

1     **Successful postnatal care of a premature orphan foal delivered by**  
2                                 **Caesarean section**

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26  
27    **Abstract**

28 A 9-year-old Shagya Arabian pregnant mare showing signs of acute colic, underwent  
29 exploratory laparotomy at our clinic. Owing to the discovery of an inoperable  
30 leiomyoma in the abdominal cavity, the animal was euthanized. The 305-day-old  
31 fetus, which showed signs of prematurity was removed by Caesarean section and  
32 resuscitated. Clinicopathologic examination revealed a low neutrophil:lymphocyte  
33 ratio. Radiographic evaluation of the carpus and tarsus was performed and showed  
34 grade 3 ossification according to the Adams-Poulos Grading System.

35 The immature gastrointestinal tract was unable to digest enteral feeding; this led to  
36 the development of enterocolitis and septicemia. Thrombophlebitis developed at the  
37 site of the long-acting intravenous catheter and methicillin-resistant *Staphylococcus*  
38 *aureus* was isolated from the exudate. The guttural pouches were empyematous.  
39 Follow-up radiographic examinations indicated improvement in bone maturation.

40 The foal was discharged in a good state of health 35 days postnatum, and apart from  
41 an easily managed respiratory tract infection, no further problems were reported.

42

43 **Keywords:** prematurity, equine neonate, septic foal, postnatal care

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47 **1. Introduction**

48 Equine prematurity is difficult to define because of differences between breeds and  
49 other influencing factors (the fetal gender, etc.). Nevertheless, according to current  
50 knowledge premature foals born after less than 280 days of gestation almost never  
51 survive and even foals delivered at 300 days tend not to be viable. The survival rate  
52 after 320 days of gestation is more satisfactory [10]. When predicting chances of  
53 survival, it is also important to consider whether the delivery occurred spontaneously  
54 or the foal was born as a result of Caesarean section owing to a sudden illness or injury  
55 to the mare. Premature foals born spontaneously as a result of chronic stress have  
56 better survival rates than foals delivered by Caesarean section at the same gestational  
57 age [24,25,26]. While the development of the fetal pituitary adrenal axis is dependent  
58 on cortisol, during the majority of gestation, the fetus is not exposed to high levels of  
59 cortisol. Fetal cortisol production increases significantly during the final 24–48 h  
60 before parturition, presumably as a result of increased adrenal 17 $\alpha$ -hydroxylase  
61 activity. This period of cortisol exposure is essential for the final maturation of the  
62 fetal pituitary adrenal axis and the respiratory system [34]. Consequently, if  
63 spontaneous delivery is brought on by chronic stress (e.g., because of placentitis, as  
64 the result of twinning, or congenital problems associated with the placenta or the foal)  
65 and stress-associated–cortisol release occurs in the mare, the foal may still undergo  
66 the necessary cortisol-related maturation to prepare it for the extrauterine environment  
67 [10,33].

68 Even when born at term, the rearing of a newborn foal poses a special challenge and  
69 if it is compounded by a mare's death, it may further reduce the chances of the foal's  
70 survival [14].

71

## 72 **2. Case history and clinical findings**

73 A Shagya Arabian mare was presented for acute colic at the Clinic for Large Animals  
74 (presently Equine Department and Clinic) in Üllő, Hungary. The 9-year-old mare  
75 weighed 445 kg and was in advanced pregnancy (gestational length: 305 days).

76 During exploratory laparotomy, the premature foal was delivered by Caesarean  
77 section. The mare was euthanized during

78 surgery because of the presence of an  
79 inoperable tumor in the abdominal  
80 cavity (Fig. 1). The weight of the tumor  
81 was 40 kg, and histological examination  
82 revealed it to be a leiomyoma.



