Synergistic effect of hepatitis virus infection and occupational exposures to vinyl chloride monomer and ethylene dichloride on serum aminotransferase activity

H-I Hsieh, J-D Wang, P-C Chen, T-J Cheng

Aims: To study the synergistic effect of occupational chemical exposure and hepatitis virus infection on serum aminotransferase activity.

Methods: A total of 568 male workers who were employed in five polyvinyl chloride (PVC) or four vinyl chloride monomer (VCM) manufacturing factories were studied. Information relating to current job title, alcohol consumption, and cigarette smoking was obtained. Exposure level of chemical mixtures was classified by hygienic effect (a summation of personal time weighted average/reference permissible exposure level of each chemical) into high, moderate, and low exposure groups. Serum aspartate aminotransferase (AST) and alanine aminotransferase (ALT), hepatitis B surface antigen (HBsAg), hepatitis B e antigen (HBeAg), and anti-hepatitis C antibody were assayed.

Results: Hepatitis virus infection and increased body mass index were associated with abnormal serum aminotransferase activity. In workers with hepatitis virus infection, those with high exposure had a higher prevalence of abnormal AST and ALT compared to low exposure; among those without hepatitis virus infection, the differences of prevalence of abnormal AST and ALT were not significant between different chemical exposure groups. There was a significant trend of increasing risks of increased AST and ALT in moderate and high exposure groups with hepatitis virus infection. Such a synergistic effect was more prominent among HBeAg-positive workers.

Conclusions: Mixed exposures to 1,2-ethylene dichloride and VCM have a positive synergistic effect with hepatitis virus infection on liver damage. Assessment of fitness for work should be considered in workers with hepatitis B and C infection, when they have potential exposure to hepatotoxins in the workplace.

Measurements of serum aspartate aminotransferase (AST) and alanine aminotransferase (ALT) have been used extensively in the assessment of liver damage. Abnormal serum aminotransferases have been associated with occupational and non-occupational factors. Among non-occupational hepatotoxins, hepatitis B and C virus infection have synergistic effects with alcohol consumption on serum aminotransferases. However, it is not clear whether occupational chemical exposure and hepatitis B and C have synergistic effects on these hepatic enzymes or not. This question needs to be resolved for the assessment of fitness for work, particularly in populations with a high prevalence of hepatitis B infection, including Taiwan.

We have studied liver disorders in vinyl chloride monomer (VCM) and polyvinyl chloride (PVC) manufacturing workers over the past decade. VCM (CAS no. 75-01-4) or 1,2-ethylene dichloride (EDC; CAS no. 107-06-2) exposure has been associated with serum aminotransferase abnormalities in our previous studies. In the above studies, it seems that exposed workers with hepatitis B infection tend to have a higher risk of abnormal serum aminotransferase compared to those without hepatitis B infection. However, it was difficult to draw a conclusion because of the small number of subjects with abnormal serum aminotransferase. To increase the power of detection for the interaction between hepatitis virus infection and occupational chemical exposure, we included workers from both PVC and VCM manufacturing plants, who received medical examinations between 1995 and 1997. Since workers with hepatitis B e antigen (HBeAg) also had a higher prevalence of serum aminotransferase abnormality, the interaction between HBeAg and occupational chemical exposure on serum aminotransferase was further assessed.

METHODS

Study population
A total of 617 workers were eligible. Because 49 workers had incomplete data, 568 (92%) workers from five PVC (n = 292) and four VCM (n = 276) manufacturing factories were included for the analysis. Among them, 11 workers were office workers, who stayed indoor during most of their working hours, and three workers were guards who stayed away from the manufacturing site. Most of these workers have been presented in previous studies to investigate the relation between external chemical exposures and liver function or genotoxicity, or the effects of metabolic genotypes on liver function or genotoxicity.

