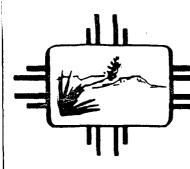
GW - 23

MONITORING REPORTS

DATE:

1989 - 1988





GARREY CARRUTHERS
Governor

DENNIS BOYD Secretary

MICHAEL J. BURKHART Deputy Secretary

RICHARD MITZELFELT
Director

October 25, 1989

Mr. Dave Boyer Oil Conservation Division State Land Office Bldg. P.O. Box 2088 Santa Fe, NM 87504

Dear Mr. Boyer:

Enclosed for your information are copies of EPA's Comprehensive Groundwater Monitoring Reports for the four Phillips Gas Plants - Artisia, Eunice, Lee and Lusk. These reports have not been reviewed by the Hazardous Waste Program and are to be considered as draft reports. At this time no further action is expected on the reports to finalize them.

If you have any questions, or need additional information please call me at 827-0170.

Sincerely,

1. Surgane More Mayor Suzanne Moore-Mayne

Water Resource Specialist II

Hazardous Waste Program

SMM/vga

Encl.



ODESSA, TEXAS 79762 4001 PENBROOK

April 5, 1989

Groundwater Monitoring Analyses
Artesia, Eunice, Lee and Lusk Plants

Mr. Dave Boyer Environmental Bureau Chief New Mexico Oil Conservation Division P. O. Box 2088 Santa Fe, New Mexico 87501

Dear Mr. Boyer:

Per your request, attached please find coppes of the fourth quarter groundwater monitoring analyses for the above referenced plants.

If you should have any questions regarding this information, please contact me at (915) 367-1316.

Very truly yours,

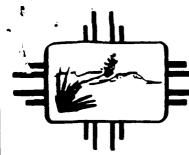
Michael D. Ford

Environmental Analyst

MDF

Attachments

See "Amalyses" Son copies New Mexico Health and Environment; Departmention



RECEIVED

°ES OCT 30 AM 11 12 Dennis Boyd

Secretary

MICHAEL J. BURKHART Deputy Secretary

RICHARD MITZELFELT

September 13, 1989

William F. Ballard, Manager Phillips Petroleum Company 12 A4 Phillips Bldg. Bartlesville, OK 74004

RE: RCRA status Artesia, Eunice, Lee and Lusk Plants-NMD000709667, NMD000709634, NMD000709675, NMD000709659

Dear Mr. Ballard:

The New Mexico Environmental Improvement Division (NMEID), accepts Phillips Petroleum Company's (Phillips') position presented in their May 17, 1989 correspondence that the four Phillips facilities in New Mexico, Artesia, Eunice, Lee and Lusk are exempt from RCRA regulation based upon EPA's Regulatory Determination of July 6, 1988 Federal Register. NMEID also accepts Phillips' Certificate of No Hazardous Waste Activity included in the May 17, 1989 correspondence.

NMEID's acceptance of Phillips' position does not remove Phillips from regulation under the Hazardous Waste Management Regulations, (HWMR-5, as amended 1989) and the New Mexico Hazardous Waste Act, New Mexico Statutes Annotated 1978, (1989, Supp.), if Phillips transports, treats, stores or disposes of hazardous wastes in the future. To the extent that Phillips generates hazardous wastes, Phillips is subject to the generator requirements of HWMR-5.

If NMEID receives any new information that indicates that Phillips has been or may be regulated under RCRA, enforcement actions will be initiated. With NMEID's acceptance of Phillips' position, compliance with the April 19, 1988 Compliance Order/Schedule is determined to be resolved. However, Phillips may still be subject to EPA enforcement actions.

Mr. Ballard September 13, 1989 Page 2

A copy of EPA's response to NMETD's request to provide an interpretation of the oil and gas exemption in the July 6, 1988 Federal Register is enclosed for Phillips' information.

If you have any questions or need additional information, please call me at (505) 827-2926.

Sincerely,

Boyd Hamilton

Program Manager Hazardous Waste Program

BH/SMM/smm

Encl.

cc: Lynn Prince, EPA Region 6
Tracy Hughes, Office of General Counsel, EID
Knut Am, Phillips Petroleum Company
Reese B. Copeland, Phillips Petroleum Company



UNITED STATES ENVIRONMENTAL PROTECTION AGENCY

REGION (

1445 ROSS AVENUE, SUITE 1200 DALLAS, TEXAS 75202

July, 18, 1989

Mr. Boyd Hamilton
Program Manager
Hazardous Waste Program
New Mexico Health and Environment Department
Harold Runnels Building
1190 St. Francis Drive
Santa Fe, New Mexico 87503

Dear Mr. Hamilton:

On June 8, 1989, you requested that the Environmental Protection Agency (EPA) provide an interpretation of the so called oil and gas exemption to the Resource Conservation and Recovery Act (RCRA) as delineated in the Regulatory Determination in the July 6, 1988, Federal Register (FR). Specifically, you asked if the exemption applied to four gas plants operated by Phillips Petroleum Company (Phillips) in eastern New Mexico. This request was prompted by Phillips' assertion, in a letter dated May 17, 1989, that the surface impoundments in question are not RCRA regulated units based on that regulatory determination. Phillips supported this position with a certificate of no hazardous waste activity for the four plants.

In EPA's regulatory determination, on Page 25454, cooling tower blowdown is specifically included in the wastes exempted from RCRA regulation. However, gas plant cooling tower cleaning wastes are specifically excluded from the exemption. These determinations are based on the three criteria included as an attachment to the June 6, 1989, letter from Dan Derkics, (Chief, Large Volume Waste Section EPA Headquarters) to Julie Wanslow, a copy of which was included in your letter to me of June 15, 1989. Mr. Derkics letter states that cooling tower blowdown "... is comprised only of water, scale or other wastes generated by the actual operation of the cooling tower." The Region interprets this to mean that corrosion inhibitors and biological control agents are included in cooling tower blowdown.

Mr. Derkics also clarifies the meaning of cooling tower cleaning wastes as those wastes which, may be generated by <u>any</u> cooling tower and includes "...solvents, scrubbing agents or <u>other cleaning materials</u> introduced

into the process solely to remove buildup or otherwise clean the equipment, and are not included as part of the functional operation of the cooling tower." Such wastes are not intrinsically derived from primary field operations for natural gas production. The Region interprets this to mean that the wastes generated during the periodic cleaning are not exempt.

In their No Hazardous Waste Activity Certificate, Phillips states that both chromate and non-chromate chemicals have been used in the cooling towers since November 19, 1980, as corrosion inhibitors at these sites. They further state that cooling towers must be cleaned on a periodic basis (approximately once every five years) and that this cleaning consists of removing the sludge by vacuum truck from the basin and removing scale from the cooling coil heads and laterals by sandblasting. Phillips also asserts that these materials have been tested and are not hazardous wastes.

One of the reasons that cleaning waste from a cooling tower may be RCRA hazardous waste is due to the chemicals added to the system for corrosion inhibition or control of biological agents. Chromate compounds have been widely used in this application as they have at the Phillips gas plants. Discarded materials generated in the cooling tower would be hazardous waste, as that term is defined in 40 CFR $\S261.3$, when the chromium concentration reaches 5.0~mg/l when tested using the procedures for EP toxicity.

If the waste generated during the periodic cleaning exceeds a concentration of 5.0 mg/l for chromium, then the waste is hazardous waste. Phillips claims the waste is tested in their certificate but they do not provide enough information for a determination of the adequacy of the testing. Should this waste be EP Toxic and should it be placed in the same surface impoundments as the cooling tower blowdown, then the units are RCRA regulated regardless of the exemption for cooling tower blowdown. If on the other hand these conditions are not met, then the material is not hazardous waste. At the very least, the coil heads and laterals have the potential of naving significant levels of chromium waste/scale which must be sandblasted off. It is this cooling tower cleaning waste that may make the units regulated, however, such a determination is not possible from the information provided in the certificate.

Some discussion is necessary about a mixture of an exempted waste and a non-exempted waste. EPA has in the past exempted some such mixtures as in the case of ash waste and flue gas emission control waste generated primarily from the combustion of coal and fossil fuels. [40 CFR 261.4(b)(4)] However, the wastes which are co-disposed and also exempt are those materials generated in conjunction with the exempted wastes. The waste materials are not segregated from the combustion wastes. Wastes which

are segregated and dissed of or treated separately fun combustion wastes and otherwise meet the definition of a hazardous waste are regulated under RCRA. This determination was made in 1981 in response to the Utility Solid Waste Activities Group.

The clearest exposition of EPA's stand regarding the applicability of the mixture rule when an exempted waste is mixed with a hazardous waste is found in the proposed rule published in the Federal Register on April 17, 1989, for mining waste.

"EPA has decided, however, that it is appropriate to revise the proposed regulatory status of some mixtures of non-excluded 'characteristic' wastes and Bevill wastes. In these instances, the mixture will be considered a hazardous waste if it exhibits one or more of the same hazardous characteristics that are exhibited by the non-excluded waste. If the mixture exhibits one or more hazardous characteristics that are exhibited by the Bevill waste but not by the non-excluded characteristic waste, then the mixture is not hazardous waste.

EPA wishes to make clear, however that in any case, mixing a characteristic hazardous waste with a Bevill waste would require a RCRA treatment, storage or disposal permit....

Although this interpretation applies to a <u>proposed</u> mining waste rule, EPA's Office of General Counsel has assured the Region that the same idea applies in the petroleum exclusion.

Clearly, if at any time the cooling tower cleaning waste meets the definition of hazardous waste and it is mixed with the exempted waste, the unit where mixing takes place is a regulated unit.

The interpretations of the exemption contained in this letter are consistent with those of EPA's Office of General Counsel.

I would suggest that EID review Philip's analysis and all available information to determine if the cooling tower cleaning waste is EP-toxic for chromium or is not. You should also determine what quantity of waste is generated and if this waste is/was placed in the surface impoundments after 1980.

Although further investigation/evidence is required to conclusively determine the regulatory status of these sites, I hope the information provided above will prove useful to your staff. If your staff has any questions, please have them call Court Fesmire at (214) 655-6775.

Sincerely,

Randall E. Brown, Chief RCRA Enforcement Branch ADDED 4TH PAGE TO 3RD PAGE SINCE IT WAS SHORT ENOUGH.

cc: Tracy Huges

Office of General Counsel

NMEID

ecol Hobbs: Oddie Seay sextor



PHILLIPS PETROLEUM COMPANY

BARTLESVILLE, OKLAHOMA 74004

918 661-6600

LEGAL

June 27, 1989

Mr. Randall E. Brown Chief, Enforcement Branch U.S. EPA Region VI 6H-C 1445 Ross Avenue, Suite 1200 Dallas, Texas 75202

Dear Mr. Brown:

I have read with interest the letter from Dan Derkics, Chief, Large Volume Waste Section, to Julie Wanslow, which was forwarded to you with Mr. Boyd Hamilton's letter of June 15, 1989, for your use in determining the regulatory status of the four Phillips Petroleum Company natural gas plants requested in Mr. Hamilton's June 8, 1989 letter.

Obviously, it is Phillips's view that Mr. Derkic's letter, read in conjunction with the matters contained in Phillips' letter of May 17, 1989, and the attachments to it, establishes that Phillips' facilities are not subject to RCRA hazardous waste regulations as a result of cooling tower blowdown being directed to the surface impoundments at the plants during normal operation of the plant and cooling towers.

If you have any questions concerning the foregoing or require anything further from Phillips Petroleum Company, please advise.

Very truly yours,

Reese B. Copeland

Attorney

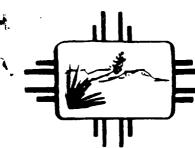
1297 Adams Building Bartlesville, Ok 74004

(918) 661-3758

RBC: am: 599

Boyd Hamilton - New Mexico Health & Env. Dept. Lynn Prince - EPA Region VI, 6H-HS

Mike Fitzpatrick - EPA Headquarters, Washington DC Tracy Hughes - Office of General Counsel, EID Cindy Smith - Phillips Petroleum Company, Bartlesville, OK Knut Am - Phillips Petroleum Company, Odessa, TX



New Mexico Health and Environment Department

Joen 1-20-57

MARALYN BUDKE Acting Secretary

CARLA L. MUTH Deputy Secretary

MICHAEL J. BURKHART Deputy Secretary

RICHARD MITZELFELT
Director

June 15, 1989

Randall E. Brown Chief, Enforcement Branch U.S. EPA Region VI 6H-C 1445 Ross Avenue, Suite 1200 Dallas, Texas 75202

Dear Mr. Brown:

I am enclosing a copy of a letter to Julie Wanslow of our staff from Dan Derkics, Chief, EPA Headquarters concerning the the oil field RCRA exemption and clarifying the differences between gas plant cooling tower wastes and cooling tower blowdown. I believe this will be useful to you in your determination of the regulatory status of the four Phillips Petroleum Company facilities as we requested in our June 8, 1989 letter.

If you have any questions or need additional information, please feel free to contact me at (505) 827-2928.

Sincerely,

Boyd Hamilton

Program Manager

Hazardous Waste Program

cc: Lynn Prince, EPA Region VI, 6H-HS

Mike Fitzpatrick, EPA Headquarters

Knut Am, Manager, Permian Basin Region,

Phillips Petroleum Company

Reese B. Copeland, Attorney for Phillips Petroleum Company

Tracy Hughes, Office of General Counsel, EID

Enclosure

BH/SMM/smm



JUN 6 1989

OFFICE OF SOLID WASTE AND EMERGENCY R

Julie Wanslow Hazardous Waste Section NMEID 1190 Saint Francis Street Santa Fe, N.M. 87503

Dear Ms. Wanslow:

Is. Wanslow:

HAZARDOUS WASTE SECTION
In response to your phone conversation of March 22, 1989 with Mike Fitzpatrick of my staff, we have prepared the following explanations to clarify the boundaries of the oil field RCRA exemption as discussed in the December 1987 EPA Report to Congress (RTC) and given final definition in the July 1988 regulatory determination.

The scope of the exemption as defined in the RTC and regulatory determination is based on the legislative history and Sections 3001(b)(2)(A) and 8002(m) of RCRA. Using these sources the Agency has identified three separate criteria to be used when defining specific waste streams that are exempt. These criteria are listed on pages II-18 and II-19 of the RTC (enclosed.)

In regard to pipeline or gathering line-related wastes, the following excerpts from the criteria in the RTC may prove helpful:

"Primary field operations encompass those activities occurring at or near the well head, but prior to the transport of oil from an individual field facility or a centrally located facility to a carrier (i.e., pipeline or trucking concern) for transport to a refinery or to a refiner.... Waste generated by the transportation process itself are not exempt because they are not intrinsically associated with primary field operations.... Transportation for the oil and gas industry may be for short or long distances." [emphasis added].

According to the Manual of Oil and Gas Terms (sixth edition) there are many terms in common usage within the industry and applied to the various pipelines associated with oil and gas production and transportation (see enclosed definition of "pipeline"). Feeder lines may or may not be exempt depending on the point of custody transfer or other

site-specific factors relating to transportation from the primary field operation as defined in the RTC. Although the Agency used the term "gathering line" in the RTC in reference to a generally small diameter pipe within a primary field operation, the term "gathering line" itself should not be used as the determining factor in defining the scope of the exemption. Rather, the applicability of the criteria in the RTC to the particular line in question should be used in determining the scope of the exemption.

As for gas plant cooling tower wastes, the July 6, 1988, regulatory determination identifies "cooling tower blowdown" as exempt and "gas plant cooling tower cleaning wastes" as non-exempt. The difference between the two is that blowdown is comprised only of water, scale or other wastes generated by the actual operation of the cooling tower; whereas cleaning wastes include any solvents, scrubbing agents or other cleaning materials introduced into the process solely to remove buildup or otherwise clean the equipment and are not included as part of the functional operation of the cooling tower. Since these cleaning wastes can come from any cooling tower, they are not intrinsically derived from primary field operations for natural gas production. The determining factor for defining the exemption is not the frequency with which the cooling tower is blown down, either with or without cleaning agents, but whether the resulting waste is solely derived from the normal operation of the tower for natural gas production or from any added cleaning materials.

I trust these explanations will enable you to better determine the scope of the RCRA exemption as applied to the specific waste streams within your jurisdiction. If you have any further questions please contact Mike Fitzpatrick at (202) 475-6783.

Sincerely,

Dan Derkics

Chief

Large Volume Waste Section

Enclosure

cc: Mike Fitzpatrick

Ivy Main, Office of General Counsel

OIL AND GAS TERMS

Sixth Edition

Annotated Manual of
Legal
Engineering
Tax Words and Phrases

HOWARD R. WILLIAMS

Robert E. Paradise Professor of Natural Resources Law, Stanford University

Stella W. and Ira S. Lillick Professor of Law, Stanford
University

and

CHARLES J. MEYERS

Formerly Richard E. Lang Professor of Law and Dean, Stanford University

1984



MATTHEW BENDER

235 E. 45TH STREET, NEW YORK, N. Y. 10017 450 SANSOME STREET, SAN FRANCISCO, CALIF. 94 11

Displaced gas

THE THE STATE OF STATE OF THE S

Transportation gas (q, ν) which has been displaced by the carrier in order to deliver Consumer gas (q, ν) and which is delivered at a later time when capacity becomes available.

Disposal well

A well employed for the reinjection of salt water produced with oil into an underground formation.

Disqualified transferor

For purposes of the Crude oil Windfall Profit Tax Act of 1980 (q.v.), this term means, with respect to any quarter, any person who: (1) had qualified production for such quarter which exceeded superson's independent producer amount for such quarter, or (2) was not an independent producer for such quarter. Internal Revenue Code § 4992.

Disrepute clause

A clause said to be included in all Petromin contracts with foreign government oil purchasers which enables the Saudi government entity to terminate the contract if the Saudis conclude that the other government acts in a manner which brings discredit to the Saudi government. Conant, "Government-to-Government Agreements," Energy Law 1981, Seminar of the International Bar Association Committee on Energy and Natural Resources at p. 8 (1981). See also, Concession.

Dissolved gas drive

energy, derived from expansion of solution gas, used in the production of oil. Syn.: Solution-gas expansion. Gas escapes from solution within the oil upon reduction of pressure and drives the oil from the reservoir into the well. This form of drive is characterized by rapidly declining pressure and an increasing amount of gas necessary to produce a barrel of oil, with rapidly increasing gas-oil ratios. See also, Reservoir energy.

Distillate

Liquid hydrocarbons, usually colorless and of high API gravity (above 60 degrees), recovered from wet gas by a separator that condenses the liquid out of the gas. This is the older name for the sub-

stance; generally at present the term Natural gasoline (q, ν) or Condensate (q, ν) is used.

Any product separated, or purified, or identified by distillation. See Asiatic Petroleum Corp. v. United States, 183 F. Supp. 275, 12 O.&G.R. 841 (Customs Court 1959).

Distillate fuel oil

A term subject to a variety of definitions. Sometimes the definition is based on the method of production (distillation), but other definitions are based on boiling range, viscosity, or use. See RESIDUAL FUEL OIL. Most commonly the term is used in connection with diesel oil and the light fuel oils used for residential heating. See Hammond, Metz, and Maugh, Energy and the Future 159 (1973). Distillates are classified in grades, called Number 1, 2, 3, 4, 5, and 6 fuels. The specific gravity of fuel oils range from 0.92 to 0.99.

As distinguished from residual fuel oils which are leftovers of refining processes, distillate fuel oils are products of distillation and are lighter. They are used for a variety of purposes, including diesel fuel and for space heating. Residual fuel oils are used under boilers in ships and in power plants. See Zimmermann, Conservation in the Production of Petroleum 85 (1957).

Distribution line

A pipeline other than a Gathering line (q.v.) or Transmission line (q.v.). 49 C.F.R. § 192.3 (1982). See Hamman v. Southwestern Gas Pipeline, Inc., 721 F.2d 140, 78 O.&G.R. 552 (5th Cir. 1983) (concerned with classification of pipeline in order to determine whether it was subject to regulation under the Natural Gas Pipeline Safety Act (NGPSA) (q.v.)

Distribution system

"... [T]he mains which are provided primarily for distributing gas within a distribution area, together with land, structures, valves, regulators, services and measuring devices, including the mains for transportation of gas from production plants or points of receipt located within such distribution area to other points therein. The distribution system owned by companies having no transmission facilities connected to such distribution system begins at the inlet side of the distribution system equipment which meters or regulates the entry of gas into the distribution system and ends with and includes property on the customer's premises. For companies which own both

e station

A location at which gas changes ownership, from one party to anter, neither of which is the ultimate consumer. Also referred to as 1Y GATE (q, ν) station, town border station. American Gas Assotion Bureau of Statistics. Glossary for the Gas Industry 26.

thering facilities

Pipe lines and other facilities used to collect gas from various wells d bring it by separate and individual lines to a central point where is delivered into a single line. In the Matter of Barnes Transportan Co., 18 F.P.C. 369, 7 O.&G.R. 1527 (1957).

the g gas

The first taking or the first retaining of possession of gas for transission through a pipe line, after the severance of such gas, and after
e passage of such gas through any separator, drip, trap or meter
at may be located at or near the well. In the case of gas containing
isoline or liquid hydrocarbons that are removed or extracted in
immercial quantities at a plant by scrubbing, absorption, compresin, or any similar process, the term means the first taking or the
st retaining of possession of such gas for transmission through a
pe line after such gas has passed through the outlet of such plant,
the act of collecting gas after it has been brought from the earth.
iturn Oil & Gas Co. v. Federal Power Comm'n, 250 F.2d 61, 8
1.6G.R. 365 (10th Cir. 1957), cert. denied, 355 U.S. 956, 8 O.&G.R.
13 (1958).

thering line

Piper used to transport oil or gas from the lease to the main pipene in the area. In the case of oil, the lines run from lease tanks to a intral pump station at the beginning of the main pipeline. In the ase of gas, the flow is continuous from the well head to the ultimate onsumer, since gas cannot be stored.

Gathering lines collect gas under fluctuating pressures which are then regulated by regulating stations before the gas is introduced to trunk or transmission lines. Smith v. Inland Gas & Oil Co., 14 v.W. R. 558, 4 O.&G.R. 937 (Sup. Ct. of Alberta 1955).

For purposes of regulation of a pipeline under the Natural Gas ipeline Safety Act, classification of the line as a gathering line, Dispersion line (q, ν) or Transmission line (q, ν) may be of signifi-

cance. See Hamman v. Southwestern Gas Pipeline, Inc., 721 F.2d 140 (5th Cir. 1983) (holding that the gathering line exception in the Act must be restricted to those pipelines that connect a transmission line to a gas well).

See also, PIPELINE

Gathering station

A compressor station at which gas is gathered from wells by means of suction because pressure is not sufficient to produce the desired rate of flow into a transmission or distribution system. America Gas Association Burcau of Statistics, Glossary for the Gas Industr, 26

Gathering system

The GATHERING LINES (q.v.), pumps, auxiliary tanks (in the case of oil), and other equipment used to move oil or gas from the well site to the main pipeline for eventual delivery to the refinery or consumer, as the case may be. In the case of gas, the gathering system includes the processing plant (if any) in which the gas is prepared for the market.

See also, Collecting system; Distribution system

Gathering tax

A tax laid on the process of gathering gas. The Gas Gathering Tax of the State of Texas was declared unconstitutional in Michigan-Wisconsin Pipe Line Co. v. Calvert, 347 U.S. 157 (1954), as a tax on interstate commerce.

Gauge pressure

See PSIG.

Gauger

A person who measures the quantity and quality of oil and/or gas produced.

Gauging a well

Measurement of such characteristics of a well as potential for purposes of prorationing.

ase-in crude

The share of Participation Crude (q.v.) which the host nation nay sell and which the operating oil company must accept. "This rovision, in effect, offers the governments an assured dump market hile they develop their own crude outlets." See "The Economics of nergy and Natural Resource Pricing," Committee Print, A Compition of Reports and Hearings of the Ad Hoc Committee on the omestic and International Monetary Effect of Energy and Other latural Resource Pricing, House Committee on Banking, Currency and Housing, 94th Cong., 1st Sess. March 1975, at p. 100.

See also, Bridging crude; Buyback oil; Concession.

ilippines National Oil Company (PNOC)

A see owned oil company. See Chandler, "Current Developents of Oil and Gas Law: The ASEAN Countries," International Barks'n, I Energy Law 1981 at p. 217.

ysical depletion

See DEPLETION.

ysical waste

Operational losses in the production of oil and gas. There are two rain divisions of loss of oil and gas, namely, surface loss and underround loss. Surface loss of oil is due principally to evaporation and urface loss of gas is due principally to burning at field flares or lowing into the atmosphere. Underground loss is due to failure to ecover the maximum quantity which theoretically could be prouced by dissipation of seservoir pressure.

