“Neonatal Thyroxine Level and Perchlorate in Drinking Water”

By

Zili Li, MD, MPH\textsuperscript{1,2}
Feng Xiao Li, MD, PhD\textsuperscript{1}
Dan Byrd, PhD\textsuperscript{3}
Gloria M. Deyhle, RN\textsuperscript{4}
David E. Sesser, BA\textsuperscript{5}
Michael R. Skeels, PhD, MPH\textsuperscript{5}
and Steven H. Lamm, MD\textsuperscript{1,2}

1 Consultants in Epidemiology and Occupational Health, Inc.
2 Preventive Medicine Residency Program, Johns Hopkins School of Hygiene and Public Health
3 Consultants in Toxicology, Risk Assessment, and Product Safety
4 Newborn Screening Program, Nevada State Health Division
5 Oregon State Public Health Laboratory

Funded by American Pacific Corporation.
Submitted to the Journal of Occupational and Environmental Medicine, Nov 15, 1999
ABSTRACT:

Environmental contamination of drinking water has been observed for perchlorate, a chemical able to affect thyroid function. This study examines whether that exposure has affected the thyroid function of newborns. Neonatal blood thyroxine (T₄) levels for days 1-4 of life have been compared for newborns from the city of Las Vegas, Nevada, which has perchlorate in its drinking water, and those from the city of Reno, Nevada, which does not [detection limit 4 ug/l (ppb)]. This study is based on blood T₄ analyses from over 23,000 newborns in these two cities during the period April 1998 through June 1999. No difference was found in the mean blood T₄ levels of the newborns from these two cities. Drinking water perchlorate levels measured monthly for Las Vegas ranged during this study period from non-detectable for eight months to levels of 9 to 15 ppb for seven months. Temporal differences in mean T₄ level were noted in both cities but were unrelated to the perchlorate exposure. This study was sufficiently sensitive to detect the effects of gender, birth weight, and the day of life on which the blood sample was taken on the neonatal T₄ level but detected no effect from environmental exposures to perchlorate that ranged up to 15 ug/l (ppb).

INTRODUCTION:

Perchlorate competitively inhibits the uptake of iodide into the thyroid gland¹ [Stanbury and Wyngaarden, 1952]. The pharmacology of perchlorate has been well described² (Wolff, 1998).
Blockade of iodine uptake leads to increased serum thyrotropin (TSH) levels and decreased blood thyroxine (T₄) levels. Perchlorate is used medically to treat hyperthyroidism of various etiologies (e.g., Graves' Disease, amiodarone toxicity) at doses of 900 mg/day or less. Studies of workers showed no effect on thyroid function at 34 mg/day³ (Lamm et al., 1999). Studies of subjects ingesting oral perchlorate doses of 10 mg/day for two weeks showed no effect on thyroid function in spite of a demonstrated partial inhibition of iodide uptake⁴ (Lawrence et al., 1999). Perchlorate has been shown in guinea pigs to cause a fetal goiter at dosages (1 % in water) that did not cause a maternal goiter⁵ (Postel, 1957).

Several public water supplies in California and southern Nevada contain perchlorate in the 4-16 ppb concentration range. Newborns are screened for metabolic diseases in hospitals and doctor's offices in a mandatory state-run program. These programs include the measurement of neonatal blood thyroxine (T₄) levels as a screening procedure for congenital hypothyroidism. An analysis of the data from these programs for California and Nevada has demonstrated that the incidence of congenital hypothyroidism among children born in areas with perchlorate in drinking water did not differ from the incidence in perchlorate-free areas⁶ [Lamm and Doemland, 1999]. The present study supplements this observation by examining for newborns in Nevada whether there is an effect on neonatal thyroid status as determined by the neonatal T₄ level associated with the presence of perchlorate the drinking water.

MATERIALS AND METHODS:

The drinking water supply for Clark County, Nevada, and its largest city Las Vegas, comes from a part of Lake Mead that is contaminated with perchlorate (ClO₄⁻). In some months, the level
is below the detection limit of 4 ppb (ug/l), and in other months the concentration is measured in
the range of 4.1 – 15 ppb. This is the only known and documented source of perchlorate
contamination of a public water supply in the State of Nevada. The city of Reno, the second largest
city in Nevada, is 450 miles away in the foothills of Sierra Nevada mountains and has an
independent source of drinking water that has no evidence of perchlorate contamination.

