

Ten-year outcomes of uveal melanoma based on The Cancer Genome Atlas (TCGA) classification in 1001 cases

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Purpose: To understand the prognostic value of The Cancer Genome Atlas (TCGA) for uveal melanoma metastasis, using a simplified 4-category classification, based on tumor DNA. **Methods:** A retrospective cohort study of 1001 eyes with uveal melanoma at a single center, categorized according to TCGA as Group A, B, C, or D (by fine-needle aspiration biopsy for DNA analysis), and treated with standard methods, was studied for melanoma-related metastasis at 5 and 10 years. **Results:** Of 1001 eyes with uveal melanoma, the TCGA categories included Group A ($n = 486$, 49%), B ($n = 141$, 14%), C ($n = 260$, 26%), and D ($n = 114$, 11%). By comparison, increasing category (A vs. B vs. C vs. D) was associated with features of older age at presentation (56.8 vs. 52.8 vs. 61.1 vs. 63.5 years, $P < 0.001$), less often visual acuity of 20/20–20/50 (80% vs. 67% vs. 70% vs. 65%, $P = 0.001$), tumor location further from the optic disc ($P < 0.001$) and foveola ($P < 0.001$), and greater median tumor basal diameter (10.0 vs. 13.0 vs. 14.0 vs. 16.0 mm, $P < 0.001$) and tumor thickness (3.5 vs. 5.2 vs. 6.0 vs. 7.1 mm, $P < 0.001$). The Kaplan–Meier (5-year/10-year) rate of metastasis was 4%/6% for Group A, 12%/20% for Group B, 33%/49% for Group C, and 60%/not available for Group D. **Conclusion:** A simplified 4-category classification of uveal melanoma using TCGA, based on tumor DNA, is highly predictive of risk for metastatic disease.

Key words: Choroid, ciliary body, cytogenetics, genetics, melanoma, outcomes, TCGA, The Cancer Genome Atlas, uvea

The Cancer Genome Atlas (TCGA) is an international collaborative project designed by the National Cancer Institute's Center for Cancer Genomics and the National Human Genome Research Institute for the investigation of human cancer-related mutations. This project explored molecular aberrations on various platforms including histologic features, chromosome copy numbers, genetic mutations, expression of RNA, DNA methylation status, proteins, biochemical pathways, and immune markers for 33 cancer types, including uveal melanoma.^[1–8] In 2017, Robertson *et al.* published TCGA results of 80 eyes with uveal melanoma and identified four distinct molecular subsets, two associated with good prognosis (disomy 3 [D3]) and two associated with poor prognosis (monosomy 3 [M3]).^[3] In 2018, Jager *et al.* commented on TCGA results for uveal melanoma in an editorial, indicating the enormity and organization of this project, and provided clarity to the 4 main prognostic categories of uveal melanoma as identified by TCGA.^[4] They remarked that these clusters included Group A (D3, disomy 8), Group B (D3, 8q gain), Group C (M3, 8q gain possible), and Group D (M3, 8q gain multiple [isochromosome for 8q]).

In 2019, Vichitvejpaisal *et al.* applied the simplified 4-group classification of TCGA to a cohort of 658 patients with uveal melanoma at a single center and found the 5-year cumulative percentage of distant metastasis (odds ratio, OR) at 4% (OR 1.0)

for Group A, 20% (OR 3.5) for Group B, 33% (OR 11.4) for Group C, and 63% (OR 26.4) for Group D.^[5] Mazloumi *et al.* later documented that TCGA classification provided superior accuracy to the American Joint Committee on Cancer (AJCC) 8th edition categories, subcategories, and stages for 5-year prediction of metastasis of uveal melanoma.^[7] Herein, we explore a larger cohort of 1001 cases with a longer follow-up period using 5-year and 10-year Kaplan–Meier analyses to further validate TCGA in the prediction of uveal melanoma metastasis.

