

Dexamethasone serum concentrations after intravenous administration in horses during race training

S. McClure^{1,2}, C. Fenger^{3*}, K. Kersh², B. Brown⁴, G. Maylin⁵, W. Duer⁶, L. Dirikolu⁷, K. Brewer⁸, J. Machin⁹ and T. Tobin⁹

¹Midwest Equine Surgery and Sports Medicine, 2615 Eastgate Drive, Boone, IA 50036, USA; ²Department of Veterinary Clinical Sciences College of Veterinary Medicine, Iowa State University, 1800 Christensen Drive, Ames, IA 50011-1134, USA; ³Equine Integrated Medicine, PLC, 4904 Ironworks Rd., Georgetown, KY 40324, USA; ⁴Bradley Brown, private practice, 425 Sacree Rd., Shelbyville, KY 40065, USA; ⁵New York Drug Testing and Research Program, Morrisville State College, 777 Warren Rd, Ithaca, NY 14853, USA; ⁶Duer Forensic Toxicology LLC, 1621 Gulf Blvd #102, Clearwater, FL 33767-2928, USA; ⁷Department of Comparative Biomedical Sciences, School of Veterinary Medicine, Louisiana State University, Skip Bertman Drive, Baton Rouge, LA 70803, USA; ⁸Kimberly Brewer, private practice, 15775 Cypress Creek Lane, Wellington, FL 33414, USA; ⁹The Maxwell H. Gluck Equine Research Center, University of Kentucky, 1400 Nicholasville Rd, Lexington, KY 40503, USA; drfenger@hotmail.com

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Abstract

Dexamethasone (DXM) sodium phosphate is a widely used corticosteroid for inflammatory conditions in horses, regulated in racing jurisdictions in the USA by a 0.005 ng/ml serum/plasma threshold. This study seeks to describe serum concentrations of DXM at 48 and 72 h after intravenous administration of 20 mg DXM sodium phosphate over 1 to 5 days, and to identify a possible source of DXM overages. 74 horses (39 Thoroughbreds, 13 Standardbreds, 22 Quarter Horses) in active race training received 20 mg DXM sodium phosphate. Serum was collected before injection, at 48 and 72 h post last injection, and analysed by LC/MS-MS (limit of quantification (LOQ) = 2.5 pg/ml). No differences were identified by ANOVA ($P \leq 0.05$) for racing breeds, age, gender or the number of days of DXM sodium phosphate administration, so data were pooled for each time point. The DXM serum concentration at 48 h (mean $\pm$ standard deviation, range) was 2.18 ± 1.56 pg/ml (<2.5 to 40 pg/ml). Summary statistics could not be derived for 72 h DXM serum concentration data owing to censored data, but ranged from <2.5 to 95.8 pg/ml. There was one extreme outlier (Tukey) at 48 h, and two extreme outliers at 72 h. A separate study was conducted using sedentary experimental horses to determine the likelihood that positive DXM samples could result from environmental transfer. Urine was collected from a mare 2 to 3 h post administration of 20 mg DXM. Hay with 100 ml of the DXM (17 ng/ml) containing urine was offered to each of six experimental horses and blood was collected at 0, 4, 8, 12, 16, 20 and 24 h. All six horses had plasma DXM concentration above the limit of detection and five of six had plasma DXM concentrations above the LOQ for at least one sample time.

Keywords: horse, dexamethasone, racing, regulatory thresholds

1. Introduction

Dexamethasone ($C_{22}H_{29}FO_5$, molecular weight 392.461, DXM) sodium phosphate is an FDA-approved, short-acting therapeutic corticosteroid used in equine practice for the prevention and treatment of inflammatory and allergic conditions. Dexamethasone for parenteral administration can be formulated as a highly water-soluble pro-drug, DXM 21-phosphate disodium salt (Soma *et al.*, 2013; Tobin, 1981). Following administration, the ester linked phosphate group

is hydrolysed by plasma esterases (Samtani and Jusko, 2005) releasing the relatively short plasma half-life DXM (Soma *et al.*, 2013).

Equine athletes are at risk of inflammatory airway disease (IAD), which affects up to 80% of 2-year-old racing horses (Christley *et al.*, 2001), resulting in impaired gas exchange and sub-optimal performance (Couëttil *et al.*, 2016). Approximately 14% of all age groups of racing horses suffer from IAD (Wood *et al.*, 2005) and there is evidence that

