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Date: September 22, 1997
Subject: Calendar of "events"
Pages (including cover): 3

Swenberg Laboratories 1997-1998

Sept. 97:	εG methods development continues DNA from 4 wk whole livers
Oct. 97:	1 st analyses rat whole liver 4 week, all doses start εA analyses in urines - <i>one response</i>
Nov. 97:	¹³ C whole liver DNA in εG method DNA from 4 wk heps and NPCs urine analyses
Dec. 97:	urine analyses ¹³ C whole liver DNA in εG method 4 wk heps and NPCs DNA in εG method
Jan. 98:	Urine analyses Abasic site measurement 4 wk whole liver ¹³ C heps and NPC DNA
Feb. 98:	Urine analyses Abasic site measurement 4 wk whole liver ¹³ C heps and NPC DNA in εG method 32P postlabeling analyses of εA and εC 4 wk whole liver
Mar. 98:	Urine analyses Abasic site measurement 4 wk whole liver ¹³ C heps and NPC DNA in εG method 32P postlabeling analyses of εA and εC 4 wk whole liver 1,N ² -εG analyses start
Apr. 98:	Abasic site measurement 4 wk whole liver ¹³ C heps and NPC DNA in εG method 32P postlabeling analyses of εA and εC continue 1,N ² -εG analyses
May 98:	Abasic site measurement 4 wk whole liver 32P postlabeling analyses of εA and εC 1,N ² -εG analyses
June 98:	MPG & AP endonuclease assays begin Abasic site measurement 4 wk whole liver 32P postlabeling analyses of εA and εC 1,N ² -εG analyses

July 98:	MPG & AP endonuclease assays begin Abasic site measurement 4 wk whole liver 32P postlabeling analyses of ϵ A and ϵ C 1,N ² - ϵ G analyses
Aug. 98:	MPG & AP endonuclease assays begin Abasic site measurement 4 wk whole liver 32P postlabeling analyses of ϵ A and ϵ C 1,N ² - ϵ G analyses
Year 2:98-99	Begin with brain
Year 3:99-00	Lung, Kidney, risk assessment analyses begin

Molecular Dosimetry of Vinyl Chloride

James A. Swenberg, D.V.M., Ph.D.

Research Report and Budget Request

July 28, 1997

Research Report

Year 1 (1996-1997 Budget)

The laboratory portion of this project began this spring by sending teams of 3-4 of our laboratory personnel to New Jersey to collect tissues and liver cell types from rats that were exposed to 0, 10, 100, or 1100 ppm vinyl chloride (VC) for 1-20 days. Procedures were rather difficult during the first set of necropsies, but went much better for subsequent procedures after we sent an additional person and shipped up our own centrifuge. We were able to harvest good yields of cells and tissues for the experiment as a whole. It was unfortunate that the [^{14}C]-VC experiment was the first to come off, as this was the least smooth running of all. Even so, adequate numbers of cells were able to be harvested to meet our objectives.

We are currently isolating DNA from the 4-week hepatocytes, as these are the most plentiful cells. A new immunoaffinity chromatograph-GC/MS method is replacing our previous low pressure strong cation exchange-GC/MS method for the analysis of ethenoguanine. After this shakedown, we will begin working with the [^{14}C]-VC liver cells. Parallel to this effort will be the analysis of urines for excretion of ethenoadenine using immunoaffinity chromatography and LC-MS. Likewise, we will begin examining the major DNA repair pathway for VC DNA adducts including methyl purine glycosylase (MPG) and AP-endonuclease (AP).

Year 2 (1997-1998 Budget)

During Year 2 we will continue to work up the many samples collected in New Jersey. Priority will be given to establishing exposure response relationships, identifying cell-specific differences in DNA repair, determining effects of exposure on endogenously formed DNA adducts, and doing initial studies related to incorporating these data into a PB/PK risk assessment model. We are also developing new methods for the $^1\text{N},2$ -ethenoguanine adduct, which has never been characterized *in vivo*.

