

1 Edited by CAS October 9, 2013 AJVR No. 13-05-0138R

2 Effect of body position on intra-abdominal pressures and abdominal perfusion pressures
3 measured at three sites in horses anesthetized with short-term total intravenous anesthesia

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8 Received May 9, 2013.

9 Accepted October 7, 2013.

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14 Presented in part in abstract form at the 2012 American College of Veterinary Internal Medicine
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17

18 **Objective**—To assess effects of body position on direct measurements of intra-abdominal
19 pressure (IAP) and abdominal perfusion pressure (APP) in horses anesthetized with total
20 intravenous anesthesia (TIVA).

21 **Animals**—9 healthy adult horses.

22 **Procedures**—Instrumentation in unsedated standing horses involved insertion of an arterial
23 catheter for blood pressure measurements and 3 intraperitoneal cannulas (left flank, right flank
24 and ventral abdomen) for IAP measurements. Baseline values were measured for heart rate,
25 respiratory rate, systolic arterial pressure, mean arterial blood pressure (MAP), diastolic arterial
26 blood pressure, and IAP. Horses were medicated with xylazine and pressures were measured
27 again. Anesthesia was induced with ketamine-diazepam and maintained with a ketamine-
28 guaifenesin infusion. Horses were positioned twice into left lateral recumbency, right lateral
29 recumbency, or dorsal recumbency. Hemodynamic pressures and accessible abdominal pressures
30 were measured for each recumbency position. The APP was calculated as $MAP - IAP$.
31 Differences in IAP, MAP, APP and sedation (standing horses) or body position (anesthetized
32 horses) were compared by repeated-measures ANOVA or paired *t* tests.

33 **Results**—Baseline hemodynamic and intra-abdominal pressures were not different after xylazine
34 administration. Ventral abdomen IAP and MAP were lower for horses in dorsal recumbency than
35 in right or left lateral recumbency. Ventral abdomen APP remained unchanged. For lateral
36 recumbencies, flank IAP was lower and APP was higher than pressure measurements at the same
37 sites for dorsal recumbency.

38 **Conclusions and Clinical Relevance**—Body position affected IAP and APP in healthy
39 anesthetized horses. These effects should be considered when developing IAP acquisition

40 methods for use in horses with abdominal disease. (*Am J Vet Res* 2014;75:xxx–xxx)

41 **Approximately 250 words** **6**

42

43 **ABBREVIATIONS** **7**

44 APP Abdominal perfusion pressure

45 CI Confidence interval

46 DAP Diastolic arterial blood pressure

47 IAH Intra-abdominal hypertension

48 IAP Intra-abdominal pressure

49 MAP Mean arterial blood pressure

50 SAP Systolic arterial blood pressure

51 TIVA Total intravenous anesthesia

52

53 Accurate measurement of IAP in human medicine has become an increasingly important

54 monitoring tool for critically ill patients. Intra-abdominal hypertension in humans is defined as

55 sustained IAP \geq 12 mm Hg.¹ Complications associated with protracted IAH include reduced

56 microcirculatory blood flow to viscera, development of organ dysfunction, and possible organ

57 failure. Abdominal perfusion pressure is a calculated index of abdominal blood flow (MAP –

58 IAP) and has been proposed as an accurate predictor of visceral perfusion and an end point for
59 resuscitation.^{2,3} Abdominal compartment syndrome describes the natural progression of pressure-
60 induced organ changes that develop if IAH is not recognized and treated in a timely manner.¹ In
61 critically ill humans, IAH is a risk factor for organ failure and fatality.⁴ Mortality rates associated
62 with abdominal compartment syndrome in critically ill adults and children range from 50% to
63 60%.⁵

64
65 Identification of IAH and abdominal compartment syndrome in any species requires accurate
66 measurement of IAP. In humans, indirectly measured intravesicular pressures are considered
67 accurate and can be used for serial acquisition of IAP.¹ Many variables, including body position,
68 directly affect abdominal pressures.⁶⁻¹¹ Therefore, it is advised that IAP measurements be
69 obtained with the person in supine recumbency.^{1,12} Body position can also affect directly
70 measured IAP in dogs anesthetized with short-term TIVA.¹³ Standardized methods of IAP
71 measurement in horses and reference range values are lacking. Previous reports^{14,15} in horses
72 have indicated that direct intraperitoneal cannulation is the most accurate method, with reduced
73 variation of IAP, compared with results for indirect intravesicular or intragastric techniques.
74 Furthermore, IAP can differ depending on the location in the abdomen of standing horses.¹⁶ To
75 our knowledge, the effect of body position on directly measured abdominal pressure has not been
76 evaluated and must be examined if IAP is to be a potentially useful monitoring tool in critically
77 ill equine patients.