"[T] the loss or destruction of oil or gas after recovery thereof such s to prevent proper utilization and beneficial use thereof, and the ass of oil or gas prior to recovery thereof by isolation or entrapnent, by migration; by premature release of natural gas from solution in oil, or in any other manner such as to render impracticable the ecovery of such oil or gas." 30 C.F.R. § 221.2(n)(1) (1980).

See also, WASTE.

ΑP

The Petroleum Import Adjustment Program (q.v.)

Pick-up oil

Oil which has escaped from a well or storage tank by overflow or seepage recovered by a Pick-up station (q, ν) .

Pick-up station

A surface pit or other type of trap utilized to gather oil which has escaped from a well or storage tank by overflow or seepage.

g

A scraping device for cleaning and testing petroleum and natural gas pipelines.

Piled steel platform

A conventional drilling and production platform for offshore drilling and production operations. A steel jacket enclosing conductor pipes is pinned to the sea bed by long steel piles and is surmounted by a sizel deck on which is located housing, a drilling rig, and other installations.

Pincher Creek Decision

The decision of the Public Utilities Board of Alberta, pursuant to an application under Section 9 of The Gas Utilities Act, relating to charges and deductions to be allowed for processing costs in determining the value of gas for royalty purposes. See Rae, "Royalty Clauses in Oil and Gas Leases," 4 Alberta L. Rev. 323 at 346 (1965); Muir, "Utilization of Alberta Gas," 13 Alberta L. Rev. 64 (1975).

inch out

A trap formed by the disappearance or wedging out of a porous, permeable rock between two layers of impervious rock.

7

PETROLEUM INCENTIVES PROGRAM (q.v.)

Pipeline

A tube or system of tubes used for the transportation of oil or gas. Types of oil pipelines include: lead lines, from pumping well to a storage tank; flow lines, from flowing well to a storage tank; lease lines, extending from the wells to lease tanks; gathering lines, extend-

ANNOLLING

ing from lease tanks to a central accumulation point; feeder lines, extending from leases to trunk lines; and trunk lines, extending from a producing area to refineries or terminals.

In the case of gas, the Gathering system (q, v) delivers the gas to the main pipeline which takes the gas directly to the distributor at the place of consumption.

Static capacity of a pipeline is calculated by multiplying the square of the pipe diameter (in inches) by .0009714 to give barrels of oil per lineal foot or by multiplying the square of pipe diameter (in inches) by .005454 to give cubic feet of gas per lineal foot. The quantity passing through the line in a given period will depend on initial pressure, flow characteristics, ground elevation, density, delivery pressure, and the booster stations employed.

Of the Lines (1979). "Today there are over 227,000 miles of operating crude and products lines (including gathering lines) in the United States, exceeding by nearly 40 percent the total miles of mainline railroad right-of-way." Id. at 26.

For a detailed examination of the history of pipeline regulation and an order prescribing new criteria for the derivation of maximum permissible rates of return, see Williams Pipe Line Co., 23 F.P.S. 5-685 (F.E.R.C. Opinion No. 154, Nov. 30, 1982).

See also the following:

Pierce, "Reconsidering the Roles of Regulation and Competition in the Natural Gas Industry," 97 Harv. L. Rev. 345 (1983) (arguing that the natural gas market would function more efficiently if Congress deregulated gas pipeline companies and required them to compete against one another);

Malet, "Oil Pipelines as Common Carriers: Issues of Form and Substance," 20 Houston L. Rev. 801 (1983);

her, "Access to Submarine Pipelines and Tariffs: The Legal Framework," [1982] I OGLTR 9;

Adams and Brock, "Deregulation or Divestiture: The Case of Petroleum Pipelines," 19 Wake Forest L. Rev. 705 (1983) (a detailed argument for requiring divestiture of pipeline ownership by integrated oil companies);

Mitchell (ed.), Oil Pipelines and Public Policy (American Enterprise Institute, 1979), discussing proposals for industry reform and reorganization.

See also, Agreement on Principles; Alaska Natural Gas Pipeline Financing Act; ANGTA; ANGTS; Big Inch Pipeline; Distribution Line; Flow Line; Gathering Line; Hepburn Act; High-Volume System; Hinshaw Pipeline; Independent Pipeline; Inter-

ġ.

MEDIATE-VOLUME SYSTEM; LINE LOSS; LINE PACK GAS; LITTLE BIG INCH PIPELINE; LOOP; LOOPING; LOOP LINES; LOW-VOLUME SYSTEM; MANDATORY CONTRACT CARRIAGE; MOTHER HUBBARD CASE; MINIMUM TENDER; NATURAL GAS PIPELINE SAFETY ACT (NGPSA); NORTHERN TIER PIPELINE CO.; PARTIAL LOOPING; PIPELINE CONSENT DECREE OF 1941; PRUDENT PIPELINE STANDARD; QUALITY BANK; SERVICE LINE; TAPLINE; TAPS AGREEMENT; THROUGHPUT AND DEFICIENCY AGREEMENT; TRANS-ALASKA PIPELINE AUTHORIZATION ACT; TRANSIT PIPELINE TREATY; TRANSMISSION LINE; TRANSMISSION SYSTEM; TRUNK LINE; TURNED INTO THE LINE; UNCONNECTED WELL; UNDIVIDED INTEREST PIPELINE; YAMBURG-URENGOI PIPELINE.

Pipeline consent decree of 1941

The agreement entered into by the Department of Justice, 20 major oil companies, and 59 pipeline companies stipulating that dividends paid by the pipeline companies to their shipper-owners would not be unlawful rebates if they did not exceed 7 percent of the Interstate Commerce Commission's valuation of the pipelines' properties. Since the dividend limit was based on valuation rather than on equity in the pipelines, debt capital has been resorted to for much of the cost of constructing pipelines, thus leading to a high rate of return on equity capital. See Report to the Congress by the Comptroller General, Petroleum Pipeline Rates and Competition 14 (July 13, 1979). Debt financing of construction costs has been facilitated by Throughput and defection of the pipeline Rates and Competition 14 (July 13, 1979). Debt financing of construction costs has been facilitated by Throughput and defection of the pipeline Rates and Competition 14 (July 13, 1979). Debt financing of construction costs has been facilitated by Throughput and defection of the pipeline Rates and Competition 14 (July 13, 1979). Debt financing of construction costs has been facilitated by Throughput and defection of the pipeline Rates and Competition 14 (July 13, 1979). Debt financing of construction costs has been facilitated by Throughput and Defection of the cost o

For a discussion of this decree see Adams and Brock, "Deregulation or Divestiture: The Case of Petroleum Pipelines," 19 Wake For est L. Rev. 705 at 729 (1983).

Pipeline gas

A term used to describe gas which has sufficient pressure to enter the high pressure lines of the purchaser for distribution to its customers without further compression and which is sufficiently dry so that the liquid hydrocarbons therefrom will not drop out in the transmission lines. Greenshields v. Warren Petroleum Corp., 241 F.2d 61, 8 O.&G.R. 937 (10th Cir. 1957), cert. denied, 355 U.S. 90; (1957).

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and for shutting in wells producing from a gas cap, called Gas cap allowable $(q, \nu,)$.

It was held in Tidewater Oil Co. v. United States, 339 F.2d 633, 21 O.&G.R. 695 (Ct. Cl. 1964), that the transferor did not have an economic interest in the oil produced by the transferee of a saltwater shut-in allowable and hence the transferee and not the transferor was entitled to the depletion allowance on the oil produced. See "Depletion on Transferred Allowables," 14 O.&G. Tax Q. 133 (1965).

Marlin Oil Corp. v. Corporation Comm'n, 562 P.2d 851, 58 O.&G.R. 225 (Okla. 1977), sustained the power of the Commission to protect correlative rights by ordering the transfer to a new well drilled on a spacing unit the accumulated underages of a prior well on the compact continuous co

See "Transferred Allowables and Substitute Royalties," 5 O.&G. Tax Q. 59 (1956). For discussion of subject with reference to Louisiana law, see Hussey, "Conservation Developments of the Year," 4 L.S.U. Min. L. Inst. 148 at 161–165 (1956).

See also, Fluid injection well; Observation well; Substitute royalty; Unit surplus well.

Transfer rule

The rule designed to prevent proliferation of the so-called independent producers exemption in Section 613A of the Internal Revenue Code. Under this exemption, independent producers continued to be entitled to percentage depletion on a limited quantity of oil. The transfer rule provides that (with limited exceptions) when an interest in and gas in proved oil and gas property is transferred, the independent producers exemption will not apply to the transfere with respect to oil or gas produced from that property. See Linden, "An Analysis of the Transfer Rule" of the Proposed Regs. on Oil and Gas Depletion," 45 J. Taxation 112 (1976).

See also, Depletion, Percentage

Transition zone

An area where the wells produce both free oil and free gas. Union Texas Petroleum v. Corporation Comm'n, 651 P.2d 652 at 665 (Okla. 1981) (dissenting opinion).

Transit Pipeline Treaty

The Agreement Between the Government of the United States of America and the Government of Canada Concerning Transit Pipelines, 28 U.S.T. 7449, T.I.A.S. No. 8720.

See also, Agreement on Principles; Pipeline.

Transmission line

A pipe line extending from a producing area to a refinery or terminal. Syn.: Trunk line.

See also, Distribution line; Gathering line; Pipeline.

Transmission system

tors, tanks, compressors, and their driving units and appurtenances, and other equipment used primarily for transmitting gas from a production plant, delivery point of purchased gas, gathering system, storage area, or other wholesale source of gas to one or more distribution areas. The transmission system begins at the outlet side of the valve at the connection to the last equipment in a manufactured gas plant, the connection to gathering lines or delivery point of purchased gas, and includes the equipment at such connection that is used to bring the gas to transmission pressure, and ends at the outlet side of the equipment which meters or regulates the entry of gas into the distribution system or into a storage area. It does not include storage land or structures." 18 C.F.R. Part 201, Definitions 26B (1980).

Transportation costs

The costs of transporting oil or gas to a market. The operator of a lease upon gaining production will seek to secure a pipe line connection at the well or lease and to make delivery of the oil or gas at such pipeline connection to the purchaser of the oil or gas. Prior to the extension of pipe lines to the lease by a purchaser, the operator of the well or lease may find it necessary to transport the product to a distant pipe line connection or to a railroad or refinery by truck or by his own pipe line.

The lessor is entitled to a royalty free and clear of costs at the wellhead; if the product cannot be disposed of at the wellhead to a purchaser, then the lessor must normally share in the expenses of transporting the product to market. Molter v. Lewis, 156 Kan. 544, 134 P.2d 404 (1943); TREATISE §§ 645-645.3. An occasional lease

Trespass

See Bad faith trespasser; Geophysical trespass; Good faith trespasser; Subsurface trespass.

Tribal lands

See Indian Lands.

Trip tank

A small calibrated tank used to measure the volume of drilling fluid required to fill the hole while pulling pipe from a well. Alberta Energy Resources Conservation Board, *Inquiry Report* 78-8, page 9 e 9, 1978).

Truman proclamation

The Proclamation of September 1945 by President Truman claiming for the United States the natural resources of the subsoil and seabed of the continental shelf beneath the high seas but contiguous to the coasts of the United States as appertaining to the United States, subject to its jurisdiction and control. United States Department of State Bulletin, No. 327 (Sept. 30, 1945), p. 485.

See also, OUTER CONTINENTAL SHELF.

Trunk line

A pipe line for the transportation of oil or gas from producing areas to refineries or terminals. Syn.: Transmission line.



See Landowners' royalty pool; Royalty trust.

Tubing

A string of pipe set into a well through which oil is produced.

Syn.: Oil STRING.

Tubing a well

Setting and sealing into the well a string of pipe, called tubing, after perforations have been made in the casing or the well has been drilled to the desired total depth. The oil or gas is produced through the tubing which may have screens at the level of the producing stratum to strain out sand and other foreign matter.

*

Tubing pressure

See SHUT-IN PRESSURE.

Tubular goods

Well casing and tubing, drill pipe, standard pipe, line pipe, etc.

Turbodrilling

A method of drilling wells in which "the bit is turned not by rotation of the drill string, as in rotary drilling, but by a downhole turbine, driven by the fluid pumped down through the drill stem. The turbodrill can thus be thought of as using a hydraulic transmission system that gets power to the bottom of the hole in contrast to the rotary drilling system's dependence on mechanical transmission." Campbell, The Economics of Soviet Oil and Gas 108 (1968). For a discussion of the development and use of this method of drilling see Id. at 108-120.

See also, Cable tool drilling; Rotary drilling.

Turkish National Petroleum Co. (TPAO)

For a discussion of this company see Shwadran, The Middle East, Oil and the Great Powers 491 (3d ed. revised and enlarged, 1973).

Turned into the line

A pipeline has begun to run oil from field tanks into which a well's production first goes, so that the well's production is being marketed. Ball, Ball, and Turner, *This Fascinating Oil Business* 109 (2d ed. 1965).

Turning to the right

A colloquialism for actual drilling of a well. Peterson, "Extensions and Suspensions of Federal Oil and Gas Leases," Rocky Mt. Min. L. Fdn. Inst. on The Overthrust Belt—Oil and Gas Legal and Land Is sues 12-1 at 12-7 (1980).

Turnkey contract

A contract in which an independent drilling contractor undertake to furnish all materials and labor and to do all the work required to complete a well in a workmanlike manner, place it on production

- 1. Exempt wastes must be associated with measures (1) to locate oil or gas deposits, (2) to remove oil or natural gas from the ground, or (3) to remove impurities from such substances, provided that the purification process is an integral part of primary field operations.⁵
- 2. Only waste streams intrinsic to the exploration for, or the development and production of, crude oil and natural gas are subject to exemption. Waste streams generated at oil and gas facilities that are not uniquely associated with the exploration, development, or production activities are not exempt. (Examples would include spent solvents from equipment cleanup or air emissions from diesel engines used to operate drilling rigs.)

Clearly those substances that are extracted from the ground or injected into the ground to facilitate the drilling, operation, or maintenance of a well or to enhance the recovery of oil and gas are considered to be uniquely associated with primary field operations. Additionally, the injection of materials into the pipeline at the wellhead which keep the lines from freezing or which serve as solvents to prevent paraffin accumulation is intrinsically associated with primary field operations. With regard to injection for enhanced recovery, the injected materials must function primarily to enhance recovery of oil and gas and must be recognized by the Agency as being appropriate for enhanced recovery. An example would be produced water. In this context, "primarily functions" means that the main reason for injecting the materials is to enhance recovery of oil and gas rather than to serve as a means for disposing of those materials.

3. Drilling fluids, produced waters, and other wastes intrinsically derived from primary field operations associated with the exploration, development, or production of crude oil, natural gas, or geothermal energy are subject to exemption. Primary field operations encompass production-related activities but not transportation or manufacturing activities. With respect to oil production, primary field operations encompass those activities occurring at or near the wellhead, but prior to the transport of oil from an individual field facility or a centrally located facility to a carrier (i.e., pipeline or trucking concern) for transport to a refinery or to a refiner. With respect to natural gas production, primary field operations are those activities occurring at or near the wellhead or at the gas plant but prior to that point at which the gas is transferred from an individual field facility, a centrally located facility, or a gas plant to a carrier for transport to market.

Thus, wastes associated with such processes as oil refining, petrochemical-related ... manufacturing, or electricity generation are not exempt because those processes do not occur at the primary field operations.

Primary field operations may encompass the primary, secondary, and tertiary production of oil or gas. Wastes generated by the transportation process itself are not exempt because they are not intrinsically associated with primary field operations. An example would be pigging waste from pipeline pumping stations.

Transportation for the oil and gas industry may be for short or long distances. Wastes associated with manufacturing are not exempt because they are not associated with exploration, development, or production and hence are not intrinsically associated with primary field operations. Manufacturing (for the oil and gas industry) is defined as any activity occurring within a refinery or other manufacturing facility the purpose of which is to render the product commercially saleable.

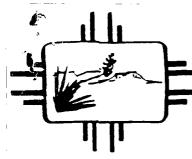
Using these definitions, Table II-1 presents definitions of exempted wastes as defined by EPA for the purposes of this study. Note that this is a partial list only. Although it includes all the major streams that EPA has considered in the preparation of this report, others may exist. In that case, the definitions listed above would be applied to determine their status under RCRA.

Waste Volume Estimation Methodology

Information concerning volumes of wastes from oil and gas exploration, development, and production operations is not routinely collected nationwide, making it necessary to develop methods for estimating these volumes by indirect methods in order to comply with the Section 8002(m) requirement to present such estimates to Congress. For this study, estimates were compiled independently by EPA and by the American Petroleum Institute (API) using different methods. Both are discussed below.

Estimating Volumes of Drilling Fluids and Cuttings

EPA considered several different methodologies for determining volume estimates for produced water and drilling fluid.



MARALYN BUDKE Acting Secretary

CARLA L. MUTH
Deputy Secretary

MICHAEL J. BURKHART

RICHARD MITZELFELT
Director

June 8, 1989

Randall E. Brown Chief, Enforcement Branch U.S. EPA Region VI, 6H-C 1445 Ross Avenue, Suite 1200

Dear Mr. Brown:

Dallas, TX 75202

Mark Con Lang

PHILLIPS PETROLEUM CO.

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The New Mexico Hazardous Waste Program is requesting an EPA interpretation of the July 6, 1988 Federal Register because the interpretation rendered will, or at least could, affect all states within Region VI. This interpretation is requested by June 30, 1989 in order for the state to determine if the facilities which raised the issue, are or are not in violation of a state issued compliance order. It should be noted that these facilities were required by the 1984 EPA consent agreements to submit to the state RCRA closure plans for review and processing which may suggest EPA involvement.

The Hazardous Waste Program met with Phillips Petroleum Company on May 1, 1989 to discuss Phillip's non-compliance with the April 19, 1988 Compliance Order issued by the EID. The purpose of the meeting with Phillips was to outline our concerns at each of the four facilities Artesia, Eunice, Lee and Lusk. Our primary concerns are that the groundwater monitoring systems at each facility are inadequate and the analytical results for the October 1988 CME indicate elevated levels of total chromium in the groundwater at all four facilities. Organic constituents such as benzene, toluene, and ethylbenzene are also present in the groundwater at all the facilities except Lusk. Phillips indicated during the meeting that they did not believe they were subject to RCRA based upon the regulatory determination published in the Federal Register for July 6, 1988, 53 Federal Register 25,446 which exempts cooling tower blowdown. Phillips also indicated that they were working with the New Mexico Oil Conservation Division to close the surface impoundments (RCRA units) de sulla wasta disposal units.

Mr. Brown June 8, 1989 Page 2

As a result of this meeting Phillips was asked to provide a letter stating their position and a statement of their past activities at the surface impoundments (RCRA units). We have received their information and are forwarding it to you with this letter.

At the time of the meeting with Phillips the Hazardous Waste Program's position was that the chromium waste that was being regulated under RCRA was the result of gas plant cooling tower cleaning wastes and therefore would not be exempt. Staff has had several conversations with Mike Fitzpatrick with U.S. EPA Headquarters (contact person for the oil and gas exemption), both before and after the meeting with Phillips. Prior to the May 1, 1989 meeting our staff's understanding of the July 6, 1988 Federal Register, the Report to Congress, and conversations with Mike Fitzpatrick was that cooling tower blowdown was a material that had not been altered by any additives or cleaning agents and that gas plant cooling tower cleaning wastes were cooling tower blowdown that had been altered by the addition of cleaning agents such as solvents or chromium.

Upon further conversations with Mike Fitzpatrick after the meeting with Phillips, and after explaining the process Phillips uses at the gas plant cooling towers, it seems that cooling tower blowdown can have additives in it which enable the gas plant to operate. If the plant is shut down for cleaning or for cleaning agents to be added to the cooling tower water, then those wastes generated from the cleaning process would be considered gas plant cooling tower cleaning wastes.

Staff has reviewed the information packet provided by Phillips which includes the discussion explaining their position and the Certificate of No Hazardous Waste Activity. At this time EID believes that the chromium in the impoundments is exempt from RCRA and therefore the four Phillips facilities Artesia, Eunice, Lee and Lusk are not regulated under RCRA.

We believe this is an extremely important issue. If gas plant cooling tower cleaning wastes includes cooling tower blowdown with hazardous constituent additives, then several hundred additional facilities would potentially be added to the RCRA system in New Mexico. If however these gas plants are exempt then several hundred facilities are potentially not being regulated for the presence of possible EP Tox levels of chromium in their surface impoundments.

Mr. Brown June 8, 1989 Page 3

Should you have any questions, please feel free to contact me at (505) 827-2928.

Sincerely,

Boyd Hamilton

Boyd Hamilton Program Manager Hazardous Waste Program

cc: Lynn Prince, EPA Region IV, 6H-HS

Mike Fitzpatrick, EPA Headquarters

Knut Am, Manager, Permian Basin Region,

Phillips Petroleum Company

Reese B. Copeland, Attorney for Phillips Petroleum Company

Tracy Hughes, Office of General Counsel, EID

Enclosures

BH/SMM/smm



May 17, 1989



MAY 22 1989

Mr. Boyd Hamilton
Program Manager
Hazardous Waste Program
New Mexico Health and
Environment Department
1190 St. Francis Drive
Santa Fe, New Mexico 87503

Re: RCRA Status Artesia, Eunice, Lusk and Lee Gas Plants - NMD000709667, NMD000709634, NMD000709675, NMD000709659

Dear Mr. Hamilton:

At the meeting held May 1, 1989 in Santa Fe, between yourself, your staff and Phillips Petroleum Company (Phillips) representatives, you requested that Phillips furnish you a written statement of its position concerning the status of the above-referenced facilities, given the EPA's Regulatory Determination of June 29, 1988 (RD); the earlier Consent Agreement and Final Order's pertaining to the facilities entered in EPA Dockets numbered RCRA VI-311-H, RCRA VI-312-H, RCRA VI-313-H, RCRA VI-314-H; and the facts concerning the operation of the cooling towers with the discharge to the surface impoundments at each of the facilities.

Phillips has reviewed the matter internally. Its position is that under the RD (Attachment I), cooling tower blowdown is expressly declared to be within the category of wastes exempted from regulation as Subtitle C wastes under Section 3001(b)(2)(A) of the 1980 Amendments to RCRA, 42 USC 6921(b)(2)(A), 53 F.R. 25453-25454.

The only waste streams directed to the surface impoundments at the four plants after November 19, 1980, which could have made the impoundments RCRA units were the cooling tower blowdown streams. The cooling tower blowdown streams would not contain chemical or other cleaning wastes (see Attachment II, Certificate No Hazardous Waste Activity Cooling Tower Usage, Artesia, Eunice, Lee and Lusk Gas Plants). Therefore, none of the surface impoundments are RCRA units, nor are they subject to regulation by the EID, but rather are solid waste disposal facilities subject to the jurisdiction of the OCD.

The Consent Agreement and Final Orders (Order) applicable to their respective plants (Attachments III, IV, V, and VI), do not alter this result. Each order recites that its effects are limited to that particular proceeding and that neither the findings of fact nor conclusions of law may be used for any other purpose or in any other proceedings except for the purposes of computing penalties for any alleged violations which represent a continuation or repetition of alleged violations contained in the findings of fact and conclusions of law in each Order. Thus, any future violation, even if a continuation or repetition, would need to be proved up independently of the Order and the recited findings of fact and conclusions of law, and all facts, including jurisdictional facts, would have to be proved up without recourse to the Order.

The Order pertaining to Lusk recites that it is specifically understood by EPA that Respondent (Phillips) is in the process of seeking to obtain closure approval for this facility from the State of New Mexico. The Orders pertaining to Artesia, Lee and Eunice, each recite that Phillips has submitted a closure plan and a post-closure plan to EPA and the State of New Mexico. These statements of fact were correct. statements do not operate to extend the effects of the Orders to closure proceedings, because the Orders recite that the fact findings and conclusions of law are limited to the individual enforcement proceeding, as stated above, and because New Mexico had acquired interim authorization to administer the RCRA program with regard to interim status closure prior to the entry of the Orders, in late August or early September 1984. Thus, interim status closure was New Mexico's responsibility, not EPA's. anything, these recitations imply an approval by EPA of the interim status closure and post-closure plans which were submitted by Phillips, (see letter dated February 15, 1984, to Frank Collis from William H. Taylor, Jr., Attachment VII, concerning the Lusk Plant).

Finally, Phillips' position is that the record reflects it effected closure in conformity with the interim status closure plans submitted. Thus, any obligations which might be improperly inferred from these factual recitations have been discharged.

Phillips has submitted the groundwater monitoring data gathered at these plants over the past year to OCD in order to close the impoundments at all four plants with such groundwater cleanup as is appropriate. It appears that groundwater cleanup will be required at Lee and perhaps Eunice plants. The cleanup does not appear to be required because of disposal activity connected to the surface impoundment at either plant.

