

Beef Session

Moderators: *Christine Navarre, Trent Fox*

Fetal programming: implications for beef cattle production

R.N. Funston, PhD; A.F. Summers

University of Nebraska, West Central Research and Extension Center, North Platte, NE 69101

Abstract

The beef cattle industry relies on the use of high-forage diets to develop replacement females, maintain the cow herd, and sustain stocker operations. Forage quantity and quality fluctuate with season and environmental conditions. Depending on class and physiological state of the animal, a forage diet may not always meet nutritional requirements, resulting in reduced average daily gain or body weight (BW) loss if supplemental nutrients are not provided. It is important to understand the consequences of such BW loss and the economics of providing supplementation to the beef production system. Periods of limited or insufficient nutrient availability can be followed by periods of compensatory BW gain once dietary conditions improve. This may have less impact on breeding animals, provided reproductive efficiency is not compromised, where actual BW is not as important as it is in animals destined for the feedlot. A rapidly evolving body of literature is also demonstrating that nutritional status of cows during pregnancy can affect subsequent offspring development and production characteristics later in life. The concept of fetal programming is that maternal stimuli during critical periods of fetal development have long-term implications for offspring. Depending on timing, magnitude, and duration of nutrient limitation or supplementation, it is possible that early measures in life, such as calf birth BW, may be unaffected, whereas measures later in life, such as weaning BW, carcass characteristics, and reproductive traits, may be influenced. This body of research provides compelling evidence of a fetal programming response to maternal nutrition in beef cattle. Future competitiveness of the US beef industry will continue to be dependent on the use of high-forage diets to meet the majority of nutrient requirements. Consequences of nutrient restriction or supplementation must be considered not only on individual animal performance, but also the developing fetus and its subsequent performance throughout life.

Key words: cattle, beef, fetal programming

Résumé

L'industrie de l'élevage bovin se fie aux diètes de fourrage pour le développement des génisses de remplacement et le maintien du troupeau de vaches et pour soutenir le parc d'élevage. La quantité et la qualité du fourrage changent avec la saison et les conditions environnementales. Selon la classe et l'état physiologique de l'animal, il est possible qu'une diète de fourrage ne réponde pas bien aux besoins nutritifs, ce qui peut entraîner une réduction du gain de poids quotidien ou du poids si des éléments nutritifs complémentaires ne sont pas disponibles. Il est important pour l'industrie de l'élevage bovin de bien comprendre les conséquences de telles pertes de poids et la rentabilité associée à l'utilisation d'éléments nutritifs complémentaires. Des périodes de disponibilité réduite ou insuffisante d'éléments nutritifs peuvent être suivies de périodes de gain de poids compensatoires lorsque les conditions d'alimentation s'améliorent. Ceci peut avoir moins d'impact sur les animaux en reproduction, en autant que la reproduction ne soit pas affectée, car contrairement aux animaux destinés à l'engraissement le poids est en fait est moins important. Un nombre d'études grandissant démontre aussi que le statut nutritionnel de la vache en gestation peut affecter le développement du veau et son profil de production plus tard dans la vie. Selon le concept de la programmation fœtale, les stimuli maternels durant les périodes critiques du développement fœtal ont des implications à long terme pour le jeune. Selon le moment, l'amplitude et la durée de la limitation ou de la supplémentation alimentaire, il est possible que des mesures prises tôt dans la vie, comme le poids du veau à la naissance, ne soient pas affectées alors que des mesures prises plus tardivement, comme le poids au sevrage, les caractéristiques de la carcasse et de la reproduction, le soient. Cet ensemble de travaux supporte bien l'idée que la nutrition maternelle chez les bovins de boucherie a un impact par le biais de la programmation fœtale. La compétitivité future de l'industrie américaine du bœuf restera liée à l'utilisation de diètes de fourrage pour répondre à la plupart des besoins nutritifs. Les conséquences de la restriction ou de la supplémentation alimentaire doivent être prises en compte non seulement pour la vache gestante elle-même mais aussi pour le fœtus en développement et sa performance à vie.

Introduction

The concept of fetal programming, also known as developmental programming, was first hypothesized using human epidemiological data in which environmental stimulus in utero resulted in altered long-term development, growth, and disease susceptibility in children from undernourished mothers during the Dutch famine.¹ Recently, literature regarding fetal programming effects in domesticated livestock has been reviewed.^{8,9}

Many factors influence livestock nutrient requirements including breed, season, and physiological function.³¹ Fetal programming responses can result from a negative nutrient environment, which can be caused by 1) breeding of young dams who compete for nutrients with rapidly growing fetal systems; 2) increased incidences of multiple fetuses or large litters; 3) selection for increased milk production, which competes for nutrients with increased energy demand from fetal and placental growth; or 4) breeding of livestock during high environmental temperatures and pregnancy occurring during periods of poor pasture conditions.^{40,54} Studies have reported instances of compromised maternal nutrition during gestation resulting in increased neonatal mortality, intestinal and respiratory dysfunction, metabolic disorders, decreased postnatal growth rates, and reduced meat quality.⁵⁴ Proper management of cow nutrition during gestation can improve progeny performance and health.