Methods

The medical records of the Ocular Oncology Service at the Wills Eye Hospital, Thomas Jefferson University, Philadelphia, Pennsylvania, USA, were retrospectively reviewed for patients with the clinical diagnosis of uveal melanoma between February 4, 2004 and June 2, 2020, who underwent genetic evaluation and assessment for The Cancer Genome Atlas (TCGA) classification. This study was approved by the Institutional Review Board of Wills Eye Hospital, adhering to the tenets of the Declaration of Helsinki, and complied with the Health Insurance Portability and Accountability Act. Informed consent was obtained from all patients.

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All patients were examined by a trained ocular oncologist (CLS, SEL, JAS) for clinical confirmation of diagnosis of uveal melanoma based on indirect ophthalmoscopy with large detailed fundus drawings and imaging. Ophthalmic imaging included fundus photography with wide-angle imaging, fundus autofluorescence, ultrasonography, optical coherence tomography (OCT), fluorescein angiography, indocyanine green angiography, and OCT angiography, as needed for documentation at the first examination and subsequent examinations. Those patients who had undergone genetic testing by fine-needle aspiration biopsy or open biopsy with results on the status of chromosomes 3 and 8 were classified according to TCGA as A, B, C, or D and included in this study.

Data were recorded at each examination and documented in patient's chart. The demographic data included age (years), sex (male, female), race (Caucasian, African American, Hispanic, Asian, others/unknown), affected eye (right, left), and visual acuity (20/20–20/50, 20/60–20/200, 20/400–no light perception [NLP]).

The tumor features at presentation included tumor location with distance from the optic disc (millimeters [mm]), distance from the foveola (mm), largest tumor basal diameter (mm), tumor thickness (mm), tumor epicenter (choroid, ciliary body, iris), anterior margin of the tumor (macula, macula to the equator, equator to ora serrata, ciliary body, iris), and posterior margin of the tumor (macula, macula to the equator, equator to ora serrata, ciliary body, iris).

Samples for genetic testing were obtained by fine-needle aspiration biopsy (FNAB), performed in the operating room at the time of uveal melanoma treatment, as described in the literature.^[5] The samples were stored in Hanks' balanced salt solution (Gibco, Life Technologies, Grand Island, NY) at 4 degrees Celsius, and DNA analysis was performed using a DNA Micro Kit (QIAGEN, Valencia, CA).^[5]

Primary outcomes included the rate and mean time to melanoma-related metastasis and death. Information on metastasis was gathered through history from the patient and correspondence from physicians. Information on death was gathered through history from the family and correspondence from physicians and family. Metastasis was further stratified based on the rate of metastasis to the liver, lung, and other systemic locations. Kaplan–Meier analysis at 1 year, 2 years, 5 years, and 10 years for each outcome of melanoma-related metastasis and death as well as metastasis to the liver, lung, and other systemic locations was stratified by TCGA classification (Group A vs. Group B vs. Group C vs. Group D). Additionally, Cox regression analyses to assess competing risks were performed but did not differ significantly from Kaplan–Meier analysis in this population.

Statistical analysis was performed using SAS Software Suite (version 9.4; SAS Institute). Continuous variables were expressed as mean (median, range). The one-sample Shapiro–Wilk test was used to assess the normality of distribution. A comparison between groups was performed using the one-way ANOVA test for continuous variables with normal distribution and the Kruskal–Wallis test for continuous variables without normal distribution. A comparison of categorical variables was performed using the likelihood ratio Chi-square test and Fisher's exact test when indicated. Binary logistic regression analysis was performed to identify factors potentially predictive of metastasis and death, which could act as confounders because of their strong correlation with the TCGA classification. Variables found to be significant in

univariate analysis at a level of $P < 0.05$ (age, largest tumor basal diameter, tumor thickness, distance from the optic disc, distance from the foveola, location of the tumor epicenter, anterior margin of the tumor, and posterior margin of the tumor) were entered into multivariate multiple regression models using the stepwise Wald method, which further excluded variables non-contributory to the fit of the model ($P > 0.05$). Odds ratios and 95% confidence intervals were generated for both univariate and multivariate regression models. Kaplan–Meier analysis was performed for metastasis (liver metastasis, lung metastasis, any metastasis) and death from uveal melanoma. A P value < 0.05 was considered statistically significant for the results of multivariate multiple regression and Kaplan–Meier analysis.

