation of solutes in the lake water

during non-overflow periods of the lake's history. During low lake level or desiccation episodes, accumulated solutes precipitated out of the water rapidly as thick sequences of evaporite minerals. Conversely, the AIR component of inflow waters had a relatively short residence time in the lake water, and therefore accumulated at a relatively constant rate through time.

Results from detailed geochemical analyses of core KM-3 were presented by Smith and others (1983). The core was divided up into 254 intervals according to lithology; geochemical analysis, including determining of acid-insoluble residue (AIR), was performed on each interval. Jannik (1989) calculated cumulative AIR versus time and concluded that the accumulation of the AIR component of the sediments varied smoothly through time. A plot of "AIR vs. Depth" displayed a "stair-step" pattern reflecting the change in sedimentation rates from pluvial (low sedimentation) to more arid (high sedimentation) periods. AIR data were assigned ages by linearly interpolating between absolutely-dated points. The AIR ages were then correlated with AIR depths to create "AIR-interpolated ages" for the lithologic intervals. Analyses of some of the intervals were combined due to similar geochemical characteristics. A total of 144 intervals were assigned AIR-interpolated ages.

It should be emphasized that the AIR values are merely

a method for interpolating ages between absolute radiometric and paleomagnetic dates. The purpose of using the AIR-interpolation procedure (instead of depth interpolation) is to compensate for drastic fluctuations in sedimentation rate caused by alternation of evaporite and marl-depositing conditions. A very high deposition rate for the Upper Salt can be demonstrated by ^{14}C dating (discussed above). It seems reasonable to infer that similar evaporites deeper in the core were deposited under similar conditions. The AIR interpolation avoids unrealistically long deposition times for these evaporites. Errors assumed for this technique are difficult to quantify. The AIR technique has, however, substantially decreased errors that would have resulted had a simple linear interpolation between absolutely-dated points been used.

Composite Chronology

With all the possible errors in mind using absolute dating techniques, the chronology for Searles Lake is remarkably detailed, especially in the upper units. Following Jannik (1989), a composite chronology of core KM-3 is presented below according to the informal (but generally accepted) stratigraphic unit names. Slight revisions of the Jannik chronology have been made, and a discussion of these revisions are included in the following section.

Overburden Mud Evidence presented by Stuiver and Smith (1979) indicates most of the mud in this unit is reworked older sediment. Therefore, no dates on carbonate minerals or organic sediments are thought to be reliable. A ^{36}Cl date of 67 ka \pm 10 kyr is clearly much too old and also will not be used. The only reliable date available is a ^{14}C date on wood at 2.4 m in depth with an age of 3.52 ka \pm 0.19 kyr. One AIR-interpolated age at the Overburden Mud/Upper Salt boundary implies an age of 6 ka.

Upper Salt Twelve radiocarbon dates (Stuiver and Smith, 1979) for this unit are from inorganic carbonates and are considered unreliable. Four radiocarbon dates on disseminated organic carbon ranged from 11.4 ka \pm 0.6 kyr near the top of the unit to 9.72 ka \pm 0.2 kyr at the base of the unit. All other dates fell within the 10 ka to 11 ka range for the bottom 80% of the unit.

As discussed by previous investigators, (Stuiver, 1964; Benson and others, in press) the reversals in age and the small range of ages throughout the bottom 80% of the Lower Salt indicate rapid precipitation of this salt unit. The AIR data also support this conclusion.

Two U-Th age-estimates (Peng and others, 1978) near the top and base of the Lower Salt zone gave "corrected" ^{230}Th ages of 5.3 ka \pm 1.2 kyr and 8.7 ka \pm 0.2 kyr respectively. The age-estimates are in general agreement with AIR and ^{14}C

dates.

Again, as stable isotope analysis was not made on samples above the Parting Mud, age-estimates for the Upper Salt are not crucial to this report. I have used an approximate average of the ^{14}C values for the Parting Mud/Upper Salt boundary, of 10 ka, which is in general agreement with all other age-estimates. AIR dates for locations within the unit have been used as they are stratigraphically consistent while the radiocarbon dates are not.

Parting Mud Thirty-nine ^{14}C dates for the Parting Mud are summarized in Stuiver and Smith (1979); fourteen radiocarbon dates were from inorganic carbonates and, therefore, unreliable. Twenty-five radiocarbon dates from disseminated organic carbon were presented; the radiocarbon age-estimates on these samples range from 10.06 ka ± 90 yr to 10.9 ka ± 90 yr at the top of the unit, and 22.4 ka ± 1.0 kyr to 24.69 ka ± 1.07 kyr at the base. Following Stuiver and Smith (1979), an age-estimate of 23 ka for the base of the Parting Mud is chosen here, although the age could range from ~21 ka to 24 ka.

Lower Salt Twenty-seven radiocarbon dates for this unit are summarized in Stuiver and Smith (1979): seven are from inorganic carbon, nineteen are from disseminated organic

carbon, and one radiocarbon date on wood is available. As before, radiocarbon-dates on inorganic carbonates are probably unreliable and will not be considered further. The uppermost radiocarbon date is on sub-unit M-7, with an age estimate of 23.75 ka \pm 400 yr. The deepest reliable date is from sub-unit M-2, with an age-estimate of 32.6 ka \pm 500 yr. Sub-unit M-3 is the most uncertain, with two disseminated organic radiocarbon dates of 32.0ka \pm 1 kyr and 29.5 ka \pm 2 kyr. The wood radiocarbon date for M-3 is 26.7 ka \pm 2 kyr. Between M-6 and M-2, several radiocarbon dates are stratigraphically inconsistent. Detailed AIR dates are available in the Lower Salt units; these data again support conclusions by previous authors that most of the salt sub-units were deposited in a very short amount of time.

Eighteen U-Th age-estimates are available for the salt sub-units of the Lower Salt; sixteen "corrected" ^{230}Th dates were published in Peng and others (1978) and one ^{230}Th age-estimate for the S-1 submit was presented in Bischoff and others (1985). A few ^{230}Th age-estimates were stratigraphically inconsistent. ^{230}Th dates ranged from 23.8 ka \pm 1.4 kyr for subunit S-7, to 33.0 ka \pm 1.5 kyr for sub-unit S-1 (Peng and others, 1978). The ^{230}Th age-estimate for S-1 in Bishoff and others (1985) was 35.3 ka $^{+2.7-2.4}$ kyr.

Selecting a chronology for the Lower Salt was problematic. Overall, both ^{14}C and ^{230}Th dates were in general agreement. However, data that were

stratigraphically reversed, in addition to doubts concerning "reservoir effects" and Peng's ^{232}Th corrections, made choosing the best dates difficult. Therefore, I have chosen to assign a "probable" age to the Lower Salt/Bottom Mud boundary of 33.0 ka. This age is within the reported confidence intervals of all ^{14}C and ^{230}Th age-estimates. Using the "probable age" of the LS/BM boundary of 33 ka, the "probable age" of the PM/LS boundary of 23 ka, and AIR data, I interpolated a chronology for the subunits of the Lower Salt. In general, the new AIR interpolated-ages for the subunits are in agreement with ^{14}C and ^{230}Th age-estimates. A problem immediately apparent with this AIR-interpolated chronology is that the AIR ages for the submit M-3 boundaries are not within the confidence interval of the radiocarbon age-estimate on the wood from submit M-3, presumably the most reliable date in the Lower Salt. Therefore, the confidence intervals for assigned sample ages within this unit should probably be as large as ± 5 kyr.

Bottom Mud Four ^{14}C age-estimates are available for the Bottom Mud; three of the four dates are on disseminated organic carbon. Two of the three organic dates are near the top of the Bottom Mud, and the previously assigned "probable age" of the LS/BM boundary is within the confidence interval of both dates.

Five ^{230}Th age-estimates for the Bottom Mud were

presented in Bischoff and others (1985). Ages ranged from 58.4 ka $^{+1.5-1.1}$ kyr near the top of the Bottom Mud, to 122.3 $^{+3.6-3.0}$ kyr approximately 30% above the base of the unit.

AIR data are available for the Bottom Mud. Of greatest interest is one point at the base of the unit. If the first ^{230}Th date of 58.4 ka is incorporated into the chronology, a sedimentation rate of less than 1 cm/100 yr is inferred from the LS/BM boundary to this sample. This sedimentation rate is more than an order of magnitude less than the rates in the intervals above and below. Therefore, only the lower four ^{230}Th age-estimates are used. AIR data are also incorporated into the Bottom Mud chronology. An age-estimate of 153.8 ka is interpolated for the BM/Mixed Layer boundary.

Mixed Layer: A+B Four ^{230}Th age-estimates for the A+B unit were presented by Bischoff and others (1985). The A+B/C boundary was dated by the ^{36}Cl method to yield an age of 286 ka ± 47 ka (Jannik, 1989). Seven AIR intervals are also available for this unit.

I have chosen not to use two of the four ^{230}Th age-estimates available for the A+B unit. The uppermost sample (Bischoff sample 7) yields an extremely low calculated sedimentation rate. The deepest ^{230}Th date (Bischoff sample 11) also yields a low sedimentation rate, and has a very large analytical uncertainty. Two ^{230}Th ages were available

for Bischoff sample 8, and, following Jannik (1989), I have averaged the two dates for an age-estimate of 160.5 ka at 70.86 m in KM-3. All AIR data were used.

Mixed Layer: Unit C Four ^{36}Cl age-estimates and twenty six AIR-interpolated ages are available for Unit C (Jannik, 1989). All available data were incorporated into the unit C chronology. The AIR interpolated age at the Unit C/D+E boundary is 598 ka.

Mixed Layer: Units D+E Four ^{36}Cl age-estimates are available for unit D+E (Phillips and others, 1983; Jannik, 1989). One of the four (SLC-6) represented a reversal, and therefore, was not included in this chronology. An ash layer at 168.6 m was correlated with the Lava Creek B ash (610 ka) (Hay and Guldman, 1986). The first three paleomagnetic boundaries in KM-3 were defined in the D+E unit (Liddicoat and others, 1980): the Brunhes-Matuyama (730 ka) at 185 m, the Matuyama-Jaramillo (900 ka) at 204.9, and the Jaramillo-Matuyama (970 ka) at 221.2 m. Thirteen AIR ages are also available in D+E (Jannik, 1989).

All available age-estimates, with the exception of the one ^{36}Cl date previously mentioned, are incorporated into the D+E chronology presented in this paper. No data are available for the D+E/F boundary, but a linear interpolation between the last absolutely-dated point at 221.2 m (970 ka)

and the AIR age below the boundary (1.06 Ma at 248.11 m) gives an approximate age of 992 ka for the D+E/F boundary.

Mixed Layer: Unit F Only seven AIR-interpolated ages (Jannik, 1989) are available for Unit F. The deepest AIR interval, located at the F/G boundary, gives an approximate age of 1.268 Ma for the stratigraphic boundary. Two paleomagnetic boundaries were defined in Unit F, but as they are not included in the Mankinen and Dalrymple (1979) time scale, they will not be used in this chronology.

Mixed Layer: Unit G This is the deepest stratigraphic unit presented in this report that was sampled for stable isotope analysis. One ^{36}Cl date and thirty-four AIR-interpolated ages are available for this unit (Jannik, 1989). Six paleomagnetic ages are also available (Liddicoat & others, 1980). Two of the six paleomagnetic ages are undefined boundaries, and therefore will not be used in this chronology. The four defined paleomagnetic boundaries that are used are 1) the Matuyama-Olduvai (1.67 Ma) at 378.6 m; 2) the Olduvai-Matuyama (1.87 Ma) at 403.25m; 3) The Matuyama-Reunion (2.01 Ma) at 408.3 m; 4) and the Reunion-Matuyama (2.04 Ma) at 424.7 m near the G/H stratigraphic boundary.

The ^{36}Cl date, the four previously mentioned paleomagnetic dates and all AIR ages are used in Unit G

chronology. An approximate age of the G/H stratigraphic boundary is 2.04 Ma.

Composite Chronology All ^{14}C , ^{230}Th , ^{36}Cl , AIR-interpolated and paleomagnetic age-estimates used in this chronology are listed in Appendix A. Graphical representations of KM-3 stratigraphy vs. core depth and age are shown in figures 5 and 6. KM-3 mud samples prepared for stable isotope analysis were assigned ages using linear interpolation between absolutely dated isotope, AIR and paleomagnetic points.

CORE KM-3 CHRONOLOGY: UPPER UNITS
Revised from Jannik (1989)

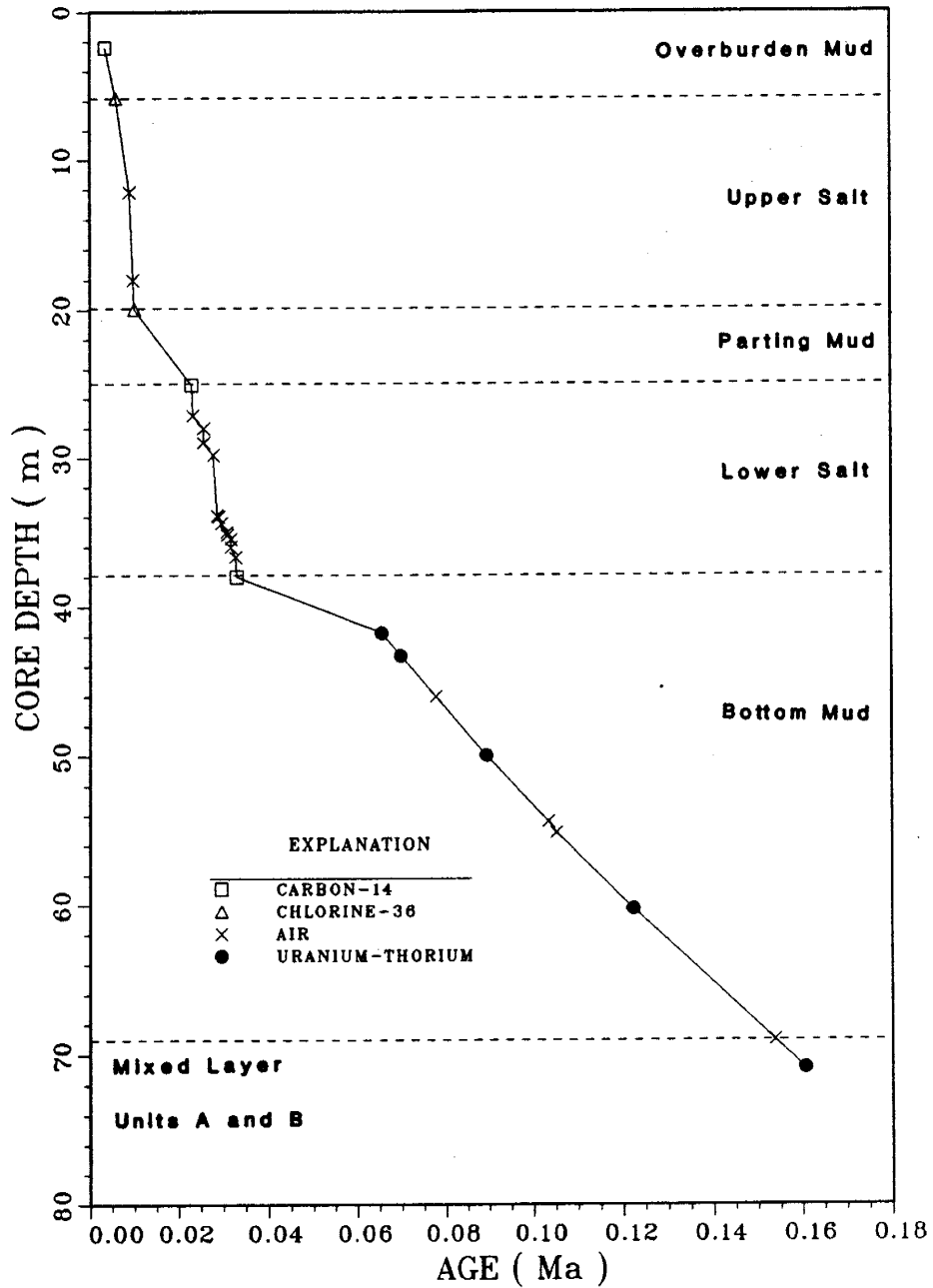


Figure 5 Age versus depth relationship for upper portion (0-80 m depth) of core KM-3, using carbon-14, thorium-230, and chlorine-36 dates and acid-insoluble residue (AIR) interpolation. Revised from Jannik (1989).

SEARLES LAKE CORE KM-3 CHRONOLOGY

Revised from Jannik (1989)

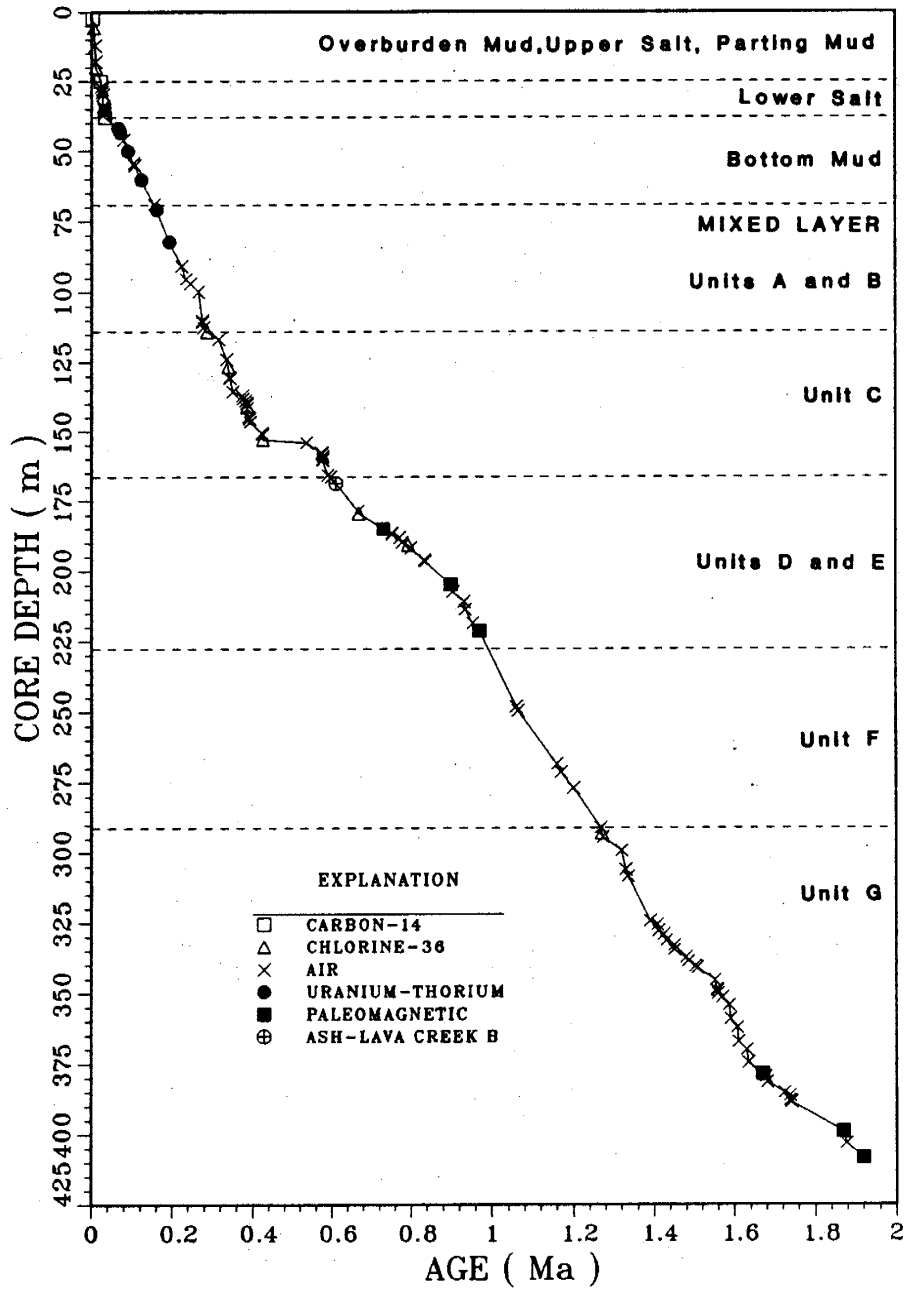


Figure 6 Age versus depth relationship for the upper 425 m of core KM-3, using carbon-14, thorium-230, chlorine-36, paleomagnetic dates, and acid-insoluble residue (AIR) interpolation. Revised from Jannik (1989).

V. METHODS AND PROCEDURES

Introduction

A detailed sampling scheme was developed in order to achieve a high time-resolution record of stable isotope variation in Searles Lake paleowaters. Care was taken in sampling the core to minimize location errors. Once the core was been sampled, the relative amounts and types of minerals were determined. Any substances which would interfere with mass spectrometer analysis of primary carbonates was removed from the samples. Following this, a methodology was developed to separate primary carbonates from each other, as each mineral has its own water-carbonate fractionation factor and phosphoric acid fractionation factor.

Sampling

247 samples were collected from core KM-3 mud units during October of 1986. The desired sampling frequency was approximately one sample every one to two thousand years. Using Smith and others' (1983) core log and calculated sedimentation rates as a guide, approximately fifty grams of sample were taken every one to two feet in most intervals from the Parting Mud through Unit G of the Mixed Layer. Sample descriptions and field notes are listed in Appendix B. Care was taken to remove large macroscopic salt crystals

and to scrape off the outside of the core to minimize contamination. Severe "telescoping" had occurred in several core boxes; the core boxes with this problem are noted in Appendix B. Sample depths are listed to the tenth of a foot, but this should not be interpreted to imply confidence in the depths to this degree, especially in intervals with severe "telescoping" problems.

Sample Preparation

Core samples contain a variety of primary and secondary authigenic minerals as described in an earlier section. In order to obtain a good stable isotopic analysis of the primary carbonate (principally dolomite), minerals which could interfere with the analysis of the carbonate had to be removed. The samples also contain detrital silicates; however, it was not necessary to remove the silicates as only carbonate minerals react with the phosphoric acid to release CO_2 in the acid extraction technique.

Both physical and chemical methods were used to achieve the separation of the primary carbonate from interfering minerals. First, macroscopic crystals of unwanted salts were removed from the sample by hand-picking. These crystals included secondary carbonate (gaylussite and pirssonite), bicarbonates, sulfates and halite. Halite needed to be removed from the samples as it was found to react with phosphoric acid to produce HCl gas; the HCl gas

creates severe problems during the mass spectrometry analysis.

Due to the much greater solubility of the unwanted minerals than the primary carbonates, water leaching was used as a second step to dissolve the remaining unwanted minerals. This procedure is described in Appendix C. X-ray diffraction (XRD) techniques were used to determine the success of the leaching procedure. Scanning parameters and XRD results are listed in Appendix B.

An unexpected problem with the leaching procedure was raised by the XRD results. Although the diagenetic carbonates identified by XRD in bulk samples dissolved, calcite appeared to have precipitated during the leaching process. An experiment was conducted to determine the origin of this calcite which I have dubbed "mongrel" calcite due to its uncertain parentage. Clean gaylussite and pirssonite crystals were placed in water to dissolve overnight. The solid collected by filtration was identified by XRD as calcite and confirmed that calcite had resulted from the dissolution of the two diagenetic carbonates. This phenomenon was previously observed by G.I. Smith (pers. comm). Whether the calcite was the result of incongruent dissolution or a later precipitate is not known.

