



Invited Review

Omega-3 fatty acid supplementation in horses

Tanja Hess¹, Trinette Ross-Jones²

¹ Colorado State University, Department of Animal Science, Fort Collins, CO, USA.

² Department of Animal Science and Wildlife Management, Tarleton State University, Stephenville, TX, USA.

ABSTRACT - Polyunsaturated omega-3 fatty acids (n-3 PUFA) are a family of essential fatty acids with many biological activities. These fatty acids are incorporated into cell membranes, changing their structural and functional characteristics. N-3 PUFA can act by modulating inflammatory responses at different levels. Omega-3 PUFA can be converted in the body to longer-chain n-3 PUFA at a limited rate and are differently converted in body systems. It appears that when specific longer-chain n-3 PUFA are desired these need to be supplemented directly in the diet. In different species some evidence indicates a potential effect on improving insulin sensitivity. Recently, a novel class of n-3 PUFA-derived anti-inflammatory mediators have been recognized, termed E-series and D-series resolvins, formed from EPA and DHA, respectively. N-3 PUFA derived resolvins and protectins are heavily involved in the resolution of inflammation. Supplementation with n-3 fatty acids in horses may help manage chronic inflammatory conditions such as osteoarthritis, equine metabolic syndrome, laminitis, and thereby help to improve longevity of sport horse.

Key Words: alpha linolenic acid, arachidonic acid inflammation, docosahexaenoic acid, eicosapentaenoic acid, linoleic acid

Dietary n-3 and n-6 long-chain polyunsaturated fatty acids (PUFA)

Dietary fats are required to support absorption of fat-soluble vitamins and provide the essential fatty acids (NRC, 2007), linoleic acid (LA) and alpha-linolenic acid (ALA). The long-chain fatty acid family of omega-6 (n-6) and omega-3 polyunsaturated fatty acids (PUFA) are essential components of the diet and are necessary in daily physiological functions as well as for fetal development and neonatal growth. The 'parent' fatty acids are linoleic acid (LA) for the n-6 family and alpha-linolenic acid (ALA) for the n-3 family. These 'parent' fatty acids are considered essential in mammalian diets; the lack of proper enzymes prevents their endogenous synthesis. The derivatives of these 'parents' are likely of even greater importance; LA can be elongated and converted to arachidonic acid (ARA), whose primary role is to produce 20 carbon-signaling molecules known as eicosanoids. Eicosanoids have short half-lives and are localized close to their production site, influencing events within and around the cells that produce them. Arachidonic acid and other eicosanoid-producing fatty

acids must be present in tissue in order for these signaling molecules to be effective. Eicosanoids are important in that they regulate a variety of cellular functions, during both physiological (normal) and inflammatory events. The most well-known classes of eicosanoids are prostaglandins (PG), thromboxanes (TX), leukotrienes (LT) and lipoxins (LX); with PG and TX being synthesized via the cyclooxygenase (COX) pathway and LT and LX being converted from ARA by lipoxigenases (Figure 1). While all classes are vital physiological components, prostaglandins are important as they are utilized by all major organ systems including reproductive, gastrointestinal and neurological. Of these active eicosanoids, prostaglandin E₂ is the primary PG, synthesized exclusively from ARA, playing an important role in the inflammatory response. In terms of joint health and the development of osteoarthritis, PGE₂ has been implicated as therapeutic target as it is elevated in early stages of the disease and contributes to downstream production of degradative cartilage enzymes (McIlwraith, 2005).