*Fig. 1. Inoperable leiomyoma in the abdominal cavity of the dam*

83 The newborn filly showed signs of

84 prematurity: underweight (30 kg) (the average weight of a newborn foal is 50 kg) [26]

85 with pliant ears, soft lips, entropion, weak musculature, and poor suckling reflex. The  
86 foal was unconscious and unresponsive, and respiration was irregular and infrequent.

87 The peripheral pulse was almost absent and heart rate was low (40 bpm) [10].

88 Mydriasis was observed in both eyes and capillary refill time was prolonged.

89 Clinicopathological examination revealed a low neutrophil:lymphocyte ratio and the

90 white blood cell count was also low. The blood lactate and creatinine levels were

91 elevated, and the packed cell volume (PCV) was also high. Although there was no

92 immediate possibility to measure IgG level, the serum globulin level was low. The

93 foal had mild hypoglycemia, which persisted and occasionally worsened over the first

94 few days post-partum. Urine output was low.

95 Radiographic evaluation of the carpus and tarsus was  
96 performed to assess the degree of ossification. The  
97 degree of incomplete ossification in the cuboidal  
98 bones was assessed as being grade 3 according to the  
99 Adams–Poulos grading system [1] (Fig. 2).



Fig. 2. Radiographic appearance of the premature carpus

### 101 3. Treatments

102 On the basis of these results and observations, resuscitation was performed immediately after birth [11]. The high creatinine level  
103 can reflect not only hypovolaemia in newborn foals, but also can be caused by other  
104 abnormal conditions, e.g. by in utero placental dysfunction. Increased PCV is also not  
105 solely indicative of hypovolemia [8], but based on the history, laboratory findings and  
106 clinical signs, emergency fluid resuscitation was initiated. A bolus of half of the shock  
107 dose rate (60 mL/kg) of balanced electrolyte solution (Lactated Ringer’s Solution,  
108 “Baxter” 1000 mL)<sup>1</sup> was administered rapidly, and the foal was reassessed: Basic  
109 clinical parameters were within normal limits (HR, 86 bpm; RR, 46/min; and T,  
110 37.4°C [under the infrared light]), the pulse was strong, and mentation was improved.  
111 To provide for maintenance requirements and ongoing losses, fluid therapy was  
112 continued at a rate of 5 mL/kg·h<sup>-1</sup>. For short-term nutritional support 30 mL of 40%  
113 Glucose (Glucose 40% inf. 500 mL)<sup>2</sup> and 30 mL of a vitamin and electrolyte solution  
114 (Duphalyte<sup>®</sup> injection A.U.V.)<sup>3</sup> were added to each liter of crystalloid infusion [8]. As  
115 part of our intensive care strategy antimicrobial therapy with ceftriaxon (Rocephin,  
116 25 mg/kg i.v. *b.i.d.*)<sup>4</sup> was started immediately after birth.

118 Because of the absence of suckling reflex, a permanent nasogastric tube (Salem Sump  
119 tube GS4018)<sup>6</sup> was placed and 1 L of stored colostrum was administered in 4 portions  
120 over the first 8 h. Frozen colostrum was used from mares at the clinic. Despite not  
121 knowing the IgG level of the foal, 3 L of plasma supplementation was also  
122 administered ( $20 \text{ mL/kg}\cdot\text{h}^{-1}$ ) to support immune function. After the supplementation,  
123 plasma total protein and the globulin level were 63.4 g/L and 35.3 g/L, respectively,  
124 suggesting that an adequate humoral immune status had been achieved [23]. Over the  
125 same period, the blood glucose level decreased in spite of the recommended amounts  
126 of good quality colostrum being fed. (Table 1.)

127 The foal did not pass meconium in the first 36 h, and subsequent treatment with an  
128 acetylcysteine retention enema was successful. Because of the risk of corneal  
129 ulceration from the congenital entropion [3], the in-turning eyelids were corrected  
130 surgically.