It is Phillips' understanding that you desire this written statement of Phillips' position in order to submit it with a write-up of your own position to Region VI of EPA to obtain EPA's view of the status of these facilities under the RD, the Orders and the facts concerning the operation of the cooling towers and the cooling tower blowdown discharges to the surface impoundments at each of the plants. It is Phillips' view that the matter is so clear on its face that no interpretation by EPA is necessary, but Phillips is willing to cooperate in seeking it. Phillips does not agree that it will accept without further resort to the courts or other legal process a determination by EPA that these surface impoundments are subject to Subtitle C regulation under RCRA. It is Phillips' understanding that you will furnish it a complete copy of the submittal when it is forwarded to EPA.

Phillips' position, summarized above, is discussed below in greater detail with respect to the development of the RD.

The Report to Congress (Report), compiled in response to Section 8002(m) of the Resource Conservation and Recovery Act (RCRA), 42 USC 6982(m) was submitted in December of 1987. EPA, in a Federal Register Notice, advised the public of the availability of the Report and especially invited public comment on the scope of the exemption as stated in the Report, as well as another subject not relevant to this discussion. 53 F.R. 81.

The Report presented EPA's tentative definition of the scope of the exemption for oil and gas drilling fluids, produced waters and other wastes associated with the exploration, development or production of crude oil or natural gas. (II-16-II-19).

The Report tentatively concluded that the "other wastes associated" meant that the wastes must be intrinsically derived from "primary field operations associated with the exploration, development or production of crude oil or natural gas", and that primary field operations, with respect to natural gas production, were those activities occurring at or near the wellhead or at the gas plant, but prior to that point at which the gas is transferred from an individual field facility, a centrally located facility, or a gas plant to a carrier for transport to market. The Report also tentatively concluded inter alia that exempt wastes must be associated with measures to remove impurities from oil or gas, provided this was an integral part of primary field operations. (Ibid.)

Table II-1 of the Report contained a partial list of exempt and non-exempt wastes. Neither cooling tower blowdown nor

Mr. Boyd Hamilton -4-May 17, 1989 gas plant cooling tower cleaning wastes are mentioned in either Volume III of the Report, in its Glossary of Terms, contains the following definitions: The emptying or depressuring of a material from a vessel. The material thus discarded.

A structure in which air contact is used to Cooling Tower: cool a stream of water that has been heated by circulating through a system. The air flows counter- or cross-currently to the water.

list.

Blowdown:

Gas Plant: An installation in which natural gas is processed to prepare it for sale to consumers. A gas plant separates desirable hydrocarbon components from impurities in natural gas.

Many members of the oil and gas industry accepted EPA's invitation to comment on the scope of the exemption, as did State and Federal authorities, the environmental community and numerous individuals.

Among the submissions available to and presumably relied upon by EPA in issuing the subsequent RD is a document titled "Production Waters Associated With The Production, Processing, Transmission and Storage of Natural Gas: Literature Survey" which EPA characterized as technical support material in the administrative record for the RD (Certified List of Documents Comprising the Adminstrative Record, Alaska Center for the Environment v. Thomas, Case No. 88-1715, U.S. Ct. of App. DC Cir. F-88-OGRA-SO528).

This document gives a detailed description of the waters generated from gas processing including cooling tower blowdown. It is noted in the discussion of the chemical composition of such waters that they typically contain chromium, zinc, and phosphate as corrosion inhibitors, and that chlorines and small concentrations of biocides may be added to these waters. (SO528 pp. 5-69 - 5-76).

Among comments the Agency received on the scope of the exemption, the Gas Processors Association urged that all gas related associated wastes, such as contaminated glycol, iron sponge, and amines should be exempt from federal regulations. The Agency, in its response, noted that as explained in the RD, certain types of gas production associated wastes have remained exempt from RCRA Subtitle C. (F-88-OGRA-S0707 p. 24).

-5-Mr. Boyd Hamilton May 17, 1989 The Interstate Natural Gas Association urged that all wastes generated by natural gas treatment and processing operations should be exempt. Waste iron sponge, glycol, corrosion inhibitors and all other associated waste should be The Agency acknowledged the comments and pointed out that, as explained in the RD, certain types of gas treatment and production associated wastes have remained exempt from RCRA Subtitle C. (SO707 p. 27). The Rocky Mountain Oil and Gas Association commented that the waste exemption should be much broader than the EPA The Agency's response was that the waste exemption has defined. been defined in the RD. (SO707 p. 28). The RD's list of exempt wastes lists cooling tower blowdown, 53 FR 25454. It is submitted that the conclusion is inescapable that on this record, cooling tower blowdown from gas plants containing treating levels of corrosion inhibitor and small concentrations of biocides is an exempt waste. I certify, under penalty of law, that I have personally examined and am familiar with the information submitted in the foregoing five pages, and the attachments hereto, and based upon my inquiry of those individuals immediately responsible for obtaining the information, I believe the submitted information is true, accurate and complete. I am aware that there are significant penalties for submitting false information, including the possibility of fine and imprisonment. Very truly yours, PHILLIPS PETROLEUM COMPANY en Manager, Permian Basin Region 4001 Penbrook Odessa, Texas 79762 KA:RBC:am:535 Enclosures

CERTIFICATE NO HAZARDOUS WASTE ACTIVITY COOLING TOWER USAGE ARTESIA, EUNICE, LEE AND LUSK GAS PLANTS

At all times during plant operations from November 19, 1980 to present, cooling towers at the above mentioned gas plants have been used to cool gas streams, gas treating solutions and, at Eunice and Lee, engine jacket water. This activity occurred prior to the point at which gas is transferred from the plant to a carrier for transport to market. Water in the towers is continuously recirculated and evaporated, which causes total dissolved solids and chloride content to steadily increase.

Total dissolved solids and chloride levels are controlled by "blowing down" or removing a portion of the total water volume in the tower and then making up the blowdown volume with fresh water; the process is one of continuous displacement. Chemicals used to control corrosion and maintain proper pH are added to the basin with the fresh water make-up. The tower is also periodically "shocked" with biocides to control algae and fungus growth to enhance the corrosion inhibition within the system. It should be noted the blowdown is normally shut in while biocides are in use. The biocides are spent before blowdown is resumed.

Blowdown water typically has a high concentration of total dissolved solids and chlorides. It will also contain the

treating level of chemicals used for corrosion inhibition. Both chromate and non-chromate corrosion inhibitors have been used since November 19, 1980. Cooling tower blowdown rates currently average 25 gallons per minute at the Artesia, Eunice and Lee Plants. Lusk is presently shut down and has been for some time.

Cooling towers must be cleaned on a periodic basis for proper operation (approximately once every five years). Cleaning consists of removing the sludge from the basin and removing scale from the cooling coil heads and laterals. Sludge is removed manually or by vacuum truck for proper disposal. The cooling coils and laterals are mechanically cleaned by sandblasting. Typically, the operations are done together. The materials have been tested and have not tested as hazardous waste.

Cooling tower basins and cooling coil heads and laterals have not been chemically cleaned in this region or at any of these plants.

Given the Regulatory Determination of June 29, 1989, which declares cooling tower blowdown to be an exempt waste and therefore not subject to regulation under Subtitle C of RCRA, there has been no hazardous waste generated, stored or disposed of at the Artesia, Eunice, Lee and Lusk natural gas plants since November 19, 1980.

I certify, under penalty of law, that I have personally examined and am familiar with the information submitted in this Certificate, and based upon my inquiry of those individuals immediately responsible for obtaining the information, I believe that the submitted information is true, accurate and complete. I am aware that there are significant penalties for submitting false information, including the possibility of fine and imprisonment.

Dated: May 17, 1989.

Knut Am

Permian Basin Region
PHILLIPS PETROLEUM COMPANY
4001 Penbrook

Odessa, Texas 79762



LEGAL

May 5, 1989

Mr. Boyd Hamilton
New Mexico Health and
Environmental Department
1190 St. Francis Drive
Santa Fe, New Mexico 87503

Re: May 1, 1989 Meeting - Lusk, Lee, Eunice, and Artesia Natural Gas Processing Plants

Dear Mr. Hamilton:

At the May 1, 1989 meeting, Phillips agreed to furnish you a written statement of its position with respect to the cooling tower blowdown production waste exemption as it applies to the above four plants. It is Phillips' understanding that you desired this statement in order to submit the same to Region VI of the United States Environmental Protection Agency, together with your position statement on those issues, in order to obtain a determination from Region VI, U.S. EPA, whether the plants are within or without the subtitle C regulations because of the chromate containing cooling tower blowdown water at the plants and the rather long and tortured procedural history involving these four plants.

It is Phillips' understanding that until the regulatory determination is received, the NOV's previously issued concerning the groundwater monitoring system will remain in abeyance.

You made it clear that if the EPA decides that the plants contain regulated units, your agency will be vigorous with respect to the groundwater monitoring program.

Phillips has agreed to furnish the statement of its position, in writing, and will do so. However, it is Phillips' position that the regulatory determination of EPA as published in the Federal Register for July 6, 1988, 53 Fed. Reg. 25,446, is clear on its face and requires no agency interpretation. In fact, if the Agency were to conclude otherwise, it is Phillips' position it would be required to go through Notice and Rulemaking.

Mr. Boyd Hamilton -2-May 5, 1989 You should have Phillips' written position statement in hand within the time frame originally agreed upon, ten working days from May 1, 1989. Thank you again for the courtesy extended to us by you and your staff. Very truly yours, Reese B. Copeland 1297 Adams Building Bartlesville, OK 74004 (918) 661-3758 RBC: am: 525



1190 St. Francis Drive Santa Fe, New Mexico 87503 GARREY CARRUTHERS
Governor
MUTH
Secretary

APR 2 4 IMCHAEL J. BURKHART Deputy Secretary OIL COLSERVATION DIV.

CERTIFIED MAIL
RETURN RECEIPT REQUESTED

Mr. William F. Ballard, Manager Phillips Petroleum Company 12 A4 Phillips Building Bartlesville, OK 74004 APR 10 1989

RE: Phillips Plants Artesia, Eunice, Lee, and Lusk
NMD000709667, NMD000709675, NMD000709659, NMD000709634

Dear Mr. Ballard:

A Compliance Order was issued to Phillips Petroleum Company (Phillips) on April 19, 1988. The Compliance Order required Phillips to install by May 1, 1988, an adequate groundwater monitoring system.

The Environmental Improvement Division (EID) of the New Mexico Health and Environment Department has determined that the groundwater monitoring systems are inadequate at the Artesia, Eunice, Lee and Lusk facilities based upon a review of four quarters of water elevation data and the site maps. Groundwater flow directions change seasonally and therefore, three wells are not always located downgradient of the RCRA units. Downgradient wells are not always located to immediately detect a release from the RCRA units. The seasonally variable groundwater flow directions indicate that the upgradient wells at some of the Phillips facilities may be affected by the facility.

In addition, EID has noted that Phillips is filtering the metals samples. The New Mexico Hazardous Waste Section requires analysis of total metals, not filtered metals. All future metal samples must be unfiltered and analyzed for total metals.

The groundwater monitoring data, including the quarterly water elevation data and the groundwater analyses, are currently being reviewed in reference to Phillips' closure plans.

Turbidity values for most wells are in excess of the recommended turbidity value of 5 nephelometric turbidity units (NTUs) for ensuring the collection of representative samples. Most of the wells have become increasingly turbid since they were installed. EID strongly suggests Phillips redevelop the wells which have turbidity values exceeding 5 NTUs. Developed water that contains hazardous constituents must be disposed of in a RCRA unit or drummed and shipped offsite.

William F. Ballard April 4, 1989 Page 2

EID requests that Phillips meet with EID staff as soon as possible and provide in writing to EID within thirty (30) days of receipt of this letter, Phillips' plans to comply with the April 19, 1988 Compliance Order.

Please contact Suzanne Moore-Mayne at (505) 827-0170 to schedule a meeting and to answer any questions or provide additional information.

Sincerely,

Boyd Hamilton

Program Manager

Hazardous Waste Program

BH/SMM/smm

cc: Janie Hernandez, U.S. EPA - Region VI

Gary McCaslin, EID District IV

Tracy Hughes, HED

Plant	ARTESIA	7

lant F	IN I ESIM					
nonitor well	MW-1	MW-Z	MW-3	MW-4		
Number ev. to top of casing (2")	36/1.47	3605.23	l _	3606.13		
et casing (2)	3611.47	3603.2	3800 0	3808.73		
8/11/88	3550.56	3550,80	3550,96	3550,50		
9/7/28		3550,55	3550, 58	3550,32		
10/26/88	3550,43	3550,47	3550,47	3550,29		
218/89	3550,04	3549,85	3549,81	3549,66		
				·		
					·	
		<u> </u>				
						
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						ĺ

Pla	nt_	_EU	W	ICE

nitor well	<u> </u>		1				T
Number	MW-1	MW-Z	MW-3	MW-4			-
casing (2")	3562.63	3561.74	3561.64	3561.87			
	_						-
8/31/88	3447,28	3443.69	1	3444.15		 	
11/2/88	3447,45	3443,67	3443,84	3444.10		<u></u>	
11/89	3447,31	3443,61	3443,69	3443,96			ļ
							
					: :		
							
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							,
					· · · · · · · · · · · · · · · · · · ·		
					•		
]

Plant	LEE					
monitor well Number	MW-1	MW-Z	Mw-3	MW-4		
of casing (2")	3979, 27	3980.59	3980.37	3980.29		
8/29/58	3884,69	3883.77	3893,75	3883.75		
10/31/88	3984,43	38p3,57	3213,56	38P3,57		
1/30/89	388S.2Y	3883.S >	3#R3,\$0	3883,51		
					-	-

Plant LUSK

Plant LU	sk					
						
monitor well Number	MW-1	MW-Z	Mw-3	MW-4		
of casing (2")	357/.6/	3566.99	3565.77	3565.86		ļ
8/15/88	353/33	3< X 88	3524,36	3525,83		
10/24/88		3529,22		3526,58		
2/13/89	3531,46	354.33	3524,71	3526,05		
		——————————————————————————————————————				
						



ODESSA, TEXAS 79762 4001 PENBROOK

April 5, 1989

Groundwater Monitoring Analyses Artesia, Eunice, Lee and Lusk Plants

Mr. Dave Boyer Environmental Bureau Chief New Mexico Oil Conservation Division P. O. Box 2088 Santa Fe, New Mexico 87501

Dear Mr. Boyer:

Per your request, attached please find copies of the fourth quarter groundwater monitoring analyses for the above referenced plants.

If you should have any questions regarding this information, please contact me at (915) 367-1316.

Very truly yours,

Michael D. Ford

Environmental Analyst

MDF

Attachments

Page

Received: 02/10/89

RAS

571N REPORT 03/09/89 16:16:17

Work Order # 89-02-119

REPORT	Phillips Petroleum	PREPARED Radian Analytical Services	`
	Radian		//
	B1. 1	PO Box 201088	6
	Austin	Austin, TX 78720-1088	CERT
ATTEN	Linda Bendele	ATTEN	
		PHONE 512-454-4797	CD
CLIENT	PHILLIPS P SAMPLES 3		
COMPANY	etroleum		
=ACILITY	Odessa, TX		ı
		Unknown compounds present in both GC samples	ples
HOBK IN	01+0001U		
TAKEN	MF.		
TRANS	UPS		
TYPE			
P. O. #	errigiumpetumminis tenishitatinis gamagikon fritaritis dirigitis kiristis da janaginia dinggarandikanikanikan		
INVOICE	under separate cover		

CERTIFIED BY

CONTACT BENDELE

|--|--|

2/89-414 Quenter

Artesia MW-

SAMPLE IDENTIFICATION

reagent blank Artesia MW-2

DG601C DG302 PA60 Silver, Digestion, method 3020 Gross alpha radiation Sodium, EPA method 602 Cadmium, Gross beta radiation Arsenic, graphite AA <u>Specific conductance</u> Mercuru, cold vapor Vitrate, langanese, ron, ICPES otal coliform luoride, igestion, hromium, hloride, graphite AA ICPES ICPES ICPES colorimetri ICPES ICPES method 6010 ICPES

TEST CODES and NAMES used on this report

XYLENE	TURB	TOX	TOC	S04 IC	SE G	RA 226	PHEN	
Xulenes, EPA 602	rbiditu	tal organic h	ta		Selenium, graphite AA	adiun	Total phenolics	化二 化二甲基 一角化的一心

Page 2 Received: 02/10/89

Austin REPORT
 Results By Test

RAS

Work Order # 89-02-119

Artesia MW-2	Artesia MW-1	SAMPLE Id	Artesia MW-1 02 Artesia MW-2	Sample Id	Artesia MW-1 02 Artesia MW-2	Sample Id
02/22/89	02/22/89	Test: D36010	<0.005 <0.005	Test: CD E	<0.03 <0.03	Test: AG E
. C.Ti	€0.04	Test: FE E	160 660	Test: CL IC	15.7(1.4) pci/L 21.7(2.5) pci/L	Test: ALPHA
2.7	1. 9	Test: F IC	210	Test: COLI I	0. 013	Test: AS G
1400 3700 3600	1400 1400 1400	Test: MHO	<0.03 <0.03	Test: CR E	0. 07 2. 0	Test: BA E
1. 9	0.09	Test: MN E	02/23/89	Test: DG3020	15.9(1.2) pci/L 32.2(2.9)	Test: BETA

Page 3 Received: 02/10/89 Samp le Sample Artesia MW-1 tesia MW-2 SAMPLE SAMPLE ស Test: DG6010 est: NA E 3 110 ug/ml RAS Austin REPORT
Results By Test Test: NO3 est: FE E < 0.020mg/L as N Test:F IC est: PB G <0.001 0.003* ug/ml mq/1 Test: PH Test: MHO Work Order # 89-02-119 Continued From Above pH units 3600 3600 6.92 6. 98 6.87 6.99 6.91 6.91 6.91 7. 12 eq/L as phenol Test: PHEN Test: ME <0.005 0.008*****

Page 4 Received: 02/10/89 Sample Id Artesia MW-1 Artesia MW-2 SAMPLE ស្ល 2 2.14(.08) pci/L 0.52(.04) pci/L Test: RA 226 RAS Austin Test: SE G <0.004 <0.004 Results By Test uq/ml Test: <u>SO4 IC</u> 8 4 Test: TOC Work Order # 89-02-119 $\ddot{\omega}$ Test: TOX 0.05* 0.04* 0.04* 0.03* 0.04* 0.05* 0.04* 0.05

Page 5 Received: 02/10/89

SAMPLE ID Artesia MW-1

RAS Austin

REPORT

Work Order # 89-02-119

NAME EPA method 602

Category

FRACTION O1J TEST CODE EPA602
Date & Time Collected 02/09/89 Results by Sample

			ANALYST JB	
108-88-2	71-43-2	CAS#	INJECTED 02/13/89	
Toluene	Benzene	COMPOUND RESULT	FILE #	
0.7* 0.20	16 0.20	RESULT DET LIMIT	UNITSUq/L	VERIFIED DMV

SURROGATES

541-73-1

1,3-Dichlorobenzene

1,4-Dichlorobenzene

Chlorobenzene-A

0.30

0.30

0.30

Ethylbenzene

1,2-Dichlorobenzene

S

0.40

0.40

95-50-1

106-46-7

108-90-7

100-41-4

8-30-86 a, a, a-Trifluorotoluene 130 0% recovery

-4		INSTRMT HACH		SAMPLE ID Artesia MW-1	(1)See Appendix			INSTRMT 403	" **********************************	SAMPLE ID Artesia MW-1	Page 6 Received: 02/10/89
Turbidity 21 1.0	ANALYTE RESULT DET LIMIT	AMALYZED 02/10/89	VE	FRACTION <u>OIA</u> TEST CODE TURB Date & Time Collected 02/09/89	A for glossary of report and data flag	Mercury ND 0.0002	ANALYTE RESULT DET LIMIT	AMALYZED 02/22/89	VΕ	FRACTION OIL TEST CODE HG C Date & Time Collected 02/09/89	RAS - Austin REPORT Results by Sample
		UNITS NTU	VERIFIED LM	JRB NAME Turbidity /89 Category	definitions.			UNITSuq/ml	VERIFIED RHH	/89 NAME Mercury, cold vapor	Work Order # 89-02-119

CORPORATION

	SAMPLE ID Artesia Mi-1	Received: 02/10/89	Page 7
		,	RAS .
Date & Time Co	FRACTION OLU	Results by Sample	- Austin
Date & Time Collected 02/09/89	TEST CODE XYLENE	Sample	REPORT
Category	NAME Xylenes, EPA 602		Work Order # 89-02-119

VERIFIED CL

CAS # COMPOUND RESULT DET LIMIT 1330-20-7 Total xylenes <u>3.1</u> <u>0.20</u>

FILE # _____

INJECTD 02/13/89

ANALYST INSTRMT

B

UNITS Ug/L

SURROGATES

a, a, a-Trifluorotoluene 130 3% recovery

8-80-86

(1)See Appendix A for glossary of report and data flag definitions.

*: *:

Page 8 Received: 02/10/89

SAMPLE ID Artesia MW-2

RAS Austin

REPORT

Work Order # 89-02-119

FRACTION 02J TEST CODE EPA602
Date & Time Collected 02/09/89 Results by Sample NAME EPA method 602

VERIFIED DMV

95-50-1	541-73-1	106-46-7	108-90-7	100-41-4	108-88-3	71-43-2	CAE#	ANALYST JB
1,2-Dichlorobenzene	1,3-Dichlorobenzene	7 1,4-Dichlorobenzene	7 Chlorobenzene-A	f Ethylbenzene	3 Toluene	Benzene	# COMPOUND	FILE #
ND 2.0	ND 2.0	ND 1.5	ND 1.5	66 1.5	2.7* 1.0	15 1.0	RESULT DET LIMIT	SLINO
								ug/L

SURROGATES

98-09-B a, a, a-Trifluorotoluene 132 Q% recovery

INTRMT HACH		(1)See Appendix A for SAMPLE ID Artesia MW-2		INSTRMT 403		SAMPLE ID Artesia MW-2	Page 9 Received: 02/10/89
ANALYTE RESULT DET LIMIT Turbidity261.0	Date & Time Collected 02/09/89 VERIFIED	dix A for glossa	ANALYTE RESULT DET LIMIT Mercury ND 0.0002	ANALYZED 02/22/89	VERIFIED	FRACTION 021 TEST CODE HG C	RAS - Austin REPORT Results by Sample
UNITS NTU	Category	NAME Turbidity		UNITS ug/ml	RHH	NAME Mercury, cold vapor	Work Order # 89-02-119

Page 10 Received: 02/10/89

SAMPLE ID Artesia MW-2

Austin

RAS

REPORT

Work Order # 89-02-119

FRACTION 02J TEST CODE XYLENE Date & Time Collected 02/09/89 Results by Sample

NAME Xulenes, EPA 602

Category

VERIFIED

ANALYST INSTRMT SB

INJECTD 02/13/89

FILE #

UNITS

J/bn

SURROGATES

a, a, a-Trifluorotoluene

144 G% recovery

RESULT DET LIMIT 6.6 0.20

1330-20-7 CAS #

COMPOUND Total xylenes

8-80-86

Page 11 Received: 02/10/89

SAMPLE ID reagent blank

P

RAS

REPORT

Work Order # 89-02-119

FRACTION 03A TEST CODE EPA602 NAME EPA method 602
Date & Time Collected not specified Category Austin REPO

			A.A.	VERIFIED CL
NALYST JB	INJECTED	ED <u>02/13/89</u> FILE # _		UNITS ug/L
	CAS#	COMPOUND	RESULT D	DET LIMIT
	71-43-2	Benzene	ND	0. 20
	108-88-3	Toluene	ND	0. 20
	100-41-4	Ethylbenzene	ND	0. 30
	108-90-7	Chlorobenzene-A	NO	0. 30
	106-45-7	1,4-Dichlorobenzene	ND ND	0.30
	541-73-i	1,3-Dichlorobenzene	NO	0.40
	95-50-1	1,2-Dichlorobenzene	N	0.40

SURROGATES

98-08-8 a, a, a-Trifluorotoluene N/A% recovery

Page 12

SAMPLE ID reagent blank Received: 02/10/89

> RAS Austin

Results by Sample REPORT

Work Order # 89-02-119

FRACTION <u>O3A</u> TEST CODE XYLENE NAME Xylenes, EPA 602

Date & Time Collected not specified Category Category

VERIFIED 5

ANALYST INSTRMT S B

INJECTD 02/13/89

FILE #

SLING Ug/L

RESULT DET LIMIT 0. 20

1330-20-7 C4S #

Total xylenes

COMPOUND

8-80-8

SURROGATES

a, a, a-Trifluorotoluene N/A% recovery

Page 13 Received: 02/10/89

RAS Austin

REPORT

Work Order # 89-02-119

Test Methodology

TEST CODE ALPHA NAME Gross alpha radiation

confidence level. expressed as: The value in parentheses is a + or - one sigma value. value (+ or -1 sigma). One sigma = one standard deviation, Results are thus 789

TEST CODE BETA NAME Gross beta radiation

extenses essential estates extenses extenses essential e The value in parentheses is a + or - one sigma value. One sigma = one standard deviation, 68% Results are thus

TEST CODE RA 226 NAME Radium 226

expressed as: 68% confidence level. The value in parenthases is a + or - one sigma value. value (+ or - one sigma). One sigma = one standard deviation, Results are thus

affed Received: 02/10/89

REPORT Phillips Petroleum

5

Radian

Austin

RAS

REPORT

Work Order # 89-02-120

03/09/89 16: 32: 02

PREPARED <u>Radian Analytical Services</u>
BY <u>8501 Mo-pac 81</u> PHONE ATTEN 512-454-4797 PO Box 201088 <u>Austin, TX 78720-1088</u>

CERTIFIED B

CONTACT BENDELE

INVOICE P. O. # TYPE under separate cover

FACILITY

Odessa.

