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ATTACHMENT 1

CHRONIC TOXICITY STUDY PERFORMED ON PRE-TECT 9002 HP

Submitted to:

Mr. Brian LaPlante Calgon Corporation P.O. Box 1346 Pittsburgh, PA 15230

Prepared by:

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Biological Monitoring, Inc. P.O. Box 184 Blacksburg, VA 24063

Phone: 703-953-2821 Fax: 703-951-1481

April 27, 1995

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BIOLOGICAL MONITORING, INC.

P.O. Box 184 · Blacksburg, Virginia 24063 · Telephone 703-953-2821 · Fax 703-951-1481

STUDY PROTOCOL

Title: Chronic Toxicity Study	1111 project Number: 2372
Purpose To determine the chronic taxic	ity of Pro-Toct 9002 HP to Ceriodaphia dubia
Project Scientist(s): Anthony Smith	
Study Director: William J. Rasnake	
Name and Address of Sponsor or Test Facility:	Mr. Brian LaPlante
	Calgon Corporation
	P.O. Box 1346
	Pittsburgh, PA 15230
Proposed Starting and Completion Dates: 4/17	7 /95 - 4 / 24 / 95
Test Mode and Justification: 7 day static with	th renewal short term chronic test
4 84	۰
Test Species Information: Species: Oeriodaphni	ia dubia Age: 18-24 h
No. per Replicate: 1 Sex: F Source	BMI Strain: NA
Experimental Design Description (SOP number or	summarize and attach detailed description)
SOP # H.1.1 EPA 600/4-89/001	
·····	
Test Article Preparation: Solvents: NA	Emulsifiers: NA
Other: Ro	ute of Exposure:
Feeding Regime and Diet: NA	
Records to Be Maintained: Daily counts and	a water chemistry
Test Endpoints and Statistical Methods to be used:	NOEC and LOEC for survival and reproduction
EPA Program	
Date of Protocol Approval by Sponsor or Regulate	bry Agency: / /
Study Director: (Signature)	
mu Senior Project Con	adinates .
(Title): wanter Froject Coor	turnator

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1.0 INTRODUCTION

Calgon Corporation, Pittsburgh, Pennsylvania, contracted Biologleal Monitoring, Inc. (BMI) of Blacksburg, Virginia, to perform a short term chronic toxicity test on the product Pro-Tect 9002 HP. The <u>Ceriodaphnia dubia</u> was used as the test organism.

2.0 MATERIALS and METHODS

2.1 Dilution Water

Blacksburg, Virginia, municipal tapwater, dechlorinated, deionized and 0.2 micron filtered (BMI-DMW) was reconstituted with Evian brand mineral water and used as the dilution water. This water is BMI's standard dilution water for chronic <u>Ceriodaphnia dubia</u> testing and is used to culture and maintain the <u>Ceriodaphnia dubia</u> at BMI. Dechlorination was accomplished by activated carbon filtration. No total residual chlorine was detected in the dilution water used in this study (detection limit = 0.01 mg/L). Deionization was accomplished using 2 mixed bed deionization tanks and a Milli-Pore. Milli-Q, UV plus system (final resistivity = 18.2 megohm).

2.2 Product Sample

The product sample was labeled Pre-Tect 9002 HP. The product was a slightly viscous liquid with a pronounced odor and was received in a 125 mL amber glass container on April 5, 1995. The producted was noted to be relatively volatile. The product sample was stored in a flammable storage cabinet at room temperature (20 \pm 1°C) until use.

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2.3 Test Organism - Ceriodephnia dubia

The invertebrate used in this study was the <u>Ceriodaphnia dubia</u>. <u>Ceriodaphnia dubia</u> are continuously cultured and maintained at BMI. 18-24 hour old neonates were obtained as per BMI

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Standard Operating Procedure (Appendix *). The results of the reference toxicant test conducted during the month of the study indicated that the <u>Ceriodaphnia dubia</u> were of normal health.

2.4 Cerlodapheia dubia Toxicity Test Methods

Test concentrations for the 7 day short-term chronic definitive study were based on a four day range-finding test. (See Appendix B). This range-finding test indicated that 0, 62.5, 125, 250, 500, and 1000 mg/L would be appropriate concentrations for the definitive test. The methods used in the test conformed to the recommended guidelines specified by EPA for chronic toxicity tests (EPA/600/4-89/001) (See BMI SOP's, Appendix A).