pulmonary inflammation contributes to the incidence of exercise-induced pulmonary haemorrhage (EIPH) (McKane and Slocombe, 2010). Because DXM has a short plasma half-life and duration of action (Soma *et al.*, 2013), it is uniquely suited for use for treatment of IAD and other inflammatory and allergic conditions of horses approaching competition. The appropriateness of this recommendation is reflected in the Canadian Para-Mutuel Agency (CPMA) recommendation of a 48 h withdrawal after 25 mg intravenous (IV), 20 mg intramuscularly (IM), or a 5 day course of 10 mg orally (PO) (Anonymous, 2016), and the Fédération Equestre Internationale which has a published detection time of 2 days after IV administration of 10 mg DXM sodium phosphate (Anonymous, 2018). Similar to equine athletes, humans competing under World Anti-Doping Agency (WADA) rules cannot compete under the influence of corticosteroids. Exceptions to this rule are peri- or intra-articular corticosteroids and systemic corticosteroids when medically necessary, as determined by a Therapeutic Use Exemption (TUE), which is an application to WADA before administration. The purpose of these TUEs is to permit the administration of appropriate therapeutic medication to athletes. Corticosteroids represent the largest category of approved TUEs in human athletes since the TUE process and its predecessor program was instituted in 1986 (Fitch, 2016).

In the United States, many racing jurisdictions have adopted a regulatory threshold of 5 pg/ml DXM in plasma or serum pursuant to a recommendation by the Racing Medication and Testing Consortium (RMTC), which is accompanied by a dose recommendation of 0.05 mg/kg of DXM sodium phosphate by IV or IM administration or DXM by oral administration, and a withdrawal recommendation of 72 h. The RMTC typically determines thresholds based on the application of a statistical method called the 95/95 tolerance (Owen, 1968), although it is not clear whether this statistical method was used in the case of DXM. The RMTC references an unpublished pharmacokinetic study in Thoroughbreds as the basis for this threshold and withdrawal (ARCI Controlled Therapeutic Medication Schedule for Horses – Version 4.1 Revised – January, 2019). Introduction of this threshold has been attendant with a large spike in racing medication overages for DXM (ARCI, 2019).

This study was undertaken in order to test the hypothesis that the RMTC-based threshold and withdrawal recommendation can be relied upon where DXM is used in fit horses in race training and to identify a possible inadvertent source of DXM overages. Specifically, we investigated the potential effect of multiple day courses of DXM sodium phosphate treatment, breed differences by including Thoroughbreds, Standardbreds and Quarter Horses under actual training conditions, and the effect on serum DXM concentration of exposure of horses to

urine from a DXM sodium phosphate treated horse. We chose 48 and 72 h post-DXM sodium phosphate administration for collection of serum samples because most regulatory agencies use either a 48 or 72 h withdrawal recommendation. Further, we investigated the possibility that hay contaminated with DXM containing urine might cause random positive tests / overages for DXM.

2. Materials and methods

Experiment 1

Study facilities and animals

Privately owned Thoroughbred, Standardbred and Quarter Horse racehorses in race training in the practice population of two of the authors (CF, BB) were used throughout. Horses were stabled on the racetrack and were housed and trained according to standard procedures at racing facilities at Prairie Meadows Racetrack (Altoona, IA, USA) and the Red Mile (Lexington, KY, USA). The feed, bedding and water sources were consistent with routine management at each facility. Training adhered to regimens consistent with the type of racing specific to the racing discipline. Informed consent was obtained for all horses enrolled. Inclusion criteria were a full clinical examination, mucus present on endoscopy, with or without trans-tracheal cytology for a diagnosis of IAD or other inflammatory or allergic condition requiring therapy with 20 mg DXM sodium phosphate (Bimeda, Oakbrook Terrace, IL, USA) IV. Treatment was based on the clinical examination and diagnosis by the examining investigator, a signed owner consent form, long-term trusted relationships between the investigator and trainer to ensure trainer compliance and active participation in racing or fast workouts in preparation for racing. Exclusion criteria were injections with betamethasone or DXM within 7 days, or any other medications within the 24 h prior to blood collection in order to minimise potential interference with the analytical method. The study was approved by the Iowa State University Animal Care and Use Committee.

Experimental design

All racehorses in two investigators' practices that fulfilled the inclusion criteria with none of the exclusion criteria were enrolled in the study. The investigators administered 20 mg DXM sodium phosphate by IV injection, and took all precautions to prevent other exposure to DXM in the training barns. Administration of topical or oral DXM or betamethasone was restricted in the training barns. In order to replicate the usual clinical usage patterns of DXM sodium phosphate, the only restrictions on co-administered medications were the restrictions within 24 h of blood collection in order to prevent interference with the analytical methodology. Any concomitant medications

were recorded. The time of day of the treatment and number of consecutive treatments were recorded. Blood samples were drawn into 10 ml serum separator vacuum tubes immediately preceding the first dose of DXM, and at 48 and 72 h after the last dose of DXM. These samples were allowed to clot, then refrigerated at 2–3 °C, centrifuged within 4 h and the serum transferred to cryovials and stored at -70 °C for batch analysis. In seven cases where pre-injection samples were damaged such that they could not be analysed, a complete review of the horse's medical record for the last month was performed to ensure that no prior injection with any DXM or betamethasone containing product had occurred.