Long-chain derivatives of ALA, specifically eicosapentaenoic acid (EPA), n-3 docosapentaenoic acid (DPA) and docosahexaenoic acid (DHA) (Figure 2), have as equally important roles as ARA in cellular function and physiologic homeostasis. While all ALA derivatives can be converted to produce eicosanoids, EPA is the most widely recognized n-3 PUFA that is a source for anti-inflammatory

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Corresponding author: tanja.hess@colostate.edu

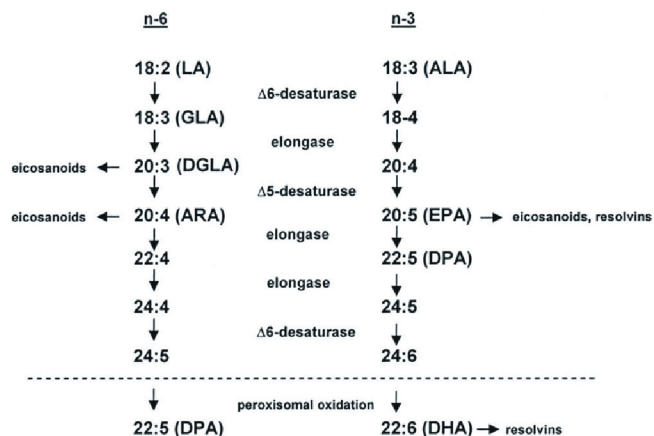
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PG, TX, LT and LX (Figure 1). However, research results indicate that n-3 DPA and DHA also play vital roles in mediating the inflammatory response in conjunction with EPA (Kaur et al., 2011). Increased intake of EPA and DHA in a dose-dependent manner has been shown to decrease ARA amounts in cell membrane phospholipids involved in inflammation (Calder, 2014). Furthermore, increased EPA intake has been shown to inhibit ARA metabolism (Calder, 2014) and decrease the expression of the pro-inflammatory gene COX-2 (Calder, 2014). One of the premier sources of EPA and DHA is marine fish oil, also known as menhaden oil. Eicosapentaenoic acid, though found in the largest quantities in marine fish oil, originates from ALA (Figure 2), which is encountered in high concentrations in plant fats. Plant oils such as linseed, soybean and flaxseed oils are all sources of ALA, therefore, when consumed, the animal may be able to synthesize EPA, DPA and DHA from these plant oils (Figure 2). Alpha-linolenic acid is converted to EPA, DPA and DHA via a desaturation and elongation pathway (Figure 2). The initial step, the addition of a double bond to ALA by the $\Delta 6$ -desaturase enzyme, is the

ARA	EPA
- Prostaglandins (PGE ₂)	- Prostaglandins (PGE ₃)
- Thromboxanes (A ₂)	- Thromboxanes (A ₃)
- Leukotrienes (4-series)	- Leukotrienes (5-series)

Figure 1 - Eicosanoid formation from arachidonic acid (ARA) and eicosapentaenoic acid (EPA).



ALA - α -linolenic acid; ARA - arachidonic acid; DGLA - dihomo- γ -linolenic acid; DHA - docosahexaenoic acid; DPA - docosapentaenoic acid; EPA - eicosapentaenoic acid; GLA - γ -linolenic acid; LA - linoleic acid.

Figure 2 - Biochemical pathway for the incorporation of n-3 and n-6 fatty acids.

Source: Reprinted from Arterburn et al. (2006), with permission.

rate-limiting step in the pathway and contributes to the reported low conversion efficiency of ALA to the longer-chain PUFA (Calder, 2013). Both LA and ALA share a need for $\Delta 6$ -desaturase (Figure 2), and while the enzyme has a higher affinity for ALA, in humans dietary fats contain a higher percentage of LA, therefore competing for the same enzyme ALA needs for conversion to its longer derivate (Tu et al., 2010). In addition, ALA has been shown to have the highest oxidation rate among all unsaturated fatty acids in human tracer studies (Nettleton, 1991), contributing to the low conversion of ALA to its longer derivate. A review of studies investigating the efficiency of dietary ALA conversion in humans reported the conversion of ALA to EPA ranged from 8-10% with conversion efficiency being as low at 4% for DHA (Williams and Burdge, 2006). Due to evidence of a low conversion rate, it is recommended to supply EPA and DHA directly in the diet (Arterburn et al., 2006).