Results

There were 1001 consecutive eyes with uveal melanoma in 999 patients that were sampled for DNA analysis of chromosomes 3 and 8 at the time of tumor management at the Ocular Oncology Service at the Wills Eye Hospital at Thomas Jefferson University, Philadelphia, Pennsylvania USA, over a 22-year period. Patients with no genetic testing and/or those with no follow-up information were not included in this analysis.

Of all 1001 eyes with uveal melanoma, TCGA categories included Group A ($n = 486$, 49%), B ($n = 141$, 14%), C ($n = 260$, 26%), and D ($n = 114$, 11%). Demographic features are listed in Table 1. By comparison, increasing category (A vs. B vs. C vs. D) was associated with older age at presentation (56.8 vs. 52.8 vs. 61.1 vs. 63.5 years, $P < 0.001$) and less often visual acuity of 20/20–20/50 (80% vs. 67% vs. 70% vs. 65%, $P = 0.001$). There was no difference in sex, race, or affected eye.

Tumor characteristics are listed in Table 2. By comparison, increasing category (A vs. B vs. C vs. D) was associated with tumor location further from the optic disc ($P < 0.001$) and foveola ($P < 0.001$), increasing tumor basal diameter (10.5 mm vs. 12.7 mm vs. 13.6 mm vs. 15.3 mm, $P < 0.001$) and tumor thickness (4.4 mm vs. 6.2 mm vs. 6.7 mm vs. 7.6 mm, $P < 0.001$), greater frequency of the anterior margin involving the ciliary body (13% vs. 28% vs. 32% vs. 41%, $P < 0.001$), and less frequency of the posterior margin in the macula (63% vs. 56% vs. 48% vs. 55%, $P < 0.001$).

Outcomes for melanoma-related metastasis and death are listed in Table 3. Overall mean follow-up duration was 41.0 months (median 30.6, range < 0.1 –184.9), and tumors with greater TCGA classification had a shorter follow-up duration (46.4 vs. 45.0 vs. 32.4 vs. 32.7, $P < 0.001$).

By comparison, increasing category (A vs. B vs. C vs. D) was associated with the increased rate of any melanoma metastasis (3% vs. 9% vs. 20% vs. 46%, $P < 0.001$), shorter mean time to any metastasis (37.4 vs. 38.7 vs. 27.7 vs. 21.5, $P = 0.009$), and specifically the increased rate of liver metastasis (2% vs. 9% vs. 20% vs. 46%, $P < 0.001$), lung metastasis ($< 1\%$ vs. 1% vs. 4% vs. 10%, $P < 0.001$), metastasis to other systemic locations (bone, brain, breast, intestine, distant lymph nodes, mesentery, muscle, skin) (1% vs. 4% vs. 5% vs. 14%, $P < 0.001$), and melanoma-related death ($< 1\%$ vs. 0% vs. 2% vs. 7%, $P = 0.003$). By comparison, increasing specific category was associated with the increased odds ratio (OR) for any metastasis (A vs. B, OR 2.48 ($P = 0.026$); A vs. C, OR 6.21 ($P = 0.027$); A vs. D, OR 22.25 ($P < 0.001$)) and for liver metastasis (A vs. B, OR 3.20 ($P = 0.026$); A vs. C, OR 7.32 ($P = 0.009$); A vs. D, OR 25.04 ($P < 0.001$)). A similar increasing risk (A vs. D) was observed for lung metastasis (OR

Table 1: Ten-Year Outcomes of Uveal Melanoma Based on The Cancer Genome Atlas (TCGA) Classification in 1001 Cases. Patient demographics