Placental Development

The bovine placenta attaches along the uterine wall at locations known as caruncles. These knob-like structures along the uterine luminal surface serve as attachment sites for the chorionic villi of the fetal placenta known as cotyledons. The caruncle-cotyledonary unit, also known as a placentome, serves as the primary functional area of physiological exchange between mother and fetus.⁹ Establishment of functional uteroplacental and fetal circulation is one of the earliest events during embryonic and placental development^{32,35} allowing for transportation of all respiratory gas, nutrient, and waste exchanges between the maternal and fetal systems.^{37,38} The efficiency of transport is related to uteroplacental blood flow,³⁷ and although placental growth slows during the last half of gestation, blood flow to the placenta increases 3- to 4-fold from mid to late gestation to support the exponential rate of fetal growth.^{7,27,36,37,42}

Due to the importance of placental development on fetal nutrient transfer, studies have been conducted to determine how maternal nutrition can influence placental development, or placental programming. Zhu et al⁵⁶ reported nutrient restriction of beef cows from day 30 to 125 of gestation resulted in reduced ($P < 0.05$) caruncular and cotyledonary weights from nutrient-restricted cows compared to control, unrestricted cows, and fetal weights from nutrient-restricted

cows tended ($P = 0.12$) to be reduced compared to control cows. Following realimentation during day 125 to 250 of gestation, caruncular and cotyledonary weights were still reduced for nutrient-restricted cows; however, fetal weight was not different. Vonnahme et al,⁵³ using the same cows, reported increased placental angiogenesis as well as angiogenic factor mRNA abundance in the caruncular and cotyledonary tissues at the end of the nutrient restriction period. It was hypothesized the lack of significant fetal weight differences in regard to maternal nutrient restriction may have resulted from the increase in cotyledonary arteriolar density allowing for adequate nutrient transfer.^{53,56}

To measure capillary vascularity of the cotyledon, 4 measurements are collected: capillary area density (**CAD**), a flow-related measure; capillary number density (**CND**), an angiogenesis-related measure; capillary surface density (**CSD**), a nutrient-exchange measure; and area per capillary (**APC**), a capillary density per cross section of muscle area. Vonnahme et al⁵³ reported no difference in these 4 measures from day 30 to day 125 of gestation; however, from day 125 to 250, there were significant differences in CAD, CND, and CSD when comparing control and nutrient-restricted cows, suggesting capillary area, numbers, and surface densities had been hindered upon realimentation. Nutrient restriction from day 30 to 125 of gestation in cows did not alter the vasculature of the bovine placenta; however, placental function must have been compromised due to reduced fetal weights.

Fetal Organ Development

Robinson et al⁴¹ reported 75% of ruminant fetal growth occurs during the last 2 months of gestation. Due to the minimal nutrient requirement during early gestation, inadequate nutrition during this time was thought to have little significance. However, during the early phase of fetal development critical events for normal conceptus development occur, including differentiation, vascularization, fetal organogenesis, and as previously mentioned, placental development.⁹

Fetal organ formation occurs simultaneously to placental development, with limb development occurring as early as day 25 of gestation. Following limb development is a sequential development of other organs including the pancreas, liver, adrenal glands, lungs, thyroid, spleen, brain, thymus, and kidneys.¹⁹ Testicle development begins by day 45 in male calves, and ovarian development begins in female calves by day 50 to 60. Another important event in female gonadal development occurs approximately day 80 of gestation, when oocyte nests break down to form primordial follicles.³⁰ These follicles represent the oocyte supply available to a female after puberty known as the ovarian reserve, which can influence her reproductive lifespan.¹⁸ A review by Caton et al³ lists fetal programming examples in livestock models of individual organs including heart,¹⁷ lung,¹³ pancreas,^{22,23} kidney,¹² placenta,³⁹ perirenal fat,^{25,26} and small intestine.¹⁴

Fetal Muscle Development

The fetal stage is also crucial for skeletal muscle development because muscle fiber numbers do not increase after birth.^{47,55} Skeletal muscle is a lower priority in nutrient partitioning compared with the brain, heart, or other organ systems,^{2,4} making it particularly vulnerable to nutrient deficiency. Thus, a decrease in nutrient availability to the dam during gestation can result in a reduced number of muscle fibers through fetal programming, reducing muscle mass and impacting animal performance. Both muscle fiber number and intramuscular adipocytes, which provide the sites for intramuscular fat accumulation or marbling formation, are influenced during fetal development.^{6,50}

Figure 1 depicts the effects of maternal nutrition on fetal skeletal muscle formation and control points in which maternal nutrition has been shown to impact fetal muscle development. Although primary muscle fibers of the bovine fetus begin forming within the first 2 months of gestation,⁴³ very limited numbers of muscle fibers are formed at this stage; thus, maternal nutrition has little influence on primary

muscle formation during this early time frame.⁶ During the second to eighth month of gestation, the majority of muscle fibers form; therefore, reduction of muscle fiber formation during this stage through any source of stimuli (e.g., maternal nutrition) has long-lasting, irreversible consequences to the offspring.⁶ The prospect of nutritional management's altering marbling may be greatest for the fetal stage, due to its importance in adipocyte formation, followed by the neonatal stage, early weaning stage (i.e., 150 to 250 days of age), and finally, weaning and older stages.⁶

