

NCTR PROTOCOL E0219001

**TWO YEAR CHRONIC TOXICOLOGY STUDY OF BISPHENOL A (BPA) [CAS # 80-05-7]
ADMINISTERED BY GAVAGE TO SPRAGUE-DAWLEY RATS (NCTR) FROM GESTATIONAL DAY 6
UNTIL BIRTH AND DIRECTLY TO F₁ PUPS FROM POSTNATAL DAY (PND) 1; CONTINUOUS AND
STOP DOSE (PND 21) EXPOSURES**

STATISTICAL REPORT

STATISTICAL ANALYSIS OF TERMINAL SACRIFICE SURVIVAL DATA

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FOR

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Statistical Analysis of Terminal Sacrifice Survival Data

Objectives

1.1 Project Objectives

The goal of this two year chronic study is to characterize the long term toxicity of orally administered BPA, including developmental exposure, in the NCTR Sprague-Dawley (CD) rat over a broad dose range.

1.2 Analysis Objectives

The goal of this analysis is to evaluate the effects of exposure to BPA in Sprague-Dawley rats on 2 year terminal sacrifice survival.

Experimental Design

The study design consisted of first generation female and male rats (F₀) for up to 600 mating pairs randomized to treatment groups in 5 litters. The goal of the F₀ matings was to obtain 352 study litters, 50 per dose group for vehicle controls and five BPA dose groups, 2.5, 25, 250, 2500, and 25000 µg/kg bw/day, and 26 for each of two EE₂ dose groups, 0.05 and 0.5 µg/kg bw/day. Dams were dosed daily from gestation day (GD) 6 until parturition. Dosing was by gavage for F₀ dams and F₁ pups, the second study generation. Litters were culled to 10 pups on PND 1. There were two study dosing arms of F₁ animals, daily continuous dosing to termination, and daily dose stopped at post-natal day (PND) 21. There was a vehicle control group and five BPA groups for each study dosing arm, and EE₂ daily dose groups for the continuous dosing arm only. From the F₁ litters, pups were allocated at weaning, PND 21, to the interim (1 year) and terminal (2 year) sacrifices for the core study. For vehicle and BPA terminal sacrifice groups, there were 50 pups each; for the interim sacrifice and the EE₂ terminal sacrifice groups, there were 20-26 pups each. Pups within litter and sex were assigned to different dosing arms and sacrifice times.

Survival Data

For this analysis, survival is followed from weaning to 2 years for the animals allocated to 2 year terminal sacrifice.

Statistical Methods

Statistical analyses were performed separately for the BPA study arms, stop dose and continuous dose, and for the EE₂ continuous dose. Animals with a disposition observed as dead or moribund were treated as uncensored observations, while those observed as reaching terminal sacrifice were considered censored.

Cox proportional hazards regression analysis was performed to compare survival of treatment groups to the control group. The survival time of each member of a population is assumed to follow its own hazard function. In Cox regression, the hazard functions of any two groups are assumed to be proportional at any particular time. Multiple comparisons of treatments to the vehicle control group were adjusted using Holm's (step-down Bonferroni) method, and all tests were performed as two-sided. Test of dose trend, increasing effect of treatment with increasing dose, was performed for the BPA and vehicle control groups.

For analysis of each endpoint, a sensitivity analysis was also performed. During initial preweaning of animals, 263 core study 2 year terminal sacrifice animals (56 in vehicle control, 175 in BPA

2.5, 25, 250, 2500, and 25000 µg/kg bw/day, and 32 in EE₂ µg/kg bw/day dose groups) were held in the same rooms as a special BPA 250,000 µg/kg bw/day high dose requested by an academic laboratory. In consultation with the Principal Investigator, to address the possibility of inadvertent exposure of the core study animals, a sensitivity analysis excluding these 263 animals was also performed to test the robustness of the results. Additional statistically significant pairwise comparisons from the sensitivity analysis are reported in the text.

Results

Results of analyses using all study animals are presented in Appendix A for Tables and in Appendix B for Figures.

4.1 BPA Treatments Stop Dose Arm

Disposition counts and proportions for the BPA stop dose arm are presented in Table 1 for females and in Table 2 for males.

The results of the proportional hazards model analysis for the BPA stop dose arm are presented in Table 3 for females and in Table 4 for males. Dose trend and hazard ratios of treatment groups to the vehicle control were not significant for females or males.

In the sensitivity analyses for the BPA stop dose arm, there were no statistically significant results for females or males.

4.2 BPA Treatments Continuous Dose Arm

Disposition counts and proportions for the BPA continuous dose arm are presented in Table 5 for females and in Table 6 for males.

The results of the proportional hazards model analysis for the BPA continuous dose arm are presented in Table 7 for females and in Table 8 for males. Dose trend and hazard ratios of treatment groups to the vehicle control were not significant for females or males.

In the sensitivity analyses for the BPA continuous dose arm, there were no statistically significant results for females or males.

4.3 EE₂ Treatments Continuous Dose

Disposition counts and proportions for the EE₂ continuous dose are presented in Table 9 for females and in Table 10 for males.

