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## Race/ethnicity and the risk childhood leukemia: a case-control study in California

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### Abstract

**Background**—We conducted a large registry-based study in California to investigate the association between race/ethnicity and childhood leukemia, focusing on two subtypes: acute lymphoblastic leukemia (ALL) and acute myeloid leukemia (AML).

**Methods**—We obtained information on 5788 cases and 5788 controls by linking California cancer and birth registries. We evaluated relative risk of childhood leukemia by race and ethnicity of the child and their parents using conditional logistic regression, with adjustment for potential confounders.

**Results**—Compared to Whites, Black children had lower risk of ALL (odds ratio [OR]=0.54, 95% CI: 0.45–0.66) as well as children of Black/Asian parents (OR=0.31, 95% CI: 0.10–0.94). Asian race was associated with increased risk of AML with OR=1.643, 95% CI: 1.10–2.46 for Asian vs. Whites and with OR=1.67, 95% CI: 1.04–2.70 for Asian/Asian vs. White/White. Hispanic ethnicity was associated with increased risk of ALL (OR=1.37, 95% CI: 1.22–1.52). A gradient in risk of ALL was observed comparing Hispanic children with both parents Hispanic, one parent Hispanic and non-Hispanic children (p-value for trend <0.0001). The highest risk of ALL was observed for children with a combination of Hispanic ethnicity and White race compared to non-Hispanic Whites (OR=1.27, 95% CI: 1.12–1.44). The lowest risk was observed for non-Hispanic Blacks (OR=0.46, 95% CI: 0.36–0.60). Associations for total childhood leukemia were similar to ALL.

**Conclusions**—Our results confirm that there are ethnic and racial differences in the incidence of childhood leukemia. These differences indicate that some genetic and/or environmental/cultural factors are involved in etiology of childhood leukemia.

### Keywords

childhood leukemia; child's race; parental race; child's Hispanic ethnicity; parental Hispanic ethnicity

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### Introduction

A limited number of studies have specifically examined race and/or ethnicity in relation to childhood leukemia risk.[1 2] Most studies on childhood leukemia have considered race and/or Hispanic ethnicity only as a covariate in their analyses. One interview-based study found that the proportion of Whites among controls was higher than among childhood leukemia cases [3], another found no association between race and childhood leukemia [4]; both studies were prone to biases. The majority of studies have shown that Black race was associated with decreased risk of childhood leukemia [1 2 5–10] compared to Whites. The definition of race/ethnicity differed in all these studies.

Fewer studies have looked at relationships between race/ethnicity and risk of major subtypes of childhood leukemia, acute lymphoblastic leukemia (ALL) and acute myeloid leukemia (AML). Most of these studies observed similar results for total childhood leukemia and ALL.[1 2 6 11–13] Two studies found no association with any subtype. [4 13] For AML findings varied: two studies found increased risk associated with White race, one study detected higher risk for Asian children [6], others did not find any association.[12]

Most studies on Hispanic origin of the child have report an increased risk of total childhood leukemia and ALL for Hispanic ethnicity [1 11 14 15]; however, some studies found no association between Hispanic origin and the risk of childhood leukemia.[4 16]

The aims of this large-scale study were to examine the relationships between race/ethnicity of child, mother and father and childhood leukemia and its subtypes. California is particularly suitable for studying racial and ethnic differences in the incidence of childhood leukemia due to its diverse racial/ethnic distribution. In addition to being one of the most diverse states in the U.S. [17], California has cancer and birth registries that have almost complete (99%) registration.[18 19]

Most previous record-based studies used a single definition of child's race and/or ethnicity. In our study we explored several definitions of child's race and ethnicity and their combination. Registry-based child's race/ethnicity may have sizeable amount of missing data. We reconstructed child's race and ethnicity from mother's and father's race and ethnicity, hence reducing missing data.

