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Short communication

Toxicokinetics of ergovaline in the horse after an intravenous administration

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Abstract – The toxicokinetics of ergovaline (an ergopeptide mycotoxin present in some grasses infected with endophytic fungus of the genus *Neotyphodium*) were studied after intravenous administration of a single dose of 15 µg/kg bwt in four gelding horses. Plasma ergovaline concentrations were measured by high performance liquid chromatography, and the kinetic data were described by a three-compartment model. The elimination half-life and the total clearance of ergovaline were found to be 56.83 ± 13.48 min and 0.020 ± 0.004 L/min·kg, respectively. According to the toxicological data previously reported in the horse, and in spite of the very low dose administered, clinical signs were observed, including excessive coolness of the ears and the nose, excessive sweating and prostration.

ergovaline / mycotoxin / horse / kinetics / *Neotyphodium*

Résumé – **Étude toxicocinétique de l'ergovaline chez le cheval après injection intraveineuse.** Les paramètres toxicocinétiques de l'ergovaline (un alcaloïde ergopeptidique présent dans les graminées porteuses d'un champignon endophyte du genre *Neotyphodium*) ont été déterminés après l'injection intraveineuse d'une dose de 15 µg/kg chez quatre chevaux hongres. Les concentrations plasmatiques d'ergovaline ont été mesurées par chromatographie liquide haute performance et les données cinétiques décrites selon un modèle à trois compartiments. La demi-vie d'élimination et la clairance totale de l'ergovaline ont été respectivement de $56,83 \pm 13,48$ min et de $0,020 \pm 0,004$ L/min·kg. Malgré la très faible dose administrée, des signes cliniques compatibles avec les données toxicologiques rapportées dans l'espèce équine ont été observés, à savoir hypothermie des oreilles et du nez, hyper-sudation et prostration.

ergovaline / mycotoxine / cinétique / cheval / *Neotyphodium*

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was removed and stored at -20°C until analysis. For each sampling time, an aliquot of plasma was analysed according to a previously described high-performance liquid chromatographic method, using ergotamine as internal standard [11]. Briefly, the method consisted in a simple extraction procedure with diethyloxide followed by an HPLC analysis on a Zorbax C_{18} column, with the excitation and emission wavelength fixed at 250 and 420 nm respectively on a fluorimetric detector. The method was improved by using 4 mL of plasma for analysis, instead of 1 mL as in [11]. After a standard validation procedure, the limit of quantification (LOQ) was lowered from 3.5 ng/mL to 0.7 ng/mL of horse plasma.

2.3. Kinetic data analysis

A kinetic analysis of the concentration profiles was performed, using a programme for the residuals method. The number of exponents needed for each set of data was determined using the SIPHAR (v. 4. 0) software [10]. The total area under the plasma concentration time curve (AUC), was calculated by linear trapezoidal rule without extrapolation to infinity. The plasma concentration at $t = 0$ (C_0) and the volume of the central compartment (V_C) were calculated with the usual equations [9].

Each individual plasma profile could be fitted to the following tri-exponential function, characterizing a three-compartment model, and a three-phase kinetic process:

$$C = A_1 e^{-\lambda_1 t} + A_2 e^{-\lambda_2 t} + A_3 e^{-\lambda_3 t}$$

where A_1 , A_2 and A_3 are ordinari intercepts, and λ_1 , λ_2 and λ_3 are slope values [9]. The tri-exponential function was chosen because it resulted in the best fitting of the experimental data (correlation coefficient of 0.9997 for the terminal phase).

3. RESULTS

The logarithmic plot of the mean plasma ergovaline concentration versus time, after intravenous administration, is shown in Figure 2. A very rapid decrease of the mycotoxin plasma levels occurs, the LOQ being reached 150 min post-dosing.

The individual values of different kinetic parameters for each horse are presented in Table I. The plasma clearance and the elimination half-life for terminal phase (λ_3) of ergovaline were found to be $0.020 \pm 0.004 \text{ L/min}\cdot\text{kg}^{-1}$ body weight and $56.83 \pm 13.48 \text{ min}$, respectively.

All animals treated showed the same clinical signs, observed about half an hour after the ergovaline injection: coolness of the ears and the nose, excessive sweating, prostration, and difficulty in urination.