Larson et al²¹ reported increased progeny birth weights from protein-supplemented dams, suggesting a potential alteration in fetal muscle growth. Greenwood et al¹⁵ reported steers from cows nutritionally restricted during gestation had reduced body weight and carcass weight at 30 months of age compared to steers from adequately fed cows. Both Larson et al²¹ and Greenwood et al¹⁵ reported a retail yield on a carcass weight basis was greater in steers from nutrient-restricted cows, indicating an increased propensity for carcass fatness was not a consequence of nutritional restriction in utero.

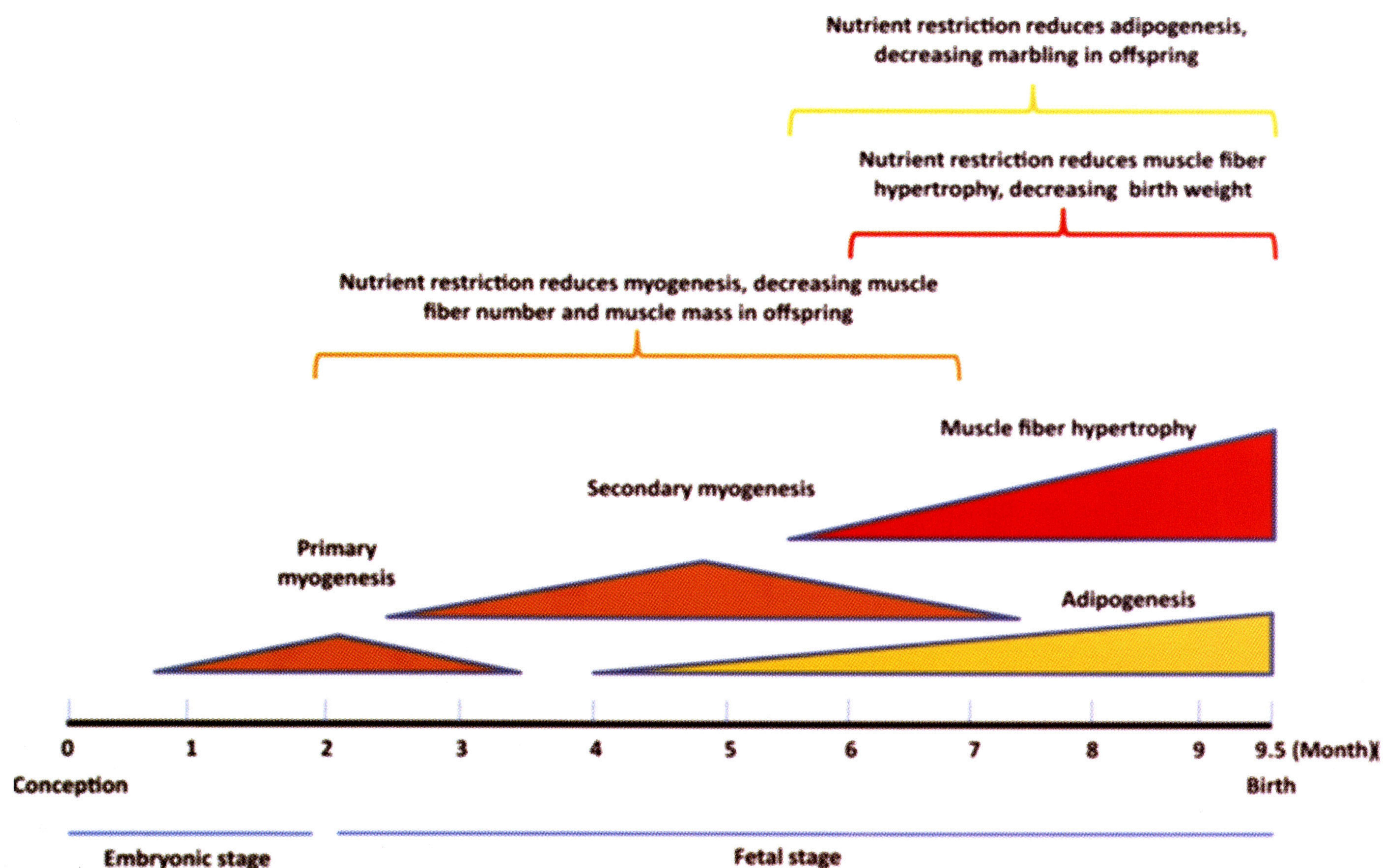


Figure 1. Effects of maternal nutrition on bovine fetal skeletal muscle development. Dates are estimated mainly based on data from studies in sheep, rodents, and humans and represent progression through the various developmental stages. Nutrient restriction during mid-gestation reduces muscle fiber numbers, whereas restriction during late gestation reduces both muscle fiber sizes and the formation of intramuscular adipocytes. From Du M, Tong J, Zhao J, Underwood KR, Zhu M, Ford SP, Nathanielsz PW. Fetal programming of skeletal muscle development in ruminant animals. *J Anim Sci* 2010; 88(E. Suppl.):E51-E60.

