

VET 406: Water Soluble Vitamins

Vitamin	Source	Species	Function	Deficiency
Niacin (B <sub>3</sub> )	<p>Good: Meat, Yeast, Leafy Greens</p> <p>Excess tryptophan</p> <p>Poor: Grains (corn and sorghum)</p>	<p>Swine</p> <p>Poultry</p> <p>Dogs</p>	<p>Coenzyme for oxidoreductase reactions</p> <p>Substrate for ADP-ribosyl transferases catalyzing the transfer of ADP-ribose units to proteins</p> <p>Tx of hyperlipidemia</p>	<p>Pellagra (humans)</p> <p>Black tongue (dogs)</p> <p>4D's</p> <p>Dermatitis</p> <p>Diarrhea</p> <p>Dementia</p> <p>Death</p> <p>Swine: diarrhea, stomatitis, dermatitis, alopecia, degenerative changes in the nervous system</p> <p>Poultry: Enlargement of the hock joint, bowing of the legs, stomatitis, glossitis, dermatitis</p> <p>Dogs: Glossitis, gingivitis, necrosis and ulceration of oral mucosa and GIT</p>
Riboflavin (B <sub>2</sub> )	<p>Good: Animal Products, green forages and vegetables</p> <p>Poor: Cereal grains, food products exposed to UV light</p>	<p>Poultry example, but other species are able to be affected</p>	<p>Prosthetic group of flavoproteins and functions in oxidoreduction reactions</p> <ol style="list-style-type: none"> <li>1. NADH Dehydrogenases</li> <li>2. Oxidases</li> <li>3. Dehydrogenases</li> </ol>	<p>Signs primarily involve the eyes, skin or nervous system</p> <p>Photophobia, burning and itching eyes, cataracts and vascularization of the cornea</p> <p>Angular stomatitis, dermatitis, and alopecia</p>

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				Neuropathy, degenerating myelin of the sciatic and brachial nerves  *Curly toe in birds
Vitamin C	Most species can synthesize vitamin C from glucose  Species that lack gluconolactone oxidase require dietary source of vit. C	Required by: Primates  Guinea pigs  Bats  Some birds  Some fish	Collagen synthesis -Required for production of hydroxylation of proline and lysine  Water soluble antioxidant and regenerates reduced vitamin E  Reduces metal ions and enhances Fe absorption  Required for formation of epinephrine from tyrosine (hydroxylation reaction)	Scurvy
Thiamin (B <sub>1</sub> )	Good: Cereal grains  Soybean meal  Greens, leafy hay  Animal products (egg yolk, liver, kidneys)	Dogs  Cats  Poultry  Sheep  Ruminant	Oxidative decarboxylation  Transketolase reaction  Nerve transmission  Reasons for low levels:  Damaged by processing	Beriberi *Star gazing  Cardiovascular and nervous system disorders  Tachycardia (or bradycardia in some species) enlarged heart, edema  Ataxia, paralysis, muscle weakness, mental confusion, severe anorexia,

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	Poor: oils, fats, refined foods		<p>Destruction of thiamin</p> <ul style="list-style-type: none"> <li>-Thiaminases in raw fish, bracken fern, horsetail, star thistle, endophyte-infected tall fescue, gut/rumen microbes</li> <li>-Antagonists, polio encephalomalacia in ruminants</li> </ul>	opisthotonos, bilateral symmetrical lesions in distinct brain regions
Vitamin B <sub>6</sub>			<p>Amino acid metabolism</p> <ul style="list-style-type: none"> <li>-Transaminases</li> <li>-decarboxylation reactions</li> <li>-side chain cleavage</li> </ul> <p>Glycogen metabolism</p> <ul style="list-style-type: none"> <li>-binds to lysine in glycogen phosphorylase to make the enzyme active</li> </ul>	
Pantothenic Acid			<p>Component of CoA required for fatty acid carbohydrate and protein metabolism.</p> <p>Component of acyl carrier protein required for fatty acid synthesis</p>	
Biotin (Vit. H)			<p>Coenzyme for four carboxylase enzymes</p> <ul style="list-style-type: none"> <li>-pyruvate carboxylase</li> </ul>	Sever dermatitis and alopecia

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			-Acetyl CoA carboxylase -Propionyl CoA carboxylase -3-methylcrotonyl CoA carboxylase	Cracked hooves, dermal lesions on the bottom of the feet
Folate (folacin)			Methyl-met formation  Methylene- DNA synthesis  Formimino- Histidine metabolism	Megaloblastic anemia and leukopenia
Vit. B <sub>12</sub>	Only synthesized by microorganisms  Requires cobalt  Animal products (meat, milk, eggs, fish)  Intrinsic factor is required for absorption in the ileum Stomach, pancreas, and ileum are all required for absorption		Methionine synthetase  Methylmalonyl CoA mutase	Methylmalonic aciduria, hyperammonemia  Wasting syndrome  Megaloblastic anemia (humans)  Normocytic, normochromic anemia (some animals)  Neutropenia