A second problem with the leaching process was the occasional persistence of pirssonite and, more often, northupite. Repeated leaching could not remove all of these

AGE (Ma)	DEPTH (m)	DEPTH (ft)	DATE TYPE
0.57680	159.000	521.652	AIR
0.57700	160.000	524.933	AIR
0.57700	160.300	525.918	AIR
0.59000	165.800	543.962	AIR
0.59800	166.420	545.996	AIR
0.61000	168.600	553.148	ASH
0.66700	178.640	586.088	AIR
0.66800	179.410	588.614	CL-36
0.73000	185.000	606.954	Paleo-Mag
0.75100	186.540	612.007	AIR
0.75200	186.900	613.188	AIR
0.77100	188.150	617.289	AIR
0.77800	189.950	623.194	AIR
0.79200	190.700	625.655	CL-36
0.79900	192.020	629.986	AIR
0.83400	196.140	643.503	AIR
0.83500	196.660	645.209	AIR
0.89600	204.520	670.996	AIR
0.90000	204.900	672.243	Paleo-Mag
0.90400	207.450	680.609	AIR
0.93200	210.920	691.993	AIR
0.93500	213.660	700.983	AIR
0.95300	218.540	716.993	AIR
0.97000	221.200	725.720	Paleo-Mag
1.06000	248.110	814.008	AIR
1.06400	249.480	818.502	AIR
1.16000	268.530	881.002	AIR
1.17000	271.420	890.484	AIR
1.20000	276.910	908.496	AIR
1.26800	291.080	954.985	AIR
1.27000	293.000	961.284	CL-36
1.27300	294.440	966.009	AIR
1.32000	299.310	981.986	AIR
1.33000	306.020	1004.001	AIR
1.33500	308.460	1012.006	AIR
1.39000	324.310	1064.007	AIR
1.40800	325.530	1068.010	AIR
1.41000	327.960	1075.982	AIR
1.42400	329.180	1079.985	AIR
1.43000	331.320	1087.006	AIR
1.45000	333.150	1093.010	AIR
1.45000	334.670	1097.996	AIR
1.48000	337.260	1106.494	AIR
1.48500	338.630	1110.989	AIR
1.50300	340.310	1116.500	AIR
1.50800	341.070	1118.994	AIR
1.55200	345.340	1133.003	AIR
1.55500	348.690	1143.994	AIR
1.55700	349.150	1145.503	AIR
1.55800	350.220	1149.013	AIR
1.57000	351.430	1152.983	AIR
1.58600	354.180	1162.005	AIR
1.59000	359.050	1177.983	AIR
1.60700	362.100	1187.990	AIR
1.61000	367.130	1204.492	AIR
1.63000	370.030	1214.007	AIR
1.63500	374.750	1229.492	AIR
1.67000	378.560	1241.992	Paleo-Mag
1.67200	379.170	1243.994	AIR
1.67800	379.630	1245.503	AIR
1.68100	381.610	1251.999	AIR
1.72400	385.270	1264.007	AIR
1.73700	386.330	1267.484	AIR
1.73800	388.010	1272.996	AIR
1.74000	388.620	1274.997	AIR
1.87000	398.980	1308.987	Paleo-Mag
1.87700	403.250	1322.996	AIR
1.92000	408.300	1339.564	Paleo-Mag

Appendix B

Core Sampling Field Notes and X-Ray Diffraction Results

CORE DEPTH (ft.)	CORE DEPTH (m)	FIELD DESCRIPTION	XRD LEACHED SAMPLES	XRD BULK SAMPLES
72.1	22.18	mud, white/green lam white lam thicker than above;	I-aragonite, TR-calcite	
72.6	22.34	mud, dk green/white lam, white lam less frequent than above (several mm apart);	I-aragonite, TR-calcite	
73.1	22.49	mud, yellow-green/white lam, lam finer than 72.6;	I-aragonite, TR-calcite, TR-dolomite	
73.6	22.65	mud, green/white lam, lam as 72.6;	I-aragonite, TR-dolomite	
74.1	22.80	mud, lt green/white lam, white lam (~2 mm) parted from green;	I-aragonite, TR-calcite	
74.6	22.95	as 74.1; possible GAY xtals	I-aragonite, TR-dolomite, TR-calcite	
75.3	23.17	as 74.1;	I-aragonite, TR-calcite	
75.8	23.32	mud, lt grey-green, friable;GAY xtals	I-calcite, I-dolomite	
80.0	24.62	mud, lt grey, friable; GAY xtals **** HAND SAMPLE ****	I-calcite, I-dolomite	
81.0	24.92	mud, lt grey,dense;	I-calcite, TR-dolomite TR-pirssonite(?)	
81.6	25.11	mud, dk green,friable; large borax xtals, small (PIRS?) xtals	I-calcite, TR-dolomite TR-pirssonite(?)	
81.9	25.20	mud, dk green;borax xtals, large (PIRS?) xtals, sm trona (?)	I-calcite, TR-dolomite TR-pirssonite? TR-trona	M-pirssonite I-trona TR-dolomite
***** BOTTOM OF PARTING MUD *****				
89.5	27.54	mud, dk green; GAY(?) xtals	M-calcite	
90.7	27.91	mud, dk green lam; GAY xtals	M-calcite	
91.5	28.15	mud,dk green; mainly HAL(?)xtals,few large borax xtals	TR-calcite, TR-aragonite(?)	
95.5	29.38	mud, lt green-yellow lam; GAY xtals	I-calcite	
96.6	29.72	mud, dk green; mainly GAY xtals	M-calcite	I-gaylussite I-burkeite

CORE DEPTH (ft.)	CORE DEPTH (m)	FIELD DESCRIPTION	XRD LEACHED SAMPLES	XRD BULK SAMPLES
104.9-115.5		****PHOTOS****	trona and halite units	
111.7	34.37	mud, lt green finely lam; no salt xtals (above large trona seam)	I-calcite	
113.3	34.86	mud, lt green finely lam;(below trona seam)	TR-aragonite, TR-calcite	
114.5	35.23	mud, very lt green lam;	I-calcite, TR-aragonite	
115.9		****HAND SAMPLE****	Algal mats(?)	
116.4	35.82	mud, green, finely lam; some small borax xtals	M-calcite	
118.5	36.46	mud ,finely lam;	I-calcite, TR-aragonite	
120.0	36.92	mud, lt grey-green; small trona xtals	I-calcite	
***** TOP OF BOTTOM MUD *****				
124.9	38.43	mud, lt green/orange lam;	M-calcite, TR-dolomite	
125.7	38.68	mud, green lam; possible borax	M-calcite	
126.3	38.86	mud, green finely lam; coarse GAY xtals	I-calcite, I-dolomite	
126.9	39.05	mud, buff, lam; GAY xtals	I-dolomite, TR-calcite	
127.7	39.29	mud, dk green, lam; GAY xtals,HAND SAMPLE	I-calcite, TR-dolomite	
128.7	39.60	mud, dk green; ~95% GAY xtals	TR-dolomite, TR-calcite	
128.9	39.66	mud, buff/green lam; NO macroscopic GAY xtals	I-dolomite, TR-calcite, TR-aragonite(?)	
129.8	39.94	mud, lt green/buff lam; small GAY xtals	M-calcite,	
130.1	40.03	mud, lt green/buff finely lam;(above THEN seam)	I-dolomite, TR-calcite	
130.4	40.12	mud, buff,crudely lam; (below THEN seam)	M-calcite, I-dolomite	I-gaylussite I-dolomite
131.5	40.46	mud, dk green/buff lam;	I-aragonite, TR-calcite TR-dolomite	

CORE DEPTH (ft.)	CORE DEPTH (m)	FIELD DESCRIPTION	XRD LEACHED SAMPLES	XRD BULK SAMPLES
132.3	40.71	mud, dk green lam; ~80% GAY xtals	TR-dolomite, TR-calcite TR-aragonite	
134.4	41.35	mud, lt green, mass;	M-calcite, TR-dolomite	M-gaylussite TR-dolomite
136.4	41.97	mud, grey, mass; THEN(?) xtals (above THEN seam)	M-calcite, TR-dolomite	
137.5	42.31	mud, green-grey finely lam; (below THEN seam)	M-calcite, TR-dolomite	
139.0	42.77	mud, dk green/white lam;	I-aragonite, TR-calcite TR-dolomite	I-aragonite TR-calcite TR-dolomite
140.5	43.23	as 139.0, lam spaced further apart; coarse GAY xtals	M-dolomite, Tr-calcite	M-dolomite TR-calcite
141.8	43.63	mud, dk green/white lam, lam spaced further apart than 140.5;	I-aragonite, I-dolomite TR-calcite	I-dolomite I-aragonite TR-calcite
142.8	43.94	mud, dk green, finely lam;	I-dolomite TR-calcite TR-aragonite	
*****		LOTS OF TELESCOPING IN CORE BOXES (150-200 ft); DEPTHS DUBIOUS! ***** (MEASUREMENTS RECORDED TO TENTH OF FOOT DOES NOT IMPLY CONFIDENCE AT THIS LEVEL!!!!!!!!!!)		
151.1	46.49	mud, grey-green; mainly GAY xtals	I-dolomite, I-calcite	
153.0	47.08	as 151.1;	I-calcite	
154.1	47.42	as 151.1;	I-calcite	
158.6	48.80	mud, lt green, finely lam;	M-dolomite	
164.0	50.46	mud, lt green, mass;	M-dolomite	
170.0	52.31	mud, dk green; small salt(?) xtals	I-dolomite	
178.5	54.92	mud, green, mass;	TR-calcite, TR-dolomite	
183.0	56.31	mud, lt green, finely lam; few salt(?) xtals	I-calcite, I-dolomite	
187.4	57.66	mud, lt to dk green, finely lam; small salt(?) xtals	I-calcite, I-dolomite	
189.4	58.28	as 187.4; more salt(?) xtals	M-calcite, TR-dolomite	

CORE DEPTH (ft.)	CORE DEPTH (m)	FIELD DESCRIPTION	XRD LEACHED SAMPLES	XRD BULK SAMPLES
191.9	59.05	mud, dk grey-green, lam; GAY xtals	M-calcite	
194.9	59.97	mud, green, lam;	I-dolomite	
199.9	61.51	as 194.9;GAY xtals	I-dolomite, TR-calcite	
205.0	63.08	mud, green,lam; GAY xtals	M-calcite, I-dolomite	
211.1	64.95	mud, green; GAY xtals	I-calcite	
218.0	67.08	mud, dk green-black; few salt(?) xtals	I-dolomite	
221.0	68.00	mud, dk green/buff lam, lam 2-10mm a; small salt(?) xtals	M-dolomite	M-dolomite
224.1	68.95	as 221.0;	I-dolomite	
226.6	69.72	mud, dk green-black, finely laminated, orange stained;	I-dolomite	
256.1	78.80	mud, green-buff; salt(?) xtals	TR-dolomite, TR-calcite	
257.0	79.08	mud, lt green, orange stained; salt(?) xtals	I-calcite, TR-dolomite	
***** LIMITED CORE RECOVERY; DEPTH DUBIOUS!!!!*****				
277.5	85.38	mud, lt grey/dk green crumbled mud; salt(?) xtals	TR-dolomite, TR-calcite	
***** END OF LIMITED CORE RECOVERY *****				
288.1	88.65	mud, lt grey-green, mass; salt(?) xtals	TR-calcite	
298.1	91.72	as 288.1, less consolidated;	I-calcite	
308.0	94.77	mud, lt green, mass; small salt(?) xtals	I-calcite	
326.5	100.46	mud, lt green, orange-stained; - 50% salts	TR-calcite, TR-aragonite(?)	I-gaylussite TR-Northupite TR-aragonite?
349.9	107.66	as 326.5; more salt	I-northupite	I-northupite
359.3	110.55	as 326.5;	TR-calcite Unknown peaks 25-28 2-theta	
369.5	113.69	mud, lt green; small salt xtals	TR-aragonite TR-calcite? unknown peaks	

CORE DEPTH (ft.)	CORE DEPTH (m)	FIELD DESCRIPTION	XRD LEACHED SAMPLES	XRD BULK SAMPLES
370.5	114.00	mud, lt green, orange staining; small salt xtals	TR-calcite	
371.8	114.40	mud, lt green; trona	TR-calcite	
373.3	114.86	as 371.8;	TR-calcite	
375.0	115.38	mud, lt green;	I-calcite	
446.1	137.26	mud, green-black; few salt(?) xtals	TR-calcite TR-thenardite?	
447.5	137.69	mud, lt green, orange stained; few salt(?) xtals	TR-calcite	
448.5	138.00	as 447.5; less salt	TR-calcite TR-thenardite?	
449.4	138.28	as 447.5;	I-calcite	
453.0	139.38	mud, dk green; mainly salt	TR-dolomite	
455.5	140.15	as 453.0;	M-northupite TR-dolomite	
457.0	140.62	mud, dk green; trona, other salts(?)	I-northupite TR-calcite?	
459.5	141.38	mud, green; salts(?)	TR-calcite	
476.0	146.46	as 459.6;	I-calcite	
481.0	148.00	mud, green-grey; more salt than 459.6;	I-calcite,	
483.4	148.74	mud, lt green, orange-stained; large salt(?) xtals	I-calcite	
484.4	149.05	as 483.4; trona	I-calcite	
485.7	149.45	mud, green, orange-stained; large salt(?) xtals	I-calcite, TR-pirssonite	
486.0	149.54	as 485.7;	I-calcite	
487.8	150.09	mud, dk green; small salt(?) xtals	I-calcite	
489.0	150.46	mud, lt green; small salt(?) xtals	I-calcite	
490.0	150.77	mud, lt green, orange-stained; large salt(?) xtals	TR-northupite TR-dolomite TR-calcite(?)	

CORE DEPTH (ft.)	CORE DEPTH (m)	FIELD DESCRIPTION	XRD LEACHED SAMPLES	XRD BULK SAMPLES
491.2	151.14	mud, lt green, orange-stained; smaller salt(?) xtals than above	M-calcite	
492.0	151.38	as 491.2; more salt(?) than 491.2	M-calcite TR-pirssonite?	
493.2	151.75	as 491.2;	I-calcite	
493.9	151.97	mud, green; from thin bed of mud in between halite seams (2-6-8 cm)	I-calcite	
502.1	154.49	mud, green; large halite xtals	TR-calcite	
546.1	168.03	mud, dk green; PIRS xtals	I-calcite	
547.4	168.43	as 546.1;	M-calcite	
547.7	168.52	mud, dk green-black; less PIRS xtals than 546.1	I-calcite	
548.5	168.77	mud, green; many small salt(?) xtals	M-calcite	
549.5	169.08	as 548.5;	I-calcite	
550.5	169.38	mud, lt green; large PIRS(?) xtals	M-calcite, TR-dolomite	I-dolomite I-pirssonite
551.7	169.75	mud, green; PIRS(?) xtals	M-calcite	
552.6	170.03	mud, dk green, orange-stained; small PIRS? xtals	I-calcite	
553.8	170.40	as 552.6;	I-calcite Tr-pirssonite?	
555.0	170.77	mud, green, finely lam; little salt	M-calcite TR-dolomite	
556.0	171.08	mud, lt green; small salt? xtals	M-calcite TR-dolomite	
557.1	171.42	mud, lt green, orange-stained; small salt? xtals	I-calcite TR-dolomite	
558.2	171.75	mud, dk green-black; small salt? xtals	I-calcite TR-dolomite	
559.2	172.06	mud, green; white powdery substance, small salt?	TR-calcite TR-dolomite	
560.1	172.34	mud, green, finely lam; some salts?	I-calcite I-dolomite TR-thenardite TR-northupite?	

CORE DEPTH (ft.)	CORE DEPTH (m)	FIELD DESCRIPTION	XRD LEACHED SAMPLES	XRD BULK SAMPLES
561.0	172.62	as 560.1;	I-calcite, TR-dolomite	
562.2	172.98	mud, dk green-black;PIRS xtals	M-calcite, TR-dolomite	
563.2	173.29	mud, green;PIRS xtals	I-calcite, TR-dolomite	
564.3	173.63	mud, dk green, finely lam;salt? xtals	I-calcite, I-dolomite	
565.3	173.94	mud, green;mainly PIRS? xtals	M-calcite, I-dolomite	I-dolomite TR-pirssonite
567.0	174.46	as 565.3;	I-calcite, TR-dolomite	
568.7	174.98	mud, dk green; mainly PIRS? xtals	I-calcite, TR-dolomite	
569.9	175.35	mud, lt green, orange-stained; few salt? xtals	M-dolomite, TR-northupite?	
571.2	175.75	mud, lt green, orange-stained; large salt? xtals	I-dolomite, TR-calcite	
572.5	176.15	as 569.9;	I-dolomite	
573.8	176.55	mud, yellow-black, finely lam;	TR-dolomite	
574.9	176.89	as 573.8;	I-dolomite	
575.4	177.05	mud, lt green-yellow, finely lam; large salt? xtals	M-dolomite, I-calcite TR-northupite	
576.4	177.35	as 575.4;	TR-dolomite	
577.6	177.72	mud, green-grey, hard,dense; small salt? xtals	M-calcite	
578.6	178.03	as 577.6, some orange-staining	I-calcite, TR-dolomite	
579.5	178.31	mud, yellow-green, hard, dense, finely lam;few small salt? xtals	I-calcite, TR-dolomite	
580.7	178.68	as 579.5;	I-calcite, TR-dolomite	
582.2	179.14	mud, yellow-black, hard, dense; white stringers?, small salt? xtals	I-calcite, TR-dolomite?	
583.3	179.48	mud, green-grey, hard,dense, some orange-staining;	TR-thenardite TR-trona Tr-calcite	

CORE DEPTH (ft.)	CORE DEPTH (m)	FIELD DESCRIPTION	XRD LEACHED SAMPLES	XRD BULK SAMPLES
584.2	179.75	mud, orange-black; some salt? xtals, trona?	TR-calcite, TR-dolomite?	
587.8	180.86	mud, dk green-black, orange-stained; large salt? xtals, (below 2ft seam of halite)	TR-calcite	
590.9	181.82	mud, yellow-dk green, lam; small salt? xtals	I-dolomite	
592.6	182.34	as 590.9; salt? xtals larger, more frequent	M-northupite, TR-dolomite, TR-calcite	
594.8	183.02	mud, lt green-yellow, salt? xtals	M-dolomite	
598.5	184.15	mud, lt green-yellow, hard, orange-stained, small salt? xtals	I-calcite, I-dolomite	I-dolomite TR-pirssonite
601.5	185.08	mud, orange-green; large salt? xtals	TR-thenardite	
603.0	185.54	as 601.5; (just below thin trona seam)	I-northupite, I-pirssonite, TR-calcite, TR-dolomite	
605.0	186.15	mud, orange-green; ~50% salt xtals	I-pirssonite, TR-calcite	
607.9	187.05	mud, green-grey; large salt? xtals	TR-northupite, TR-calcite, TR-dolomite?	
609.9	187.66	mud, orange-green; ~50% salt? xtals	M-pirssonite, M-northupite, TR-dolomite	
611.5	188.15	as 609.9; (0.7 ft above trona seam)	TR-thenardite, TR-dolomite	
615.0	189.23	mud, orange-green; small xtals	TR-northupite	
624.0	192.00	as 615.0; small PIRS? xtals	I-calcite	
631.8	194.40	mud, dk green-orange, hard, dense; small salt? xtals	TR-calcite	
634.2	195.14	mud, dk green-black, hard, dense, orange staining; small salt? xtals	I-calcite, TR-dolomite	
636.4	195.82	as 634.2; less salt?	TR-calcite, TR-dolomite	

CORE DEPTH (ft.)	CORE DEPTH (m)	FIELD DESCRIPTION	XRD LEACHED SAMPLES	XRD BULK SAMPLES
638.5	196.46	mud, lt green, hard, dense, orange stained; salt? stringers	TR-northupite TR-calcite	
641.1	197.26	as 638.5;	TR-dolomite, TR-calcite	
642.5	197.69	mud, lt green, less dense than above; ~50% salts	M-calcite, TR-dolomite	
645.2	198.52	mud, green, crumbly; ~30% salts (below 1.5 ft halite seam)	M-calcite, TR-dolomite	
647.4	199.20	mud, dk green; ~20% salts	M-calcite, TR-pirssonite	
651.4	200.43	as 647.4;	M-calcite	
653.6	201.11	mud, grey; little salt	M-calcite I-dolomite	I-dolomite I-pirssonite
659.2	202.83	mud, green-grey; little salt	TR-calcite TR-dolomite	
661.7	203.60	as 659.2;	TR-calcite TR-dolomite	
664.8	204.55	mud, dk green; few salt? xtals	I-dolomite	
666.5	205.08	as 664.8;	TR-calcite TR-dolomite TR-northupite?	
668.4	205.66	mud, green-grey, orange-stained; few salts	I-calcite I-dolomite	
670.2	206.22	mud, dk green-orange; few salt stringers	TR-dolomite TR-northupite?	
671.5	206.62	mud, grey-black; small salt? xtals	TR-dolomite? TR-northupite?	
680.7	209.45	mud, yellow-green; little salt	M-dolomite	
684.7	210.68	mud, white, soft; no salt	M-dolomite	
690.3	212.40	mud, lt green-orange; PIRS? xtals	TR-dolomite TR-calcite	
701.2	215.75	mud, green-brown; ~60% salts?(just below coarse halite)	I-northupite I-dolomite	
706.2	217.29	mud, green; large salt? xtals	M-calcite TR-pirssonite	
710.2	218.52	as 706.2;	M-calcite TR-pirssonite	

CORE DEPTH (ft.)	CORE DEPTH (m)	FIELD DESCRIPTION	XRD LEACHED SAMPLES	XRD BULK SAMPLES
713.4	219.51	as 706.2;	I-calcite TR-pirssonite	
717.0	220.62	mud, dk green; salt? xtals	I-calcite TR-pirssonite	
724.9	223.05	as 717.0;	I-calcite Tr-pirssonite	
727.5	223.85	mud, green, crumbly; small PIRS? xtals	I-calcite	
747.2	229.91	mud, lt grey- orange, lam w/ salts;(just above minor halite seam)	I-dolomite TR-calcite	
750.0	230.77	mud, lt green; little salt	I-dolomite	
752.5	231.54	mud, buff/black lam; salt? xtals	TR-calcite TR-northupite TR-dolomite	TR-pirssonite TR-northupite TR-dolomite
758.0	233.23	as 752.5;less salt	I-dolomite TR-calcite	
762.4	234.58	mud, grey/buff lam; little salt	M-dolomite	
766.5	235.85	mud, green/buff lam; more salt than 762.4	M-calcite TR-dolomite	I-pirssonite TR-dolomite
772.2	237.60	mud, buff/orange lam; little salt	TR-calcite TR-dolomite	
775.8	238.71	mud, green-orange, lam; minor salt	I-dolomite	
780.8	240.25	mud, green-dk orange; PIRS? xtals	M-calcite I-dolomite	
785.9	241.82	mud, green, mass;	I-dolomite	
794.2	244.37	mud, green-orange, mass;	I-dolomite	
798.7	245.75	as 780.8;	I-calcite I-dolomite	
805.0	247.69	mud, dk green-orange; minor salts	I-calcite TR-dolomite	
814.0	250.46	mud, lt green; large amounts of salts?	I-calcite TR-dolomite	
818.2	251.75	mud, green-orange, hard, dense; minor amounts of salt	M-calcite TR-dolomite	M-calcite I-pirssonite I-dolomite
823.6	253.42	as 818.2;	TR-calcite	

CORE DEPTH (ft.)	CORE DEPTH (m)	FIELD DESCRIPTION	XRD LEACHED SAMPLES	XRD BULK SAMPLES
827.0	254.46	mud, dk green/orange lam, hard; minor salts	TR-calcite TR-dolomite	
831.7	255.91	mud, dk green/orange lam, hard, dense; little salt	TR-dolomite TR-calcite	
837.0	257.54	mud, green/buff lam; no salt ***HAND SAMPLE****	M-dolomite	M-dolomite
841.6	258.95	mud, green-orange; minor salt	I-dolomite TR-calcite	
846.6	260.49	mud, buff;mainly PIRS xtals	M-calcite TR-dolomite TR-pirssonite	I-pirssonite I-dolomite TR-calcite
859.8	264.55	mud, green-orange, hard,dense; salt stringers	TR-calcite TR-dolomite	
864.0	265.85	mud, lt green-buff; minor salt	I-dolomite	
876.2	269.60	mud, lt green/buff lam; no salt	M-dolomite	
887.6	273.11	mud, green-orange, hard, dense; large amounts of salt	M-calcite TR-dolomite	
893.9	275.05	mud, dk green, hard, dense,orange-stained; minor salt	I-calcite	
898.9	276.58	as 893.9;	I-calcite	
903.1	277.88	mud, green-orange; large amount of salt	I-calcite I-dolomite	
908.6	279.57	mud, green-orange, hard, dense;minor salts	I-calcite TR-dolomite	
917.1	282.18	mud, grey-buff, hard, dense; salt stringers	M-calcite	
922.0	283.69	mud, grey-orange,hard dense; minor salt	M-calcite TR-dolomite	M-calcite TR-dolomite
930.0	286.15	mud, lt green-orange; minor salt	TR-calcite TR-dolomite	
935.7	287.91	as 930.0;	TR-calcite TR-dolomite	
939.9	289.20	mud, green/orange lam; hard, dense; salt xtals	TR-calcite TR-dolomite	
948.0	291.69	mud, green-orange, hard, dense; salt xtals	M-calcite	