Patient Demographics	TCGA Class					Total Population (n=1001) [n (%)]
	A (n=486) [n (%)]	B (n=141) [n (%)]	C (n=260) [n (%)]	D (n=114) [n (%)]	P	
Age						
Mean (years) (median, range)	56.8 (58.0, 10.0 - 90.0)	52.8 (54.0, 13.0 - 83.0)	61.1 (62.5, 12.0 - 88.0)	63.5 (64.0, 28.0 - 94.0)	<0.001	58.1 (59.0, 10.0 - 94.0)
Sex						
Male	266 (55)	72 (51)	121 (47)	57 (50)	0.194	516 (52)
Female	220 (45)	69 (49)	139 (53)	57 (50)		485 (48)
Race						
Caucasian	472 (97)	130 (92)	251 (97)	112 (98)	0.088	965 (96)
African American	0 (0)	2 (1)	0 (0)	0 (0)		2 (<1)
Hispanic	10 (2)	3 (2)	5 (2)	2 (2)		20 (2)
Asian	1 (<1)	3 (2)	1 (<1)	0 (0)		5 (<1)
Other/unknown	3 (1)	3 (2)	3 (1)	0 (0)		9 (1)
Affected eye						
Right	249 (51)	77 (55)	143 (55)	60 (53)	0.760	529 (53)
Left	237 (49)	64 (45)	117 (45)	54 (47)		472 (47)
Visual Acuity						
20/20 - 20/50	389 (80)	94 (67)	182 (70)	74 (65)	0.001	739 (74)
20/60 - 20/200	55 (11)	29 (21)	53 (20)	28 (25)		165 (16)
20/400 - NLP	42 (9)	18 (13)	25 (10)	12 (11)		97 (10)

Bold values indicate significant P. TCGA=The Cancer Genome Atlas; NLP=No light perception

Table 2: Ten-Year Outcomes of Uveal Melanoma Based on The Cancer Genome Atlas (TCGA) Classification in 1001 Cases. Tumor features

Tumor Features	TCGA Class					Total Population (n=1001) [n (%)]
	A (n=486) [n (%)]	B (n=141) [n (%)]	C (n=260) [n (%)]	D (n=114) [n (%)]	P	
Distance to optic disc (mm) [mean (median, range)]	3.9 (3.0, 0.0 - 20.0)	4.7 (4.0, 0.0 - 18.0)	5.5 (5.0, 0.0 - 17.0)	4.9 (5.0, 0.0 - 18.0)	<0.001	4.5 (3.5, 0.0 - 20.0)
Distance to foveola (mm) [mean (median, range)]	3.6 (2.3, 0.0 - 18.4)	4.1 (3.3, 0.0 - 15.0)	5.4 (4.0, 0.0 - 18.0)	4.7 (3.0, 0.0 - 17.0)	<0.001	4.3 (3.0, 0.0 - 18.4)
Largest basal diameter (mm) [mean (median, range)]	10.5 (10.0, 1.0 - 22.0)	12.7 (13.0, 2.0 - 22.0)	13.6 (14.0, 2.0 - 24.0)	15.3 (16.0, 6.0 - 24.0)	<0.001	12.1 (12.0, 1.0 - 24.0)
Thickness (mm) [mean (median, range)]	4.4 (3.5, 1.0 - 14.0)	6.2 (5.2, 1.3 - 15.0)	6.7 (6.0, 0.7 - 16.0)	7.6 (7.1, 2.1 - 20.4)	<0.001	5.6 (4.7, 0.7 - 20.4)
Tumor epicenter						
Choroid	457 (94)	126 (89)	227 (87)	103 (90)	0.008	913 (91)
Ciliary body	20 (4)	11 (8)	26 (10)	11 (10)		68 (7)
Iris	9 (2)	4 (3)	7 (3)	0 (0)		20 (2)
Anterior margin						
Macula	32 (7)	9 (6)	6 (2)	4 (4)	<0.001	51 (5)
Macula to equator	245 (50)	40 (28)	64 (25)	22 (19)		371 (37)
Equator to ora serrata	131 (27)	44 (31)	81 (31)	38 (33)		294 (29)
Ciliary body	62 (13)	40 (28)	82 (32)	47 (41)		231 (23)
Iris	16 (3)	8 (6)	27 (10)	3 (3)		54 (5)
Posterior margin						
Macula	305 (63)	79 (56)	124 (48)	63 (55)	<0.001	571 (57)
Macula to equator	156 (32)	54 (38)	115 (44)	48 (42)		373 (37)
Equator to ora serrata	10 (2)	2 (1)	12 (5)	3 (3)		27 (3)
Ciliary body	7 (1)	2 (1)	5 (2)	0 (0)		14 (1)
Iris	8 (2)	4 (3)	4 (2)	0 (0)		16 (2)

Bold values indicate significant P. TCGA=The Cancer Genome Atlas

Table 3. Ten-Year Outcomes of Uveal Melanoma Based on The Cancer Genome Atlas (TCGA) Classification in 1001 Cases. Outcomes.