131 Biochemistry performed 2 days after delivery showed a low neutrophil count, marked  
132 hypoglycemia, arterial hypoxaemia, and metabolic acidosis. Clinical examination at  
133 this point showed the foal to be hyperthermic, depressed, and diarrheic. According to  
134 the Foal Sepsis Score Sheet, the foal had a total of 23 points, which is predictive of  
135 sepsis in 93% of the time [4,7,13]. Thoracic and abdominal radiography showed  
136 increased pulmonary interstitial pattern, and distended intestines which were also  
137 visible on ultrasound. Observation of comet-tail echoes and inflammatory areas on  
138 ultrasonographic examination of the lungs confirmed the presence of pneumonia.  
139 When enterocolitis developed, total parenteral nutrition with appropriate amino acid,  
140 lipid, and vitamin content was considered but not administered owing to financial  
141 limitations. During the period of diarrhea nutrition was provided both parenterally and

142 peroral. The maintenance fluid requirement ( $100 \text{ mL/kg}\cdot\text{day}^{-1}$ ) was supplemented  
143 with Glucose (Glucose 40% inf. 500 mL)<sup>2</sup> and a vitamin and electrolyte solution  
144 (Duphalyte<sup>®</sup> injection A.U.V.)<sup>3</sup>. The reduced oral nutrition consisted of overly diluted  
145 milk replacer (Salvana Fohlenmilch)<sup>5</sup> and the concentration was increased as the foal  
146 accepted more intensive peroral feeding. The foal was initially fed every hour and  
147 then gradually less frequently as larger volumes could be given [10]. In spite of regular  
148 feeding and parenteral energy supplementation, the foal had extremely high  
149 hypertriglyceridemia ( $9.2 \text{ mmol/L}$ ). In addition to the glucose infusion, recombinant  
150 human regular insulin (Actrapid Penfill)<sup>6</sup> was administered by continuous rate  
151 infusion (dose rates ranged between  $0.0016$  and  $0.018 \text{ IU/kg}\cdot\text{h}^{-1}$ ) [22,29]. Blood  
152 glucose level was regularly checked. When the foal was able to utilize a sufficient  
153 amount of orally administered feed (as seen by increased blood glucose and reduced  
154 triglyceride levels), intravenous fluid and energy supplementation were discontinued.  
155 Non-steroidal anti-inflammatory drugs (Neoprogen 10% inj.,  $2.2 \text{ mg/kg i.v. s.i.d.}$ )<sup>7</sup> [2]  
156 were administered during the period of diarrhea and, in order to safeguard against  
157 possible side effects, renal function was monitored and proton pump inhibitors were  
158 given (Omeprazole,  $4 \text{ mg/kg p.o. s.i.d.}$ )<sup>8</sup>. With the exception of the first 2 days after  
159 delivery, the levels of creatinine and BUN were within the normal ranges. Despite a  
160 negative blood culture the spectrum of antimicrobial therapy was extended with  
161 penicillin (Tardomyocel comp. III susp.,  $3 \text{ mL/50 kg i.m. s.i.d.}$ )<sup>9</sup>, metronidazole  
162 (Klion,  $25 \text{ mg/kg p.o. b.i.d.}$ )<sup>10</sup> and amikacin (Likacin inj.,  $25 \text{ mg/kg i.v. s.i.d.}$ )<sup>11</sup> [9]. A  
163 probiotic preparation (Pro-Paste for horses,  $2 \text{ mL/foal, p.o. b.i.d.}$ )<sup>12</sup> was also  
164 administered.

165 Treatments were supplemented with kaolin bolus (Bolus adstringens, 1 tablet/10 kg  
 166 p.o. *b.i.d.*)<sup>13</sup>, Psyllium (Sand Clear, 1 scoop/day, p.o. *b.i.d.*)<sup>14</sup>, activated carbon (Carbo  
 167 Activatus 500 g)<sup>15</sup> and paraffin oil (Mol White Oil M 46)<sup>16</sup> and diarrhea was solved.  
 168 In addition to the antimicrobial therapy, the lower airways were supported with  
 169 Vitamin C (Acidum ascorbicum)<sup>15</sup> and bromhexine (Bisolvon, 0.25 g/10 kg p.o.  
 170 *b.i.d.*)<sup>17</sup>.

