Neuropsychological findings in a non-clinical sample of workers exposed to solvents

I Reinvang, H M Borchgrevink, O Aaserud, V Lie, U F Malt, P Nakstad, P G Larsson, L Gjerstad

Abstract
The risk of long-term damage to the CNS after exposure to mixed solvents in work environments is controversial. Thirty-six workers were studied who had been exposed to organic solvents for more than 10 years (mean 24.5 years) in a working environment. The workers and unexposed controls were studied with a battery of neuropsychological and cognitive tests. Significant group differences were observed for the Wechsler adult intelligence scale (WAIS) digit span and symbol digit substitution, and on paired associate learning and continuous word recognition. The results suggest that long-term work-related exposure to organic solvents may have chronic toxic effects.

Subjects and methods
METHODS
Behavioural tests were chosen to be representative of earlier studies and recommendations from international review groups. The so-called WHO battery is recommended by an international review group to provide a common reference for cooperative studies.7 Tests belonging to this battery are marked (WHO) in the following.

Subtests from the Wechsler adult intelligence scale (WAIS) included similarities, block design, digit span (WHO), and digit symbol substitution (WHO). Memory tests included the paired associate learning test and symbol digit recall test used in previous neurotoxicological studies.8 Standard clinical neuropsychological tests9 included a non-verbal clinical memory test, Kimura recurring figures, with a recognition response. The trail-making test, parts A and B, was included as a measure of cognitive speed, and the grooved pegboard test as a measure of dexterity and coordination.

A continuous word recognition task10 was adapted for use in this study. A series of 90 words is shown on a computer screen with three-second intervals. Thirty words are shown only once and 30 words are shown twice. The number of words intervening between recurring words may be two, four, or six (short interval), or 12, 24, or 36 (long interval). The reaction time to yes/no button-press and accuracy is measured.

Reaction time (WHO) is a sensitive measure of acute neurobehavioural effects and is measured in this study with the simple reaction time and continuous performance tests from the NES 2 computerised battery.11 Tests of manual dexterity and coordination (WHO) include a computerised test, hand-eye coordination from NES 2. The WHO battery includes a self-rating of mood and this study uses the version of profile of mood states from the NES 2.

SUBJECTS
Thirty-six male workers exposed to organic solvents for more than 10 years in a working environment.
environment were studied. They were recruited from military sites on the basis of available exposure records of work tasks including construction painting, degreasing, or handling of jet fuel. All of the workers had been exposed to mixtures of solvents, mainly white spirit and various paints, and the same worker may have been exposed to different combinations of solvents at different times.

The workers were all fully employed at the time of the study, some of them still working with solvents, but with low exposure levels in recent years. None was seeking compensation for work-related injury at the time of entering the study. The mean (SD) age was 44·5 (8·0) years, and the mean (SD) length of exposure was 24·2 (7·5) years. Most (80%) had an educational level of less than 12 years of general education with additional work-related training.

Controls were selected as pairs matched for sex, age, and level of education from unexposed workers at the same work sites by the local medical officer. If a certain site had five workers satisfying the exposure inclusion criteria for the study, then the local medical officer also recruited five controls from the same site on a best-match basis. The resulting control group (mean (SD) age 44·1 (8·1) years) was well matched to the exposed group on demographic variables. The study group and controls were screened for medical history and alcohol consumption. Subjects were excluded if they had a history of head trauma with unconsciousness on admission to hospital, had other known brain disease, diabetes, or a history of drug or alcohol abuse.

Neuropsychological and other studies were performed blindly, without knowledge of the exposure status of the subjects, and the group membership code was not broken until data collection for the whole study was complete.

### Results

The test variables showed distributions of varying degrees of normality and to use a consistent statistic, the Wilcoxon non-parametric two-sample test for non-independent samples was reported. Test results are consistently reported as raw scores, and all standardised tests were scored according to published standard procedures for scoring.

The groups were comparable on general measures of verbal and visuospatial function from the WAIS, but significant differences were observed for digit span and symbol digit substitution (table 1).

On neuropathological and performance tests significant differences were found on number of trials before reaching criterion on paired associate learning (table 2), and for number of items correctly identified on continuous word recognition. Significant differences on computerised measures of reaction time or visuo-motor coordination were not found, and indications of mood differences were not found on the total score nor on any of the subscales.

A further analysis of subscales in tests showing significant findings is of interest as a basis for interpreting the results. Thus paired associate learning was analysed on the first recall attempt as well as on total trials to criterion and delayed recall. All measures showed significant group differences, but only for difficult items on the delayed recall measure. Continuous word recognition was categorised into performance on words with six or less intervening items \( v \) words with 12 or more intervening items (see table 2).