The results of the proportional hazards model analysis for the EE₂ continuous dose are presented in Table 11 for females and in Table 12 for males. Hazard ratios of treatment groups to the vehicle control were not significant for females or males.

In the sensitivity analyses for the EE₂ continuous dose, there were no statistically significant results for females or males.

Conclusions

5.1 BPA Treatments Stop Dose Arm

There were no significant differences in survival for treatments in the BPA stop dose arm compared to vehicle control for females or males.

5.2 BPA Treatments Continuous Dose Arm

There were no significant differences in survival for treatments in the BPA continuous dose arm compared to vehicle control for females or males.

5.3 EE₂ Treatments Continuous Dose

There were no significant differences in survival for EE₂ continuous dose treatments compared to vehicle control for females or males.

Appendices

A. Statistical Tables

a) *BPA Treatments Stop Dose Arm*

Table 1. Disposition and Censoring of Animals for Terminal Sacrifice Female Bisphenol-A Stop Dose Arm

<i>Dose ($\mu\text{g}/\text{kg}\cdot\text{BW}/\text{day}$)</i>	<i>N</i>	<i>Dead</i>	<i>Moribund</i>	<i>Terminal Sacrifice</i>	<i>Censored</i>	<i>Uncensored</i>	<i>Proportion Censored¹</i>
0	50	3	36	11	11	39	0.220
2.5	50	6	32	12	12	38	0.240
25	48	3	32	13	13	35	0.271
250	50	2	35	13	13	37	0.260
2500	50	3	30	17	17	33	0.340
25000	46	2	31	13	13	33	0.283

¹ Uncensored animals include those that were moribund or dead; censored animals include those that reached terminal sacrifice.

Table 2. Disposition and Censoring of Animals for Terminal Sacrifice Male Bisphenol-A Stop Dose Arm

<i>Dose ($\mu\text{g}/\text{kg}\cdot\text{BW}/\text{day}$)</i>	<i>N</i>	<i>Dead</i>	<i>Moribund</i>	<i>Terminal Sacrifice</i>	<i>Censored</i>	<i>Uncensored</i>	<i>Proportion Censored¹</i>
0	50	13	20	17	17	33	0.340
2.5	48	12	20	16	16	32	0.333
25	48	8	24	16	16	32	0.333
250	50	8	29	13	13	37	0.260
2500	50	8	27	15	15	35	0.300
25000	46	8	29	9	9	37	0.196

¹ Uncensored animals include those that were moribund or dead; censored animals include those that reached terminal sacrifice.

Table 3. Cox Proportional Hazards Analysis for Terminal Sacrifice Female Bisphenol-A Stop Dose Arm

<i>Dose ($\mu\text{g}/\text{kg}_{\text{BW}}/\text{day}$)¹</i>	<i>Hazard Ratio²</i>	<i>P-value³</i>
0	-	0.203
2.5	1.038	1.000
25	0.971	1.000
250	0.883	1.000
2500	0.756	1.000
25000	0.863	1.000

¹ P-value for dose trend is shown for vehicle control.

² Hazard ratios are relative to vehicle control.

³ P-values for dose comparisons to control are adjusted using Holm's method.

Table 4. Cox Proportional Hazards Analysis for Terminal Sacrifice Male Bisphenol-A Stop Dose Arm

<i>Dose ($\mu\text{g}/\text{kg}_{\text{BW}}/\text{day}$)¹</i>	<i>Hazard Ratio²</i>	<i>P-value³</i>
0	-	0.053
2.5	1.096	1.000
25	1.032	1.000
250	1.473	0.424
2500	1.081	1.000
25000	1.628	0.209

¹ P-value for dose trend is shown for vehicle control.

² Hazard ratios are relative to vehicle control.

³ P-values for dose comparisons to control are adjusted using Holm's method.

b) BPA Treatments Continuous Dose Arm

Table 5. Disposition and Censoring of Animals for Terminal Sacrifice Female Bisphenol-A Continuous Dose Arm

<i>Dose (µg/kg_{BW}/day)</i>	<i>N</i>	<i>Dead</i>	<i>Moribund</i>	<i>Terminal Sacrifice</i>	<i>Censored</i>	<i>Uncensored</i>	<i>Proportion Censored¹</i>
0	50	6	28	16	16	34	0.320
2.5	48	1	28	19	19	29	0.396
25	46	1	31	14	14	32	0.304
250	49	5	31	13	13	36	0.265
2500	50	7	33	10	10	40	0.200
25000	46	3	35	8	8	38	0.174

¹ Uncensored animals include those that were moribund or dead; censored animals include those that reached terminal sacrifice.

