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by Stalberg E, Hilton-Brown P, Kolmodin-Hedman B, Holmstedt B, Augustinsson K-B

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Effect of occupational exposure to organophosphorus insecticides on neuromuscular function

by ERIK STÅLBERG, PER HILTON-BROWN, BIRGITTA KOLMODIN-HEDMAN, BO HOLMSTEDT and KLAS-BERTIL AUGUSTINSSON

STÅLBERG, E., HILTON-BROWN, P., KOLMODIN-HEDMAN, B., HOLMSTEDT, B. and AUGUSTINSSON, K.-B. Effect of occupational exposure to organophosphorus insecticides on neuromuscular function. Scand. j. work environ. & health 4 (1978) 255—261. Neurophysiological investigations and determinations of cholinesterase activity on plasma and erythrocytes were carried out on 11 Swedish spraymen exposed to bromophos, diazinon, dursbane, and malathion. Plasma cholinesterase activity was significantly reduced after work, while erythrocyte cholinesterase activity was unchanged. In none of the workers with a decreased plasma cholinesterase activity after work could any related acute neuromuscular disturbance be detected when the men were tested with repetitive nerve stimulation and with single fiber electromyography. Signs of subclinical neuropathy were present as a slight reduction in sensory conduction velocity and increased fiber density in some workers.

Key words: cholinesterase activity, electromyography, neuromuscular transmission, neuropathy, organophosphorus compounds, pesticides, single fiber electromyography.

Electromyography (EMG) has been used on an increasing scale in occupational medicine over the past years (14, 16). One example is the study of effects on nerve and muscle in workers handling different kinds of pesticides such as organophosphorus compounds (11, 15). The cholinesterase inhibition of these compounds might possibly affect neuromuscular transmission in exposed workers. Roberts et al. (11, 15) suggested that a relatively simple EMG technique was very useful and, to some extent, superior to cholinesterase measurements when the effects of exposure to organophosphorus compounds are being monitored (6, 10, 11, 15).

In the present investigation neuromuscular transmission and cholinesterase activity have been studied with more sensitive methods than were used in earlier investigations. It is an attempt to determine whether a group of Swedish spraymen was exposed to organophosphorus pesticides to such a degree that EMG tests or cholinesterase determinations showed abnormal values and, if so, to correlate the...
methods used. In addition to measuring the decremental response of the surface EMG recording and nerve conduction velocity, we performed single fiber EMG (SFEMG) (17). This method is particularly sensitive to slight neuromuscular disturbances and changes in the terminal innervation pattern. Both plasma and erythrocyte cholinesterase activity was measured.

SUBJECTS AND EXPOSURE

Eleven men from a company handling the commercial application of insecticides were examined. They were all spraymen, working daily with solutions of 4 \% lindane, 0.2 \% pyrethrum, 0.125 \% piperonylbutoxide, 2.5 \% malathion or 2 \% bromophos in kerosene. Intermittently they were exposed to 0.5 diazinon and dursbaine. The employment period varied between 1 and 24 years. They were equipped with routine protective devices such as respirators, boots, overalls and gloves. The men were examined clinically, neurophysiologically, and with measurements of blood cholinesterase activity. They also formed their own reference group since measurements were also made after 1—4 weeks of non-exposure (preexposure values). The measurements following exposure were generally made within 1—24 h after a period of spraywork. Exposure values were taken during the spring season, and the preexposure values after the summer vacation. Blood samples for cholinesterase activity analysis were taken immediately before each EMG investigation.

DETERMINATION OF
CHOLINESTERASE ACTIVITIES

The cholinesterase activity of plasma (BuChE) and erythrocytes (AChE) was determined with a gasometric technique using whole blood samples (50 \( \mu l \)) applied and dried on filter paper) and butyrylcholine iodide and acetyl-\( \beta \)-methylcholine iodide as the selective substrates for the two activities (3, 4). The activity was expressed in \( b_{30} \) values, i.e., microliters of carbon dioxide evolved in 30 min/50 \( \mu l \) blood. These values can easily be converted to nanomoles of substrate hydrolyzed per second per liter of blood (which is the new unit, called katal, recommended by the International Enzyme Commission) with the following formula:

\[
1 \text{ nkat} = 0.5 \times b_{30}.
\]

NEUROPHYSIOLOGICAL
INVESTIGATIONS

Nerve conduction studies

Motor nerve conduction velocity measurements were made with the standard technique with surface recording (metal discs) and stimulating electrodes (DISA, 13 K 62). The filter setting of the amplifier was 2 Hz to 20 kHz. The ulnar nerve was stimulated on both sides. The active recording electrode was placed over the belly of the abductor digitii minimi (ADM) muscle and the indifferent electrode over the fifth metacarpophalangeal joint.