78

79 Manipulation of body position in horses requires anesthetic immobilization. Sedation and TIVA
80 are frequently used in horses to facilitate minor procedures performed by practitioners in both
81 field and hospital settings. In humans with IAH, administration of sedatives is a nonsurgical
82 method for lowering IAP through abdominal wall relaxation.^{17,18} Investigators in a recent study¹⁶
83 found that right flank IAP in clinically normal unsedated standing horses was similar to that
84 reported in another study¹⁴ in which horses were sedated by administration of detomidine
85 administered 30 minutes before IAP measurement. This indicated that sedative administration
86 may have negligible effects on IAP in clinically normal horses. Although the effects of sedation
87 and various anesthetic regimens on systemic hemodynamic variables have been characterized in
88 healthy horses,¹⁹⁻²² IAP and APP in sedated horses and horses anesthetized with TIVA and
89 placed in various body positions are currently unknown.

90

91 The purpose of the study reported here was to determine whether body position influences IAP
92 and APP in healthy horses under clinically relevant conditions. Results of the study may be
93 useful in subsequently developing standardized methods for acquisition of IAP measurements
94 that can be applied in both field and hospital settings to horses with abdominal disease. We
95 hypothesized that direct measurement of abdominal pressures (IAP and APP) at 3 sites in healthy
96 standing horses will not be affected by sedation achieved by IV administration of xylazine but
97 that abdominal pressures at the 3 sites will be affected by body position of horses anesthetized
98 with TIVA. Specifically, an increase in IAP and decrease in APP is anticipated whenever the
99 respective site of measurement is closer to the ground, compared with values when the
100 measurement site is farther from the ground.

101

102 **Materials and Methods**

103 **Animals**—Nine university-owned adult (> 1 year old) female horses of various breeds were
104 included in the study. Horses were considered free of abdominal disease on the basis that no
105 abnormalities were detected during physical examination and per rectal examination, there was
106 no history of colic or abdominal surgery during the preceding 6 months, and no abnormalities
107 were detected during transabdominal or transthoracic ultrasonography. The animal use protocol
108 and all experimental procedures were approved by the institutional animal care and use
109 committee of The Ohio State University as well as the institutional clinical trials office and
110 hospital clinical research advisory committee. All procedures complied with the National
111 Institutes of Health standards for the ethical treatment of animals.

112

113 **Instrumentation**—In an attempt to standardize the amount of material in the gastrointestinal
114 tract among the study population, 4 L of mineral oil were administered via nasogastric intubation
115 and food was withheld for 24 hours before instrumentation. Water was withheld for 6 hours
116 before instrumentation and any experiments. Horses were housed in temperature-controlled
117 (25°C) indoor stalls. All horses were weighed and assigned a body condition score²³ immediately
118 prior to instrumentation.

119

120 Nonsedated horses were placed in a standing position in stocks for instrumentation. Sites for
121 catheter insertion were clipped and aseptically prepared by use of chlorhexidine gluconate and

122 isopropyl alcohol. Local anesthesia was achieved by SC injection **8** of mepivacaine,^a**9** and a 14-
123 gauge, 5.25-inch catheter^b was placed in the left jugular vein and secured with 2-0 polypropylene
124 suture^c to provide venous access in all horses. Local anesthesia was achieved by SC injection
125 **8** of mepivacaine, and a 20-gauge, 1.25-inch catheter^d was placed in a transverse facial artery or
126 facial artery for direct measurement of arterial blood pressures (SAP, MAP, and SAP). The
127 arterial catheter was secured to skin with cyanoacrylate glue and connected to an 84-cm-long
128 extension set that was filled with heparinized saline (0.9% NaCl) solution; the extension set was
129 attached to the horse's halter.

130

131 Intra-abdominal cannulation was performed as a modified abdominocentesis at 3 locations in the
132 abdomen (right flank, left flank, and ventrum). The flank sites for cannulation were midway
133 between the center of the tuber ischii and the cranial eminence of the greater tubercle of the
134 humerus at a point 12 cm caudal to the last rib.¹⁴ The ventral abdominal cannula site was
135 identified by visual inspection of the standing horse, and the shortest ground-to-abdomen
136 distance was determined with a tape measure (typically on the linea alba at a point 10 to 15 cm
137 caudal to the xiphoid process). A 5 X 5-cm area at each cannulation site was clipped and
138 aseptically prepared by use of chlorhexidine gluconate and isopropyl alcohol. Mepivacaine (8
139 mL) was locally infiltrated, and a No.15 scalpel blade was used to make a stab incision into the
140 skin and subcutis.