A source of ALA in the equine diet is forage (hay or pasture), which is the largest portion of equine diets. Many horses are fed forage alone; however, performance horses require additional energy-dense feeds such as grains or oils due to their higher energy demands. Vegetable oils are also a source of ALA, such as flaxseed and linseed oil, which can be top-dressed onto feed. Linoleic acid is very abundant in cereal grains such as corn and barley, and makes up the majority of the fat in corn oil as well. Algae produce the n-3 PUFA, EPA and DHA; therefore they are present in fish because algae is an ordinary diet of fish. For that reason, marine foods and fish oil are a good source of EPA and DHA and may be consumed by humans and other carnivores or omnivores. Additionally, they have been used as supplements in equine diets. Horse studies indicated that supplementation with ALA would not lead to increases in circulating DHA (Hansen et al., 2002; Vineyard et al., 2010), only increases in circulating EPA.

Inflammation and PUFA

Cellular activities involved in inflammatory responses are designed to be harmful to pathogens; however, they can cause damage to the host tissues (Calder, 2014). Inflammation usually is self-limiting, and resolves rapidly due to the activation of negative feed-back mechanisms like secretion of anti-inflammatory cytokines or pro-resolving lipid mediators, shedding of receptors for inflammation, and activation of regulatory cells (Calder, 2014). Loss of this regulatory process can result in excessive, inappropriate or chronic inflammation that can cause damage to the host organism (Calder, 2014). As part of these anti-inflammatory and pro-resolving mediators, a novel class

of n-3 PUFA-derived anti-inflammatory mediators have been recognized, termed E-series and D-series resolvins, formed from EPA and DHA, respectively (Calder, 2009). Protectins are another class of mediators produced from DHA (Calder, 2014). While information is limited, it appears n-3 PUFA-derived resolvins and protectins are heavily involved in the resolution of inflammation (Kohli and Levy, 2009). Protectins D1 are produced from DHA, and known as protectins D1, owing to their protecting activity in inflammatory and neural systems. Biological effects of resolvins and protectins have been studied in cell culture and animal models of inflammation and have been shown to stimulate resolution and reduce magnitude of inflammatory response *in vivo* (Serhan et al., 2008). Resolvin E1 reduces inflammation *in vivo* and blocks human transendothelial migration (Serhan et al., 2004). Resolvin E2 reduces zymosan initiated neutrophil infiltration (Tjonahem et al., 2006). Resolvin D1 has been shown to be a potent regulator of mouse and human neutrophils (Serhan et al., 2004; Sun et al., 2007). Resolvin E1 also has been shown to initiate resolution of inflammation causing a decrease in the number of neutrophils in exsudates sooner than during spontaneous resolution. Resolvin E1 and resolvin D1 prevented the infiltration of neutrophils into sites of inflammation, and inhibited IL-1 β production (Calder, 2014). Protectin D1 also blocked T cell migration *in vivo*, reducing TNF- α (tumor necrosis factor alpha), IL-1 β , interferon- γ (IF- γ) secretion, and promoting T cell apoptosis (Ariel et al., 2006). Protectin D1 shifted the onset of resolution to an earlier time point in addition to shortening the time to reduce the number of maximum neutrophils by half (Serhan et al., 2008).

Other anti-inflammatory effects of n-3 PUFA from plant and animal sources include a reduced cytokine production *in vitro* (De Caterina et al., 1994) and *in vivo* (Meydani et al., 1993; Grimm et al., 1994; McCann et al., 2000).