171

172 **Table 1. Clinicopathological data of the foal pre-, during and after treatment**

| AGE (day)                             | 0      | 1     | 1 (afternoon) | 2     | 5     | 8     | 11    | 15    | 18    | 21    | 25    | 32    |
|---------------------------------------|--------|-------|---------------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| WBC (x10 <sup>9</sup> /L)             | 4.28   | 3.07  | 1.77          | 3.72  | 33.3  | 10.00 | 24.6  | 28.3  | 8.97  | 7.79  | 9.55  | 7.90  |
| N:L                                   | 0.28   | 0.78  | 0.84          | 6.76  | 9.56  | 5.7   | 27.58 | 12.78 | 3.47  | 1.49  | 1.72  | 1.73  |
| Albumin (g/L)                         | 27.83  |       | 28.71         |       |       | 25.98 | 23.23 | 23.86 | 22.37 | 23.59 | 25.61 | 23.90 |
| Total protein (g/L)                   | 42.00  |       | 63.40         |       |       | 50.30 | 46.30 | 49.70 | 52.30 | 57.20 | 48.60 | 49.80 |
| Globulin (g/L)                        | 11.36  |       | 35.30         |       |       | 24.32 | 20.36 | 22.05 | 29.93 | 32.61 | 19.93 | 25.90 |
| Glucose (mmol/L)                      | 3.12   | 2.30  | 1.59          | 2.21  |       | 8.43  | 10.43 | 8.06  | 6.46  | 7.70  | 4.61  | 7.53  |
| Tryglicerid (mmol)                    | 3.28   | 4.40  | 7.76          | 9.20  |       | 2.45  | 0.59  | 1.27  | 0.48  | 0.68  | 0.91  | 0.82  |
| Lactate (mmol/L)                      | 13.55  |       | 3.34          | 10.84 |       | 3.37  | 2.29  | 1.44  | 2.31  | 1.65  | 1.39  | 2.06  |
| BUN (mmol/L)                          | 11.10  |       | 10.60         |       |       | 3.80  | 1.90  |       |       | 1.50  | 2.30  | 1.60  |
| Creatinin (µmol/L)                    | 225.00 |       | 198.20        |       |       | 72.00 | 74.20 |       |       | 79.00 | 80.00 | 81.00 |
| pH                                    |        | 7.26  |               | 7.27  | 7.27  | 7.30  |       |       |       |       |       | 7.37  |
| p <sub>a</sub> CO <sub>2</sub> (mmHg) |        | 52.80 |               | 51.10 | 45.30 | 44.60 |       |       |       |       |       | 44.20 |
| p <sub>a</sub> O <sub>2</sub> (mmHg)  |        | 60.00 |               | 64.00 | 77.00 | 79.90 |       |       |       |       |       | 94.00 |
| HCO <sup>3-</sup> (mmol/L)            |        | 22.70 |               | 23.10 | 24.70 | 25.50 |       |       |       |       |       | 31.10 |

173

174

175 The foal was recumbent and depressed during the first week and was first able to stand  
 176 without assistance 10 days after delivery. The follow-up radiographic assessment of  
 177 the limbs and cuboidal bones showed at that time improvement in terms of  
 178 ossification. Exercise was gradually increased as the foal became stronger [16].  
 179 Strengthening of the soft tissues of the limbs was supported by the use of bandaging  
 180 and splinting on all 4 legs. This limb support was removed for part of each day (about



181 12 h/day) to allow for loading of the tendons and the periarticular soft-tissue  
182 structures.