Analytical methods

The analytical procedure followed was the ISO 17025/RMTC accredited quantitative analytical procedure for DXM in place in the New York Drug Testing and Research Laboratory. The analytical reference standard for DXM was purchased from Sigma Aldrich (St. Louis, MO, USA) and d4-dexamethasone (DXM-d4) from CDN Isotopes (Pointe-Claire, Canada), respectively. Stock solutions of DXM and DXM-d4 were prepared at 1 mg/ml in methanol. Acetonitrile and methanol were purchased from EMD Millipore (Sigma Millipore, St. Louis, MO, USA), and methyl tert-butyl ether and ammonium formate were purchased from Fisher Scientific (Lenexa, KS, USA). Deionised water was filtered onsite to the specification of 18.2 meg Ω . Ethanol was purchased from Pharmco-AAPER (Brookfield, CT, USA). All reagents were of HPLC grade or better.

Working DXM solutions were prepared by dilution of the 1 mg/ml stock solution with ethanol to concentrations of 1 and 100 pg/ μ l. Plasma calibrators in concentrations of 2.5, 5 and 10 pg/ml were prepared by the addition of the working standard solution to plasma harvested from experimental horses known to be drug-free. Calibration curves and negative control samples were prepared fresh for each quantitative assay.

The 2.0 ml aliquots of samples were prepared alongside calibration curve and negative control samples. Internal standard was added to each tube. Samples were mixed by vortex, and 5 ml of methyl tert-butyl ether was added. Samples were mixed by rotation for 10 min, centrifuged at 2,400 $\times$ g for 5 min, the emulsion broken, and centrifuged again at 2,400 $\times$ g for 5 more min. The top ether layer was removed and dried under nitrogen. Samples were dissolved in 50 μ l of equal parts acetonitrile, methanol, and deionised water. 2.5 μ l was injected into the LC-MS/MS system (Agilent Technologies, Inc, Santa Clara, CA, USA) coupled with a U/HPLC chromatography system (Agilent).

The concentration of DXM was measured in plasma by LC-MS/MS using positive electrospray ionisation with Agilent Jet Stream technology. Chromatography employed a Zorbax SB-C18 (Agilent) column with a length of 100 mm and a pore size of 3.0 μ m. The beginning mobile phase composition was 50% 5 mM ammonium formate in de-ionised water and 50% acetonitrile (ACN). The initial ACN concentration was held at 50% for 2.75 min, ramped to 95% until 3.25 min, and held at that concentration for until a runtime of 4.5 min at which the mobile phase composition was reset to the initial settings.

Detection and quantification were conducted using selective reaction monitoring (SRM) of initial precursor ion for DXM (mass-to-charge ratio 393.5 m/z) and the internal standard (397.5 m/z). The response for the product ions for DXM (m/z 373, 355, 337) and the internal standard (m/z 377) were plotted and peaks at the proper retention time integrated using MassHunter software (Agilent). MassHunter software was used to generate calibration curves and quantitate DXM in all samples by linear regression analysis.

The validation of the method employed for the analysis of DXM contained a calibration curve performed encompassing 2.5, 5.0 and 10.0 pg/ml DXM. The response was linear and gave correlation coefficients (R^2) of 0.99 or better. Quality control sample replicates were performed ($n=7$). The inter-day accuracy was 1.9% for 5 pg/ml DXM. The intra-day accuracy was 8.3% for 5 pg/ml DXM. The inter-day precision was 8.4% for 100 pg/ml DXM. The intraday precision was 10.6% for 100 pg/ml DXM. The technique was optimised to provide a limit of quantitation (LOQ) of 2.5 pg/ml. The limit of detection (LOD) was 1.0 pg/ml.

Data analysis

The 48 h and 72 h post administration serum DXM concentrations were analysed for percentage of censored (below LOQ) data, effects of number of consecutive days of treatment using Robust Regression on Order and General Linear Model statistical methods, with statistical significance at $P<0.05$ (Helsel, 2012). The 48 and 72 h datasets were first analysed for percent censored data, then normality tests (Shapiro-Wilk, Anderson-Darling, Lillefors and Jarque-Bera) were performed on uncensored (above LOQ) data [using R-programming language] in order to determine the most appropriate statistical analysis for threshold determination. Where data were normally distributed, the effects of age, gender, breed and number of days treated were analysed by ANOVA [XLSTAT®, ADDInsoft 2016 <https://www.xlstat.com/en/> as an Excel® for Mac 2011, Microsoft add-in]. Summary statistics for each data subgroup were obtained for each group using RROS (Helsel, 2012) in R-programming language and bootstrap analyses with 100,000 resamplings [XLSTAT®, ADDInsoft 2016 <https://www.xlstat.com/en/> as an Excel®