Although fetal adipocyte development begins early in gestation, the majority of fetal adipose tissue is not deposited until the final few weeks of gestation.⁴⁸ Adipose tissue growth occurs through preadipocyte proliferation, impacting formation of new mature adipocytes (hyperplasia); and increased size and lipid storage capacity of mature adipocytes (hypertrophy). By feeding ewes 150% of National Research Council (NRC) nutrient requirements, Tong et al^{49,50} reported increased adipogenesis in fetal skeletal muscle. In a review on fetal programming of skeletal muscle, Du et al⁶ reported when University of Wyoming scientists fed beef cattle 1 of 3 diets (100%, 70% of NRC nutrient requirements,³¹ or 70% of NRC nutrient requirements plus supplementation of ruminal bypass protein from day 60 to 180 of gestation), steer progeny from dams fed 70% nutrient requirements plus supplement had numerical decreases in marbling scores when compared to steers from dams fed 100% of requirements. Underwood et al⁵¹ also reported increased tenderness in steers from dams grazed on improved pasture compared to steers from dams grazed on native range during mid-gestation.

Heifer Progeny Performance

Data regarding the effect of late-gestation protein supplementation on heifer progeny performance are reported in Table 1. Martin et al²⁴ conducted a study with cows grazing dormant Sandhills range during late gestation. One group received a 42% CP (DM basis) cube offered 3 times weekly at the equivalent of 1.0 lb (0.45 kg)/day while another group re-

ceived no supplement. Calf birth weight between heifer progeny from supplemented and non-supplemented dams was not different; however, heifer progeny from supplemented cows had increased adjusted 205-day weaning weights, prebreeding weight, weight at pregnancy diagnosis, and improved pregnancy rates compared to heifers from non-supplemented dams. Martin et al²⁴ also reported after a subset of these heifers were placed in a Calan gate individual feeding system, dry matter intake (DMI), average daily gain (ADG), and residual feed intake between heifer progeny from supplemented and non-supplemented dams was not different.

Funston et al,¹⁰ using the same cow herd, offered a distillers based supplement (28% CP, DM basis) 3 times weekly at the equivalent of 1.0 lb (0.45 kg)/day, or no supplement during late gestation as cows grazed either dormant Sandhills range or corn crop residue. Calf weaning weight was greater ($P = 0.04$) for heifers from protein-supplemented dams, whereas Martin et al²⁴ reported a trend ($P = 0.12$) for increased weaning weight for heifers from protein-supplemented dams. Funston et al¹⁰ also reported a decreased age at puberty for heifers from protein-supplemented cows and a trend ($P = 0.13$) for higher pregnancy rates when compared to heifers from non-supplemented dams, possibly related to decreased age at puberty. Similarly, Corah et al⁵ reported heifers born to primiparous heifers fed 100% of their dietary energy requirement during the last 90 days of gestation were pubertal 19 days earlier than heifers born to primiparous heifers fed 65% of their dietary energy requirement.

Funston et al¹⁰ reported no differences in heifer weight at prebreeding and no differences in calf birth weight, calf production, or second calf rebreeding when comparing heifer progeny from supplemented and non-supplemented cows. Gunn et al¹⁶ reported a decrease in the proportion of singleton, and an increase in the proportion of multiple births over 3 parities in progeny born to ewes offered a protein supplement while grazing native pastures during the last 100 days of gestation compared to progeny from non-supplemented ewes. Late-gestation supplementation did not alter the proportion of barren ewe progeny.¹⁶ Martin et al²⁴ reported a 28% increase in the proportion of heifers calving in the first 21 days of the calving season from protein-supplemented dams compared to heifers from non-supplemented dams. Pryce et al³³ reported no difference in progeny heifer reproductive performance when considering dairy cow maternal nutritional status, determined by body condition score (BCS), DMI, and milk yield of fat and protein.

Steer Progeny Performance

As previously mentioned, studies have reported improved muscle development in steers from adequately fed dams when compared to progeny from nutrient-restricted dams. Underwood et al⁵¹ reported increased weight gains, final weight, and hot carcass weight in steers from cows grazing improved pasture from day 120 to 180 of gestation

Table 1. Effect of maternal protein supplementation on heifer progeny performance.

Item	Dietary treatment			
	Martin et al ²⁴		Funston et al ¹⁰	
	NS	SUP	NS	SUP
Weaning BW, lb	456	467	492 ^a	511 ^b
Adj. 205-d wt, lb	481 ^a	498 ^b	470	478
DMI, lb/d	14.39	14.88	20.89	20.50
ADG, lb/d	0.90	0.88	1.86 ^x	1.74 ^y
Residual feed intake	-0.12	0.07	0.08	-0.04
Age at puberty, d	334	339	365 ^x	352 ^y
Pregnant, %	80 ^a	93 ^b	83	90

¹NS = dams did not receive protein supplement while grazing dormant Sandhills range during the last third of gestation; SUP = dams were supplemented 3 times per week with the equivalent of 1.0 lb (0.45 kg)/d of 42% CP cube (DM basis) while grazing dormant Sandhills range during the last third of gestation.