Table 6. Disposition and Censoring of Animals for Terminal Sacrifice Male Bisphenol-A Continuous Dose Arm

<i>Dose (µg/kg_{BW}/day)</i>	<i>N</i>	<i>Dead</i>	<i>Moribund</i>	<i>Terminal Sacrifice</i>	<i>Censored</i>	<i>Uncensored</i>	<i>Proportion Censored¹</i>
0	50	11	24	15	15	35	0.300
2.5	48	16	16	16	16	32	0.333
25	48	4	27	17	17	31	0.354
250	50	15	21	14	14	36	0.280
2500	50	10	24	16	16	34	0.320
25000	46	8	27	11	11	35	0.239

¹ Uncensored animals include those that were moribund or dead; censored animals include those that reached terminal sacrifice.

Table 7. Cox Proportional Hazards Analysis for Terminal Sacrifice Female Bisphenol-A Continuous Dose Arm

<i>Dose ($\mu\text{g}/\text{kg}_{\text{BW}}/\text{day}$)¹</i>	<i>Hazard Ratio²</i>	<i>P-value³</i>
0	-	0.071
2.5	0.860	1.000
25	1.209	1.000
250	1.127	1.000
2500	1.467	0.502
25000	1.225	1.000

¹ P-value for dose trend is shown for vehicle control.

² Hazard ratios are relative to vehicle control.

³ P-values for dose comparisons to control are adjusted using Holm's method.

Table 8. Cox Proportional Hazards Analysis for Terminal Sacrifice Male Bisphenol-A Continuous Dose Arm

<i>Dose ($\mu\text{g}/\text{kg}_{\text{BW}}/\text{day}$)¹</i>	<i>Hazard Ratio²</i>	<i>P-value³</i>
0	-	0.327
2.5	0.892	1.000
25	0.933	1.000
250	1.034	1.000
2500	0.985	1.000
25000	1.221	1.000

¹ P-value for dose trend is shown for vehicle control.

² Hazard ratios are relative to vehicle control.

³ P-values for dose comparisons to control are adjusted using Holm's method.

c) *EE₂ Treatments Continuous Dose*

Table 9. Disposition and Censoring of Animals for Terminal Sacrifice Female Ethinyl Estradiol Dose

<i>Dose</i> ($\mu\text{g}/\text{kg}\cdot\text{BW}/\text{day}$)	<i>N</i>	<i>Dead</i>	<i>Moribund</i>	<i>Terminal Sacrifice</i>	<i>Censored</i>	<i>Uncensored</i>	<i>Proportion Censored</i> ¹
0	50	6	28	16	16	34	0.320
0.05	26	1	18	7	7	19	0.269
0.5	26	4	18	4	4	22	0.154

¹ Uncensored animals include those that were moribund or dead; censored animals include those that reached terminal sacrifice.

Table 10. Disposition and Censoring of Animals for Terminal Sacrifice Male Ethinyl Estradiol Dose

<i>Dose</i> ($\mu\text{g}/\text{kg}\cdot\text{BW}/\text{day}$)	<i>N</i>	<i>Dead</i>	<i>Moribund</i>	<i>Terminal Sacrifice</i>	<i>Censored</i>	<i>Uncensored</i>	<i>Proportion Censored</i> ¹
0	50	11	24	15	15	35	0.300
0.05	26	3	14	9	9	17	0.346
0.5	26	4	10	12	12	14	0.462

¹ Uncensored animals include those that were moribund or dead; censored animals include those that reached terminal sacrifice.

Table 11. Cox Proportional Hazards Analysis for Terminal Sacrifice Female Ethinyl Estradiol Dose

<i>Dose</i> ($\mu\text{g}/\text{kg}\cdot\text{BW}/\text{day}$)	<i>Hazard Ratio</i> ¹	<i>P-value</i> ²
0.05	1.275	0.396
0.5	1.584	0.188

¹ Hazard ratios are relative to vehicle control.

² P-values for dose comparisons to control are adjusted using Holm's method.

Table 12. Cox Proportional Hazards Analysis for Terminal Sacrifice Male Ethinyl Estradiol Dose

<i>Dose</i> ($\mu\text{g}/\text{kg}\cdot\text{BW}/\text{day}$)	<i>Hazard Ratio</i> ¹	<i>P-value</i> ²
0.05	0.956	0.879
0.5	0.672	0.419

¹ Hazard ratios are relative to vehicle control.

² P-values for dose comparisons to control are adjusted using Holm's method.

B. Figures

Figure 1. Kaplan-Meier Survival Curve for Terminal Sacrifice Female BPA Stop Dose Arm (Weeks 4-104)