4. DISCUSSION

The plasma clearance and the elimination half-life for terminal phase (λ_3) of ergovaline obtained in horse are of the same order as those obtained in sheep and goats ($0.020 \text{ L/min}\cdot\text{kg}^{-1}$ body weight ; 23.6 min, and $0.034 \text{ L/min}\cdot\text{kg}^{-1}$; 32.4 min, respectively) using a two-compartment model [7, 11].

The clinical signs, similar to those previously observed in goats [7] and in horses [14], are in accordance with the known acute effects of the toxin [13], particularly those linked to the peripheral vasoconstrictive activity of ergot alkaloids.

The rapid decrease of ergovaline in blood, observed after intravenous administration in horses, is of particularly great interest: it indicates that low mycotoxin plasma levels should be expected in animals, after ingestion of grass infected with *Neotyphodium*. A more complete kinetic study should be undertaken in order to determine the paths (urine or/and bile) which are involved in the elimination of ergovaline in

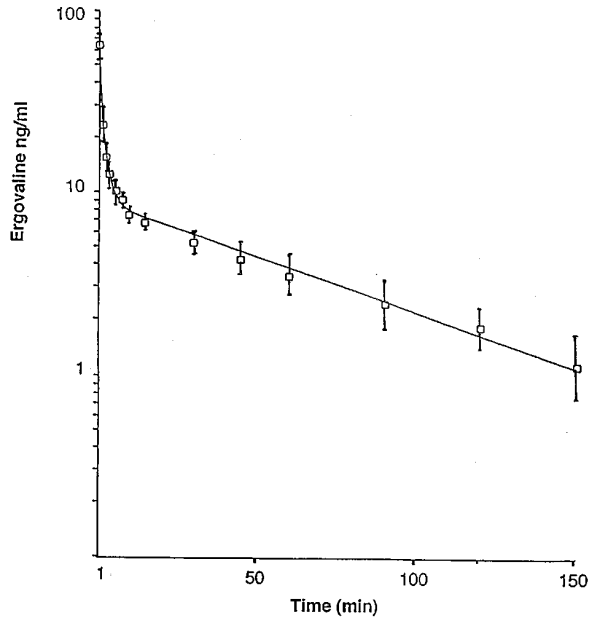


Figure 2. Plasma log.ergovaline concentration versus time profile in horses (mean \pm sd, $n = 4$), after an intravenous administration of a single dose of 15 $\mu\text{g}/\text{kg}\cdot\text{bwt}$.

Table I. Toxicokinetic parameters describing the disposal kinetics of ergovaline in plasma after an intravenous administration (15 $\mu\text{g}/\text{kg}\cdot\text{bwt}$) in four horses.

<i>Kinetic parameters (units)</i>	<i>Horse 1</i>	<i>Horse 2</i>	<i>Horse 3</i>	<i>Horse 4</i>	<i>Mean</i>	<i>SD</i>
λ_1 (per min)	1.93	2.04	1.29	1.50	1.690	0.354
λ_2 (per min)	0.41	0.22	0.15	0.17	0.237	0.118
λ_3 (per min)	0.011	0.014	0.015	0.009	0.012	0.002
A_1 (ng/mL)	278.42	389.34	160.67	179.54	251.99	105.12
A_2 (ng/mL)	22.52	19.27	9.17	6.32	14.32	7.79
A_3 (ng/mL)	8.24	7.12	7.28	7.17	7.60	0.83
$t_{1/2\alpha}$ (min)	58.89	48.91	44.58	74.94	56.83	13.48
Cl (L/min·kg)	0.017	0.020	0.026	0.018	0.020	0.004
AUC (ng·min/L)	949.80	779.11	653.80	932.42	828.78	139.61
MRT (min)	67.43	46.11	46.85	90.22	62.65	20.86
Vc (L/kg)	0.048	0.036	0.085	0.078	0.062	0.024
Co (ng/mL)	309.78	415.73	177.12	193.03	273.92	115.52

A_1, A_2, A_3 = pre-exponential constants; $\lambda_1, \lambda_2, \lambda_3$, exponential constants for triexponential equation; $t_{1/2\alpha}$, half-life for terminal phase; Cl = body clearance; AUC = area under the plasma concentration curve; MRT = mean residence time calculated until the last sampling time; Vc = volume of the central compartment ($Vc = X_0/Co$ where X_0 is the administered dose); $Co = A_1 + A_2 + A_3$.

the equine species. Moreover, the dose of 15 µg/kg could be used as a toxicity threshold, in the absence of quantitative toxicological data (NOAEL: no observed adverse effect level, LOEL: lowest observed effect level) concerning ergovaline.

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