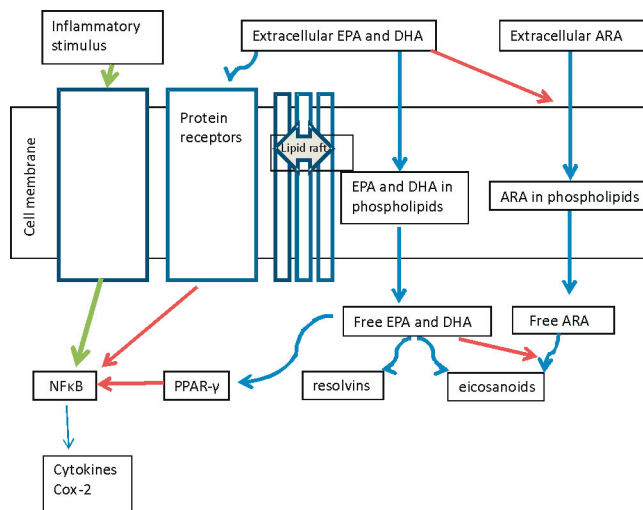
Sources of fatty acids in the diet

In humans, supplementation with long-chain n-3 PUFA has been shown to improve inflammatory status, prevent cardiovascular diseases (Calder, 2001), and reduce pain and inflammation in patients with rheumatoid arthritis (MacLean et al., 2004). In arthritic horses, supplementation with EPA and DHA increased stride length (Woodward et al., 2005) and reduced inflammatory markers (Manhart et al., 2009). In horses, feeding Menhaden fish oil modulated leucotriene synthesis influencing inflammatory conditions (Hall et al., 2004).

Proposed mechanisms of EPA and DHA on inflammation

Several mechanisms have been proposed regarding the processes by which EPA and DHA act on the inflammatory response in tissues. It is well established that these lipids act on both a direct (by alteration of eicosanoid production via cyclooxygenase and lipoxygenase pathways) and indirect (modification of gene transcription) mechanism (Calder, 2006). Direct modification of prostaglandin and leukotriene synthesis was outlined previously. Supplementation of n-3 PUFA in feeds will also exert an effect on the expression of inflammatory genes (Renier et al., 1993; Curtis et al., 2000; Wallace et al., 2001).

It is hypothesized that particular fatty acids, such as EPA and/or DHA, may modify transcription factors in the nucleus and thus influence cytokine and eicosanoid production at the level of gene expression (Figure 3). Another theory is that n-6 (i.e., ARA) and n-3 (i.e., EPA and DHA) fatty acids modify protein synthesis of inflammatory mediators via modification of cell surface receptors on lipid rafts or within the cell by suppression of nuclear receptor activation (Chapkin et al., 2009). Additional regulation comes in the form of peroxisome proliferator-activated receptors, which are key nuclear receptors that regulate transcription of genes through ligand binding with a variety



EPA - eicosapentaenoic acid; DHA - docosahexaenoic acid; ARA - arachidonic acid; COX - cyclooxygenase; NFκB - nuclear factor kappa B; PPAR - peroxisome proliferator activated receptor.

Solid red lines indicate inhibition.

Solid blue lines indicate sources of n-3 PUFA.

Solid green lines indicate inflammatory activation.

Figure 3 - Summary of the anti-inflammatory actions of n-3 polyunsaturated fatty acids; modified from Calder (2013).

of lipophilic metabolites, having a high affinity for PUFA, in particular DHA (Stulnig, 2003). Peroxisome proliferator-activated receptors usually have three isoforms; α , β , γ and are believed to be potent regulators of adipocyte function as well as immune molecules such as lymphocytes and macrophages (Marx et al., 2002) influencing downstream transcription of inflammatory cytokines (Figure 3).

Insulin sensitivity and n-3 PUFA

Insulin resistance (decreased insulin sensitivity) in horses has been linked to the development of laminitis, osteochondrosis, and metabolic syndrome (Coffman and Coles, 1983; Ralston, 1996; Treiber et al., 2006; Frank et al., 2010) and is therefore considered a problem in the equine industry. These diseases can result in a loss of function and performance of the horse. Several factors may predispose a horse to developing insulin resistance, such as diet, age, breed/genetics and obesity (Jeffcot et al., 1986; Treiber et al., 2006; Vick et al., 2007). Supplementation with certain dietary components could increase insulin sensitivity in animals that have insulin resistance, reducing the risk for the development of diseases such as metabolic syndrome and laminitis.