WORK ID

Artesia

TAKEN

TRANS

UPS

COMPANY

CLIENT

PHILLIPS P

SAMPLES

Phillips Petroleum

ATTEN

Linda Bendele

Austin

SAMPLE IDENTIFICATION

<u>-</u> 의에의의의 Artesia MW-3 reagent blank Artesia MW-4 duplicate Artesia MW-4 trip blank

COLI ALPHA BA E AS G S S S FE E EPA602 BETA 066010 063020 7 Sodium, Specific conductance Barium, Silver, Mercuru, cold yapor EPA method 602 <u>Gross beta radiation</u> Yanganese, Cadmium, ICPES Arsenic, <u>Gross alpha radiation</u> <u> [otal coliform</u> hloride, itrate, colorimetric luoride, igestion, igestion, method hromium, graphite AA ICPES ICPES **ICPES** ICPES graphite AA 7 5 ICPES method ICPES 601C 3020

TEST CODES and NAMES used on this report

PHEN
RA 226
SE G
SO4 IC
TOC TURB XOI XYLENE Radium 226 Sulfate, Selenium, graphite AA Total phenolics Xulenes, Total organic <u>fotal organic</u> urbiditu EPA 602 halides carbon

Page 2 Received: 02/10/89

RAS - Austin REPORT Results By Test

Work Order # 89-02-120

Artesia MW-4	SAMPLE 1 Artesia MW-3	Artesia MW-4 du	SAMPLE I	Artesia MW-3 02 Artesia MW-4	SAMPLE I
02/24/89	Test: <u>D36010</u> date complete 02/24/89	<0.005 <0.005	Test: CD E	<0.03 <0.03	Test: AC E
<0.04	Test: <u>FE E ug/m1</u> <0.04	160 250	Test: CL IC	22.7(2.4) pci/L 16.8(2.1) pci/L	Test: ALPHA
2 1	Test: F IC	20	Test:COLI T	0. 079 0. 013	Test: AS G
2500 2400 2300	Test: MHD	<0.03 <0.03 <0.03	Test: CR E	<0.01 <0.01	Test: BA E
<0.01	Test: MN E	02/23/89	Test: DG3020	21. 6(2.3) pci/L 14. 4(2.3)	Test: BETA

02 85 0.35 (0.001	SAMPLE Test: NO3 Test: PB G Sample Id	03 02/24/89	Page 3 Received: 02/10/89 Results By Test
6. 82 6. 84 01 6. 75 <0. 005	Test:	2400 2300 2300	Work Order # 89-02-120 Continued From Above IC Test: MHO Test: MN E

Page 4 Received: 02/10/89 RAS Austin REPORT
 Results By Test

Work Order # 89-02-120 Continued From Above

Sample Id Sample Id On 01 0.	Test: RA 226 0.48(.04) pci/L	Test: <u>SE G</u>	Test: <u>SO4 IC</u> mq/L as <u>SO4</u> 670	6. 99 Test: TOC 22 25
<u> </u>	.48(.04) pci/L	<0.004	670	22
				24
				23
82	0.40(.04)	<0.004	580	9
Artesia MW-4	bC1/L			10
			·	12
				11

Page 5 Received: 02/10/89

SAMPLE ID Artesia NW-3

RAS Austin REPO

REPORT

Work Order # 89-02-120

FRACTION 01J TEST CODE EPA602 NAME EPA method 602

Date & Time Collected 02/09/89 Category

VERIFIED

ANALYST JB	INJECT	INJECTED 02/13/89		UNITS Ug/L
	C + 0 +	COMPOUND	RESULT DE	DET LIMIT
	71-43-2	Benzene	Ü.N	0. 20
	108-88-3	Toluene	div	0. 20
	100-41-4	Ethylbenzene	0 8 *	0. 30
	108-90-7	Chlorobenzene-A	NO.	0, 30
	106-45-7	1,4-Dichlorobenzene	ON	0. 30
	541-73-1	1,3-Dichlorobenzene	UN	0.40
	95-50-1	1,2-Dichlorobenzene	ND	0.40

SURROGATES

8-80-8 a, a, a-Trifluorotoluene 113% recovery

	SAMPLE ID Artesia MW-3	Received: 02/10/89	Page 6
			RAS
Date & Time Collected	FRACTION 011	Results by	- Austin
	TEST CODE HG C	by Sample	REPORT
Category	NAME Mercury, cold vapor		Work Order # 89-02-120

SAMPLE ID <u>Artesia NW-3</u>	(1)See App			INSTRMT 403	
<u></u>	endix A fo	Mercury	ANALYTE	Anu	
FRACT Date	r glossary of	D	RESULT	AMALYZED <u>02/22/89</u>	
FRACTION <u>OIA</u> TEST CODE TURB Date & Time Collected <u>O2/09/89</u>	(1)See Appendix A for glossary of report and data fl	0.0002	DET LIMIT	89	
)E TURB 2/09/89	flag definitions.			_ SLINO	VERIFIED R
NAME Turbidity Category				uq/ml	RHH

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<u> _</u> S

н<u>асн</u>

A: ALYZED 02/10/89

CNITS NTO

ANALYTE RESULT DET LIMIT

Turbidity

46

1.0

Page 7 SAMPLE ID Artesia Mi-3 Received: 02/10/89

> RAS Austin

REPOST

Work Order # 89-02-120

FRACTION O1J TEST CODE XYLENE Date & Time Collected 02/09/89 Results by Sample

NAME Xulenes, EPA 602

Category

VERIFIED

ANALYST INSTRMT B

INJECTD 02/13/89

FILE #

UNITS **UQ/L**

8-80-85

Total xylenes COMPOUND

1330-20-7 CAS #

RESULT DET LIMIT

a, a, a-Trifluorotoluene 113% recovery

SURROGATES

Page 8 Received: 02/10/89

SAMPLE ID Artesia MW-4

RAS

REPORT

Work Order # 89-02-120

Austin REPO

FRACTION <u>02J</u> TEST CODE <u>EPA602</u>

Date & Time Collected <u>02/09/89</u> NAME EPA method 602 Category

ANALYST 541-73-1 108-86-2 106-46-7 108-90-7 100-41-4 95-50-1 71-43-2 45YO INJECTED 02/13/89 1,2-Dichlorobenzene 1,3-Dichlorobenzene 1,4-Dichlorobenzene Chlorobenzene-A Ethylbenzene COMPOUND Benzene Toluene RESULT DET LIMIT 0.6% 0.5% 5 VERIFIED 0. 20 0.20 0.40 0.40 0.30 0. 30 0.30 STINO 5

SURROGATES

8-80-86 a, a, a-Trifluorotoluene 116% recovery

SAMPLE ID Artesia MW-4 Page 9 Received: 02/10/89 RAS FRACTION <u>021</u> TEST CODE HG C Date & Time Collected <u>02/09/89</u> Austin Results by Sample REPORT NAME Mercury, cold vapor Work Order # 89-02-120 Category

VERIFIED RHH

ANALYST KCP

ANALYZED 02/22/89

UNITS ug/ml

ANALYTE RESULT

Mercury

R

DET LIMIT

0.0002

(1)See Appendix A for glossary of report and data flag definitions

SAMPLE ID Artesia NW-4

FRACTION <u>O2A</u> TEST CODE TURB
Date & Time Collected <u>02/09/89</u>

NAME Turbidity

Category

TRMT HACH

AHALYZED 02/10/89

.

UNITS

VERIFIED

ANALYTE RESULT DET LIMIT

Turbidity 28 1

(1)See Appendix A for glossary of report and data flag definitions

· ** ** • · ·

Page 10 Received: 02/10/89

SAMPLE ID Artesia MW-4

RAS Austin

REPORT

Work Order # 89-02-120

NAME Xulenes, EPA 602

Category

FRACTION 021 TEST CODE XYLENE Date & Time Collected 02/09/89 Results by Sample

VERIFIED CF

ANALYST INTRMT B

INJECTD 02/13/89

FILE #

UNITS ug/L

1330-20-7 CAS # Total xylenes COMPOUND RESULT DET LIMIT

SURROGATES

a, a, a-Trifluorotoluene 116% recovery

8-80-86

Page 11 Received: 02/10/89

RAS Austin REPO

REPORT

Work Order # 89-02-120

SAMPLE ID Artesia MW-4 duplicate FRACTION 03B TEST CODE EPA602
Date & Time Collected 02/09/89 NAME EPA method 602 Category

					ERIFIEDC	CF.
NALYST	e B	INJECT	FILE #		ONITS	ug/L
`		CAS#	COMPOUND	RESULT	DET LIMIT	
		71-43-2	Benzene	5.6	0.20	
		108-88-3	Toluene	ND	0. 20	
		100-41-4	Ethylbenzene	ON	0.30	
		108-90-7	Chlorobenzene-A	NO	0. 30	
		106-45-7	1,4-Dichlorobenzene	ND	0. 30	
		541-73-1	1,3-Dichlorobenzene	ND	0.40	
		95-50-1	1,2-Dichlorobenzene	NO	0.40	

SURROGATES

8-80-86 a, a, a-Trifluorotoluene 100% recovery

Page 12 Received: 02/10/89

SAMPLE ID Artesia MW-4 duplicate

Austin

REPORT

Work Order # 89-02-120

FRACTION 03B TEST CODE XYLENE NAME Xylenes, EPA 602

Date & Time Collected 02/09/89

Category Results by Sample

Category

VERIFIED 2

ANALYST INSTRMT B

CAS # 1330-20-7

COMPOUND Total xylenes

8-80-85

INJECTD 02/13/89

FILE #

SLING

SURROGATES

RESULT DET LIMIT

(1)See Appendix A for glossary of report and data flag definitions.

a, a, a-Trifluorotoluene

108% recovery

Page 13 Received: 02/10/89

SAMPLE ID trip blank

RAS Austin REPO

REPORT

Work Order # 89-02-120

FRACTION 04A TEST CODE EPA602 NAME EPA method 602

Date & Time Collected not specified Category Category

						ANALYST JB	
106-46-7	108-90-7	100-4:-4	108-88-3	71-43-2	CAS#	IHJECTE	÷
1,4-Dichlorobenzene	Chlorobenzene-A	Ethylbenzene	Toluene	Benzene	COMPOUND	FILE # PILE #	
ND	GN	מו	div	ND	RESULT DET LIMIT		VE
0.30	0. 30	0. 30	0.20	0. 20	ET LIMIT	UNITSuq/L	VERIFIEDCL

SURROGATES

541-73-1

1,3-Dichlorobenzene

1,2-Dichlorobenzene

S

0.40

0.40

95-50-1

8-98-8 a, a, a-Trifluorotoluene 110% recovery

Page 14 Received: 02/10/89

SAMPLE ID trip blank

Results by Sample REPORT

RAS

Austin

Work Order # 89-02-120

FRACTION <u>04A</u> TEST CODE <u>XYLENE</u> NAME <u>Xylenes</u>, <u>EPA 602</u>

Date & Time Collected <u>not specified</u> Category Category

VERIFIED P

ANALYST I TRMT 旧

INJECTD 02/13/89

FILE #

UNITS

CAS # 1330-20-7

Total xylenes

COMPOUND

RESULT DET LIMIT

98-08-8

SURROGATES

a, a, a-Trifluorotoluene 110% recovery



Page 15 Received: 02/10/89

SAMPLE ID reagent blank

Austin REPO

RAS

REPORT

Work Order # 89-02-120

FRACTION <u>05A</u> TEST CODE <u>EPA602</u> Note & Time Collected <u>not specified</u> TEST CODE EPA602 NAME EPA method 602 Category

ANALYST INGTRMT 108-90-7 100-41-4 108-88-3 71-43-2 CASH INJECTED 02/13/89 Chlorobenzene-A Ethylbenzene COMPOUND Benzene FILE # Toluene RESULT DET LIMIT **VERIFIED** 0.30 0.20 0.20 0.30 UNITS 2

SURROGATES

541-73-1

1,3-Dichlorobenzene

1,2-Dichlorobenzene

8

0.40

K

0.40

0.30

1,4-Dichlorobenzene

95-50-1

106-45-7

98-08-E a, a, a-Trifluorotoluene N/A% recovery

Page 16 Received: 02/10/89 SAMPLE ID reagent blank

RAS

REPORT

Work Order # 89-02-120

Austin REPO

FRACTION 05A TEST CODE XYLENE NAME Xylenes, EPA 602

Date & Time Collected not specified Category

VERIFIED

ANALYST B

INJECTD 02/13/89

FILE #

UNITS

ug/L

8-80-86

Total xylenes SURROGATES

COMPOUND

1330-20-7

RESULT DET LIMIT

a, a, a-Trifluorotoluene N/A% recovery

Received: 02/10/89

RAS Austin

REPORT

Work Order # 89-02-120

Test Methodology

TEST CODE ALPHA NAME Gross alpha radiation

expressed as: confidence level. The value in parentheses is a + or - one sigma value. value (+ or - 1 sigma). One sigma = one standard deviation, 68% Results are thus

TEST CODE BETA NAME Gross beta radiation

confidence level. value in parentheses is a + or - one sigma value. ¹essed as: value (+ or − 1 sigma). One sigma = one standard deviation, 68% Results are thus

TEST CODE RA 226 NAME Radium 226

expressed as: 68% confidence level. The value in parentheses is a + or - one sigma value. value (+ or - one sigma). One sigma = one standard deviation, Results are thus

-MW-1
MW=2
4 MM-3
MW-4
02-122

RESULTS	IN	ug/L
---------	----	------

CAS #	COMPOUND	01B	02B	03B	04B
58-89-9	gamma-BHC,(Lindane)	0.16*	<0.038	<0.019	<0.039
72-20-8	Endrin	<0.038	<0.038	<0.019	<0.039
8001-35-2	Toxaphene	<1.9	<1.9	<0.95	<2.0
72-43-5	Methoxychlor	<0.19	<0.19	<0.095	<0.20

SURROGATE RECOVERIES	(results in % recovery)				
Dibutylchlorendate	110	111	102	112	
2,4,5,6-Tetrachloro-m-xylene	121	113	96	139	

(1) See Appendix A glossary of report and data flag definitions.

Pertreiles 2/89 4th Quarter

RADIAN

05B REAGENT BLANK

Client: Phillips Petroleum

Radian Bl.1 Austin

EPA METHOD 8080

Lab No: 89-02-122

RESULTS IN ug/L

CAS #	COMPOUND	05B
58-89-9	gamma-BHC, (Lindane)	<0.002
72-20-8	Endrin	<0.002
8001-35-2	Toxaphene	<0.10
72-43-5	Methoxychlor	<0.010

SURROGATE RECOVERIES (results in % recovery)

Dibutylchlorendate 116 2,4,5,6-Tetrachloro-m-xylene 83

(1) See Appendix A glossary of report and data flag definitions.

Page 1

Client: Radian

B1.1

Austin

Ola MW-1 ARTESIA MW-1 02A MW-2 ARTESIA MW-2 03A MW-3 ARTESIA MW-3

04A MW-4 ARTESIA MW-4

EPA METHOD 8150

Lab No: 89-02-122

RESULTS IN ug/L

CAS ‡	COMPOUND	OlA	02A	03A	04A
94-75-7	2,4-D	5.6 C	<1.4	<1.4	<1.4
93-72-1	2,4,5-TP (Silvex)	<0.14	<0.42	<0.42	<0.43

SURROGATE RECOVERIES (results in % recovery)

2,4-Dichlorophenyl acetic acid 122

166QC

132

108

NOTES AND DEFINITIONS FOR THIS REPORT.

QC = OUTSIDE CONTROL LIMITS.

* = LESS THAN 5 TIMES THE DETECTION LIMIT.

C = RESULT CONFIRMED BY SECOND COLUMN ANALYSIS.

ND = NOT DETECTED AT DETECTION LIMIT.

NA = NOT ANALYZED.

 $N\setminus A = NOT AVAILABLE.$

NS = NOT SPIKED.

Page 2

05A REAGENT BLANK

Client: Radian

B1.1

Austin

EPA METHOD 8150

Lab No: 89-02-122

RESULTS IN ug/L

CAS # COMPOUND 05A

94-75-7 2,4-D

<0.50

93-72-1 2,4,5-TP (Silvex)

<0.15

SURROGATE RECOVERIES (results in % recovery)

2,4-Dichlorophenyl acetic acid 102

NOTES AND DEFINITIONS FOR THIS REPORT.

QC = OUTSIDE CONTROL LIMITS.

* = LESS THAN 5 TIMES THE DETECTION LIMIT.

C = RESULT CONFIRMED BY SECOND COLUMN ANALYSIS.

ND = NOT DETECTED AT DETECTION LIMIT.

NA = NOT ANALYZED.

N/A = NOT AVAILABLE.

NS = NOT SPIKED.

Page 2

06A RECOVERY CHECK

Client: Radian

Bl.1

Austin

EPA METHOD 8150

Lab No: 89-02-122

RESULTS IN %

CAS # COMPOUND 06A

94-75-7 2,4-D 101

93-72-1 2,4,5-TP (Silvex) 99

SURROGATE RECOVERIES (results in % recovery)

2,4-Dichlorophenyl acetic acid 97

NOTES AND DEFINITIONS FOR THIS REPORT.

QC = OUTSIDE CONTROL LIMITS.

* = LESS THAN 5 TIMES THE DETECTION LIMIT.

C = RESULT CONFIRMED BY SECOND COLUMN ANALYSIS.

ND = NOT DETECTED AT DETECTION LIMIT.

NA = NOT ANALYZED.

N A = NOT AVAILABLE.

NS = NOT SPIKED.

REPORT FLAGS

- R Indicates that the matrix spike recovery for this analysis is not within acceptable limits indicating an interferent present.
- + Indicates that the RPD between the percent recoveries of the MS and MSD is not within acceptable limits.
- @ Indicates that the analytical spike recovery for this analysis is not within acceptable limits indicating an interferent present.
- B Sample result was greater than four times the spike added concentration, therefore the spike recovery should not be considered.
- A Indicates the RPD of the duplicate analysis is not within acceptable limits.
- Indicates that the value obtained is less than five times the detection limit. Potential error for such low values range between 50 % and 100 %.
- E Indicates that the reported value is estimated due to the presence of an interference.
- s Indicates that a value is determined by Method of Standard Addition.
- Q Daily EPA QC recovery outside 95% confidence limit.
- ** Surrogate recovery is not within acceptable limits. This indicates that a possible interference is present.

Work Order # P8-05-56

7 7		~		
00000000000000000000000000000000000000	01 SAMPLE 02 880 512 03 880 5112 04 880 5112	WORK ID TAKEN TYPE	ATTEN CLIENT COMPANY FACILITY	REPORT TO
0952		Geosciences W S Dubyk Aqueous Agueous BB-0190-700 under separate cover	W.S. Dubyk GEOSCIENCE Geoscience Consultants, Ltd. 500 Copper NW Albequerque, NM 87102	Mike Selke Geoscience Consultants, Ltd. Albequerque, NM 87102
	OC TMO	1 1 1 1 1 1 1 1 1 1	g g	72

PREPARED Radian Analutical BY BIdg. 900 Perimet eviously Reported on 06/13/88 ATTEN PHDNE 919-481-0212

CONTACT M DAY

TEST CODES and NAMES used on this report

Page 2 • comp Réceived: 05/12/88

RAS Perimeter REPORT

Work Order # P8-05€36

TEST CODE default units EXT GC date completed	TEST CODE default units date completed	TEST CODE EXT GC date completed
Sample 11 (entered units) 05/20/88	Sample 06 (entered units) 05/17/88	Sample 01 (entered units) 05/18/88
Sample 12 (entered units) 05/20/88	Sample 07 (entered units) 05/17/88	Sample 02 (entered units) 05/18/88
Sample 13 (entered units) 05/18/88	Sample 08 (entered units) 05/17/88	Sample 03 (entered units) 05/17/88
Sample 15 (entered units) 05/17/88	Sample 09 (entered units) 05/18/88	Sample 04 (entered units) 05/17/88
	Sample 10 (entered units) 05/20/88	Sample 05 (entered units) 05/17/88

Page 3 Page 3 Received: 05/12/88

RADIA

RAS Perimeter

Results By Test

Work Order # P8-05-55

SAMPLE Sample Id Sample Id 01 8805121240 02	Test: <u>EXT GC</u> date completed 05/18/88	
8805121042 03	05/17/88	
8805111656 04	05/17/88	
8805111812 05	05/17/88	
8805110958 06	05/17/88	
8805110852 07	05/17/88	
8805110922 08	05/17/88	
8805110938 09	05/18/88	
8805121844 10	05/20/88	
	05/20/88	
12	05/20/88	
عبيو	05/18/88	
Method Blank #2 15	05/17/88	
Method Blank #1 :		

RAS Perimeter Results by Sample

REPORT

Work Order # P8-05. 26

SAMPLE ID 8805121240 SAMPLE # 01 FRACTIONS: A
Date & Time Collected 05/12/88

EXT GC 05/18/88

Category

Page 6 Comp Received: 05/12/88

RAS Perimeter REPORT

Work Order # P8-05 36

Category

SAMPLE ID 8805121042 SAMPLE # 02 FRACTIONS: A
Date % Time Collected 05/12/88

EXT GC 05/18/88

RAS Perimeter

REPORT

Work Order # P8-05-036

Category

SAMPLE ID 8805111656 SAMPLE # 03 FRACTIONS: A
Date & Time Collected 05/11/88

EXT_GC 05/17/88

Results by Sample

Work Order # P8-05-36

SAMPLE ID 8805111812

RAS Perimeter REPORT

Results by Sample

SAMPLE # 04 FRACTIONS: A
Date % Time Collected 05/11/88

Category

EXT GC 05/17/88

Work Order # P8-05-036

SAMPLE ID 8805110958

EXT GC 05/17/88

RAS Perimeter
Results by Sample

SAMPLE # 05 FRACTIONS: A
Date & Time Collected 05/11/88

Category

Page 14 Comme Received: 05/12/88

RAS Perimeter REPORT
Results by Sample

Work Order # P8-05-536

Category

SAMPLE ID 8805110852 SAMPLE # 06 FRACTIONS: A
Date & Time Collected 05/11/88

EXT_GC_05/17/88

Page 16 Comp Received: 05/12/88

RAS Perimeter
Results by Sample

Work Order # P8-05-036

Category

SAMPLE ID 8805110922 SAMPLE # 07 FRACTIONS: A
Date & Time Collected 05/11/88

EXT GC 05/17/88

RAS Perimeter Results by Sample

Work Order # P8-05 36

SAMPLE ID 8805110938

EXT GC 05/17/88

SAMPLE # 08 FRACTIONS: A
Date & Time Collected 05/11/88

Category

RAS Perimeter
Results by Sample

Work Order # P8-05 Vd6

SAMPLE ID 8805121844

EXT GC 05/18/88

SAMPLE # 09 FRACTIONS: A Date % Time Collected 05/12/88

Category

RAS Perimeter

ter Results by Sample

Work Order # P8-05 36

Category

SAMPLE ID 8805130903 EXT_GC_05/20/88 SAMPLE # 10 FRACTIONS: A
Date & Time Collected 05/13/88

RAS Perimeter ע אברע Results by Sample

REPORT

Work Order # P8-0, €036

SAMPLE ID 8805130949 EXT GC 05/20/88

SAMPLE # 11 FRACTIONS: A
Date & Time Collected 05/13/88

Category

RAS Perimeter

Work Order # P8-05-36

Category

SAMPLE ID 8805131420

EXT_GC_05/20/88

SAMPLE # 12 FRACTIONS: A
Date & Time Collected 05/13/88 Results by Sample

RAS Perimeter REPORT

Work Order # P8-05-36

Category

SAMPLE ID Method Blank #2 SAMPLE # 13 FRACTIONS: A

Date & Time Collected not specified

EXT GC 05/18/88

RAS Perimeter

REPORT

Work Order # PB-05-36

SAMPLE ID Method Blank #2 Results by Sample

FRACTION 13A TEST CODE 509B No Date & Time Collected not specified 系統 GC of Herbicides Category

DRGANICS ANALYSIS DATA SHEET HERBICIDES

INSTRMT BLACKLEY 93-76-5 94-75-7 93-72-1 CAS # EXTRCTD 05/18/88 INJECTD 05/26/88 2,4,5-TP (Silvex) COMPOUND 2,4,5-T 2,4-D FILE # RESULT VERIFIED DET. LIMIT 0 10 0.50 0.10 팢 SLINO

NOTES ITIONS FOR THIS REPORT.