## Materials and methods

Eligible childhood leukemia cases included in this analysis were diagnosed between 1988 and 2008 in California-born children younger than 16 years who resided in California at the time of diagnosis. Information about cases, cancer types and characteristics was extracted from the population-based California Cancer Registry (CCR). Cancer registry was linked to the California Birth Registry (CBR) to select controls and to obtain information on socio-demographic and other factors. Paper birth certificates were obtained from CBR for years prior 1997 when no electronic birth records existed. Due to high cost of each certificate and low number of cases prior 1986, we restricted selection of cases and controls to 1986–2007 birth years. Controls were selected randomly from CBR and matched to cases (1 to 1) on date of birth ( $\pm 6$  months) and sex.

### Measures

In CCR child's race and ethnicity was available for cases only; maternal and paternal race was not available. Therefore, we used CBR to extract information on child's and parental race/ethnicity. Racial groups for parents available from CBR were as follows: White, Black, American Indian, Asian-unspecified and Asian-specified, Chinese, Japanese, Korean, Vietnamese, Cambodian, Thai, Laotian, Indian, Filipino, Guamanian, Samoan, Eskimo, Aleut, Pacific Islander, Hawaiian, and Other. Following the Surveillance, Epidemiology and End Results (SEER) classification, these groups were combined into five main racial categories as follows: White (White), Black (Black), Asian (Asian-unspecified and Asian-specified, Chinese, Japanese, Korean, Vietnamese, Cambodian, Thai, Laotian, Indian, & Filipino), American Indian and Alaskan (Eskimo, Aleut) and Pacific Islanders (PI) and Other (Guamanian, Samoan Hawaiian & all others).[20] These five categories were used in analysis for mother's and father's race.

Although child's race was available in CBR, more than 30% of values were missing. Significantly fewer subjects were missing information on parental race (see Table 2).

We, therefore, created classifications of child's race based on race of birth parents as recorded in child's birth certificate. We used several alternative approaches to classify child's race based on afore-mentioned five categories. By the reconstructed classification, a child was considered White if both parents were White; Black if either parent was Black; Asian if both parents were Asian or if one of the parents was White and another was Asian; American Indian and Alaskan if both parents were American Indian or Alaskan or if one parent was American Indian/Alaskan and the other was either White, or Asian, or PI & Other; and Pacific Islander (PI) and Other if both parents were PI or Other, if one parent was PI or Other and the second was either White or Asian.

To check how sensitive our results were to differences in classification of race in some analyses we used the original child's race from birth records categorized into the same five racial groups. Findings did not vary; therefore, we present results for the reconstructed child race.

Similar to a system used by Chow et al. [16], we constructed a 14-category variable for child's race consisting of the combinations of parental races: White/White, Black/Black, Asian/Asian, American Indian/American Indians, PI & Other/PI & Other, White/Black, White/Asian, White/American Indian, White/PI & Other, Black/Asian, Black/American Indian, Black/PI & Other, Asian/American Indian, Asian/PI & Other (see Table 1).

Parental Hispanic ethnicity was considered as a dichotomous variable separately from racial categorization: Hispanic (Mexican, Puerto Rican, Cuban, Central/South American, Other Hispanic) and non-Hispanic. Hispanic origin of child was not recorded in registries and was derived from parental Hispanic ethnicity from CBR. A child was classified as Hispanic if either parent was Hispanic. Analyses were also conducted that classified children of Hispanic ethnicity as having one or both parents Hispanic.

For some analyses, we combined child's race and Hispanic ethnicity to create a variable with ten categories: Non-Hispanic White, Hispanic White, Non-Hispanic Black, Hispanic Black, Non-Hispanic Asian, Hispanic Asian, Non-Hispanic American Indian, Hispanic American Indian, Non-Hispanic PI & Other and Hispanic PI & Other.