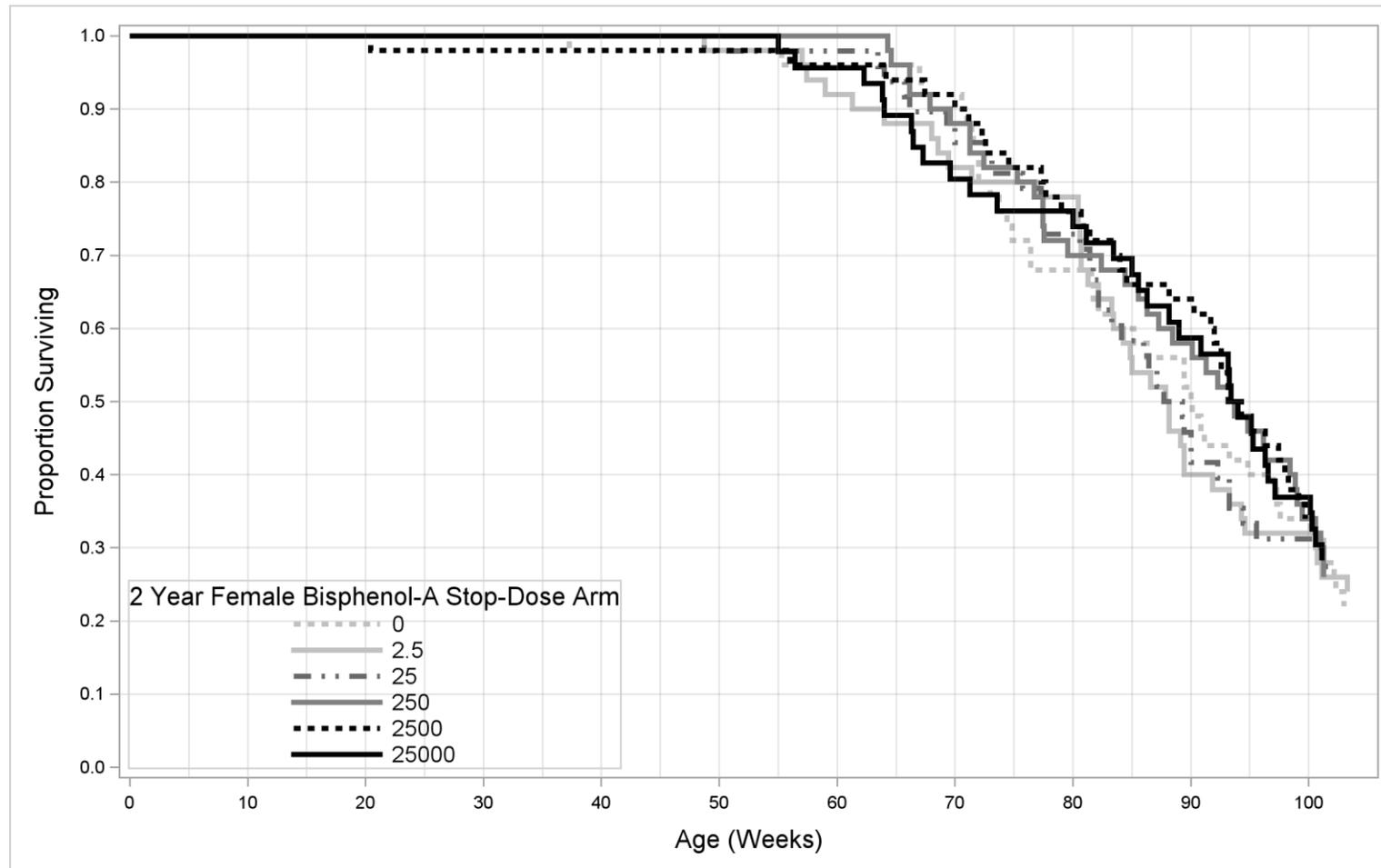


Figure 2. Kaplan-Meier Survival Curve for Terminal Sacrifice Male BPA Stop Dose Arm (Weeks 4-104)

