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Short Communication Perinatal risk factors for hepatoblastoma

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We confirmed the strong association of hepatoblastoma with very low birth weight (relative risk $< 1000 \text{ g vs} \ge 2000 \text{ g} = 25.6$; 95% confidence interval: 7.70–85.0) and demonstrated independent associations with congenital abnormalities and maternal Asian race in a population-based Minnesota study that included 36 cases and 7788 controls.

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Hepatoblastoma is a rare embryonal tumour that comprises most cases of liver cancer in children aged 0-5 years in the United States (Ries *et al*, 1999). Although its causes are mostly obscure, it is apparent that very low birth weight (VLBW), generally defined as <1500 g, is a potent risk factor (Tanimura *et al*, 1998; Reynolds *et al*, 2004; Ansell *et al*, 2005; McLaughlin *et al*, 2006). Hepatoblastoma incidence doubled between 1975 and 1999 (Spector *et al*, 2004), possibly related to the concomitant rise in prevalence of VLBW infants and a marked drop in their mortality (CDC, 2002). These observations may indicate that some component of treatment for prematurity is carcinogenic or, alternatively, that the aetiology may overlap with that of VLBW. To expand a sparse literature, we examined the relation between hepatoblastoma and birth characteristics in Minnesota.

MATERIALS AND METHODS

The methods used in this case-cohort study have previously been described (Puumala *et al*, 2008). Briefly, we matched Minnesota Cancer Surveillance System (MCSS) records of incident first cancers diagnosed in children aged 28 days to 14 years during 1988–2004 to birth records using probabilistic record linkage (Jaro, 1995). For each of the 2188 successfully linked cases (out of 2655 total) we randomly selected four birth records of children born in the same year and who survived at least 28 days past birth; this comparison group of 8752 subjects is referred to as the subcohort. In this analysis, matching cases were compared to all subcohort members born in 1982 (i.e., the earliest year a case would have been born) or later to improve study power.

Exposure variables derived from Minnesota birth records are shown in Table 1. Some variables were not available during the entire study period. Birth weight was divided into categories of <1000, 1000-1999, and ≥ 2000 g to maximize the number of cases

in each stratum. Race was classified as white or non-white. Other variables were categorized using customary cut offs (Ries *et al*, 1999).

We calculated hazard ratios (HRs) and 95% confidence intervals (CIs) using stratified Cox regression models using SAS 9.1 (SAS institute Inc., Cary, NC, USA) (Langholz and Jiao, 2007). The number of cases precluded a full multivariate model. However, birth weight is a known risk factor; all variables were modelled adjusting for birth weight as well as for year of birth and sex.

RESULTS

Of 39 cases of hepatoblastoma identified by MCSS, 36 (92.3%) linked to birth records. Cases were compared to 7788 members of the subcohort. Exposure frequencies and adjusted HRs are presented in Table 1; where cell size was <4, we reported frequencies but did not calculate HRs. There were strong associations with birth weight <1000 g (HR = 25.6; 95% CI: 7.70-85.0) and 1000-1999 g (HR = 9.15; 95% CI: 3.09-27.1) compared to \geq 2000 g, their magnitude with each category of low birth weight was lessened, but remained highly significant, when adjusting for covariates (data not shown). Significant univariate associations were noted with maternal and paternal Hispanic ethnicity, maternal birthplace outside the US, intrauterine procedures during pregnancy, male sex, gestational age <37 weeks, 1 and 5 min apgar scores, assisted ventilation, and congenital abnormalities (data not shown). However, only associations with congenital abnormalities (HR = 5.87; 95% CI: 1.88-18.3), paternal Hispanic ethnicity (HR = 4.18; 95% CI: 1.22-14.3), and maternal birthplace outside the US (HR = 3.55; 95% CI: 1.51-8.32) remained significant after adjustment. The association of foreign maternal birthplace reflected the disproportionate number of cases with mothers from Southeast Asia. Accordingly, there was a significant adjusted association of maternal Asian race with hepatoblastoma (HR = 3.86; 95% CI: 1.30 – 11.52).