We here compare for the period April 1998 through June 1999 the mean monthly T\textsubscript{4} levels
of newborns in two urban areas in Nevada, one (Las Vegas) with perchlorate in the drinking water
supply and one (Reno) without perchlorate in the drinking water supply. The population of Las
Vegas is about three times that of Reno. The study cohorts consist of all newborns who had blood
samples submitted within four days of birth, who had birth weights between 2,500 and 4,500
grams, and who had not been admitted to a Neonatal Intensive Care Unit (NICU) by the time of
blood sample collection. Demographic data on the newborns included date of birth, date of sample
collection, gender, birth weight, birthplace, and office address of each newborn's pediatrician.

Outcome Variable

The outcome of interest was neonatal blood T\textsubscript{4} level, as measured by the Oregon State
Public Health Laboratory (OSPHL) for the Nevada State Health Department. All states measure
neonatal thyroid status (either T\textsubscript{4} or TSH) in heelstick blood obtained in the hospital during the first
few days of life, as part of the legally-mandated neonatal metabolic diseases screening program. In
Nevada, over 80 % of newborns also have a later blood sample submitted for T\textsubscript{4} analysis from the
physician’s office at a follow-up visit.
OSPHL provides these analyses for the Nevada State Health Division program, including births in both Las Vegas and Reno. The laboratory used a radio-immunoassay method\textsuperscript{7} (Murphy, 1964) to analyze T\textsubscript{4} levels in the congenital hypothyroidism screening program. The blood T\textsubscript{4} data were analyzed as a continuous variable. Blood thyroxine (T\textsubscript{4}) levels depend on the stability of the particular assay method in a laboratory. The Oregon State Public Health Laboratory established stabilization of their T\textsubscript{4} assay most recently in March 1998, thus defining the time period of observation for this study as beginning in April 1998.

Exposure Variable

The exposure factor of primary interest was perchlorate in drinking water, which occurred in Las Vegas and not in Reno. Lake Mead is the sole source for the public water supply of Las Vegas. Perchlorate in the intake of water to the city of Las Vegas occurred when the turbulent conditions of Lake Mead overcame the thermal stratification of the lake and thus presented contaminated water to the intake of the water supply. Monthly measurements of the perchlorate levels in Las Vegas finished water have been made since July 1997 by the Southern Nevada Water Authority, using a method with a detection limit of 4 ppb (ug/l) developed by the California Department of Health Services, Sanitation and Radiation Laboratory in April 1997\textsuperscript{8} (Okamoto et al, 1999). Perchlorate was detected in the Las Vegas drinking water during seven of the 15 months in this study period. Analyses have been performed comparing the T\textsubscript{4} levels of the children born in the seven study months in which perchlorate was detected in the Las Vegas water supply (time period A) and those born in the eight study months in which perchlorate was not detected in the Las
Vegas water supply (time period B). Perchlorate was also detected in the Las Vegas water supply during the nine months prior to the study period.

The water supply in Reno had no connection with Lake Mead or the sources of perchlorate in Lake Mead. Instead, Reno derives 80 % of its supply from the high mountains via Lake Tahoe and the Truckee River and 20% from local wells in the Reno area. Tests of the water sources for Reno, using the same laboratory method, detected no perchlorate\(^9\) (Auckly, 1999).

Statistical Methods

The distributions of demographic variables of the newborns of the two cities were compared using a chi-square test for categorical variables and a t-test for continuous variables. The mean T4 levels between the 17,308 newborns in Las Vegas and the 5,882 newborns in Reno were compared in a univariate analysis, both crude and stratified by time period. A multivariate analysis was performed with T4 level as the outcome variable, city and time period as the main effect variables, and gender, birth weight and age at time of sample collection as the covariates. An interaction term between city and time period was included in the model as a marker for perchlorate exposure. All statistical tests were two-sided, and the cut-off probability for a type I error was 0.05. All analyses used Stata statistical software (version 5.0).

Initial analyses used the T4 levels of the blood samples collected in the hospital up through the newborn’s fourth day of life and the perchlorate level in the Las Vegas drinking water during the month of birth, expressed in ug/l. Subsequent analyses used the same T4 levels and the
cumulative perchlorate levels, expressed in ug/l-months, for either the nine months of pregnancy or the time period equivalent to the first trimester of pregnancy. Additional analysis used the T₄ levels of the bloods submitted from the physician’s offices and examined the differences in the mean T₄ level of the infants of the two cities by the age of the newborn in days for the first sixty post-natal days.