141

142 **Measurement of IAP**—Direct measurement of IAP was obtained via a 3-way stopcock attached

143 to a sterile 10-gauge 10-cm metal teat cannula filled with heparinized saline solution. Sterile
144 water-based lubricant was applied at the site of cannula insertion to prevent entry of air and
145 development of pneumoperitoneum. The cannula was inserted through the body wall and
146 peritoneum into the intra-abdominal space and held in position by an assistant as reported
147 elsewhere.^{14,16} Placement through the peritoneum was confirmed by obtaining peritoneal fluid or
148 by a lack of resistance to flushing with sterile saline solution (< 2 mL). The cannula was
149 connected to an 84-cm-long extension set filled with heparinized saline solution. The other end
150 of the extension set was attached to a pressure transducer^e and electronic manometer^f for data
151 collection as reported elsewhere.^{16,24} The transducer was checked against other transducers prior
152 to experimentation and the manometer was calibrated annually. For IAP measurement, the
153 transducer was set to zero at the level of cannula insertion into the abdomen, and the 3-way
154 stopcock then was turned to the open position at the cannula end.^{14-16,24} Each pressure was
155 recorded in triplicate at the end of expiration as is standard in human medicine.¹

156

157 Arterial blood pressures were obtained by connecting the arterial catheter to a pressure
158 transducer. The transducer for blood pressure was set to zero at the level of the point of the
159 shoulder (estimated level of the right atrium) for standing horses and horses in dorsal
160 recumbency, and at the level of the sternum for horses in right or left lateral recumbency. Direct
161 measurements of arterial pressure were recorded simultaneously with IAP measurements. The
162 transducer for arterial pressures was reset to zero after anesthetized horses were repositioned. All
163 pressure recording systems involved polypropylene tubing **10** filled with heparinized saline
164 solution, which was visually assessed for the presence of air bubbles prior to connection to a

165 horse. When air bubbles were detected, the tubing was with heparinized saline solution until
166 bubbles were no longer evident. All pressure recording systems were assessed for dampening
167 with the square-wave flush test and visual inspection of the pressure waveform for
168 underdampening or overdampening, whereby no appreciable effect of dampening was
169 observed.²⁵

170

171 Hemodynamic and intra-abdominal variables assessed during the experiments included heart rate
172 determined by ECG, respiratory rate, SAP, MAP, DAP, and IAP at each of the 3 abdominal
173 locations (left flank, right flank and ventral). The APP was calculated for each site of IAP
174 measurement by use of the following equation: $APP_x = MAP_x - IAP_x$, where x is the abdominal
175 location (ie, left flank, right flank, or ventral).

176

177 **Experimental procedures**—After instrumentation was completed, horses were allowed to stand
178 in the stocks uninterrupted for 10 minutes. Baseline hemodynamic and intra-abdominal pressures
179 then were obtained. The intraperitoneal cannulas were removed. Horses were moved to an
180 induction stall and medicated with xylazine hydrochloride^g (1.1 mg/kg, IV). The intraperitoneal
181 cannulas were replaced. Five minutes after xylazine administration, horses were assessed to
182 determine adequate sedation (lowered head and minimal response to external stimuli) and all
183 variables were measured. Intraperitoneal cannulas were again removed. Anesthesia was induced
184 by IV administration of ketamine hydrochloride^h (2.2 mg/kg) and diazepamⁱ (0.075 mg/kg).
185 Anesthesia was maintained with a continuous infusion of ketamine (2 mg/mL of solution) and
186 guaifenesin guacolate in 5% dextrose solution (50 mg of guaifenesin/mL of solution). Rate of the

187 ketamine-guaifenesin infusion was adjusted to maintain a light plane of anesthesia (minimal to
188 no nystagmus and no spontaneous movements of the limbs, head, or neck) to facilitate animal
189 positioning. Horses were intubated with a 26-mm cuffed orotracheal tube and allowed to
190 spontaneously breathe room air (fraction of inspired oxygen, 0.21).

191

192 The order of recumbency positions for each horse was determined with a randomization
193 procedure (a web-based random number generator^j). Each horse was placed in each recumbency
194 position twice during an experiment. If the same position was designated consecutively, then the
195 horse was placed in a different recumbency (but without obtaining measurements) before being
196 returned to the designated recumbency. For example, when a specific lateral recumbency
197 position was designated consecutively, then the default different recumbency was the opposite
198 lateral side (ie, if right lateral recumbency were consecutively assigned, the measurements were
199 obtained with the horse in right lateral recumbency, the horse then was placed in left lateral
200 recumbency but no measurements were obtained, and the horse then was repositioned into right
201 lateral recumbency and measurements were obtained). When dorsal recumbency was
202 consecutively designated, then the different recumbency was the opposite lateral recumbency to
203 that which had been most recently used (ie, if a horse had most recently been positioned in right
204 lateral recumbency before consecutive designations of dorsal recumbency, then measurements
205 were obtained with the horse in dorsal recumbency, the horse was positioned in left lateral
206 recumbency but no measurements were obtained, and the horse then was repositioned in dorsal
207 recumbency and measurements were obtained). Horses in dorsal recumbency were manually
208 supported by 4 assistants; one was positioned at each limb. **11**

209

210 Measurements were obtained, cannulas were removed, horses were repositioned, and cannulas
211 then were reinserted. Horses were manually rolled from one body position to the next. A 2-
212 minute period was allowed after manipulation of a horse into a new body position prior to data
213 collection. Ventral IAP was obtained for all 3 recumbency positions. The IAP at the left flank
214 was obtained when horses were in right lateral and dorsal recumbency, whereas IAP at the right
215 flank was obtained when horses were in left lateral and dorsal recumbency.