After informed consent was obtained, all study subjects were surveyed by an interviewer administered questionnaire to obtain information on smoking, alcohol consumption, medicines, and medical and occupational histories. Alcohol consumption was calculated from drinking frequency and alcohol content of each beverage consumed. Those who drank at least once and had alcohol consumption with a

Abbreviations: anti-HCV, anti-hepatitis C antibody; ALT, alanine aminotransferase; AST, aspartate aminotransferase; BMI, body mass index; EDC, 1,2-ethylene dichloride; GSH, glutathione; HBeAg, hepatitis B e antigen; HBsAg, hepatitis B surface antigen; PEL, permissible exposure level; PVC, polyvinyl chloride; TWA, time weighted average; VCM, vinyl chloride monomer
Chemical-viral interaction on serum aminotransferases

Minimum of 80 g intake per week in the past one month were defined as having drinking habit. Smoking behaviour was defined as having smoked at least one cigarette per day within the preceding six months of data collection.

Exposure assessment

EDC is used in the production of VCM, and VCM is subsequently used for the polymerisation to manufacture PVC. In VCM manufacturing plants, workers were exposed to both EDC and VCM, while workers in PVC plants were exposed to VCM only. Detailed occupational history included job title, daily activity, and use of respirators in the current and previous jobs. Personal samplings were conducted to calculated EDC and VCM time weighted average (TWA) for each category of work.91 If personal sampling data were not available, data of area sampling was used. Office workers and guards were presumably exposed to extreme low concentrations of chemicals, thus 0 ppm of VCM and EDC were assumed as their TWA. To consider the combined effect of EDC and VCM, the hygienic effect was calculated by using the model of \( \frac{C_1}{T_1} + \frac{C_2}{T_2} \), where \( C \) was the measured TWA and \( T \) was the permissible exposure levels or equivalents for each chemical. One ppm was used for both EDC and VCM in this study, which has been adopted by many institutions. Our previous study also found that EDC and VCM cause abnormal liver aminotransferase around 1 ppm.7 Workers with hygienic effects below 1 were classified into the low exposure group. Workers with hygienic effects between 1 and 5 were classified into the moderate exposure group, and workers with exposures greater than 5 were classified into the high exposure group.

Biochemical tests and hepatitis virus markers

Markers of liver damage, including AST and ALT, were analysed with a Hitachi 7050 autoanalyzer (Hitachi Co., Tokyo, Japan) at National Taiwan University Hospital (NTUH). Hepatitis B virus surface antigen (HBsAg) and anti-hepatitis C virus antibody (anti-HCV) were determined by enzyme linked immunoassay (EIA, Abbott Laboratories, Chicago, IL, USA), respectively. HBeAg was also assayed by ELISA in workers with positive HBsAg. Abnormal results for serum aminotransferases were defined as having values greater than the reference provided by NTUH. In 1995, it was 31 for both ALT and AST. In 1996 and 1997, it was 37 for AST, and 41 for ALT. Subjects with positive hepatitis B infection were defined as having positive HBsAg, and subjects with positive hepatitis C infection were defined as having positive anti-HCV. Since the number of subjects exhibiting a positive titre for anti-HCV was small, HBsAg and anti-HCV were grouped together as hepatitis virus infection. Body mass index (BMI) was calculated as weight in kilograms divided by the square of height in metres.

Statistical analysis

The PC/SAS statistical package (SAS Institute Inc., Cary, NC, USA) was used for the statistical analysis. The \( \chi^2 \) test was used to compare the differences of age, employment duration, hepatitis virus infections, body mass index, and alcohol consumption between different exposure groups. Crude comparisons of abnormal AST and ALT by variables of interests were conducted in the univariate analysis. Subsequently, a multiple logistic regression model was used to determine the odds ratio (OR) of abnormal AST and ALT levels for different exposure groups (high, moderate, and low chemical exposure), hepatitis virus infection (yes and no), body mass index (\( >25.0 \) and \( \leq 25.0 \) kg/m\(^2\)), and habitual drinking (yes and no). OR of abnormal AST and ALT levels on different exposure groups stratified by hepatitis B and C virus infection was also calculated after controlling for potential confounders including age, BMI, and alcohol drinking. OR of abnormal AST and ALT on the chemical exposure (low, moderate, and high) was further calculated by HBeAg and HBsAg status (–/−, –/+ , and +/+ , respectively). All p values were quoted two sided, and those values <0.05 were regarded as statistically significant.