DISCUSSION

Very low birth weight has emerged over the past 15 years as a potent risk factor for hepatoblastoma (Feusner *et al*, 1998;

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$\begin{array}{c} 0 \\ -2 \\ \geqslant 3 \\ \text{transmitted} \text{since letting bith (upper)} \end{array}$	3062 (39.9) 3799 (49.5) 808 (10.5)	11 (30.6)
Interval since last live birth (years) No prior birth ≤3 >3	3062 (40.1) 2966 (38.9) 1600 (21.0)	18 (50.0)
Prior fetal losses None Any	5919 (77.5) 1717 (22.5)	
Parental demographics Mother's age (years) <25 25-29 ≥30 Father's age (years) <30 ≥30 Mother's race White Non-white	2358 (30.3) 2708 (34.8) 2722 (35.0) 3347 (47.5) 3693 (52.5) 7010 (90.9) 700 (9.1)	10 (27.8) 15 (41.7) 11 (33.3) 22 (66.7)
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istics with hepatoblastoma adjusted for birth weight, birth year, and sex

Subcohort

(n = 7788)

N (%)

7606 (98.0)

124 (1.6)

31 (0.4)

3778 (48.5)

4010 (51.5)

6888 (91.6)

631 (8.4)

252 (3.4)

5427 (72.4)

1821 (24.3)

7592 (97.5)

196 (2.5)

5725 (76.6)

1746 (23.4)

7192 (96.4)

265 (3.6)

4549 (98.3)

79 (1.7)

7676 (98.6)

||2 (|.4)

4530 (67.9)

1488 (22.3)

658 (9.9)

1029 (26.7)

1295 (33.6)

1526 (39.6)

7271 (97.4)

193 (2.6)

3439 (71.6)

1361 (28.4)

6162 (82.0)

1355 (18.0)

Cases

(n = 36)

N (%)

28 (77.8)

4 (11.1)

4 (11.1)

|| (30.6)

25 (69.4)

25 (69.4)

11 (30.6)

0 (0.0)

29 (80.6)

7 (19.4)

34 (94.4)

2 (5.6)

19 (52.8)

17 (47.2)

29 (80.6)

25 (86.2)

4 (13.8)

31 (86.1)

5 (13.9)

24 (75.0)

4 (12.5)

4(12.5)

7 (25.9)

10 (37.0)

10 (37.0)

32 (88.9)

4(||.|)

20 (66.7)

10 (33.3)

26 (72.2)

10 (27.8)

7 (19.4)

HR^a

9.15

2.03

1.70

1.99

2.24

2.00

5.87

0.61

1.11

1.36

1.40

2.39

1 32

126

0.62

2.07

0.60

1.27

1.31

1.55

1.56

25.6

95% CI

3.09-27.1

7.70-85.0

0.98-4.19

0.56-5.13

0.95-4.16

0.70-7.11

049-819

1.88-18.3

0.21-1.78

0.36-3.43

0.48-3.83

0.48 - 4.07

0.68-8.41

0.60 - 2.93

0.58 - 2.77

0.28 - 1.35

0.87 - 4.92

0.24 - 1.49

0.53 - 3.08

057 - 301

0.74-3.25

0.63-3.89

Minnesota, 1982-2004

Characteristic

Birth characteristics Birth weight (g) > 2000

< 1000

Male

 ≥ 37 < 37

Average

Large Multiple birth

No

Yes

>7

≼7

>7

<7

Yes

No

Yes

Sex Female

1000-1999

Gestational age (weeks)^b

Size for gestational age Small

Apgar score (1 min)

Apgar score (5 min)

Assisted ventilation No

Congenital abnormality

Index pregnancy history Adequacy of prenatal care

Intermediate

Weight gain (pounds)

Intrauterine procedures

Caesarean section

Maternal reproductive history

Induction of labor

Type of delivery

Prior live births

Vaginal

Inadequate

≤24

> 30

No

Yes

No

Yes

25 - 30

Adequate

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Table I Associations of demographic, pregnancy, and birth character-**Table I** (Continued)

Characteristic	Subcohort (n = 7788) N (%)	Cases (n=36) N (%)	HRª	95% CI
Mother's ethnicity ^c				
Non-Hispanic	4475 (96.5)	26 (86.7)	1	
Hispanic	160 (3.5)	4 (13.3)	2.96	0.88-10.0
Mother's birthplace	()			
United States	7297 (93.8)	28 (77.8)	1	
Elsewhere	485 (6.2)	8 (22.2)	3.55	1.51-8.32
Father's race				
White	6381 (92.5)	28 (87.5)	1	
Nonwhite	521 (7.5)	4 (12.5)	1.44	0.49-4.29
Father's ethnicity ^c	· · · ·	. ,		
Non-Hispanic	3996 (97.0)	24 (85.7)	1	
Hispanic	125 (3.0)	4 (14.3)	4.18	1.22-14.3
Mother's education (years)				
<12	782 (10.6)	8 (22.9)	1.60	0.64-4.02
12	2706 (36.8)	14 (40.0)	1	
> 12	3867 (52.6)	13 (37.1)	0.61	0.28-1.32
Father's education (years)	()	. /		
≤12	2840 (44.2)	13 (43.3)	I	
>12	3587 (55.8)	17 (56.7)	1.03	0.49-2.16

Abbreviations: CI = confidence interval; HR = Hazard ratios; LNMP = last normal menstrual period ^aHazard ratios adjusted for birth weight, birth year, and sex. ^bImputed gestational age based on LNMP when available and physician's estimate when LNMP was missing. ^cCollected in birth years 1989-2004.