CORE DEPTH (ft.)	CORE DEPTH (m)	FIELD DESCRIPTION	XRD LEACHED SAMPLES	XRD BULK SAMPLES
955.0	293.85	mud, dk green, hard, dense; minor salt	I-calcite TR-dolomite	
959.0	295.08	mud, green, hard, dense; minor salt	M-calcite	
966.0	297.23	mud, green/buff lam; salt stringers	M-calcite	M-calcite
972.1	299.11	mud, dk green, mass;	TR-dolomite	
977.0	300.62	mud, green/buff lam;	I-calcite TR-dolomite	I-calcite TR-dolomite
986.0	303.38	mud, green, mass;	I-calcite	
1006.0	309.54	mud, green, mass; minor salt	I-aragonite? TR-calcite TR-dolomite?	
1012.0	311.38	mud, dk green, mass;	I-calcite	
1017.0	312.92	as 1012.0;	I-calcite I-dolomite	
1034.2	318.22	mud, green, mass; large HAL xtals	M-dolomite	
1068.0	328.62	mud, dk green, hard, dense; large HAL xtals	TR-dolomite TR-calcite	
1076.0	331.08	mud, green/orange lam; hard, dense; some salt	I-calcite	I-calcite
1080.4	332.43	mud, green, hard, dense; minor salt	TR-calcite	
1089.0	335.08	mud, green, lam;	I-calcite	
1093.0	336.31	mud, green-orange, hard, dense; large HAL xtals	M-calcite	
1099.0	338.15	mud, green, hard, dense;	TR-calcite	
1104.0	339.69	mud, grey/buff lam; minor salt	TR-calcite	
1106.5	340.46	mud, grey-buff, lam; minor salt	TR-calcite	
1110.6	341.72	mud, grey-black lam; little salt	TR-calcite	
1115.3	343.17	mud, grey-buff; **HAND SAMPLE**	TR-calcite	
1120.0	344.62	mud, grey lam; minor salt	I-calcite	
1125.5	346.31	mud, green-brown; no salt	TR-calcite	

CORE DEPTH (ft.)	CORE DEPTH (m)	FIELD DESCRIPTION	XRD LEACHED SAMPLES	XRD BULK SAMPLES
1130.5	347.85	mud, white/green lam, orange-stained; minor halite	M-calcite I-dolomite	M-calcite I-dolomite
1132.7	348.52	mud, green, lam; minor halite	M-dolomite TR-calcite	
1156.5	355.85	mud, grey/brown lam; minor salt	TR-calcite	
1210.7	372.52	mud, grey-buff, hard, dense; no salt	I-calcite	
1294.0	398.15	mud, grey-buff, hard, dense; **HAND SAMPLE**	TR-calcite	

Appendix C

Water-leaching Procedure

APPENDIX C

WATER-LEACHING PROCEDURE

Materials needed:

tweezers
Mortar and Pestle
50 ml centrifuge tubes with caps
distilled water
centrifuge
watch glasses
oven
2-drams vials with lids

Procedure

- 1) Carefully inspect each sample for macroscopic salts crystals (gaylussite, pirssonite, trona, thenardite, borax, northupite, halite, etc.). Remove all macroscopic salts crystals with tweezers.
- 2) Grind remaining sample in mortar until it would pass ~100 mesh.
- 3) Place approximately five grams of ground mud sample into a 50 ml centrifuge tube. Fill tube with distilled water and cap (no air pockets should be in the tube).
- 4) Shake centrifuge tube. Leave sample for 24 hours, shaking periodically, to dissolve soluble salts.
- 5) After ~24 hours, check sample. If the sample is cloudy, centrifuge sample until liquid is clear. Carefully decant liquid. Add fresh distilled water, cap, and shake sample to rinse mud. Centrifuge sample until liquid is clear and, again, carefully decant liquid. Repeat the rinsing process at least three times or until liquid shows no signs of color (the 'leachate' was usually a dark yellow to brown color).
- 6) Carefully scrape the centrifuged mud (not easy) from the bottom of the centrifuge tube. Place mud on a watch glass. [NOTE: At this point the sample was checked to make sure no tiny salt crystals were visible. If there were crystals visible, the entire leaching process for that sample was repeated.] Place in the oven overnight at a low temperature to dry the mud.
- 7) Remove sample from oven. [NOTE: again, another check for salt was made. If an efflorescence was

observed, the entire leaching process was repeated.]
Remove dried sample from watch glass and crush in
mortar until the sample will pass ~ 200 mesh. Place
sample in vial for mass spectrometric analysis.

Appendix D

EDTA Dissolution Procedure

APPENDIX D

EDTA DISSOLUTION PROCEDURE AND RESULTS

Introduction

Following Glover (1961) and Bodine and Fernalld (1973), an attempt was made to separate calcite from dolomite using a high pH aqueous solution of the complexing agent EDTA (ethylenedinitrilotetraacetic acid). The chemical processes governing the dissolution of alkaline earth carbonates by EDTA can be found in Bodine and Fernalld (1973; after Welcher, 1958).

Materials Needed

Tetrasodium EDTA dihydrate powder
50 ml vials (centrifuge tubes used here) with caps
pH paper or meter
shaker
watch glasses
centrifuge
distilled water
oven
containers for leached sample (2-dram vials used here)

Procedure Followed

- 1) A .25M (near saturation) solution of EDTA (using tetrasodium EDTA, pH should be ~10.5) was prepared.
- 2) Searles Lake sample 127.7 (a fairly equal mixture of "mongrel calcite and dolomite, and about 30 weight percent carbonate) was selected as a test sample. Four aliquots of water-leached mud sample (passing 100 mesh) at ~two grams each were placed into 50 ml centrifuge tubes. ~50 ml of EDTA solution were added to the sample, and the centrifuge tube was capped.
- 3) The four samples were placed on a shaker for 1,2,4, and

8 hours respectively at room temperature ($\sim 20^{\circ} \text{C}$). High temperature used by previous investigators were not used due to the extreme rapidity of the complexing reaction.

- 4) After each time interval had passed, one sample was removed from the shaker. The pH was measured (all samples had a pH between 9.5-10.5).
- 5) The sample was placed in a centrifuge briefly to spin down solids. The EDTA solution was carefully decanted. The sample was rinsed with distilled water several times to remove all EDTA. The mud was carefully removed from the centrifuge tube and placed on a watch glass. The watch glass was placed in a low temperature oven overnight to dry.
- 6) Mortar and pestle were used to crush the dried sample until it passed 200 mesh. Samples were placed in vials for XRD analysis.
- 7) The steps one through six were repeated until eight samples were created (A and B): two 1-hour, two 2-hour, two 4-hour, and two 8-hour samples.
- 8) X-Ray Diffraction analysis was performed on the eight samples. The samples were scanned from 25-32 degrees $2-\theta$ Cu $K\alpha$ for the primary carbonate peaks. The calcite peak gradually disappeared, and the primary dolomite peak diminished slightly by 4 hours and was significantly reduced at 8 hours.
- 9) The eight samples were analyzed for ^{18}O and ^{13}C content at the University of New Mexico (Albuquerque) isotope lab following the methods of McCrea (1950). The UNM lab reported difficulties in the mass spectrometric analyses of these samples: samples 1A, 4A, and 8B were reported "contaminated". The stable isotope content of the remaining samples is shown graphically in fig. 34.

No acid fraction factors were assigned to the samples. The stable isotope content of the bulk water-leached sample and the dolomite (determined later) values are included in fig. 34 for illustrative purposes.

Discussion

The isotopic content of the 1-hour sample is very similar to that of the bulk sample, as would be expected.

TETRASODIUM EDTA MINERAL SEPARATIONS

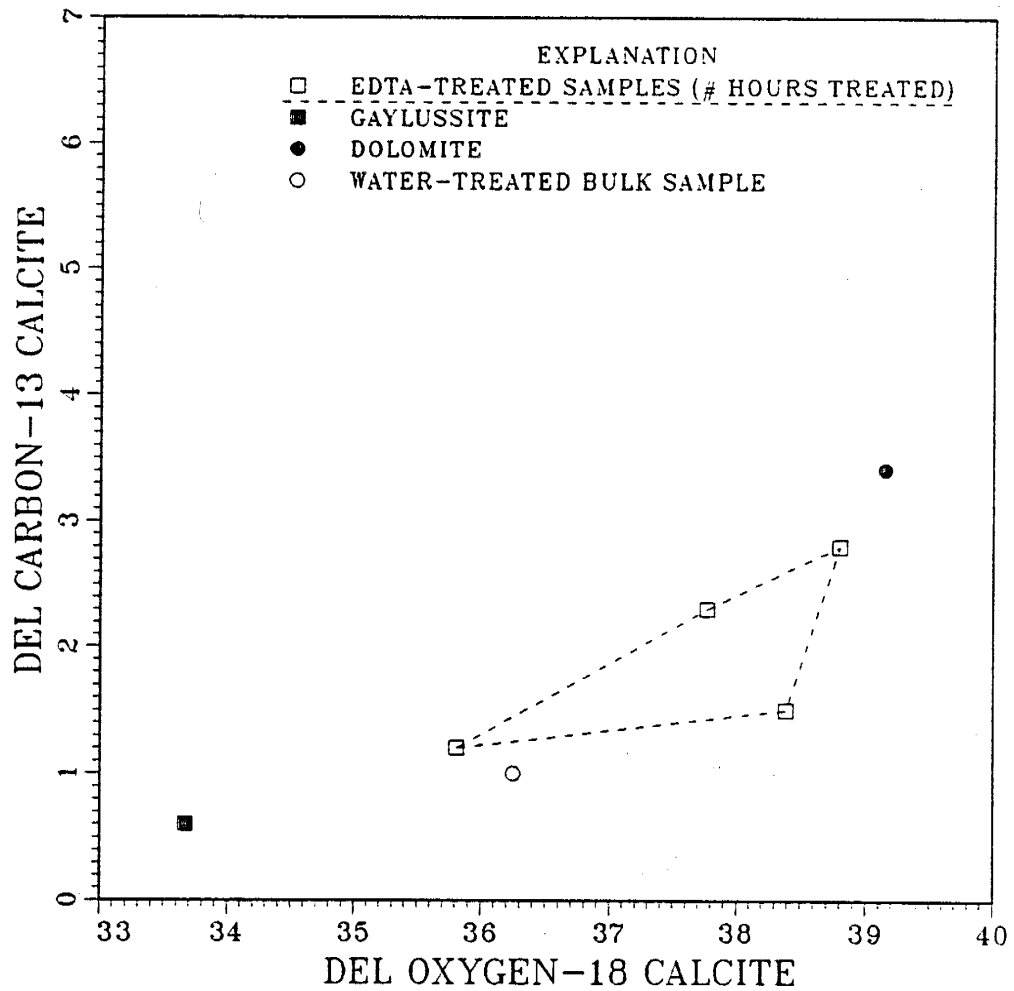


Figure 34. EDTA chemical separation of KM-3 sample 127.7 (feet).

The isotopic content of the 2-hour sample approaches the dolomite values. However, it appears that the EDTA had begun to complex with the dolomite by the 4-hour sample, and by eight hours the isotopic content of the sample had essentially returned to that of the 1-hour sample. It appears that the same relative proportion of calcite/dolomite existed after eight hours of EDTA dissolution as had existed in the 1-hour sample.

The money available for the stable isotope analysis did not permit the detailed mole% calcite and dolomite determination nor exhaustive grain size analysis that would have been required to calibrate this procedure. Further, the 'dolomite endpoint' might not be achieved as the EDTA began to complex with the dolomite in the sample.

Appendix E

Hierarchical Statistics Analysis of Uncertainty
for ^{18}O and ^{13}C Analyses

C
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PROGRAM RUNS ANALYSIS OF VARIANCE FOR DATA SET

```
program anova
character fname*80
character fname2*20
character desc*50
dimension yhia(1000),yhiabar(1000)
dimension yhi(1000),yhibar(1000),nhi(1000)
dimension yh(1000),yhbar(1000),nh(1000)
dimension min(1000),maj(1000)
DIMENSION EHI(1000),THI(1000),RMHI(1000)

write(6,*)'enter the input file name'
read(5,'(a)')fname
open(unit=99,file=fname,status='old')
write(6,*)'enter output file name'
read(5,'(a)')fname2
open(unit=96,file=fname2,status='new')

read(99,*)n,nmhi,nmh
if(n.gt.1000 .or.nmhi.gt.1000 .or. nmh.gt.1000)then
  write(6,*)'program dimensioned to 1000',n,nmhi,nmh
  stop
endif
write(6,*)'input number of cycles per minor group'
read(5,*)k

do 100 i = 1,n
  read(99,*)yhia(i),min(i),maj(i)
  nhi(min(i)) = nhi(min(i)) + 1
  nh(maj(i)) = nh(maj(i)) + 1
100 continue

write(96,*)
write(96,*)
write(96,*)
write(96,*)
write(96,*)
write(96,*)
write(6,*)'enter decription'
read(5,'(a)')desc
write(96,*)desc
write(96,*)'_____ '
write(96,*)
write(96,*)'number of points:      ',n
write(96,*)'number of minor groups:',nmhi
write(96,*)'number of major groups:',nmh
write(96,*)

C data has been input, compute the totals for each minor and major group

do 200 i = 1,n
  yhi(min(i)) = yhi(min(i)) + yhia(i)
  yh(maj(i)) = yh(maj(i)) + yhia(i)
  y = y + yhia(i)
200 continue
DO 300 I = 1,NMHI*nmh
  YHIBAR(I) = YHI(I) / FLOAT(NHI(I) )
  WRITE(96,10)' MINOR GROUP #, SIZE, TOTAL, MEAN',
1      I,NHI(I),YHI(I),YHIBAR(I)
300 CONTINUE
WRITE(96,*)
DO 400 I = 1,NMH
  YHBAR(I) = YH(I) / FLOAT(NH(I) )
  WRITE(96,10)' MAJOR GROUP #, SIZE, TOTAL, MEAN',
1      I,NH(I),YH(I),YHBAR(I)
```


SAMPLE 690.3 OXYGEN-18

number of points: 150
 number of minor groups: 5
 number of major groups: 5

MINOR GROUP #, SIZE, TOTAL, MEAN	1	6	1.4449E+02	2.4082E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	2	6	1.4366E+02	2.3943E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	3	6	1.4320E+02	2.3866E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	4	6	1.4308E+02	2.3846E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	5	6	1.4321E+02	2.3868E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	6	6	1.4750E+02	2.4583E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	7	6	1.4807E+02	2.4679E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	8	6	1.4814E+02	2.4690E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	9	6	1.4813E+02	2.4689E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	10	6	1.4813E+02	2.4688E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	11	6	1.5090E+02	2.5150E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	12	6	1.4941E+02	2.4901E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	13	6	1.4892E+02	2.4820E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	14	6	1.4892E+02	2.4820E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	15	6	1.4907E+02	2.4845E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	16	6	1.4845E+02	2.4742E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	17	6	1.4764E+02	2.4607E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	18	6	1.4765E+02	2.4608E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	19	6	1.4739E+02	2.4565E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	20	6	1.4747E+02	2.4578E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	21	6	1.3855E+02	2.3091E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	22	6	1.3964E+02	2.3274E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	23	6	1.3950E+02	2.3249E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	24	6	1.3935E+02	2.3225E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	25	6	1.3914E+02	2.3191E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	1	30	7.1763E+02	2.3921E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	2	30	7.3997E+02	2.4666E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	3	30	7.4721E+02	2.4907E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	4	30	7.3859E+02	2.4620E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	5	30	6.9618E+02	2.3206E+01	
TOTAL RUN	SIZE, TOTAL, MEAN	1	150	3.6396E+03	2.4264E+01

ANOVA

	SUM SQUARES	DEG FROM	MEAN SUM SQUARES	VARIANCE
BTWN SAMPLES	58.1719	4	14.5430	0.4831
BTWN GAS ALIQUOTS	1.0000	20	0.0500	0.0072
BTWN MACHINE CYCLES	0.8359	125	0.0067	0.0067

RAW OXYGEN_18 CYCLE DATA: SAMPLE 690.3

CYCLE	MINOR GROUP	MAJOR GROUP
24.087	1	1
24.069	1	1
23.995	1	1
24.125	1	1
24.216	1	1
24.001	1	1
23.652	2	1
24.025	2	1
24.019	2	1
23.910	2	1
23.974	2	1
24.076	2	1
23.867	3	1
23.742	3	1
23.907	3	1
23.824	3	1
23.909	3	1
23.949	3	1
23.815	4	1
23.842	4	1
23.769	4	1
23.779	4	1
24.010	4	1
23.863	4	1
23.839	5	1
23.857	5	1
23.955	5	1
23.968	5	1
23.856	5	1
23.730	5	1
24.461	6	2
24.547	6	2
24.769	6	2
24.512	6	2
24.532	6	2
24.676	6	2
24.563	7	2
24.701	7	2
24.676	7	2
24.758	7	2
24.665	7	2
24.710	7	2
24.686	8	2
24.722	8	2
24.725	8	2
24.695	8	2
24.650	8	2
24.659	8	2
24.684	9	2
24.699	9	2
24.692	9	2
24.754	9	2
24.695	9	2
24.608	9	2
24.697	10	2
24.625	10	2
24.704	10	2
24.635	10	2
24.728	10	2
24.738	10	2
25.194	11	3
25.190	11	3
25.063	11	3
25.060	11	3
25.209	11	3
25.184	11	3

CYCLE	MINOR GROUP	MAJOR GROUP
24.848	12	3
24.823	12	3
24.866	12	3
24.977	12	3
24.947	12	3
24.946	12	3
24.728	13	3
24.911	13	3
24.786	13	3
24.866	13	3
24.771	13	3
24.857	13	3
24.841	14	3
24.838	14	3
24.775	14	3
24.767	14	3
24.869	14	3
24.830	14	3
24.867	15	3
24.865	15	3
24.846	15	3
24.760	15	3
24.795	15	3
24.935	15	3
24.800	16	4
24.779	16	4
24.771	16	4
24.690	16	4
24.629	16	4
24.780	16	4
24.610	17	4
24.574	17	4
24.642	17	4
24.615	17	4
24.733	17	4
24.465	17	4
24.561	18	4
24.618	18	4
24.654	18	4
24.603	18	4
24.512	18	4
24.699	18	4
24.520	19	4
24.518	19	4
24.688	19	4
24.563	19	4
24.493	19	4
24.609	19	4
24.573	20	4
24.576	20	4
24.659	20	4
24.604	20	4
24.577	20	4
24.476	20	4
22.855	21	5
22.988	21	5
23.179	21	5
23.160	21	5
23.159	21	5
23.206	21	5
23.287	22	5
23.102	22	5
23.209	22	5
23.331	22	5
23.288	22	5
23.425	22	5

CYCLE	MINOR	GROUP	MAJOR	GROUP
23.287		23		5
23.383		23		5
23.291		23		5
22.991		23		5
23.284		23		5
23.259		23		5
23.241		24		5
23.220		24		5
23.203		24		5
23.167		24		5
23.303		24		5
23.218		24		5
23.223		25		5
23.214		25		5
23.173		25		5
23.109		25		5
23.242		25		5
23.182		25		5

SAMPLE 690.3 CARBON-13

number of points: 150
 number of minor groups: 5
 number of major groups: 5

MINOR GROUP #, SIZE, TOTAL, MEAN	1	6	2.6780E+02	4.4633E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	2	6	2.6750E+02	4.4583E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	3	6	2.6751E+02	4.4585E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	4	6	2.6750E+02	4.4583E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	5	6	2.6793E+02	4.4655E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	6	6	2.6807E+02	4.4679E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	7	6	2.6854E+02	4.4756E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	8	6	2.6837E+02	4.4729E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	9	6	2.6890E+02	4.4817E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	10	6	2.6856E+02	4.4759E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	11	6	2.6943E+02	4.4905E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	12	6	2.6906E+02	4.4844E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	13	6	2.6917E+02	4.4862E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	14	6	2.6887E+02	4.4812E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	15	6	2.6898E+02	4.4830E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	16	6	2.6845E+02	4.4742E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	17	6	2.6838E+02	4.4730E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	18	6	2.6854E+02	4.4757E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	19	6	2.6857E+02	4.4762E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	20	6	2.6851E+02	4.4752E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	21	6	2.6615E+02	4.4358E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	22	6	2.6691E+02	4.4485E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	23	6	2.6672E+02	4.4453E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	24	6	2.6689E+02	4.4482E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	25	6	2.6669E+02	4.4448E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	1	30	1.3382E+03	4.4608E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	2	30	1.3424E+03	4.4748E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	3	30	1.3455E+03	4.4851E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	4	30	1.3425E+03	4.4749E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	5	30	1.3334E+03	4.4445E+01	
TOTAL RUN	SIZE, TOTAL, MEAN	1	150	6.7020E+03	4.4680E+01

ANOVA

	SUM SQUARES	DEG FRDM	MEAN SUM SQUARES	VARIANCE
BTWN SAMPLES	3.0000	4	0.7500	0.0247
BTWN GAS ALIQUOTS	0.1563	20	0.0078	0.0008
BTWN MACHINE CYCLES	0.4063	125	0.0033	0.0033

RAW CARBON-13 CYCLE DATA: SAMPLE 690.3

CYCLE	MINOR GROUP	MAJOR GROUP
44.667	1	1
44.531	1	1
44.551	1	1
44.678	1	1
44.690	1	1
44.684	1	1
44.542	2	1
44.575	2	1
44.519	2	1
44.506	2	1
44.690	2	1
44.666	2	1
44.578	3	1
44.536	3	1
44.498	3	1
44.608	3	1
44.712	3	1
44.578	3	1
44.639	4	1
44.557	4	1
44.617	4	1
44.478	4	1
44.635	4	1
44.572	4	1
44.625	5	1
44.608	5	1
44.643	5	1
44.702	5	1
44.646	5	1
44.706	5	1
44.654	6	2
44.666	6	2
44.726	6	2
44.671	6	2
44.638	6	2
44.716	6	2
44.704	7	2
44.765	7	2
44.747	7	2
44.749	7	2
44.755	7	2
44.815	7	2
44.733	8	2
44.729	8	2
44.708	8	2
44.776	8	2
44.705	8	2
44.721	8	2
44.800	9	2
44.816	9	2
44.844	9	2
44.846	9	2
44.789	9	2
44.804	9	2
44.744	10	2
44.712	10	2
44.672	10	2
44.808	10	2
44.816	10	2
44.803	10	2
44.937	11	3
44.938	11	3
44.908	11	3
44.911	11	3
44.881	11	3
44.855	11	3

CYCLE	MINOR GROUP	MAJOR GROUP
44.791	12	3
44.765	12	3
44.849	12	3
44.914	12	3
44.949	12	3
44.794	12	3
44.838	13	3
44.922	13	3
44.812	13	3
44.896	13	3
44.849	13	3
44.853	13	3
44.825	14	3
44.808	14	3
44.846	14	3
44.846	14	3
44.783	14	3
44.766	14	3
44.860	15	3
44.892	15	3
44.792	15	3
44.830	15	3
44.792	15	3
44.816	15	3
44.690	16	4
44.790	16	4
44.757	16	4
44.711	16	4
44.768	16	4
44.737	16	4
44.739	17	4
44.687	17	4
44.711	17	4
44.708	17	4
44.789	17	4
44.745	17	4
44.739	18	4
44.695	18	4
44.761	18	4
44.790	18	4
44.772	18	4
44.783	18	4
44.702	19	4
44.735	19	4
44.829	19	4
44.809	19	4
44.740	19	4
44.758	19	4
44.775	20	4
44.805	20	4
44.710	20	4
44.720	20	4
44.789	20	4
44.711	20	4
44.227	21	5
44.353	21	5
44.374	21	5
44.361	21	5
44.444	21	5
44.390	21	5
44.401	22	5
44.518	22	5
44.538	22	5
44.441	22	5
44.521	22	5
44.493	22	5

CYCLE	MINOR GROUP	MAJOR GROUP
44.353	23	5
44.492	23	5
44.535	23	5
44.449	23	5
44.483	23	5
44.405	23	5
44.432	24	5
44.420	24	5
44.549	24	5
44.605	24	5
44.459	24	5
44.429	24	5
44.426	25	5
44.522	25	5
44.398	25	5
44.374	25	5
44.485	25	5
44.483	25	5