for Mac 2011, Microsoft add-in]. Bootstrap with resampling was performed to estimate population summary statistics. Thresholds for each dataset were determined using two different methods; (95/95) tolerance interval (Owen, 1968) and Gauss-Camp-Meidell (Savage, 1961) (GCM), as previously described (Machin *et al.*, 2019). The 95/95 tolerance is calculated by:

$$x_{\text{tol}} = m + k s_x$$

Where x_{tol} is the 95/95 tolerance threshold, m is the mean and s_x is the standard deviation of the sample population. The k value is a tolerance limit factor equivalent to a z -score corrected for sample size (Owen, 1968). The GCM is a probability inequality that provides bounds for the probability that the plasma concentration will fall within a given P -value, and requires only a unimodal distribution (Savage, 1961).

Experiment 2

Study facilities and animals

Six sedentary Thoroughbreds owned by Equine Integrated Medicine, PLC (5 mares, 1 stallion, mean $\pm$ standard deviation (SD), range 10 $\pm$ 5 years old, 3 to 17 years) were used in the DXM urine environmental exposure study. Horses were stabled in the research facility of Equine Integrated Medicine, PLC, and all procedures approved by its Animal Care and Use Committee (# AAW0012015). Feed and water sources were consistent with routine management at the facility, horses were bedded on dried grass bedding, fed a 12% Complete Feed mixture, and offered a timothy, orchard grass and bluegrass mixture *ad libitum*. Prior to the onset of the study, the stalls were stripped, and cleaned by pressure washing and phenolic antiseptic, allowed to completely dry for a minimum of one week, and bedded with fresh grass bedding. No medications were administered within 1 month of the study, and the horses were not in proximity to any other horses during the study.

Experimental design

A mixed breed mare was administered 0.05 mg/kg DXM IV. A Foley catheter (#J0447H, Jorgensen Laboratories, Inc, Loveland, CO, USA) was placed at the time of administration, and the bladder evacuated. The Foley catheter was left in place, and urine collected at 3 h post administration. The six experimental horses were stall housed and in addition to the *ad libitum* mixture of grass hay, 2.27 kg of Lucerne hay was offered which was sprinkled with 100 ml of the DXM containing urine. Blood samples were collected from these horses at 0, 1, 3, 6, 12, and 24 h and the samples were analysed as described above. Because

of the expected presence of lower serum concentrations of DXM in the contamination experiment, the LCMS method was optimised for more sensitive detection of DXM by the addition of a 1.0 pg/ml standard. The LOD was 0.5 pg/ml.

3. Results

Experiment 1

74 horses met the inclusion criteria (Thoroughbreds, $n=39$, Standardbreds $n=13$, Quarter Horses $n=22$). An ANOVA showed no difference between age, gender, breed or number of daily treatments, so serum concentrations of all horses were pooled for threshold determination analysis for each of the 48 and 72 h datasets. Three additional horses were excluded for pre-treatment DXM serum concentrations (33 pg/ml, 4.5 pg/ml and 4.0 pg/ml) above the LOQ, despite no identifiable exclusion criteria. Although these horses were excluded, evaluation of the 48 and 72 h data with these horses included did not change the results.

The number of horses that received each series of consecutive daily treatments, number censored and presence of outliers is shown in Table 1. Means and standard deviations were not determined because the numbers within each group were insufficient to employ RROS. 67 horses had pre-injection serum samples below the LOQ of the analytical method for DXM. Seven horses did not have pre-injection serum samples analysed, but a review of the medical history for these horses showed no previous DXM administration or exposure, so these horses were not excluded from the analysis. Of these seven horses, five had DXM serum concentrations below the LOQ at both 48 and 72 h. One had DXM serum concentration below the LOQ at 48 h, and 4.1 pg/ml DXM serum concentration at 72 h. One had DXM serum concentration below the LOQ at 48 h, and 95.8 pg/ml at 72 h. Because the 48 h DXM concentrations were below the LOQ for all of the horses without pre-injection serum sample analyses, they were not excluded from the analysis. Three horses had DXM concentrations above the LOQ in the pre-injection samples with no history or evidence of DXM or betamethasone exposure within the preceding 14 days. These horses were excluded from the analysis, although the 48 and 72 h DXM concentrations would not have affected the analysis, had they not been excluded. Two of these horses had censored DXM concentrations at both 48 and 72 h, and the third horse had 6.4 pg/ml of DXM at 48 h and censored concentration at 72 h.

Concomitant medications were administered to 54 horses, and are summarised in Table 2. The total number of horses receiving each concomitant medication was insufficient to apply any statistical methods, so these data were pooled for statistical analysis. One extreme outlier was evident in a horse that received both phenylbutazone and flunixin.