Outcomes	TCGA Class				P	Total Population (n=1001) [n (%)]
	A (n=486) [n (%)]	B (n=141) [n (%)]	C (n=260) [n (%)]	D (n=114) [n (%)]		
Follow-up (months) (median, range)	46.4 (40.1, <0.1 - 161.0)	45.0 (33.9, 0.1 - 184.9)	32.4 (23.9, 0.1 - 173.1)	32.7 (28.0, 0.0 - 117.4)	<0.001	41.0 (30.6, <0.1 - 184.9)
Metastasis	14 (3)	12 (9)	53 (20)	52 (46)	0.026 0.027 <0.001 0.009	131 (13)
Mean time to metastasis (months) (median, range)	37.4 (40.2, 0.5 - 81.0)	38.7 (38.3, 1.1 - 77.3)	27.7 (19.1, 0.2 - 85.6)	21.5 (18.0, 0.0 - 107.2)	---	27.3 (19.9, 0.0 - 107.2)
Liver metastasis	10 (2)	12 (9)	51 (20)	52 (46)	0.026 0.009 <0.001 0.032	125 (12)
Mean time to metastasis (months) (median, range)	30.2 (22.2, 0.5 - 65.7)	38.9 (38.3, 1.1 - 77.3)	26.7 (18.7, 0.2 - 85.6)	21.9 (18.0, 0.0 - 107.2)	---	26.2 (19.5, 0.0 - 107.2)
Lung metastasis	1 (<1)	2 (1)	10 (4)	11 (10)	0.615 0.047 <0.001 0.896	24 (2)
Mean time to metastasis (months) (median, range)	28.4 (28.4, 28.4 - 28.4)	47.4 (47.4, 17.5 - 77.3)	31.5 (25.1, 1.3 - 65.7)	36.8 (37.5, 5.6 - 72.3)	A vs. B: 4.91 (0.44 - 55.23) A vs. C: 11.50 (1.43 - 92.32) A vs. D: 34.75 (4.40 - 274.59)	34.8 (32.9, 1.3 - 77.3)
Other metastasis*	5 (1)	5 (4)	13 (5)	16 (14)	0.659 0.709 <0.001 0.263	39 (4)
Mean time to metastasis (months) (median, range)	48.2 (53.9, 0.5 - 81.0)	42.8 (50.7, 6.9 - 65.9)	41.7 (41.8, 13.6 - 73.7)	35.3 (35.2, 3.6 - 72.3)	---	40.2 (43.4, 0.5 - 81.0)
Death from UM metastasis	2 (<1)	0 (0)	4 (2)	8 (7)	-- A vs. B: na A vs. C: 2.26 (0.36 - 14.07) A vs. D: 9.65 (1.67 - 55.73)	14 (1)
Mean time to death (months) (median, range)	48.0 (48.0, 27.2 - 68.8)	0.0 (0.0, 0.0 - 0.0)	47.4 (46.0, 38.2 - 59.4)	26.5 (24.3, 3.5 - 50.5)	---	35.5 (38.3, 3.5 - 68.8)

Bold values indicate significant P. UM = Uveal melanoma, NA= not available. *Sites of other metastasis include bone, brain, breast, intestine, distant lymph nodes, mesentery, muscle, skin

34.75, $P < 0.001$) and metastasis to other locations (OR 7.30, $P < 0.001$).

Kaplan–Meier analysis of outcomes of metastasis and death is listed in Table 4 [Fig. 1]. By comparison, increasing category (A vs. B vs. C vs. D) was associated with greater risk for any melanoma-related metastasis ($P < 0.001$) at 1-year (1% vs. 2% vs. 7% vs. 15%), at 2-years (2% vs. 3% vs. 15% vs. 36%), at 5-years (4% vs. 12% vs. 33% vs. 60%), and at 10-years (6% vs. 20% vs. 49% vs. not available). A similar increasing risk for liver ($P < 0.001$), lung ($P < 0.001$), and other metastases as well as melanoma-related death ($P < 0.001$) over time were documented [Table 4 and Fig. 1].