216

217 After data collection was completed, intraperitoneal cannulas were removed and the skin
218 incisions were stapled. Horses remained anesthetized and were immediately enrolled into another
219 unrelated study. Horses were euthanized after completion of that unrelated study.

220

221 **Statistical analysis**—All hemodynamic and intra-abdominal pressure variables were measured
222 in triplicate for each site of measurement and each recumbency. The mean of the 3 values was
223 used for statistical analysis. Variability of IAP and MAP obtained for the duplicated
224 recumbencies (ie, twice each for dorsal, left lateral, and right lateral recumbency) was calculated
225 by use of the following equation:

226
$$\text{Variability} = \frac{([Y1 \text{ sample mean}] - [Y2 \text{ sample mean}])}{([\{Y1 \text{ sample mean} + Y2$$

227
$$\text{sample}\}/2])} \times 100$$

228 where Y1 and Y2 are the first and second measurements obtained for a body position (left
229 lateral, right lateral, or dorsal recumbency). An arbitrary clinical cutoff value of $\leq 12\%$ was
230 considered acceptable variability for both IAP and MAP.

231

232 Statistical testing was performed with commercial software programs.^{k,l} Data were assessed for
233 normality with the Shapiro-Wilk and D'Agostino & Pearson omnibus normality tests and found
234 to have a Gaussian distribution. Data were reported as mean \pm SD or 95% CI unless otherwise
235 stipulated. Effect of xylazine on all variables (baseline value versus value after xylazine
236 medication) was assessed with paired *t* tests. Intra-abdominal pressure, MAP, and APP were
237 obtained for 2 cannula locations when horses were in lateral recumbency and 3 locations when
238 horses were in dorsal recumbency, which resulted in an unbalanced incomplete block design.
239 The effect of body position on IAP, MAP, and APP for the ventral cannula site was assessed
240 with a repeated-measures 1-way ANOVA with Holm-Sidak post hoc testing. The effect of body
241 position on IAP, MAP, and APP for the flank cannula sites was assessed with paired *t* tests.
242 Significance was set at values of $P < 0.05$ for all analyses.

243

244 **Results**

245 **Horses**—The median age of the 9 horses was 21 years (interquartile range, 13 to 25 years).
246 There were 5 geldings and 4 mares; none of the mares was pregnant. Breeds included were
247 Quarter Horse (n = 5), Thoroughbred (2), Standardbred (1), and Rocky Mountain Horse (1).
248 Median body condition score (scale of 1 to 9) was 5 (range, 3 to 8). Median body weight of the
249 horses was 485 kg (interquartile range, 439 to 528 kg).

250

251 All horses completed the study. Mean \pm SD duration of anesthesia for data collection was $27 \pm$
252 3.4 minutes. Median rate for infusion of the ketamine-guaifenesin solution was 2.06 mL/kg/h
253 (95% CI, **12** 1.89 to 2.28 mL/kg/h). Median rate for administration of ketamine and guaifenesin
254 guacolate was 4.12 mg/kg/h (95% CI, **12** 3.79 to 2.28 mg/kg/h) and 103.09 mg/kg/h (95% CI, **12**
255 94.70 to 113.90 mg/kg/h), respectively. One horse received an additional 200 mg of ketamine
256 IV during the experiment.

257

258 **Effect of xylazine on hemodynamic and intra-abdominal pressures in horses**—At 5 minutes
259 after injection of xylazine, all horses were clinically sedated (to a level adequate for anesthetic
260 induction). The head was lowered and the horse was minimally responsive to external stimuli.
261 Sedation did not have a significant effect on any measured variable (**Table 1**).

262

263 **Effect of body position on abdominal pressures in anesthetized horses**—There was $\leq 10\%$
264 variability in mean IAP values obtained for each cannula site for each duplicate recumbency (eg,
265 between the first and second positioning in left lateral recumbency). There was $\leq 12\%$ variability
266 in the MAP obtained for each duplicate recumbency.

267

268 Ventral IAP was significantly ($P < 0.001$) lower when horses were in dorsal recumbency,
269 compared with values obtained when horses were in left or right lateral recumbency. Left flank

270 IAP and right flank IAP were significantly ($P < 0.001$) higher when horses were in dorsal
271 recumbency, compared with values obtained when horses were in lateral recumbency. Directly
272 measured MAP was significantly ($P < 0.001$) lower when horses were positioned in dorsal
273 recumbency, compared with MAP when horses were positioned in left or right lateral
274 recumbency. Ventral APP did not differ significantly ($P = 0.23$) among the 3 recumbency
275 positions. The APP was significantly lower for the left flank ($P < 0.001$) and right flank ($P =$
276 0.002) when horses were positioned in dorsal recumbency, compared with values measured
277 when horses were in lateral recumbency (Table 2).