— detection limit.
detected at specified detection
required for analysis. limit

W

minimum detection limit

RAS Perimeter
Results by Sample

Work Order # P8-05-05

Category

SAMPLE ID Method Blank SAMPLE # 15 FRACTIONS: A
Date & Time Collected not specified

EXT GC 05/17/88

RATION RAS P

RAS Perimeter

REPORT

Work Order # P8-05-536

SAMPLE ID Method Blank #1

INSTRMT BLACKLEY

Results by Sample

FRACTION 15A TEST CODE 509B NAME Date & Time Collected not specified

8

of Herbicides

Category

DRGANICS ANALYSIS DATA SHEET HERBICIDES

93-76-5 93-72-1 94-75-7 EXTRCTD 05/17/88 INJECTD 05/24/88 CAS # (Silvex) COMPOUND 2, 4, 5-1 2,4-D FILE # RESULT VERIFIED 급 S DET. 0.10 0.50 O LIMIT SLINO

NOTES ITIONS FOR THIS REPORT.

= detection limit.
detected at specified detection limit.
required for analysis.
und peak saturated. minimum detection limit.

RAS Perimeter REPORT

NonReported Work

FRACTION AND TEST CODES FOR WORK NOT REPORTED ELSEWHERE 16A : HOLD

Work Order # P8-05-06

REPORT

Mike Selke

5

Geoscience Consultants,

Albequerque,

Z

87102

RAS Perimeter

Work Order # P8-05-035

Received: 05/12/88

06/13/88 16:14:44

PREPARED Radian Analytical Services Bldg. 900 Perimeter Park Morrisville. NC 27560

PHONE ATTEN 919-481-0212

CERT

FIED

벌

CONTACT M DAY

Matrix interference

FACILITY

500 Capper NW

Geoscience Consultants,

Albequerque,

3

87102

COMPANY

CLIENT

GEDSCIENCE

SAMPLES

ATTEN

Mike Selke

INVOICE TYPE TRANS TAKEN 88-0190-700 PHILLIPS under separate Aqueous Fed Ex (see file Dubuk COVET for #'s

그것의 4岁 4 2 2 2 2 3 8805121239 8805110920 8805110851 8805110955 8805111811/ 8805121041 3805111655 205110936 ARTIGUA ARTISSIA 9RTGS/B CUSK *lusk* SS X DW. m4-2 1 × 100 MKmW-MH-3 MW-E-MW

Method Blank

3805131420 3805130948 3805130902

DW. DW-MM - 2 MW-3

505121843

TEE

rip Blank

EST CODES and NAMES used on this report

BOBO EXT GC Pesticides/PCBs xtraction for GC

Received: 05/12/88

RAS Perimeter
Results By Test

Work Order # P8-05-035

EXT GC	TEST CODE	EXT GC	TEST CODE	EXT GC	TEST CODE
05/16/88	Sample 11	05/16/88	Sample 06	05/16/88	Sample 01
05/16/88	Sample 12 (entered units)	05/16/88	Sample 07	05/16/88	Sample 02 (entered units)
05/16/88	Sample 16 (entered units)	05/16/88	(entered units)	05/16/88	Sample 03 (entered units)
		05/16/88	Sample 09	05/16/88	Sample 04 (entered units)
		05/16/88	Sample 10 (entered units)	05/16/88	Sample 05 (entered units)

Work
Order
:# :
P8-05-055

'age 3 () = 8	RAS Perimeter RAS Perimeter R	Results By Test	Work Order # P8-05-055
SAMPLE Id	Test: EXT GC		
01			
8805121239 02	05/16/88		
BC	05/16/88		
04	05/16/88		
9805111811	05/16/88		
905110755	05/16/88		
8803110831	05/16/88		
80 277000	05/16/88		
09	05/16/88		
10	05/16/88	,	
200130702	05/16/88		
8805130748 12	05/16/88		
16	05/16/88		

Page 4 Teceived: 05/12/88

RAS Perimeter

Results by Sample

Work Order # P8-05-035

Category

SAMPLE ID 8805121239 SAMPLE # 01 FRACTIONS: A
Date & Time Collected 05/12/88

EXT GC 05/16/88

Page 7 Teceived: 05/12/88

RAS Perimeter

Results by Sample

Work Order # P8-05-055

SAMPLE ID 8805121041

EXT GC 05/16/88

SAMPLE # 02 FRACTIONS: A
Date & Time Collected 05/12/88

Category

Page 10 25/12/88

RAS Perimeter
Results by Sample

Work Order # P8-05-035

Category

SAMPLE ID 8805111655 EXT GC 05/16/88 SAMPLE # 03 FRACTIONS: A Date & Time Collected 05/11/88

Page 13 ■ Complex Received: 05/12/88

Work Order # P8-05-033

SAMPLE ID 8805111811

EXT GC 05/16/88

RAS Perimeter

Results by Sample

SAMPLE # 04 FRACTIONS: A
Date & Time Collected 05/11/88

Category

Page 16 Teceived: 05/12/88

SAMPLE ID 8805110955

EXT GC 05/16/88

RAS Perimeter
Results by Sample

Work Order # P8-05-035

Category

SAMPLE # 05 FRACTIONS: A
Date & Time Collected 05/11/88

RAS Perimeter REPORT

Work Order # P8-05-035

Category

SAMPLE ID 8805110851 SAMPLE # 06 FRACTIONS: A
Date & Time Collected 05/11/88

EXT GC 05/16/88

RAS Perimeter
Results by Sample

Work Order # P8-05-035

SAMPLE ID 8805110920	SAMPLE # 07 FRACTIONS: A	
	Date & Time Collected 05/11/88	Category
2		
EX 67 03/16/88	5	
date completed		

Page 25 ■ ***
Received: 05/12/88

SAMPLE ID 8805110936

EXT GC 05/16/88

Work Order # P8-05-035

Category

RAS Perimeter
Results by Sample

SAMPLE # 08 FRACTIONS: A
Date & Time Collected 05/11/88

Work Order # P8-05-05-

Category

SAMPLE ID 8805121843

EXT GC 05/16/88

RAS Perimeter
Results by Sample

SAMPLE # 09 FRACTIONS: A Date & Time Collected 05/12/88

REPORT

RAS Perimeter Results by Sample

Work Order # P8-05-035

Category

SAMPLE ID 8805130902 SAMPLE # 10 FRACTIONS: A
Date & Time Collected 05/13/88

EXT GC 05/16/88

Page 34 Pereived: 05/12/88

EXT GC 05/16/88

RAS Perimeter

Results by Sample

Work Order # P8-05-035

SAMPLE ID 8805130948

SAMPLE # 11 FRACTIONS: A
Date & Time Collected 05/13/88

Category

EPORT

Work Order # P8-05-

SAMPLE ID 8805131420

EXT GC 05/16/88

RAS Perimeter Results by Sample SAMPLE # 12 FRACTIONS: A
Date & Time Collected 05/13/88

Category

Page 40 Teceived: 05/12/88

RAS Perimeter
Results by Sample

Work Order # P8-05

SAMPLE ID Method Blank EXT GC 05/16/88 SAMPLE # 16 FRACTIONS: A Date & Time Collected not specified

Category

RAS Perimeter

REPORT

Work Order # P8-05

SAMPLE ID Method Blank

Results by Sample

PRACTION 16A TEST CODE 8080 No Date & Time Collected not specified NAME Pesticides/PCBs.

Category

DRGANICS ANALYSIS DATA SHEET PESTICIDES by METHOD 8080

VERIFIED

11096-82-5	11097-69-1	12672-29-6	53469-21-9	11141-16-5	11104-28-2	12674-11-2	8001-35-2	57-74-9	1031-07-B	7421-93-4	50-29-3	33213-65-9	72-54-8	72-20-8	60-57-1	72-55-9	959-98-8	1024-57-3	6	319-86-8	76-44-8	319-85-7	58-89-9	319-84-6		CAS #		ANALYST BLACKLEY
PCB-1260	PCB-1254	PCB-1248	PCB-1242	PCB-1232	PCB-1221	PCB-1016	toxaphene	chlordane	endosulfan sulphate	endrin aldehyde	4, 4'-DDT	endosulfan II	4, 4'-ppp	endrin	dieldrin	4, 4 '-DDE	endosulfan I	heptachlor epoxide	aldrin	delta-BHC	heptachlor	beta-BHC	gamma-BHC (lindane)	alpha-BHC		COMPOUND	INJECTD 06/05/88	EXTRCTD <u>05/16/88</u>
NA NA	N/R	273	NA R	•	NR	NR	ND	ND	ND	ND	•	Z	ND NO	•	•		ND	ND	ND	ND	ND	ND	•	ND	, · · · · · · · · · · · · · · · · · · ·	RESULT DE		FILE #
0. 20	0.20	0.10	0. 10	0. 20	0.20	0. 10	0. 50	0.050	0.050	0.020	0.020	0. 030	0.010	0. 010	0.010	0.010	0.010	0.010	0.010	0.010	0.010	0.010	0, 010	0.010		ET LIMIT	UNITS U9/L	* The state of the

'age 42 'eceived: 05/12/88

RAS Perimeter

REPORT

Work Order # P8-05 Continued From Above

NAME Pesticides/PCBs.

Category

AMPLE ID Method Blank

FRACTION 16A TEST CODE 8080 No Date & Time Collected not specified Results by Sample

SURROGATE RECOVERY

COMPOUND

RECOVERY

dibutyl chlorendate 100 %

tetrachlorometaxylene 85 ×

NOTES AND DEFINITIONS FOR THIS REPORT:

ND = not detected at specified detection limit

NR = not required for analysis.

compound peak saturated.

∠ ≡ estimated value less than 3 x minimum detection limit

RAS Perimeter

REPORT

NonReported Work

RACTION AND TEST CODES FOR WORK NOT REPORTED ELSEWHERE

ISA ! HOLD

Work Order # P8-05-

RADIAN ANALYTICAL SERVICES QUALITY CONTROL DATA SUMMARY

Form II

Client	ID	Resocience Consultar	ئە
Workord	ler	88-05-067	

Compiled	Sqc 6-29-88
Submitted	-

INITIAL AND CONTINUING CALIBRATION VERIFICATION

Units _____/ml

			2-11		T			· 0-1		0.	
Parameter	Run Date	True	ial Cali Found		True	Found		uing Cal Found	bratio %R	n Found	% R
Ag	6-13-88		4.78	96	5.00	4.89	98	4.74	95	Ourid	
Ba			4.92	98		5.03	101	5.02	100		
Ca			4.91	98	·	5.00	100	4.91	98		
Cr			4.76	95	V	4.78	96	4.71	ं १५		
Fe		20.0	19.5	98	20.0	19.6	98	19.3	97	,	
Mn		5.00	4.78	96	S 00	4.82	96	4.69.	94		
Na		20.0	20.4	102	20.0	17.9	90	20.9:	105		
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	·			9				.,			
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							114			<u> </u>	
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						1		1			

1.	Control	Limits	for	3R:	ICF	ES	
					AA	F	

RADIAN ANALYTICAL SERVICES QUALITY CONTROL DATA SUMMARY FORM ILL

Client ID	Georgiance Consultants
Workorder	88-05-067

Compiled System Submitted

BLANKS

Units _____

,		<u>initial</u> Calibration	Cont	inuing Call Blank Val		Preparat Blank Va	
Parameter	IDL	Blank Value	11	2	3		2
Ag	0.003	(0.003	(0.003	<0.003		(0.003	
Ba	0.003	10.003	(0.003	(0.003		0.003*	
Cd	600.0	(0.003	40.003	40.003		(0.003	
Cr	0.003	<0.003	40.003	<0.003		< 0.003	
<u>Le</u>	8.003	0.016	10.003	<0.003		0.031	
ma	100.0	40.001	40.001	<0.001		0.002	
Na	0.021	0.036*	40.021	40.021		3.17	
				4			
							
		,					
							
		<u> </u>		1	1		
	1	 		1	†	-	
	†				 		
	†			1	-		
	+	 				#	
			#	 			

^{1.} IDL = Instrument Detection Limit

^{*} Indicates value is less than 5X the IDL.

QUALITY CONTROL DATA SUMMARY

FORM Y

Client H	0	Georgiance Consultanto
Workorde	_	88-05-067

_	
	Compile

Compiled __

Ser 6-29-88

Submitted

Matrix

Blank Spik

SPIKED SAMPLE RECOVERY

Spiking method agreeus

Units welme

arameter	Sample No.	Control Limit	Spiked Sample Result (SSR)	Sample Result(SR)	Spike Added (SA)	D.F.	SR ¹	SR Flag ²
Δg	Blank	75-125	0.856	R	1.00		86	
Ba			0.875				88	
Ca			0.888				89	
Cr			0.856		V		86	
Fe			8.77		10.0	J	88	
Mn			0.861	Į.	1.00		86	
Na			8.92		10.0		89	
	,							
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Indicates value is less than 5% iDL. (IDL=Instrument detection limit)

1. \$R = Percent Recovery = [(SSR - SR)/SA] X 100

2. Refor metrix spike: SR was not within control limit, which may indicate matrix interferences.

B-Sample result was greater than 4 times spike added concentration, therefore spike added concentration is considered insignificant.

B- Blank result is subtracted out from the samples.

Compiled_ Submitted_ 58x 6-25-88

Workorder PA-05-067

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Shamish noted

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		CAL	CAL IBRATION				1							
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		7.48	8.00	94										
		7.60	8.00	95										
ग ू	69-1-8	+ · o+	4.00	101	N 02	1.74	1.75	-	B) 02	2.65	121	1.00	91	
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•		3 . 88	4.00	97									_	
204	1-88 ·	39.2	40. o	98	A) 02	9.4	46	0	NO2	55.8	46	10.0	98	
		39.3	40.0	98										
		39.1	40.0	98										
֡														

 $RPD = [(1S-D1)/((S+D)/2)] \times 100$ NC = Not catculable due to a value RFD = Relative Percent Difference less than five times the IDL

* " Value is less than five times SPIKE JR = [(SSR-SR)/SA] × 100 IDL = Instrument Detection Limit the instrument detection limit

A = Analytical P = Predigestion

SSR = Spiked Sample Result SA = Spiked Added SR = Sample Result

Submitted Sex 6-21-88

Norkorder 88-05-067

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Otherwise noted

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P) .002														
9 <0:002							1,64		,	104	0.100	a. 104		
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₹) <0.0∞1										2	0.0042 0.0040	2400.0	elm C	001
60.000								, , ,		110	0400.0 PHO0.0	5.0044 PHO0 '9	5-23-88	Ha
BLANKS	188	AS	SI2	SSR	SAMP!	RPD	Tand		SAMP!	\$R	FOUND TRUE SR	COUND	DATE	PARAMETER
		₹	RECOVE RY	SPIKE			AMALYSIS	DUPLICATE A	DÚPL	TOS.	CALIBRATION ST	VERLE I	AMLYSIS	
	I V							+						

RPD = [(IS-DI)/((S+D)/2)] × 100
RPD = Relative Percent Difference
NC = Not calculable due to a value
less than five times the IDL

SPIKE \$R = [(SSR-SR)/SA] × 100

* Value is less than five times
the instrument detection limit
IDL = instrument Detection Limit

A = Analytical
P = Predigestion
SSR = Spiked Sample Result
SR = Sample Result
SA = Spiked Added

Submitted Syk 6-29-88

Workorder 88-05-067

Citent Seacence Congultonto

Otherwise notes

101 = 0.003 Ryported Phenolica 101=0.005 Dr 20.01 m Tox PARAMETER Se 1 ì 5-18-88 oH unit 89-14-8 88-4-4 5-27-88 5-17-88 AMLYSIS DATE 0.263 949 0.929 158.0 0.276 22.0 24.0 0.054 0.049 0.275 7.00 COMPO 7.00 7.00 MERIEICATION STOS. CALIBRATION 0.050 0.050 IBUE 2.55 0.050 0.250 1.00 0.750 7.00 0.250 55.2 7.00 7.00 B 501 110 100 23 801 100 78 もっ 98 98 9 00 CHNS 9 Ms + mso þ ō DUPLICATE AMILYSIS 68.3 SAMS 7.19 _ _ 66.9 DUP 7.19 801 **E** 4 œ 0 48.64 3 3020 5/16 Blank SAMP 0 111 0.012 DIC 1:1 SPIKE 358 801 RECOVER 67.6 67.6 60.0 60.0 0.020 167R 82 0 Ħ 115 oras (0.003 9 × 6 003 \$0.005 40.003 BLANKS

RPD = [(IS-DI)/((S+D)/2)] × 100
RPD = Relative Percent Difference
NC = Not calculable due to a value
, less than five times the IDL

SPIKE \$R = [(SSR-SR)/SA] x 100

* = Value is less than five times
the instrument detection limit
IDL = instrument Detection Limit

A = Analytical
P = Predigestion
SSR = Spiked Sample Result
SR = Sample Result
SA = Spiked Added

Workorder E	Compiled_	Submitte
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SPIKE															
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RPD = Relative Percent Difference $RPD = [(1S-D1)/((S+D)/2)] \times 100$ NC = Not calculable due to a value less than five times the IDL

the instrument detection limit SPIKE JR = [(SSR-SR)/SA] × 100 IDL = Instrument Detection Limit

A = Analytical
P = Predigestion SA = Spiked Added SR = Sample Result SSR = Spikad Sample Result

26 6-29-88

DAILY QUALITY CONTROL RAS GC LAB

DATE:	5-17-88	SPIKED VALUE (ug/L)	Analyzed Yalus	Recovery	Analyzed Value	\$ Recovery
	INSTRUMENT		D	D	ري	G
TEST METHOD	COMPOUNO		*********	20444444		*********
EPA 601	EPA WP 483 CONC. 2					
	Chloroform	12.0				
	1.2-Dichiorgethane	2.0				
	1.1.1-Trichioroethane	1.4				
^	Carbon Tetrachioride	2.6				1
	Bromodichioromethane	2.0			·	
	Trichlaroethene	2.9				
	Dibromochioromethane	2.6				
	Brometerm	2.9				
	Tetrachioroethene	1.6	·			
EPA 602	EPA - WP 879 CONC.1					
	Benzene	30.7	26.2	85	29.4	96
	Taluene	4.1	4.0	98	4.9	120
	Ethylhenzene	11.5	9.1	79	10.4	90
	P-Xylene	19.1	18.6	97	21.9	115
	M-Xylene	42.6	385	90	42.3	99
	0-Xviene	10.6	10.5	99	12.6	119
EPA 608	Arocior 1242	(ug/g) 58.7				
	Arociar 1260	56.8				i

VOA RESULTS

AB # < >	sta Bunk		
SAMPLE ID			
3/MFLE IU			24222222222222
EPA METHOD 601	DATE: ANALYST: INSTRUMENT:	EPA METHOO 602	DATE:5/17/1 ANALYST:4 INSTRUMENTAL
COMPOUND	CONCENTRATION (ug/L)	COMPOUND	CONCENTRATION (ug/L)
Chloromethane		Benzene	W
Bromomethane		Toluene	
Vinvl Chloride		Ethyl benzene	·
Chloroethane		Chlorobenzene	
Methylene chloride		1.4-Dichlorobenzene	
Trichlorofluorometha	ane	1.3-Dichlorobenzene	
1.1-Dichloroethene		1.2-Dichlorobenzene	
1.1-Dichloroethane		P-Xylene	
Trans-1.2-Dichloroe	thene	M-Xvlene	
Chloroform		0-Xylene	
1.2-Dichloroethane			
1.1.1-Trichloroethan]	
Carbon tetrachloride			
Bromodichloromethane			
1.2-Dichloropropane		SURROGATE RECOVER	RIES:
Trans-1.3-Dichloron	copene	601	
Trichloroethene		Bromochloromethan	
<u>Dibromochloromethan</u>		2-Bromo-1-Chlorop	
1.1.2-Trichloroetha		4-Dichlorobutar	
cis-1.3-Dichloroprop		4-Bromofluorobenz	tene
2-Chloroethylvinyl	ether	4	
Bromoform		602	
1.1.2.2-Tetrachloro		a,a,a,-Trifluorot	roluene
Tetrachioroethylene		4	
Chlorobenzene		4	
1.3-Dichiorobenzene		4	
1.2-Dichlorobenzene		4	
1.4-Dichlorobenzene		4	
			•

YOA RESULTS

COMPOUND CONCENTRATION (ug/L) Chloromethane Bromomethane Toluene Vinyl Chloride Chlorosthane Chlorosthane Chlorosthane Chlorosthane Trichlorofluoromethane 1.3-Dichlorobenzene 1.1-Dichlorosthane Trans-1.2-Dichlorosthane Trans-1.2-Dichlorosthane Trans-1.2-Dichlorosthane Trans-1.3-Dichlorosthane Trans-1.3-Dichlorosthane Trans-1.3-Dichlorosthane Trans-1.3-Dichlorosthane Trans-1.3-Dichloromethane	EPA METHOO 601	DATE: ANALYST: INSTRUMENT:	EPA METHOD 602	DATE: 1768 ANALYST: G INSTRUMENT ()
Richloromethane Toluene Vinyl Chloride Ethyl henzene Chlorobenzene Methylene chloride 1.4-Dichlorobenzene Trichlorofiluoromethane 1.3-Dichlorobenzene 1.1-Dichloroethane 1.2-Dichlorobenzene 1.1-Dichloroethane 1.2-Dichlorobenzene 1.1-Dichloroethane 1.2-Dichlorobenzene 1.1-Dichloroethane M-Xylene Chloroform O-Xylene 1.2-Dichloroethane 1.1-Trichloroethane SURROGATE RECOVERIES: Trans=1.3-Dichloromethane Carbon tetrachloride Bromodichloromethane Carbon tetrachloride Bromodichloromethane Carbon tetrachloropropane 1.1-2-Trichloroethane Carbon tetrachloropropane 1.1-2-Trichloroethane Carbon tetrachloromethane 1.1-2-Trichloroethane Carbon tetrachloropropane 1.1-2-Trichloroethane Carbon tetrachloropropane 1.1-2-Trichloroethane Carbon tetrachloropropane 1.1-2-Trichloroethane Carbon tetrachloroethane 1.1-2-Trichloroethane Carbon tetrachloroethane 1.1-2-Trichloroethane Carbon tetrachloroethylvinyl ether Bromoform Carbon tetrachloroethane Carbon tetrachloroethylvinyl ether Bromoform Carbon tetrachloroethylvinyl ether Carbon tetrachloroethylvinyl eth	COMPOUND		COMPOUND	
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Chlorosthane Mathylane chloride Mathylane chloride Mathylane chloride Mathylane chloride Mathylane chloride Mathylane chloride Mathylane chlorosethane Mathylane chlorosethane Mathylane M				
Mathylene chloride Trichlorofluoromethane 1.3-Dichlorobenzene 1.1-Dichloroethane 1.1-Dichloroethane 1.2-Dichloroethane Trichloroethane 1.2-Dichloroethane 1.2-Dichloroethane 1.1-Trichloroethane 1.1-Trichloroethane 1.2-Dichloromethane 1.3-Dichloromethane 1.4-Dichloromethane 1.4-Dichlorobutane 4-Bromofluorobenzene 1.1-2-Tetrachloroethane 1.1-2-Tetrachloroethane 1.1-2-Tetrachloroethane 1.1-2-Tetrachloroethane 1.1-2-Dichlorobenzene 1.3-Dichlorobenzene 1.3-Dichlorobenzene 1.3-Dichlorobenzene 1.3-Dichlorobenzene		·		
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1.2-DichloropropaneSURROGATE RECOVERIES:Trans-1.3-Dichloropropene601TrichloroetheneBromochloromethaneDibromochloromethane2-Bromo-1-Chloropropane1.1.2-Trichloroethane1,4-Dichlorobutanecis-1.3-Dichloropropene4-Bromofluorobenzene2-Chloroethylvinyl ether602Bromoform6021.1.2.2-Tetrachloroethanea,a,a,-TrifluorotolueneTetrachloroethyleneChlorobenzene1.3-Dichlorobenzene1.2-Dichlorobenzene			+	•
Trans-1.3-Dichloropropene Trichloroethene Dibromochloromethane 1.1.2-Trichloroethane cis-1.3-Dichloropropene 2-Chloroethylvinyl ether Bromoform Tetrachloroethylene Chlorobenzene 1.3-Dichlorobenzene 1.3-Dichlorobenzene 1.2-Dichlorobenzene 1.2-Dichlorobenzene			CURROCATE RECOVERY	Ec.
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1.3-Dichlorobenzene 1.2-Dichlorobenzene			7	
1.2-Dichlorobenzene			1	
	1.4-Dichlorobenzene		3	



REPORT FLAGS

- R Indicates that the matrix spike recovery for this analysis was not within acceptance limits indicating an interferent present.
- + Indicates that the RPD between the MS and MSD is not wihin control limits.
- O Indicates that the analytical spike recovery for this analysis was not within acceptance limits indicating an interferent present.
- B Sample result was greater than four times the spike added concentration, therefore the spike recovery should not be considered.
- ∧ Indicates duplicate analysis is not within control limits.
- * Indicates that the value obtained was less than 5 times the detection limit. Potential error for such low values range between 50 % and 100 %.
- E The reported value is estimated due to the presence of a interference.