Discussion

The Cancer Genome Atlas (TCGA) has provided a comprehensive “cancer atlas” through wide-ranging multi-platform analyses of over 30 human cancers, including uveal melanoma.^[2] This project involved a multicenter organization including tissue source sites for the collection of blood and tissue samples; biospecimen core resources for coordination of sample delivery and cataloging; genome sequencing centers for high-throughput sequencing and identification of DNA alterations; cancer genome characterization centers for description of alterations in miRNA, gene expression,

single nucleotide polymorphisms, and others; proteome characterization centers for the identification of cancer-specific proteins; data coordinating centers for the collection and transfer of data to public databases; cancer genomics hub for storage, cataloging, and access to information; and genome data analysis centers for the development of informatics tools for processing data across the entire genome.^[2] This unprecedented effort was then made available for public access, providing researchers the opportunity to evaluate this data.

In 2017, Robertson *et al.* were the first to publish results of TCGA regarding uveal melanoma in 80 cases.^[3] They identified four molecularly-distinct groups of uveal melanoma, with two groups demonstrating favorable prognosis related to disomy 3 (D3) and two groups showing poor prognosis related to monosomy 3 (M3). Within each of the four groups there were unique gene alterations, DNA methylation, mRNA expression levels, and other findings that accounted for increasing the risk for melanoma metastasis. Jager *et al.* later clarified this distinct molecular grouping of uveal melanoma as A, B, C, and D, whereby the criteria for Group A included D3 and disomy of chromosome 8, Group B showed D3 and 8q gain, Group C showed M3 and often with 8q gain, and Group D showed M3 with multiple 8q gains manifesting as an isochromosome for 8q.^[4] The simplicity of this 4-category

Table 4: Ten-Year Outcomes for Uveal Melanoma Based on The Cancer Genome Atlas (TCGA) Classification in 1001 Cases. Event-Free Survival Analysis of Metastasis and Death

Event-Free Survival	TCGA Class				Total Population (n=1001) [n (%)]
	A (n=486) [n (%)]	B (n=141) [n (%)]	C (n=260) [n (%)]	D (n=114) [n (%)]	
Overall Metastasis					
1 Year	394 (99)	123 (98)	185 (93)	83 (85)	785 (96)
2 Years	315 (98)	92 (97)	118 (85)	52 (64)	577 (91)
5 Years	160 (96)	40 (88)	41 (67)	12 (40)	254 (82)
10 Years	16 (94)	6 (80)	5 (51)	NA	27 (75)
Liver Metastasis					
1 Year	394 (99)	123 (98)	185 (93)	83 (86)	785 (96)
2 Years	315 (98)	92 (97)	118 (85)	52 (65)	577 (91)
5 Years	160 (98)	40 (88)	41 (67)	12 (42)	254 (82)
10 Years	17 (96)	6 (80)	5 (55)	NA	28 (77)
Lung Metastasis					
1 Year	394 (100)	123 (100)	195 (99)	91 (99)	801 (>99)
2 Years	316 (100)	91 (>99)	130 (97)	64 (95)	601 (99)
5 Years	161 (99)	41 (99)	42 (93)	13 (76)	257 (96)
10 Years	17 (99)	6 (95)	5 (89)	NA	28 (94)
Other Metastasis*					
1 Year	394 (>99)	121 (99)	196 (100)	92 (99)	802 (>99)
2 Years	316 (>99)	91 (99)	131 (99)	64 (93)	604 (99)
5 Years	160 (98)	40 (95)	42 (88)	14 (71)	256 (93)
10 Years	16 (97)	6 (90)	5 (83)	NA	27 (90)
Death from Uveal Melanoma					
1 Year	394 (100)	123 (100)	196 (100)	92 (99)	802 (>99)
2 Years	316 (100)	92 (100)	131 (100)	66 (96)	604 (99)
5 Years	162 (>99)	41 (100)	42 (93)	14 (85)	259 (97)
10 Years	17 (99)	6 (100)	5 (93)	NA	28 (97)