183 Recurrent mild impaction, which resulted in abdominal distention, was presumed to  
184 be caused by the immaturity of the gastrointestinal tract. It was treated using  
185 prokinetics such as metoclopramide (Cerucal inj., 0.05 mg/kg i.m.)<sup>18</sup> and neostigmine  
186 (Konstigmin inj., 0.0044 mg/kg s.c.)<sup>19</sup> [31], and also seemed to be alleviated when the  
187 foal was encouraged to move.

188 Despite using a long-acting intravenous catheter (Equivet HiFlow LongTerm IV  
189 Catheter 14 G × 3.5 cm, Langeskov, Denmark)<sup>20</sup> and antibiotics, thrombophlebitis  
190 developed at 2 weeks of age. The foal became recumbent, hyperthermic, and  
191 developed severe tachycardia. The left jugular vein was warm and appeared painful,  
192 and a small amount of purulent discharge was present at the site of the intravenous  
193 catheter from which methicillin resistant *Staphylococcus aureus* was isolated. Both  
194 guttural pouches were empyematous. Over this period, in order to overcome the  
195 multiple bacterial resistances antibiotic treatment was changed to a combination of  
196 erythromycin (Meromycin, 37.5 mg/kg p.o. *b.i.d.*)<sup>21</sup> and rifampicin (Rifamed, 5-10  
197 mg/kg p.o. *s.i.d.*)<sup>22</sup> [22]. To prevent transmission to other animals the foal was isolated  
198 and was handled by only a small number of people. Precautionary measures were used  
199 to minimize pathogen spread and avoid zoonotic transmission [27]. Aggressive wound  
200 cleaning was maintained until the infection resolved and the wound healed. The  
201 guttural pouches were flushed with sterile fluids endoscopically twice a day until there  
202 was no discharge.

203 One week later, the results of physical examination and clinicopathological findings  
204 were within normal limits. The antimicrobial and supportive treatments were

205 continued for a further 7 days after which only probiotic therapy was continued. The  
206 nasogastric tube was removed at this time and the foal was trained to drink from a  
207 bucket.

208

#### 209 **4. Outcome**

210 Thirty-five days after delivery, the foal was discharged in a good state of health to the  
211 Babolna National Stud farm. The stud farm veterinary surgeon reported that the foal,  
212 aged 1.5 years at the time of writing, had suffered a respiratory tract infection 3  
213 months after discharge which was treated successfully, and has since displayed no  
214 other health issues.

215

#### 216 **5. Discussion**

217 In order to assess the level of prematurity and the presence and degree of failure of  
218 passive transfer (FPT) an adrenocorticotrophic hormone (ACTH) stimulation test, and  
219 cortisol- and IgG level determination should be performed shortly after birth. In  
220 premature foals, FPT cannot only be the result of inadequate colostrum intake, but can  
221 also reflect gastrointestinal absorption abnormalities. That a combination of these  
222 factors was involved in the development of FPT in this case is supported by the lack  
223 of increase in blood glucose after peroral colostrum feeding. Plasma globulin status  
224 was improved when, in addition to the recommended volume of colostrum, the foal  
225 also received plasma supplementation. In this case, use of the ACTH stimulation test  
226 to determine the degree of prematurity was unnecessary, as this was apparent from  
227 the physical appearance as well as knowledge of the gestational age of the foal.

228 Caesarean section has been associated with neonate immaturity because of the  
229 absence of the pre-parturient endogenous steroid release. It is debated as to whether  
230 the use of corticosteroids in premature foals is beneficial as these immunosuppressant  
231 agents have been shown to have a harmful effect on the immune system of equine  
232 neonates [20]. However, the immunosuppressive activity is predominantly restricted  
233 to cell-mediated immunity, with only a minimal inhibitory effect on humoral  
234 immunity. The real immunologic and clinical effects of a hydrocortisone therapy are  
235 still not clear [17] and this treatment was, therefore, not used in this case. It is possible  
236 that the presence of the abdominal tumor was a source of chronic stress for the mare  
237 and the fetus which could have resulted in untimely endogenous steroid release. The  
238 Adams–Poulos Grade 3 ossification of the cuboidal bones was higher than could be  
239 expected at 305 days of gestation. However, the intestinal dysfunction, and the  
240 external appearance of the foal were suggestive of prematurity, the foal was probably  
241 more developed than a normal 305-day-old fetus. However it was still considered to  
242 be premature.