Dietary supplementation with n-3 PUFA has been shown to increase insulin sensitivity in pigs and rats (Behme, 1996; Luo et al., 1996). Dietary EPA and DHA incorporate into cell membranes increasing membrane fluidity due to greater unsaturation of the membrane improving glucose transport function (Lardinois et al., 1987; Zhao et al., 2008). Incorporation of EPA and DHA in muscle cell membrane has been shown to increase binding of insulin (Storlein et al., 1991) and m-RNA expression of GLUT-4 transporters in rats (Figueiras et al., 2010). Supplementation with EPA and DHA has shown to improve insulin sensitivity in rats (Storlein et al., 1991), pigs (Behme, 1996), and in humans (Rasic-Milutinovic et al., 2007).

n-3 PUFA supplementation and exercise

Supplementation with EPA and DHA in horses has also been shown to lower heart rate, plasma glycerol, free fatty acids and cholesterol during an exercise test compared to horses supplemented with corn oil (O'Connor et al., 2004). In human trained athletes supplemented with fish oil, heart rate and oxygen consumption was lower than in subjects supplemented with olive oil (Peoples et al., 2008). Incorporation of n-3 PUFA to muscle membranes increased insulin sensitivity (Pan et al., 1995) and resulted in increased ability of skeletal muscle to take up glucose.

Furthermore, studies have shown that lower proportions of long-chain PUFA and higher proportions of saturated fatty acids in skeletal muscle phospholipids are associated with insulin resistance. Skeletal muscle characteristics (i.e., fatty acid profile) have some genetic influence (Baur et al., 1999) but diet and physical activity also influence skeletal muscle fatty acid profile in rats, humans, and horses (Ayre et al., 1996; Andersson et al., 2000; Hess et al., 2012).

Specific horse studies

Effects of two different dietary sources of n-3 PUFA on incorporation into the plasma, red blood cell, and skeletal muscle in horses (Hess et al., 2012)

In humans, the PUFA DHA and EPA need to be supplemented in order for them to be incorporated into tissues. There is limited conversion of ALA to DHA in humans (Burdge et al., 2001; Arterburn et al., 2006). As stated before, in horses, supplementation with ALA did not lead to increases in circulating DHA (Hansen et al., 2002; Vineyard et al., 2010), only EPA. Feeding of n-3 PUFA to horses may increase circulating levels and increase incorporation into muscle tissue. This could potentially improve chronic inflammatory conditions in horses; however, the optimal type of fatty acid to be supplemented needed to be investigated.

In a related study in our laboratory (Hess et al., 2012) in order to compare different sources of dietary omega-3 fatty acid supplementations on plasma, red blood cell and skeletal muscle fatty acid compositions in horses, three (alfalfa/bromegrass) hay and barley diets were fed: one was supplemented with an algae and fish oil containing DHA and EPA (MARINE; Magnitude; JBS United, Sheridan, IN); another group was supplemented with the same amount of ALA in flaxseed (FLAX) as FLAX, and a third group (control; CON) was fed hay and barley to make all diets have the same calorie amount. Treatments were supplemented for 90 d. Blood samples and muscle middle gluteal biopsies were collected on d 0, 30, 60, and 90 of supplementation.

Direct supplementation of EPA and DHA through a marine source increased PUFA concentrations in the plasma, red blood cell and muscle tissue of equines (Hess et al., 2012). Although present in muscle tissue at baseline, EPA and DHA increased in horses supplemented with a marine source containing these specific fatty acids. In plasma and red blood cells, EPA and DHA were below detection levels in all groups and increased only in MARINE-supplemented horses. Supplementation with dietary sources containing

EPA and DHA may be indicated when increased incorporation of these n-3 fatty acids to muscle and red blood cells is desired. Conversion of ALA from flaxseed and forages to EPA and DHA occurs in the skeletal muscle, as seen in this study (Hess et al., 2012) by the detection of these fatty acids in equine muscles at baseline. Supplementation with ALA through flaxseed lead to higher incorporation of muscle n-3 DPA, a derivative of EPA, compared to MARINE supplementation. Some positive effects of n-3 DPA on inflammation have been reported; however, the effects of such increases should be evaluated in further studies addressing inflammatory responses and compared to EPA and DHA in horses (Hess et al., 2012). In this study, conversion of ALA to EPA and DHA did not occur after supplementation with extra ALA (above the control dietary n-3 PUFA level) from FLAX.