TCGA=The Cancer Genome Atlas, NA=not available. *Sites of other metastasis include bone, brain, breast, intestine, distant lymph nodes, mesentery, muscle, skin

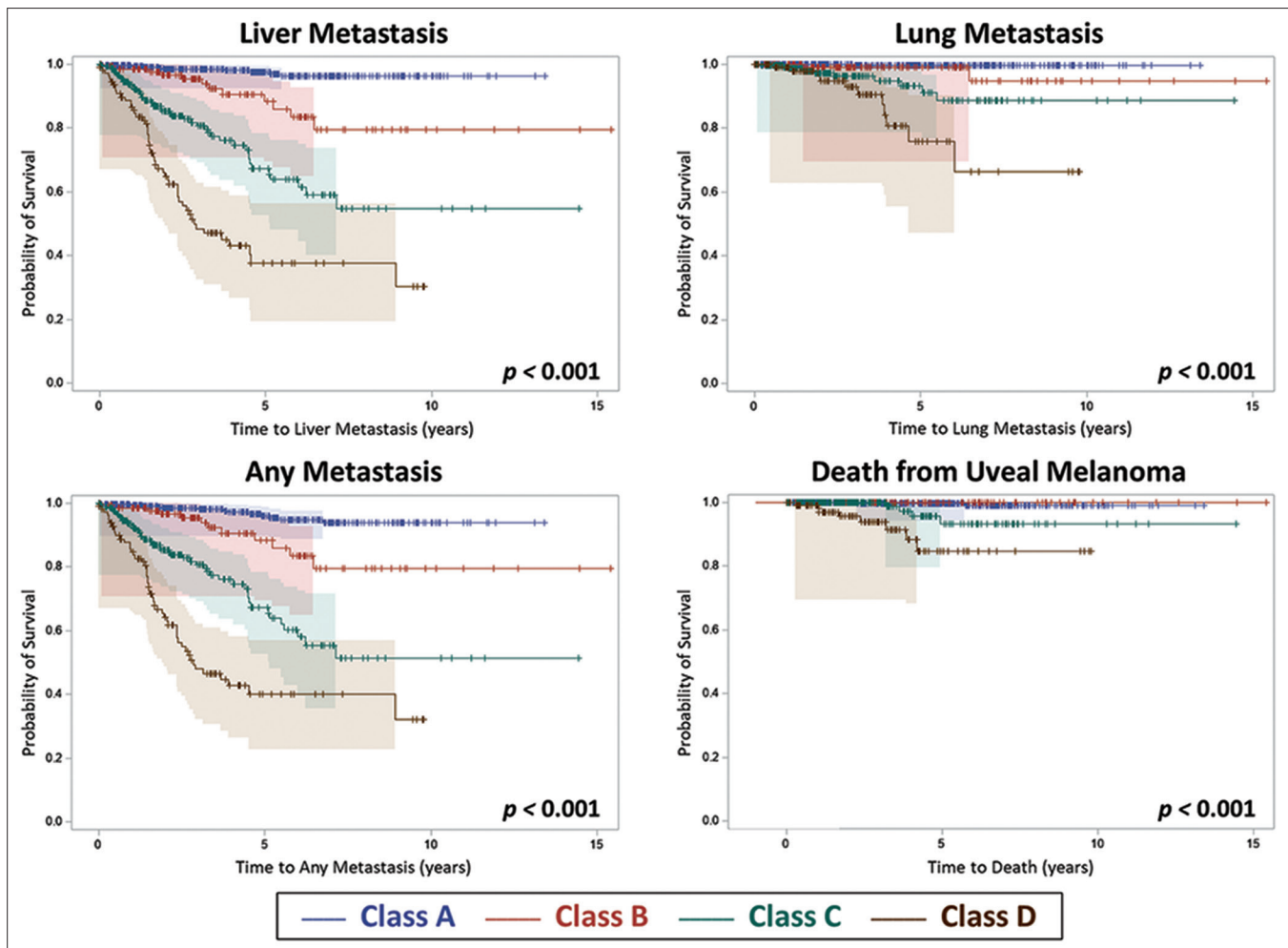


Figure 1: Kaplan–Meier estimates of metastasis and survival according to The Cancer Genome Atlas (TCGA) classification into Group A, B, C, or D. With increasing group, there was increasing risk for liver metastasis ($P < 0.001$), lung metastasis ($P < 0.001$), any metastasis ($P < 0.001$), and death ($P < 0.001$)