278

279 Discussion

280 Direct techniques for measurement of IAP in horses are repeatable and are currently considered
281 to be the most accurate method of acquisition.^{14,15} Values for IAP obtained from the flank and
282 ventral aspect of the abdomen of standing horses in the present study were comparable to those
283 reported by use of the same anatomic landmarks for identification in healthy horses.¹⁴⁻¹⁶

284

285 In the present study, we found that hemodynamic and intra-abdominal pressures were unchanged
286 in healthy standing horses in response to IV administration of xylazine. A reduction in heart rate
287 and respiratory rate as well as a brief period of hypertension (followed by hypotension) has been
288 reported after administration of α_2 -receptor agonists,^{21,26} which is in contrast to the results
289 obtained for the study reported here in which no change in heart rate, respiratory rate, or blood
290 pressure was observed after horses were sedated. Although the horses in the present study were
291 conditioned to their environment and did not display excitement or anxious behavior prior to

292 xylazine administration, we speculate that the process of instrumentation (without sedation) as
293 well as movement to the induction stall may have acted as a stimulus to the sympathetic nervous
294 system that affected the hemodynamic response to xylazine.

295

296 To our knowledge, the effect of IV administration of xylazine on IAP has not been evaluated in
297 horses. Sedative administration is used as one type of medical management for IAH in humans,
298 which results in an increase in abdominal compliance and subsequent reduction in IAP.^{17,18}
299 Therefore, results of the present study are in contrast with those reported in the human literature;
300 however, in contrast to critically ill humans whereby patients are in a supine position during IAP
301 measurement, the study population for the present study comprised standing healthy horses
302 without preexisting abdominal distension. We speculate that the effects of xylazine and other
303 sedatives may differ in recumbent horses with abdominal disease and potential IAH; however,
304 additional studies are needed to make that determination. Species differences in abdominal
305 conformation, effect of the amount of material in the gastrointestinal tract, body condition score,
306 and IAP acquisition method may also have accounted for these contrasting findings. Analysis of
307 the limited data for horses suggests that there is an effect of body weight on directly measured
308 IAP,¹⁴ and given the wide range of body condition scores of horses in the present study, future
309 studies designed to specifically investigate the effect of body weight and body condition on IAP
310 are warranted.

311

312 The common use of sedatives in equine practice suggests that strategic sedation might offer a
313 practical method for medical management of IAH in a clinical setting, assuming sedation is

314 found to be efficacious for reducing IAP in horses with abdominal disease. Further investigation
315 is required in this area. The use of sedatives in humans reportedly reduces variations in IAP
316 measurements by reducing fluctuations in abdominal wall compliance.²⁷ We did not observe
317 substantial variation in the horses of the present study; however, we were investigating the
318 effects of xylazine in clinically normal standing horses. Abdominal pain in horses and
319 consequent distension of the abdomen might contribute to wider variations in IAP. Evaluations
320 of horses with IAH to investigate IAP variation would be a logical step to determine optimum
321 conditions for the development of standardized methods for the acquisition of IAP. We did not
322 find a significant decrease in IAP after IV administration of a 1.1 mg/kg dose of xylazine to
323 healthy horses, despite the observed clinical effects of sedation; however, further studies are
324 required before definitive conclusions can be made in this area.

325

326 In the present study, IAP measured at 3 locations in the abdomen of horses resulted in
327 differences between the values obtained for each site, in response to changes in body position.
328 IAP was increased at the ventral site when horses were positioned in lateral recumbencies and
329 decreased when horses were positioned in dorsal recumbency. The IAP obtained from the flank
330 positions of cannulation was high when horses were positioned in dorsal recumbency but lower
331 when horses were in lateral recumbencies. In humans, the effect of body position on IAP is
332 clearly established.¹ Studies in humans⁶⁻¹¹ reveal that lateral recumbency and various
333 semirecumbent positions (supine positioning with head-of-bed elevation) result in significant
334 increases in IAP, compared with values obtained for patients strictly in a supine position. Values
335 for IAP in humans reportedly are highest when a patient assumes an upright position.²⁸ The
336 results from these humans studies were all obtained by use of the human consensus intravesicular

337 acquisition method for IAP.¹ In contrast to results for humans,^{29,30} indirect methods of IAP
338 measurement in horses are poorly correlated with direct intraperitoneal cannulation.^{14,15} Body
339 position also affects IAP in dogs.¹³ In that study,¹³ investigators found that direct measurement
340 of IAP with an intraperitoneal catheter filled with saline solution yielded higher values with dogs
341 in upright, lateral, and prone positions, compared with results for dogs in a supine position. To
342 our knowledge, the effect of body position on IAP in horses has not been investigated
343 previously, but the findings for the present study are in concordance with those of other species.
344 Body position is a variable that must be considered when developing a standardized, reliable, and
345 repeatable method for abdominal pressure acquisition.