RESULTS

Descriptive statistics

Table 1 summarises the basic characteristics of study subjects stratified by different exposure groups. The median TWA of VCM was 0.67 (range 0.0–73.8) ppm and of EDC was 0.35 (range 0.0–30.5) ppm. Most workers (83.6%) were less than 50 years of age, 29.8% of workers had BMI greater than 25, 11.1% of workers consumed more than 80 g of alcohol per week, 17.3% of workers had HBsAg, and 3.5% of workers had HBeAg. The high exposure group had more habitual drinkers than the moderate and low exposure groups. The low exposure group was older than the moderate and high exposure groups. All other characteristics of cigarette smoking, BMI, HBeAg, HBsAg, and Anti-HCV were not statistically significant between these three exposure groups.

Overall, 112 workers (19.7%) showed increases of AST or ALT. There were 22.4%, 20.4%, and 18.6% of workers with abnormal AST or ALT among the high, moderate, and low exposure groups. Workers with high exposure had more cases of abnormal AST or ALT when compared with other workers, but this did not reach statistical significance.

Multiple logistic regression analysis

Multiple logistic regression analysis (see table 2) revealed that AST was associated with BMI and hepatitis virus infection. There were similar findings with ALT, but the association with hepatitis C infection and ALT did not reach statistical significance. Increased chemical exposure was also associated with abnormal ALT or AST, but this association did not reach statistical significance.
Interaction analysis of factors on serum aminotransferase

The interactions between chemical exposure and each potential factor (hepatitis virus infection, BMI, and alcohol drinking) on serum aminotransferase were calculated. Significant interactions were observed for chemical exposure and hepatitis virus infection (table 3). When workers did not have hepatitis virus infection, all serum aminotransferases showed no difference among different exposure groups. Dose dependent effects of chemicals on AST and ALT were found when workers had hepatitis virus infection. Workers who had high chemical exposure and hepatitis virus infection had the highest risk of abnormal AST (OR 10.6; 95% CI 3.6 to 32.9) and ALT (OR 6.4; 95% CI 2.1 to 19.1) when compared with workers who had low chemical exposure and lacked hepatitis virus infection. If we confined the analysis to those with workers who had positive HBsAg and HBeAg. Among workers who had positive HBsAg but did not have positive HBeAg, a higher risk of abnormal AST (OR 9.1; 95% CI 2.4 to 34.1) and ALT (OR 2.9; 95% CI 0.9 to 10.1) compared to those without hepatitis virus infection and habitual drinking. The positive interaction effect of hepatitis virus infection and alcohol consumption on AST reached statistical significance.

**Table 1** Frequency distribution of basic characteristics in 568 male workers stratified by exposure categories

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>High (≥5)</th>
<th>Moderate (≥1 and &lt;5)</th>
<th>Low (&lt;1)</th>
<th>Total (n = 568)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Manufacturing factory</td>
<td>67.1</td>
<td>64.4</td>
<td>39.2</td>
<td>51.4***</td>
</tr>
<tr>
<td>PVC</td>
<td>32.9</td>
<td>25.6</td>
<td>60.8</td>
<td>48.6***</td>
</tr>
<tr>
<td>VCM</td>
<td>46.1</td>
<td>46.1</td>
<td>59.1</td>
<td>53.0</td>
</tr>
<tr>
<td>Age ≥40 years</td>
<td>43.4</td>
<td>45.0</td>
<td>55.5</td>
<td>50.9*</td>
</tr>
<tr>
<td>Duration of employment ≥15 years</td>
<td>29.0</td>
<td>33.0</td>
<td>27.9</td>
<td>29.8</td>
</tr>
<tr>
<td>Body mass index ≥25.0 kg/m²</td>
<td>48.7</td>
<td>43.5</td>
<td>36.9</td>
<td>40.7</td>
</tr>
<tr>
<td>Current cigarette smoking (yes)</td>
<td>23.7</td>
<td>8.4</td>
<td>9.6</td>
<td>11.1**</td>
</tr>
<tr>
<td>Alcohol drinking (yes)</td>
<td>19.7</td>
<td>15.7</td>
<td>17.6</td>
<td>17.3</td>
</tr>
<tr>
<td>Positive hepatitis B surface antigen (HBsAg)</td>
<td>7.9</td>
<td>1.6</td>
<td>3.7</td>
<td>3.5</td>
</tr>
<tr>
<td>Positive hepatitis B e antigen (HBeAg)</td>
<td>1.3</td>
<td>4.7</td>
<td>3.7</td>
<td>3.7</td>
</tr>
</tbody>
</table>

*p<0.05; **p<0.01 by χ² test.