DNA-based classification scheme provides a straightforward approach for stratifying uveal melanoma prognosis, as previous investigations have included complex multiple factors^[9–14] such as over 50 combinations of DNA alterations in chromosomes 3, 6, and 8 giving a stepwise, graded prognosis,^[15,16] whereas others have blended in the AJCC classification^[17–19] with DNA alterations^[20,21] further refining prognosis, and still others have combined DNA alterations of chromosomes 3 and 8 plus mitotic activity, closed loops, epithelioid cells, basal tumor diameter, extraocular spread, and optic disc and ciliary body involvement in the equation for an all-inclusive prognosis.^[22,23]

In 2019, Vichitvejpaisal *et al.* validated the 4-category TCGA classification in an analysis of 658 cases, demonstrating that the 5-year Kaplan–Meier cumulative percentage of distant metastasis based on DNA results for Group A was 4%, B was 20%, C was 33%, and D was 63%.^[5] In that cohort, longer-term, 10-year data was not available. In this current analysis, we explore a larger cohort of 1001 cases of uveal melanoma with more robust 10-year Kaplan–Meier outcomes. Based on TCGA classification using DNA results, we found that the 5-year Kaplan–Meier rate of any distant metastasis for Group A was 4%, B was 12%, C was 33%, and D was 60%, whereas the

10-year Kaplan–Meier rate of any distant metastasis for Group A was 6%, B was 20%, C was 49%, and D was not available due to the small cohort number. The odds ratios for any distant metastasis (vs. Group A) were 2.48 for Group B, 6.21 for Group C, and 22.25 for Group D. Furthermore, the 10-year Kaplan–Meier rates of specific liver metastasis revealed Group A at 4%, B at 20%, C at 45%, and D not available with odds ratios for liver metastasis (vs. Group A) at 3.20 for Group B, 7.32 for Group C, and 25.04 for Group D.

Understanding the uveal melanoma prognosis is an important driver for adjuvant therapies to prevent metastasis. Before TCGA data were released, a previous study of adjuvant sunitinib^[24] for patients at a high-risk of metastases, defined as M3 and 8q gain or M3 and large tumor size^[15] or gene expression profiling of Class 2,^[25,26] revealed a better overall survival in the treatment arm, particularly if the patient was younger than 60 years of age. Currently, we employ TCGA classification for high-risk adjuvant trials, and those that meet the criteria for Group C or Group D are considered for adjuvant therapy.

There are limitations to this study including its retrospective nature and the rarity of uveal melanoma. However, in our practice of ocular oncology, we specialize in the management of uveal

melanoma and have offered FNAB for DNA prognostication to all patients undergoing therapy, with subsequent monitoring of patient outcomes for decades. When we began our FNAB program for genetic testing in the mid-2000s, analysis was performed only on chromosome 3; thus the data from these patients could not be incorporated into TCGA as information on chromosome 8 is required. Nowadays, we sample for multiple chromosomes and our large cohort of 1001 eyes with uveal melanoma includes information on chromosomes 3 and 8 and provides a robust sample for validation of outcomes. Another limitation of this dataset is that outcomes for metastasis and death were per report by the patient, family, or physician. We realize that there can be gaps with this method of information collection, but this represents “real world” data analysis. Some metastatic and death events could be under-reported.

Conclusion

In conclusion, we have updated our database regarding prognostic classification of uveal melanoma by the 4-category TCGA and now provide 10-year outcomes for any metastasis and specifically for metastasis to the liver, lung, and other sites. We have shown that increasing TCGA grouping leads to a significantly ($P < 0.001$) increased risk for metastatic events and death over time. This classification system is practical and highly predictive of uveal melanoma metastatic risk.

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Conflicts of interest

There are no conflicts of interest.

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