### Discussion

Group differences were found in several tests of memory and speed, whereas cognitively complex tasks measuring verbal and non-verbal abilities did not show significant differences. All the observed differences are in the direction of indicating poorer test results in the group of workers exposed to solvents, and may thus indicate a harmful effect of work-related solvent exposure. The results thus confirm previous findings while avoiding the bias towards self-referred subjects in several reported studies. This study may also contain a selection bias: the healthy worker effect. If workers who have experienced symptoms have left their jobs and thus not been included in the study, this bias would decrease the likelihood of finding significant deficits in active workers. The fact that our study design is conservative tends to strengthen the findings.

The issue of controlling for pre-morbid ability differences in neuropsychological research has been controversial. Tests of vocabulary and verbal conceptual reasoning may show deficits in some studies of workers exposed to solvents, and the differences may reflect selection factors to certain professions. In this study a test of verbal conceptual reasoning (definition of similarities) was chosen as a measure of verbal ability. This test is

### Table 1

<table>
<thead>
<tr>
<th>Test</th>
<th>Exposed (n = 36)</th>
<th>Unexposed (n = 36)</th>
<th>Z value</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Digit span—forward</td>
<td>5 (1)</td>
<td>6 (2)</td>
<td>2·27</td>
<td>0·023</td>
</tr>
<tr>
<td>Digit span—backward</td>
<td>4 (2)</td>
<td>4·5 (1)</td>
<td>2·30</td>
<td>0·021</td>
</tr>
<tr>
<td>Digit symbol</td>
<td>38·5 (13·5)</td>
<td>43·5 (15)</td>
<td>2·90</td>
<td>0·004</td>
</tr>
<tr>
<td>Similarities</td>
<td>17·5 (6)</td>
<td>18·6 (6)</td>
<td>0·96</td>
<td>0·33</td>
</tr>
<tr>
<td>Block design</td>
<td>37·5 (11)</td>
<td>37·5 (14)</td>
<td>0·60</td>
<td>0·55</td>
</tr>
</tbody>
</table>

### Table 2

<table>
<thead>
<tr>
<th>Test</th>
<th>Exposed (n = 36)</th>
<th>Unexposed (n = 36)</th>
<th>Z value</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Peg-board dominant</td>
<td>68 (14)</td>
<td>65·5 (14·5)</td>
<td>0·95</td>
<td>0·34</td>
</tr>
<tr>
<td>Peg-board non-dominant</td>
<td>70 (14·5)</td>
<td>69·5 (17·5)</td>
<td>0·39</td>
<td>0·69</td>
</tr>
<tr>
<td>Trail-making A</td>
<td>39 (21)</td>
<td>35·5 (17·5)</td>
<td>1·48</td>
<td>0·13</td>
</tr>
<tr>
<td>Trail-making B</td>
<td>108 (45·5)</td>
<td>98·5 (57·5)</td>
<td>0·96</td>
<td>0·33</td>
</tr>
<tr>
<td>Digit symbol reproduction</td>
<td>6 (3·5)</td>
<td>7 (3)</td>
<td>1·34</td>
<td>0·18</td>
</tr>
<tr>
<td>Kimura recognition</td>
<td>31·5 (12)</td>
<td>33·5 (11)</td>
<td>0·60</td>
<td>0·55</td>
</tr>
<tr>
<td>Paired associate learning</td>
<td>29·5 (21·5)</td>
<td>19·5 (20)</td>
<td>2·81</td>
<td>0·005</td>
</tr>
</tbody>
</table>

*RT = Reaction time; POMS = profile of mood states.
recommended in neuropsychology because the results are less dependent on schooling than tests of factual knowledge. As the groups in this study did not differ significantly on this test, it can be argued that the group selection criteria have functioned appropriately. There was a non-significant trend towards lower scores in the exposed group on Similarities, but repeated statistical analyses with score on Similarities as covariate resulted in only minor adjustments of the significance levels, and all major findings, including differences on paired associate learning, held up. The study by Morrow et al also reports a deficit in Paired Associate learning and Digit Span.

Performance measures that have been found to be sensitive in epidemiological research were included in addition to more traditional neuropsychological tests. We found no evidence that computerised tasks based on subtle reaction time measures, as in the NES 2 battery, have a higher sensitivity than neuropsychological tests.

Earlier reports indicate that subjective reports of neurasthenic symptoms are sensitive to mild toxic effects. In this study no differences in reported symptoms were found on mood scales corresponding closely to WHO standards. It could be argued that the significant findings on cognitive tests in our study are caused by acute effects. It is therefore worth noting that we did not observe group differences on other measures assumed to be more sensitive to acute effects.

Several memory tests in our study show significant deficits. The continuous recognition memory paradigm was included in the study to facilitate a theoretically motivated analysis of memory mechanisms. The results on this paradigm as well as the other memory tests may indicate that short- and long-term storage mechanisms are affected in the study group.

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