RESULTS:

There were 23,305 newborns from Las Vegas and Reno who met the inclusion criteria during the 15-month period between April 1, 1998 and June 30, 1999, and 23,190 of them (99.5%) had valid neonatal T₄ measurements. The values for 115 newborns were excluded either because of unsatisfactory blood samples (N = 113) or invalid T₄ measurement (N = 2). The final study cohort consisted of 17,308 newborns from Las Vegas and 5,882 from Reno.

Table 1 displays the distributions and corresponding p-values of three potential confounding factors for the two cities. The newborns from Las Vegas and Reno did not differ in the gender distribution but did differ in average birth weight and in age in days at time of sample collection. The percentage of males in Las Vegas resembled that in Reno, 51.3% vs 51.1% (p=0.779). The average birth weight of 3,379 grams in Las Vegas exceeded the average birth weight of 3,365 grams in Reno (p=0.033). The mean number of days between the day of birth and the day of sample collection also differed significantly between Las Vegas and Reno, 1.20 vs 1.45 days (p<0.001).
Figure 1 displays for the fifteen month period (April 1998 – June 1999) the monthly mean neonatal T₄ values for Las Vegas and for Reno. The monthly mean neonatal T₄ values are about 17 ug/dl with little difference between the two cities and little variation across the time period. No particular seasonal variation is evident. Figure 1 also displays the perchlorate level in the Las Vegas drinking water for each study month. Perchlorate was detected in the drinking water in seven of the fifteen months ranging between 9 and 15 ppb (ug/l). Perchlorate was not detected in the other eight months. The limit of detection was 4 ppb. Figure 1 shows no co-variation between the perchlorate level in the Las Vegas drinking water and the mean monthly neonatal T₄ level in either Las Vegas or Reno.

Figure 1 shows that some months lacked detectable perchlorate in the Las Vegas water, which allowed the introduction of a second exposure-related variable. We designated time period A as the months with detectable perchlorate in Las Vegas water and time period B as the months without detectable perchlorate in Las Vegas water.

Table 2 shows that the mean T₄ level was 17.11 ug/dl in Las Vegas and 17.12 ug/dl in Reno (p = 0.901) in the analysis of the crude data. Stratification of the data by time period (period A and period B) showed no difference between the means of the two cities during either period A or period B, i.e., no place effect. For each city, there was a difference in their mean T₄ levels for period A and period B, i.e., a period effect. The period effect was similar in both cities (p = 0.699).

Table 3 displays the results of a multivariate analysis. As in the crude analysis, the multivariate analysis demonstrated no statistically significant difference in mean T₄ values with
respect to place (i.e., Las Vegas vs Reno; p = 0.407); However, it did show a significant difference with respect to period (i.e., B vs. A; p = 0.000). Since there was no significant interaction between place and period, the period effect could not be explained by the presence of perchlorate in only one of the two places. The above analysis was controlled for gender, birth weight, and age at time of sample collection, all of which were significant co-variables (p = 0.000 for each).

The above analyses have used the mean monthly T4 level as the summary statistic to represent the T4 distribution in either city for any particular month. This statistic represents the central tendency of the distribution. The lower tail (or extreme) of the distribution might also be used to represent the T4 distribution in either city for any particular month. We have examined this by comparing the 10th percentile levels for either city during periods A and B. We similarly found that the period effects in the two cities were indistinguishable. Thus, the two cities did not differ in their prevalences of low neonatal blood T4 levels with respect to presence of perchlorate in the Las Vegas drinking water.

Perchlorate concentration data exist for Las Vegas for the nine months preceding this study period. An estimate can thus be made on a monthly basis of the perchlorate exposure those newborns would have had during their full period of pregnancy. The Las Vegas newborns during this study period would have had perchlorate exposures that ranged between 9 ppb-months (ug/l-months) and 83 ppb-months (mean = 48 ppb-months), and the Reno newborns during this period are presumed to have had zero ppb-months. For each month, the difference between mean neonatal T4 levels of the two cities was examined along with the cumulative perchlorate exposure estimated to have been experienced by those newborns. Linear regression analysis (Figure 2) showed no
evidence of an association (slope = -0.0003; $R^2 = 0.002$). Similarly, when estimated exposure during the first trimester of pregnancy was used as the exposure variable (0.0-36 ppb-months; mean = 20 ppb-months), linear regression analysis also showed no evidence of an association (slope = 0.0008; $R^2 = 0.002$).