We used parental education as a proxy for SES at the individual level. Although maternal education was available from CBR, it was missing for about 60% of subjects. Therefore, we used paternal education to adjust for SES. It was categorized into four levels: <12 years, 12 years, 13–16 years, and 17 years and more. We used a measure of community-based SES derived from U.S. Census data using principal components analysis based on seven indicator variables at a census block level.[21] Available to us were component scores from principal component analysis for SES index categorized into quintiles. We adjusted models for father's education and census-based SES separately.

### Statistical Analysis

The primary analysis method was conditional logistic regression utilizing the matched case-control pairs. Several models using different subsets of covariates were fitted and checked for potentially influential observations. The models chosen based on information on known or potential confounders and model fit statistics; the most parsimonious models with the lowest Akaike information criterion values are presented. Models for child, maternal and paternal race/ethnicity were fit separately to avoid close correlations. We have also considered models with interaction between child's race and other variables in the model.

Large sample size of the study allowed us to conduct analysis by two main subtypes of childhood leukemia (ALL and AML) using the same models.

Despite the large number of cases and controls, sample sizes for some analyses were reduced due to missing data. Due to differences in data collection by year the pattern of missingness varied by year but no differences in patterns of missingness were detected between cases and controls. Under a missing at random assumption, multivariate imputation techniques were used to impute missing values for all variables in models by the MI procedure in SAS. [22–24] Analyses were repeated using the multiply imputed data using the MIANALYZE procedure.[24]

Analyses were conducted using SAS 9.3.[24]

The study was approved by University of California, Los Angeles Office for the Protection of Research Subjects and California Committee for the Protection of Human Subjects.

## Results

A total of 6645 childhood leukemia cases were identified from the California Cancer Registry. Linkage to birth records was successful for 87.1% of cases. Of the 5788 cases (55.8% males and 44.2% females) included in this analysis, 4721 were ALL cases, 852 were AML cases, and 215 were other types. The median age at diagnosis was 3.8 years (range 0–15.4) with the peak for ALL at 2–5 years of age and at 0–2 years of age for AML. Table 2 shows other characteristics of study subjects.

Results of conditional logistic regression analyses assessing the association of childhood leukemia and child's and parental races presented in the Table 3 indicate that Black race of child and of mother and father were each associated with a decreased risk of childhood leukemia compared to White race. Similar findings were observed for ALL. For AML, increased but imprecise risk was observed for Black children, Asian children, and children of Asian fathers. Adjusted analysis of a child's race as defined by father's and mother's races combined showed similar results to the main analysis with reconstructed 5-category child race.

We repeated the analyses adjusting for census-based SES instead of father's education. Results were similar, except the association of Asian race with AML was more precise. Children of Asian fathers had higher risk of AML compared to children of White fathers (OR=1.75, 95% CI: 1.13–2.72). Asian children had higher risk of AML in all analyses (OR=1.64, 95% CI: 1.10–2.46 for Asian vs. Whites and OR=1.67, 95% CI: 1.04–2.70 for Asian/Asian vs. White/White children).

Analysis with Hispanic ethnicity considered independent of race revealed that Hispanic origin of child and of parents was associated with increased risk of total childhood leukemia and ALL. Estimates and confidence intervals for parental Hispanic ethnicity were almost identical to those for child thus, in Table 4 we presented results for child's Hispanic ethnicity only.

A trend in risk of total childhood leukemia and ALL was observed comparing Hispanic children with both parents Hispanic, one parent Hispanic and non-Hispanic children, with the highest risk observed for children with both parents Hispanic (p-value for the trend < 0.0001).

Models with interactions between child's race and birth order, mother's age, SES proxies and child's Hispanic ethnicity were also considered. Interactions were detected between child's race and ethnicity (p-value=0.02). For further investigation, we combined child's race and Hispanic ethnicity. The highest OR for total childhood leukemia and for ALL was observed in Hispanic White and the lowest in non-Hispanic Black children compared to

non-Hispanic Whites. Non-Hispanic Asians were at slightly increased risk of total leukemia, ALL and AML, but with imprecise estimates (Table 5).

After performing complete case analyses, analyses were repeated using multiply imputed data. Results were very similar to complete case analysis. Some of these results are presented in Appendix Table 6 online.