Sensory nerve conduction velocity measurements were made from the right sural nerve with antidromic stimulation between the two heads of the gastrocnemius muscle and the recording of the sural nerve action potential at the lateral malleolus with fill pad electrodes (DISA 13 K 62).

Furthermore the amplitude of the compound nerve action potential was measured with fill pad electrodes on both sides at the sulcus ulnaris when the ulnar nerve was stimulated at the level of the wrist.

Before all nerve conduction studies we checked that the extremity was warm. The temperature of the laboratory was
kept constant at 22°C in all the investigations.

Muscle response

The change in the compound muscle action potential was measured from the *abductor digiti minimi* muscle with repetitive ulnar nerve stimulation. For the recording and stimulating the setting was the same as that used for the measurements of the motor conduction velocity of the ulnar nerve. The amplitude (negative peak) of the first response and the change in amplitude and integrated surface (the whole response) between the first and fourth response at repetitive 2-Hz nerve stimulation (decrement) was measured on-line with a computer (PDP 11/40). The accuracy of the amplitude measurements was 0.1 mV. On different occasions the amplitude may vary up to 2 mV (range) in a normal subject. The decrement measurements were made with an accuracy of about 1 %, and a value exceeding 5 % (amplitude and surface) was considered abnormal. These parameters were measured at rest and immediately after 20 s of maximal voluntary activation of the muscle (postactivity facilitation). A final test was made 5 min after the activation period (postactivity exhaustion). Care was taken that the hand muscle was warm during the investigation.

Single fiber electromyography

SFEMG was performed in the *extensor digitorum communis* (EDC) muscle and measurements were made of the jitter and fiber density.

**Jitter.** Jitter recordings were made from the slightly voluntarily activated muscle as previously described (7). The electrode was positioned to record simultaneously activity from two or more muscle fibers belonging to the same motor unit. There is a variability in the time interval between two action potentials from muscle fibers in the same motor unit at consecutive discharges, called the jitter, which is on the order of 5—55 \( \mu s \) in a normal EDC muscle. This jitter is mainly due to a variability in the neuromuscular transmission time (17). The jitter is increased in cases with disturbed neuromuscular transmission even before impulse blocking occurs, i.e., before decrement studies or clinical tests show any abnormalities. The jitter analysis was made on a computer (PDP 11/40) off-line. Exceptionally manual analysis was made from measurements of recordings on film or a storage oscilloscope. Recordings were made from 10—20 potential pairs (corresponding to 20—40 motor end-plates) at each investigation.

**Fiber density.** Fiber density is the average number of muscle fibers from one motor unit within the SFEMG electrode uptake area (radius about 270 \( \mu m \)), obtained from 20 recording sites. The method and normal results for different ages have been presented earlier (18).

In case of reinnervation (posttraumatic, polyneuropathy, etc.) the organization of muscle fibers in the motor unit is changed, due to collateral sprouting, and causes an increased fiber density.

**Statistical methods.** Student's t-test of paired differences with the one-tailed test was used for the statistical analysis.

RESULTS

Clinical investigation

None of the examined subjects reported clinical symptoms of intoxication indicating a reaction to organophosphate exposure. Apart from occasional headache, sometimes related to work, the subjects had no symptoms or signs of any neurological disturbance.

Cholinesterase activities

The cholinesterase activities of the 11 spraymen before and the mean difference between the values after and before exposure are presented in table 1. The values
### Table 1. Cholinesterase values and neurophysiological data.

| Variable                              | Preexposure value | Difference between pre- and postexposure values | p values |
|---------------------------------------|-------------------|-----------------------------------------------|----------|
|                                       | Mean (SD)         | Mean (SD)                                     |          |
| Erythrocyte cholinesterase (AChE) (b30) | 33.3 (1.8)        | —2.04 (8.8)                                   | p > 0.2  |
| Plasma cholinesterase (BuChE) (b30)   | 77.6 (8.7)        | —6.14 (6.3)                                   | p < 0.01 |
| Motor conduction velocity (m/s)       | 58.6 (5.3)        | 1.6 (4.0)                                     | p > 0.1  |
| Sensory conduction velocity (m/s)     | 42.4 (3.6)        | —1.3 (2.4)                                    | p = 0.05 |
| Compound nerve action potential (mean values for right and left arm) (μV) | 81.6 (18.2)      | —3.6 (15.1)                                   | p > 0.2  |
| Muscle response (mV)                  | 10.6 (1.8)        | 1.1 (1.6)                                     | p > 0.2  |
| Fiber density                         | 1.6 (0.2)         | 0.04 (0.2)                                    | p > 0.2  |
| Mean jitter (μs)                       | 29.7 (7.3)        | 1.08 (5.4)                                    | p > 0.2  |

were all within the normal range reported for the technique used (3). The results revealed that plasma cholinesterase activity was decreased 8 % on the average (range 0–25 %) after exposure (p < 0.01). Only one subject had a decrease in plasma activity exceeding the normal intrindividual variation. In this connection it may be pertinent to report that 10 of the 11 workers had a lower plasma cholinesterase activity after exposure than before. The erythrocyte cholinesterase was not affected (p > 0.2).