 The analysis of a serial dilution did not agree within 10 % of the original determination.
- s Indicates value determined by Method of Standard Addition.
- Q Daily EPA QC recovery outside 95% confidence limit.
- ** Surrogate recovery outside control limits indicating possible interferences.

DATA SUMMRY

88-P6-9 pottladus

188 199 OF

Workorder 88-05-147 Compiled_

PNN 7

units See below

			204			•	Thas		Ξ				CL			187_	#	PARAMETER		
			6-15-88		,		88-9-9		88-9-9				6-6-88		0	men soon	6-13-88	DATE	AMILYSIS	
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			1323				4,68		3.10						0.006	<0.200		ES	RECOYER	
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			:								,	:			(CO)	2.62	20.07	BLANKS		

RPD = [(IS-DI)/((S+D)/2)] x 100 RPD = Relative Percent Difference NC = Not calculable due to a value less than five times the IDL

SPIKE \$R = [(SSR-SR)/SA] x 100

* = Value is less than five times IDL = Instrument Detection Limit the instrument detection limit

SR = Sample Result
SA = Spiked Added SSR = Spiked Sample Result P = Prodigestion A = Analytical

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Compiled 6-29-88

Norkorder 86 - 05-147

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					•				S.,	1				
	A LINE IN COLOR	CAL	CALIBRATION STDS	ns.		ICATE AI	SISATW	ļ		SPIKE	RECOVERY			,
PARAMETER	DATE	FOUND	TRUE	€		1	Land	RPD T	JAHVS	SSR	BS	VS	SR	BLANKS
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						1			,					
Noz	6-10-88	1:27	1.25	401	@ E	C0.07	2002	NC	310	0.088	40.02	O.	58 ^P	
デト・	D. 03-49/ml	الهوا	Seil	103					,					7 9 7
	<i>I</i> * -													,

RPD = [(IS-DI)/((S+D)/2)] x 100
RPD = Relative Percent Difference
NC = Not calculable due to a value
less than five times the IDL

SPIKE \$R = [(SSR-SR)/SA] x 100

* * Value is less than five times

the instrument detection limit

IDL = instrument Detection Limit

A = Analytical P = Predigestion SSR = Spiked Sample Result

SA = Sample Result
SA = Spiked Added

DATA SUPPLARY

Submitted 6-29-88

Spassionca

J wod

0.00 1	0.050
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0-100 100 100 110	0.050
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٥٠١٥٥ (١٥٥)	J 0. 104
	6-7-88 9:100
TRIE I'S SAMPI SAMP	FOUND TRUE
CATION STDS. DIPLICATE ANALYSIS SPIKE	_ 9

RPD = [(15-D1)/((5+D)/2)] x 100
RPD = Relative Percent Difference
NC = Not calculable due to a value
less than five times the IDL

SPIKE \$R = [(SSR-SR)/SA] x 100

* = Value is less than five times
the instrument detection [imit
IDL = instrument Detection Limit

A = Analytical
P = Predigestion
SCD = Solid Score

SSR = Spiked Sample Result
SR = Sample Result
SA = Spiked Added

Norkorder 8805147 Compiled. Submitted 5-25-88

PNN7

	AMLYSIS	CAL	CALIBRATION SIDS.	DS.	Tano		SISAT			SPIRE	RECOVERY			
PARAMETER	DATE	FOUND	TRUE	ÎR.	SHE	SAMS	DIE	N S	JAMS	SSR	35	8	5	BLANKS
TOC	6-14-85	36.3	0.416	105	40 02 03	47.1	ان ان ##	8	250m	47.1	21.4	25	100	1
してっ	Thorassil.	2500 24-0	0-tre	101		n ng			500	for that	4.16	35	18	
	0	25.1	0.mc	185	,	, ,	-4							
			Ç.											
791	88-11-9	(-0d	0-00	101	015	66.6	63.9	4	ms OIE	121	66.6	60	9	_
i))C		60.7	60-0		astasu O I	اها	14	Ö	MSJ 61 E-	اجا	6.6	60	<u>අ</u>	
		0.49	0.0	163			. 1				·			
·														
Tox	88-8-9	0.924	1-00	92	450	0.137	0-11-3	F	4460	8HE.0	0.137	0.100	8	251.0
181	o. et malad	0.833	(-grg	83										
						``							·	
TOX	6-7-88	20.1	9	201		***								
10 (0.01 uslml	0.86	1-00	8			,	·						
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100	5.38-86													
SLIND	NTY					,	, r							
19.	1-00								·	• .				:
				;		ν.								,

RPD = [(15-D1)/((5+D)/2)] × 100 RPD = Relative Percent Difference NC = Not calculable due to a value less than five times the IDL

SPIKE SR = [(SSR-SR)/SA] × 100 IDL = Instrument Detection Limit * = Yalue is less than five times the instrument detection limit

SA = Spiked Added SR = Sample Result SSR = Spiked Sample Result P = Predigestion A = Analytical

RADIAN ANALYTICAL SERVICE QUALITY CONTROL DATA SUMMARY

Form II

Client ID	Geoscience
Workorder	88-05-147

Compiled PNN
Submitted 6-29-88

INITIAL AND CONTINUING CALIBRATION VERIFICATION 1

Units ug/ml

										0'	
_	Run		<u>ial Cali</u>					uing Cal			
Parameter N ₋	Date 6-15-68	True	Found 4.96	5 R	True 5-00	Found 4.86	\$ R	Found 4-97	\$ R	Found	\$R 94
49	6-15-60	5.00	5.11	10.5	5.00	5·11	102	5.17	103	5.12	
67					1						102
9		5.00	5.18	104	5.00	5.21	104	5.27	105	5.23	105
Cd Cr Fe		5.00	4.99	100	5.00	4.90	98	5.67	101	4.93	97
		20.0	20.0	100	20.0	19.7		20.2	101	19.9	100
Mo		5.80	5.00	100	5.00	4.97	99	5.02	100	#.95	99
No	₩.	20.0	21.2	106	20.0	20.1	101	31.3	106	20.9	105
		ļ			·				<u> </u>		
											1
	<u> </u>										1
	Con	i hu	na	(a	lub	ta to	n				
			4			·					
Aq	6-15-88				5-00	A.96	99				
1 28					500	5.18	104		1		
G					B.00	5.31	106		ì		
Cr					5.00	5.05	101				1
fe			7		20.0	20.1	101				
Ma					5-00	5.07	101				
Na				1	20.0	21.6	108				
109		1		<u> </u>			170				1
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	<u> </u>	<u> </u>	1	<u> </u>	H	<u> </u>	<u> </u>	<u> 1</u>	<u> </u>	1	<u>l</u>

1.	Control Limits	for	% R:	ICPES	
				AA F	

RADIAN ANALYTICAL SERVICES QUALITY CONTROL DATA SUMMARY

Form III

Client ID Geoscience Workorder 88-05-14

Compiled PNN Submitted 6-29

BLANKS

Units ____

	T	initial	Cont	inu ing Calib	ration	Prepar	ation
		Calibration		Blank Valu		Blank	
Parameter	IDL	Blank Value	11	2	3	1	2
Ae	0.003	40.003	KO·001	20.001	0.004	20.003	
Ae Be Cd	0.003	40.002	(0.001	40.003	<0.003	0.004	
Cd .	0.003	(D.003	60.001	CO.003	40.003	€0.003	
Cr	0.003	40.030	0.031	0.00%	0.014	0.006	
G Fe	0.003	0.085	0.132	0.049	0.081	0.035	
Ma	0.001	0.002	0.003	40.001	0.002	0.001	
Na	D.021	<0.013	4.508	40.021	<0.021	D. 044	
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^{1.} IDL = Instrument Detection Limit

^{*} indicates value is less than 5X the IDL.

RADIAN ANALYTICAL SERVICES QUALITY CONTROL DATA SUMMARY

Form VI

Client ID <u>Sesscience</u>
Workorder <u>88-05-147</u>

Complied

PNN

Submitted

6-29-88

Matrix

Aqueens

DUPLICATES
Type Pre Digeoted

Units usimal

Parameter	Sample No.	Control Limit	Sample (S)	Duplicate (D)	RPD 1	RPD Flag
) a	DDI	RPD < 20	40,003	0.001	NC	
Ba		'	0.011	0.089	68^	
Cd			<0.003	0.001	NC	
Ag Ba Cd Cr Fe			0.037	0.016	791	
Fe			10.2	10.2	0	
Mn			0.494	0.131	116	
Na		V	449	447	41	
1				8		
			<u> </u>			
			 			
		······································				
						-
	-					-

^{*} Indicates value is less than 5% IDL. (IDL=Instrument detection limit)

^{1.} RPD=Relative percent difference=[|S-D|/((S+D)/2)]X100.

^{2.} NC, =Not calculable due to a value less than 5X the IDL.

NC =Not calculable due to a value less than the CRDL. (Contract Required Detection Limit)

⁼RPD out of control limit for matrix duplicate, which may indicate non-homogeneity of the sample.

VOA RESULTS

LAB #			,			
CLIENT NAME						
SAMPLE ID	ゆんこくよう					
**************		************	**********			
EPA METHOD 601	DATE: ANALYST: INSTRUMENT:	EPA METHOD 602	DATE: CICA ANALYST: BAA INSTRUMENT LO			
COMPOUND	CONCENTRATION (ug/L)	COMPOUND	CONCENTRATION (ug/L)			
Chloromethane		Benzene	246			
Bromomethane		Toluene	44.6			
Vinvi Chloride		Ethvi benzene	37.9			
Chloroethane		Chlorobenzene				
Methylene chloride		1.4-Dichiorobenzene				
Trichlorofluorometh	ane	1.3-Dichlorobenzene	,			
1.1-Dichloroethene		1.2-Dichlorobenzene	•			
1.1-Dichloroethane		P-Xvlene	19.4			
Trans-1.2-Dichloroe	thene	M-Xvlene	20.2			
Chloroform		0-Xviene	237			
1.2-Dichloroethane						
1.1.1-Trichloroetha	ne		,			
Carbon tetrachiorid	e					
Bromodichloromethan	е] '				
1.2-Dichloropropane		SURROGATE RECOVERIES:				
Trans-1.3-Dichloron	ropene	601				
Trichloroethene		Bromochloromethane				
Dibromochloromethan	е	2-Bromo-1-Chloropropane				
1.1.2-Trichloroetha		1,4-Dichlorobutane				
cis-1.3-Dichloropro		4-Bromofluorobenzene				
2-Chloroethylyinyl]				
Bromoform		602	ŋ			
1.1.2.2-Tetrachloro	ethane	a,a,a,-Trifluorotoluene 187.				
Tetrachioroethylene		الملاماما وصدها	NAC DE LES MARTE			
Chlorobenzene			S LUKER CAME.			
1.3-Dichlorobenzene						
1.2-Dichlorobenzene						
1.4-Dichlorobenzene						

1:5

YOA RESULTS

LAB #X 5051	10.092		
CLIENT NAME	Godena		
SAMPLE ID	EUNICE 14		
EPA METHOD 601	DATE: ANALYST: INSTRUMENT:	EPA METHOD 602	DATE: 6/18 ANALYST: By INSTRUMENT &
COMPOUND	CONCENTRATION (ug/L)	COMPOUND	CONCENTRATION (ug/L)
Chloromethane		Benzene	0:61+/0.61+
Bromomethane		Toluene	0.12+/0.53*
Vinvi Chloride	٠.	Ethyl benzene	1
Chloroethane		Chiorobenzene	
Methylene chloride		1.4-Dichlorobenzene	
Trichiorofluorometh	ane	1.3-Dichlorobenzene	
1.1-Dichloroethene		1.2-Dichlorobenzene	
1.1-Dichloroethane		P-Xviene	
Trans-1.2-Dichloroe	thene	M-Xvlene	
Chloroform		0-Xviene	
1.2-Dichloroethane 1.1.1-Trichloroetha Carbon tetrachlorid Bromodichloromethan 1.2-Dichloropropane Trans-1.3-Dichlorop Trichloroethene Dibromochloromethan 1.1.2-Trichloroetha cis-1.3-Dichloropro 2-Chloroethylvinyi Bromoform 1.1.2.2-Tetrachloro Tetrachloroethylene Chlorobenzene 1.3-Dichlorobenzene 1.2-Dichlorobenzene 1.4-Dichlorobenzene	e ropene e ne pene ether ethane	SURROGATE RECOVER 601 Bromochloromethan 2-Bromo-1-Chlorop 1,4-Dichlorobutan 4-Bromofluorobenz 602 a,a,a,-Trifluorot	nenene

YOA RESULTS

LAB # XZOSH	リンク3A					
CLIENT NAME	<i>Gحو</i> ن(بدیمدذ					
SAMPLE ID	TIP BLANK					
EPA METHOD 601	DATE: ANALYST: INSTRUMENT:	EPA METHOD 602	DATE: Ulds ANALYST: DM INSTRUMENT; Oll			
COMPOUND	CONCENTRATION (ug/L)	COMPOUND	CONCENTRATION (ug/L)			
Chloromethane		Benzene	No			
Bromomethane		Toluene				
Vinyl Chloride		Ethyl benzene				
Chloroethane		Chlorobenzene				
Methylene chioride		1.4-Dichlorobenzene				
Trichlorofluoromethan	ne	1.3-Dichiorobenzene				
1.1-Dichloroethene	· · · · · · · · · · · · · · · · · · ·	1.2-Dichlorobenzene				
1.1-Dichloroethane		P-Xviene				
Trans-1.2-Dichioroeti	nene	M-Xylene				
Chloroform		0-Xviene	<u> </u>			
1.2-Dichloroethane						
1.1.1-Trichioroethans						
Carbon tetrachloride						
Bromodichioromethane						
1.2-Dichloropropane		SURROGATE RECOVER	ES:			
Trans-1.3-Dichloropre	opene	601				
Trichloroethene		Bromochloromethane	·			
Dibromochioromethane		2-Bromo-1-Chloropropane				
1.1.2-Trichloroethan		1,4-Dichlorobutane				
cis-1.3-Dichloroprop		4-Bromofluorobenze	ne			
2-Chloroethylvinyl e	ther		ń			
Bromoform		602	00)			
1.1.2.2-Tetrachloroe	thane	a,a,a,-Trifluoroto	oluene 110			
Tetrachioroethylene						
Chlorobenzene						
1.3-Dichlorobenzene		1				
1.2-Dichlorobenzene		4				
1.4-Dichlorobenzene		-				
·						

VOA RESULTS

EPA METHOD 601 COMPOUND Chloromethane Bromomethane Vinyl Chloride Chloroethane Methylene chloride Irichlorofluoromethane 1.1-Dichloroethane 1.1-Dichloroethane Irans-1.2-Dichloroethane Chloroform	DATE: ANALYST: INSTRUMENT: CONCENTRATION (ug/L)	EPA METHOD 602 COMPOUND Benzene Toluene Ethyl benzene Chiorobenzene 1.4-Dichiorobenzene 1.2-Dichiorobenzene P-Xylene M-Xylene	DATE: UNION ANALYST: ACINSTRUMENT ON (Ug/L) CONCENTRATION (Ug/L)			
COMPOUND Chloromethane Bromomethane Yinyl Chloride Chloroethane Methylene chloride Trichlorofluoromethane 1.1-Dichloroethane 1.1-Dichloroethane Trans-1.2-Dichloroethane Chloroform	ANALYST: INSTRUMENT: CONCENTRATION (ug/L)	COMPOUND Benzene Toluene Ethyl benzene Chiorobenzene 1.4-Dichiorobenzene 1.2-Dichiorobenzene P-Xylene	ANALYST: AO INSTRUMENT DO CONCENTRATION			
COMPOUND Chloromethane Bromomethane Vinvi Chloride Chloroethane Methylene chloride Trichlorofluoromethane 1.1-Dichloroethane Trans-1.2-Dichloroethene	ANALYST: INSTRUMENT: CONCENTRATION (ug/L)	COMPOUND Benzene Toluene Ethyl benzene Chiorobenzene 1.4-Dichiorobenzene 1.2-Dichiorobenzene P-Xylene	ANALYST: AO INSTRUMENT DO CONCENTRATION			
Chloromethane Bromomethane Vinvi Chloride Chloroethane Methylene chloride Trichlorofluoromethane 1.1-Dichloroethane 1.1-Dichloroethane Trans-1.2-Dichloroethane Chloroform	(ug/L)	Benzene Toluene Ethyl benzene Chiorobenzene 1.4-Dichiorobenzene 1.2-Dichiorobenzene P-Xylene				
Vinyl Chloride Chloroethane Methylene chloride Trichlorofluoromethane 1.1-Dichloroethane 1.1-Dichloroethane Trans-1.2-Dichloroethane Chloroform		Toluene Ethyl benzene Chiorobenzene 1.4-Dichiorobenzene 1.3-Dichiorobenzene 1.2-Dichiorobenzene P-Xylene				
Chloroethane Methylene chloride Trichlorofluoromethane 1.1-Dichloroethane 1.1-Dichloroethane Trans-1.2-Dichloroethane Chloroform		Ethyl benzene Chiorobenzene 1.4-Dichiorobenzene 1.3-Dichiorobenzene 1.2-Dichiorobenzene P-Xylene				
Chloroethane Methylene chloride Trichlorofluoromethane 1.1-Dichloroethane 1.1-Dichloroethane Trans-1.2-Dichloroethane Chloroform		Chiorobenzene 1.4-Dichiorobenzene 1.3-Dichiorobenzene 1.2-Dichiorobenzene P-Xylene				
Methylene chloride Trichlorofluoromethane 1.1-Dichloroethane 1.1-Dichloroethane Trans-1.2-Dichloroethane Chloroform		1.4-Dichiorobenzene 1.3-Dichiorobenzene 1.2-Dichiorobenzene P-Xylene				
Trichlorofluoromethane 1.1-Dichloroethane 1.1-Dichloroethane Trans-1.2-Dichloroethane Chloroform		1.3-Dichlorobenzene 1.2-Dichlorobenzene P-Xylene				
1.1-Dichloroethene 1.1-Dichloroethene Trans-1.2-Dichloroethene Chloroform		1.2-Dichiorobenzene P-Xviene				
1.1-Dichloroethane Trans-1.2-Dichloroethene Chloroform		P-Xviene				
Trans-1.2-Dichloroethene Chloroform						
Chioroform	, , , , , , , , , , , , , , , , , , ,	1 M_V1				
		M-VATEUR				
1 2-Dichiocoethana		0-Xylene	4)			
1.1.1-Trichloroethane						
Carbon tetrachloride]				
Bromodichloromethane]				
1.2-Dichloropropane		SURROGATE RECOVERIES:				
Trans-1.3-Dichloroproper	ıe	601				
Trichioroethene		Bromochloromethane				
Dibromochloromethane		2-Bromo-1-Chloropropane				
1.1.2-Trichloroethane		1,4-Dichlorobutane				
cis-1.3-Dichioropropene		4-Bromofluorobenzene				
2-Chloroethylvinyl ether	•					
Bromoform		602				
1.1.2.2-Tetrachloroethan	ne	a,a,a,-Trifluorotoluene				
Tetrachioroethylene		4				
Chlorobenzene		4				
1.3-Dichlorobenzene		4				
1.2-Dichlorobenzene		4				
1.4-Dichlorobenzene		4				

88-26-9

DAILY QUALITY CONTROL RAS GC LAB

DATE:		SPIKED VALUE (ug/L)	Analyzed Value	\$ Recovery	Analyzed \$
	INSTRUMENT		0	D	
TEST METHOD	COMPOUND		*******		***********
EPA 601	EPA WP 483 CONC. 2	•			
	Chloroform	12.0			
	1.2-Dichloroethane	2.0			•
	1.1.1-Trichioroethane	1.4			
	Carbon Tetrachioride	2.6			
	Browodichloromethane	2.0			
	Trichloroethene	2.9			
	Dibromochloromethane	2.6			
	Bromoform	2.9			
	Tetrachioroethene	1.6			
EPA 602	EPA - WP 879 CONC.1	30.7	28.8	94	
	Toluene	4.1	4.4	107	
yndg	Ethylbenzene	11.5	9-6	83	
	P-Xviene	19.1	19.0	99	
	M-Xviene	42.6	39.2	92	
	0-Xviene	10.6	10.5	99	
EPA 608		(ug/g)			
	Arocior 1242	58.7			
1	Arociar 1260	56.8		.	

RADIAN ANALYTICAL SERVICES QUALITY CONTROL DATA SUMMARY

Form II

			<u>FOLILLIT</u>	
Client ID	Georcience Ce	<u>msult</u> ants	Compiled	Ser 6-28-88
Workorder	88-05-050		Submitted	
	88-05-057 88-05-058	INITIAL AN	D CONTINUING CALIBRATION VERIFICATION	
			Units	refore

• ٧	Run Initial Calib			Continuing Calibration							
Parameter	Date	True	Found	%R	True	Found	S R	Found	%R	Found	% R
Ag	6-13-88	5.00	4.78	96	5.00	4.89	98	4.74	95		
<u> (</u>	,		4.92	98		5.03	101	5.02	100		
Ca			4.91	98		5.00	100	4.91	198		
Cr			4.76	95		4.78	96	4.71	94		
Fe		20.0	19.5	98	20.0	19.6	98	19.3	97		
mn		5.03	4.78	96	5.00	4.82	96	4.69	94		
72		20.0	20.4	١٥٧	20.0	17.9	90	20.9	105		
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1. Control Limits for \$R: ICPES _ AA F _

RADIAN ANALYTICAL SERVICES QUALITY CONTROL DATA SUMMARY FORM LLL

Client ID	Georgiance Commeltano
Workorder	88-05-050
	88-05-057
	09-05-058

Submitted See 6-28-88

Matrix acres

BLANKS

Units _____

,		<u>initial</u> Calibration	Cont	inuing Calibr Blank Value		Prepar Blank	
Parameter	IDL	Blank Value	11_	2	3		2
Aa	0.003	60.003	(0.003	<0.003		(0.003	
<u> </u>	0.003	(0.003	40.003	(0.003		0.003*	
Cd	5.003	(0.003	< 0.003	40.003		KO 003	
C-	0.003	40.003	< 0.003	40.003		(0:003	
Fe	0.003	0.156	(0.003	<0.003		0.031	
mn	0.001	(0.001	(0.001	40.001		0.002	•
Na '	0.021	0.036	10.021	140.021		3.17	
						`	
						`	#90 <u>.</u>
							·
	·				,		
					-		
				·			

^{1.} IDL = Instrument Detection Limit

^{*} indicates value is less than 5X the IDL.

KALIAN ANALYTICAL SERVICES

QUALITY CONTROL DATA SUMMERY

Form Y

Client	10	Ger	حى	ence Consultants
		00		

₩orkorder <u>88-05-050</u>

88-92-028

Submitted

Matrix

Agreems

Agreems

Matrix

SPIKED SAMPLE RECOVERY

Spiking method predicention

Units _____ /ne

Parameter	Sample No.	Control Limit ot\$R	Spiked Sample Result (SSR)	Sample Result(SR)	Spike Added (SA)	D.F.	sR [†]	SR Flag
Aq	Blan 16	75-125	0.856	B	1.00	,	86	
Aq Ba			0.875			1	88	
cd			0.888				89	
Cr			0.856				86	
Fe	4	·	8.77		10.0		88	
Mn			0.861	, ,	1.00		96	
Na			8.92		10.0	_	89	
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Indicates value is less than 5x IDL. (IDL=Instrument detection limit)

2. R=For matrix spike: SR was not within control limit, which may indicate matrix interferences.

B-Sample result was greater than 4 times spike added concentration, therefore spike added concentration is considered insignificant.