Table 1. The number of horses receiving each group of consecutive daily treatments, number censored, and presence of outliers.

Number of days	Number of horses	48 h		72 h	
		Number censored	Outlier (pg/ml)	Number censored	Outlier (pg/ml)
1	24	17	None	19	95.8 & 10
2	14	10	None	14	None
3	17	15	None	16	None
4	6	5	40.6	4	None
5	13	7	None	12	None

Table 2. The number of horses receiving concomitant medications, number censored, and presence of outliers.

Concomitant medication	Number of horses	48 h		72 h	
		Number censored	Outlier (pg/ml)	Number censored	Outlier ³ (pg/ml)
Phenylbutazone	13	10	None	11	95.8*
Flunixin	4	4	None	3	95.8*
Ceftiofur crystalline free acid [Excede®] ¹	1	1	None	1	None
Gentamicin	10	9	None	8	None
Trimethoprim-sulfadiazine	6	5	None	6	None
Oxytetracycline	10	7	None	9	None
Acepromazine	1	1	None	0	None
Dantrolene	1	1	None	0	None
Polysulfated Glucosaminoglycan [Adequan®] ²	3	1	None	2	None
Dimethylsulfoxide	2	1	None	2	None
Neomycin-Polymixin-Bacitracin Ophthalmic ointment	1	1	None	1	None
Vitamin E/Selenium	2	2	None	2	None

¹ Zoetis, Parsippany, NJ, USA.² American Regent Animal Health, Shirley, NY, USA.³ Some horses received more than one concomitant medication. The highest outlier horse (*) at 72 h received two concomitant medications, phenylbutazone and flunixin.

The serum concentrations of DXM for the 48 h samples are presented in Figure 1. The presence of one outlier in the 48 h dataset (40.6 pg/ml) and two outliers in the 72 h dataset (10 and 95.8 pg/ml) are noted. The percent censored data (below the LOQ of 2.5 pg/ml) for the 48 and 72 h datasets were 74 and 90%, respectively. Normality tests (Shapiro-Wilk, Anderson-Darling, Lilliefors and Jarque-Bera) for each dataset indicated that non-censored data (serum concentrations above the LOQ) were normally distributed.

The 48 h serum concentration (mean $\pm$ SD) outlier omitted) for the 48 h group was 2.18 ± 1.56 pg/ml. The tolerance limits or thresholds generated using 95/95 or GCM ($P=0.05$, $P=0.01$) for the 48 h samples are presented

in Table 3. Thresholds could not be determined for the 72 h data because the 90% censored data precludes the determination of a threshold from these data (Helsel, 2012). Nine of the 74 horses had measurable DXM concentrations at 72 h. Of those nine, seven horses, including two extreme outliers, had higher DXM concentrations at 72 than 48 h.

Experiment 2

Results of the second experiment are shown in Table 4. Three of six horses had quantifiable concentrations of DXM, five of six had DXM concentrations above the LOD, and only one horse had no detectable DXM during the 24 h collection period.

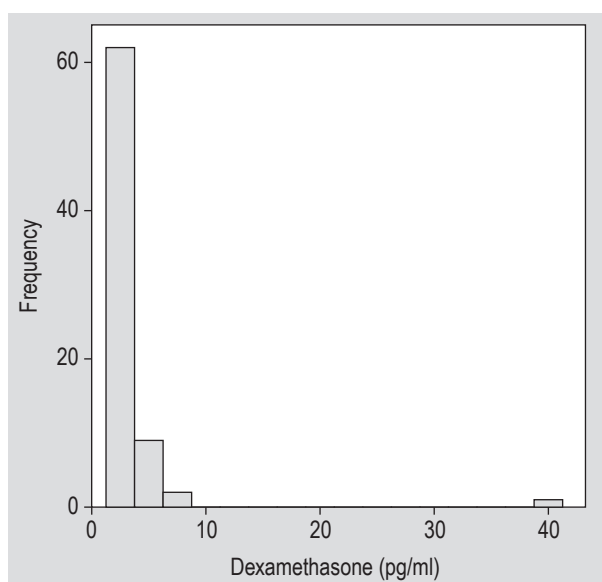


Figure 1. Frequency histogram for dexamethasone (DXM) serum concentrations 48 h post 1-5 daily doses of 20 mg DXM sodium phosphate in Thoroughbred, Standardbred and Quarter Horse racehorses in race training.

Table 3. Percent censored data, and threshold concentrations in pg/ml as determined by 95/95 tolerance, Gauss-Camp-Meidell (GCM) at $P=0.05$, corresponding to 1 in 20 risk of violating the threshold, and $P=0.01$, corresponding to a 1 in 100 risk of violating the threshold.