346

347 It has been postulated that the human abdominal cavity behaves as a homogenous hydraulic fluid
348 system in accordance with the dynamics of Pascal's law.^{1,9,31} Pascal's law states that the pressure
349 exerted anywhere in a confined noncompressible fluid is transmitted equally in all directions
350 throughout the fluid, such that the pressure ratio (initial difference) remains the same. Such a
351 hypothesis means that IAP should remain constant, regardless of body position; however, several
352 human studies^{1,6-11} as well as the present study in horses provide contradictory findings. In a
353 study¹³ in dogs, investigators found that 3 factors (gravity, visceral shear deformation, and
354 visceral compression) are involved in the determination of IAP. Forces of gravity and visceral
355 shear are considered negligible for human patients lying in a supine position. In such
356 circumstances, visceral compression will correlate directly with intravesicular pressure and IAP.⁷
357 Therefore, recumbency in a supine position allows the abdomen to behave as a hydraulic system.
358 In other body positions, however, shape-unstable viscera (ie, the bladder) become deformed and
359 change abdominal pressure dynamics away from a simple hydrostatic system.⁸

360

361 Exact reasons why changes in body position alter IAP remain incompletely defined, but several
362 interacting forces may explain the heterogeneous behavior of the abdomen when patients are
363 placed in different recumbencies. Manipulating recumbency positions in horses changes the
364 intra-abdominal cannula height relative to the bulk of abdominal mass and will increase or
365 decrease gravitational and shear forces accordingly for each of the fixed cannula sites used for
366 this study protocol. The intraperitoneal cannulas were presumed to be in direct contact with
367 gastrointestinal viscera, and we propose that the main effect of body position on IAP was related
368 to movement of viscera and forces of gravity as recumbency was manipulated.

369

370 The effects of body position on cardiopulmonary variables have been reported in horses for
371 various conditions, including prolonged anesthesia,^{32,33} inhalation anesthesia,³²⁻³⁶ and positive-
372 pressure ventilation.^{36,37} However, the effect of changing recumbency positions on directly
373 measured IAP and calculated APP have not been reported. The MAP is one of the variables
374 required to calculate APP, and we found in the present study that MAP was lower when horses
375 were in dorsal recumbency, compared with MAP when horses were in lateral recumbency. The
376 same effect on blood pressure has been reported in halothane-anesthetized ponies with
377 comparable manipulation of position.³⁶ Another study³⁸ performed in dogs anesthetized by IV
378 administration of anesthetic agents (a proportion of which were spontaneously breathing room
379 air) revealed that blood pressure and systemic vascular resistance were significantly lower when
380 dogs were placed in a supine position, compared with results when dogs were in lateral
381 recumbency. We speculate that the change in MAP identified by this study³⁸ in dogs is the direct

382 result of changes in body position and may be attributable to the weight of abdominal organs
383 compressing the caudal vena cava. Such compression could lead to a decrease in venous return to
384 the heart, which would be followed by a reduction in cardiac output and a subsequent decrease in
385 arterial blood pressure. Similar physiologic processes have been described for pregnant women
386 in the context of human aortocaval compression syndrome.³⁹ It is important that these changes in
387 MAP with alterations in body position are considered because MAP is integral for the calculation
388 and interpretation of APP.

389

390 Calculated APP is a concept analogous to the widely accepted notion of cerebral perfusion
391 pressure (ie, the difference between MAP and intracranial pressure).¹² However, the boundaries
392 of the skull are nonpliable, whereas the abdominal wall is typically compliant. Therefore, it is
393 possible that APP has a wider range of reference values and more variation in healthy subjects
394 than does cranial perfusion pressure. Given the physiologic processes of pressure and high
395 compliancy of the equine abdomen, we speculate that APP ranges in horses might be more
396 expansive than other species. **13** This point of discussion warrants further investigation and
397 should include comparative evaluation among species, in addition to consideration of anatomic
398 features such as diaphragmatic shape.

399

400 In the present study, we found that when the ventral cannula site was used as a point of
401 reference, APP remained unchanged regardless of body position. This is because ventral IAP and
402 MAP increased or decreased by comparable magnitudes in response to manipulation of
403 recumbency position. When the left or right flank cannula sites were used as reference points for

404 abdominal pressures, IAP and MAP increased or decreased in opposite directions in response to
405 a change in body position; therefore, values of calculated perfusion pressure were different for
406 lateral versus dorsal recumbency. It has been proposed that APP can be used as a predictor of
407 fatality and may serve as an optimal endpoint for fluid resuscitation in human critical care
408 medicine; however, its usefulness has not yet been fully determined.² It is important to remember
409 that APP is a calculated estimation of visceral perfusion. Because location of IAP acquisition and
410 body position both appear to be variables affecting calculated values in horses, interpretation and
411 clinical importance of APP is currently unknown. A standardized method for IAP acquisition is
412 required, which should then be followed by comparison of the resultant APP value with results
413 for other quantifiable methods of visceral perfusion. Investigating IAP measurements in horses
414 with clinical abdominal disease by use of the same 3 locations that were used in the present study
415 might prove useful for determining the utility of obtaining a pressure from the ventral location
416 during routine abdominocentesis in order to identify horses with IAH. Moreover, use of the
417 ventral location to monitor the response to medical or surgical treatments would aid in the
418 understanding of abdominal pressure dynamics in horses with colic.