Tanimura et al, 1998; Reynolds et al, 2004; Ansell et al, 2005; McLaughlin et al, 2006). We have now confirmed this association in the paediatric population of Minnesota. Unadjusted relative risks between 16 and 70 comparing very low to moderate birth weight children have been reported in studies from the United Kingdom, Japan, and now three US states (Figure 1). This striking association thus appears across industrialized nations.

Rather than being causative per se, VLBW likely signals the involvement of correlated factors. Limited multivariate analyses in this study and others (Reynolds et al, 2004; McLaughlin et al, 2006) have begun to tease apart the role of other characteristics from that of VLBW. Notably, preterm birth (<37 weeks) was not an independent risk factor in any of the studies after adjustment.

We found a strong association with congenital abnormalities, which was attenuated but still present when controlling for VLBW. One specific abnormality, an omphalocele, was recorded, which may be indicative of Beckwith-Wiedemann syndrome, an overgrowth disorder which is known to increase the risk of hepatoblastoma (Ranzini et al, 1997; DeBaun and Tucker, 1998). The remaining five abnormalities were nonspecific (one central nervous system, one urogenital, and three 'others'). Higher than expected proportions of congenital abnormalities have previously been noted among cases (Narod et al, 1997; Ansell et al, 2005), but without factoring in birth weight.

The association of maternal Asian race, specifically of Southeast Asian ancestry, with hepatoblastoma was unexpected. Similar findings were not noted in studies in California (Reynolds et al, 2004) and New York (McLaughlin et al, 2006), and there was not an elevated rate in Ho Chi Minh City, Vietnam (Nguyen et al, 2000). Our observation was therefore novel and independent of birth weight. We also observed an independent association of paternal, but not maternal, Hispanic ethnicity. This finding may have been an artifact, as its significance was dependent on two cases for which paternal ethnicity was missing.

Maternal hypertension (Roberts et al, 2003), maternal tobacco use (Horta et al, 1997), and conception by assisted reproductive technology (Schieve et al, 2002) are known to reduce birth weight and have been examined in other studies. An excess of maternal



Study *n* cases Comparison

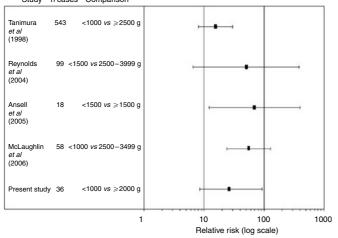


Figure I Relative risk of hepatoblastoma comparing very low to moderate birth weight children in five studies.

pre-eclampsia, without adjustment for VLBW, has been noted (Ansell *et al*, 2005). Three previous studies have found that maternal smoking significantly raised offspring risk (Pang *et al*, 2003; Sorahan and Lancashire, 2004; McLaughlin *et al*, 2006), whereas a fourth did not (Buckley *et al*, 1989); associations remained after adjustment for VLBW in two studies (Pang *et al*, 2003; Spector and Ross, 2003; McLaughlin *et al*, 2006). Lastly, a ninefold increased risk of hepatoblastoma was reported among children born following infertility treatment, which was independent of birth weight (McLaughlin *et al*, 2006). That maternal smoking and conception by assisted reproductive technology

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remain as risk factors after adjustment for VLBW suggests that if causal relationships exist, they operate independently of birth weight.

Use of population-based registry data was the major strength of this study, as any misclassification would most likely be nondifferential and HRs would be underestimated. Birth characteristics and delivery methods are reliably recorded in birth records whereas other factors, including congenital abnormalities, are substantially underreported (Northam and Knapp, 2006), contributing to sparse data for several variables. Although cases that occurred among outmigrating children or those who resided in Minnesota during 1982–1987 would have been missed, this occurrence is unlikely given the very low incidence of hepatoblastoma. Lastly, the small number of cases resulted in a reduced ability to perform multivariate analyses.

The strong association of hepatoblastoma with VLBW has been robust to adjustment for other factors. That control for several prenatal determinants of small infant size has not explained the VLBW association may indicate that postnatal treatment is the causative correlate. Although three very small casecontrol studies of neonatal medical history among VLBW infants preliminarily suggest greater oxygen exposure in cases (Maruyama *et al*, 1999, 2000; Oue *et al*, 2003), larger studies are plainly required. Therefore, a multicenter case-control study has been initiated (National Institutes of Health Grant R01CA111355; L Spector, Principal Investigator) that will examine risk factors for hepatoblastoma, with a special focus on VLBW infants.

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