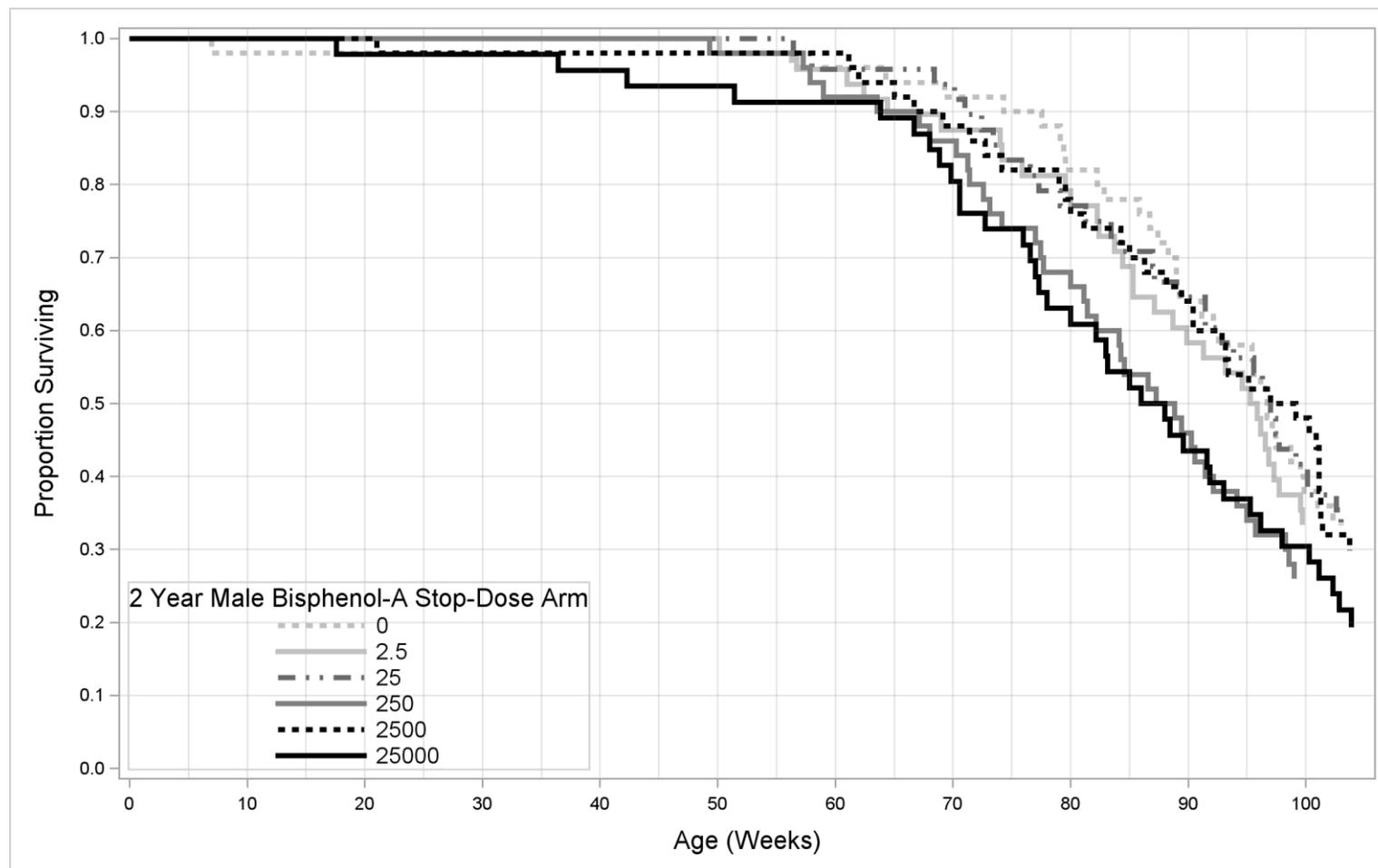


Figure 3. Kaplan-Meier Survival Curve for Terminal Sacrifice Female BPA Continuous Dose Arm (Weeks 4-104)