243 The prematurity-related intestinal dysfunction and absorption deficiency that  
244 represented the major problems during the first 2 weeks of life were compounded by  
245 the development of septicaemia and enterocolitis. It is not clear whether the FPT led  
246 to sepsis and subsequent enterocolitis or whether enterocolitis, caused by enteral  
247 feeding of the immature gastrointestinal tract, led to the development of septicaemia  
248 as a result of the excessive translocation of bacteria across the gut. Besides these  
249 eventualities hypoxemia, and consequent tissue hypoxia, gastroduodenal ulceration,  
250 lactose intolerance, infections (e.g. *Clostridium difficile*) or other factors may have  
251 played also a role in the apparent diarrhoea. Some of these possibilities were unlikely,

252 since the foal had normal faeces several times between the period of meconium  
253 impaction and diarrhoea. Clostridium infection was also not expected, because the  
254 occurrence of *Cl. difficile* caused enterocolitis is very uncommon in Hungary. The  
255 presence of gastroduodenal ulceration could not be ruled out, but Omeprazole was  
256 administered.

257 Conflicting opinions arose in connection with the administration of Omeprazole.  
258 Javsicas and Sanches (2010) [21] investigated the effect of Omeprazole (4 mg/kg p.o.)  
259 on intragastric pH in critically ill neonates. The intragastric pH was significantly  
260 higher in the post treatment period compared to the pretreatment period. Furr et al.  
261 (2012) [12] evaluated the influence of anti-ulcer medications on the development of  
262 undifferentiated or infection caused diarrhea in compromised neonatal foals. The  
263 importance of gastric acidity in protecting against bacterial translocation was also  
264 investigated. In the examined 1102 foals the occurrence of diarrhea was significantly  
265 higher with the use of any anti-ulcer medication, including sucralfate treatment.  
266 However the study was designed retrospectively and the influence of the different  
267 hospitals, clinicians and the original disease of the foals may not have been completely  
268 eliminated [12]. In this premature case, the used nonsteroidal anti-inflammatory drug  
269 (ketoprofen) was a nonselective cyclo-oxygenase (COX) inhibitor; therefore the  
270 synthesis of prostaglandins was also inhibited. Besides this and the possible perinatal  
271 transient tissue hypoxia, pH dependent gastric ulceration also had to be taken into  
272 account.

273 The use of nonsteroidal anti-inflammatory drugs in critically ill neonates raises many  
274 issues and it is a frequently studied topic also in equine and in human medicine  
275 [5,28,30]. Morris et al. [28] discussed the importance of the prostaglandin system in

276 the healthy development of human neonates. In this study the role of selective cyclo-  
277 oxygenase type 2 (COX-2) inhibitors arose especially in connection with the  
278 gastrointestinal adverse effects. However it was also emphasized, that the possibility  
279 of investigations in this topic is still far from being exhausted [28]. Several years later,  
280 Raidal et al. [30] tested the use of meloxicam (0.6 mg/kg; p.o.) in foals less than 6  
281 weeks of age. No threatening side effects were revealed even at higher dose (1.8  
282 mg/kg; p.o.) of administration. Nevertheless only healthy foals were used in the  
283 experiment, hence the extrapolation of these results to compromised neonates requires  
284 special caution [30]. These data [30] were not available at the time of the admission  
285 of the presented foal, but the main advantages of the relatively COX-2 selective  
286 meloxicam were already known. Using a nonselective COX inhibitor (ketoprofen)  
287 was a necessary decision in this case, since selective COX-2 inhibitor was not  
288 available in Hungary.