B) - Blank result is subtracted out from the somple result.

^{1. \$}R = Percent Recovery = [(SSR - SR)/SA] X 100

QUALITY CONTROL DATA SUMMARY

Form VI

Client 10 Gerociance Consultents		Compiled	Fyr 6-28
Workorder 89-05-050	•	Submitted	
		Matrix	agueron
	DUDI ICATEC		D

DUPLICATES

Type medigention

Units we/ml

					0		
Parameter	Sample No.	Control Limit	Sample (S)	Ouplicate (D)	RPD 1	RPD Flag ²	
<u> </u>	05050-01	RPD = 20	<0.003	<0.003	· NC		
Ba			0.056	0.056	0		
Cal			0.016	0.016	0		
Cr			0.024	0.033	32	^	
Fe		<u>.</u>	0.069	0.106	42	^	
Mn			0.279	0.778	0		
Na		Į.	227	225			
				:		·	
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					1		
			1				
·						 	
				 			
	 					 	
	-						
	 		-			 	
	 					 	
	 		+	 		 	
	1			 		 	

- * Indicates value is less than 5% IDL. (IDL=Instrument detection limit)
- 1. RPD=Relative percent difference=[IS-D]/((S+D)/2)]X100.
- 2. NC, =Not calculable due to a value less than 5X the IDL.
 - NC =Not calculable due to a value less than the CRDL. (Contract Required Detection Limit.
 - =RPD out of control limit for matrix duplicate, which may indicate non-homogeneity of the sample.

DAILY QUALITY CONTROL RAS GC LAB

DATE:	5-12-88	SPIKED VALUE (ug/L)	Analyzed Yalue	Recovery		Analyzed Value	\$ Recovery
	INSTRUMENT		D	D		G	G-
TEST METHOD	COMPOUND		*********		, 48.88		
EPA 601	EPA WP 483 CONC. 2						
	Chloroform	12.0				,	
	1.2-0 ich ioroethane	2.0					
	1.1.1-Trichiorosthans	1.4					
·	Carbon Tetrachioride	2.6					1
	Bromodichioromethane	2.0					
	Trichioroethene	2.9	·		,		
	Dibromochioromethane	2.6					
	Bromeferm	2.9		,			
,	Tetrachloroethene	1.6					
EPA 602	EPA - WP 879 CONC.1						
	Benzene	30.7	26.5	86		22".7	74
	Toluene	4.1	4.4	107		4.5	110
	Ethylbenzene	11.5	9.7	84		9.3	81
	P-Xviene	19.1	19:7	103		19.0	99
	M-Xvlene	42.6	38.8	91		35.5	83
	G-Xviene	10.6	10.4	98		10.7	101
EPA 608		(ug/g)					
	Arociar 1242	58.7				 	
	Arociar 1260	56.8	I	1		1	1

DAILY QUALITY CONTROL RAS GC LAB

DATE:	5-16-88	SPIKED VALUE (ug/L)	Analyzed Value	\$ Recovery		Analyzed Value	\$ Recovery
	INSTRUMENT		В	R		૯	6
TEST METHOD	COMPOUND				*****	*******	
EPA 601	EPA WP 483 CONC. 2						
	Chloroform	12.0	11.7	98			
	1.2-Bichiomethans	2.0	1.9	95			
	1.1.1-Trichiorosthans	1.4	2.1	150	<u>.</u>		
1	Carbon Tetrachionide	2.6	2.8	108			1
	Bromodichioromethane	2.0	2.3	115			
	Trichioroethene	2.9	3.0	103			
	Dibromochioromethane	2.6	4.5	173			
	Bromeform	2.9	2.2	76			
	Tetrachloroethene	1.6	1.7	106			
EPA 602	EPA - WP 879 CONC.1		D	D			
	Benzene	30.7	27.8	91	1	30.1	98
	Toluene	4.1	4.7	115		5.4	132
	Ethylbenzene	11.5	9.8	85		10.5	91
	P-Xviene	19.1	20.6	108		22.5	1180
	M-Xvlene	42.5	40.1	94		42.5	100
	Q-Xviene	10.6	11.1	105		12.5	118
EPA 608		(ug/g)				}	
	Arociar 1242	58.7			1		
	Arocior 1260	56.8		1		1	i

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	TM RIANK		
CLIENT NAME			
SAMPLE ID			
EPA METHOD 601	DATE: ANALYST: INSTRUMENT:	EPA METHOD 602	DATE: SILAS ANALYST: AC INSTRUMENTO
COMPOUND	CONCENTRATION (ug/L)	COMPOUND	CONCENTRATION (ug/jL)
Chloromethane	•	Benzene	$\mathcal{N}_{\mathcal{D}}$
Bromomethane	• 1	Toluene	
Vinyl Chloride		Ethyl benzene	
Chloroethane		Chlorobenzene	
Methylene chioride		1.4-Dichlorobenzene	
Trichiarofluorometha	ne	1.3-Dichlorobenzene	
1.1-Dichioroethene	·	1.2-Dichlorobenzene	
1.1-Dichloroethane		P-Xvlene	
Trans-1.2-Dichloroe	thene	M-Xylene	
Chioroform		0-Xylene	19
1.2-Dichloroethane		·	·
1.1.1-Trichloroetha			
Carbon tetrachlorid		·	
<u>Bromodichioromethan</u>			.50
1.2-Dichloropropane		SURROGATE RECOVER	IES:
Trans-1.3-Dichloron	ropene	601	
Trichloroethene		Bromoch Loromethan	
Dibromochloromethan		2-Bromo-1-Chlorop	
1.1.2-Trichloroetha		1,4-Dichlorobutan	
cis-1.3-Dichloropro		4-Bromofluorobenz	916
2-Chloroethylvinyl	etn e r	602	
Bromoform 1.1.2.2-Tetrachioro		602	en lucan
Tetrachioroethylane		a,a,a,-Trifluorot	Oldaua
Chlorobenzene		1	
1.3-Dichlorobenzene		4	
1.2-Dichlorobenzene		4	•
1.4-Dichlorobenzene		1	
A TOUT ON SHARING			
		•	
		1	

	ENT RUNK-UFW	J	
CLIENT NAME			
SAMPLE ID			
**************		****************	
EPA METHOD	DATE:	EPA METHOD	DATE: 5/12/38
601	ANALYST:	602	ANALYST: 4
	INSTRUMENT:		INSTRUMENT!
COMPOUND	CONCENTRATION (ug/L)	COMPOUND	CONCENTRATION (ug/L)
Chioromethane		Benzene	\mathcal{N}
romomethane	,	Toluene	
Vinvi Chloride		Ethvi benzene	
hloroethane		Chlorobenzene	
fethviene chloride		1.4-Dichlorobenzene	
[richlorofluorometha	ane	1.3-Dichlorobenzene	
1.1-Dichloroethene		1.2-Dichlorobenzene	
.1-Dichloroethane		P-Xviene	
Trans-1.2-Dichloroe	thene	M-Xvlene	
Chloroform		0-Xviene	
1.2-Dichloroethane			
1.1.1-Trichioroetha	18		
Carbon tetrachloride			
Bromodichloromethan	9]	•
1.2-Dichloropropane		SURROGATE RECOVER	RIES:
Trans-1.3-Dichloron	ropene	601	
Trichloroethene		Bromochloromethan	le
Dibromochloromethan		2-Bromo-1-Chlorop	
1.1.2-Trichioroetha		1,4-Dichlorobutar	
cis-1.3-Dichloropro	pene	4-Bromofluorobenz	eneene
2-Chloroethylvinyl	ether	1	
Bromoform		602	
1.1.2.2-Tetrachloro		a,a,a,-Trifluorot	oluene
Tetrachloroethylene		_	
Chlorobenzene		1	
1.3-Dichlorobenzene		4	
1.2-Dichlorobenzene		4	
1.4-Dichlorobenzene			
		1	

	sten Bime		
CLIENT NAME			
SAMPLE ID			

EPA METHOD	DATE:	EPA METHOD	DATE: 3/3/06
601	ANALYST:	602	ANALYST: MO
	INSTRUMENT:	· ·	INSTRUMENT
COMPOUND	CONCENTRATION	COMPOUND	CONCENTRATION
	(ug/L)		(ug/L)
)			10
Chloromethane Bromomethane		Benzene Toluene	γ-
Vinyl Chlorida		Ethyl benzene	
Chioroethane		Chlorobenzene	
Methylene chloride		1.4-Dichiorobenzene	
Trichlorofluorometha	ne	1.3-Dichlorohenzene	
1.1-Dichloroethene		1.2-Dichlorobenzene	
1.1-Dichloroethane		P-Xvlene	
Trans-1.2-Dichloroet	hene "	M-Xviene	1/
Chloroform		0-Xvlene	
1.2-Dichloroethane			
1.1.1-Trichloroethar			•
Carbon tetrachloride		1	
<u>Bromodichloromethane</u>		1	
1.2-Dichioropropane		SURROGATE RECOVER	IES:
Trans-1.3-Dichlorops	Copene	601	
Trichloroethene		Bromoch (oromethan	4,4,
Dibromochloromethane		2-Bromo-1-Chlorop	
1.1.2-Trichloroethan		1,4-Dichlorobutan	
cis-1.3-Dichloroprop		4-Bromofluorobenz	ene
2-Chloroethylvinyl	ether	+	
Bromoform		602	
1.1.2.2-Tetrachioro	etnane	a,a,a,-Trifluorot	Oldeue
Tetrachloroethylene		4	
Chiorobenzene		4	
1.3-Dichlorohenzene		4	
1.2-Dichlorobenzene		4	
1.4-Dichlorobenzene		-	
	·		
		•	

	ethera Bianc of	~	
CLIENT NAME			
SAMPLE ID			
· 医非毛球病疾病 经有效的 化二甲基甲基甲基甲基甲基甲基甲基甲基甲基甲基甲基甲基甲基甲基甲基甲基甲基甲基甲基	· 高速电影等高速电影电影电影电影电影	医聚聚式医聚聚素或食物聚素或医聚聚素等	*********
EPA METHOD	DATE:	EPA METHOD	DATE: 5/13/2
601	ANALYST:	602	ANALYST: C
	INSTRUMENT:		INSTRUMENT
COMPOUND	CONCENTRATION	COMPOUND	CONCENTRATION
	(ug/L)		(ug/L)
Chioromethane		Benzene	N
Bromomethane		Toluene	7
Vinvi Chioride		Ethyl benzene	
Chloroethane		Chlorobenzene	
Methylene chloride		1.4-Dichlorobenzene	
Trichlorofluorometha	ane	1.3-Dichiorobenzene	
1.1-Dichloroethene		1.2-Dichlorobenzene	
1.1-Dichtoroethane		P-Xvlene	
Trans-1.2-Dichloroet	thene	M-Xvlene	7/
Chioroform		0-Xviene	
1.2-Dichioroethane			
1.1.1-Trichloroethan	ne	·	
Carbon tetrachioride	9		
Bromodichioromethane	9	<u>}</u>	
1.2-Dichloropropane		SURROGATE RECOVER	HES:
Trans-1.3-Dichloron	ropene	601	
Trichloroethene		Bromochloromethan	le
Dibromochioromethane	9	2-Bromo-1-Chlorop	ropane
1.1.2-Trichloroethau		1,4-Dichlorobutan	le
cis-1.3-Dichloropro		4-Bromofluorobenz	ene
2-Chloroethylvinyl	ether	<u>}</u>	
Bromoform		602	
1.1.2.2-Tetrachioro	ethane	a,a,a,-Trifluorot	oluene
Tetrachioroethylene			
Chlorobenzene		1	
1.3-Dichlorobenzene			
1.2-Dichiorobenzene			
1.4-Dichlorobenzene			

Compiled_ 88-82-9 - 455 potitionans

Workorder 68 - 05 - 050

Client Geoscience Consultant

Units we for under

					So		·	77				C	c	Proposed in	Calibra		101-0.002-	As	PARAMETER		
					5-31-88			5-31-88				5-31-88		colonies/100ml	5-30-88		e la constant	5-17-88	DAIL	VMTASIS	
			39.4	39.2	39.0	3.99	3.99	4.01		7.86	7.83	7.79		Oml			0.086	P 60.0	COMPO	CAL	
		,	40.0	40.0	40.0	H.00	4.00	4.00		8.00	8.00	8.00					0.086 0.100	0.100	FOUND TRUE SA	CALIBRATION S	
			99	98	98	100	8	100		98	98	47						1			
	,																		SAMP	our i	
																			Tang akvs	ICATE AN	
	. 4	-							,	.,			·						DUP	SISAT	
																			RPD	·	
																		3020 5/16 Blank	MAN		
																		0.019	HSS	SPIKE	
																			SE	RECOYERY	P T
																	_	0.020	AS		otherwise.
Ą																0>	6 33	900	SR BLA		worked.
		,		,												9 (0.002	3020 5/14	\$0.00 Z	SXIV		Ø

RPD = [(IS-DI)/((S+D)/2)] × 100
RPD = Relative Percent Difference NC = Not calculable due to a value less than five times the IDL

SPIKE $$R = [(SSR-SR)/SA] \times 100$ IDL = Instrument Detection Limit * - Value is less than five times the instrument detection limit

> SSR = Spiked Sample Result A = Analytical P = Predigestion

SA = Spiked Added SR = Sample Result

DINTITLY (

Workorder 88-05-050 Comp 11ed Submitted Ses 6-28-88

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Unit's aug of whole

		2	ADDAT ION					; "	7		oth	other Company	noted	D
	AMLYSIS	YERIEL	CALIBRATION S	DS.	Land	CATE	SISTIMIA				RECOVERY			;
PARAMETER	DATE	FOUND	FOUND TRUE IR		MAPA	PMVS	1400	RPD	/JANYS	SSR	S)E	AS.	SA B	SXI
Ha	88-61-5	0.0036	0.0036 D.0040	90								į.	√G	\$0.000
101 - 0.0001 -	a ford	D. 0034	D. 0034 0.0040	58			٠					-	VC	100000
		0.0038	0.0038 0.000	95									V/2	(0.000)
		,	·										,	
NO	5-24-88	1.35	1.25	108										
10c = 0.07 -w	170	1.35	1.25	108										
<u></u>	-	1.35	1.25	08				•						
	,						,							
Pb	5-18-88	P69.0	0.100	કત		*							200	(0.001
18L=0.002 -	me	0.096	0.100	96		-							0 %	0.003
		-					,						र्जे र	(0.00Z
PH	5-12-88	6.97	7.00	100	9 01 A	7.29	7.29	0						
Reported	~ PH units.	7.02	7.00	100	010	7.23	7.26	0						
-		6.98	7.00	100									-	
												-		
Pheno Rics	5-27-88	0.263	0.250	105	·								_	
101=0.005 mg/mg	e/me	0.276	0.250	110									_	
							٠							
												,	_	

. NC = Not calculable due to a value $RPD = [(1S-D1)/((S+D)/2)] \times 100$ RPD = Rejative Percent Difference less than five times the IDL

IDL = Instrument Detection Limit * = Value is less than five times SPIKE \$R = [(SSR-SR)/SA] × 100 the instrument detection limit

SR = Sample Result SSR = Spiked Sample Result SA = Spiked Added P = Predigestion A = Analytical

Compiled_ Workorder 88-05-050 Submitted 584 - 6-28-88

Cillent beoggiance

Units us for where Otherwise noted

المراجعة											0156	ロナスマイン・	3	Q
	ANALYSIS	CAL	CALIBRATION S	SIDS	2 1	CATE	ANN YSIS	•		SPIKE	A8 3A 0.338			
 PARAMETER	DATE	FOUND TRUE	TRUE		JAMS	SAMO	DuPl	RPD	1 AMS	SSR	SIR	SA	AR	BLANKS
Se	5-18-88	0.049	0.050	8			4		3020 5/16 Blank	0.022	١	0.020		50003
CO3 _	عہ امید	0.053	0.050	106		ţ		,	•					800.05
	0		-	v		,	7	,						
 Toc	6-1-88	34.5	32.8	105	DOIF	8.24	7.48	w	MS 01 E	39.6	8.11	30.0	501	9.10
 101=1.0 we/m	Q	32.2	32.8	48	ms + msp	39.6	62.6	+	M 50 F	62.6	8.11	30.0	182	
 0				• "	ms theo	39·b	38.9	2	01 F	38.9	8.11	30.0	103	
	·		·	,	1	,		•			·			
 Tox	5-18-88	0.957	1.00	96										
 101 = 0 01 m	lose	0.627	1.00	63		<i>y</i>	,	·	·		·			
 >	\ \				- 		. •							
 Tox	5-19-88	0.878	1.00	88	A) 01 (F	0.103	180.0	6	B 0 I	0.203	0.053	0.100	- 0	
 1DL =0.01 mg	Ina	D.963	.00	98			**							
				ÿ	·	ye ug N	÷							
 Turbidita	5-12-88				8) OI	17.0	18.0	c						0.7
			,				·							0.7
				ì		يوام								
 MHO	5-12-88	1389	1413	96	8) 01 D	265	270	2						
		1359	1413	96			. ,							
		1374	1413	97								_		

NC = Not calculable due to a value $RPD = [(1S-D1)/((S+D)/2)] \times 100$ NPD = Relative Percent Difference less than five times the IOL

" = Value is less than five times IDL = Instrument Detection Limit SPIKE JR = [(SSR-SR)/SA] × 100 the instrument detection limit

A = Analytical P = Predigestion

SA = Spiked Added SR = Sample Result SSR = Spiked Sample Result

DAILY QUALITY CONTROL RAS GC LAB

DATE:	5-13-88	SPIKED VALUE (ug/L)	Analyzed Yalue	Recovery	Anal yzed Yalue	\$ Recovery
	INSTRUMENT		B	В	<u> </u>	G
TEST METHOD	COMPOUND			*********	**********	*********
EPA 601	EPA WP 483 CONC. 2			,		
	Chloroform	12.0	11.7	98		
	1.2-0 ich lorgethane	2.0	1.9	95		
	1.1.1-Trichioroethane	1.4	2.1	150		
	Carbon Tetrachioride	2.6	2.8	108		1
	Bromodichioromethane	2.0	2. 3	115	·	
	Trichioroethene	2.9	3.0	103		
	Dibromochioromethane	2.6	4.50	175	·	
	Bromeform	2.9	2. 2	76		
	Tetrachloroethene	1.6	1.7	106		
EPA 602	EPA - WP 879 CONC.1		D	D		
	Benzene	30.7	27.8	91	30.1	98
	Totuene	4.1	4.7	115	5.4	132
	Ethylbenzene	11.5	9.8	85	10.5	91
	P-Xviene	19.1	20.6	108	22.5	1118 e
	M-Xylene	42.6	40.1	94	42.5	100
	0-Xviene	10.6	11.1	105	12.5	118
EPA 608		(ug/g)				}
	Arociar 1242	58.7				
	Arociar 1260	56.8				1

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VOA SPIKE RESULTS

LAB # 88-05-057 SAMPLE ID 02 T METHOD EPA 602

UNITS___/

COMPOUND	SPIKED SAMPLE RESULT (SSR)	SAMPLE RESULT (SR)	SPIKE ADDED (SA)	≸ R
Benzene	35.0	15.3	3ი. ე.	114
Toluene	5.2		ч :1	127
Ethyl benzene		46.1	11.5	97
Chiorobenzene				
1.4-Dichlorobenzene	·			
1.3-Dichlorobenzene	·			
1.2-Dichlorobenzene				,
P-Xylene		24.4	19.1	128 @
M-Xylene		47.3	42.6	111 @
0-Xylene		12.0	10.6	113

Spike of deluted sample. Onea counts outotracted out: \$ R = [(SSR-SR)/SA] × 100

Bromomethane Vinvi Chloride Chloroethane Methylene chloride Trichlorofluoromethane 1.1-Dichloroethane 1.1-Dichloroethane Trans-1.2-Dichloroethane Chloroform 1.2-Dichloroethane	DATE: ANALYST: INSTRUMENT: CONCENTRATION (ug/L)	EPA METHOD 602 COMPOUND Benzene Toluene	DATE:5//3/86 ANALYST: ACINSTRUMENT; CONCENTRATION (ug/L)
COMPOUND Chloromethane Bromomethane Vinyl Chloride Chloroethane Methylene chloride Trichlorofluoromethane 1.1-Dichloroethene	ANALYST: INSTRUMENT: CONCENTRATION	COMPOUND Benzene	INSTRUMENT O
Chloromethane Bromomethane Vinyl Chloride Chloroethane Methylene chloride Trichlorofluoromethane 1.1-Dichloroethane 1.1-Dichloroethane Trans-1.2-Dichloroethane Chloroform 1.2-Dichloroethane		Benzen e	
Bromomethane Vinvi Chloride Chloroethane Methylene chloride Trichlorofluoromethane 1.1-Dichloroethane 1.1-Dichloroethane Trans-1.2-Dichloroethane Chloroform 1.2-Dichloroethane			110
Vinvi Chloride Chloroethane Methylene chloride Trichlorofluoromethane 1.1-Dichloroethane 1.1-Dichloroethane Trans-1.2-Dichloroethane Chloroform 1.2-Dichloroethane		Toluene	
Chloroethane Methylene chloride Trichlorofluoromethane 1.1-Dichloroethane 1.1-Dichloroethane Trans-1.2-Dichloroethane Chloroform 1.2-Dichloroethane		فالمستحد والمستحد	
Methylene chloride Trichlorofluoromethane 1.1-Dichloroethane 1.1-Dichloroethane Trans-1.2-Dichloroethane Chloroform 1.2-Dichloroethane		Ethyl benzene	
Trichlorofluoromethane 1.1-Dichloroethene 1.1-Dichloroethane Trans-1.2-Dichloroethene Chloroform 1.2-Dichloroethane		Chlorobenzene	
1.1-Dichioroethene 1.1-Dichioroethane Trans-1.2-Dichioroethene Chioroform 1.2-Dichioroethane		1.4-Dichlorobenzene	
1.1-Dichioroethane Trans-1.2-Dichioroethene Chioroform 1.2-Dichioroethane		1.3-Dichlorobenzene	
Trans-1.2-Dichloroethens Chloroform 1.2-Dichloroethane		1.2-Dichlorobenzene	
Chioroform 1.2-Dichioroethane		P-Xvlene	
1.2-Dichloroethane	<u> </u>	M-Xviene	
		0-Xylene	<u> </u>
1 1 1-Trichloroothana		4	
		4	
Carbon tetrachloride		4	
Bromodichloromethane		<u>.</u>	•
1.2-Dichloropropane	·	SURROGATE RECOVER	ES:
Trans-1.3-Dichloroprope	ne	_ 601	
Trichloroethene		_ Bromochloromethane	
Dibromochloromethane		2-Bromo-1-Chloropr	
1.1.2-Trichloroethane		1,4-Dichlorobutane	
cis-1.3-Dichloropropene		4-Bromofluorobenze	ne
2-Chloroethylvinyl ether	£	4	
Bromoform		602	
1.1.2.2-Tetrachioroetha	ne	a,a,a,-Trifluoroto	luene
Tetrachloroethylene		4	
Chlorobenzene		4	
1.3-Dichlorobenzene		-{	
1.2-Dichlorobenzene		4	
1.4-Dichlorobenzene		1	
•			

CLIENT NAMESAMPLE ID	a Blank of		
EPA METHOD 601	DATE: ANALYST: INSTRUMENT:	EPA METHOD 602	DATE: \$\ist\ \ \ ANALYST: CY INSTRUMENT: \(\)
COMPOUND	CONCENTRATION (ug/L)	COMPOUND	CONCENTRATION (ug/L)
Chioromethane ·		Benzene	No
Bromomethane		Toluene	
Vinvl Chloride		Ethyl benzene	
Chloroethane		Chlorobenzene	
Methylene chloride		1.4-Dichiorobenzene	
Trichlorofluorometha	ne	1.3-Dichiorobenzene	
1.1-Dichloroethene		1.2-Dichlorobenzene	
1.1-Dichloroethane		P-Xvlene	
Trans-1.2-Dichloroet	hene	M-Xviene	
Chloroform		0-Xylene	
1.2-Dichloroethane 1.1.1-Trichloroethane 1.1.1-Trichloroethane Carbon tetrachloride Bromodichloromethane 1.2-Dichloropropane Trans-1.3-Dichloropro Trichloroethane Dibromochloromethane 1.1.2-Trichloroethan cis-1.3-Dichloroprop 2-Chloroethylvinyl e Bromoform 1.1.2.2-Tetrachloroe Tetrachloroethylene Chlorobenzene 1.3-Dichlorobenzene 1.2-Dichlorobenzene 1.4-Dichlorobenzene	opene e ene ther	SURROGATE RECOVER! 601 Bromochloromethane 2-Bromo-1-Chloropr 1,4-Dichlorobutane 4-Bromofluorobenze 602 a,a,a,-Trifluoroto	opane