SAMPLE 375.0 OXYGEN-18

number of points: 150
 number of minor groups: 5
 number of major groups: 5

MINOR GROUP #, SIZE, TOTAL, MEAN	1	6	1.0872E+02	1.8120E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	2	6	1.0946E+02	1.8243E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	3	6	1.0917E+02	1.8194E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	4	6	1.0845E+02	1.8074E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	5	6	1.0858E+02	1.8097E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	6	6	1.1005E+02	1.8341E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	7	6	1.0977E+02	1.8295E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	8	6	1.0965E+02	1.8275E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	9	6	1.0930E+02	1.8217E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	10	6	1.1006E+02	1.8344E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	11	6	1.1317E+02	1.8861E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	12	6	1.1392E+02	1.8987E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	13	6	1.1268E+02	1.8780E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	14	6	1.1264E+02	1.8774E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	15	6	1.1347E+02	1.8912E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	16	6	1.0823E+02	1.8039E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	17	6	1.0870E+02	1.8117E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	18	6	1.0996E+02	1.8327E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	19	6	1.0874E+02	1.8124E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	20	6	1.0990E+02	1.8316E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	21	6	1.1293E+02	1.8821E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	22	6	1.1407E+02	1.9011E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	23	6	1.1332E+02	1.8887E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	24	6	1.1437E+02	1.9062E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	25	6	1.1418E+02	1.9031E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	1	30	5.4437E+02	1.8146E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	2	30	5.4883E+02	1.8294E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	3	30	5.6589E+02	1.8863E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	4	30	5.4554E+02	1.8185E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	5	30	5.6887E+02	1.8962E+01	
TOTAL RUN	SIZE, TOTAL, MEAN	1	150	2.7735E+03	1.8490E+01

ANOVA

	SUM SQUARES	DEG FRDM	MEAN SUM SQUARES	VARIANCE
BTWN SAMPLES	18.3867	4	4.5967	0.1515
BTWN GAS ALIQUOTS	1.0430	20	0.0521	0.0030
BTWN MACHINE CYCLES	4.2969	125	0.0344	0.0344

RAW OXYGEN-18 CYCLE DATA: SAMPLE 375.0

CYCLE	MINOR GROUP	MAJOR GROUP
18.209	1	1
18.111	1	1
18.073	1	1
18.211	1	1
18.033	1	1
18.085	1	1
17.883	2	1
18.127	2	1
18.149	2	1
19.185	2	1
18.015	2	1
18.100	2	1
17.917	3	1
18.198	3	1
18.361	3	1
18.364	3	1
18.231	3	1
18.094	3	1
17.969	4	1
17.848	4	1
18.147	4	1
18.297	4	1
18.202	4	1
17.983	4	1
18.134	5	1
18.328	5	1
18.216	5	1
17.804	5	1
18.156	5	1
17.942	5	1
18.555	6	2
18.128	6	2
18.443	6	2
18.179	6	2
18.317	6	2
18.424	6	2
18.089	7	2
18.356	7	2
18.332	7	2
18.389	7	2
18.341	7	2
18.262	7	2
18.104	8	2
18.276	8	2
18.301	8	2
18.424	8	2
18.352	8	2
18.194	8	2
17.988	9	2
18.349	9	2
18.258	9	2
18.325	9	2
18.330	9	2
18.049	9	2
18.317	10	2
18.254	10	2
18.205	10	2
18.444	10	2
18.307	10	2
18.536	10	2
18.463	11	3
18.849	11	3
18.718	11	3
19.181	11	3
18.929	11	3
19.026	11	3

CYCLE	MINOR GROUP	MAJOR GROUP
18.859	12	3
18.759	12	3
19.057	12	3
19.130	12	3
18.959	12	3
19.159	12	3
18.811	13	3
18.727	13	3
18.985	13	3
18.954	13	3
18.555	13	3
18.649	13	3
18.844	14	3
18.819	14	3
18.744	14	3
18.750	14	3
18.768	14	3
18.718	14	3
18.952	15	3
18.922	15	3
18.835	15	3
18.866	15	3
18.907	15	3
18.991	15	3
17.900	16	4
17.868	16	4
18.235	16	4
18.126	16	4
17.883	16	4
18.221	16	4
17.962	17	4
17.952	17	4
18.165	17	4
18.382	17	4
18.139	17	4
18.101	17	4
18.069	18	4
18.226	18	4
18.636	18	4
18.429	18	4
18.414	18	4
18.190	18	4
17.966	19	4
18.070	19	4
18.440	19	4
18.014	19	4
18.015	19	4
18.238	19	4
18.176	20	4
18.083	20	4
18.373	20	4
18.603	20	4
18.512	20	4
18.151	20	4
18.683	21	5
18.796	21	5
19.017	21	5
18.727	21	5
18.751	21	5
18.952	21	5
18.895	22	5
19.014	22	5
19.148	22	5
19.189	22	5
18.808	22	5
19.015	22	5

CYCLE	MINOR	GROUP	MAJOR	GROUP
18.562		23		5
18.877		23		5
18.914		23		5
18.827		23		5
18.913		23		5
19.226		23		5
18.876		24		5
19.187		24		5
19.057		24		5
18.921		24		5
19.170		24		5
19.158		24		5
19.330		25		5
19.061		25		5
18.906		25		5
19.231		25		5
18.746		25		5
18.910		25		5

SAMPLE 375.0 CARBON-13

number of points: 150
 number of minor groups: 5
 number of major groups: 5

MINOR GROUP #, SIZE, TOTAL, MEAN	1	6	2.6720E+02	4.4533E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	2	6	2.6723E+02	4.4538E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	3	6	2.6732E+02	4.4553E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	4	6	2.6719E+02	4.4532E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	5	6	2.6734E+02	4.4557E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	6	6	2.6840E+02	4.4734E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	7	6	2.6818E+02	4.4696E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	8	6	2.6804E+02	4.4674E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	9	6	2.6830E+02	4.4717E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	10	6	2.6826E+02	4.4709E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	11	6	2.6906E+02	4.4844E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	12	6	2.6979E+02	4.4964E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	13	6	2.6929E+02	4.4882E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	14	6	2.6945E+02	4.4908E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	15	6	2.6958E+02	4.4930E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	16	6	2.6799E+02	4.4665E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	17	6	2.6917E+02	4.4862E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	18	6	2.6844E+02	4.4740E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	19	6	2.6862E+02	4.4770E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	20	6	2.6903E+02	4.4839E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	21	6	2.6998E+02	4.4997E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	22	6	2.6968E+02	4.4947E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	23	6	2.7016E+02	4.5027E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	24	6	2.7000E+02	4.5000E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	25	6	2.7042E+02	4.5070E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	1	30	1.3363E+03	4.4543E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	2	30	1.3412E+03	4.4706E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	3	30	1.3472E+03	4.4906E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	4	30	1.3433E+03	4.4775E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	5	30	1.3502E+03	4.5008E+01	
TOTAL RUN	SIZE, TOTAL, MEAN	1	150	6.7181E+03	4.4787E+01

ANOVA

	SUM SQUARES	DEG FRDM	MEAN SUM SQUARES	VARIANCE
BTWN SAMPLES	3.7813	4	0.9453	0.0312
BTWN GAS ALIQUOTS	0.1875	20	0.0094	-0.0011
BTWN MACHINE CYCLES	2.0313	125	0.0162	0.0162

RAW CARBON-13 CYCLE DATA: SAMPLE 375.0

CYCLE	MINOR GROUP	MAJOR GROUP
44.462	1	1
44.607	1	1
44.542	1	1
44.465	1	1
44.504	1	1
44.621	1	1
44.449	2	1
44.655	2	1
44.417	2	1
44.583	2	1
44.497	2	1
44.627	2	1
44.526	3	1
44.588	3	1
44.567	3	1
44.580	3	1
44.541	3	1
44.515	3	1
44.603	4	1
44.435	4	1
44.448	4	1
44.566	4	1
44.627	4	1
44.513	4	1
44.487	5	1
44.603	5	1
44.563	5	1
44.421	5	1
44.606	5	1
44.660	5	1
44.904	6	2
44.659	6	2
44.876	6	2
44.588	6	2
44.651	6	2
44.727	6	2
44.651	7	2
44.591	7	2
44.745	7	2
44.714	7	2
44.661	7	2
44.814	7	2
44.679	8	2
44.573	8	2
44.698	8	2
44.651	8	2
44.769	8	2
44.674	8	2
44.572	9	2
44.651	9	2
44.776	9	2
44.676	9	2
44.800	9	2
44.825	9	2
44.566	10	2
44.638	10	2
44.647	10	2
44.740	10	2
44.781	10	2
44.884	10	2
44.955	11	3
44.777	11	3
44.603	11	3
44.975	11	3
44.854	11	3
44.897	11	3

CYCLE	MINOR GROUP	MAJOR GROUP
45.105	12	3
44.927	12	3
44.831	12	3
44.964	12	3
44.905	12	3
45.053	12	3
44.766	13	3
44.851	13	3
44.955	13	3
44.972	13	3
44.959	13	3
44.787	13	3
44.857	14	3
45.029	14	3
45.007	14	3
44.840	14	3
44.774	14	3
44.941	14	3
44.990	15	3
44.949	15	3
44.821	15	3
45.025	15	3
44.994	15	3
44.802	15	3
44.486	16	4
44.525	16	4
44.570	16	4
44.672	16	4
45.005	16	4
44.733	16	4
44.839	17	4
44.887	17	4
44.920	17	4
45.018	17	4
44.802	17	4
44.704	17	4
44.806	18	4
44.802	18	4
44.997	18	4
44.784	18	4
44.612	18	4
44.440	18	4
44.599	19	4
44.783	19	4
44.747	19	4
45.033	19	4
44.826	19	4
44.633	19	4
44.936	20	4
44.783	20	4
44.793	20	4
44.873	20	4
44.949	20	4
44.698	20	4
44.814	21	5
45.220	21	5
44.953	21	5
44.790	21	5
45.008	21	5
45.194	21	5
45.018	22	5
45.041	22	5
44.753	22	5
44.870	22	5
44.927	22	5
45.072	22	5

CYCLE	MINOR	GROUP	MAJOR	GROUP
44.865	23		5	
44.779	23		5	
45.200	23		5	
45.077	23		5	
45.088	23		5	
45.155	23		5	
44.964	24		5	
45.082	24		5	
45.037	24		5	
44.994	24		5	
44.978	24		5	
44.946	24		5	
45.060	25		5	
44.969	25		5	
44.875	25		5	
45.485	25		5	
45.243	25		5	
44.786	25		5	

SAMPLE 127.7- OXYGEN-18

number of points: 150
 number of minor groups: 5
 number of major groups: 5

MINOR GROUP #, SIZE, TOTAL, MEAN	1	6	1.7868E+02	2.9780E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	2	6	1.7832E+02	2.9719E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	3	6	1.7861E+02	2.9768E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	4	6	1.7869E+02	2.9782E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	5	6	1.7874E+02	2.9791E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	6	6	1.7879E+02	2.9798E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	7	6	1.7864E+02	2.9773E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	8	6	1.7855E+02	2.9758E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	9	6	1.7844E+02	2.9740E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	10	6	1.7843E+02	2.9738E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	11	6	1.8324E+02	3.0540E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	12	6	1.8350E+02	3.0583E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	13	6	1.8339E+02	3.0565E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	14	6	1.8310E+02	3.0517E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	15	6	1.8324E+02	3.0539E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	16	6	1.7972E+02	2.9954E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	17	6	1.8004E+02	3.0006E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	18	6	1.8010E+02	3.0017E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	19	6	1.7962E+02	2.9937E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	20	6	1.8000E+02	3.0000E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	21	6	1.7953E+02	2.9922E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	22	6	1.7956E+02	2.9926E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	23	6	1.7898E+02	2.9830E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	24	6	1.7892E+02	2.9819E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	25	6	1.7907E+02	2.9845E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	1	30	8.9304E+02	2.9768E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	2	30	8.9284E+02	2.9761E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	3	30	9.1647E+02	3.0549E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	4	30	8.9949E+02	2.9983E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	5	30	8.9606E+02	2.9869E+01	
TOTAL RUN	SIZE, TOTAL, MEAN	1	150	4.4979E+03	2.9986E+01

ANOVA

	SUM SQUARES	DEG FRDM	MEAN SUM SQUARES	VARIANCE
BTWN SAMPLES	12.9375	4	3.2344	0.1076
BTWN GAS ALIQUOTS	0.1563	20	0.0078	0.0005
BTWN MACHINE CYCLES	0.5781	125	0.0046	0.0046

RAW OXYGEN-18 CYCLE DATA: SAMPLE 127.7

CYCLE	MINOR GROUP	MAJOR GROUP
29.818	1	1
29.858	1	1
29.864	1	1
29.700	1	1
29.745	1	1
29.697	1	1
29.863	2	1
29.752	2	1
29.618	2	1
29.587	2	1
29.776	2	1
29.721	2	1
29.667	3	1
29.790	3	1
29.830	3	1
29.760	3	1
29.816	3	1
29.747	3	1
29.746	4	1
29.820	4	1
29.872	4	1
29.761	4	1
29.759	4	1
29.734	4	1
29.841	5	1
29.866	5	1
29.822	5	1
29.743	5	1
29.792	5	1
29.680	5	1
29.822	6	2
29.751	6	2
29.853	6	2
29.819	6	2
29.720	6	2
29.820	6	2
29.742	7	2
29.749	7	2
29.774	7	2
29.815	7	2
29.775	7	2
29.786	7	2
29.720	8	2
29.818	8	2
29.736	8	2
29.752	8	2
29.705	8	2
29.818	8	2
29.841	9	2
29.703	9	2
29.740	9	2
29.746	9	2
29.716	9	2
29.693	9	2
29.809	10	2
29.750	10	2
29.717	10	2
29.693	10	2
29.699	10	2
29.762	10	2
30.444	11	3
30.522	11	3
30.521	11	3
30.605	11	3
30.570	11	3
30.578	11	3

CYCLE	MINOR GROUP	MAJOR GROUP
30.636	12	3
30.540	12	3
30.683	12	3
30.668	12	3
30.441	12	3
30.530	12	3
30.495	13	3
30.584	13	3
30.519	13	3
30.618	13	3
30.609	13	3
30.566	13	3
30.617	14	3
30.460	14	3
30.473	14	3
30.492	14	3
30.539	14	3
30.521	14	3
30.411	15	3
30.395	15	3
30.648	15	3
30.587	15	3
30.592	15	3
30.603	15	3
29.976	16	4
29.931	16	4
29.953	16	4
29.901	16	4
29.992	16	4
29.971	16	4
29.997	17	4
30.019	17	4
30.001	17	4
29.988	17	4
30.032	17	4
29.998	17	4
30.057	18	4
30.084	18	4
30.017	18	4
30.023	18	4
29.897	18	4
30.026	18	4
29.902	19	4
29.853	19	4
30.067	19	4
29.958	19	4
29.868	19	4
29.975	19	4
29.876	20	4
30.016	20	4
30.074	20	4
29.970	20	4
30.062	20	4
30.002	20	4
29.848	21	5
29.870	21	5
29.947	21	5
29.984	21	5
29.970	21	5
29.911	21	5
29.816	22	5
29.842	22	5
29.973	22	5
29.996	22	5
29.986	22	5
29.943	22	5
29.766	23	5

CYCLE	MINOR	GROUP	MAJOR	GROUP
29.868		23		5
29.824		23		5
29.874		23		5
29.817		23		5
29.833		23		5
29.819		24		5
29.746		24		5
29.726		24		5
29.945		24		5
29.816		24		5
29.863		24		5
29.849		25		5
29.962		25		5
29.886		25		5
29.885		25		5
29.823		25		5
29.668		25		5

SAMPLE 127.7- CARBON-13

number of total points: 150
 number of minor groups/major groups: 5
 number of major groups: 5

MINOR GROUP #, SIZE, TOTAL, MEAN	1	6	2.6201E+02	4.3668E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	2	6	2.6203E+02	4.3672E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	3	6	2.6172E+02	4.3620E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	4	6	2.6225E+02	4.3708E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	5	6	2.6232E+02	4.3719E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	6	6	2.6214E+02	4.3689E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	7	6	2.6225E+02	4.3708E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	8	6	2.6232E+02	4.3720E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	9	6	2.6209E+02	4.3682E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	10	6	2.6212E+02	4.3687E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	11	6	2.6093E+02	4.3488E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	12	6	2.6089E+02	4.3481E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	13	6	2.6080E+02	4.3467E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	14	6	2.6094E+02	4.3489E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	15	6	2.6093E+02	4.3489E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	16	6	2.6221E+02	4.3702E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	17	6	2.6239E+02	4.3732E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	18	6	2.6222E+02	4.3703E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	19	6	2.6201E+02	4.3669E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	20	6	2.6204E+02	4.3673E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	21	6	2.6203E+02	4.3672E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	22	6	2.6218E+02	4.3696E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	23	6	2.6201E+02	4.3668E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	24	6	2.6181E+02	4.3635E+01	
MINOR GROUP #, SIZE, TOTAL, MEAN	25	6	2.6191E+02	4.3652E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	1	30	1.3103E+03	4.3677E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	2	30	1.3109E+03	4.3697E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	3	30	1.3045E+03	4.3483E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	4	30	1.3109E+03	4.3696E+01	
MAJOR GROUP #, SIZE, TOTAL, MEAN	5	30	1.3099E+03	4.3665E+01	
TOTAL RUN	SIZE, TOTAL, MEAN	1	150	6.5465E+03	4.3644E+01

ANOVA

	SUM SQUARES	DEG FROM	MEAN SUM SQUARES	VARIANCE
BTWN SAMPLES	1.0938	4	0.2734	0.0090
BTWN GAS ALIQUOTS	0.0938	20	0.0047	0.0007
BTWN MACHINE CYCLES	0.0938	125	0.0008	0.0008

RAW CARBON-13 CYCLE DATA: SAMPLE 127.7

CYCLE	MINOR GROUP	MAJOR GROUP
43.690	1	1
43.683	1	1
43.629	1	1
43.607	1	1
43.699	1	1
43.700	1	1
43.672	2	1
43.691	2	1
43.673	2	1
43.671	2	1
43.691	2	1
43.634	2	1
43.595	3	1
43.546	3	1
43.671	3	1
43.610	3	1
43.653	3	1
43.643	3	1
43.723	4	1
43.686	4	1
43.725	4	1
43.669	4	1
43.757	4	1
43.690	4	1
43.735	5	1
43.748	5	1
43.688	5	1
43.684	5	1
43.746	5	1
43.714	5	1
43.703	6	2
43.678	6	2
43.689	6	2
43.735	6	2
43.689	6	2
43.641	6	2
43.748	7	2
43.704	7	2
43.626	7	2
43.686	7	2
43.763	7	2
43.720	7	2
43.770	8	2
43.717	8	2
43.700	8	2
43.668	8	2
43.711	8	2
43.756	8	2
43.770	9	2
43.621	9	2
43.673	9	2
43.667	9	2
43.721	9	2
43.637	9	2
43.716	10	2
43.756	10	2
43.732	10	2
43.678	10	2
43.586	10	2
43.656	10	2
43.415	11	3
43.478	11	3
43.509	11	3
43.546	11	3
43.481	11	3
43.498	11	3

CYCLE	MINOR GROUP	MAJOR GROUP
43.550	12	3
43.527	12	3
43.431	12	3
43.468	12	3
43.433	12	3
43.479	12	3
43.426	13	3
43.387	13	3
43.543	13	3
43.495	13	3
43.450	13	3
43.500	13	3
43.493	14	3
43.503	14	3
43.562	14	3
43.473	14	3
43.419	14	3
43.485	14	3
43.464	15	3
43.460	15	3
43.557	15	3
43.472	15	3
43.411	15	3
43.571	15	3
43.668	16	4
43.712	16	4
43.675	16	4
43.684	16	4
43.712	16	4
43.762	16	4
43.758	17	4
43.750	17	4
43.665	17	4
43.750	17	4
43.732	17	4
43.739	17	4
43.709	18	4
43.717	18	4
43.660	18	4
43.705	18	4
43.739	18	4
43.688	18	4
43.633	19	4
43.715	19	4
43.691	19	4
43.634	19	4
43.654	19	4
43.684	19	4
43.645	20	4
43.664	20	4
43.695	20	4
43.601	20	4
43.726	20	4
43.708	20	4
43.585	21	5
43.686	21	5
43.715	21	5
43.682	21	5
43.648	21	5
43.718	21	5
43.696	22	5
43.712	22	5
43.734	22	5
43.682	22	5
43.675	22	5
43.678	22	5
43.681	23	5

CYCLE	MINOR	GROUP	MAJOR	GROUP
43.669	23		5	
43.681	23		5	
43.636	23		5	
43.658	23		5	
43.681	23		5	
43.657	24		5	
43.696	24		5	
43.627	24		5	
43.588	24		5	
43.634	24		5	
43.607	24		5	
43.663	25		5	
43.686	25		5	
43.659	25		5	
43.597	25		5	
43.609	25		5	
43.700	25		5	

Appendix F

Searles Lake core KM-3 Isotopic Results and
Isotopic Correction Factors

APPENDIX F

SEARLES LAKE CORE KM-3: DEPTH, AGE, O-18 & C-13 RESULTS and CORRECTIONS
 SAMPLES MARKED WITH 'XX' NOT USED IN RECONSTRUCTION DUE TO MINERALOGY

CORE DEPTH (ft)	CORE DEPTH (m)	C-13 DOL. (PDB)	O-18 DOL. (SMOW)	C-13 CORRECTIONS (per mil)	O-18	C-13 FINAL (PDB)	O-18 FINAL (SMOW)	AGE (Ma)
65.6	19.99	0.004	30.018	0.800	2.800	0.804	32.818	0.0102
66.1	20.15	2.383	36.988	-----	1.000	2.383	37.988	0.0106
66.6	20.30	2.905	34.463	-----	-----	2.905	34.463	0.0109
68.1	20.76	0.331	33.558	-----	1.000	0.331	34.558	0.0121
68.6	20.91	1.657	34.889	-----	-----	1.657	34.889	0.0125
69.1	21.06	0.677	35.569	-----	-----	0.677	35.569	0.0129
69.6	21.21	1.047	32.623	-----	-----	1.047	32.623	0.0133
70.1	21.37	1.357	34.228	-----	-----	1.357	34.228	0.0137
70.6	21.52	0.965	32.802	-----	-----	0.965	32.802	0.0141
71.0	21.64	1.961	32.964	-----	-----	1.961	32.964	0.0144
71.6	21.82	2.188	32.704	0.800	2.800	2.988	35.504	0.0149
72.1	21.98	2.701	33.525	0.800	2.800	3.501	36.325	0.0153
72.6	22.13	2.804	34.929	0.800	2.800	3.604	37.729	0.0156
73.1	22.28	3.122	35.847	-----	-----	3.122	35.847	0.0160
73.6	22.43	3.386	36.617	-----	-----	3.386	36.617	0.0164
74.1	22.59	3.424	34.055	0.800	2.800	4.224	36.855	0.0168
74.6	22.74	3.845	36.239	-----	-----	3.845	36.239	0.0172
75.3	22.95	4.491	37.373	0.800	2.800	5.291	40.173	0.0178
75.8	23.10	3.416	32.154	-----	-----	3.416	32.154	0.0182
80.0	24.38	3.515	37.715	-----	-----	3.515	37.715	0.0214
81.0	24.69	4.370	39.140	-----	-----	4.370	39.140	0.0222
81.6	24.87	3.719	35.356	-----	-----	3.719	35.356	0.0227
81.9	24.96	1.719	31.829	-----	-----	1.719	31.829	0.0229
89.5	27.28	3.537	32.878	0.800	2.800	4.337	35.678	0.0236
90.7	27.65	3.764	35.804	0.800	2.800	4.564	38.604	0.0247
91.5	27.89	3.890	36.372	0.800	2.800	4.690	39.172	0.0253
95.5	29.11	2.617	28.411	0.800	2.800	3.417	31.211	0.0260
96.6	29.44	1.830	26.935	0.800	2.800	2.630	29.735	0.0269
111.7	34.05	3.152	32.440	0.800	2.800	3.952	35.240	0.0290
113.3	34.53	3.799	33.907	0.800	2.800	4.599	36.707	0.0298
114.5	34.90	4.754	32.632	0.800	2.800	5.554	35.432	0.0306
116.4	35.48	4.245	33.476	0.800	2.800	5.045	35.276	0.0317
118.5	36.12	3.861	35.465	0.800	2.800	4.661	38.265	0.0319
120.0	36.58	4.081	35.993	0.800	2.800	4.881	38.793	0.0326
124.9	38.07	4.006	38.129	-----	1.000	4.006	39.129	0.0335
125.7	38.31	4.706	32.652	0.800	2.800	5.506	35.452	0.0356
126.3	38.50	4.070	38.909	-----	-----	4.070	38.909	0.0372
126.9	38.68	4.354	40.755	-----	-----	4.354	40.755	0.0388
127.7	38.92	3.412	38.345	-----	-----	3.412	38.345	0.0409
128.7	39.23	2.841	35.474	-----	-----	2.841	35.474	0.0435
128.9	39.29	3.775	39.432	-----	-----	3.775	39.432	0.0440
129.8	39.56	3.980	36.587	0.800	2.800	4.780	39.387	0.0464
130.1	39.65	3.412	39.934	-----	-----	3.412	39.934	0.0471
130.4	39.75	3.962	40.348	-----	-----	3.962	40.348	0.0480
131.5	40.08	3.575	37.927	-----	-----	3.575	37.927	0.0508
132.3	40.33	1.831	37.379	-----	-----	1.831	37.379	0.0530
134.4	40.97	3.659	36.075	-----	1.000	3.659	37.075	0.0585
136.4	41.57	3.407	37.320	-----	1.000	3.407	38.320	0.0637
137.5	41.91	4.278	37.353	-----	1.000	4.278	38.353	0.0659
139.0	42.37	2.226	36.486	-----	-----	2.226	36.486	0.0672
140.5	42.82	3.124	40.365	-----	-----	3.124	40.365	0.0685
141.8	43.22	2.885	37.364	-----	-----	2.885	37.364	0.0696
142.8	43.53	3.078	40.448	-----	-----	3.078	40.448	0.0705
151.1	46.06	4.981	39.621	-----	-----	4.981	39.621	0.0779
153.0	46.63	3.571	33.188	0.800	2.800	4.371	35.988	0.0796
154.1	46.97	3.589	33.052	0.800	2.800	4.389	35.852	0.0806
158.6	48.34	6.557	41.891	-----	-----	6.557	41.891	0.0846
164.0	49.99	6.237	41.768	-----	-----	6.237	41.768	0.0894
170.0	51.82	4.521	40.515	-----	-----	4.521	40.515	0.0952