289 Glucose infusions, given for the apparently impaired gastrointestinal absorption, are  
290 probably only suitable for 12–24 h of nutritional support when not combined with  
291 lipid and protein supplementation [10]. The preterm gut is very sensitive to enteral  
292 feeding which may either promote gut adaptation and health, or induce gut  
293 dysfunction, bacterial overgrowth and inflammation. The interaction between gut  
294 bacteria and host tissue in a newborn compromised intestine has been studied in many  
295 species, especially in infants and piglets. Enteral feeding induced bacterial  
296 colonization stimulates structural, functional and immunological maturation of the  
297 intestinal tissue [19]. Nevertheless, enteral feeding has been associated with  
298 necrotizing enterocolitis (NEC) in preterm piglets [32]. The effect of minimal enteral  
299 nutrition combined with parenteral nutrition was evaluated by Cilieborg et al. [6] in

300 piglets. Minimal enteral colostrum feeding improved intestinal structure, function,  
301 and NEC resistance. For this reasons, despite the risk of excessive bacterial  
302 translocation across the immature gut, oral feed intake was given at a reduced rate and  
303 not completely withdrawn.

304 The presence of sepsis, combined with the impaired absorption resulted in an  
305 inadequate nutritional status, and extremely high triglyceride levels.  
306 Hypertriglyceridemia has been associated with many complications in septic human  
307 patients, including immunosuppression, increased production of inflammatory  
308 mediators (such as interleukins), lipid intolerance, allergic reactions,  
309 thrombocytopenia, cholestasis, increased parameters in liver function tests and fat  
310 embolism, the latter occurring especially in premature neonates [15,18]. The  
311 complications seen in this case were not attributed to the high triglyceride level, but  
312 probably also delayed the healing process.

313 It is important to note that handling of a premature foal is almost the most important  
314 factor in the later clinical history. For this reason a good knowledge of the availability  
315 of equipment and medication is necessary to allow rapid decision making. Delay in  
316 the treatment of the compromised neonate is an important risk factor contributing to  
317 poor survival rate [25]. For example, without financial constraints, it may have been  
318 possible to avoid the development of hypertriglyceridemia in this case with adequate  
319 parenteral nutrition. Accordingly, the planning of the intensive care of a premature,  
320 orphan foal should start with consideration of both the economic benefits and  
321 responsibilities arising from each clinical decision as considerable financial resources  
322 may be necessary to achieve a favorable outcome. With respect to the equine neonate,

323 welfare of the patient, the end use of the animal, and the emotional and financial  
324 considerations of the owner might be the overriding these concerns.

325

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332

#### 333 **5. Manufacturers**

- 334 1. Baxter Hungary Ltd., Budapest, Hungary
- 335 2. TEVA Pharmaceutical cPlc, Debrecen, Hungary
- 336 3. Fort Dodge Veterinaria, S.A., Vall de Bianya, Spain
- 337 4. Roche, Budaörs, Hungary
- 338 5. Salvana Tiernahrung GmbH, Elmshorn, Germany
- 339 6. Novo Nordisc A/S, Chartres, France
- 340 7. Kela N.V., Hoogstraten, Belgium
- 341 8. Ratiopharm Hungaria, Budapest, Hungary
- 342 9. Bayer Hungaria, Budapest, Hungary
- 343 10. Richter Gedeon Plc, Budapest, Hungary
- 344 11. Lisapharma s.p.a., Erba, Italy
- 345 12. Protexin veterinary, Probiotics International Ltd, Somerset, United  
346 Kingdom,

- 347 13. EGIS, Budapest, Hungary  
348 14. Farnam Companies, Osborn, USA  
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350 16. MOL-LUB Ltd., Almásfüzitő, Hungary  
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