²NS = dams did not receive protein supplement while grazing dormant Sandhills range or corn residue during the last third of gestation; SUP = dams were supplemented 3 times per week with the equivalent of 1.0 lb (0.45 kg)/d of a 28% CP cube (DM basis) while grazing dormant Sandhills range or corn residue during the last third of gestation.

^{a,b}Means within a study with different superscripts differ ($P \leq 0.05$).

^{x,y}Means within a study with different superscripts differ ($P \leq 0.10$).

when compared to progeny from cows grazing native range during that same time (Table 2). Steers from cows grazing improved pasture had increased back fat and tended to have improved marbling scores compared to steers from cows grazing native range.

To determine the effect dietary energy source had on progeny calf performance, Radunz³⁴ offered cows 1 of 3 diets during gestation beginning on approximately day 209 of gestation: hay (fiber), corn (starch), or distillers grains with solubles (fiber plus fat). Corn and distillers grains diets were limit-fed to ensure isocaloric intake among treatments. Results indicated reduced birth weights for calves from dams fed grass hay when compared to calves from the other 2 groups (Table 2), with an increase ($P \leq 0.05$) in calf body weight reported through weaning when comparing calves from corn-fed dams to hay-fed dams. Feedlot performance among treatments was not different; however, calves from hay-fed dams required 8 and 10 more days on feed to reach a similar fat thickness when compared to calves from distillers and corn-fed dams, respectively.

Stalker et al^{45,46} reported steer progeny from dams supplemented the equivalent of 1.0 lb (0.45 kg)/day (42% CP, DM basis) cube during late gestation had no difference in calf birth weight when compared to steers from non-supplemented dams. Conversely, Larson et al²¹ using the same cow herd, reported an increase in calf birth weight when comparing calves born to dams supplemented the equivalent of 1.0 lb (0.45 kg)/day (28% CP, DM basis) cube during late gestation to calves from non-supplemented dams. In the study reported by Stalker et al,⁴⁵ cows were utilized in a switchback design, whereas cows utilized by Larson et al²¹ remained on the same treatment over the 3-year study.

Protein supplementation during late gestation increased weaning weight, ADG to weaning, and proportion of calves weaned when comparing calves from supplemented to non-supplemented dams grazing dormant winter range^{21,45,46}

(Table 3). Stalker et al⁴⁵ reported no differences in steer progeny feedlot performance and carcass characteristics when comparing progeny from supplemented and non-supplemented dams. However, Larson et al²¹ reported increased ADG, HCW, and marbling scores in steers from supplemented dams. Furthermore, a greater proportion of steers from supplemented dams graded USDA Choice and USDA Choice or greater when compared to steers from non-supplemented dams. Non-supplemented cows in Larson et al²¹ may have been under greater nutritional stress than Stalker et al⁴⁵ as average weaning date was approximately 1 month later and possibly had greater impact on fetal development.

Influence of Maternal Nutrition on Progeny Health

Several reports have linked maternal nutrition during gestation to calf health, including Corah et al,⁵ indicating increased morbidity and mortality rates in calves born to primiparous heifers receiving 65% of their dietary energy requirement over the last 90 days of gestation compared to calves from primiparous heifers receiving 100% of their energy requirement. One factor contributing to increased morbidity and mortality is decreased birth weight. Calves born to nutrient-restricted dams were 4.5 lb (2.04 kg) lighter at birth compared to calves from dams receiving adequate nutrition.⁵ Similarly, Moule²⁸ reported as birth weight increased from 4.5 to 9 lb (2.04 kg to 4.08 kg), mortality decreased dramatically in lambs.

Mulliniks et al²⁹ and Larson et al²¹ indicated reduced proportions of steers treated in the feedlot from cows supplemented with protein compared to calves from nonsupplemented dams. Stalker et al⁴⁵ reported increased proportions of live calves weaned to dams offered supplement during late gestation; however, there was no difference in the number of treated calves prior to weaning or in the feedlot. Furthermore, Larson et al²¹ reported no difference in the number of steer calves treated for respiratory disease prior to weaning.

Table 2. Effect of maternal nutrition on steer progeny performance.

Item	Dietary treatment				
	Underwood et al ¹		Radunz ²		
	NR	IP	Hay	Corn	DDGS
Birth BW, lb	85	81	86 ^a	95 ^b	91 ^b
Weaning BW, lb	534 ^a	564 ^b	580 ^a	607 ^b	591 ^{a,b}
ADG, lb/d	3.28 ^a	3.65 ^b	3.37	3.46	3.41
HCW, lb	726 ^a	768 ^b	688	688	675
12th rib fat, in	0.49 ^a	0.65 ^b	0.48	0.50	0.51
Marbling score ³	420	455	549 ^a	506 ^b	536 ^{ab}

¹NR = dams grazed native range from day 120 to 180 of gestation; IP = dams grazed improved pasture from day 120 to 180 of gestation.