Year 3 (1998-1999)

We expect to complete this research during the third year. At this time, we will have analyzed all of the relevant tissues and cells for DNA adducts, will establish the role of defects in DNA repair and the effect of induction of DNA repair on exogenous and endogenous DNA adducts, and incorporate these data into a biologically-based risk assessment model.

Research Budget

My understanding of the funding available for this project is that between \$50,000-75,000 will be available per year for three years. The cost of this research project will greatly exceed this, with personnel costing ~\$140,000 this year. However, since VC has been and continues to be one of the chemicals that we are investigating under our NIEHS Superfund Basic Research Program Project, I can offset part of the cost, providing that the CMA agrees to share acknowledgement with NIEHS. Should we need additional [^{14}C]-VC exposures, they can be done at UNC, since the EOIC has agreed to fund new research that includes funds to set up a small nose-only exposure system for similar studies on ethylene oxide. Thus, we can maximize our future research dollars to fund critical studies. We need to move forward as fast as possible, as the EPA has already developed a new risk assessment on VC that suggests increased risk compared to their previous estimates. The new assessment incorporates metabolism data, but not information on DNA adducts or DNA repair. Finally, we incurred major expenses during the four trips to New Jersey that were not included in Year 1's budget. These are listed under sample collection in the budget.

1997-1998 Budget request

Sample collection		\$20,695
Travel expenses	\$8172	
Shipping	752	
Supplies	11771	
Research budget		\$66,500
Personnel	35000	
Supplies	24000	
LC-MS/MS time (\$25/hr)	2500	
Travel	3000	
Equipment maintenance agreements	2000	
TOTAL BUDGET REQUEST		\$87,195

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To: Bob Venezia

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From: James A. Swenberg, D.V.M., Ph.D.

Date: May 7, 1996

Bob:

Here is the research proposal/protocol. I made a few changes from our discussions in New Jersey. I sent a copy to Jim Knaak for his comments. He said that the issue of dynamic versus static exposures will be determined in the future. Dr. Rudy Jager has some dynamic exposure systems that will use even less than the 27 gm. As I do not have access to the needed information, I cannot go any farther today. I hope that this is what you need. If you have any questions, please give me a call tomorrow.

PROTOCOL FOR MOLECULAR DOSIMETRY STUDIES ON VINYL CHLORIDE

James A. Swenberg, D.V.M., Ph.D.

University of North Carolina

May 7, 1996

Listed below is a series of studies on the formation and repair of DNA adducts induced by vinyl chloride (VC) that will provide a vastly improved understanding of the mechanisms of vinyl chloride carcinogenesis. These studies will provide information relevant to cancer risk assessments of vinyl chloride. The studies would be conducted in a manner that will complement the two generation reproductive and developmental toxicity studies that will be conducted at Huntingdon Life Sciences Laboratory.

Dose-response Studies

Presently, we have data showing that preweanling rats are more susceptible than adults to VC carcinogenesis, that preweanlings develop 3-fold greater numbers of VC DNA adducts, and that preweanlings have 3-fold higher expression of CYP 2E1. The adduct and carcinogenesis data are all from "high" exposures of ~500 ppm. No information exists on lower exposures. In addition, we have clearly demonstrated that DNA adducts identical to those formed by VC are formed endogenously in unexposed rats, mice and humans. We have developed ultrasensitive and highly specific assays for the DNA adducts of VC. These include immunoaffinity/³²P-postlabeling methods for 1,N⁶-ethenodeoxyadenosine (EdA) and 3,N⁴-ethenodeoxycytidine (EdC), and GC/MS methods for N²,3-ethenoguanine (EG). By adding animals to the reproductive and developmental toxicity studies, we will be able to assess the effect of exposure concentration on 1) the molecular dose of etheno adducts, the primary promutagenic DNA adducts of VC; 2) determine if there are differences in the dose-response between adult and weanling rats; 3) examine the major DNA repair pathways for VC DNA adducts; and 4) characterize the utility of using DNA adducts excreted in the urine as a biomarker of exposure.