419

420 The data obtained in the present study are applicable to systemically healthy horses that may be
421 subjected to short-term anesthesia with TIVA. The impact of inhalation anesthesia, positive-
422 pressure ventilation, and longer durations of anesthesia on hemodynamic and abdominal
423 perfusion indices remains to be determined, especially for those patients considered to be at high
424 risk for IAH (ie, surgical colic) positioned in dorsal recumbency. These data may serve as
425 preliminary reference values for future studies conducted to investigate the role of APP in the
426 context of IAH, colic, and other abdominal disturbances of horses.

427

428 Limitations of the study included the use of TIVA to enable us to manipulate body position.
429 However, it would not have been possible to obtain these data without anesthetizing the horses.
430 The anesthetic drugs chosen for use were designed to minimize cardiovascular instability. All
431 horses were orotracheally intubated to minimize any increase in airway resistance associated
432 with anesthesia that could lead to increases in IAP as a result of high inspiratory airway pressure.
433 The effects of skeletal muscle relaxants used in the study may have impacted abdominal
434 compliance (and caused decreases in IAP); however, almost all anesthetic protocols include a
435 component to induce muscle relaxation as a typically desired effect.

436

437 The anesthesia used in the present study provides only initial information on IAP and APP, and
438 the impact of different anesthetic protocols on abdominal pressure variables requires further
439 investigation. The effect of anesthesia over time on hemodynamic and intra-abdominal pressures
440 may also have introduced bias to the results we obtained; however, patient positioning was
441 randomized such that the order of recumbencies was not predetermined or consistent for every
442 horse. The variation in pressure measurements was good (< 10 %) but not perfect. The 2-minute
443 period between changes in recumbency positions and subsequent measurements may have been
444 insufficient to achieve a steady-state IAP. Further studies with different intervals would be
445 needed to determine whether it was sufficient.

446

447 We did not measure variables in horses positioned in sternal recumbency. Arterial blood
448 pressures in horses positioned in sternal recumbency during isoflurane-induced anesthesia are

449 similar to those when horses are positioned in dorsal and lateral recumbency.³⁴ We intended to
450 determine changes in measured pressures in response to clinically relevant body positions. It is
451 uncommon for a horse to be positioned in sternal recumbency for a procedure. Dorsal-to-lateral
452 recumbency and lateral recumbency alone are extremely common positions and were the focus
453 of the experiments.

454

455 Another area for consideration includes the fact that IAP was measured at end expiration, rather
456 than end inspiration, as has been the case for some of the evaluations of IAP in horses.¹⁴ The
457 human literature reports that IAP values can be increased at end inspiration and with positive-
458 pressure ventilation⁴⁰ (because of a corresponding increase in thoracic cavity pressure, which
459 results in transfer of pressure to the abdomen); thus, the current consensus for critically ill people
460 is to measure IAP at end expiration to attain the most accurate and repeatable values.¹ To our
461 knowledge, there is no information currently available in the veterinary literature to support the
462 use of measurements obtained during either phase of the respiratory cycle; therefore, we chose a
463 protocol in concordance with the consensus for humans. It is possible that the optimal time
464 within the respiratory cycle for measurement of IAP differs between horses and humans because
465 of species differences in the respiratory pattern; however, further investigation is required in this
466 area. The authors propose that this variable is more important in horses with abdominal
467 distension and in those with an increased rate or depth of breathing than in healthy horses.

468

469 The fact that intra-abdominal cannulas were removed and reinserted during the transition
470 between body positions is another point of discussion for the present study. This method was

471 required during movement of horses and manipulation of body position to prevent damage to the
472 abdominal wall or organs and breakage of the cannulas. Walking and changes in body position
473 resulted in substantial discordant movement of the penetrated soft tissue; thus, reinsertion of the
474 cannulas ensured instruments were correctly positioned and that there was direct communication
475 with the peritoneal cavity. The depth of the abdominal wall and overlying skin resulted in closure
476 of the penetrating tract once cannulas were removed. The small areas of hair that were clipped at
477 the cannula insertion sites during patient preparation allowed for easy identification of these
478 locations for cannula reinsertion. Repeatability of IAP measurements was good, which indicated
479 that previous cannulation of the abdomen did not affect values obtained during the second
480 measurement. Thus, the authors believe that the method was appropriate and that cannula
481 manipulation did not result in bias during data acquisition.

482

483 For the present study, we did not detect significant changes in IAP, MAP, or APP in response to
484 sedation achieved by administration of xylazine in healthy standing horses. We found that
485 manipulation of body position in horses anesthetized with TIVA significantly changed IAP and
486 APP obtained at various sites throughout the abdomen in addition to lowering MAP for horses
487 positioned in dorsal recumbency. Standardized protocols for the measurement of IAP in horses
488 have not yet been developed and are required before use in a clinical setting. The repeatability of
489 measurements and simplicity of the modified-abdominocentesis technique for the study reported
490 here lends itself to use in further investigations. Such investigations should include the
491 evaluation of abdominal pressures in horses with abdominal disease (eg, colic) to determine the
492 prevalence of IAH and the efficacy of therapeutic interventions.