²Hay = dams offered a diet of grass hay beginning on day 209 of gestation; Corn = dams offered limit-fed diet of corn beginning on day 209 of gestation; DDGS = cows offered a limit-fed diet of distillers grains with solubles beginning on day 209 of gestation.

³Where 400 = Small^o.

^{a,b}Means within a study with different superscripts differ ($P \leq 0.05$).

Similarly, Funston et al¹⁰ reported no differences in illness in cohort heifers.

Snowder et al⁴⁴ reported disease incidence is more likely after 5 days on feed and remains high through the first 80 days in the feedlot. Furthermore, steers were more likely to become sick compared to heifers in the feedlot. Post-weaning stress is a factor influencing calf health. As mentioned earlier, Funston et al¹⁰ did not report any differences in heifer calf health. These heifers, unlike their steer cohorts, remained at the ranch post-weaning and were maintained on a forage based diet, likely reducing the amount of stress placed on the animal when compared to their steer cohorts who were transported to the feedlot 2 weeks post-weaning and adapted to a concentrate-based diet.

Conclusion

Management of maternal diet beginning during early gestation will ensure proper placental programming resulting in adequate nutrient transfer to the fetus. Maternal nutrition later in gestation has been reported to influence fetal organ development, muscle development, postnatal calf performance, carcass characteristics, and reproduction. Although the mechanisms by which placental and fetal programming occur are not clear, managing resources to ensure proper cow nutrient intake during critical points of gestation can improve calf performance and health.

References

1. Barker DJP, Martyn CN, Osmond C, Hales CN, Fall CHD. Growth in utero and serum cholesterol concentrations in adult life. *BMJ* 1993; 307:1524-1527.

2. Bauman DE, Eisemann JH, Currie WB. Hormonal effects on partitioning of nutrients for tissue growth: role of growth hormone and prolactin. *Fed Proc* 1982; 41:2538-2544.
3. Caton JS, Grazul-Bilska AT, Vonnahme KA, Luther JS, Lardy GP, Hammer CJ, Redmer DA, Reynolds LP. Nutritional management during gestation: impacts on lifelong performance. *Proceedings. 18th Annu Fla Rumin Nutrition Symposium, Gainesville, FL. 2007*; 1-20.
4. Close WH, Pettigrew JF. Mathematical models of sow reproduction. *J Reprod Fertil Suppl* 1990; 40:83-88.
5. Corah LR, Dunn TG, Kaltenbach CC. Influence of prepartum nutrition on the reproductive performance of beef females and the performance of their progeny. *J Anim Sci* 1975; 41:819-824.
6. Du M, Tong J, Zhao J, Underwood KR, Zhu M, Ford SP, Nathanielsz PW. Fetal programming of skeletal muscle development in ruminant animals. *J Anim Sci* 2010; 88(E. Suppl.):E51-E60.
7. Ferrell CL. Placental regulation of fetal growth. In: Campion DR, Hausman GJ, Martin RJ, eds. *Animal growth regulation*. New York: Plenum, 1989; 1-19.
8. Ford SP, Long NM. Evidence for similar changes in offspring phenotype following either maternal undernutrition or overnutrition: potential impact on fetal epigenetic mechanisms. *Reprod Fertil Dev* 2012; 24:105-111.
9. Funston RN, Larson DM, Vonnahme KA. Effects of maternal nutrition on conceptus growth and offspring performance: implications for beef cattle production. *J Anim Sci* 2010; 88(E. Suppl.):E205- E215.
10. Funston RN, Martin JL, Adams DC, Larson DM. Winter grazing system and supplementation of beef cows during late gestation influence heifer progeny. *J Anim Sci* 2010; 88:4094-4101.
11. Gardner DS, Lea RG, Sinclair KD. Developmental programming of reproduction and fertility: what is the evidence? *Anim* 2008; 2:1128-1134.
12. Gilbert JS, Ford SP, Lang AL, Pahl LR, Drumhiller MC, Babcock SA, Nathanielsz PW, Nijland MJ. Nutrient restriction impairs nephrogenesis in a gender specific manner in the ovine fetus. *Pediatr Res* 2007; 61:42-47.
13. Gnanalingham MG, Mostyn A, Dandrea J, Ykubu DP, Symonds ME, Stephenson T. Ontogeny and nutritional programming of uncoupling protein-2 and glucocorticoid receptor mRNA in the ovine lung. *J Physiol* 2005; 565:159-169.
14. Greenwood PL, Bell AW. Consequences of intrauterine growth retardation for postnatal growth, metabolism and pathophysiology. *Reprod Suppl* 2003; 61:195-206.
15. Greenwood PL, Café LM, Hearnshaw H, Hennessy DW, Morris SG. Consequences of prenatal and preweaning growth for yield of beef primal cuts from 30-month-old Piedmontese and Wagyu-sired steers. *Anim Prod Sci* 2009; 49:468-478.