## Discussion

Our results on the relationships between childhood leukemia and parental or child's race indicate that, compared to White race, being Black was highly protective for the development of childhood leukemia, particularly for ALL. This association was observed for paternal, maternal and child's races, regardless classification used. Although there were several studies that observed similar results [1 2 5–8 11 16], our study had a larger sample size, used several classification for child race, did not have issues with subject selection and, additionally, looked at maternal and paternal races.

One of possible explanation of this association, as some researchers have suggested, could be underlying SES [25 26]; others suggest the associations may be explained low birth weight among Blacks.[27–30] Many studies have shown that high birth weight was associated with increase in risk of childhood leukemia [31 32]; and consequently, low birth weight could have a protective effect on incidence of childhood leukemia. However, since we controlled for SES and for birth weight in our study, these factors are unlikely to account for the finding.

We observed an association between Asian race and AML after adjusting for census-based SES. Elevated risk of AML for Asian children was less precise after adjustment for father's education. The association of AML with Asian race was also observed by Reynolds et al (2002) in unadjusted analysis, and the estimate became imprecise when adjusted for father's education.[6] We cannot offer any specific explanation of this findings, but as noted by many researchers, AML may have different risk factors than ALL.

We observed that Hispanic ethnicity of parents and child was associated with approximately 1/3 increase in risk of total childhood leukemia and ALL. We observed a trend in the risk of total childhood leukemia and ALL when we compared Hispanic children with both parents Hispanic, one parent Hispanic and non-Hispanic children, with the highest risk observed for children with both parents Hispanic. Our findings are in line with results of other studies showing increased risk of childhood leukemia for children of Hispanic origin. [15] Incidence in several Latin American countries is the highest in the world: 5.65 cases/100,000 in Costa Rica, 5.54/100,000 in Ecuador, 4.43/100,000 in Uruguay.[33] Analysis with combined race and Hispanic ethnicity showed that non-Hispanic Black children had the lowest risk of childhood leukemia compared to White non-Hispanics. The highest risk was observed in White Hispanic children.

Some researchers have suggested that nutrition and diet could contribute to racial and ethnic differences in cancer incidence.[34] Emerging evidence suggests that genetic risk factors may also explain the markedly different risk of childhood leukemia in Hispanics and Blacks.



A recent study by Xu (2013) detected new susceptibility variants at 10p12.31–12.2 of the BMI1-PIP4K2A gene. This polymorphism is common in Hispanic ethnic groups and rare in Black populations; it could, at least partially, explain the findings.[35]

Another potential explanation of observed associations for race, ethnicity and childhood leukemia could be population mixing, in which immunologically naive susceptible individuals experience an increase in leukemia incidence as population increases.[20 36–41] Muirhead (1995) reported increasing incidence rates of childhood leukemia with increasing population density in three US metropolitan areas, including San Francisco, California.[42] During the years covered by this study California experienced population growth [43], potentially allowing the opportunity for susceptible individuals to be exposed to some new infectious agents for which their immune systems have not been modulated, resulting in a rare abnormal response of childhood leukemia.[44]

A major strength of our study was that data were obtained from nearly complete population registries and controls were randomly selected from the birth registry. Interview-based case-control studies are prone to selection bias. Meta-analysis by Slusky et al. (2012) has shown that interview-based case-control studies of childhood cancer could suffer from overrepresentation of Whites and under-representation of other races in participating controls.[45] This is not a case for registry-based non-contact studies. Since birth and cancer registries were independent of each other and participation of subjects was not required, selection bias due to participation was unlikely.

We matched on age and gender and adjusted for other potential confounders available in registries; however, we cannot completely exclude residual confounding due to other unknown/unmeasured factors.

Another advantage of this study was that the large sample size allowed us to carry out analyses for two main subtypes of childhood leukemia, ALL and AML. The risk patterns observed for AML were different from the risk patterns of ALL, and included higher risk for AML for Asian children. This highlights the importance of conducting disaggregated analyses by subtype.