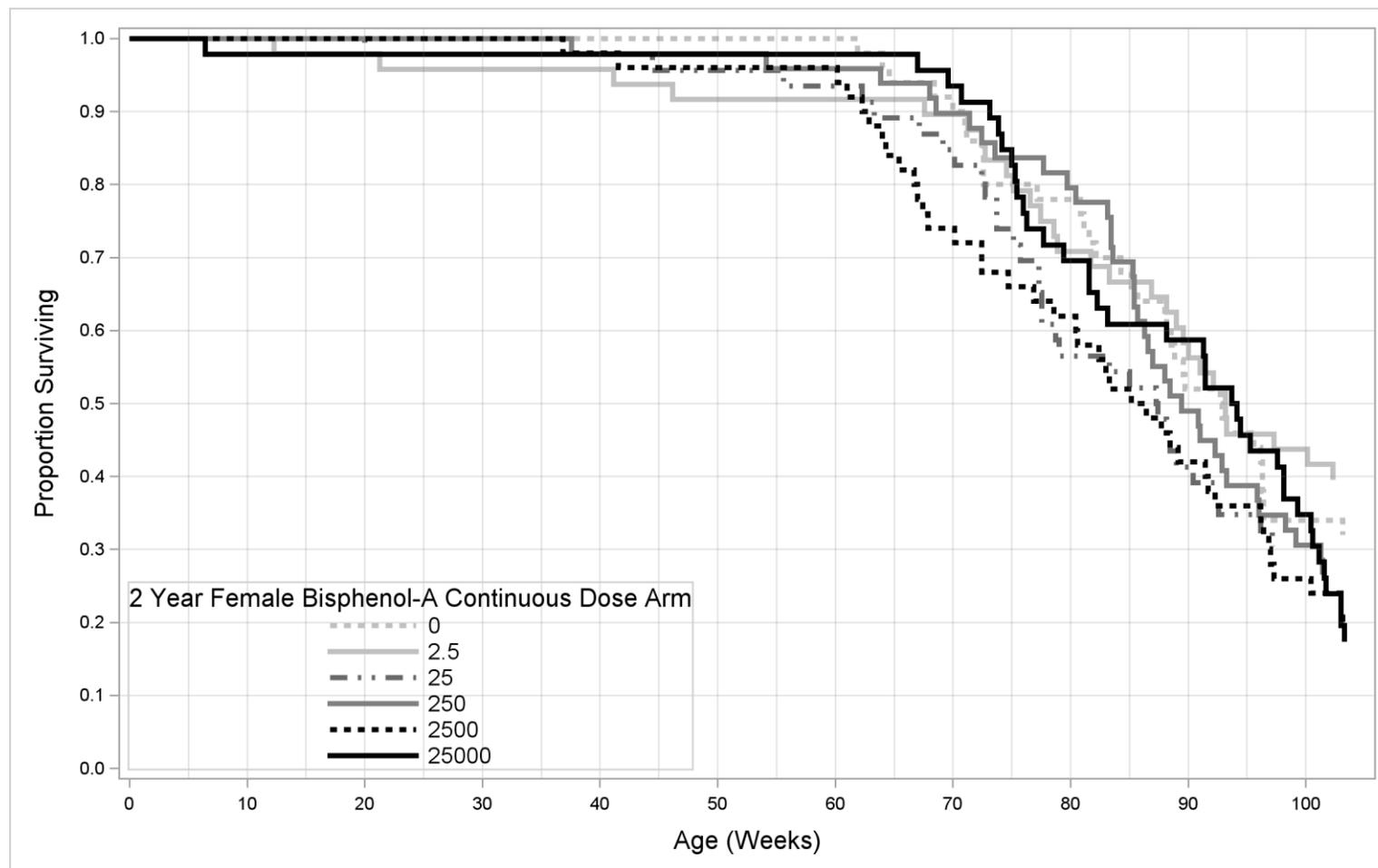


Figure 4. Kaplan-Meier Survival Curve for Terminal Sacrifice Male BPA Continuous Dose Arm (Weeks 4-104)

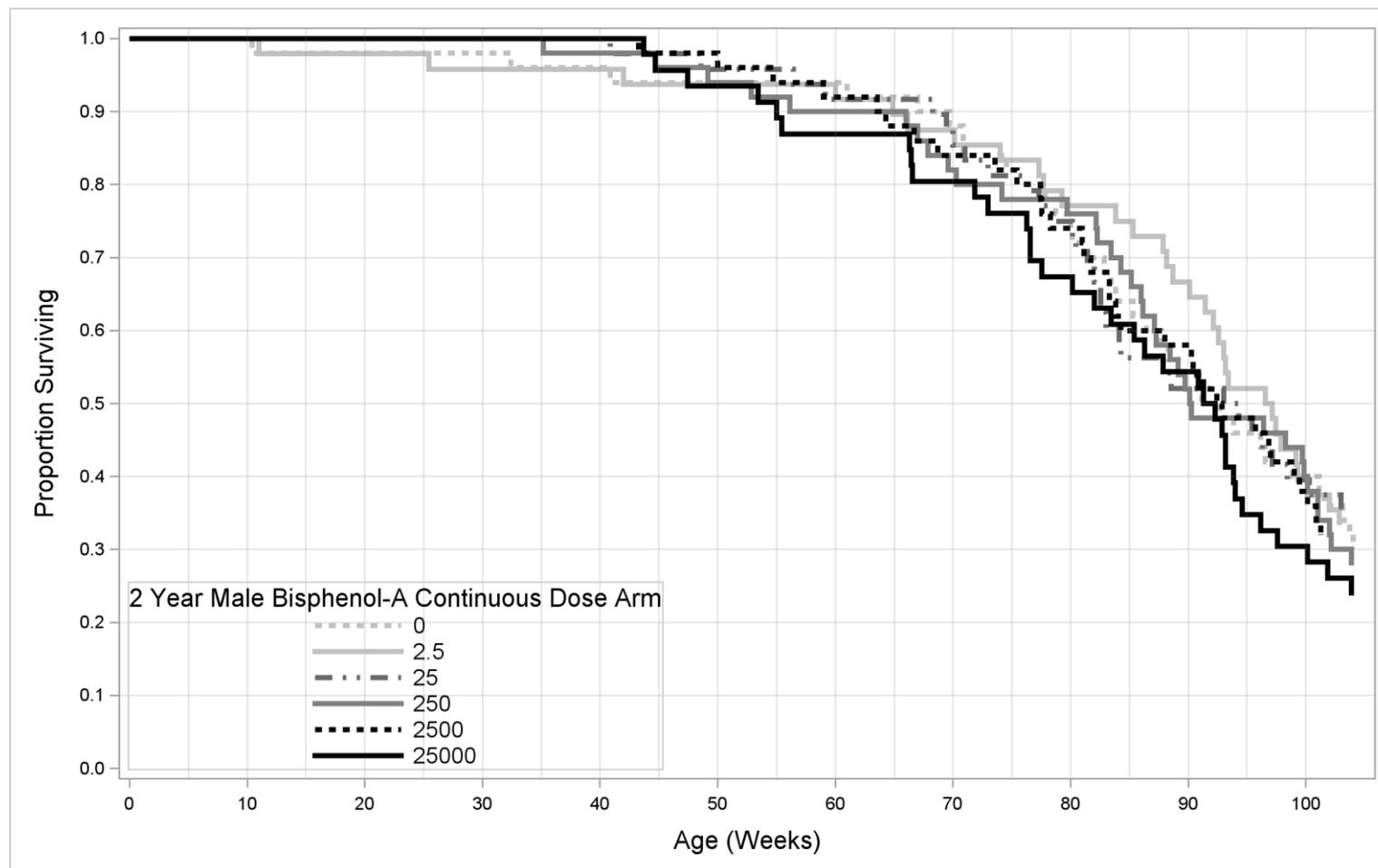


Figure 5. Kaplan-Meier Survival Curve for Terminal Sacrifice Female EE_2 Continuous Dose (Weeks 4-104)