For the definitive test, ten organisms, one organism per replicate, were randomly selected and exposed to each treatment of the definitive test. Twenty-two mL polystyrene cups, rinsed with dilution water, were used as test chambers and the test volume was 15 mL per replicate. The test temperature was 25 ± 1 °C. The test was renewed dally with a freshly prepared stock solution of the product, Pre-Tect 9002 HP. All test organisms were fed 0.1 mL YCT (See BMI SOPs F.1.1) and 0.1 mL Algae (Selanzstrum capricornutum from outside source) once per day at the renewal.

Physiochemical measurements, such as temperature, Dissolved Oxygen (mg/L), pH, and Conductivity (umbos), were made daily on all test concentrations, both before and after renewal. Alkalinity and hardness (mg/L CaCO₃) of the diluent were measured on each dilution water batch used. The number of surviving organisms, the total number of neonates produced, and time of renewal were recorded daily. (See Appendix C).

3.0 RESULTS and DISCUSSION

Table 1 presents a summary of the results of the short term chronic <u>Ceriodaphnia dubia</u> toxicity test. The 7 day NOEC (No Observed Effect Concentration) for product Pre-Tect 9002 HP, for survival and reproduction, was 1000 mg/L. This indicates that the survival and reproduction of all test concentrations were not significantly lower than survival and reproduction in the control. The chronic toxicity of Pre-Tect 9002 HP to vertebrates is not known, further testing would be required.

4.0 QUALITY CONTROL/QUALITY ASSURANCE

To ensure that the test organisms were of normal health, chronic reference toxicant tests are performed. These tests are conducted at least once a month on the fathead minnows and <u>Cerlodanhnia</u> <u>dubia</u> cultured at BMI. Fish or other organisms received from outside suppliers are also tested with a reference toxicant (usually sodium chloride) (See Appendix D). For the <u>Ceriodaphnia dubia</u> reference test performed during the month of this study, the chronic value fell well within the appropriate range of acceptability.

The Study Protocols prepared for this toxicity study are included as Page i and ii. These summarize the test conditions and protocols followed.

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Prepared by Jennifer Maloney

BIOLOGICAL MONITORING, INC. Toxicity Test Condition Summary

Clicat: Calgon Corporation

VPDES Permit #: N/A

Experiment ID# CAL041795-1

Test Organism: Ceriodaphnia dubia

Test Type: Short -term Chronic

Organism Age at Start of Test: 18-24h

Sample Tested: Pre-Tect 9002 HP

Sample Type: Compound

Sample Collection Dates and Times: Compound stock prepared daily

Sample Collector: N/A

Delivered by: Overnight Courier

No. of Organisms per Replicate: 1

Test Temperature: 25 ± 1C

Feeding Regime: 1 x Daily

Test Volume:15 mL

Test Duration: 7d

Time: 1435

Time: 1035

Test Solution Renewal Frequency: Daily

Dilution Water Used: DMW

No. of Replicates per conc.: 10

Chamber Size: 22 mL PS

Feeding prior to test: None

Photo Period: 16h light/8h dark

Start of Test: Date: 04/17/95

End of Test: Date: 04/24/95

Equipment:

pH Meter: SA 720 (c) DO Meter: YSI 58 (A) SCT Motor: YSI 33 (c) °C Measurement: Calibrated Thermomotor Salinity: SCT Meter Chlorine: Fisher/Portor Amperometric Titrator

Test Method Reference: U.S. EPA. 1989. Short-term methods for estimating the Chronic toxicity of effluents and receiving waters to freshwater organisms. EPA/600/4-89/001.

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TABLE 1 CONT.