Although there were two other studies on childhood leukemia conducted in California, our study does not have a large overlap with those. One of the two studies (Reynolds et al., 2001) included only children born in San Diego county during limited number of years (1988–1994) and had very limited sample size (90 cases/349 controls).[4] The Northern California Childhood Leukemia Study enrolled only newly diagnosed patients from hospitals in northern part of California since 1995.[46] Although cases have some overlap with our study, it is unlikely that we had a large overlap for controls because of their random selection in both studies. Our study had a larger sample size, included all leukemia cases from the whole California from 1988 to 2008 and did not have any subject selection issues. The results of the study are generalizable in any racially/ethnically heterogeneous population.

A potential bias could arise if controls of a particular race/ethnicity selectively moved out of state before their “pseudo-diagnosis” date and became cases there. For example, if Hispanic



children moved out of state and became leukemia cases outside of California, we would underestimate the association. These scenarios are very unlikely because the probability of controls moving out of California and subsequently developing childhood leukemia, a rare cancer, is quite low. Additionally, the literature indicates higher mobility of childhood leukemia cases compared to controls and not vice versa, which could not affect our results. [3 47 48]

One of a study limitations is potential misclassification on variables of interest and covariates. Misclassification of the outcome was unlikely due to the completeness and high accuracy of the CCR. Misclassification of race/ethnicity may have happened because it could be reported by parents, abstracted from a medical chart, or recorded by hospital staff based on their own observations.[16] Nonetheless, in validation studies in California where birth certificate data were compared with structured post-partum interviews, the sensitivity of birth records to correctly identify most racial and ethnic groups was greater than 94% with the exception of Native Americans.[49] Some misclassification of race and ethnicity is still possible due to categorization of these variables. We attempted to address this issue by examining sensitivity to re-classification, which, reassuringly, did not alter our results. Even if misclassification was present, we believe that it was not different for cases and controls, which would pull the estimates toward the null in the case of binary categorization.[50]

Another weakness of the study was missing data. To address the issue of missing values, at least partially, instead of only using records containing child's race that was missing for about a third of subjects, we constructed a classification of child's race using more complete information on mother's and father's races. For a majority of factors, missingness did not vary considerably by race and ethnicity. Black and Other races of children had slightly higher missing data on father's education. However, since information was missing mainly due to differences in the collected information between years rather than non-response and did not differ between cases and controls, the potential for biases was probably small, and the impact was mainly on the precision of estimates. We re-analyzed data using multiple imputations and obtained similar results with slightly narrower confidence intervals.

In summary, we found that children of Black race were at lower risk of childhood leukemia and ALL. Hispanic ethnicity was associated with high risk of childhood leukemia and ALL. A new finding was the association of Asian race and AML. Such ethnic and racial differences in incidence of childhood leukemia indicate that some genetic and environmental/cultural factors may be involved in etiology of childhood leukemia.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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### Summary box

#### What is already known on this subject?

Several studies noted decreased risk for childhood leukemia for Black children compared to Whites and increased risk for children of Hispanic ethnicity compared to non-Hispanics. Many of these studies had problems with subject selection, misclassification of race/ethnicity, and missing data.

#### What does this study add?

This registry-based study examined relationships between race/ethnicity and childhood leukemia in California. The large scale of the study allowed us to look at the association of race/ethnicity and the major subtypes of childhood leukemia (ALL and AML) which was not possible in previous research. The risk pattern observed for these two subtypes was quite different and revealed new associations.

Our study, being registry-based, did not suffer from selection bias as the majority of previous studies.

The study addressed potential misclassification of race/ethnicity by using several classifications for those.

Missing data was at least partially addressed by using not only the original child's race from birth records but also a reconstructed child's race/ethnicity based on mother's and father's race/ethnicity.