**Table 2** Odds ratios (OR) with 95% confidence intervals (CI) of multiple logistic regression modelling adjusted for major determinants, including body mass index (BMI), chemical exposure, hepatitis B surface antigen (HBsAg), anti-hepatitis C antibody (Anti-HCV), drinking, and smoking

<table>
<thead>
<tr>
<th>Determinants</th>
<th>Definition</th>
<th>AST (OR 95% CI)</th>
<th>ALT (OR 95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>≥40.0 v &lt;40.0</td>
<td>0.8 [0.5 to 1.3]</td>
<td>0.6 [0.4 to 1.0]</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>≥25.0 v &lt;25.0</td>
<td>2.2 [1.2 to 3.9]*</td>
<td>3.5 [2.2 to 5.9]*</td>
</tr>
<tr>
<td>Chemical exposure</td>
<td>High v low</td>
<td>1.3 [0.6 to 2.7]</td>
<td>1.4 [0.7 to 2.6]</td>
</tr>
<tr>
<td></td>
<td>Moderate v low</td>
<td>0.8 [0.4 to 1.5]</td>
<td>1.0 [0.6 to 1.6]</td>
</tr>
<tr>
<td>HBsAg</td>
<td>Positive v negative</td>
<td>3.5 [1.9 to 6.4]*</td>
<td>2.5 [1.5 to 4.2]*</td>
</tr>
<tr>
<td>Anti-HCV</td>
<td>Positive v negative</td>
<td>5.9 [2.2 to 15.9]*</td>
<td>2.3 [0.8 to 6.3]</td>
</tr>
<tr>
<td>Drinking</td>
<td>Yes v no</td>
<td>1.3 [0.6 to 3.0]</td>
<td>0.9 [0.5 to 1.9]</td>
</tr>
<tr>
<td>Smoking</td>
<td>Yes v no</td>
<td>1.1 [0.6 to 2.1]</td>
<td>1.0 [0.6 to 1.6]</td>
</tr>
</tbody>
</table>

*p<0.01 by χ² test.
Hepatitis B and/or C infection have been associated with increased serum aminotransferase activities. Our results reveal similar findings, although the association between anti-HCV and ALT was not statistically significant. This is probably due to small numbers of workers with anti-HCV.

Increased BMI is also associated with increased ALT and AST in our study (table 2). Increased BMI is a common aetiology of abnormal liver function tests for healthy workers. Our results corroborate such an association. Alcohol consumption has been reported to be associated with abnormal liver function, but our results did not show this relation. This is most likely due to relative small amount of alcohol consumption by these workers. Lack of association between alcohol consumption with abnormal ALT and AST was also observed in several studies conducted in Taiwanese workers. A study conducted in Italy showed that those who consumed 80 g each day had a greater risk of developing abnormal liver function among chronic symptomless HBV carriers. Positive synergism was also observed between HBV and HCV infection and alcohol consumption. Our study also revealed that HBV and HCV infection exacerbated the effect of alcohol on AST, although the effects of alcohol on ALT was less prominent. Again, this could result from the small amount of alcohol consumption in our study subjects. Additionally, AST increase is usually more prominent than ALT in alcoholic hepatitis. Thus, the relation between abnormal serum aminotransferases and non-occupational factors in our study is consistent with previous studies.

