Effect of PON1 on dichlorvos toxicokinetics

Na-Na Wang, Li Yuan, Heng Dai, Zhen-Kun Han, Min Zhao

ABSTRACT
Objectives To provide toxicokinetic and clinical evidence of the hydrolytic effect of paraoxonase-1 (PON1) on acute organophosphate poisoning in rats.

Methods 40 male Wistar rats were randomised into four equal groups. Dichlorvos administration group (A group) underwent dichlorvos injection (dissolved in corn oil) using intraperitoneal (ip) dose of 10 mg/kg. PON1 pretreatment group (B group) was injected with PON1 in the tail vein (intravenous), dose 9600 U/kg, 30 min prior to dichlorvos administration. In the treatment group (C group), atropine 0.05 mg/kg and pyraloxime chloride (PAM-CI) 120 mg/kg were injected intravenously within 2 min after dichlorvos administration. Finally, in the co-treatment group (D group), PON1 was injected intravenously with a dose of 9000 U/kg 30 min prior to dichlorvos administration; atropine 0.05 mg/kg and PAM-CI 120 mg/kg were injected intravenously within 2 min after dichlorvos administration. Blood was collected after administration. Plasma dichlorvos concentration was detected by liquid chromatography-mass spectra (LC-MS) method and clinical signs were observed. Toxicokinetic parameters were calculated in a statistical moment model.

Results AUC (0→∞) in group B was statistically different from that in groups A and C (p<0.05), while it was not different from group D (p>0.05); there was no statistical difference between group A and group C (p>0.05). The statistical results of Cmax were the same as those of AUC (0→∞). There were no differences of MRT between four groups (p>0.05). Clinical signs can be improved by PON1 and atropine + PAM-CI, and co-treatment can relieve signs more effectively.

Conclusion PON1 can decrease the amount of dichlorvos that entered the blood, lowered the peak concentration and relieved clinical signs.

Dose schedule and sample collection
40 male Wistar rats were randomised into four equal groups. The dichlorvos administration group (A group) underwent dichlorvos injection (dissolved in corn oil) using intraperitoneal (ip) dose of 10 mg/kg. The PON1 pretreatment group (B group) was injected with PON1 in the tail vein (intravenous), dose 9600 U/kg, 30 min prior to dichlorvos administration (10 mg/kg ip). In the treatment group (C group), atropine 0.05 mg/kg and pyraloxime chloride (PAM-CI) 120 mg/kg were injected intravenously within 2 min after dichlorvos administration (10 mg/kg ip). In the co-treatment group (D group), PON1 was injected intravenously in a dose of 9000 U/kg 30 min prior to dichlorvos administration (10 mg/kg ip); atropine 0.05 mg/kg and PAM-CI 120 mg/kg were injected intravenously within 2 min after dichlorvos administration.

Blood of 0.1 ml was collected at 5 min, 10 min, 20 min, 30 min, 1 h, 2 h, 4 h and 6 h after administration from the eye veins. Blood was centrifuged at 5000 r for 3 min and the supernatant was collected and stored in −80°C for dichlorvos detection.

Clinical signs after dichlorvos poisoning were observed. According to Gaidukov,6 all clinical signs noted following dichlorvos intoxication were categorised to mild, moderate or severe reactions. Mild reactions were characterised by ataxia and/or dyspnoea, while animals with severe reactions exhibited ventral position and/or tremors as well. The overall reactions observed following dichlorvos intoxication were scored using semi-quantitative grading of five grades (0–4), taking into account the severity of symptoms and the level of impairment.
consideration the severity of the reactions (0=mild reactions, 2=moderate reactions, 3=severe reactions, 4=mortality). Because there were no deaths in the experiment, no rats scored 4.

**Purification of rabbit serum PON1**  
Paraoxonase was purified from rabbit serum as previously described in detail. The collected protein was subjected to centrifugal ultrafiltration on centrulps YM-10 (Millipore, Billerica, Massachusetts, USA) so as to concentrate to >5000 U/ml. The enzyme was dissolved in buffer solution (136 mM NaCl, 0.1 mM CaCl₂) for injection before usage.

**LC-MS method**  
**Sample preparation**  
Plasma aliquot of 50 μl was added to 2.5 μl of the internal standard indomethacin solution that was prepared in methanol of 50 ng/ml. Then, acetonitrile (100 μl) was added to the mixture. The resulting mixture was vortex-mixed for 1 min and then centrifuged at 12000 r for 5 min. The supernatant was filtered through a 0.45-μm Millex肠胃®-LH filter, and 20 μl of the filtrate was injected into the LC-APCI-MS.

**LC-MS condition**  
The HPLC analysis was performed on a Diamonsil C18 column (150×4.6 mm inside diameter, 5 μm.) equipped with a phenex C18 guard column (4×3.0 mm inside diameter) at room temperature. Fluid phase was used with solvent consisting of acetonitrile–water–methanoic acid (90:10:0.2, v/v/v) at a flow rate of 0.45 ml/min. The solutions were filtered through a 0.45 μm Millex肠胃®-LH filter before use.