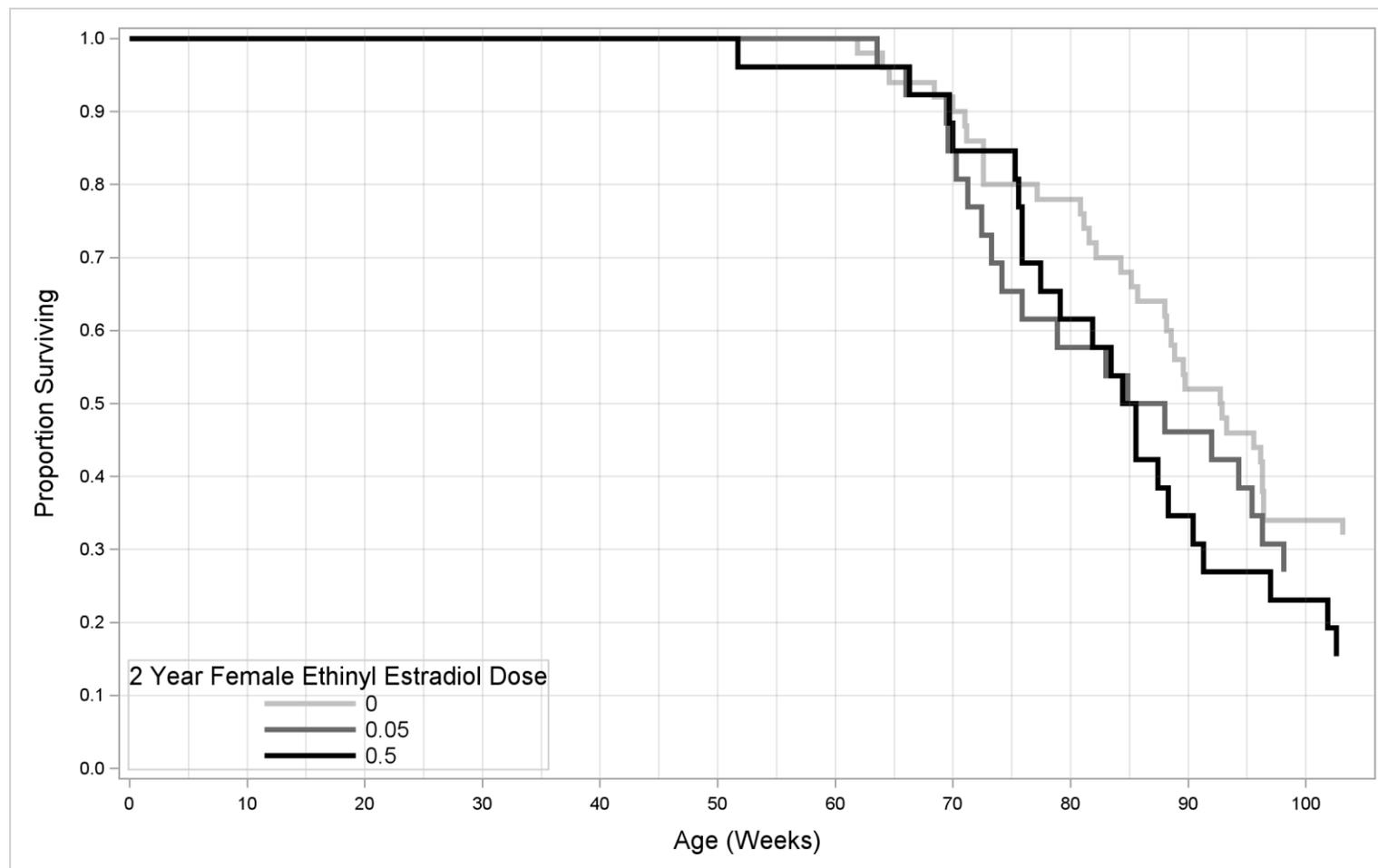
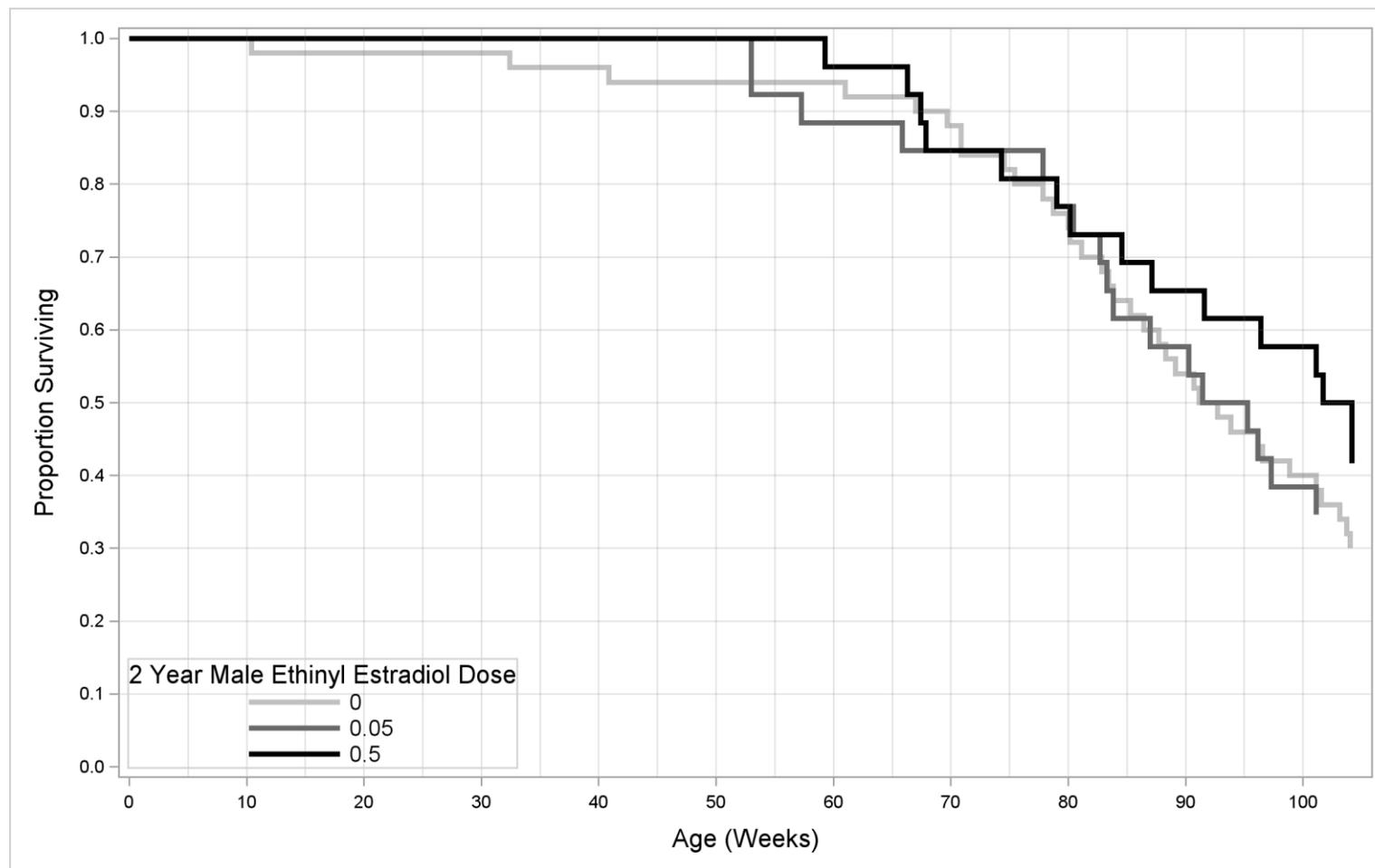