A recent study also showed that the relation between occupational dimethylformamide exposure and abnormal liver function was enhanced in those with HBV infection. Here, we showed that VCM and EDC together could also have a more than additive interaction with HBV and HCV infection. Further analysis in our study indicated that workers with HBeAg were more likely to have abnormal ALT and AST compared to those with HBsAg alone, when they were exposed to occupational chemicals. As both VCM and EDC were reported to be hazardous to the liver, detection

### Table 3

<table>
<thead>
<tr>
<th>Category</th>
<th>AST (n = 58)</th>
<th>ALT (n = 103)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>%</td>
<td>OR (95% CI)†</td>
</tr>
<tr>
<td>HBsAg (+) or anti-HCV (+)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>High exposure</td>
<td>56.3</td>
<td>10.6 (3.6 to 31.5)*</td>
</tr>
<tr>
<td>Moderate exposure</td>
<td>23.5</td>
<td>3.0 (1.1 to 8.0)</td>
</tr>
<tr>
<td>Low exposure</td>
<td>15.9</td>
<td>2.3 (0.9 to 5.4)</td>
</tr>
<tr>
<td>HBsAg (−) and anti-HCV (−)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>High exposure</td>
<td>1.7</td>
<td>0.5 (0.1 to 1.5)</td>
</tr>
<tr>
<td>Moderate exposure</td>
<td>5.7</td>
<td>0.6 (0.3 to 1.6)</td>
</tr>
<tr>
<td>Low exposure</td>
<td>8.8</td>
<td>1.0 (referent)</td>
</tr>
</tbody>
</table>

*P < 0.01 by Mantel-Haenszel χ² test for trend analysis.
†The referent group (OR = 1) refers to workers with low chemical exposure and negative hepatitis virus infection.
‡The referent group (OR = 1) refers to workers with low chemical exposure and positive hepatitis virus infection.

Figure 1

Odds ratio (OR) with 95% confidence intervals (CI) of abnormal AST (top) and ALT (bottom) among workers with different categories of exposure and hepatitis B infection; adjusted for age, drinking, anti-HCV, and BMI. The reference group (OR = 1.0) refers to workers with low chemical exposure and negative hepatitis B infection.
of such an effect is not surprising, and the mechanism should be clarified.

Previous human and animal studies indicated that glutathione (GSH) depletion can be caused by hepatitis virus infection.14–18 Glutathione S-transferases (GST) and glutathione play an important role in the metabolism of EDC and VCM, of which the electrophilic intermediate metabolites are conjugated with GSH to be detoxified.19–20 Thus, GSH depletion caused by hepatitis virus infection may lead to an accumulation of active intermediate metabolites of EDC and VCM, then exacerbate EDC and VCM induced hepatotoxicity. Previous studies conducted with 1,1-dichloroethylene which showed correlations between hepatocellular damage and magnitudes of both covalent binding and GSH depletion also supported this proposed hypothesis.21 Our recent study also suggests that the GSTT1 genotype may play an important role in liver aminotransferase abnormality caused by vinyl chloride.4 In patients with positive HBeAg, there is more active HBV replication and inflammation, which can reduce the level of GSH.22 Therefore, they are at a high risk of active HBV replication and inflammation, which can reduce the level of GSH. Therefore, they are at a high risk of showing increases in AST and ALT, as shown in fig 1. We conclude that subjects with hepatitis B and/or C virus infections are more likely to be damaged by hepatotoxic agents; such a potential synergistic effect may be caused by GSH depletion after hepatitis virus infection.

Studies conducted in humans and rats also found that ethanol significantly decreased glutathione concentrations,23–25 of which the synergistic hepatotoxic effect between alcohol and hepatitis virus infection could also have resulted from potential overloading of the oxidative damage through the generation of reactive intermediates and decreased radical scavenging.

We are concerned that VCM or EDC workers with HBV and/or HCV infections may not be well protected under current occupational standards. We suggest that workers with HBeAg should not be exposed to hepatotoxins in their work. We also advise workers with anti-HCV and abnormal serum aminotransferases, not to be exposed to hepatotoxins. Serums of workers with positive HBsAg but negative HBeAg, need to be closely monitored if exposed to higher levels of VCM or EDC. Furthermore, a more stringent occupational standard is needed to protect workers exposed to hepatotoxins in countries where hepatitis B and C virus infection is prevalent.

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