Molecular Dosimetry Studies in Nonparenchymal Cells (NPC) versus Hepatocytes

Understanding why VC targets the endothelial cell for its carcinogenic effect is critical for proper risk assessment. We have preliminary data on EG in vinyl fluoride-exposed rats (2500 ppm, 6 hr/day, 5 days/wk, 4 wks) which shows that the number of adducts is ~2.5-fold higher in NPC than hepatocytes, even though metabolism of VF to its electrophile is thought to only take place in hepatocytes. Since the NPC are exposed by diffusion of VC or VF electrophiles from neighboring hepatocytes and therefore have lower exposures, this suggested that NPC were deficient in DNA repair activity for EG. To test this hypothesis, we have developed a RT-PCR method for quantitating methylpurine-DNA glycosylase (MPG) in tissues and cells. Preliminary data have shown that NPC have <20% of the MPG activity present in hepatocytes. In addition, we have shown that NPC have 5-fold greater numbers of abasic sites in their DNA. Abasic sites are formed when MPG excises a damaged base, such as EG, EA and EC. These data suggest that NPC are deficient in a second step in the DNA repair of VC adducts. We think that the most likely candidate is a deficiency in AP endonuclease. The difference between hepatocyte and NPC abasic sites was seen in rats exposed to VF, but not in controls. The combination of two defects in DNA repair in the target cells for VC carcinogenesis provides a strong scientific basis for the induction of hemangiosarcomas by this important chemical. In view of the fact that the increase in abasic sites was only seen in highly exposed rats shows the need to understand the dose response relationship for this endpoint. By conducting 4 week exposures to 0, 10, 100, and 1000 ppm VC and measuring etheno adducts, MPG expression and abasic sites, we will be able to determine if nonlinearities exist in the molecular dose of VC, in MPG expression, and in the number of abasic sites present in the DNA. Our hypothesis is that EG will show a supralinear dose response; that MPG will be similar in exposed and unexposed hepatocytes, but lower in both exposed and unexposed NPC; but that abasic sites will exhibit a sublinear dose response characteristic of saturation of DNA repair in NPC, but not hepatocytes.

Studies Using [$^{13}\text{C}_2$]-VC

Our data showing endogenous formation of etheno DNA adducts that are identical to those induced by VC have important implications. We had previously published a paper that concluded that the DNA adducts of VC were highly persistent. We now believe that this was not correct, in that what we were calling persistent DNA adducts were actually the endogenously formed adducts at steady-state. By utilizing [$^{13}\text{C}_2$]-VC, the formation and repair of VC-induced DNA adducts can be studied relative to those adducts formed endogenously. I believe that these studies will critically impact on any risk assessment dealing with low exposures, such as might be associated with accidental releases that reach the fence line.

We will examine the amount of [$^{13}\text{C}_2$]-VC induced EG in rats exposed to 10, 100, or 1000 ppm for 5 days (6 hrs/day). These same animals will be housed in metabolism cages for 3 days prior to exposure and during each day's post exposure holding period (days 1-4) so that urine can be collected. The urine will be analyzed for endogenous and induced DNA adducts that have been repaired by the MPG pathway using GC/MS. We will also develop immunoaffinity-LC/MS methods for all three etheno bases. In addition, it may be possible to identify and quantitate [$^{13}\text{C}_2$]-VC metabolites in the urine using NMR. A second set of [$^{13}\text{C}_2$]-VC exposures is contemplated that would consist of single exposures. These will be designed after we get data from the above study.