CORE DEPTH (ft)	CORE DEPTH (m)	C-13 DOL. (PDB)	O-18 DOL. (SMOW)	C-13 CORRECTIONS (per mil)	O-18 CORRECTIONS (per mil)	C-13 FINAL (PDB)	O-18 FINAL (SMOW)	AGE (Ma)
178.5	54.41	4.568	37.689	-----	-----	4.568	37.689	0.1034
183.0	55.78	4.749	37.390	-----	-----	4.749	37.390	0.1072
187.4	57.12	5.592	38.764	-----	-----	5.592	38.764	0.1117
189.4	57.73	4.347	34.780	-----	1.000	4.347	35.780	0.1138
191.9	58.49	2.984	30.941	0.800	2.800	3.784	33.741	0.1163
194.9	59.41	5.319	40.123	-----	-----	5.319	40.123	0.1194
199.9	60.93	5.999	41.715	-----	-----	5.999	41.715	0.1247
205.0	62.48	5.098	38.463	-----	-----	5.098	38.463	0.1303
211.1	64.34	3.507	33.198	0.800	2.800	4.307	35.998	0.1370
218.0	66.45	5.956	40.727	-----	-----	5.956	40.727	0.1446
221.0	67.36	5.763	40.267	-----	-----	5.763	40.267	0.1479
224.1	68.31	6.425	41.175	-----	-----	6.425	41.175	0.1513
226.6	69.07	6.566	42.131	-----	-----	6.566	42.131	0.1540
256.1	78.06	5.590	38.712	-----	-----	5.590	38.712	0.1806
257.0	78.33	5.282	36.223	-----	-----	5.282	36.223	0.1814
277.5	84.58	4.921	34.702	-----	-----	4.921	34.702	0.2006
288.1	87.81	6.406	23.423	0.800	2.800	7.206	26.223	0.2122
298.1	90.86	2.369	24.906	0.800	2.800	3.169	27.706	0.2231
308.0	93.88	3.768	30.120	0.800	2.800	4.568	32.920	0.2303
XX 326.5	99.52	4.465	26.841	0.800	2.800	5.265	29.641	0.2629
XX 349.9	106.65	4.730	32.428	-----	-----	4.730	32.428	0.2710
XX 359.3	109.51	5.381	29.988	-----	-----	5.381	29.988	0.2735
369.5	112.62	4.890	23.163	0.800	2.800	5.690	25.963	0.2788
370.5	112.93	3.558	23.247	0.800	2.800	4.358	26.047	0.2804
371.8	113.32	4.073	21.541	0.800	2.800	4.873	24.342	0.2825
373.3	113.78	4.732	31.709	0.800	2.800	5.532	34.509	0.2849
375.0	114.30	4.965	27.420	0.800	2.800	5.765	30.220	0.2893
446.1	135.97	4.102	30.735	0.800	2.800	4.902	33.535	0.3552
447.5	136.40	3.892	28.274	0.800	2.800	4.692	31.074	0.3620
448.5	136.70	3.560	28.514	0.800	2.800	4.360	31.314	0.3668
449.4	136.98	3.599	26.006	0.800	2.800	4.399	28.806	0.3711
453.0	138.07	6.623	35.706	-----	-----	6.623	35.706	0.3760
XX 455.5	138.84	5.429	37.563	-----	-----	5.429	37.563	0.3810
XX 457.0	139.29	3.624	29.094	-----	-----	3.624	29.094	0.3828
459.5	140.06	4.951	26.749	0.800	2.800	5.751	29.544	0.3852
476.0	145.09	4.457	31.139	0.800	2.800	5.257	33.939	0.3905
481.0	146.61	3.899	32.837	0.800	2.800	4.699	35.637	0.3930
483.4	147.34	4.741	30.302	0.800	2.800	5.541	33.102	0.3985
484.4	147.65	4.372	30.112	0.800	2.800	5.172	32.912	0.4008
485.7	148.04	4.883	25.332	0.800	2.800	5.683	28.132	0.4038
486.0	148.13	4.848	28.408	0.800	2.800	5.648	31.208	0.4045
487.8	148.68	4.377	28.848	0.800	2.800	5.177	31.648	0.4087
489.0	149.05	3.761	31.085	0.800	2.800	4.561	33.885	0.4115
XX 490.0	149.35	4.079	31.244	-----	-----	4.079	31.000	0.4138
491.2	149.72	3.755	31.014	0.800	2.800	4.555	33.814	0.4165
492.0	149.96	2.985	30.480	0.800	2.800	3.785	33.280	0.4184
493.2	150.33	3.429	30.686	0.800	2.800	4.229	33.486	0.4212
493.9	150.54	2.711	35.253	0.800	2.800	3.511	38.053	0.4228
502.1	153.04	3.311	30.935	0.800	2.800	4.111	33.735	0.4301
546.1	166.45	2.765	31.910	0.800	2.800	3.565	34.710	0.5982
547.4	166.85	2.214	32.187	0.800	2.800	3.014	34.987	0.6004
547.7	166.94	1.945	32.628	0.800	2.800	2.745	35.428	0.6009
548.5	167.18	1.455	35.075	0.800	2.800	2.255	37.875	0.6022
549.5	167.49	2.285	33.221	0.800	2.800	3.085	36.021	0.6039
550.5	167.79	1.591	33.215	-----	1.000	1.591	33.415	0.6056
551.7	168.16	3.607	31.163	0.800	2.800	4.407	33.963	0.6076
552.6	168.43	1.878	28.600	0.800	2.800	2.678	31.400	0.6091
553.8	168.80	3.271	31.311	0.800	2.800	4.071	34.111	0.6111
555.0	169.16	1.634	32.290	-----	1.000	1.634	33.290	0.6132
556.0	169.47	3.003	32.705	-----	1.000	3.003	33.705	0.6149
557.1	169.80	4.053	38.224	-----	-----	4.053	38.224	0.6168
558.2	170.14	1.280	27.901	-----	-----	1.280	27.901	0.6187
559.2	170.44	1.964	32.186	-----	-----	1.964	32.186	0.6205
560.1	170.72	3.577	32.874	-----	-----	3.577	32.874	0.6220
561.0	170.99	2.147	35.478	-----	-----	2.147	35.478	0.6236

CORE DEPTH (ft)	CORE DEPTH (m)	C-13 DOL. (PDB)	O-18 DOL. (SMOW)	C-13 CORRECTIONS (per mil)	O-18	C-13 FINAL (PDB)	O-18 FINAL (SMOW)	AGE (Ma)
562.2	171.36	1.121	34.479	-----	1.000	1.121	35.479	0.6257
563.2	171.66	1.296	30.667	-----	-----	1.296	30.667	0.6274
564.3	172.00	2.333	32.120	-----	-----	2.333	32.120	0.6293
565.3	172.30	2.461	31.986	-----	-----	2.461	31.986	0.6310
567.0	172.82	2.027	33.316	-----	-----	2.027	33.316	0.6340
568.7	173.34	3.073	35.171	-----	-----	3.073	35.171	0.6369
569.9	173.71	6.209	40.770	-----	-----	6.209	40.770	0.6390
571.2	174.10	3.738	34.551	-----	-----	3.738	34.551	0.6412
572.5	174.50	4.834	39.142	-----	-----	4.834	39.142	0.6435
573.8	174.89	4.646	37.815	-----	-----	4.646	37.815	0.6457
574.9	175.23	5.332	39.691	-----	-----	5.332	39.691	0.6476
575.4	175.38	4.182	35.143	-----	-----	4.182	35.143	0.6485
576.4	175.69	4.677	37.685	-----	-----	4.677	37.685	0.6502
577.6	176.05	4.320	36.589	0.800	2.800	5.120	39.389	0.6523
578.6	176.36	4.971	30.339	-----	-----	4.971	30.339	0.6540
579.5	176.63	4.037	32.522	-----	-----	4.037	32.522	0.6556
580.7	177.00	4.475	34.149	-----	-----	4.475	34.149	0.6577
582.2	177.45	4.201	33.392	0.800	2.800	5.001	36.192	0.6603
XX 583.3	177.79	4.681	31.669	-----	-----	4.681	31.669	0.6622
XX 584.2	178.06	3.562	21.455	-----	-----	3.562	21.455	0.6637
587.8	179.16	2.644	32.359	0.800	2.800	3.444	35.159	0.6697
590.9	180.11	5.838	41.693	-----	-----	5.838	41.693	0.6794
XX 592.6	180.62	4.184	33.944	-----	-----	4.184	33.944	0.6848
594.8	181.30	5.488	42.246	-----	-----	5.488	42.246	0.6917
598.5	182.42	4.162	30.279	-----	-----	4.162	30.279	0.7034
XX 601.5	183.34	6.636	23.732	-----	-----	6.636	23.732	0.7128
XX 603.0	183.79	5.326	36.755	-----	-----	5.326	36.755	0.7175
XX 605.0	184.40	4.765	27.429	-----	-----	4.765	27.429	0.7238
XX 607.9	185.29	5.768	34.523	-----	-----	5.768	34.523	0.7339
XX 609.9	185.90	6.254	39.206	-----	-----	6.254	39.206	0.7422
XX 611.5	186.39	6.349	27.093	-----	-----	6.349	27.093	0.7489
XX 615.0	187.45	6.030	31.304	-----	-----	6.030	31.304	0.7604
624.0	190.20	4.279	28.077	0.800	2.800	5.079	30.877	0.7826
631.8	192.57	3.304	31.200	0.800	2.800	4.104	34.000	0.8037
634.2	193.30	4.220	25.842	-----	-----	4.220	25.842	0.8099
636.4	193.98	4.679	26.805	-----	-----	4.679	26.805	0.8156
XX 638.5	194.62	4.470	28.589	-----	-----	4.470	28.589	0.8210
641.1	195.41	5.353	30.612	-----	-----	5.353	30.612	0.8278
642.5	195.83	4.143	32.740	-----	1.000	4.143	33.740	0.8314
645.2	196.66	4.076	32.534	-----	1.000	4.076	33.534	0.8350
647.4	197.33	2.675	35.727	0.800	2.800	3.475	38.527	0.8402
651.4	198.55	3.440	36.050	0.800	2.800	4.240	38.850	0.8496
653.6	199.22	4.500	35.557	-----	-----	4.500	35.557	0.8548
659.2	200.92	4.736	33.899	-----	-----	4.736	33.899	0.8681
661.7	201.69	4.498	31.425	-----	-----	4.498	31.425	0.8740
664.8	202.63	5.707	38.055	-----	-----	5.707	38.055	0.8813
666.5	203.15	3.960	35.178	-----	-----	3.960	35.178	0.8854
668.4	203.73	5.562	39.513	-----	-----	5.562	39.513	0.8899
670.2	204.28	6.449	33.316	-----	-----	6.449	33.316	0.8941
671.5	204.67	3.988	31.171	-----	-----	3.988	31.171	0.8976
680.7	207.48	5.768	38.904	-----	-----	5.768	38.904	0.9042
684.7	208.70	6.016	39.346	-----	-----	6.016	39.346	0.9141
690.3	210.40	4.673	32.509	-----	-----	4.673	32.509	0.9278
XX 701.2	213.73	5.930	38.156	-----	-----	5.930	38.156	0.9352
706.2	215.25	3.374	34.371	0.800	2.800	4.174	37.171	0.9409
710.2	216.47	2.598	33.392	0.800	2.800	3.398	36.192	0.9454
713.4	217.44	3.035	36.275	0.800	2.800	3.835	39.075	0.9490
717.0	218.54	2.861	28.766	0.800	2.800	3.661	32.566	0.9530
724.9	220.95	3.403	29.145	0.800	2.800	4.203	31.945	0.9684
727.5	221.74	3.625	24.522	0.800	2.800	4.425	27.322	0.9718
747.2	227.75	6.008	34.406	-----	-----	6.008	34.406	0.9919
750.0	228.60	5.971	41.628	-----	-----	5.971	41.628	0.9948
XX 752.5	229.36	4.160	24.314	-----	-----	4.160	24.314	0.9973

CORE DEPTH (ft)	CORE DEPTH (m)	C-13 DOL. (PDB)	O-18 DOL. (SMOW)	C-13 CORRECTIONS (per mil)	O-18 CORRECTIONS (per mil)	C-13 FINAL (PDB)	O-18 FINAL (SMOW)	AGE (Ma)
758.0	231.04	6.272	39.973	-----	-----	6.272	39.973	1.0029
762.4	232.38	6.394	40.565	-----	-----	6.394	40.565	1.0074
766.5	233.63	3.581	35.114	-----	1.000	3.581	36.114	1.0116
772.2	235.37	6.234	38.277	-----	-----	6.234	38.277	1.0174
775.8	236.46	5.773	38.988	-----	-----	5.773	38.988	1.0211
780.8	237.99	4.861	30.802	-----	-----	4.861	30.802	1.0261
785.9	239.54	5.964	38.269	-----	-----	5.964	38.269	1.0313
794.2	242.07	6.178	40.195	-----	-----	6.178	40.195	1.0398
798.7	243.44	6.063	38.766	-----	-----	6.063	38.766	1.0444
805.0	245.36	4.262	32.830	-----	-----	4.262	32.830	1.0508
814.0	248.11	4.546	33.949	-----	-----	4.546	33.949	1.0600
818.2	249.39	5.270	33.377	-----	1.000	5.270	34.377	1.0637
823.6	251.03	4.384	32.283	0.800	2.800	5.184	35.083	1.0718
827.0	252.07	5.173	33.282	-----	-----	5.173	33.282	1.0771
831.7	253.50	5.348	36.596	-----	-----	5.348	36.596	1.0843
837.0	255.12	6.809	37.489	-----	-----	6.809	37.489	1.0924
841.6	256.52	6.166	37.373	-----	-----	6.166	37.373	1.0995
846.6	258.04	4.465	35.381	-----	1.000	4.465	36.381	1.1072
859.8	262.07	4.366	33.866	-----	-----	4.366	33.866	1.1274
864.0	263.35	6.830	40.501	-----	-----	6.830	40.501	1.1339
876.2	267.07	6.735	38.187	-----	-----	6.735	38.187	1.1526
XX 880.9	268.50	NO CARBONATE		-----	-----	-----	-----	1.1598
887.6	270.54	4.786	33.328	-----	1.000	4.786	34.328	1.1670
893.9	272.46	4.384	31.279	0.800	2.800	5.184	34.079	1.1757
898.9	273.99	4.473	29.276	0.800	2.800	5.273	32.076	1.1840
903.1	275.27	6.496	36.371	-----	-----	6.496	36.371	1.1910
XX 908.6	276.94	4.099	33.032	-----	-----	4.099	33.032	1.2002
917.1	279.53	5.022	34.559	0.800	2.800	5.822	37.359	1.2126
922.0	281.03	4.839	34.751	-----	1.000	4.839	35.751	1.2198
930.0	283.46	4.408	34.554	-----	-----	4.408	34.554	1.2315
935.7	285.20	2.905	35.058	-----	-----	2.905	35.058	1.2398
939.9	286.48	2.759	32.046	-----	-----	2.759	32.046	1.2459
948.0	288.95	2.330	34.153	0.800	2.800	3.130	36.953	1.2578
955.0	291.08	3.307	35.125	-----	-----	3.307	35.125	1.2680
959.0	292.30	1.909	35.548	0.800	2.800	2.709	38.348	1.2693
966.0	294.44	0.290	32.647	0.800	2.800	1.090	35.447	1.2730
972.1	296.30	2.091	29.356	-----	-----	2.091	29.356	1.2909
977.0	297.79	0.439	33.098	-----	-----	0.439	33.098	1.3053
986.0	300.53	-0.045	34.680	0.800	2.800	0.755	37.480	1.3218
1006.0	306.63	3.369	33.397	0.800	2.800	4.169	36.197	1.3312
1012.0	308.46	-0.076	31.653	0.800	2.800	0.724	34.453	1.3350
1017.0	309.98	4.825	31.282	-----	-----	4.825	31.282	1.3403
1034.2	315.22	5.640	36.016	-----	-----	5.640	36.016	1.3585
1068.0	325.53	-9.314	23.058	-----	-----	-9.314	23.058	1.4080
1076.0	327.97	-1.436	29.713	0.800	2.800	-0.636	32.513	1.4101
1080.4	329.31	-2.507	27.552	0.800	2.800	-1.707	30.352	1.4244
1089.0	331.93	-0.672	31.240	0.800	2.800	0.128	34.040	1.4366
1093.0	333.15	4.686	40.200	0.800	2.800	5.486	43.000	1.4500
1099.0	334.98	-5.207	28.305	0.800	2.800	-4.407	31.105	1.4535
1104.0	336.50	-5.748	26.033	0.800	2.800	-4.948	28.833	1.4712
1106.5	337.26	-4.479	25.626	0.800	2.800	-3.679	28.426	1.4800
1110.6	338.51	-6.709	28.633	0.800	2.800	-5.909	31.433	1.4846
1115.3	339.94	-4.257	24.794	0.800	2.800	-3.457	27.594	1.4991
1120.0	341.38	-3.085	27.424	0.800	2.800	-2.285	30.224	1.5112
1125.5	343.05	-0.470	29.556	0.800	2.800	0.330	32.356	1.5284
1130.5	344.58	3.095	33.892	-----	-----	3.095	33.892	1.5441
1132.7	345.25	3.498	34.063	-----	-----	3.498	34.063	1.5510
1156.5	352.50	-0.116	29.908	0.800	2.800	0.684	32.708	1.5762
1210.7	369.02	0.422	32.659	0.800	2.800	1.222	35.459	1.6230
1294.0	394.41	-1.131	28.458	0.800	2.800	-0.331	31.258	1.8127

Appendix G

Searles Lake Core KM-3 Isotopic Results for
Bulk Leached Samples Run at University of Missouri-Columbia

 *
 * APPENDIX G *
 *
 * SEARLES LAKE CORE KM-3: DEPTH, AGE, O-18 & C-13 RESULTS FOR BULK *
 * LEACHED SAMPLES RUN AT UNIVERSITY OF MISSOURI-COLUMBIA. *
 *

CORE DEPTH (ft)	CORE DEPTH (m)	C-13 DOL (PDB)	O-18 DOL (SMOW)	AGE (Ma)
96.6	29.44	2.38	29.59	.0269
124.9	38.07	4.06	34.99	.0335
125.7	38.31	4.04	31.161	.0356
126.3	38.50	2.52	33.34	.0372
126.9	38.68	3.93	38.92	.0388
127.7	38.92	2.964	36.033	.0409
128.7	39.29	2.83	34.22	.0440
130.1	39.65	2.51	28.35	.0471
130.4	39.75	3.6	37.65	.0480
132.3	40.33	1.75	33.21	.0530
134.4	40.97	3.40	34.44	.0585
136.4	41.57	2.63	35.71	.0637
137.5	41.91	4.05	34.61	.0659
139.0	42.37	1.52	34.15	.0672
140.5	42.82	2.79	38.084	.0685
141.8	43.22	3.77	35.93	.0696
142.8	43.53	2.82	38.97	.0705
481.0	146.61	3.34	31.97	.3930
481.0	146.61	3.36	31.36	.3930
492.0	149.96	1.88	29.21	.4148
493.9	150.54	2.14	32.24	.4228
502.1	153.04	2.204	28.63	.4301
546.1	166.45	2.56	30.94	.5982
547.4	166.85	2.00	30.14	.6004
547.7	166.94	1.364	30.88	.6009
548.5	167.18	1.62	31.95	.6022
548.5	167.18	1.36	32.11	.6022
549.5	167.49	1.63	30.76	.6039
550.5	167.79	1.518	29.558	.6056
551.7	168.16	3.36	30.15	.6076
552.6	168.43	2.54	29.08	.6091
553.8	168.80	2.85	29.29	.6111
555.0	169.16	1.76	30.61	.6132
556.0	169.47	2.88	28.56	.6149
557.1	169.80	3.71	35.12	.6168
558.2	170.14	1.07	27.13	.6187
559.2	170.44	1.34	21.52	.6205
560.1	170.72	3.32	30.79	.6220
561.0	170.99	1.13	30.81	.6236
562.2	171.36	2.31	33.41	.6257
562.2	171.36	2.58	33.14	.6257
563.2	171.66	1.04	28.27	.6274
564.3	172.00	1.56	28.19	.6293
564.3	172.00	1.475	28.924	.6293
565.3	172.30	2.67	30.98	.6310
567.0	172.82	1.73	29.70	.6340
568.7	173.34	2.90	31.64	.6369
569.9	173.71	7.19	41.13	.6390
571.2	174.10	1.88	28.45	.6412
572.5	174.50	4.94	38.40	.6435
574.9	175.23	4.51	36.31	.6476
575.4	175.38	3.19	30.94	.6485
576.4	175.69	4.58	36.02	.6502
577.6	176.05	3.68	33.66	.6523
578.6	176.36	4.19	27.37	.6540
578.6	176.36	4.46	37.68	.6540
580.7	177.00	3.95	30.20	.6577

CORE DEPTH (ft)	CORE DEPTH (m)	C-13 DOL (PDB)	O-18 DOL (SMOW)	AGE (Ma)
582.2	177.45	3.62	31.19	.6603
583.3	177.79	6.05	32.22	.6622
584.2	178.06	2.56	19.57	.6637
587.8	179.16	2.36	28.75	.6697
590.9	180.11	5.38	39.7	.6794
590.9	180.11	5.73	40.35	.6794
592.6	180.62	3.426	30.436	.6848
636.4	193.98	1.28	23.00	.8156
636.4	193.98	2.82	25.15	.8156
638.5	194.62	3.49	27.60	.8210
641.1	195.41	4.21	28.59	.8278
642.5	195.83	2.94	31.28	.8314
651.4	198.55	3.32	32.73	.8496
653.6	199.22	3.17	30.02	.8548
659.2	200.92	4.06	29.87	.8681
661.7	201.69	3.72	29.62	.8740
666.5	203.15	4.36	31.58	.8854
668.4	203.73	4.52	35.82	.8899
671.5	204.67	3.64	28.63	.8976
701.2	213.73	4.51	34.32	.9352
706.2	215.25	1.43	28.98	.9409
710.2	216.47	1.87	30.40	.9454
713.4	217.44	2.95	42.30	.9490
750.0	228.60	5.80	40.00	.9948
766.5	233.63	2.25	30.48	1.0116
772.2	235.37	5.17	35.01	1.0174
798.7	243.44	4.74	34.43	1.0444
1006.0	306.63	3.84	33.11	1.3312
1012.0	308.46	0.78	32.05	1.3350
1017.0	309.98	4.17	29.86	1.3403
1076.0	327.97	-0.95	30.24	1.4101
1080.4	329.31	-1.91	29.03	1.4244
1089.0	331.93	0.63	33.13	1.4366
1093.0	333.15	4.75	38.95	1.4500
1099.0	334.98	-5.14	27.06	1.4535

Appendix H

Analytical-Solution Lake Isotope Evolution Model and Results

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C*****
C*
C* THIS PROGRAM COMPUTES FRED'S ANALYTICAL SOLUTIONS FOR CLOSED-BASIN
C* LAKE VOLUMES AND ISOTOPIC COMPOSITION. YOU WILL BE ASKED TO INPUT
C* ALL PARAMETERS- PAY CAREFUL ATTENTION TO UNITS!
C*
C*****
PROGRAM CBLAKES
CHARACTER FNAME*50,ANS*1,ANS2*1

C
WRITE(6,*)'THIS PROGRAM COMPUTES FREDS ANALYTICAL
1 SOLUTIONS FOR'
WRITE(6,*)'CLOSED-BASIN LAKE VOLUMES AND ISOTOPIC COMPOSITION.'
WRITE(6,*)'YOU WILL BE ASKED TO INPUT ALL PARAMETERS- PAY
1 CAREFUL'
WRITE(6,*)' ATTENTION TO UNITS!'