BIOLOGICAL MONITORING, INC. Chronic Toxicity Test Data Summary

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second dependences which is second	and the part of the second secon		- Contraction of the second	The second se

Client Calgon Corporation		the log produced in the second		24
Name and Address of Street, or			N	A
Test Cercodaphnia dubra		1 Permit #		
Organism			Date	Time
Experiment	CAL041795-1			
ID		Start	04/17/95	1435
Sample	Sample Pre-Tect 9002 HD			
Tested		Ead	04/24/95	1035

RESULTS

() Genc.	lauge)					Surv	ival and Rep	production
(4)	Temp. ('C)	0.0. (my/1)	pH	Initial Alkelinity mg/1 es Caco,	Initial Hardrass mg/1 as Cx00.	Cond. (jampou)	40h / 7d	Total Offerring
62.5 125 250 500 1005	25-26 25-26 25-26 25-26 25-26 25-26 25-26	7.2-7.9 7.2-7.9 7.2-7.9 7.2-7.9 7.2-7.9 7.2-6.0 7.2-6.0	7.1-7.9 7.1-7.9 7.2-0.1 7.2-0.3 7.2-0.6 7.3-0	80	90	170-190 175-163 180-190 180-200	100/\$0 100/100 100/100 100/100	200 220 15P 202

STATISTICAL ANALYSES

Test Method		P-d D-'					
Ftsher's	Survival	Survival End Point					
Dunnett's Perioduci	LOEC= N/A						
NOTO	NOEC = 100%		LORG				
MOSC " NO ODS	erved Effect Concent:	ration LOEC = Lowert of	LUEL=N/A				
JURVIVAL DATA	7	Annual State of the State of th	served Effect Concentration				
1. No Transformati	ion was used.						
2. <u>Ceriodaphnia du</u>	bia survival in all effluer	Concentrations use					
significantly low	er than survival in the co	atrol using Fichada Tant					
REPRODUCTION	LATA	the distug Tishers Test, (p)	P0.05).				
1. No transformatic	In Was need						
2. Data PASS norm	olity tous used,		· · · · · · · · · · · · · · · · · · ·				
3. Data PASS home	arry lest using Chi-Squa	red Test.					
4. Ceriodanhaia	genetty test using Bartle	tt's Test.					
neproduction in a	bia reproduction in the al	I effluent concentrations was	s pot signification				
-oproduction in th	ne control using Dunnent	's Test, (p>0.05)	s not significantly lower than				
A							

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MATERIAL SAFETY DATA SHEET

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P.O. Box 1346 Pittsburgh, PA 15230-1346

24 Hour Emergency Telephone-(412)777-8000

Section 1. PRODUCT IDENTIFICATION PRODUCT NAME: Pre-Tect 9002 CHEMICAL DESCRIPTION: Aqueous amine solution PRODUCT CLASS: Boiler water treatment MSDS CODE: 0P04 Section 2. HAZARDOUS INGREDIENTS AND EXPOSURE LIMITS CAS % by Chemical Name Number Weight OSHA PEL ACCIH TLV Dimethylamine (DMA) TWA 10 ppm, 18 124-40-3 2 TWA 5 ppm, 9.2 me/m³ mg/m³; STEL 15 ppm, 27.6 mg/m³ Section 3. HAZARDS IDENTIFICATION ********************* . EMERGENCY OVERVIEW *** DANGERI May cause severe eye and skin damage. May be harmful if swallowed. May cause allergic skin maction. May cause respiratory tract irritation. Combustible liquid and vapor. May form suspected cancer-causing nitrosamines if mixed with nitrites. ***** PRIMARY ROUTES OF ENTRY: Eye and skin contact, inhalation, skin absorption, ingestion TARGET ORGANS: Eye, skin, lung, mucous membranes, liver MEDICAL CONDITIONS AGGRAVATED BY EXPOSURE: · Asthma Sidn disorders and allergies · Chronic respiratory disease, e.g., bronchitis, emphysema · Eye disease

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POTENTIAL HEALTH EFFECTS:

- This product may cause irreversible eye damage upon contact depending on the length of exposure, solution concentration and first aid measures. Product vapor in EYE CONTACT: low concentrations can cause tearing, conjunctivitis and corneal edema when absorbed into the tissue of the eye from the atmosphere. Corneal edema may give rise to a perception of "blue haze" or "log" around lights. The condition is transient and has no known residual effect.
- SKIN CONTACT: This product may produce burns upon contact with the skin. The severity of the burn is generally determined by the concentration of the solution and the duration of the exposure. The vapors may also be irritating to the skin. DMA may cause an allergic skin reaction and may be absorbed through the skin causing nausea, headache, and general discomfort.
- Ingestion of this product may cause severe initation or burns of the mucous INGESTION: membranes of the mouth, throat, csophagus and stomach.
- DMA vapors are initiating to the respiratory tract. Initialation of vapors may INHALATION: produce chemical pneumonitis, pulmonary edema, and delayed scarring of the airway and other affected organs. Repeated and/or prolonged exposure to vapors may cause duronic irritation of the respiratory tract, b onchogneumonia, and other adverse respiratory effects such as cough, tightness of chest, or shortness of breath.