To our knowledge, this type of analysis was never applied in previous research. In addition, we repeated our analyses using multiply imputed datasets which also helped in addressing missing data issue.

**Table 1**  
Race of child based on combination of parental races, California birth registry, 1986–2007.

Mother's race	Father's race						Total
	Missing	White	Black	Asian	Am. Indian	PI & Other	
Missing	105	3	1	0	0	0	109
White	312	8889	147	114	44	28	9534
Black	41	41	556	5	2	1	646
Asian	36	217	19	852	1	3	1128
Am. Indian	4	48	4	0	14	0	70
PI & Other	8	22	5	3	0	51	89
<b>Total</b>	506	9220	732	974	61	83	11576

**Table 2**

Characteristics of study subjects, California birth registry, 1986–2007.

Variables	Cases (%)	Controls (%)	ALL # cases/controls <sup>a</sup>	AML # cases/controls <sup>a</sup>
<b>All</b>	<b>5788</b>	<b>5788</b>	<b>4721</b>	<b>852</b>
<b>Birth weight</b>				
< 2000 g	73 (1.3)	119 (2.1)	56/96	13/21
2000–3000 g	1009 (17.4)	1071 (18.5)	796/872	177/159
3000–4000 g	3941 (68.1)	3953 (68.3)	3232/3217	565/590
4000 g	764 (13.2)	644 (11.1)	636/535	97/82
Missing	1	1	1/1	0/0
<b>Birth order</b>				
First	2223 (38.5)	2333 (40.4)	1848/1896	309/356
Other	3559 (61.5)	3449 (59.7)	2878/2819	542/496
Missing	6	6	3/6	1/0
<b>Mother's age</b>				
< 25 years	1885 (32.6)	2067 (35.7)	1530/1689	296/301
25–35 years	3025 (52.3)	2973 (51.4)	2495/2430	415/435
35–45 years	863 (14.9)	740 (12.8)	687/594	136/116
45 years	14 (0.2)	7 (0.1)	9/7	0/0
Missing	1	1	0/1	0/0
<b>Father's education</b>				
<12 years	2552 (62.3)	2510 (61.7)	2056/2034	405/378
12 years	641 (15.7)	649 (15.9)	544/548	79/83
13–16 years	646 (15.8)	661 (16.2)	554/562	68/76
17 years	258 (6.3)	251 (6.2)	213/208	35/36
Missing (Not collected)*	1691 (1509)	1717 (1505)	1354/1369	265/279
<b>Socio-economic status (SES)</b>				
Low	2390 (50.4)	2380 (50.1)	1941/1923	351/377
Middle	969 (20.4)	990 (20.9)	816/809	128/135
High	1385 (29.2)	1377 (29.0)	1116/1131	209/191
Missing	1044	1041	848/858	164/149
<b>Child's race</b>				
White	4550 (81.9)	4339 (78.8)	3777/3559	610/622
Black	290 (5.2)	490 (8.9)	198/399	76/68
Asian	614 (11.1)	569 (10.3)	480/462	105/87
Am. Indians & Alaskan	81 (1.5)	82 (1.5)	67/67	12/9
PI & Other	24 (0.4)	27 (0.5)	19/22	5/4
Missing	229	281	180/212	44/62