Significance

We have developed highly sensitive and specific assays for the promutagenic DNA adducts of vinyl chloride over the past 10 years. By exposing additional animals in the Huntingdon study, we can use these methods to answer several questions related to improved risk assessment of VC. First, the molecular dosimetry data will be the only information of its type that addresses dose-response. All other data have come from high exposures of 500-600 ppm. Second, we will address the issue of differences between weanling and adult animals. While we already know that preweanling rats are more sensitive than adults to the formation of DNA adducts and carcinogenesis at high concentrations, it is not known if this is true for low exposures. Since the only VC exposures that children encounter would be extremely low, the 10 ppm data are of particular importance. Just

because preweanling rats exposed to concentrations that saturate metabolism develop more adducts does not necessarily mean that young animals exposed to low concentrations will also develop more DNA adducts than adults. Our recent development of an assay for MPG and the finding that NPC are deficient in this DNA repair pathway provides the first mechanistic data that explains why VC causes hemangiosarcomas. The additional finding of greatly increased numbers of abasic sites in the NPC DNA further highlights the importance of cell-specific differences in DNA repair. The experiments outlined above will provide compelling data on the importance of DNA repair in VC carcinogenesis. Finally, the use of $[^{13}\text{C}_2]$ -VC will allow us to differentiate between induced and endogenous DNA adduct formation. If large numbers of induced EG are formed, they may lead to increased numbers of endogenous adducts due to competition for DNA repair. On the other hand, we will be able to determine the extent of increased DNA damage relative to endogenous adducts for all three exposure groups. For example, we might find that 10 ppm VC for 5 days only doubles the amount of EG present endogenously. Such a finding should have great impact on low dose risk assessments. We would then look at various single exposures to simulate fenceline exposure scenarios to determine if there was any detectable increase in risk. Similar studies could be used to mimic occupational or environmental exposures to VC.

Protocols

Study I. Adult Rats Exposed to Vinyl Chloride for 1 Week

Route: Whole body exposure conducted during the two generation animal portion of the main study

Exposure groups: 0, 10, 100, or 1000 ppm

No. of rats/group: 32

Total: 128

Exposure duration: 1 week (6 hr/day, 5 days)

Species/strain: CD Rat

Age: Approximately 12 weeks of age

Sex: Males

Sacrifice: 8 rats/ group at end of last exposure (non-perfused), and 8 rats/ group at end of last exposure and at 3 and 7 days post-exposure (perfused)

Urine collections: during Days 1-4 of exposure (collected during nonexposure times)

Study II. Weanling Rats Exposed to Vinyl Chloride for 1 Week

Route: Whole body exposure conducted during the two generation animal portion of the main study

Exposure groups: 0, 10, 100, or 1000 ppm

No. of rats/group: 32

Total: 128

Exposure duration: 1 week (6 hr/day, 5 days/week)

Species/strain: CD Rat

Age: Approximately 21 days of age

Sex: Males

Sacrifice: 8 rats/ group at end of last exposure (non-perfused), and 8 rats/ group at end of last exposure and at 3 and 7 days post-exposure (perfused)

Urine collections: during Days 1-4 of exposure (collected during nonexposure times)

Study III. Adult Rats Exposed to Vinyl Chloride for 4 Weeks

Route: Whole body exposure conducted during the two generation animal portion of the main study

Exposure groups: 0, 10, 100, or 1000 ppm

No. of rats/group: 32

Total: 128

Exposure duration: 4 weeks (6 hr/day, 5 days/week)

Species/strain: CD Rat

Age: Approximately 12 weeks of age

Sex: Males

Sacrifice: 8 rats/ group at end of last exposure (non-perfused), and 8 rats/ group at end of last exposure and at 3 and 7 days post-exposure (perfused)

Urine collections: during last week of exposure (collected during nonexposure times)

Study IV. Adult Rats Exposed to [¹³C₂]-Vinyl Chloride for 1 Week

Route: Nose only exposure

Exposure groups: 10, 100, or 1000 ppm

No. of rats/group: 8

Total: 24

Exposure duration: 5 days (6 hr/day)

Species/strain: CD Rat

Age: Approximately 12 weeks of age

Sex: Males

Sacrifice: 8 rats/ group at end of last exposure (perfused)

Urine collections: 3 days prior to exposure and Days 1-4 during exposure (collected during nonexposure times)