	48 h	72 h
% censored data points	74%	90%
95/95 tolerance	6.61	na ¹
GCM ($P=0.05$)	6.77	na
GCM ($P=0.01$)	12.4	na

¹ na = not applicable.

Table 4. Serum concentrations (pg/ml) of dexamethasone (DXM) detected in experimental horses after exposure to DXM containing urine contaminated lucerne hay.¹

Time after hay added (h)	Horse 1	Horse 2	Horse 3	Horse 4	Horse 5	Horse 6
0	NDD	NDD	NDD	NDD	NDD	NDD
1	NDD	NDD	>LOD	NDD	>LOD	NDD
3	NDD	NDD	NDD	>LOD	NDD	NDD
6	NDD	1.7	1.0	>LOD	NDD	–
12	NDD	>LOD	NDD	NDD	NDD	NDD
24	4.4	NDD	1.5	NDD	NDD	NDD

¹ NDD = no DXM detected; LOD = limit of detection. One 6 h sample could not be collected from horse 6, indicated by '–'.

4. Discussion

This population study in racing horses was undertaken to evaluate the post DXM sodium phosphate administration serum concentrations of DXM under field conditions, and to identify possible sources of blood DXM concentrations above the regulatory threshold of 5 pg/ml. No difference was found among the different racing breeds and disciplines which permitted pooling of the data for the purpose of calculating a 48 h threshold. Different racing breeds have previously been shown to exhibit some different pharmacokinetic parameters for glycopyrrolate (Rumpler *et al.*, 2014), differences that may be associated with differences in study design, pharmacogenetics, or physiologic differences based on characteristics such as renal or splanchnic blood flow associated with the type of exercise. In this study, no effects of different breed or training regimes were evident for either the 48 or 72 h time DXM serum concentrations.

No difference was found among serum concentrations of DXM post single or multiple day treatments, indicative of no significant accumulation of DXM in horses, consistent with previous published literature. Haspel *et al.* (2018) found the $AUC_{0-24\text{ h}}$ after a single dose to be almost identical to $AUC_{0-\infty}$, indicating minimal accumulation after the conservative dose of 5 mg used in their study. Accumulation ratios (AR) calculated from the pharmacokinetic data presented in previous studies using a low dose (AR of -0.01, Haspel *et al.*, 2018) as well as a higher therapeutic dose of 0.05 mg/kg (AR of -0.002, Grady *et al.*, 2010) fall well below an AR of 1.2, confirming the lack of accumulation (Li *et al.*, 2012) of DXM in horses dosed at 24 h intervals throughout the typical therapeutic dose range. In addition to the extreme outliers, of the nine uncensored 72 h DXM serum concentrations, four serum concentrations exceeded the 48 h DXM serum concentration from the same horse, but were below 5 pg/ml.

Three samples had detectable DXM in the pre-injection samples, despite no history of DXM or other corticosteroid injection. It is possible that if the DXM sodium phosphate is withdrawn from the vial, and any bubbles in the solution are expressed from the syringe in the vicinity of the blood collection tubes, sufficient DXM can be aerosolised to cause contamination of the rubber stoppers of the collection tubes, and subsequent measurable concentrations in the blood. These samples were excluded from the analysis, but would have had no effect on the analyses, had they been included.

Three samples in our population present as extreme outliers, defined as values that exceed 1.5 interquartile ranges above the third quartile (Tukey's Outlier Test). One outlier at 48 h was 40.6 pg/ml and two outliers at the 72 h point were 10.1 and 95.8 pg/ml. The horse that had the 10.1 pg/ml DXM serum concentration at 72 h had a serum concentration at 4.4 pg/ml at 48 h, and the horse with the 95.8 pg/ml DXM serum concentration at 72 h had a serum concentration below the LOQ at 48 h. While unexpected, these unusually high concentration data parallel the similar presence of high plasma concentrations in three previous pharmacokinetic papers (Soma *et al.*, 2005, 2013; Symonds *et al.*, 2019). At 48 h in Soma *et al.* (2005), two of six horses had plasma DXM concentrations of 600 pg/ml and 700 pg/ml after the IV administration of 0.05 mg/kg, a slightly higher dose than employed in this study. In a later paper, Soma *et al.* (2013) reported a single outlier of 57 pg/ml at 48 h among six horses after IV administration of 0.05 mg/kg DXM. While Soma did not identify this sample as an outlier, it clearly exceeds 1.5 interquartile ranges above the third quartile in his study since all of the other 48 h samples fell below the LOQ of the study, 10 pg/ml, fulfilling Tukey's criteria as an outlier. High concentrations of DXM at 48 h may represent individual variability in DXM metabolism, but the study design employed in the second Soma *et al.* study (2013) controlled for this possibility. Soma *et al.* (2013) utilised a crossover design, wherein each horse was administered DXM by four different routes, intravenous, intramuscular, oral and intra-articular. Despite the multiple routes of administration, one horse exhibited the 48 h 57 pg/ml DXM plasma concentration only after IV administration and not when DXM was administered by IM or oral routes. Symonds *et al.* (2019) administered 0.04 mg/kg DXM sodium phosphate by nebulization, and had a single 100 pg/ml outlier at 96 h.