493

494 **Footnotes** **14**

- 495 a. Carbocaine, Hospira, Lake Forest, Ill.
- 496 b. Angiocath, Becton Dickinson, Franklin Lakes, NJ.
- 497 c. Surgipro, Covidien, Mansfield, Mass.
- 498 d. Surflo, Terumo Medical, Somerset, NJ.
- 499 e. Truwave pressure transducer, model PX36N, Edward Lifesciences, Irvine, Calif.
- 500 f. Datascope Passport, Maquet GmbH & Co KG, Rastatt, Germany.
- 501 g. Anased, Akorn Inc, Decatur, Ill.
- 502 h. Ketaset, Fort Dodge Animal Health, Fort Dodge, Iowa.
- 503 i. Diazepam, Hospira, Lake Forest, Ill.
- 504 j. RANDOM.ORG **Title of webpage**. Available at: www.random.org. Accessed March, 21-
- 505 31, 2011 **month, day, year**. **15**
- 506 k. Prism, version 5.0, GraphPad Software Inc, San Diego, Calif.
- 507 l. Excel, Microsoft Corp, Mountain View, Calif.

508

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609

610

611 Table 1—Mean (95% CI) values for hemodynamic and abdominal pressure variables before and
 612 after administration of a sedative preanesthetic medication in 9 healthy standing horses. **18**

Variable	Baseline	After preanesthetic medication	<i>P</i> value*
IAP (mm Hg)			
Left flank	−3.2 (−5.2 to −3.2)	−5.1 (−7.9 to −2.3)	0.12
Right flank	−5.1 (−7.5 to −2.8)	−5.1 (−7.0 to −3.3)	0.51
Ventral	25 (22 to 27)	25 (22 to 28)	0.54
SAP (mm Hg)	134 (126 to 141)	140 (120 to 160)	0.56
MAP (mm Hg)	103 (92 to 113)	100 (86 to 113)	0.71
DAP (mm Hg)	80 (72 to 88)	81 (63 to 99)	0.92
Heart rate (beats/min)	35 (30 to 40)	36 (33 to 39)	0.56
Respiratory rate (breaths/min)	14 (13 to 16)	14 (13 to 15)	0.53
APP (mm Hg)†			
Left flank	106 (95 to 117)	103 (88 to 118)	0.88
Right flank	108 (97 to 119)	103 (89 to 118)	0.71
Ventral	78 (67 to 89)	73 (59 to 87)	0.65

613
 614 After instrumentation was completed, horses were allowed to stand in the stocks uninterrupted
 615 for 10 minutes; baseline hemodynamic and intra-abdominal pressures then were obtained. Horses
 616 were moved to an induction stall and medicated with xylazine hydrochloride (1.1 mg/kg, IV). All
 617 variables were measured again 5 minutes after xylazine administration.

618 *Considered significant at $P < 0.05$. †Calculated as MAP – IAP.

619 Table 2—Mean \pm SD (95% CI) values of directly measured IAP, MAP, and calculated APP for 9
 620 healthy anesthetized horses placed in various positions. **18**

Cannula location	Left lateral recumbency			Right lateral recumbency			Dorsal recumbency		
	IAP	MAP	APP	IAP	MAP	APP	IAP	MAP	APP
Left flank	NA	NA	NA	-5.4 ± 3.3 (-8.0 to -2.9)	95.0 ± 19.3 (80.1 to 110.0)	100.0 ± 21.3 (84.1 to 116.9)	$15.3 \pm 4.6^*$ (11.8 to 18.8)	69.0 ± 14.2 (58.2 to 79.8)	$54.0 \pm 17.3^*$ (40.5 to 67.0)
Right flank	-8.1 ± 2.1 (-9.7 to -6.5)	82.5 ± 19.2 (67.8 to 97.3)	91.0 ± 18.8 (76.2 to 105.1)	NA	NA	NA	$15.3 \pm 5.6^*$ (11.1 to 19.5)	70.4 ± 11.9 (61.3 to 79.6)	$55.0 \pm 14.6^*$ (43.9 to 66.4)
Ventral	11.3 ± 3.3 (8.8 to 13.9)	83.6 ± 14.9 (72.3 to 95.1)	72.0 ± 15.7 (60.2 to 84.4)	13.0 ± 7.1 (7.5 to 18.5)	93.4 ± 17.0 (80.3 to 106.5)	80.0 ± 20.9 (64.3 to 96.5)	$-6.7 \pm 2.4^*$ (-8.5 to -4.9)	$68.8 \pm 11.8^*$ (59.8 to 77.9)	76.0 ± 13.4 (65.2 to 85.2)

632
 633 *Within a row, value differs significantly ($P < 0.05$) from the corresponding value for other body
 634 positions.