Table 3. Effect of maternal protein supplementation on steer progeny performance.

Item	Dietary treatment					
	Stalker et al ¹		Stalker et al ¹		Larson et al ²	
	NS	SUP	NS	SUP	NS	SUP
Weaning BW, lb	463 ^a	489 ^b	463 ^a	476 ^b	514 ^a	529 ^b
DMI, lb/d	24.6 ^a	26.6 ^b	18.7	18.8	19.8 ^x	20.3 ^y
ADG, lb/d	3.53	3.70	3.46	3.44	3.66	3.75
Feed:gain	6.97	7.19	5.41	5.46	5.37	5.38
HCW, lb	765 ^a	805 ^b	800	814	805 ^a	822 ^b
Choice, %	-	-	85	96	71	85
Marbling score ³	449	461	467	479	445 ^a	492 ^b

¹NS = dams did not receive protein supplement while grazing dormant Sandhills range during the last third of gestation; SUP = dams were supplemented 3 times per week with the equivalent of 1.0 lb (0.45 kg)/d of 42% CP cube (DM basis) while grazing dormant Sandhills range during the last third of gestation.

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³Where 400 = Small^o.

^{a,b}Means within a study with different superscripts differ ($P \leq 0.05$).

^{x,y}Means within a study with different superscripts differ ($P \leq 0.10$).