Variables	Cases (%)	Controls (%)	ALL # cases/controls <sup>a</sup>	AML # cases/controls <sup>a</sup>
<b>All</b>	<b>5788</b>	<b>5788</b>	<b>4721</b>	<b>852</b>
<b>Mother's race</b>				
White	4859 (84.7)	4675 (81.6)	4020/3815	665/686
Black	230 (4.0)	416 (7.3)	154/342	62/56
Asian	578 (10.1)	550 (9.6)	457/452	94/79
Am. Indians & Alaskan	33 (0.6)	37 (0.7)	28/31	5/5
PI & Other	37 (0.6)	52 (0.9)	30/41	7/8
Missing	51	58	32/40	19/18
<b>Father's race</b>				
White	4706 (84.6)	4514 (81.9)	3910/3697	630/649
Black	274 (4.9)	458 (8.3)	186/375	73/62
Asian	507 (9.1)	467 (8.5)	389/381	91/71
Am. Indians & Alaskan	31 (0.6)	30 (0.5)	27/26	4/2
PI & Other	43 (0.8)	40(0.7)	31/32	10/6
Missing	227	279	178/210	44/62
<b>Hispanic origin of child</b>				
Both parents Hispanics	2481 (43.7)	2204 (39.1)	2074/1809	331/314
One parent Hispanic	633 (11.1)	603 (10.7)	513/485	93/94
Both parents non-Hispanic	2569 (45.2)	2827 (50.2)	2047/2304	411/417
Missing	105	154	87/123	17/27
<b>Hispanic origin of mother</b>				
Hispanic	2858 (49.6)	2559 (44.5)	2380/2082	387/381
Non-Hispanic	2899 (50.4)	3190 (55.5)	2317/2607	458/464
Missing	31	39	24/32	7/7
<b>Hispanic origin of father</b>				
Hispanic	2737 (49.1)	2452 (44.3)	2281/2021	368/341
Non-Hispanic	2842 (50.9)	3087 (55.7)	2269/2501	451/459
Missing	209	258	171/199	33/52

<sup>a</sup>Number of cases and controls for ALL and AML do not add up for the total number of cases and controls for childhood leukemia because there were few other subtypes in the dataset.

Conditional odds ratios (95% CIs) for childhood leukemia and race of child (N=7982), mother (N=8096) and father (N=7984) matched on child's age and sex and adjusted for birth order, birth weight, mother's age, and father's education. California birth registry, 1986–2007.

**Table 3**

	All types			AML		
	OR	95% CI	OR	95% CI	OR	95% CI
<b>Child's race (reconstructed)</b>						
White	1.00	-	1.00	-	1.00	-
Black	0.54	0.45 0.66	0.45	0.36 0.57	1.19	0.75 1.90
Asian	0.99	0.85 1.16	0.94	0.80 1.12	1.26	0.85 1.89
Am. Indian & PI	0.96	0.65 1.40	0.99	0.66 1.50	1.22	0.40 3.73
Other	0.75	0.37 1.49	0.79	0.36 1.69	0.74	0.12 4.51
<b>Child's race (combined parental race)</b>						
White/White	1.00	-	1.00	-	1.00	-
Black/Black	0.51	0.40 0.64	0.40	0.30 0.53	1.35	0.77 2.37
Asian/Asian	1.04	0.87 1.24	0.98	0.81 1.20	1.34	0.83 2.17
Am. Indian/Am. Indian	0.36	0.07 1.80	0.19	0.02 1.57	<i>a</i>	<i>a</i>
PI & Other/PI & Other	0.76	0.38 1.51	0.80	0.37 1.72	0.75	0.12 4.58
White/Black	0.74	0.52 1.05	0.69	0.46 1.04	1.10	0.50 2.39
White/Asian	0.89	0.68 1.17	0.85	0.63 1.15	1.13	0.57 2.21
White/Am. Indian	1.29	0.79 2.11	1.31	0.78 2.20	1.46	0.23 9.14
White/PI & Other	0.59	0.28 1.25	0.58	0.24 1.42	0.93	0.20 4.21
Black/Asian	0.31	0.10 0.94	0.23	0.07 0.80	<i>b</i>	<i>b</i>
Black/Am. Indian	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>
Black/PI & Other	0.31	0.03 2.82	0.30	0.03 2.73	<i>a</i>	<i>a</i>
Asian/Am. Indian	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	<i>b</i>	<i>b</i>
Asian/PI & Other	2.93	0.30 28.35	<i>a</i>	<i>a</i>	<i>b</i>	<i>b</i>
<b>Mother's race</b>						
White	1.00	-	1.00	-	1.00	-
Black	0.53	0.43 0.66	0.43	0.33 0.56	1.24	0.74 2.08
Asian	0.98	0.84 1.15	0.95	0.80 1.12	1.13	0.74 1.73