A Thermo Finnigan TSQ Quantum Ultra tandem mass spectrometer equipped with an APCI interface operated in the positive ion mode. The capillary voltage was set to 4.2 kV. The capillary temperature was 270°C, and the desolvation temperature was 420°C. Nitrogen was used as desolvation gas (pressure 30 Arb) and assist gas (pressure 5 Arb), and argon was used as collision gas at a pressure of 1.2 mTorr. Detection was performed in selected reaction monitoring (SRM) mode. The collision-induced dissociation voltages were 23 eV (dichlorvos) and 44 eV (indomethacin), respectively. Ionic reaction for quantitative analysis was m/z 221 → m/z 109 (dichlorvos) and m/z 358 → m/z 111 (indomethacin), respectively. Scan time was 0.3 s. Following analysis by an LC-APCI-MS system, the retention time for dichlorvos was 4.07 s, and that for indomethacin was 4.92 s (figure 1).

**Linearity**  
Added dichlorvos standards to blank plasma detected the ratio of peak area of 6 μg/ml, 5 μg/ml, 2 μg/ml, 1 μg/ml, 0.5 μg/ml and 0.2 μg/ml of dichlorvos to internal standard peak area.

With defined dichlorvos standards peak area as X-axis, ratio of dichlorvos standards peak area to internal standard peak area as Y-axis, we got the regression line of dichlorvos: Y=-0.0404282 +0.16048*X, R²=0.99945.  
The linearity was 0.2–6 μg/ml.

**Statistics**  
A statistical moment model was used to calculate toxicokinetics parameters for dichlorvos by SPSS16.0 software. Statistical analyses were conducted using SPSS16.0 software.

---

**RESULTS**  
**Toxicokinetics of dichlorvos**  
**Dichlorvos concentration**  
The concentrations of dichlorvos in B group were statistically different from A group (p<0.05), while atropine + PAM-CI did not alter dichlorvos concentration statistically (p>0.05) (table 1).

**Dichlorvos toxicokinetics parameters**  
The toxicokinetics parameter differences between four groups were performed by ANOVA. There were no statistical differences between AUC (0-∞) of groups A and C, but the effect of PON1 was obvious in groups B and D when compared with group A. There were no statistical differences between MRT (0-∞) of the four groups. Cmax was obtained by observing the peak concentrations in table 1. The statistical result coincided with that of AUC (0-∞) (table 2).

**Clinical signs**  
The clinical signs were alleviated by PON1, atropine+ PAM-CI and co-treatment, and the effect of PON1 was not different with atropine+ PAM-CI statistically. However, co-treatment achieved the best effect (table 3).

**DISCUSSION**  
The 1/2 LD₅₀ (LD₅₀, 18–20 mg/kg by ip) of dichlorvos was selected so that the most severe signs of poisoning would be elicited but no death arose through preliminary experiment. The
The results of the study suggested that PON1 can effectively decrease blood dichlorvos concentration, reduce the peak concentration of dichlorvos and lessen the amount that enters the blood. It is possible that a high degree of protection was provided by PON1 pretreatment in animals challenged with organophosphates by measuring cholinesterase activities in different tissues.\textsuperscript{10–12} Our results provide further direct evidence by showing changes in concentration with PON1 pre-treatment. We also compared the hydrolytic effect of PON1 with atropine + PAM, the most widely used therapy clinically. The results of the study show that atropine + PAM does not affect the metabolism of dichlorvos, which is consistent with the findings of a previous study.\textsuperscript{13} Co-treatment does not alter the impact of PON1 on dichlorvos concentrations, which implies that there is no interaction between PON1 and atropine + PAM-CI; therefore, it is postulated that co-treatment may be feasible in the clinical treatment of human organophosphate-related toxicity.

We also investigated the clinical signs of the four groups. Decreased blood dichlorvos concentration in the PON1 pretreatment group resulted in better clinical outcome when compared with the dichlorvos administration group. At this dose, the PON1 effect is comparable to that of atropine + PAM-CI, which is the most common antidotal treatment used clinically. In addition, another novel finding in our study was that the clinical signs were relieved the most by co-treatment. This may have important implications on the future treatment of organophosphates poisoning.

Interest in PON1 arises from the hypothesis that individuals with low serum activity of this enzyme would be expected to have a diminished ability to metabolise organophosphates. This hypothesis implies that serum PON1 has a pivotal role in the detoxification of the organophosphate. After the hypothesis was raised, some studies have been designed to evaluate the protective effect of PON1 by pretreating animals with PON1 before organophosphates administration. Such studies are needed before any definite inference can be drawn on the role that serum paraoxonase levels have in the protective effect on insecticides. However, in the treatment of human organophosphate-related toxicity, antidotes are administered after exposure, so further investigations of PON1 should be conducted with PON1 administration after organophosphate exposure on the theoretical basis of protective function.

In conclusion, PON1 can hydrolyse organophosphates in vivo and improve clinical signs of organophosphate toxicity. The use of PON1 in organophosphates poisoning requires further investigations.

### REFERENCES