Most newborns in Nevada (82 %) have a blood specimen taken in the physician’s office for a follow-up T4 measurement, 99 % of which are within the first 60 post-natal days. Figure 3 shows on a daily basis the difference between the mean T4 levels of the infants from Reno and those from Las Vegas. Throughout the first 60 days, there appears to be no systematic difference between the mean T4 levels in the two cities.

DISCUSSION:

The primary purpose of the study design was to compare the neonatal T4 levels in Las Vegas, where perchlorate was present in the drinking water, to the neonatal T4 levels in Reno, where perchlorate was not present. This comparison showed that perchlorate at environmental levels of up to 15 ppb did not affect neonatal T4 levels during the study period. Our study design with respect to T4 levels eliminated city/rural confounders by restricting the study populations to urban births, inter-laboratory variation by using data from the same laboratory, and seasonal effects by taking contemporaneous data. Our analysis controlled for the confounders gender, birth weight, and age at time of blood sampling. We found no difference in the mean T4 levels between the two cities, though we did find a period effect in the multivariate analysis. Since the period effect was
essentially the same in Las Vegas and in Reno, it could not be explained by the presence of perchlorate only in the Las Vegas drinking water. These findings demonstrate the importance of having a comparison population. The most probable explanation relates to minor temporal changes in laboratory procedures or materials or seasonal variation.

Data do not exist on how many of the 23,190 women consumed public water during their pregnancy. However, there is no reason to believe that that consumption would have changed during the time period of this study. Therefore, it is assumed that the temporal pattern of perchlorate concentration in the Las Vegas drinking water mirrors the temporal pattern of perchlorate consumption by the maternal population in Las Vegas. EPA has traditionally assumed that each adult consumes two liters of water per day, without making any specific distinction as to whether they are pregnant or not. We used the same two-liter per day assumption in our calculations. Any variation from that assumption, however, should be uniform over the time period of our study and not cause a differential effect on the analysis. Nonetheless, dose reconstruction can be performed using these assumptions. The cumulative exposures ranged from 0.9 to 4.2 mg with a mean of 2.2 mg during the pregnancy (i.e., 2.2 mg perchlorate per gestational period) and from 0 to 1.7 mg with a mean of 0.9 mg during the first trimester.

There is little literature on the effects of perchlorate exposure during human pregnancy. Our earlier paper\textsuperscript{6} (Lamm and Doemland, 1999) demonstrated no increase in the incidence of congenital hypothyroidism in areas of California and Southern Nevada with perchlorate in the drinking water, also at levels up to 16 ppb. These two papers have shown no effect from low environmental contamination levels with exposures of up to 32 \mu g/day. On the other end of the fetal
exposure spectrum is the experience with perchlorate as a therapeutic agent. In 1960, Crooks and Wayne\textsuperscript{10} reported in Lancet their results from treating 12 women with thyrotoxicosis of pregnancy with potassium perchlorate at exposures of 600 to 2000 mg/day. They noted that “one of the infants had a very slight enlargement of the thyroid gland which disappeared within 6 weeks. The remainder showed no abnormality of any kind.”

While no analysis can rule out a small effect of these perchlorate exposures on neonatal T\textsubscript{4} levels, the current study does rule out a significant effect. As this study did find that mean blood T\textsubscript{4} levels differed significantly with sex, birth weight, and time after birth, it likely had sufficient statistical power to detect any significant change induced by these perchlorate exposures. We conclude that perchlorate in drinking water at a level of up to 15 ppb has had no detectable effect on neonatal T\textsubscript{4} levels in this population.
REFERENCES


4. Lawrence JS, Lamm SH, Pino S, and Braverman LE. The Effect of Short-Term Low Dose Perchlorate on Various Aspects of Thyroid Function. (Under review).