	All types			ALL			AML		
	OR	95% CI	OR	95% CI	OR	95% CI	OR	95% CI	
Am. Indian & PI	0.73	0.41	1.30	0.65	0.35	1.21	1.22	0.20	7.55
Other	0.67	0.38	1.18	0.72	0.39	1.35	0.69	0.16	2.92
<b>Father's race</b>									
White	1.00	-	-	1.00	-	-	1.00	-	-
Black	0.55	0.45	0.67	0.45	0.36	0.57	1.30	0.80	2.11
Asian	1.01	0.86	1.20	0.95	0.79	1.14	1.42	0.92	2.19
Am. Indian & PI	1.07	0.57	2.01	1.18	0.60	2.31	1.03	0.13	8.17
Other	0.83	0.49	1.42	0.89	0.49	1.62	0.93	0.24	3.52

<sup>a</sup>Not estimable due to small cell counts

<sup>b</sup>No observations

Conditional odds ratios (95% CIs) for childhood leukemia and Hispanic origin of child matched on child's age and sex and adjusted for birth order, birth weight, mother's age, and father's education (N=8078). California birth registry, 1986–2007.

**Table 4**

	All types			ALL			AML		
	OR	95% CI	OR	95% CI	OR	95% CI	OR	95% CI	
<b>Hispanic ethnicity of child</b>									
Non-Hispanic	1.00	-	1.00	-	1.00	-	1.00	-	
Hispanic	1.29	1.17 1.43	1.37	1.22 1.52	0.98	0.74 1.30			
<b>Parental ethnicity</b>									
Non-Hispanic	1.00	-	1.00	-	1.00	-	1.00	-	
One parent Hispanic	1.17	1.00 1.38	1.20	1.01 1.44	1.08	0.70 1.64			
Both parents Hispanics <sup>a</sup>	1.33	1.20 1.49	1.42	1.26 1.60	0.96	0.71 1.29			

<sup>a</sup> trend test p-value < 0.0001

**Table 5**

Conditional odds ratios (95% CIs) for childhood leukemia and combined child's race and Hispanic ethnicity, adjusted for birth order, birth weight, mother's age, father's education and matched on child's age and sex (N=7968). California birth registry, 1986–2007.

Combined child's race and Hispanic ethnicity	All types			ALL			AML			
	OR	95% CI	OR	95% CI	OR	95% CI	OR	95% CI	OR	95% CI
Non-Hispanic White	1	-	1	-	1	-	1	-	1	-
Hispanic White	1.23	1.1 1.38	1.27	1.12 1.44	1.44	1.11 0.8	1.11	0.8 1.53		
Non-Hispanic Black	0.58	0.46 0.72	0.46	0.36 0.6	0.6	1.23 0.72	1.23	0.72 2.11		
Hispanic Black	1.05	0.65 1.69	0.95	0.55 1.65	1.65	1.66 0.55	1.66	0.55 4.98		
Non-Hispanic Asian	1.16	0.97 1.37	1.1	0.92 1.33	1.39	0.88 2.19	1.39	0.88 2.19		
Hispanic Asian	0.91	0.58 1.42	0.76	0.45 1.27	1.53	0.55 4.28	1.53	0.55 4.28		
Non-Hispanic Am. Ind.	1.13	0.65 1.96	1.2	0.67 2.15	2.17	0.39 12.08	2.17	0.39 12.08		
Hispanic Am. Ind.	1.12	0.65 1.91	1.48	0.74 2.98	0.81	0.17 3.79	0.81	0.17 3.79		
Non-Hispanic PI &Other	0.72	0.32 1.64	0.85	0.31 2.33	0.33	0.03 3.26	0.33	0.03 3.26		
Hispanic PI &Other	1.32	0.35 4.98	0.92	0.39 2.15	2.15	a	a	a	a	a