SUBCHRONIC, CHRONIC:

DMA added to the diet of rats at 150 mg/kg for 3.5 months increased liver demethylase activity even in the presence of the enzyme inducer casein. In a subchronic study, 15 rats, 15 guinea pigs, 3 rabbits, 2. dogs, and 3 monkeys were exposed continuously by inhalation at approximately 5 ppm of DMA for 90 days. There were no deaths or signs of toxicity and all hematologic values were normal. On histopathologic examination, interstitial inflammatory changes were noted in the lungs of each species. Further, the 3 rabbits and 2 monkeys showed dilatation of the bronchi.

In a 2-year inhalation study, groups of 95 male and 95 female rats and mice were exposed 6 hours/day, 5 days/week at 10, 50, or 175 ppm of DMA. Concentration-dependent toxicity was characterized by decreased body weight (175 ppm only) and progressive inflammatory, degenerative, and hyperplastic lesions of the nasal passages. Nasal toxicity was similar in both rats and mice (no sex differences) affecting respiratory and olfactory epithelia. Lesions were severe at 175 ppm, moderate at 50 ppm, and focal and mild at 10 ppm.

CARCINOGENICITY:

NTP.

"No ingredients listed in this section" LARC:

"No ingredients listed in this section" OSHA:

"No ingredlents listed in this section"

Section 4. FIRST AID MEASURES

EYE CONTACT:

In case of contact, immediately flush eyes with plenty of water for at least 15 minutes. Seek medical aid immediately.

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SKIN CONTACT: In case of contact, immediately flush skin with plenty of water for at least 15 minutes while removing contaminated dothing and shoes. Seek medical aid immediately. Wash clothing before reuse. Destroy contaminated leather apparel. Victims with orajor skin contact should be maintained under medical observation for at least 24 hours due to the possibility of delayed reaction.

INGESTION:

If swallowed, do NOT induce vomiting. Give large quantities of water. Seek medical aid immediately. Never give anything by mouth to an unconscious person.

Note to Physicians: This product is highly injurious to all tissues, similar to that of ammonia or ammonia gas. Chemical pneumonitis, putmonary edema, laryngeal edema and delayed scarring of the airway or other affected tissues may occur following exposure. There is no specific treatment. Clinical management is based on supportive treatment, which is similar to that for thermal burns.

INHALATION: If inhaled, remove to fresh air. If not breathing, give artificial respiration. If breathing is difficult, give okygen. Seek medical aid. Prevent aspiration of vomit. Turn victim's head to the side. Assure mucous does not obstruct airway.

Section 5. FIRE-FIGHTING MEASURES

FLASH POINT: 127 °F (COC) This product is a combustible liquid.

LOWER FLAMMABLE LIMIT: Not	available UPPER FLAMMABLE LIMIT: Not available
AUTO-IGNITION TEMPERATURE	Not available
EXTINGUISHING MEDIA: Use C	D2, dry chemical, alcohol foam.
FIRE-FIGHTING INSTRUCTIONS:	Exercise caution when fighting any chemical fire. A self-contained breathing apparatus and protective clothing are essential. Use water to keep fire-exposed containers cool.
FIRE & EXPLOSION HAZARDS:	Product emits toxic gases under fire conditions. Product vapors are heavier than air and may travel a considerable distance to a source of ignition and flash back. Vapors may collect in closed spaces such as sewers, caves or closed structures.
DECOMPOSITION PRODUCTS:	Upon decomposition, ammonia vapors are liberated. the presence of sufficient oxygen, product generates. monoxide, carbon dioxide, and nitrogen oxide gases. Nitrogen oxide can react with water vapor to yield nitric acid. Combustion of product under oxygen-starved conditions can be expected to produce numerous toxic products including: nitriles, cyanic acid, isocyanates, cyanogens, nitrosamines, amides, carbamates.
NFPA RATINGS: Health = 3	Flammability = 2 Reactivity = 0 Special Hazard = None