C
C
C
WRITE(6,*)'ENTER OUTPUT FILE NAME'
READ(5,'(A)')FNAME
OPEN(UNIT=95,FILE=FNAME,STATUS='NEW',CARRIAGECONTROL='LIST')

10 WRITE(6,*)'INPUT TYPE OF EQUATIONS USED (R,I,E)'
WRITE(6,*)' R= RAMP INFLOW EQUATIONS'
WRITE(6,*)' I= INSTANTANEOUS INFLOW EQUATIONS'
WRITE(6,*)' E= EXIT PROGRAM'
READ(5,'(A)')ANS
IF((ANS.EQ.'R').OR.(ANS.EQ.'r'))THEN
WRITE(6,*)'STATE SITUATION (I,D,E)'
WRITE(6,*)' I= INCREASING OR DECREASING INFLOW'
WRITE(6,*)' D= DESSICATION:INITIAL AND FINAL INFLOW=0'
WRITE(6,*)' E= EXIT RAMP EQUATIONS'
READ(5,'(A)')ANS2
IF((ANS2.EQ.'I').OR.(ANS2.EQ.'i'))THEN
CALL RAMP
ELSEIF((ANS2.EQ.'D').OR.(ANS2.EQ.'d'))THEN
CALL DESSICATE
ELSEIF((ANS2.EQ.'E').OR.(ANS2.EQ.'e'))THEN
GO TO 10
ENDIF
ELSEIF((ANS.EQ.'I').OR.(ANS.EQ.'i'))THEN
WRITE(6,*)'STATE SITUATION (I,D,E)'
WRITE(6,*)' I= INCREASING OR DECREASING INFLOW'
WRITE(6,*)' D= DESSICATION:INITIAL AND FINAL INFLOW=0'
WRITE(6,*)' E= EXIT INSTANTANEOUS INFLOW EQUATIONS'
READ(5,'(A)')ANS2
IF((ANS2.EQ.'I').OR.(ANS2.EQ.'i'))THEN
CALL INSTANT
ELSEIF((ANS2.EQ.'D').OR.(ANS2.EQ.'d'))THEN
CALL DESSICATE
ELSEIF((ANS2.EQ.'E').OR.(ANS2.EQ.'e'))THEN
GO TO 10
ENDIF
ELSEIF((ANS.EQ.'E').OR.(ANS.EQ.'e'))THEN
100 ENDIF
STOP
END

C
C*****
C
SUBROUTINE RAMP
COMMON /STUFF/ TEMP,ELVS,EVAP,AREA,VOL,DK,H,DELO,VO,DELC,CAREPS
COMMON /RAMPSTUFF/ QIO,QF,LRAMP,BETA
COMMON /BOTH/ DELI
CC
C CHARACTER TITLE*60,ANS3*1
C DIMENSION RVOL(0:1000),DLAKE(0:1000),DELOF(0:1000)
C DIMENSION CARB(0:1000)

```

```

C      COMING TO A THEATRE NEAR YOU...
      RETURN
      END
C
C*****
C*****
      SUBROUTINE INSTANT
      COMMON /STUFF/ TEMP,ELVS,EVAP,AREA,VOL,DK,H,DELO,VO,DELC,CAREPS
      COMMON /QINSTANT/ QI
      COMMON /BOTH/ DELI
C
      CHARACTER TITLE*60,ANS2*1
      DIMENSION RVOL(0:3000),DLAKE(0:3000),DELOF(0:3000)
      DIMENSION CARB(0:3000)
C
      WRITE(6,*)'DESCRIBE SITUATION (60 CHARACTERS OR LESS)!'
      READ(5, '(A)')TITLE
      WRITE(95,*)'*****!'
      WRITE(95,*)
      WRITE(95,*)TITLE
      WRITE(95,*)
      WRITE(95,*)'*****!'
C
      CALL PARAMS
C
C
C      ***** DETERMINE IF OVERFLOW WILL OCCUR WITH INFLOW *****
C
12     FORMAT(1x,A5,7x,A6,7x,A12,4x,A8)
13     FORMAT(1x,15,3x,1e13.7,5x,f11.7,9x,f8.5)
C
      V=QI/((1-H)*DK)
      IF(VOL.EQ.VO)THEN
      KOFTIM=LOG((VOL-V)/(VO-V))/(-(1-H)*DK)
      WRITE(5,*)'LAKE WILL OVERFLOW'
      WRITE(95,*)
      WRITE(95,12)'TIME ', 'VOLUME', 'DEL 0-18 H2O', ' 0-18 CARB'
      WRITE(95,12)'(yrs)', ' (m^3)', ' (SMOW) ', ' (SMOW) '
      WRITE(95,*)
      WRITE(95,*)
      WRITE(95,*)'***** OVERFLOW *****!'
      WRITE(95,*)
C
C
C      COMPUTE ISOTOPIC COMPOSITION OF OVERFLOWING LAKE
C
      QO=QI-DK*VOL*(1-H)
      KTIM=0
      P1=DK/(ELVS+1) + QO/VOL
      P2=QI*DELI/VOL + DK*(ELVS/(ELVS+1) + H*DELC)
      DO 250 J=0,500,10
      RVOL(J)=(VO-V)*EXP(-(1-H)*DK*KTIM)+V
      DELOF(J)=(P2/P1 + (DELO-P2/P1)
      *EXP((-P1)*KTIM))*1000.
      CARB(J)=CAREPS + DELOF(J)
      WRITE(95,13)KTIM,VOL,DELOF(J),CARB(J)
      KTIM=KTIM+10
      IF(RVOL(J).LT.VOL)THEN
      DELBSW=DELOF(J)/1000.
      KBSW=2000-KTIM+10
      GO TO 251
C
      ENDIF
250    CONTINUE
C
C      COMPUTE LAKE VOLUME AND ISO COMP. LAKE BENEATH SPILLWAY
C
251    WRITE(95,*)
      WRITE(95,*)'***** LAKE LEVEL BENEATH SPILLWAY *****!'
      WRITE(95,*)
      KTIM=0
      A=(ELVS+1)/DK

```



```

      B=(ELVS+DELC*H*(ELVS+1))/(1-(ELVS+1)*(1-H))
      D2=QI*(DELI*A+B*(VO-QI*A))
DO 260 J=0,KBSW,100
      RVOL(J)=(VO-V)*EXP(-(1-H)*DK*KTIM) + V
      C=EXP(-(DK*KTIM/(ELVS+1)))
      DLAKE(J)=((QI*A*(DELI-B)+(VO*(DELBSW-B)-QI*A
1          *(DELI-B))*C)/RVOL(J)+B)*1000.
      CARB(J)=CAREPS + DLAKE(J)
      WRITE(95,13)KTIM,RVOL(J),DLAKE(J),CARB(J)
      KTIM=KTIM+100
      IF(KTIM.GE.KBSW)THEN
          RVOL(KBSW)=(VO-V)*EXP(-(1-H)*DK*KBSW) + V
          C=EXP(-(DK*KBSW/(ELVS+1)))
          DLAKE(KBSW)=((QI*A*(DELI-B)+(VO*(DELBSW-B)
1          -QI*A*(DELI-B))*C)/RVOL(KBSW)+B)*1000.
          CARB(KBSW)=CAREPS+DLAKE(KBSW)
          WRITE(95,13)KBSW,RVOL(KBSW),DLAKE(KBSW),
1          CARB(KBSW)
      ENDIF
260      CONTINUE
      COMPUTE STEADY STATE DEL LAKE
      FV=QI/((1-H)*DK)
      SS1=QI*DELI*(ELVS+1)/(DK*FV) + ELVS + H*DELC*(ELVS+1)
      SS=SS1*1000.
      DOLSS=CAREPS + SS
      WRITE(95,*)'          STEADY STATE VALUES:NON-OVERFLOW'
      WRITE(95,*)'          LAKE VOLUME',FV
      WRITE(95,*)'          STEADY STATE DEL H2O',SS
      WRITE(95,*)'          STEADY STATE CARBONATE',DOLSS
      ELSEIF(V.GT.VOL)THEN
      KOFTIM=LOG((VOL-V)/(VO-V))/(-(1-H)*DK)
      WRITE(5,*)'LAKE WILL OVERFLOW'
      COMPUTE LAKE VOLUME AND ISO COMPOSITION UNTIL OVERFLOW
      WRITE(95,*)
      WRITE(95,12)'TIME ', 'VOLUME', 'DEL 0-18 H2O', ' 0-18 CARB'
      WRITE(95,12)'(yrs)', '(m^3)', '(SMOW) ', '(SHOW) '
      WRITE(95,*)
      KTIM=0
      A=(ELVS+1)/DK
      B=(ELVS+DELC*H*(ELVS+1))/(1-(ELVS+1)*(1-H))
DO 150 J=0,KOFTIM+1,10
      RVOL(J)=(VO-V)*EXP(-(1-H)*DK*KTIM) + V
      C=EXP(-(DK*KTIM/(ELVS+1)))
      IF((VO.EQ.0).AND.(KTIM.EQ.0))THEN
          DLAKE(0)=DELI*1000.
          CARB(0)=CAREPS+DLAKE(0)
      ELSE
1          DLAKE(J)=((QI*A*(DELI-B)+(VO*(DELO-B)-QI*A
          *(DELI-B))*C)/RVOL(J)+B)*1000.
          CARB(J)=CAREPS+DLAKE(J)
      ENDIF
      WRITE(95,13)KTIM,RVOL(J),DLAKE(J),CARB(J)
      KTIM=KTIM+10
      IF(KTIM.GE.KOFTIM+1)THEN
          RVOL(KOFTIM+1)=(VO-V)*
1          EXP(-(1-H)*DK*(KOFTIM+1))+V
          C=EXP(-(DK*(KOFTIM+1)/(ELVS+1)))
          DLAKE(KOFTIM+1)=((QI*A*(DELI-B)+(VO*(DELO
1          -B)-QI*A*(DELI-B))*C)/RVOL(KOFTIM+1)+B)
1          *1000.
          CARB(KOFTIM+1)=CAREPS+DLAKE(KOFTIM+1)
          WRITE(95,13)KOFTIM,RVOL(KOFTIM+1),
1          DLAKE(KOFTIM+1),CARB(KOFTIM+1)
          DELOV=(DLAKE(KOFTIM+1))/1000.
      ENDIF

```

```

150      CONTINUE
C
C      COMPUTE ISOTOPIIC COMPOSITION OF OVERFLOWING LAKE
C
      WRITE(95,*)
      WRITE(95,*)' ***** OVERFLOW *****'
      WRITE(95,*)
      QO=QI-DK*VOL*(1-H)
      KTIM=0
      P1=DK/(ELVS+1) + QO/VOL
      P2=QI*DELI/VOL + DK*(ELVS/(ELVS+1) + H*DELC)
      DO 160 J=0,500,50
1         DELOF(J)=(P2/P1 + (DELOV-P2/P1)
              *EXP((-P1)*KTIM))*1000.
              CARB(J)=CAREPS+DELOF(J)
              WRITE(95,13)KTIM,VOL,DELOF(J),CARB(J)
              KTIM=KTIM+50
160      CONTINUE
C
C      COMPUTE STEADY STATE DEL LAKE-OVERFLOW
C
      SS1=QI*DELI + DK*VOL*((ELVS/(ELVS+1)) + H*DELC)
      SS2=QI + DK*VOL*(H-1+(1/(ELVS+1)))
      SSDELO=(SS1/SS2)*1000.
      DOL=CAREPS+SSDELO
      WRITE(95,*)'          STEADY STATE VALUES-OVERFLOW'
      WRITE(95,*)
      WRITE(95,*)'          OVERFLOW TIME',KOFTIM
      WRITE(95,*)'          LAKE VOLUME',VOL
      WRITE(95,*)'          STEADY STATE DEL H2O',SSDELO
      WRITE(95,*)'          STEADY STATE CARBONATE',DOL
      WRITE(95,*)'          OUTFLOW RATE',QO
C
C      COMPUTE LAKE VOLUME FOR NON-OVERFLOWING LAKE'
C
      ELSE
      WRITE(95,*)
      WRITE(95,12)'TIME ', 'VOLUME', 'DEL 0-18 H2O', ' 0-18 CARB'
      WRITE(95,12)'(yrs)', '(m^3)', '(SHOW) ', '(SHOW) '
      WRITE(95,*)
      KTIM=0
      A=(ELVS+1)/DK
      B=(ELVS+DELC*H*(ELVS+1))/(1-(ELVS+1)*(1-H))
      DO 170 J=0,2500,100
1         RVOL(J)=(VO-V)*EXP(-(1-H)*DK*KTIM) + V
              C=EXP(-(DK*KTIM/(ELVS+1)))
              IF((VO.EQ.0).AND.(KTIM.EQ.0))THEN
              DLAKE(J)=DELI*1000.
              CARB(J)=CAREPS+DLAKE(J)
              ELSE
              DLAKE(J)=((QI*A*(DELI-B) + (VO*(DELO-B)-
              QI*A*(DELI-B))*C)/RVOL(J) + B)*1000.
              CARB(J)=CAREPS+DLAKE(J)
              ENDIF
              WRITE(95,13)KTIM,RVOL(J),DLAKE(J),CARB(J)
              KTIM=KTIM+100
              IF(KTIM.GT.2500)THEN
              ENDIF
170      CONTINUE
C
C      COMPUTE STEADY STATE DEL LAKE
C
      FV=QI/((1-H)*DK)
      SS1=QI*DELI*(ELVS+1)/(DK*FV) + ELVS + H*DELC*(ELVS+1)
      SSDELO=SS1*1000.
      DOL=CAREPS+SSDELO
      WRITE(95,*)'          STEADY STATE VALUES:NON-OVERFLOW'
      WRITE(95,*)

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        WRITE(95,*)'
        WRITE(95,*)'
        WRITE(95,*)'
                                LAKE VOLUME',FV
                                STEADY STATE DEL H2O',SSDELO
                                STEADY STATE CARBONATE',DOL
    ENDIF
    RETURN
    END
C*****
C*****
    SUBROUTINE DESSICATE
    COMMON /STUFF/ TEMP,ELVS,EVAP,AREA,VOL,DK,H,DELO,VO,DELC,CAREPS
C
    CHARACTER TITLE*60
    DIMENSION RVOL(0:2100),DLAKE(0:2100),CARB(0:2100)
C
    WRITE(6,*)'DESCRIBE SITUATION (60 CHARACTERS OR LESS)'
    READ(5,'(A)')TITLE
    WRITE(95,*)'*****
    WRITE(95,*)
    WRITE(95,*)TITLE
    WRITE(95,*)
    WRITE(95,*)'*****
C
    CALL PARAMS
C
C
C
C
    COMPUTE VOLUME AND ISO COMP OF DESSICATING LAKE
12  FORMAT(1x,A5,7x,A6,7x,A12,4x,A8)
13  FORMAT(1x,I5,3x,1e13.7,5x,f11.7,9x,f8.5)
    KTIM=0
    A=(ELVS+DELC*H*(ELVS+1))/(1-(ELVS+1)*(1-H))
    WRITE(95,*)
    WRITE(95,12)'TIME ', 'VOLUME', 'DEL O-18 H2O', ' O-18 CARB'
    WRITE(95,12)'(YRS)', ' (M^3)', ' (SMOW) ', ' (SMOW) '
    WRITE(95,*)
    DO 180 J=0,2000,100
        RVOL(J)=VO*EXP(-(1-H)*DK*KTIM)
        DLAKE(J)=((1/RVOL(J))*(VO*(DELO-A)*EXP(-DK*KTIM/(ELVS+1
1          ))) + A)*1000.
        CARB(J)=CAREPS+DLAKE(J)
        WRITE(95,13)KTIM,RVOL(J),DLAKE(J),CARB(J)
        KTIM=KTIM+100
180  CONTINUE
    RETURN
    END
C*****
C*****
C
    SUBROUTINE PARAMS
C
    COMMON /STUFF/ TEMP,ELVS,EVAP,AREA,VOL,DK,H,DELO,VO,DELC,CAREPS
    COMMON /RAMPSTUFF/ QIO,QF,LRAMP,BETA
    COMMON /QINSTANT/ QI
    COMMON /BOTH/ DELI
C
    CHARACTER ANS*1,ANS3*1,ANS4*1,ANS5*1
C
C
C
C
    INPUT TEMPERATURE AND COMPUTE ELV ACCORDING TO BOTTINGA AND CRAIG(1969)
    WRITE(6,*)'ENTER TEMPERATURE IN DEGREES CELCIUS'
    READ(5,*)TEMP
    write(95,*)'TEMPERATURE IN DEGREES CELCIUS IS',TEMP
    TEMP=TEMP + 273.4
    ELV=1.534*(1e6/TEMP**2)-3.206*(1e3/TEMP)+2.644
    WRITE(95,*)'ELV =',ELV
    WRITE(6,*)'IS THIS RUN ASSUMING DELC=DELI?(Y,N)'
    READ(5,'(A)')ANS5
        IF((ANS5.EQ.'Y').OR.(ANS5.EQ.'y'))THEN
    WRITE(6,*)'ENTER ISOTOPIC COMP. OF BACK CONDENS. FLUX(PER MIL)'
    READ(5,*)DELC
        ELSEIF((ANS5.EQ.'N').OR.(ANS5.EQ.'n'))THEN

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WRITE(6,*)'ENTER ISOTOPIC COMPOSITION OF ATMOSPHERE'
READ(5,*)DELA
DELC=ELV*(1+(DELA/1000))+DELA
ENDIF
WRITE(95,*)'ISOTOPIC COMP. OF BACK CONDENSATION FLUX IS ',DELC
DELC=DELC/1000.
WRITE(6,*)'ENTER RELATIVE HUMIDITY'
READ(5,*)H
WRITE(95,*)'RELATIVE HUMIDITY IS',H
ALPHA=EXP(ELV/1000)
ELVS=((1.0/ALPHA)*(1.0 - 0.0069)/(1.0-0.0069*H))-1
ELVS=ELVS*(-1)
WRITE(95,*)'ELV-STAR =',ELVS
C
WRITE(6,*)' COMPUTE CARBONATE VALUES ALONG WITH WATER? (Y,N)'
READ(5,'(A)')ANS3
IF((ANS3.EQ.'Y').OR.(ANS3.EQ.'y'))THEN
WRITE(6,*)'CALCITE OR DOLOMITE? (C,D)'
READ(5,'(A)')ANS4
IF((ANS4.EQ.'D').OR.(ANS4.EQ.'d'))THEN
WRITE(95,*)'COMPUTED CARBONATE VALUES FOR DOLOMITE'
C
COMPUTE DOLOMITE ELV ACCORDING TO NORTHROP & CLAYTON (1966)
C
CAREPS=3.20*(1.E6)/(TEMP**2) - 1.50
C
ELSEIF((ANS4.EQ.'C').OR.(ANS4.EQ.'c'))THEN
WRITE(95,*)'COMPUTED CARBONATE VALUES FOR CALCITE'
C
COMPUTE CALCITE EPSILON ACCORDING TO O'NEIL,CLAYTON & MAYEDA(1969)
C
CAREPS=2.78*(1.E6)/(TEMP**2)-2.89
ENDIF
ELSEIF((ANS3.EQ.'N').OR.(ANS3.EQ.'n'))THEN
ENDIF
C
INPUT OTHER INITIAL PARAMETERS
C
WRITE(6,*)'ENTER EVAPORATION RATE (m/yr)'
read(5,*)EVAP
WRITE(95,*)'EVAPORATION RATE IS (m/yr) ',EVAP
C
INPUT LAKE GEOMETRY
C
WRITE(6,*)'ENTER MAX. LAKE SURFACE AREA(EST. HIGH LAKE STAND)(m^2)'
read(5,*)AREA
WRITE(95,*)'ESTIMATED MAXIMUM LAKE SURFACE AREA IS(m^2)',AREA
WRITE(6,*)'ENTER MAX. LAKE VOLUME (EST. HIGH LAKE STAND)(m^3)'
READ(5,*)VOL
WRITE(95,*)'ESTIMATED MAXIMUM LAKE VOLUME IS (m^3)',VOL
C
COMPUTE K
C
DK=(AREA/VOL)*EVAP
WRITE(95,*)'K= ',DK
C
ENTER YET MORE INITIAL PARAMETERS
C
WRITE(6,*)'ENTER INITIAL LAKE ISO COMPOSITION (PER MIL)'
READ(5,*)DELO
WRITE(95,*)'INITIAL LAKE ISO COMPOSITION IS(PER MIL)',DELO
DELO=DELO/1000.
WRITE(6,*)'ENTER INITIAL LAKE VOLUME(FLUID IN LAKE)(m^3)'
READ(5,*)VO
WRITE(95,*)'INITIAL LAKE VOLUME (m^3) IS',VO
C
C
WRITE(6,*)'INPUT TYPE EQUATIONS USED ( R,I,D )'
WRITE(6,*)' R = RAMP EQUATIONS'

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WRITE(6,*)' I = INSTANTANEOUS INFLOW'
WRITE(6,*)' D = DESSICATION:INITIAL & FINAL INFLOW=0'
READ(5,'(A)')ANS
IF((ANS.EQ.'R').OR.(ANS.EQ.'r'))THEN
  WRITE(6,*)'ENTER INITIAL INFLOW RATE (m^3/yr)'
  read(5,*)QIO
  WRITE(95,*)'INITIAL INFLOW RATE IS (m^3/yr)',QIO
  WRITE(6,*)'ENTER FINAL INFLOW RATE (m^3/yr)'
  READ(5,*)QF
  WRITE(95,*)'FINAL INFLOW RATE IS (m^3/yr)',QF
  WRITE(6,*)'ENTER TIME RAMP (YEARS)'
  READ(5,*)LRAMP
  WRITE(95,*)'TIME RAMP IN YEARS IS',LRAMP