### Nerve conduction studies

In the nerve conduction studies (table 1) the motor nerve conduction velocity was above the lower normal limits (> 45.0 m/s) in all workers, and no significant decrease could be detected between results on the two occasions (p > 0.1). The compound nerve action potential, normally showing a great scatter because of technical difficulties, did not show any significant reduction in amplitude (p > 0.2). The sensory conduction velocity was slightly reduced (3 %) after the exposure period (p = 0.05); reduction was noted in 7 of the 11 workers.

### Muscle response

No decremental response (table 1) and no postactivity facilitation or exhaustion was observed in any of the workers. The decrement was less than 3 %, No signs of repetitive discharges were found. The recorded compound muscle action potential from the ADM muscle ranged from 7.6 to 14 mV and showed no significant difference before and after the exposure (p > 0.2).

### Jitter

Some individual motor end-plates showed an increased jitter before or after exposure (table 1). No consistent difference in the jitter with exposure was found.

### Fiber density

For the group, no difference was seen between pre- and postexposure mean values of the fiber density (table 1). It was unchanged and within normal limits, 1.44 ± 0.15 (18) in seven of the workers (fig. 1). In four cases the fiber density exceeded normal values by 2 SD before exposure. Two had increased fiber density on both the test occasions.
There was no correlation between cholinesterase activities in plasma or red cells and the neurophysiological parameters (conduction velocities, mean jitter value, number of recordings with increased jitter, fiber density). Neither was there any correlation between changes in cholinesterase activity and changes in the neurophysiological parameters.

DISCUSSION

This investigation was performed to determine any disturbance in neuromuscular function and possible correlation with a decrease in cholinesterase activity in connection with work with organophosphorus compounds. Earlier reports (11) have indicated that neurophysiological methods could reveal such disturbances even when blood cholinesterase activity was normal. The EMG surface response was said to show a decrement at repetitive nerve stimulation, a decrease after muscle activity, and a lower amplitude after exposure to organophosphorus compounds (11, 15). The two first findings, and possibly also the third, would indicate disturbed neuromuscular transmission. If this is the case, the decrease in amplitude of the recorded action potential should be proportional to the number of blocked motor end-plates.

Our neurophysiological investigations with surface recordings showed normal response amplitudes without decrement or postactivity exhaustion. However, these studies can be technically difficult, and there are many possibilities of artifacts giving false positive or possibly negative results (e.g., improper electrode placement, movement of the electrodes during stimulation, change in skin resistance, intramuscular temperature, characteristics of the amplifiers, e.g., filter settings). It is particularly difficult to use the absolute amplitude for comparison between recordings made on different occasions. For the decrement studies it is very important, in situations of minor changes, that changes in both amplitude of the response and its integrated surface be compared. Whether these factors account for some of the abnormal findings reported earlier (10, 11, 15) cannot be determined since some of the information needed for such an evaluation is missing. In the present study these factors were standardized and kept as constant as possible between the investigations. By means of a computer the measurements were made with a high accuracy.

To determine whether any neuromuscular disturbances were present despite our normal surface recordings, we performed SFEMG, which is a sensitive method for the study of the transmission in single motor end-plates in situ and may show abnormalities in muscles having non-decreasing surface responses. In none of the cases did the mean jitter change after a period of work. The investigation thus

![Fig. 1. Fiber density in the extensor digitorum muscle. Open symbols represent normal material (18). Mean value and standard deviation for each decade indicated. Filled symbols represent the spraymen, each represented twice, before (*) and after (■) work.](image-url)
did not reveal any acutely disturbed neuromuscular transmission. If the surface response shows a low amplitude or decrement due to neuromuscular disturbances, there is always an increased jitter and blockings in the SFEMG recordings.

The slight changes in plasma cholinesterase activity were not correlated to disturbed neuromuscular transmission in the subjects.

In our study we found lower sensory nerve conduction velocities after work in seven subjects. The average reduction for the group was very small (3 %) but statistically significant (p = 0.05). Reduced conduction velocity is usually a sign of neuropathy, which thus may be present in some of our workers. The increase in fiber density in four subjects is interpreted as a sign of peripheral reinnervation after denervation. Such a lesion would most likely be peripherally localized since the motor nerve conduction velocities and compound nerve action potential amplitudes were normal.

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