Figure 1:

* Detection limit was 4 ppb (ug/l).
Figure 2

Difference in Mean Neonatal T4 Levels (ug/ml) between Reno and Las Vegas by Estimated 9 month Cumulative Perchlorate Exposure in Las Vegas

\[ y = -0.0003x \]

\[ R^2 = 0.002 \]
Figure 3

Difference in Mean T4 levels (µg/ml) between Reno and Las Vegas by Age in Days (1-60)
Table 1:

<table>
<thead>
<tr>
<th></th>
<th>Las Vegas (n=17,308)</th>
<th>Reno (n=5,882)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gender (% Male)</td>
<td>51.3%</td>
<td>51.1%</td>
<td>0.779</td>
</tr>
<tr>
<td>Mean birth weight (grams)</td>
<td>3,379</td>
<td>3,365</td>
<td>0.033*</td>
</tr>
<tr>
<td>Mean age at time of sample collection (days)</td>
<td>1.20</td>
<td>1.45</td>
<td>0.0000*</td>
</tr>
</tbody>
</table>

* indicate that the differences between the pairs are statistically significant at less than p= 0.05 level

Table 2:
Analytic Comparisons of Mean Monthly Neonatal T4 Levels of Las Vegas and Reno, Nevada (April 1998-June 1999), directly and stratified by time periods

<table>
<thead>
<tr>
<th>City-specific mean T₄ levels (ug/dl)</th>
<th>Place</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Las Vegas</td>
<td>Reno</td>
</tr>
<tr>
<td>Total sample</td>
<td>17.11</td>
<td>17.12</td>
</tr>
<tr>
<td>Stratified by time Period*</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Time Period A (7 months)</td>
<td>16.78</td>
<td>16.77</td>
</tr>
<tr>
<td>Time Period B (8 months)</td>
<td>17.37</td>
<td>17.41</td>
</tr>
<tr>
<td>Difference between period A &amp; B</td>
<td>-0.59</td>
<td>-0.64</td>
</tr>
</tbody>
</table>

* During the 15-month study period, time period A was defined as a combination of those months at which perchlorate levels were detected in the drinking water in Las Vegas, Nevada whereas time period B included the remaining months at which perchlorate level were not detected in the drinking water in Las Vegas, Nevada.
Table 3.

Multivariate analysis of effect of city, time period and their interaction on neonatal T₄ levels after adjusting for gender birth weight, and age at time of sample collection, based on a cohort of 23,190 newborns delivered in Las Vegas and Reno, Nevada during a 15-month period between April 1, 1998 and June 30, 1999

<table>
<thead>
<tr>
<th>Main effects and their interaction</th>
<th>Difference in mean T₄ (ug/dl)</th>
<th>95% CI LCL</th>
<th>UCL</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>City (Las Vegas vs Reno)</td>
<td>-0.069</td>
<td>-0.232</td>
<td>0.094</td>
<td>0.407</td>
</tr>
<tr>
<td>Time Period (B vs A)*</td>
<td>0.595</td>
<td>0.484</td>
<td>0.706</td>
<td>0.000</td>
</tr>
<tr>
<td>Interaction (City &amp; Time Period)</td>
<td>0.021</td>
<td>-0.198</td>
<td>0.240</td>
<td>0.850</td>
</tr>
<tr>
<td>Control variables</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Gender (female vs male)</td>
<td>0.727</td>
<td>0.632</td>
<td>0.823</td>
<td>0.000</td>
</tr>
<tr>
<td>Birth Weight (per 1,000 gram)</td>
<td>0.850</td>
<td>0.735</td>
<td>0.964</td>
<td>0.000</td>
</tr>
<tr>
<td>Age at time of sample collection**</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>First Day vs Fourth Day</td>
<td>-1.275</td>
<td>-1.552</td>
<td>-0.999</td>
<td>0.000</td>
</tr>
<tr>
<td>Second Day vs Fourth Day</td>
<td>0.408</td>
<td>0.206</td>
<td>0.610</td>
<td>0.000</td>
</tr>
<tr>
<td>Third Days vs Fourth Day</td>
<td>0.758</td>
<td>0.538</td>
<td>0.978</td>
<td>0.000</td>
</tr>
</tbody>
</table>

* During the 15-month study period, time period A was defined as a combination of those months at which perchlorate levels were detected in the drinking water in Las Vegas, Nevada whereas time period B included the remaining months at which perchlorate level were not detected in the drinking water in Las Vegas, Nevada.

** The variable was treated as a categorical variable in the multivariate analysis.