Hazard rating scale: 0= Minimal 1= Slight 2= Moderate 3= Serious 4= Severe

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Section 6. ACCIDENTAL RELEASE MEASURES

STEPS TO BE TAKEN IF MATERIAL IS RELEASED OR SPILLED: Ventilate area of spill. Eliminate all ignition sources. Approach release from upwind. Use water spray to cool and disperse vapors, protect personnel, and dilute spills to form nonflammable mixtures. Five percent sulfuric acid may be used to neutralize diluted pools. Wearing appropriate personal protective equipment, contain spill, collect onto noncombustible absorbent like sand or earth and place into suitable container. Vapors tend to remain close to the ground and collect in out-of-the-way places. Use non-sparking blowers or ventilation facilities to remove potential explosive or toxic accumulations.

Section 7. HANDLING AND STORAGE

HANDLING: Do not get in eyes, on skin or clothing. Avoid breathing vapor or mist. Keep away from heat and flame. Use with adequate ventilation. Wash thoroughly after handling. Keep container closed when not in use. Remove all equipment which may be a source of ignition from vicinity while handling. Empty containers may contain explosive vapors. Flush empty containers with water to remove residual flammable liquid and vapors.

STORAGE: Keep away from oxidizers, heat or flames. Store away from ignition sources. Ground all containers during transfer. Store in steel containers preferably located outdoors, above ground, and surrounded by dikes to contain spills or leaks. Electrical instillations should be in accordance with Article 501 of the National Electrical Code for Class I Division 2 locations.

Section 8. EXPOSURE CONTROLS / PERSONAL PROTECTION

PERSONAL PROTECTIVE EQUIPMENT:

EYE/FACE PROTECTION: Chemical splash goggles and face shield SKIN PROTECTION: Chemical resistant gloves and protective clothing RESPIRATORY PROTECTION: If airborne concentrations exceed published exposure limits, use a NIOSH approved respirator in accordance with OSHA respiratory protection requirements (29 CFR 1910.134).

ENGINEERING CONTROLS: Local exhaust ventilation may be required in addition to general room ventilation to maintain airborne concentrations below exposure limits.

WORK PRACTICES: Eye wash station and safety shower should be accessible in the immediate area of use.

UNSATISFACTORY MATERIALS OF CONSTRUCTION: DMA corrodes copper, aluminum, zinc, and galvanized surfaces.

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Section 9. PHYSICAL AND CHEMICAL PROPERTIES

BOILING POINT: Not available

VAPOR PRESSURE: 2 torr @ 10 °C (for DMA)

VAPOR DENSITY (air = 1): 1.55 (for DMA)

% VOLATILE BY WEIGHT: 100

SOLUBILITY IN WATER: Complete SPECIFIC GRAVITY: No: available

pH: Basic

FREEZING POINT: Not available

AFPEARANCE AND ODOR: Clear, colorless liquid with ammoniacal/fishy odor.

Section 10. STABILITY AND REACTIVITY

CHEMICAL STABILITY: Stable . HAZARDOUS POLYMERIZATION: Will not occur

CONDITIONS TO AVOID: Keep away from heat and flame.

INCOMPATIBILITY: Strong oxidizers, acids, copper, aluminum, zinc and galvanized surfaces.

DECOMPOSITION PRODUCTS: Upon decomposition, ammonia vapors are liberated. Upon combustion in the presence of sufficient oxygen, product generates harmful carbon monoxide, carbon dioxide, and nitrogen oxide gases. Nitrogen oxide can react with water vapor to yield nitric acid. Combustion of product under oxygen-starved conditions can be expected to produce numerous toxic products including: nitriles, cyanic acid, isocyanates, cyanogens, nitrosamines, amides, carbamates.

Section 11. TOXICOLOGICAL INFORMATION

ON PRODUCT:

See the following information on active ingredient.