An alternative explanation for the high plasma DXM concentrations in the previous Soma *et al.* studies (2005, 2013), as well as the 48 h outlier in this study would be an inadvertent subcutaneous deposition of DXM sodium phosphate at the time of the intravenous administration. Such administration would have been expected to have resulted in a sustained absorption phase, which could not be identified with our study design, but would likely have

been identifiable by the pharmacokinetic parameters of the individual horses in the previous Soma studies. However, the plasma DXM outlier of 100 pg/ml at 96 h in the Symonds *et al.* (2019) study could not be explained by subcutaneous administration, because the study used a single nebulised dose of DXM sodium phosphate. The outliers at 72 h in our study similarly could not be explained by subcutaneous administration, because both horses were below the LOQ at 48 h. The findings of these three previous papers and the results of our field study argue against individual metabolic variability as the source of these outliers, requiring the investigation of an alternative source of these outliers.

The horses in experiment 1 were under the control of the investigators, and received no alternative source of DXM, such as oral powders, topical creams or other compounds. As a prescription medication, DXM is only available through veterinarians, so additional unrecorded intentional administration of DXM is unlikely. However, the racetrack environment is commonly contaminated with therapeutic medications (Barker, 2008), and previous contamination of the stalls by concentrated sources of DXM, such as residual powder or compounds cannot be ruled out. Using the previously published DXM pharmacokinetics (Soma *et al.*, 2013), our highest outlier would have only required 0.14 mg of DXM for a blood concentration of 95.8 pg/ml at maximal blood concentration.

Alternatively, these outliers could have resulted from recycling of excreted drug. Transfer from the environment has been reported for other medications, such as flunixin (Popot *et al.*, 2011), naproxen (Wennerlund *et al.*, 2000) and isoxsuprine (Russell and Maynard, 2000), among others. Dexamethasone is excreted in urine in concentrations as high as 50 ng/ml after low 5 mg dose administration (Chen *et al.*, 1996) making concentrations as high as 200 ng/ml likely after a dose of 20 mg, such as used in our study. Further, some of the DXM dose may be excreted in manure, such as is seen in cattle (Vanhaecke *et al.*, 2011), leaving multiple possible routes of inadvertent ingestion from recycling of administered drug. Using the highest attained plasma concentration and bioavailability from Soma *et al.* (2013) rather than the average values, a volume of distribution equal to the central compartment and assuming that the 95.8 pg/ml outlier was a peak serum concentration, only 19.4 ml (3.9 µg DXM) of urine would need to be consumed to produce this high outlier serum DXM concentration. While this requires many assumptions, the actual bioavailability, absorption and metabolism of DXM in the horse that experienced the 95.8 pg/ml 72 h serum DXM concentration are unknown. However, DXM metabolism and disposition is highly variable (Soma *et al.*, 2013), and could produce a relatively high serum concentrations of DXM if the urine with DXM were considered to be the source. Further, when water evaporates from urine, it becomes a concentrated source

of salt, which may attract some horses. The horses in this field study were protected from exogenous exposure to DXM or betamethasone, and experimental horses in three previous studies (Soma *et al.*, 2005, 2013; Symonds, 2019) experienced similar outlier plasma concentrations, indicating that such extreme outlier plasma concentrations may be a risk among horses exposed to DXM.

One of the three extreme outlier DXM serum concentrations was from a horse administered concomitant medications. This horse received both phenylbutazone and flunixin after its DXM administration. It is possible that factors related to the combination of these two non-steroidal anti-inflammatory drugs contributed to the high 72 h DXM serum concentration. Two other horses in this study received both phenylbutazone and flunixin, one of which had a 4.1 pg/ml serum DXM concentration at 72 h, and the other which had both its 48 and 72 h DXM concentrations censored. No conclusions can be drawn about any possible interactions, because of the small numbers of horses that received these concomitant medications.