Effects of n-3 PUFA supplementation on insulin sensitivity in horses (Hess et al., 2013)

Dietary supplementation with n-3 PUFA has been shown to increase insulin sensitivity as described above. In order to test the hypothesis that supplementation with n-3 PUFA would improve insulin sensitivity, the same diets described for the previous study were fed to a group of 21 adult mares to test glucose and insulin dynamics. Specific tests to determine insulin sensitivity (frequent sampling intravenous glucose tolerance tests) were performed on days 0, 30, 60, and 90 of supplementation (Hess et al., 2013).

No overall treatment effect was observed when all mares within each treatment were compared among themselves. However, when treatments were compared among mares with the lowest quintile in insulin sensitivity (Treiber et al., 2005) (11 mares) there was a trend ($P = 0.08$) for a treatment effect, where MARINE ($n = 5$) and FLAX ($n = 3$) horses had higher insulin sensitivity ($SI = 1.18 \pm 0.16$ in FLAX, and 1.05 ± 0.16 in MARINE compared to 0.59 ± 0.16 in CON, $n = 3$). Although a small number of insulin resistant mares were compared, further studies should be performed in a larger group of insulin resistant horses. If proven effective, supplementation with omega-3 fatty acids could help to reduce problems associated with insulin resistance in horses (Hess et al., 2013).

Evaluation of synovial fluid in horses fed two different sources n-3 PUFA: a pilot study (Ross-Jones et al., 2014)

Elevating n-3 PUFA in mammalian diets may downregulate inflammatory processes in the joint

(Proudman et al., 2008) and has been shown to have symptom-modifying effects in inflammatory diseases (Lau et al., 1993). One hypothesized mechanism is the potential of long-chain n-3 PUFA to reduce the production of potent inflammatory eicosanoids (Chapkin et al., 2009), primarily PGE_2 . In order to test the effects of different sources of n-3 PUFA on synovial fluid composition, the diets described on the first study (Hess et al., 2012) were supplemented for 90 days (Ross-Jones et al., 2014).

On day 90 of supplementation, approximately 3 mL of synovial fluid were extracted from the right carpus of each horse. Fluid was analyzed for fatty acid concentration and PGE_2 concentration. Synovial fluid samples from the MARINE group exhibited EPA and DHA, whereas the FLAX and CON groups did not express detectable concentrations. Synovial prostaglandin E_2 concentration in the MARINE group tended to be lower compared to CON horses ($P < 0.10$).

Synovial fluid fatty acid levels for EPA and DHA were significantly higher in the MARINE group compared to either CON or FLAX groups, indicating that direct supplementation of EPA and DHA is required if higher fluid concentrations of those fatty acids are desired (Ross-Jones et al., 2014). A small difference between treatment and control in synovial fluid PGE_2 concentration in the current study may be due to all horses being healthy and free of existing joint inflammation or disease. Inducing experimental inflammation in healthy animals receiving a dietary n-3 PUFA supplement may cause measurable differences in eicosanoid levels.

Results indicated a possible inhibition of inflammatory eicosanoid prostaglandin E_2 production in equine synovial fluid by oral supplementation of the polyunsaturated fatty acids EPA and DHA. Further research is needed to determine if oral n-3 PUFA supplementation can be therapeutically advantageous in horses experiencing joint inflammation.

Implications

Based on results from several studies, supplementation with n-3 polyunsaturated fatty acids has the potential to benefit humans and horses in diverse ways. Future studies should address the effects of n-3 polyunsaturated fatty acid supplementation on inflammation and specifically in horses with chronic inflammatory diseases such as laminitis, metabolic syndrome, pituitary pars intermedia disease and osteoarthritis improving the horses' health and wellbeing.

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