		- BLMK	LAB #
			SAMPLE ID
DATE:5/13/2	EPA METHOD	DATE:	EPA METHOD
ANALYST: AC	602	ANALYST:	601
INSTRUMENT		INSTRUMENT:	331
Q li		111011101161117	
CONCENTRATION	COMPOUND	CONCENTRATION	COMPOUND
(ug/L)	•	(ug/L)	
		J ·	
ND	Benzene		Chloromethane
	Toluene		Bromomethane
· · · · · · · · · · · · · · · · · · ·	Ethyl benzene		Vinyl Chloride
	Chlorobenzene		Chioroethane
ene	1.4-Dichlorobenzene		Methylene chloride
	1.3-Dichlorobenzene		Trichlorofluoromethane
	1.2-Dichlorobenzene		1.1-Dichloroethene
	P-Xylene		1.1-Dichloroethane
	M-Xviene	ane	<u> Trans-1.2-Dichloroether</u>
- V	0-Xylene		Chloroform
•			1.2-Dichloroethane
			1.1.1-Trichloroethane
	1		Carbon tetrachloride
			Bromodichloromethane
VERIES:	SURROGATE RECOVER!		1.2-Dichioropropane
	601	oene	Trans-1.3-Dichloroprope
	Bromoch loromethane		Trichioroethene
	2-Bromo-1-Chloropr		Dibromochloromethane
	1,4-Dichlorobutane		1.1.2-Trichloroethane
enzene	4-Bromofluorobenze		cis-1.3-Dichloropropene
	+	ner	2-Chloroethylvinyl ethe
	602		Bromoform
protoluene	a,a,a,-Trifluoroto	nane	1.1.2.2-Tetrachloroetha
	4		Tetrachioroethylene
	4		Chlorobenzene
	• •		1.3-Dichlorobenzene
	+		1.2-Dichlorobenzene
	4		1.4-Dichlorobenzene

SAMPLE ID	************		***************
EPA METHOD 601	DATE: ANALYST: INSTRUMENT:	EPA METHOD 602	DATE: 5/13/88 ANALYST: () INSTRUMENT:
COMPOUND	CONCENTRATION (ug/L)	COMPOUND	CONCENTRATION (ug/L)
Chioromethane		Benzene	
Bromomethane		Toluene	
Vinyl Chloride		Ethyl benzene	
Chloroethane		Chiorobenzene	
Methylene chloride		1.4-Dichlorobenzene	
Trichlorofluorometha		1.3-Dichlorobenzene	
1.1-Dichloroethene		1.2-Dichlorobenzene	
1.1-Dichloroethane	,	P-Xvlene	10
Trans-1.2-Dichloroet	hene	M-Xv lene	
Chioroform		0-Xylene	<i>V</i>
1.2-Dichloroethane			
1.1.1-Trichloroethan			
Carbon tetrachloride		1	
Bromodichloromethane			.50
1.2-Dichloropropane		SURROGATE RECOVER	ES:
Trans-1.3-Dichloropr	Opene	601	_
Trichloroethene		Bromoch loromethane	9
Dibromochloromethane		2-Bromo-1-Chiloropi	-opane
1.1.2-Trichloroethan		1,4-Dichlorobutane	
cis-1.3-Dichloropror		4-Bromofluorobenze	ne
2-Chloroethylvinyl	Ther	1 600	
Bromoform		602	-1
1.1.2.2-Tetrachioroe	Thane	a,a,a,-Trifluoroto	ordeue
Tetrachioroethylene Chiosphospas		4	
Chlorobenzene 1.3-Dichlorobenzene		+	
1.2-Dichiorobenzene		4	
1.4-Dichlorobenzene		4	
		1	

Submitted Sign 6-28-88

Morkorder 88-05-057

Cilent Capocance Consultanto

Units we me malace

Propose in 101.002 Californ 105 PARAMETER T colfernoso 5-17-88 5-31-88 5-31-88 5-31-88 S-30-88 SISATMV 0.086 980.0 0.094 4.01 7.86 39.4 7.79 39.2 3.99 FOUND | IRUE 40.0 3.99 7.83 VERIFICATION SIDS. CALIBRATION 0.100 H.00 4.00 0.100 0.100 40.0 4.00 8.00 40.0 8.8 8.00 40.0 100 001 98 99 100 98 93 00 99 19 86 9 DUPLICATE ANALYSIS P10.0 0.014 RPD b 3020 S/16 マイマンス 0.019 SPIKE 355 RECOVER O themine noted 0.020 3.5 A P 200 002 BLANKS 3020 5/16 <0.002 <0.002

RPD = [(IS-DI)/((S+D)/2)] × 100
RPD = Relative Percent Difference
NC = Not calculable due to a value
less than five times the IDL

SPIKE \$R = [(SSR-SR)/SA] x 100

** Value is less than (ive times the instrument detection limit in the instrument Detection Dete

A = Analytical P = Predigestion SSR = Spiked Sample Result

SR = Sample Result
SA = Spiked Added

Submitted Str 6-28-88

Workorder 88-05-057

Cilent beggieres Consultand

Otherware noted

	SISATWA	CAL VERJE I	CALIBRATION WERIFICATION STDS	TOS.	DUPLI	ICATE A	ICATE ANALYSIS			SPIKE	RECOVE RY	D themores	- noted
PARAMETER	DATE	COUND	IRUE	SR	JAMVS	JHVS	Tang	RPD	JAMVS	SSR	SIR	X	SH BLANKS
Ha	5-13-68	0.0036		Ô			·	1					21
<u>0</u>	elme	6.0034	0.0040	85								-	P)
1	G	0.0038		95									0.000
			_					,`					
340	5-17-88	1320	1413	93	8) 02 B	3430	OGHE						
Ruported in	sumbos unito	1344	1413	95					·			·	
_		·				· ·							
76 20	5-23-88	1.26	1.25	101		,							
02	120	1.26	1.25	101						,			
						,	,						
Ьb	5-18-88	P60.0	0.100	94	ပ ွဲ့	500.002	(0.002	NC	3020 5/16 Blonk	0.012		0.020	(10 Ke-po2
106: 0:002	FR.	0.096	0.160	46									0.003
		0.104	0.100	104		ر د <u>د</u>							p) <0.∞2
	,			,a,	·		*						(A) √0.∞2
PH	5-17-88	7.00	7.00	100	A) 02 B	7.24	7.23	0					
Reported in	DH wite.	7.00	7.00	100									
a		7.00	2.00	100									
	,					.,	,						

RPD = [(15-D1)/((5+D)/2)] × 100
RPD = Rejative Percent Difference
NC = Not calculable due to a value
less than five times the IDL

SPIKE \$R = [(SSR-SR)/SA] × 100

** - Value is less than five times

the instrument detection limit

10L = instrument Detection Limit

A = Analytical P = Predigestion SSR = Spiked Sample Result

SR = Sample Result SA = Spiked Added

Submitted Sylve 6-28-88
Compiled

Workorder 88-05-057

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1		CA	CALIBRATION	3							240	Otherbonne maked	, ×et	E
	SISATMNV	VERIFI	CALION SI	DS.	DUPL	CAIL	SISTANA			SPIKE	RECOYERY			
PARAMETER	DATE	FOUND	IRUE SA	SR	JAMS	SAMP	DUP	RPD CPS	1/2HVS	HSS	SIR	NS.	S R	BLANKS
Phanodics	5-27-88		0.250	1 05		,	٠.	,						
S	me/ne	9.276		- 0		1, 1		. 4						
	Q													
Se	5-18-88	0.049	0.050	á8	I10 (8	₹0.00	<0.003	NC	3020 8/16 Blenk	0.022		0.020	10	P) <0.003
0.003	re/me	0.053	0.050	106									A	३०२० ज्ञा ४०.००३
	O	0.049	0.050	98 ⁻	4× -	- 4		·					•	50003
							ć	•						
Toc	6-1-88	34.5	32.8	los	6214 6211.2	5.hb	2.46	0	ms 02 H	176	9 4.3	60.0	136 T	۵۰۱۶
l Barrie	. •	32,2	32.8	98	MS tmsD	176	175		W 20 W 20 W 30 W 30 W 30 W 30 W 30 W 30 W 30 W 3	175	94.3	60.0	135 135 135 135 135 135 135 135 135 135	9.1.0
٥		34·3	32.8	105		۱ نه. و	. 4							
					-									
Tox	5-23-66	1.08	, 00	801	N 026	0.114	901.0	7	·					
مد ده	he	0.869	1.00	87										
DY.	5-24-88	0.930	1.00	93	8026	0.142	0.131	00	P02 H	0.181	0.694	0.100	87	-
101 - 20.02 - 101	met.	0.841	1.00	48										1.6
\triangle				V		~ .			·					
Turbidity					ا <i>و</i> (چه	H.3	4.2	2						
٠		4							ŕ			_		
			,	. 4	1.5	,	į		< !					

RPD = [(15-D1)/((5+D)/2)] × 100
RPD = Relative Percent Difference
NC = Not calculable due to a value
less than five times the IDL

SPIKE JR = [(SSR-SR)/SA] x 100

* * Value is less than five times
the instrument detection limit
IDL = instrument Detection Limit

A = Analytical
P = Predigestion

SSR = Spiked Sample Result SR = Sample Result SA = Spiked Added

DAILY QUALITY CONTROL RAS GC LAB

DATE:	5-13-88	SPIKED VALUE (ug/L)	Analyzed Value	Recovery	Analyzed Value	\$ Recovery
	INSTRUMENT	:	B	B	6	G
TEST METHOD	COMPOUND		78778688481	**********	*********	24444444
EPA 601	EPA WP 483 CONG. 2				1	
	Chloroform	12.0	11.7	98		
	1.2-Dichiocoethane	2.0	1.9	95		
	1.1.1-Trichioroethane	1.4	2.1	150	i.	
	Carbon Tetrachionide	2.6	2.8	108		1
	Bromodichioromethane	2.0	2.3	115		
	Trichiarosthens	2.9	3.0	103		
	Dibromochiocomethane	2.6	4.5 1	173		
	Bronoform	2.9	2.2	76		
	Tetrachioroethene	1.6	1.7	106		
EPA 602	EPA - WP 879 CONC.1		D	D		
	Benzene	30.7	27.8	91	30.1	.98
	Toluene	4.1	4.7	115	5.4	132
	Ethylbenzene	11.5	9.8	85	10.5	91
	P-Xviene	19.1	20.6	108	22.5	1189
`	M-Xylene	42.5	40.1	94	42.5	100
	0-Xviene	10.6	11.1	105	12.5	118
EPA 608	Arocior 1242	(ug/g)				
	Arociar 1260	56.8				i .

^{1 -} interference

DAILY QUALITY CONTROL RAS GC LAB

DATE:	5-17-88	SPIKED VALUE (ug/L)	Anaiyzed Value	\$ Recovery	Analyzed	\$ Recovery
	LINSTRUMENT		D	D	G	G.
TEST METHOD	COMPOUND	********				
EPA 601	EPA WP 483 CONC. 2					
	Chloroform	12.0				
	1.2-Dichiorosthane	2.0				
	1.1.1-Trichiorosthane	1.4				
	Carbon Tetrachionide	2.6				
	Bromodichloromethane	2.0				
	Trichiocosthese	2.9				
	Dibromochloromethane	2.6				
	Bromoform	2.9				
	Tetrachioroethene	1.6	,			- Inde
EPA 602	EPA - WP 879 CONC.1				•	
	Benzene	30.7	26.2	85	29.4	96
	Toluene	4.1	4.0	98	4.9	120
	Ethylbenzene	11.5	9 - 1	79	10.4	90
	P-Xviene	19.1	18.6	97	21.9	115 0
	M-Xviene	42.5	38.5	90	42.3	99
	0-Xviene	10.6	10.5	99	12.6	119
EPA 608		(ug/g)				
	Arociar 1242	58.7		-		1
<u> </u>	Aroclar 1260	56.8		1		1

rt igne Name			
CLIENT NAME			
SAMPLE ID			
EPA METHOD 601	DATE: ANALYST: INSTRUMENT:	EPA METHOD 602	DATE: 5/3/36 ANALYST: AO INSTRUMENT: Ju
COMPOUND	CONCENTRATION (ug/L)	COMPOUND	CONCENTRATION (ug/L)
Chloromethane ·		Benzene	
Bromomethane		Toluene	79
Vinvi Chloride		Ethyl benzene	.']
Chloroethane	· ·	Chlorobenzene	
Methylene chloride		1.4-Dichlorobenzene	
Trichlorofluoromethan	е	1.3-Dichlorobenzene	<i>T</i> ·
1.1-Dichloroethene		1.2-Dichtorobenzene	
1.1-Dichloroethane		P-Xvlene	
Trans-1.2-Dichloroeth	ene	M-Xylene	\sim \sim
Chicroform 1.2-Dichicroethane		0-Xylene	\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \
1.1.1-Trichloroethane Carbon tetrachloride Bromodichloromethane 1.2-Dichloropropane Trans-1.3-Dichloropro Trichloroethane Dibromochloromethane 1.1.2-Trichloroethane cis-1.3-Dichloroprope 2-Chloroethylvinyl et Bromoform 1.1.2.2-Tetrachloroet Tetrachloroethylene Chlorobenzene 1.3-Dichlorobenzene 1.2-Dichlorobenzene 1.2-Dichlorobenzene 1.4-Dichlorobenzene	nene	SURROGATE RECOVER 601 Bromochloromethane 2-Bromo-1-Chioropy 1,4-Dichlorobutane 4-Bromofluorobenze 602 a,a,a,-Trifluorote	opane

LAB & PLEAGE	rt Bunk of		
CLIENT NAME			
SAMPLE ID			
EPA METHOD 601	DATE: ANALYST: INSTRUMENT:	EPA METHOD 602	DATE: 5/3/32 ANALYST: CO INSTRUMENT Sym
COMPOUND	CONCENTRATION (ug/L)	COMPOUND	CONCENTRATION (ug/L)
Chioromethane ·		Benzene	
Bromomethane		Toluene	71
Vinvi Chloride		Ethyl benzene	
Chloroethane		Chlorobenzene	
Methylene chioride	!	1.4-Dichlorobenzene	
Trichlorofluorometha	ne	1.3-Dichlorobenzene	
1.1-Dichloroethene		1.2-Dichlorobenzene	
1.1-Dichloroethane		P-Xviene	10-
Trans-1.2-Dichloroet	hen e	M-Xvlene	
Chloroform		0-Xviene	Y
1.2-Dichloroethane			
1.1.1-Trichloroethan	e		
Carbon tetrachioride		1	, 1
Bromodichioromethane		1	
1.2-Dichloropropane		SURROGATE RECOVER	IES:
Trans-1.3-Dichloronr	opene	601	į
Trichloroethene		Bromochioromethane	9
Dibromochloromethane		2-Bromo-1-Chloropi	ropane
1.1.2-Trichloroethan		1,4-Dichiorobutane	ə <u></u>
cis-1.3-Dichloroprop		4-Bromofluorobenze	ene
2-Chloroethylvinyl e	ther	1	
Bromoform		」602	
1.1.2.2-Tetrachloroe	thane	a,a,a,-Trifluorote	oluene
Tetrachioroethylene		4	· · · · · · · · · · · · · · · · · · ·
Chlorobenzene			
1.3-Dichlorobenzene		4	
1.2-Dichlorobenzene		4	
1.4-Dichlorobenzene			
; ,			,
•		1	

LAB # Sy	con Reme		
CLIENT NAME			
SAMPLE ID			
*************	***********	· · · ·	
EPA METHOD	DATE:	EPA METHOD	DATE:5/17/8
601	ANALYST:	602	ANALYST: 10
	INSTRUMENT:	,	INSTRUMENTAL)
			V.C.V.
COMPOUND	CONCENTRATION	COMPOUND	CONCENTRATION
	(ug/L)		(ug/L)
			7/
Chloromethane		Benzene	100
Bromomethane		Toluene	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,
Vinyl Chloride		Ethyl benzene	
Chloroethane		Chlorobenzene	
Methylene chloride		1.4-Dichlorobenzene	·
Trichlorofluorometha	16	1.3-Dichlorobenzene	
1.1-Dichloroethene		1.2-Dichlorobenzene	
1.1-Dichloroethane		P-Xvlene	
Trans-1.2-Dichloroet	hene	M-Xvlene	\'/
Chloroform		0-Xylene	V
1.2-Dichloroethane			
1.1.1-Trichloroethan	9]	
Carbon tetrachloride]	
Bromodichloromethane]	•
1.2-Dichloropropane		SURROGATE RECOVER	ES:
Trans-1.3-Dichloronr	opene	601	
Trichloroethene		Bromoch loromethane	
Dibromochloromethane	,	2-Bromo-1-Chloropr	opane
1.1.2-Trichloroethan	e	1,4-Dichlorobutane	
cis-1.3-Dichloroprop		4-Bromofluorobenze	ne
2-Chloroethylvinyl e	ther	4	
Bromoform		602	•
1.1.2.2-Tetrachloroe	thane	a,a,a,-Trifluoroto	luene
Tetrachioroethylene		4	
Chlorobenzene		4	
1.3-Dichlorobenzene		4	
1.2-Dichlorobenzene		4	
1.4-Dichlorobenzene			
!			
•			
•			•
		1	

	acad Bunt of		
CLIENT NAME			
SAMPLE ID	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~		
***************	*************		
EPA METHOD	DATE:	EPA METHOD	DATE: 5/17/88
601	ANALYST:	. 602	ANALYST: 4
	INSTRUMENT:	· :	INSTRUMENT:
COMPOUND	CONCENTRATION	COMPOUND	CONCENTRATION
	(ug/L)		(ug/L)
Chloromethane		Benzene	JO
Bromomethane		Toluene	
Vinvi Chloride		Ethyl benzene	
Chloroethane		Chlorobenzene	<u> </u>
Methylene chloride		1.4-Dichiorobenzene	
Trichlorofluorometha	ne	1.3-Dichiorobenzene	
1.1-Dichloroethene		1.2-Dichlorobenzene	
1.1-Dichloroethane		P-Xviene	-
Trans-1.2-Dichloroet	hene	M-Xylene	1
Chloroform		Q-Xviene	\mathcal{V}
1.2-Dichioroethane			
1.1.1-Trichloroethan	8		
Carbon tetrachloride			
Bromodichloromethane			
1.2-Dichloropropane		SURROGATE RECOVER	RIES:
Trans-1.3-Dichloropr	opene	601	
Trichloroethene		Bromochioromethan	10
Dibromochloromethane	,	2-Bromo-1-Chiorop	ropane
1.1.2-Trichloroethan	e	1,4-0 ich lorobutar	ne
cis-1.3-Dichloroprop	ene	4-Bromofluorobena	zene
2-Chioroethylvinyl e	ther		·
Bromoform		602	
1.1.2.2-Tetrachioroe	thane	a,a,a,-Trifluorot	toluene
Tetrachloroethylene			
Chlorobenzene			
1.3-Dichlorobenzene		_	
1.2-Dichlorobenzene			
1.4-Dichlorobenzene			
			`

	a material state		
	1378 BLUE		
CLIENT NAME			
SAMPLE ID			
EPA METHOD 601	DATE: ANALYST: INSTRUMENT:	EPA METHOD 602	DATE: 5/7/38 ANALYST: 01 INSTRUMENT: 01
COMPOUND	CONCENTRATION (ug/L)	COMPOUND	CONCENTRATION (ug/L)
Chloromethane		Benzene	W
Bromomethane		Toluene	
Vinyl Chioride		Ethyl benzene	
Chloroethane		Chlorobenzene	
Methylene chloride		1.4-Dichlorobenzene	
Trichlarofluorometha	ine	1.3-Dichlorobenzene	
1.1-Dichloroethene		1.2-Dichlorobenzens	
1.1-Dichloroethane		P-Xviene	
Trans-1.2-Dichloroet	hens	M-Xviene	
Chloroform		0-Xylene	V
1.2-Dichloroethane			
1.1.1-Trichioroethan	18	1	
Carbon tetrachloride]	
Bromodichioromethane			
1.2-Dichloropropane		SURROGATE RECOVER	NES:
Trans-1.3-Dichloron	opene	601	··· ·· ·· · · ·
Trichloroethene		Bromochloromethan	l e
Dibromochloromethane	<u> </u>	2-Bromo-1-Chlorop	
1.1.2-Trichloroethar		1,4-Dichlorobutan	
cis-1.3-Dichloroproc		4-Bromofluorobenz	
2-Chloroethylvinyl			
Bromoform		.602	
1.1.2.2-Tetrachloroe	thane	a,a,a,-Trifluorot	roluene
Tetrachioroethylene			· • · • • · · · · · · · · · · · · · · ·
Chiorobenzene		1	
1.3-Dichlorobenzene		1	
1.2-Dichlorobenzene		1	-
1.4-Dichlorobenzene		1	
The state of the s		†	
•			

LAB # CERG	ian Bume of 1		
CLIENT NAME			
SAMPLE ID			
*************	********	***************	
EPA METHOD	DATE:	EPA METHOD	DATE: 5/17/8
601	ANALYST:	. 602	ANALYST: 4
	INSTRUMENT:		INSTRUMENT OU
COMPOUND	CONCENTRATION	COMPOUND	CONCENTRATION
	(ug/L)		(ug/L)
Oh I a annahbana		8	1 A A ()
Chloromethane		Benzene	/ \\\\
Scomomethane Vinvl Chloride		Toluene Ethyl benzene	<u>`-</u>
Chloroethane		Chiorobenzene	
Methylene chloride	· · · · · · · · · · · · · · · · · · ·	1.4-Dichlorobenzene	
Trichiorofluoromethan		1.3-Dichlorobenzene	/
1.1-Dichioroethene		1.2-Dichlorobenzene	
1.1-Dichloroethane		P-Xviene	
Trans-1.2-0 ich loroett	\	M-Xylene	N
Chloroform	19119	0-Xviene	W
1.2-Dichloroethane		O A TIGIO	
1.1.1-Trichloroethane		1	
Carbon tetrachloride		1	
Bromodich loromethane		7	
1.2-Dichloropropane	`	SURROGATE RECOVER I	ES:
Trans-1.3-Dichloropro	pene	601	
Trichloroethene		Bromoch loromethane	
Dibromochloromethane		2-Bromo-1-Chloropr	
1.1.2-Trichloroethane	9	1,4-Dichlorobutane	
cis-1.3-Dichloronrone		4-Bromofluorobenze	
2-Chloroethylvinyl et			
Bromoform		602	
1.1.2.2-Tetrachloroe	thane	a,a,a,-Trifluoroto	luene
Tetrachioroethylene			
Chlorobenzene		_	
1.3-Dichlorobenzene		1	
1.2-Dichlorobenzene		4	
1.4-Dichlorobenzene		4	
	`		

Submitted Sys 6-28-88

Compiled

Workorder 88-05-058

Hent Geneciance Commende

Otherwise noted

				50 H			٠.	T			Ce		1	Californ			200.02701	A	PARAMETER		
				88-1-8		:	; ;	88-1-9			88-1-9	5	colones 100 m	5-30-88		C	eme	5-17-88	DAIE	SISTIMA	
		39.1	39. 3	39.2		3.68	3.84	4.04	7.58	7.48	7.86				1		0.086	0.094	FOUND	VERIFI	
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RPD = [(15-D1)/((5+D)/2)] × 100
RPD = Rejetive Percent Difference
NC = Not calculable due to a value
less than (ive times the IDL

SPIKE #R = [(SSR-SR)/SA] x 100

** Value is jess than five times

the instrument detection limit

IDL = instrument Detection Limit

A = Analytical P = Predigestion SSR = Spiked Sample Result

SR = Sample Result SA = Spiked Added

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RPD = [(IS-DI)/((S+D)/2)] × 100
RPD = Relative Percent Difference
NC = Not calculable due to a value
less than five times the IDL

SPIKE \$R = [(SSR-SR)/SA] × 100

* = Value is less than five times

the instrument detection limit

IDL = instrument Detection Limit

A = Analytical P = Predigestion

P = Predigestion SSR = Spiked Sample Result SR = Sample Result

SA = Spiked Added

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RPD = [(15-D1)/((5+D)/2)] x 100
RPD = Rejetive Percent Difference
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SPIKE \$R = [(SSR-SR)/SA] × 100 * = Value is joss than five times the instrument detection limit IDL = instrument Detection Limit

A = Analytical
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