  COMPUTE BETA

  BETA=(QF-QIO)/LRAMP
  WRITE(95,*)'BETA =',BETA
  WRITE(6,*)'INPUT ISOTOPIIC COMP.
6          OF INFLOW WATER(PER MIL)'
  READ(5,*)DELI
  WRITE(95,*)'ISOTOPIIC COMP OF INFLOW WATER IS',DELI
  DELI=DELI/1000.
ELSEIF((ANS.EQ.'I').OR.(ANS.EQ.'i'))THEN
  WRITE(6,*)'ENTER INFLOW RATE (m^3/yr)'
  read(5,*)QI
  WRITE(95,*)' INFLOW RATE IS (m^3/yr)',QI
  WRITE(6,*)'INPUT ISOTOPIIC COMP.OF INFLOW WATER(PER MIL)'
  READ(5,*)DELI
  WRITE(95,*)'ISOTOPIIC COMP OF INFLOW WATER IS',DELI
  DELI=DELI/1000.
ELSEIF((ANS.EQ.'D').OR.(ANS.EQ.'d'))THEN
ENDIF
RETURN
END

```

C
C
C

C*****

INFILL OWENS FROM DRY LAKE; DELI=DELC

TEMPERATURE IN DEGREES CELCIUS IS 15.50000
 ELV = 9.926092
 ISOTOPIC COMP. OF BACK CONDENSATION FLUX IS -16.25000
 RELATIVE HUMIDITY IS 0.2780000
 ELV-STAR = 1.4819145E-02
 COMPUTED CARBONATE VALUES FOR DOLOMITE
 EVAPORATION RATE IS (m/yr) 1.177000
 ESTIMATED MAXIMUM LAKE SURFACE AREA IS (m^2) 6.9400000E+08
 ESTIMATED MAXIMUM LAKE VOLUME IS (m^3) 3.0020000E+10
 K= 2.7209796E-02
 INITIAL LAKE ISO COMPOSITION IS (PER MIL) -16.25000
 INITIAL LAKE VOLUME (m^3) IS 0.0000000E+00
 INFLOW RATE IS (m^3/yr) 3.7000000E+09
 ISOTOPIC COMP OF INFLOW WATER IS -16.25000

TIME (yrs)	VOLUME (m^3)	DEL O-18 H2O (SMOW)	O-18 CARB (SMOW)
0	0.0000000E+00	-16.2500000	20.59025
8	0.3052221E+11	-14.5784540	22.26180

***** OVERFLOW *****

0	0.3002000E+11	-14.5784540	22.26180
50	0.3002000E+11	-13.2548151	23.58543
100	0.3002000E+11	-13.2528658	23.58738
150	0.3002000E+11	-13.2528629	23.58739
200	0.3002000E+11	-13.2528629	23.58739
250	0.3002000E+11	-13.2528629	23.58739
300	0.3002000E+11	-13.2528629	23.58739
350	0.3002000E+11	-13.2528629	23.58739
400	0.3002000E+11	-13.2528629	23.58739
450	0.3002000E+11	-13.2528629	23.58739
500	0.3002000E+11	-13.2528629	23.58739

STEADY STATE VALUES-OVERFLOW

OVERFLOW TIME 8
 LAKE VOLUME 3.0020000E+10
 STEADY STATE DEL H2O -13.25286
 STEADY STATE CARBONATE 23.58739
 OUTFLOW RATE 3.1102428E+09

INFILL CHINA LAKE FROM DRY LAKE; DELI=DEL C

TEMPERATURE IN DEGREES CELCIUS IS 18.40000
 ELV = 9.672875
 ISOTOPIC COMP. OF BACK CONDENSATION FLUX IS -13.25286
 RELATIVE HUMIDITY IS 0.2320000
 ELV-STAR = 1.4882922E-02
 COMPUTED CARBONATE VALUES FOR DOLOMITE
 EVAPORATION RATE IS (m/yr) 1.413000
 ESTIMATED MAXIMUM LAKE SURFACE AREA IS (m^2) 1.5500000E+08
 ESTIMATED MAXIMUM LAKE VOLUME IS (m^3) 6.9600000E+08
 K= 0.3146767
 INITIAL LAKE ISO COMPOSITION IS (PER MIL) -13.25286
 INITIAL LAKE VOLUME (m^3) IS 0.0000000E+00
 INFLOW RATE IS (m^3/yr) 3.1100001E+09
 ISOTOPIC COMP OF INFLOW WATER IS -13.25286

TIME (yrs)	VOLUME (m^3)	DEL O-18 H2O (SMOW)	O-18 CARB (SMOW)
0	0.0000000E+00	-13.2528601	22.82910
0	0.2762729E+10	-11.1157265	24.96623

***** OVERFLOW *****

0	0.6960000E+09	-11.1157265	24.96623
50	0.6960000E+09	-12.2491808	23.83278
100	0.6960000E+09	-12.2491808	23.83278
150	0.6960000E+09	-12.2491808	23.83278
200	0.6960000E+09	-12.2491808	23.83278
250	0.6960000E+09	-12.2491808	23.83278
300	0.6960000E+09	-12.2491808	23.83278
350	0.6960000E+09	-12.2491808	23.83278
400	0.6960000E+09	-12.2491808	23.83278
450	0.6960000E+09	-12.2491808	23.83278
500	0.6960000E+09	-12.2491808	23.83278

STEADY STATE VALUES-OVERFLOW

OVERFLOW TIME 0
 LAKE VOLUME 6.9600000E+08
 STEADY STATE DEL H2O -12.24918
 STEADY STATE CARBONATE 23.83278
 OUTFLOW RATE 2.9417966E+09

INFILL SEARLES LAKE FROM DRY LAKE; DELI=DEL C

TEMPERATURE IN DEGREES CELCIUS IS 19.10000
 ELV = 9.613042
 ISOTOPIC COMP. OF BACK CONDENSATION FLUX IS -12.24918
 RELATIVE HUMIDITY IS 0.2210000
 ELV-STAR = 1.4898777E-02
 COMPUTED CARBONATE VALUES FOR DOLOMITE
 EVAPORATION RATE IS (m/yr) 1.685000
 ESTIMATED MAXIMUM LAKE SURFACE AREA IS (m^2) 9.9400000E+08
 ESTIMATED MAXIMUM LAKE VOLUME IS (m^3) 8.5279998E+10
 K= 1.9639893E-02
 INITIAL LAKE ISO COMPOSITION IS (PER MIL) -12.24918
 INITIAL LAKE VOLUME (m^3) IS 0.0000000E+00
 INFLOW RATE IS (m^3/yr) 2.9417999E+09
 ISOTOPIC COMP OF INFLOW WATER IS -12.24918

TIME (yrs)	VOLUME (m^3)	DEL 0-18 H2O (SMOW)	0-18 CARB (SMOW)
0	0.0000000E+00	-12.2491798	23.65311
10	0.2727811E+11	-10.8798704	25.02242
20	0.5068638E+11	-9.6160879	26.28621
30	0.7077383E+11	-8.4522257	27.45007
38	0.8640402E+11	-7.4856467	28.41665

***** OVERFLOW *****

0	0.8528000E+11	-7.4856467	28.41665
50	0.8528000E+11	-5.2432985	30.65900
100	0.8528000E+11	-4.9169765	30.98532
150	0.8528000E+11	-4.8694882	31.03281
200	0.8528000E+11	-4.8625770	31.03972
250	0.8528000E+11	-4.8615713	31.04072
300	0.8528000E+11	-4.8614249	31.04087
350	0.8528000E+11	-4.8614039	31.04089
400	0.8528000E+11	-4.8614006	31.04089
450	0.8528000E+11	-4.8614001	31.04089
500	0.8528000E+11	-4.8614001	31.04089

STEADY STATE VALUES-OVERFLOW

OVERFLOW TIME 38
 LAKE VOLUME 8.5279998E+10
 STEADY STATE DEL H2O -4.861401
 STEADY STATE CARBONATE 31.04089
 OUTFLOW RATE 1.6370606E+09

SEARLES OVERFLOW TO BENEATH SPILLWAY; DELI=DELC

TEMPERATURE IN DEGREES CELCIUS IS 19.10000
 ELV = 9.613042
 ISOTOPIC COMP. OF BACK CONDENSATION FLUX IS -12.24918
 RELATIVE HUMIDITY IS 0.2210000
 ELV-STAR = 1.4898777E-02
 COMPUTED CARBONATE VALUES FOR DOLOMITE
 EVAPORATION RATE IS (m/yr) 1.685000
 ESTIMATED MAXIMUM LAKE SURFACE AREA IS (m^2) 9.9400000E+08
 ESTIMATED MAXIMUM LAKE VOLUME IS (m^3) 8.5279998E+10
 K= 1.9639893E-02
 INITIAL LAKE ISO COMPOSITION IS (PER MIL) -4.861401
 INITIAL LAKE VOLUME (m^3) IS 8.5279998E+10
 INFLOW RATE IS (m^3/yr) 1.0000000E+09
 ISOTOPIC COMP OF INFLOW WATER IS -12.24918

TIME (yrs)	VOLUME (m^3)	DEL 0-18 H2O (SMOW)	0-18 CARB (SMOW)
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***** OVERFLOW *****

0	0.8528000E+11	-4.8614011	31.04089
10	0.8528000E+11	-3.3052216	32.59707

***** LAKE LEVEL BENEATH SPILLWAY *****

0	0.8528000E+11	-3.3052227	32.59707
100	0.6967496E+11	2.5928617	38.49516
200	0.6629573E+11	2.7469881	38.64928
300	0.6556398E+11	2.5650747	38.46737
400	0.6540551E+11	2.4936721	38.39597
500	0.6537120E+11	2.4736226	38.37592
600	0.6536377E+11	2.4686158	38.37091
700	0.6536216E+11	2.4674423	38.36974
800	0.6536181E+11	2.4671741	38.36947
900	0.6536174E+11	2.4671144	38.36941
1000	0.6536172E+11	2.4670997	38.36939
1100	0.6536172E+11	2.4670959	38.36939
1200	0.6536172E+11	2.4670959	38.36939
1300	0.6536172E+11	2.4670959	38.36939
1400	0.6536172E+11	2.4670959	38.36939
1490	0.6536172E+11	2.4670959	38.36939

STEADY STATE VALUES:NON-OVERFLOW

LAKE VOLUME 6.5361715E+10
 STEADY STATE DEL H2O 2.467098
 STEADY STATE CARBONATE 38.36939

SEARLES HIGH LAKE LEVEL TO LOW LEVEL; DELI=DELC

TEMPERATURE IN DEGREES CELCIUS IS 19.10000
ELV = 9.613042
ISOTOPIC COMP. OF BACK CONDENSATION FLUX IS -12.24918
RELATIVE HUMIDITY IS 0.2210000
ELV-STAR = 1.4898777E-02
COMPUTED CARBONATE VALUES FOR DOLOMITE
EVAPORATION RATE IS (m/yr) 1.685000
ESTIMATED MAXIMUM LAKE SURFACE AREA IS (m²) 9.9400000E+08
ESTIMATED MAXIMUM LAKE VOLUME IS (m³) 8.5279998E+10
K= 1.9639893E-02
INITIAL LAKE ISO COMPOSITION IS (PER MIL) 2.467098
INITIAL LAKE VOLUME (m³) IS 6.5361719E+10
INFLOW RATE IS (m³/yr) 3.0000000E+08
ISOTOPIC COMP OF INFLOW WATER IS -12.24918

TIME (yrs)	VOLUME (m ³)	DEL O-18 H2O (SMOW)	O-18 CARB (SMOW)
0	0.6536172E+11	2.4670959	38.36939
100	0.2951623E+11	8.6809998	44.58329
200	0.2175400E+11	5.5103006	41.41259
300	0.2007311E+11	3.3718011	39.27409
400	0.1970912E+11	2.6946478	38.59694
500	0.1963030E+11	2.5206320	38.42293
600	0.1961323E+11	2.4792888	38.38158
700	0.1960954E+11	2.4698229	38.37212
800	0.1960874E+11	2.4676993	38.37000
900	0.1960856E+11	2.4672225	38.36952
1000	0.1960852E+11	2.4671257	38.36942
1100	0.1960852E+11	2.4671032	38.36940
1200	0.1960851E+11	2.4670959	38.36939
1300	0.1960851E+11	2.4670959	38.36939
1400	0.1960851E+11	2.4670959	38.36939
1500	0.1960851E+11	2.4670959	38.36939
1600	0.1960851E+11	2.4670959	38.36939
1700	0.1960851E+11	2.4670959	38.36939
1800	0.1960851E+11	2.4670959	38.36939
1900	0.1960851E+11	2.4670959	38.36939
2000	0.1960851E+11	2.4670959	38.36939

STEADY STATE VALUES:NON-OVERFLOW

LAKE VOLUME 1.9608515E+10
STEADY STATE DEL H2O 2.467099
STEADY STATE CARBONATE 38.36939

DESSICATE SEARLES LAKE; DELI=DEL C

TEMPERATURE IN DEGREES CELCIUS IS 19.10000
ELV = 9.613042
ISOTOPIC COMP. OF BACK CONDENSATION FLUX IS -12.24918
RELATIVE HUMIDITY IS 0.2950000
ELV-STAR = 1.4394820E-02
COMPUTED CARBONATE VALUES FOR DOLOMITE
EVAPORATION RATE IS (m/yr) 1.685000
ESTIMATED MAXIMUM LAKE SURFACE AREA IS (m^2) 9.9400000E+08
ESTIMATED MAXIMUM LAKE VOLUME IS (m^3) 8.5279998E+10
K= 1.9639893E-02
INITIAL LAKE ISO COMPOSITION IS (PER MIL) 2.467098
INITIAL LAKE VOLUME (m^3) IS 1.9608515E+10

TIME (YRS)	VOLUME (M^3)	DEL O-18 H2O (SMOW)	O-18 CARB (SMOW)
0	0.1960851E+11	2.4670997	38.36939
50	0.9812506E+10	10.9500923	46.85239
100	0.4910380E+10	17.3886890	53.29099
150	0.2457255E+10	22.2755814	58.17788
200	0.1229661E+10	25.9847355	61.88703
250	0.6153478E+09	28.7999821	64.70228
300	0.3079328E+09	30.9367580	66.83905
350	0.1540960E+09	32.5585709	68.46087
400	0.7711277E+08	33.7895241	69.69182
450	0.3858883E+08	34.7238235	70.62612
500	0.1931064E+08	35.4329491	71.33524
550	0.9663446E+07	35.9711800	71.87347
600	0.4835787E+07	36.3796997	72.28200
650	0.2419927E+07	36.6897621	72.59206
700	0.1210982E+07	36.9250984	72.82739
750	0.6060003E+06	37.1037254	73.00602
800	0.3032550E+06	37.2392998	73.14159
850	0.1517550E+06	37.3421974	73.24449
900	0.7594140E+05	37.4202995	73.32259
950	0.3800263E+05	37.4795799	73.38187
1000	0.1901730E+05	37.5245743	73.42687

INFILL OWENS FROM DRY LAKE; DELI > DELC

TEMPERATURE IN DEGREES CELCIUS IS 15.50000
 ELV = 9.926092
 ISOTOPIC COMP. OF BACK CONDENSATION FLUX IS -21.68460
 RELATIVE HUMIDITY IS 0.2780000
 ELV-STAR = 1.4819145E-02
 COMPUTED CARBONATE VALUES FOR DOLOMITE
 EVAPORATION RATE IS (m/yr) 1.177000
 ESTIMATED MAXIMUM LAKE SURFACE AREA IS (m^2) 6.9400000E+08
 ESTIMATED MAXIMUM LAKE VOLUME IS (m^3) 3.0020000E+10
 K= 2.7209796E-02
 INITIAL LAKE ISO COMPOSITION IS (PER MIL) -16.25000
 INITIAL LAKE VOLUME (m^3) IS 0.0000000E+00
 INFLOW RATE IS (m^3/yr) 3.7000000E+09
 ISOTOPIC COMP OF INFLOW WATER IS -16.25000

TIME (yrs)	VOLUME (m^3)	DEL O-18 H2O (SMOW)	O-18 CARB (SMOW)
0	0.0000000E+00	-16.2500000	20.59025
8	0.3052221E+11	-14.7542467	22.08600

***** OVERFLOW *****

0	0.3002000E+11	-14.7542467	22.08600
50	0.3002000E+11	-13.5698185	23.27043
100	0.3002000E+11	-13.5680752	23.27217
150	0.3002000E+11	-13.5680723	23.27218
200	0.3002000E+11	-13.5680723	23.27218
250	0.3002000E+11	-13.5680723	23.27218
300	0.3002000E+11	-13.5680723	23.27218
350	0.3002000E+11	-13.5680723	23.27218
400	0.3002000E+11	-13.5680723	23.27218
450	0.3002000E+11	-13.5680723	23.27218
500	0.3002000E+11	-13.5680723	23.27218

STEADY STATE VALUES-OVERFLOW

OVERFLOW TIME 8
 LAKE VOLUME 3.0020000E+10
 STEADY STATE DEL H2O -13.56807
 STEADY STATE CARBONATE 23.27218
 OUTFLOW RATE 3.1102428E+09

INFILL CHINA LAKE FROM DRY LAKE; DELI > DELC

TEMPERATURE IN DEGREES CELCIUS IS 18.40000
 ELV = 9.672875
 ISOTOPIC COMP. OF BACK CONDENSATION FLUX IS -13.75154
 RELATIVE HUMIDITY IS 0.2320000
 ELV-STAR = 1.4882922E-02
 COMPUTED CARBONATE VALUES FOR DOLOMITE
 EVAPORATION RATE IS (m/yr) 1.413000
 ESTIMATED MAXIMUM LAKE SURFACE AREA IS(m^2) 1.5500000E+08
 ESTIMATED MAXIMUM LAKE VOLUME IS (m^3) 6.9600000E+08
 K= 0.3146767
 INITIAL LAKE ISO COMPOSITION IS(PER MIL) -13.56807
 INITIAL LAKE VOLUME (m^3) IS 0.0000000E+00
 INFLOW RATE IS (m^3/yr) 3.1100001E+09
 ISOTOPIC COMP OF INFLOW WATER IS -13.56807

TIME (yrs)	VOLUME (m^3)	DEL O-18 H2O (SMOW)	O-18 CARB (SMOW)
0	0.0000000E+00	-13.5680704	22.51389
0	0.2762729E+10	-11.4379120	24.64405

***** OVERFLOW *****

0	0.6960000E+09	-11.4379120	24.64405
50	0.6960000E+09	-12.5676632	23.51430
100	0.6960000E+09	-12.5676632	23.51430
150	0.6960000E+09	-12.5676632	23.51430
200	0.6960000E+09	-12.5676632	23.51430
250	0.6960000E+09	-12.5676632	23.51430
300	0.6960000E+09	-12.5676632	23.51430
350	0.6960000E+09	-12.5676632	23.51430
400	0.6960000E+09	-12.5676632	23.51430
450	0.6960000E+09	-12.5676632	23.51430
500	0.6960000E+09	-12.5676632	23.51430

STEADY STATE VALUES-OVERFLOW

OVERFLOW TIME 0
 LAKE VOLUME 6.9600000E+08
 STEADY STATE DEL H2O -12.56766
 STEADY STATE CARBONATE 23.51430
 OUTFLOW RATE 2.9417966E+09

INFILL SEARLES LAKE FROM DRY LAKE; DELI > DELC

TEMPERATURE IN DEGREES CELCIUS IS 19.10000
ELV = 9.613042
ISOTOPIC COMP. OF BACK CONDENSATION FLUX IS -13.80998
RELATIVE HUMIDITY IS 0.2210000
ELV-STAR = 1.4898777E-02
COMPUTED CARBONATE VALUES FOR DOLOMITE
EVAPORATION RATE IS (m/yr) 1.685000
ESTIMATED MAXIMUM LAKE SURFACE AREA IS (m^2) 9.9400000E+08