Figure 6. Kaplan-Meier Survival Curve for Terminal Sacrifice Male EE_2 Continuous Dose (Weeks 4-104)



C. Data

Survival data were extracted from the Genesis database using SAS Proc SQL, utilizing the Vortex ODBC driver.

Quality Control

1. *Data Verification*

The extraction of the data into SAS was verified by the statistical reviewer by review of the SAS code used to extract and verify the data.

2. *Computer Program Verification*

SAS programs were used to extract the data, explore the distributional properties of the data, and perform the statistical analysis.

The SAS programs were verified by detailed review of the program code, the program log, and the program output.

3. *Statistical Report Review*

3.1 *Statistical Report Text*

The statistical report was reviewed for logic, internal completeness, technical appropriateness, technical accuracy, and grammar. Technical appropriateness was reviewed based on statistical expertise.

Comments and questions were provided from the reviewer to the statistician. The statistician made appropriate changes and returned the report to the reviewer for final verification.

The text of the final statistical report was considered by the reviewer to be logical, internally complete, and technically appropriate and accurate. The statistical results stated in the text accurately presented those in the tables.

3.2 *Table Verification*

Analysis results were output from SAS to an .rtf file using PROC REPORT, which were then copied into the statistical report.

Statistical report tables were verified by checking the procedure used to create the tables and, additionally, by checking numbers sufficiently to conclude that the tables are correct.

3.3 *Graph Verification*

Graphs were verified by review of the SAS code used to generate them, and by calculation of summary statistics and checking numbers sufficiently to conclude that the graphs are correct. Graphs appear to be appropriate and correct.

4. *Conclusions*

The final statistical report has been fully reviewed and is considered by the reviewer to be logical, internally complete, and technically appropriate and accurate.