The second experiment was performed to investigate the possibility that DXM plasma concentrations may result from recycled urinary DXM. Table 4 shows that five of six horses offered DXM containing urine had detectable plasma DXM during at least once time point in a 24 h period. The highest plasma concentration was 4.4 pg/ml, 24 hours after exposure to the contaminated hay, and 12 h after the previous plasma sampling. While this plasma DXM concentration was below the ARCI threshold of 5 pg/ml DXM, the peak plasma concentration could have occurred at any time in the preceding ~11 h (Soma *et al.*, 2013). Using the PK parameters from Soma *et al.* (2013), the peak plasma concentration in this horse (Horse A) could have been anywhere from 4.4 pg/ml to 24 pg/ml. The DXM concentration in the urine used to contaminate hay for our experimental horses was only 17 ng/ml, and we used only 100 ml of this urine. It is clear from these data that the plasma concentrations found in our field study horses, as well as the study horses of Soma *et al.* (2005, 2013) and Symonds *et al.* (2019) could have resulted from recycled DXM from urine.

The 48 h serum threshold concentrations were 6.61 pg/ml using the 95/95 tolerance method, 6.77 pg/ml using GCM at $P=0.05$ and 12.4 pg/ml using GCM at $P=0.01$. The 95/95 tolerance methodology provides a 95% confidence that 95% of the values will fall below the calculated threshold, where the underlying data are normally distributed (Owen, 1968). A number of assumptions are made in the calculation of 95/95 tolerance, which are previously discussed (Owen, 1968). The GCM is a probability inequality that guarantees that no more than a certain fraction of values will exceed a specified distance from the mean. The chief difference between the threshold determinations is that GCM is a

distribution free method that does not require normally distributed data, only a unimodal distribution (Savage, 1961). In the case of both methods of calculating thresholds at a risk of 1 in 20, 2 of the 74 member 48 h data set (not including the extreme outlier), or ~3% exceeded the threshold, exemplifying the point that both methods carry an approximately 5% risk of a violation where the withdrawal is carefully followed. Because the underlying assumptions for both methods of threshold determination are met, the threshold for both 95/95 tolerance and GCM ($P=0.05$) are comparable in the case of this DXM study. Where the underlying assumptions are not met, the most appropriate threshold determination methodology should be chosen (Machin *et al.*, 2019). When the GCM methodology using a 1 in 100 risk of a positive ($P=0.01$) is employed instead, the threshold becomes 12.40 pg/ml, which exceeds all values in the dataset, other than the extreme outlier.

The statistical methods employed for threshold determination in this study for the 48 h dataset require several assumptions. The RROS methodology of estimating values for censored data requires that the DXM concentrations below the LOQ follow the same distribution as the measured concentrations. Therefore, the statistical moments calculated from those RROS estimates must be considered in light of these assumptions, where 72% of the data cannot be measured. However, as discussed in the preceding paragraph, this assumption is likely to be accurate, because when thresholds are calculated by two different methods that carry a 5% risk of a violation, about 3% of the population exceeds the threshold.

Statistics to determine threshold values cannot be applied where censored data exceed 80% of the dataset (Helsel, 2012). Therefore, we could not calculate thresholds by any methodology for our 72 h dataset. However, all but the two extreme outliers fell below the recommended 5 pg/ml threshold in place in racing in the United States. Since both horses with outlier DXM concentrations at 72 h were below 5 pg/ml at 48 h, inadvertent subcutaneous drug administration or idiosyncratic metabolism are unlikely sources of these high DXM serum concentrations. Nonetheless, it is clear that a small number of horses may be at risk of exceeding the threshold at 72 h despite falling below the threshold at 48 h.

The pharmacokinetic studies performed by Soma *et al.* (2005, 2013) were performed exclusively in research horses, which received no concurrent medications, and may or may not have been exercised at a level comparable to race training. Pharmacokinetics differ not only between Standardbreds and Thoroughbreds, but also between animals that are fit or sedentary (Khazaenia *et al.*, 2000). No previous studies have been performed to differentiate drug disposition among horses that are in Thoroughbred, Standardbred or Quarter Horse race training.

This study provides several important guidelines for veterinarians using DXM for the treatment of IAD, other inflammatory or allergic conditions of equine athletes that perform in a regulated environment. First, this study demonstrates that there is no accumulation of DXM used at a dose of 20 mg IV as the sodium phosphate solution once daily, so the withdrawal is the same after the final dose of intravenous DXM sodium phosphate, whether it is used for a single day or multiple days. Second, this study supports the 5 pg/ml threshold at 72 h with 95% confidence, because the two outlier concentrations out of 74 horses represents a 2.7% risk of a positive test. In this field study of seventy-four horses sampled at 2 time points, there were 3 extreme outliers, representing a risk of violation of approximately 1 in 50. In the Soma *et al.* studies (2005, 2013), and the Symonds study (2019), there were also extreme outliers, despite a controlled environment. Therefore, withdrawal guidelines should include warnings that there is a risk of violating the threshold, even if the withdrawal time is strictly followed. Further studies are warranted to investigate the possible sources of these outlier high serum concentrations which have presented in a number of studies.

Conflict of interest

The authors declare no conflict of interest.

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