ESTIMATED MAXIMUM LAKE VOLUME IS (m^3) 8.5279998E+10
K= 1.9639893E-02
INITIAL LAKE ISO COMPOSITION IS (PER MIL) -12.56766
INITIAL LAKE VOLUME (m^3) IS 0.0000000E+00
INFLOW RATE IS (m^3/yr) 2.9417966E+09
ISOTOPIC COMP OF INFLOW WATER IS -12.56766

TIME (yrs)	VOLUME (m^3)	DEL 0-18 H2O (SMOW)	0-18 CARB (SMOW)
0	0.0000000E+00	-12.5676603	23.33463
10	0.2727808E+11	-11.2246838	24.67761
20	0.5068633E+11	-9.9852495	25.91705
30	0.7077375E+11	-8.8438129	27.05848
38	0.8640392E+11	-7.8958535	28.00644

***** OVERFLOW *****

0	0.8528000E+11	-7.8958535	28.00644
50	0.8528000E+11	-5.6966844	30.20561
100	0.8528000E+11	-5.3766456	30.52565
150	0.8528000E+11	-5.3300710	30.57222
200	0.8528000E+11	-5.3232932	30.57900
250	0.8528000E+11	-5.3223071	30.57999
300	0.8528000E+11	-5.3221631	30.58013
350	0.8528000E+11	-5.3221426	30.58015
400	0.8528000E+11	-5.3221393	30.58015
450	0.8528000E+11	-5.3221388	30.58016
500	0.8528000E+11	-5.3221388	30.58016

STEADY STATE VALUES-OVERFLOW

OVERFLOW TIME 38
LAKE VOLUME 8.5279998E+10
STEADY STATE DEL H2O -5.322138
STEADY STATE CARBONATE 30.58016
OUTFLOW RATE 1.6370573E+09

SEARLES OVERFLOW TO BENEATH SPILLWAY; DELI > DELC

TEMPERATURE IN DEGREES CELCIUS IS 19.10000
 ELV = 9.613042
 ISOTOPIC COMP. OF BACK CONDENSATION FLUX IS -13.80998
 RELATIVE HUMIDITY IS 0.2210000
 ELV-STAR = 1.4898777E-02
 COMPUTED CARBONATE VALUES FOR DOLOMITE
 EVAPORATION RATE IS (m/yr) 1.685000
 ESTIMATED MAXIMUM LAKE SURFACE AREA IS (m^2) 9.9400000E+08
 ESTIMATED MAXIMUM LAKE VOLUME IS (m^3) 8.5279998E+10
 K= 1.9639893E-02
 INITIAL LAKE ISO COMPOSITION IS (PER MIL) -5.322138
 INITIAL LAKE VOLUME (m^3) IS 8.5279998E+10
 INFLOW RATE IS (m^3/yr) 1.0000000E+09
 ISOTOPIC COMP OF INFLOW WATER IS -12.56766

TIME (yrs)	VOLUME (m^3)	DEL O-18 H2O (SMOW)	O-18 CARB (SMOW)
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***** OVERFLOW *****

0	0.8528000E+11	-5.3221383	30.58016
10	0.8528000E+11	-3.7959268	32.10637

***** LAKE LEVEL BENEATH SPILLWAY *****

0	0.8528000E+11	-3.7959256	32.10637
100	0.6967496E+11	1.9885786	37.89087
200	0.6629573E+11	2.1397359	38.04203
300	0.6556398E+11	1.9613206	37.86361
400	0.6540551E+11	1.8913001	37.79359
500	0.6537120E+11	1.8716305	37.77393
600	0.6536377E+11	1.8667244	37.76902
700	0.6536216E+11	1.8655733	37.76787
800	0.6536181E+11	1.8653088	37.76760
900	0.6536174E+11	1.8652492	37.76754
1000	0.6536172E+11	1.8652343	37.76753
1100	0.6536172E+11	1.8652306	37.76752
1200	0.6536172E+11	1.8652306	37.76752
1300	0.6536172E+11	1.8652306	37.76752
1400	0.6536172E+11	1.8652306	37.76752
1490	0.6536172E+11	1.8652306	37.76752

STEADY STATE VALUES:NON-OVERFLOW

LAKE VOLUME 6.5361715E+10
 STEADY STATE DEL H2O 1.865230
 STEADY STATE CARBONATE 37.76752

SEARLES HIGH LAKE LEVEL TO LOW LEVEL; DELI > DELC

TEMPERATURE IN DEGREES CELCIUS IS 19.10000
 ELV = 9.613042
 ISOTOPIC COMP. OF BACK CONDENSATION FLUX IS -13.80998
 RELATIVE HUMIDITY IS 0.2210000
 ELV-STAR = 1.4898777E-02
 COMPUTED CARBONATE VALUES FOR DOLOMITE
 EVAPORATION RATE IS (m/yr) 1.685000
 ESTIMATED MAXIMUM LAKE SURFACE AREA IS (m^2) 9.9400000E+08
 ESTIMATED MAXIMUM LAKE VOLUME IS (m^3) 8.5279998E+10
 K= 1.9639893E-02
 INITIAL LAKE ISO COMPOSITION IS (PER MIL) 1.865231
 INITIAL LAKE VOLUME (m^3) IS 6.5361715E+10
 INFLOW RATE IS (m^3/yr) 3.0000000E+08
 ISOTOPIC COMP OF INFLOW WATER IS -12.56766

TIME (yrs)	VOLUME (m^3)	DEL O-18 H2O (SMOW)	O-18 CARB (SMOW)
0	0.6536172E+11	1.8652267	37.76752
100	0.2951623E+11	7.9594703	43.86176
200	0.2175400E+11	4.8498325	40.75213
300	0.2007311E+11	2.7525201	38.65482
400	0.1970912E+11	2.0884016	37.99070
500	0.1963030E+11	1.9177384	37.82003
600	0.1961323E+11	1.8771924	37.77949
700	0.1960954E+11	1.8679053	37.77020
800	0.1960874E+11	1.8658266	37.76812
900	0.1960856E+11	1.8653609	37.76765
1000	0.1960852E+11	1.8652604	37.76756
1100	0.1960852E+11	1.8652380	37.76753
1200	0.1960851E+11	1.8652343	37.76753
1300	0.1960851E+11	1.8652343	37.76753
1400	0.1960851E+11	1.8652343	37.76753
1500	0.1960851E+11	1.8652343	37.76753
1600	0.1960851E+11	1.8652343	37.76753
1700	0.1960851E+11	1.8652343	37.76753
1800	0.1960851E+11	1.8652343	37.76753
1900	0.1960851E+11	1.8652343	37.76753
2000	0.1960851E+11	1.8652343	37.76753

STEADY STATE VALUES:NON-OVERFLOW

LAKE VOLUME 1.9608515E+10
 STEADY STATE DEL H2O 1.865231
 STEADY STATE CARBONATE 37.76752

DESSICATE SEARLES LAKE; DELI > DELC

TEMPERATURE IN DEGREES CELCIUS IS 19.10000
ELV = 9.613042
ISOTOPIC COMP. OF BACK CONDENSATION FLUX IS -13.80998
RELATIVE HUMIDITY IS 0.2950000
ELV-STAR = 1.4394820E-02
COMPUTED CARBONATE VALUES FOR DOLOMITE
EVAPORATION RATE IS (m/yr) 1.685000
ESTIMATED MAXIMUM LAKE SURFACE AREA IS (m²) 9.9400000E+08
ESTIMATED MAXIMUM LAKE VOLUME IS (m³) 8.5279998E+10
K= 1.9639893E-02
INITIAL LAKE ISO COMPOSITION IS (PER MIL) 1.865231
INITIAL LAKE VOLUME (m³) IS 1.9600001E+10

TIME (YRS)	VOLUME (M ³)	DEL O-18 H2O (SMOW)	O-18 CARB (SMOW)
0	0.1960000E+11	1.8652306	37.76752
50	0.9808245E+10	10.0981112	46.00040
100	0.4908248E+10	16.3468742	52.24917
150	0.2456189E+10	21.0896816	56.99198
200	0.1229128E+10	24.6894760	60.59177
250	0.6150807E+09	27.4217167	63.32401
300	0.3077991E+09	29.4954910	65.39779
350	0.1540291E+09	31.0694866	66.97178
400	0.7707929E+08	32.2641487	68.16644
450	0.3857208E+08	33.1708984	69.07320
500	0.1930226E+08	33.8591194	69.76141
550	0.9659251E+07	34.3814774	70.28378
600	0.4833687E+07	34.7779503	70.68024
650	0.2418876E+07	35.0788765	70.98117
700	0.1210457E+07	35.3072739	71.20957
750	0.6057372E+06	35.4806290	71.38293
800	0.3031233E+06	35.6122055	71.51450
850	0.1516891E+06	35.7120743	71.61437
900	0.7590842E+05	35.7878723	71.69017
950	0.3798613E+05	35.8454056	71.74770
1000	0.1900904E+05	35.8890686	71.79137

Appendix I

Climatological Data for Southeastern California
Weather Stations

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* This file contains climatological data for SE California weather
* stations east of the Sierra Nevada divide.
*
* Data from White Mountain stations are from info compiled by UC-Davis
* for the BARCROFT(white Mountain2) 1953-1973 and CROOKED CREEK
* (white mountain1) 1949-1973 reseach stations.
*
* Relative humidity data for Bishop is from:
* Ruffner,J.A. (ed.),1985,CLimates of the States:Gale Research Co.
* Detroit, Michigan. Vol. 1
* (compilation from NOAA climatological data QE 983 .C56 1985 v.1)
*
* All other elevation,temp and precip data are from "Monthly normals of
* Temperature, Precipitation, and Heating and Cooling Degree Days
* 1951-1980 no.81 (California):NOAA,National Climatic Center, Asheville,
* North Carolina,1982."
*
* Evaporation data (following Smith,G.I. and Street-Perrott,F.A.) are
* interpolations between "evaporation from lake surfaces-Plate 3"
* IN Meyers,J.S.,1962, Evaporation from the 17 Western States, USGS
* Professional Paper 272-D: USGS etc....
*
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STATION NAME /LOCATION	ELEVATION (METERS)	TEMP DEG C 30*YR MEAN	ANNUAL PRECIP 30yr MEAN (cm)	REL HUMIDITY 30yr av annual
DEATH VALLEY	-59.69	24.83	5.16	----
TRONA	521.54	19.0	10.03	----
INYOKERN	750.77	17.39	10.57	----
INDEPENDENCE	1215.39	-----	13.69	----
BISHOP	1264.0	13.33	----	0.29
MONO LAKE	1984.62	8.72	----	----
WHITE MOUNTAIN1	3123.08	1.28	34.70	0.529
WHITE MOUNTAIN2	3836.92	-2.44	49.61	0.558

LOCATION	ELEVATION (METERS)	EVAPORATION FROM LAKE SURFACES ANNUAL GROSS (m/yr)
LAKE MANLY	-86	2.13
LAKE PANAMINT	317	1.93
SEARLES LAKE	493	1.78
CHINA LAKE	657	1.52
OWENS LAKE	1081	1.33
HAIWEE RESEV	1177	1.025
INDEPENDENCE	1215.39	1.025
BISHOP	1264.0	0.77

ELEVATION vs. 30-yr MEAN ANNUAL TEMPERATURE
OWENS VALLEY AREA WEATHER STATIONS

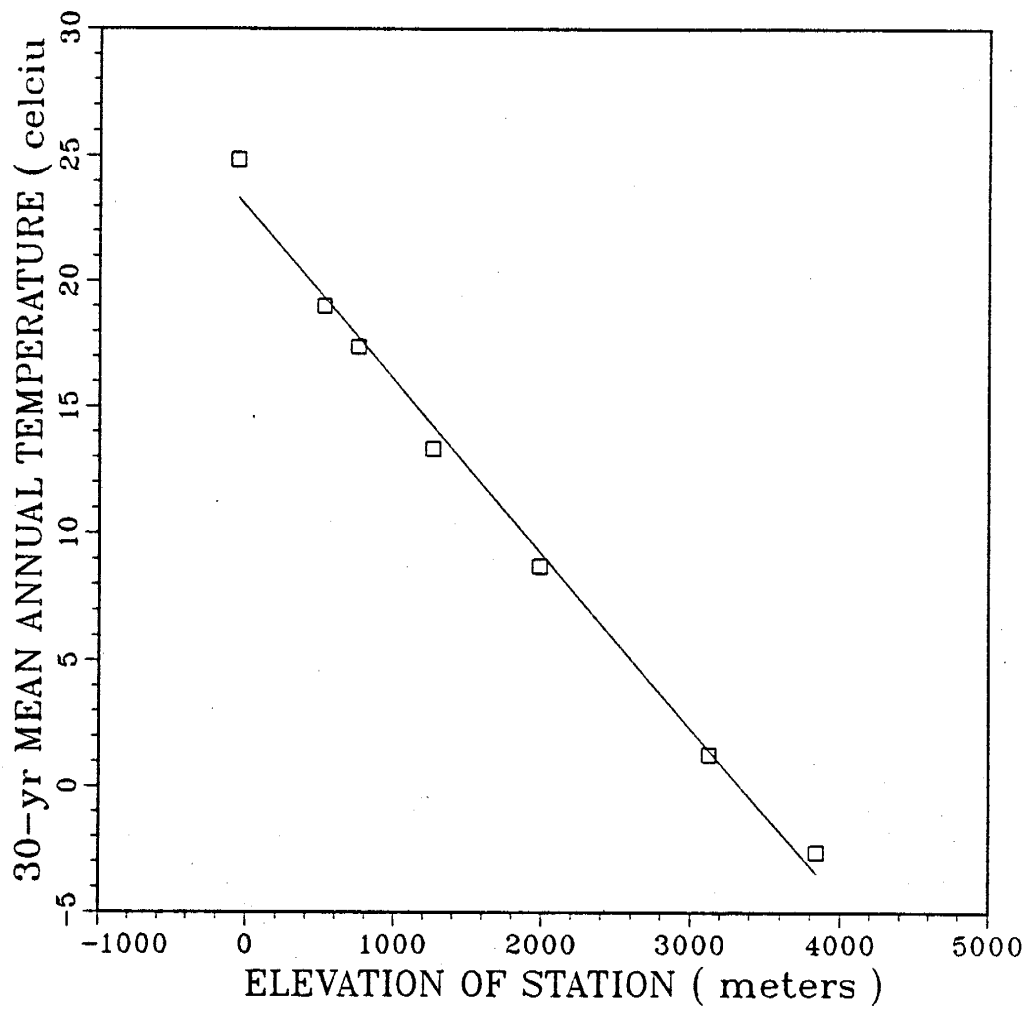


Figure 35. Linear regression of temperature as a function of elevation for Owens Valley area weather stations.

FOR YOUR CHOICE: "LINE" $Y=A+B*X$
"A"= 2.2934E+01
"B" = -6.8820E-03

THE REGRESSION COEFFICIENT "R" = 0.99251

THE CALCULATED VALUES:

X	Y	YCALC	% DIFF
-5.9692E+01	2.4830E+01	2.3345E+01	5.98
5.2154E+02	1.9000E+01	1.9345E+01	-1.82
7.5077E+02	1.7390E+01	1.7768E+01	-2.17
1.2640E+03	1.3330E+01	1.4236E+01	-6.79
1.9846E+03	8.7200E+00	9.2763E+00	-6.38
3.1231E+03	1.2800E+00	1.4414E+00	-12.61
3.8369E+03	-2.6100E+00	-3.4713E+00	-33.00

THE NONLINEAR CORRELATION COEFFICIENT: $R^{**2}= 0.99251443$
the ave. % diff= 9.82137108

ELEVATION vs. 30-yr MEAN ANNUAL PRECIPITATION
OWENS VALLEY AREA WEATHER STATIONS

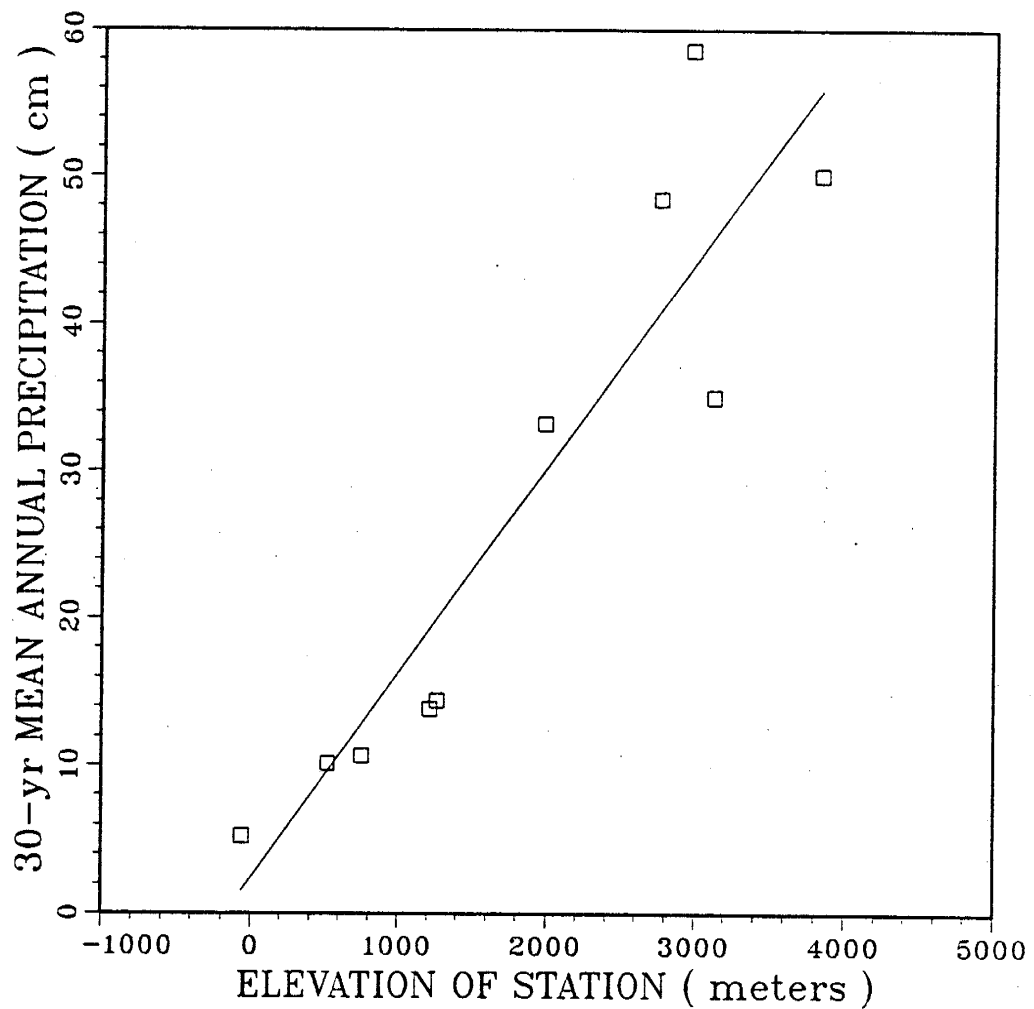


Figure 36. Linear regression of precipitation as a function of elevation for Owens Valley area weather stations.

FOR YOUR CHOICE: "LINE" $Y=A+B*X$
 "A"= 2.3442E+00
 "B" = 1.3947E-02

THE REGRESSION COEFFICIENT "R" = 0.84895

THE CALCULATED VALUES:

X	Y	YCALC	% DIFF
-5.9692E+01	5.2050E+00	1.5117E+00	70.96
5.2154E+02	1.0128E+01	9.6180E+00	5.04
7.5077E+02	1.0667E+01	1.2815E+01	-20.14
1.2154E+03	1.3821E+01	1.9295E+01	-39.61
1.2640E+03	1.4385E+01	1.9973E+01	-38.85
1.9846E+03	3.3231E+01	3.0023E+01	9.65
2.7600E+03	4.8436E+01	4.0837E+01	15.69
2.9677E+03	5.8590E+01	4.3734E+01	25.36
3.1231E+03	3.5026E+01	4.5901E+01	-31.05
3.8369E+03	5.0077E+01	5.5857E+01	-11.54

THE NONLINEAR CORRELATION COEFFICIENT: R^{**2} = 0.84895211
 the ave. % diff= 26.78702736

ELEVATION vs. ANNUAL EVAPORATION RATES
EVAPORATION OVER LAKE SURFACES
OWENS VALLEY AREA WEATHER STATIONS

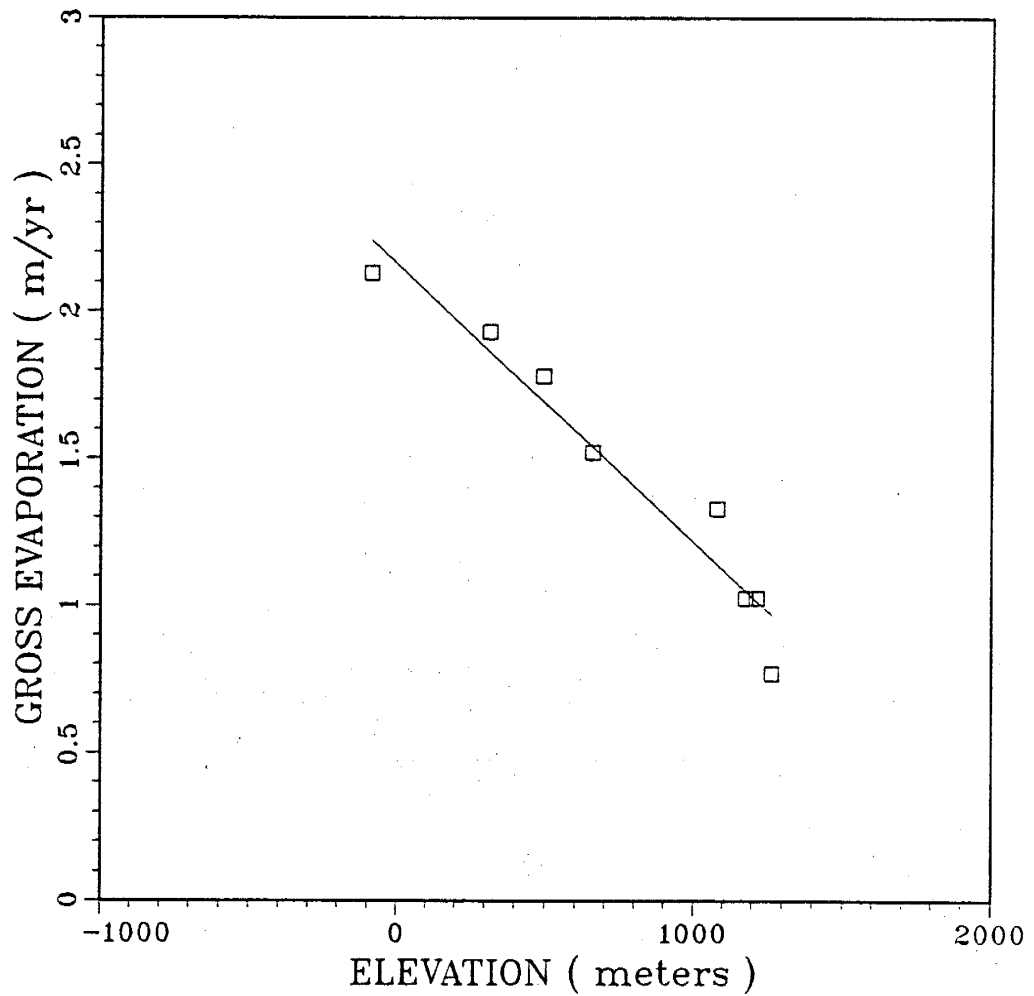


Figure 37. Linear regression of gross evaporation as a function of elevation for Owens Valley area weather stations.

FOR YOUR CHOICE: "LINE" $Y=A+B*X$
 "A"= 2.1607E+00
 "B" = -9.4398E-04

THE REGRESSION COEFFICIENT "R" = 0.93882

THE CALCULATED VALUES:

X	Y	YCALC	% DIFF
-8.6000E+01	2.1300E+00	2.2418E+00	-5.25
3.1700E+02	1.9300E+00	1.8614E+00	3.55
4.9300E+02	1.7800E+00	1.6953E+00	4.76
6.5700E+02	1.5200E+00	1.5405E+00	-1.35
1.0810E+03	1.3300E+00	1.1402E+00	14.27
1.1770E+03	1.0250E+00	1.0496E+00	-2.40
1.2150E+03	1.0250E+00	1.0137E+00	1.10
1.2640E+03	7.7000E-01	9.6747E-01	-25.64

THE NONLINEAR CORRELATION COEFFICIENT: $R^{**2}= 0.93882328$
 the ave. % diff= 7.29056263

ELEVATION vs. 30-yr MEAN ANNUAL RELATIVE HUMIDITY
OWENS VALLEY AREA WEATHER STATIONS

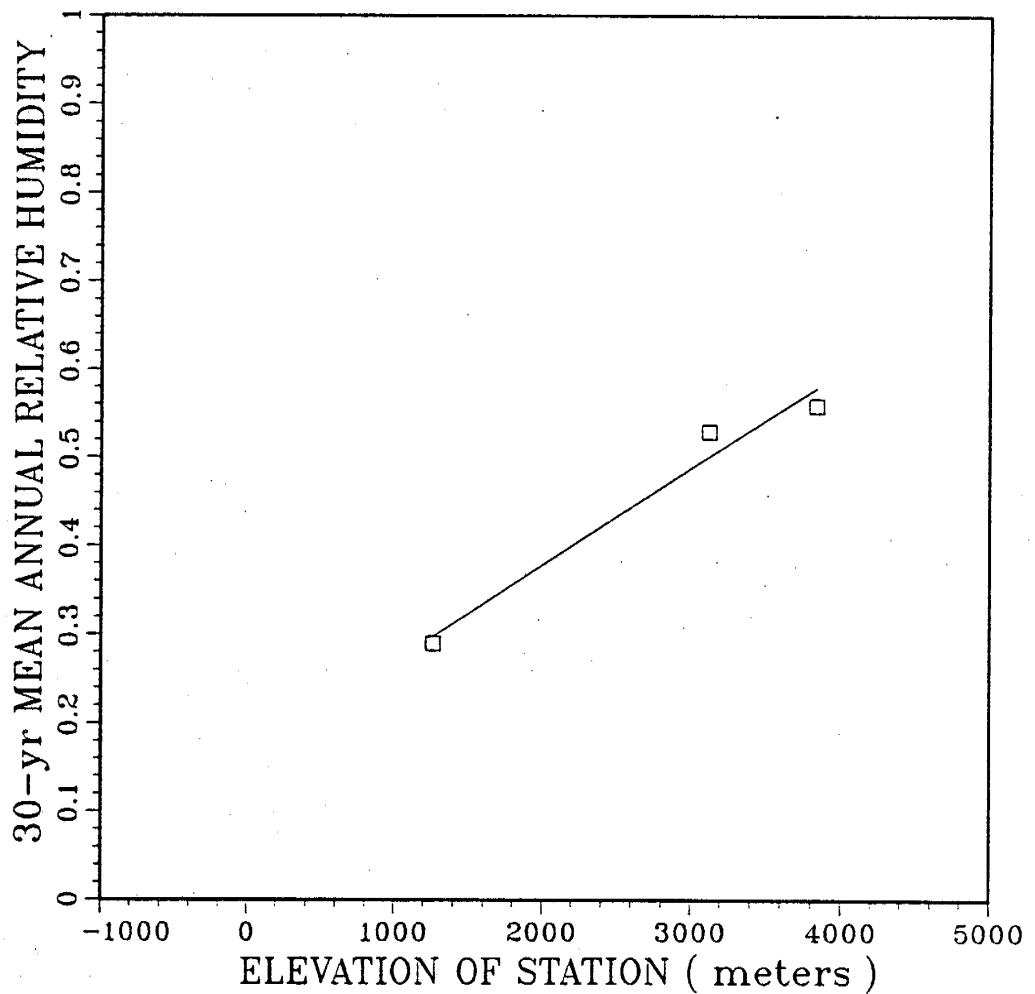


Figure 38. Linear regression of relative humidity as a function of elevation for Owens Valley area weather stations.

FOR YOUR CHOICE: "LINE" $Y=A+B*X$

"A"= 1.6001E-01

"B" = 1.0907E-04

THE REGRESSION COEFFICIENT "R" = 0.97026

THE CALCULATED VALUES:

X	Y	YCALC	% DIFF
1.2640E+03	2.9000E-01	2.9787E-01	-2.71
3.1231E+03	5.2900E-01	5.0064E-01	5.36
3.8369E+03	5.5800E-01	5.7849E-01	-3.67

THE NONLINEAR CORRELATION COEFFICIENT: $R^{**2}= 0.97026372$

the ave. % diff= 3.91607213

Appendix J

Bathymetric Data for Lakes in the Paleo-Owens River System

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* This file contains routing data for the Owens River Lake System
* Data for Owens and Searles are from Smith( 1979 )
* Data for China Lake from Jannik (1989)
* Data for Panamint is from Smith, R.S.U. ( )
* Data from Death Valley is from Jannik (1989)
*
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CUMULATIVE DEPTH (m)	CUMULATIVE AREA (x10 ⁹ m ²)	LAKE AREA (x10 ⁹ m ²)	ADDED VOLUME m ³)	CUMULATIVE LAKE VOL. (x10 ⁹ m ³)	K AREA/VOLUME
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0.0	0.0	0.0	0.0	0.0	0
2.4	0.20	0.20	0.16	0.16	1.25
14.0	0.29	0.29	2.99	3.15	0.092
64.0	0.694	0.694	26.87	30.02	0.023

***** OWENS OVERFLOW *****

68.0	0.721	0.027	0.036	0.036	0.75
72.0	0.849	0.155	0.660	0.696	.223

***** CHINA OVERFLOW *****

97.0	1.094	0.245	2.04	2.04	.120
129.0	1.149	0.300	8.71	10.75	.028
160.0	1.199	0.350	10.07	20.82	.017
192.0	1.291	0.442	12.64	33.46	.013
219.0	1.382	0.533	13.14	46.60	.011

250.0	1.564	0.715	19.27	65.87	.011
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***** China coalesces with Searles *****

272.0	1.688	0.994	18.71	85.28	.012
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***** SEARLES OVERFLOW *****

290.0	1.806	0.118	0.71	0.71	.166
310.0	1.863	0.175	2.91	3.62	.048
321.0	1.877	0.189	2.00	5.62	.034
351.0	1.930	0.242	6.45	12.07	.020
382.0	1.977	0.289	8.22	20.29	.014
412.0	2.017	0.329	9.26	29.55	.011
443.0	2.057	0.369	10.81	40.36	.009
478.0	2.116	0.428	13.93	54.29	.008
498.0	2.176	0.488	9.15	63.44	.008
513.0	2.212	0.524	7.59	71.03	.007
533.0	2.256	0.568	10.92	81.95	.007
559.0	2.326	0.638	15.67	97.62	.007
574.0	2.415	0.727	10.23	107.85	.007

***** PANAMINT OVERFLOW *****

613.0	2.465	0.050	0.65	0.65	
786.0	2.998	0.583	46.35	47.0	