ON INGREDIENTS:

Chemical Name	Chemical Name (rat)	Dermal LD ₅₀ (rabbit)	Inhalation LC ₅₀ (rat)
Dimethylamine (DMA)	698 mus/kg	Not available	4540 ppa/6H

Section 12. ECOLOGICAL INFORMATION

ON PRODUCT:

Aquatic toxicity data on a 10% solution of DMA: 48 hr LC_{50} (Daphnia magna): 675 ppm 96 hr LC_{50} (fathead minnow): > 1000 ppm

96 hr LC50 (bluegill sunfish): > 1000 ppm

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Section 13. DISPOSAL CONSIDERATIONS

RCRA STATUS: The EPA Hazardous Waste Number is U092.

DISPOSAL: Dispose of in accordance with local, state and federal regulations. Incineration is acceptable and the preferred method of disposal. However, nitrogen oxide emission controls may be required to meet specifications. Chemical and/or biological degradation is feasible. A suitable industrial or municipal waste treatment system can be used depending on the quality and quantity of waste to be treated, the treatment plant capability, and discharge water quality standards. Incinerate in an open container. Do not dump into municipal sewers or enclosed drains that present a fire or explosion hazard.

Section 14. TRANSPORT INFORMATION

DOT CLASSIFICATION: Class/Division: 3 Proper Shlpping Name: Flammable liquid, corrosive, n.o.s. (Dimethylamine) Label: Flammable liquid, Corrosive Packing Group: III ID Number: UN 2924

Section 15. REGULATORY INFORMATION

OSHA Hazard Communication Status: Hazardous

TSCA: The ingredients of this product are listed on the Toxic Substances Control Act (TSCA) Chemical Substances Inventory.

CERCLA reportable quantity of EPA hazardous substances in product:

Chemical Name			-	RO
Dimethylamine (DMA)	4	х.		1000 fb
	1.00			

Product RQ: 50,000 lb

(Notify EFA of product spills exceeding this amount.)

SARA TITLE III:

Section 302 Extremely Hazardous Substances:

Section 311 and 312 Health and Physical Hayards:

Chemical Name	CAS #	RO	TPO
"No ingredients listed in this section"			

Immediate [yes]	Delayed [yes]	Fire [yes]	Pressure [no]	Reactivity [no]
Section 313 Toxic Chemica	da:			
Chemical Name	the contract	CAS #		% by Weight

"No ingredients listed in this section"

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Section 16. OTHER INFORMATION

HMIS RATINGS: Health = 3* Flammability = 2 Reactivity = 0 Personal Protective Equipment = X (to be specified by user depending on use conditions)

"There are potential chronic health effects to consider.

Hazard rating scale: 0 = Minimal 1= Slight 2= Moderate 3 = Serious 4 = Severe :

MSDS REVISION SUMMARY: Not applicable

While this information and recommendations set forth herein sie believed to be accurate as of the date hereof, CALGON CORPORATION MAKES NO WARRANTY WITH RESPECT MERETO AND DISCLAIMS ALL LIABILITY FROM RELIANCE THEREOM.

PREPARED BY: P.J. Maloney

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Commissioner: Douglas E. Bryant

Board John H Llurriss, Chairman Sandra J Molander, Secretary

Promoting Health, Protecting the Environment

Richard F. Jabbour, DDS William M. Hull, Jr., MD. Roger Leaks, Jr.

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June 28, 1995	ELMIRONIMERTAL PROTECTION SECTION				
	JUL 0 3 1995				
	D	TICKLER DATE			
		СОРҮ			
		ROUTE			
Maintenance Chemical Request Duke PWR/Oconee Nuclear Stati	on				

Re: Oconee County

Dear Mr. Wylie:

Mr. Robert R. Wylie Duke Power Company

13339 Hagers Ferry Rd. Huntersville, NC 28078-7929

Electric System Support Department

This letter is in regards to your May 30, 1995 request to use Calgon Corporation's Pre-Tect 9002. HP in Oconee Nuclear Station's feedwater system. Based upon the results of the toxicity tests you have submitted, use of this product is approvable.

This Office approves the use of Pre-Tect 9002 HP, with the condition that the concentration does not exceed the NOEC of 1000 ppm at Outfall(s) 002 and 004.

Should you have any questions, please contact me at 734-5248.

Sincerely,

William C. Batts

William C. Botts Industrial & Agricultural Wastewater Division Bureau of Water Pollution Control

cc: George Tomlin, App I EQC Vernon Beaty, WQAE