16. Gunn RG, Sim DA, Hunter EA. Effects of nutrition in utero and in early life on the subsequent lifetime reproductive performance of Scottish Blackface ewes in two management systems. *Anim Sci* 1995; 60:223-230.
17. Han HC, Austin KJ, Nathanielsz PW, Ford SP, Nijland MJ, Hansen TR. Maternal nutrient restriction alters gene expression in the ovine fetal heart. *J Physiol* 2004; 558:111-121.
18. Hirshfield AN. Relationship between the supply of primordial follicles and the onset of follicular growth in rats. *Biol Reprod* 1994; 50:421-428.
19. Hubbert WT, Stalheim OHV, Booth GD. Changes in organ weights and fluid volumes during growth of the bovine fetus. *Growth* 1972; 36:217-233.
20. Kelly AP, Janzen ED. A review of morbidity rates and disease occurrence in North American feedlot cattle. *Can Vet J* 1986; 27:496-500.
21. Larson DM, Martin JL, Adams DC, Funston RN. Winter grazing system and supplementation during late gestation influence performance of beef cows and steer progeny. *J Anim Sci* 2009; 87:1147-1155.
22. Limesand SW, Jensen J, Hutton JC, Hay Jr WW. Diminished beta-cell replication contributes to reduced beta-cell mass in fetal sheep with intrauterine growth restriction. *Am J Physiol Regul Integr Comp Physiol* 2005; 288:1297-1305. Epub 2005 Jan 13.
23. Limesand SW, Rozance PJ, Zerbe GO, Hutton JC, Hay Jr WW. Attenuated insulin release and storage in fetal sheep pancreatic islets with intrauterine growth restriction. *Endocrinology* 2006; 147:1488-1497. Epub 2005 Dec 8.
24. Martin JL, Vonnahme KA, Adams DC, Lardy GP, Funston RN. Effects of dam nutrition on growth and reproductive performance of heifer calves. *J Anim Sci* 2007; 85:841-847.
25. Matsuaki M, Milne JL, Aitken RP, Wallace JM. Overnourishing pregnant adolescent ewes preserves perirenal fat deposition in their growth-restricted fetuses. *Reprod Fertil Dev* 2006; 18:357-364.
26. McMillin IC, Muhlhausler BS, Duffield JA, Yuen BS. Prenatal programming of postnatal obesity: fetal nutrition and the regulation of leptin synthesis and secretion before birth. *Proc Nutr Soc* 2004; 63:405-412.
27. Metcalfe J, Stock MK, Barron DH. Maternal physiology during gestation. In: Knobil E, Neill J, Ewing JJ, eds. *Physiology of reproduction*. New York: Raven Press, 1988; 2145-2176.
28. Moule GR. Some problems of sheep husbandry in tropical Australia. *Aust Vet J* 1956; 32:289-298.
29. Mulliniks JT, Cox SH, Ivey SL, Mathis CP, Sawyer JE, Petersen MK. Cow nutrition impacts feedlot pull rate. *Proc West Sec Am Soc Anim Sci* 2008; 59:91-94.
30. Nilsson EE, Skinner MK. Progesterone regulation of primordial follicle assembly in bovine fetal ovaries. *Mol Cell Endocrinol* 2009; 313:9-16.
31. NRC. *Nutrient requirements of beef cattle*. 7th ed. Washington DC: National Academy Press, 2000.
32. Patten BM. *Foundations of embryology*. 2nd ed. New York: McGraw-Hill, 1964.
33. Pryce JE, Simm G, Robinson JJ. Effects of selection for production and maternal diet on maiden dairy heifer fertility. *Anim Sci* 2002; 74:415-421.
34. Radunz AE. Effects of prepartum dam energy source on progeny growth, glucose tolerance, and carcass composition in beef and sheep. PhD Dissertation. The Ohio State University, Columbus, 2009.
35. Ramsey EM. *The placenta, human and animal*. New York: Praeger, 1982.
36. Reynolds LP, Ferrell CL, Robertson DA, Ford SP. Metabolism of the gravid uterus, foetus and uteroplacenta at several stages of gestation in cows. *J Agric Sci* 1986; 106:437-444.
37. Reynolds LP, Redmer DA. Utero-placental vascular development and placental function. *J Anim Sci* 1995; 73:1839-1851.
38. Reynolds LP, Redmer DA. Angiogenesis in the placenta. *Biol Reprod* 2001; 64:1033-1040.
39. Reynolds LP, Caton JS, Redmer DA, Grazul-Bilska AT, Vonnahme KA, Borowicz PP, Luther JS, Wallace JM, Wu G, Spencer TE. Evidence for altered placental blood flow and vascularity in compromised pregnancies. *J Physiol* 2006; 572:51-58.
40. Reynolds LP, Borowicz PP, Caton JS, Vonnahme KA, Luther JS, Hammer CJ, Maddock Carlin KR, Grazul-Bilska AT, Redmer DA. Developmental programming: the concept, large animal models, and the key role of uteroplacental vascular development. *J Anim Sci* 2010; 88(E. Suppl.):E61-E72.
41. Robinson JJ, McDonald I, Fraser C, McHattie I. Studies on reproduction in prolific ewes. I. Growth of the products of conception. *J Agric Sci Camb* 1977; 88:539-552.
42. Rosenfeld CR, Morriss FH, Makowski EL, Meschia G, Battaglia FC. Circulatory changes in the reproductive tissues of ewes during pregnancy. *Gynecol Invest* 1974; 5:252-268.
43. Russell RG, Oteruelo FT. An ultrastructural study of the differentiation of skeletal muscle in the bovine fetus. *Anat Embryol (Berl)* 1981; 162:403-417.
44. Snowden GD, Van Vleck LD, Cundiff LV, Bennett GL. Bovine respiratory disease in feedlot cattle: environmental, genetic, and economic factors. *J Anim Sci* 2006; 84:1999-2008.
45. Stalker LA, Adams DC, Klopfenstein TJ, Feuz DM, Funston RN. Effects of pre- and postpartum nutrition on reproduction in spring calving cows and calf feedlot performance. *J Anim Sci* 2006; 84:2582-2589.
46. Stalker LA, Ciminski LA, Adams DC, Klopfenstein TJ, Clark RT. Effects of weaning date and prepartum protein supplementation on cow performance and calf growth. *Rangeland Ecol Manage* 2007; 60:578-587.
47. Stickland NC. A quantitative study of muscle development in the bovine foetus (*Bos indicus*). *Anat Histol Embryol* 1978; 7:193-205.
48. Symonds ME, Stephenson T, Gardner DS, Budge H. Long-term effects of nutritional programming of the embryo and fetus: mechanisms and critical windows. *Reproduction Fertility and Development* 2007; 19:53-63.
49. Tong J, Zhu MJ, Underwood KR, Hess BW, Ford SP, Du M. AMP-activated protein kinase and adipogenesis in sheep fetal skeletal muscle and 3T3-L1 cells. *J Anim Sci* 2008; 86:1296-1305.
50. Tong JF, Yan X, Zhu MJ, Ford SP, Nathanielsz PW, Du M. Maternal obesity downregulates myogenesis and β -catenin signaling in fetal skeletal muscle. *Am J Physiol Endocrinol Metab* 2009; 296:E917-E924.
51. Underwood KR, Tong JF, Price PL, Roberts AJ, Grings EE, Hess BW, Means WJ, Du M. Nutrition during mid to late gestation affects growth, adipose tissue deposition and tenderness in cross-bred beef steers. *Meat Sci* 2010; 86:588-593.
52. Vogel GJ, Parrott C. Mortality survey in feedyards: the incidence of death from digestive, respiratory, and other causes in feedyards on the Great Plains. *Comp Cont Ed Prac Vet* 1994; 16:227-234.
53. Vonnahme KA, Zhu MJ, Borowicz PP, Geary TW, Hess BW, Reynolds LP, Caton JS, Means WJ, Ford SP. Effect of early gestational undernutrition on angiogenic factor expression and vascularity in the bovine placenta. *J Anim Sci* 2007; 85:2464-2472.
54. Wu G, Bazer FW, Wallace JM, Spencer TE. Board invited review. Intrauterine growth retardation: implications for the animal sciences. *J Anim Sci* 2006; 84:2316-2337.
55. Zhu MJ, Ford SP, Nathanielsz PW, Du M. Effect of maternal nutrient restriction in sheep on the development of fetal skeletal muscle. *Biol Reprod* 2004; 71:1968-1973.
56. Zhu MJ, Du M, Hess BW, Means WJ, Nathanielsz PW, Ford SP. Maternal nutrient restriction upregulates growth signaling pathway in the cotyledonary artery of cow placentomes. *Placenta* 2007; 28:361-368.