# **Choline: Needed for Normal Development of Memory**

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Key words: choline, development, pregnancy, eggs, brain

Choline is a dietary component essential for normal function of all cells. It, or its metabolites, assures the structural integrity and signaling functions of cell membranes; it is the major source of methyl-groups in the diet (one of choline's metabolites, betaine, participates in the methylation of homocysteine to form methionine); and it directly affects nerve signaling, cell signaling and lipid transport/metabolism. In 1998, the National Academy of Sciences, USA, issued a report identifying choline as a required nutrient for humans and recommended daily intake amounts. Eggs are an excellent dietary source of choline.

Pregnancy and lactation are periods when maternal reserves of choline are depleted. At the same time, the availability of choline for normal development of the brain is critical. When rat pups received choline supplements (in utero or during the second week of life), their brain function changed, resulting in the lifelong memory enhancement. This change in memory function appears to be due to changes in the development of the memory center (hippocampus) in the brain. The mother's dietary choline during a critical period in brain development of her infant influences the rate of birth and death of nerve cells in this center. These changes are so important that we can pick out the groups of animals whose mothers had extra choline even when these animals are elderly. Thus, memory function in the aged rat is, in part, determined by what the mother ate. This is not the first example of a critical nutrient that must be present at a specific time in brain development. If folate isn't available in the first few weeks of pregnancy, the brain does not form normally. Thus, we suggest that pregnancy is a period when special attention has to be paid to dietary intake.

### **Key teaching points:**

- · Choline has been identified as a required nutrient.
- · Studies indicate that choline plays a role in development of memory.
- Choline intake during pregnancy may be important for brain development.
- Eggs are a good source of choline in the diet.

The Institute of Medicine (IOM) of the US National Academy of Sciences recently made recommendations for dietary choline requirements [1]. Choline in the diet is important for many reasons: it is needed for synthesis of the phospholipids in cell membranes, methyl metabolism, cholinergic neurotransmission, transmembrane signaling, and lipid-cholesterol transport and metabolism [2]. Most choline in the body is found in phospholipids such as phosphatidylcholine and sphingomyelin. Though representing a smaller proportion of the total choline pool, important metabolites of choline include platelet-activating factor, acetylcholine, choline plasmalogens, lysophosphatidylcholine, phosphocholine, glycerophosphocholine and betaine. There are

several comprehensive reviews of the metabolism and functions of choline [1–3].

Male humans [4] and many species of animals, including the baboon, fed a choline deficient diet deplete choline stores and develop liver dysfunction [5–9]. Some humans (male and female) fed with total parenteral nutrition (TPN) solutions devoid of choline, but adequate for methionine and folate, develop fatty liver and liver damage that resolves when a source of dietary choline is provided [10–15]. Fatty liver occurs because choline is required to make the phosphatidylcholine portion of the very-low-density lipoprotein (VLDL) particle that is needed to export triacylglycerol from the liver

Presented, in part, at a meeting sponsored by the American Egg Board and Egg Nutrition Center held at Amelia Island, FL on February 25–27, 2000. An honorarium was provided for support of this manuscript by the Egg Nutrition Center.

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Journal of the American College of Nutrition, Vol. 19, No. 5, 528S–531S (2000) Published by the American College of Nutrition

[16,17]. Animals fed a choline deficient diet may also develop growth retardation, renal dysfunction and hemorrhage, or bone abnormalities [8,18,19]. There is no doubt that cells in culture absolutely require choline [20] and die by programmed cell suicide when deprived of this nutrient [21-23]. The exact amount of choline that the human diet must contain to sustain life is modulated by the pathway (most active in the liver) for the de novo biosynthesis of the choline moiety via the sequential methylation of phosphatidylethanolamine using S-adenosylmethionine as the methyl donor [24]. This ability to form choline moiety de novo means that some of the demand for choline can, in part, be met by using methyl-groups derived from one carbon metabolism (via methyl-folate and methionine). Because of this metabolic interrelationship, choline deficiency depletes cells of methyl-folate and methionine, and increases intracellular S-adenosylhomocysteine and homocysteine concentrations [25,26]. As our understanding of the importance of folate and homocysteine nutrition increases, there should be increased interest in how choline interacts with folate and homocysteine metabolism.

The IOM recommended an Adequate Intake (AI) estimation for choline in the diet. The IOM report cautioned, 'this amount will be influenced by the availability of methionine and methylfolate in the diet. It may be influenced by gender, and it may be influenced by pregnancy, lactation, and stage of development. Although AIs are set for choline, it may be that the choline requirement can be met by endogenous synthesis at some of these stages.' The IOM recommendations are shown in Table 1.

Almost no information is available about the choline content of foods. Eggs have an especially high choline moiety content (about 300 mg choline/egg, mostly in the form of phosphatidylcholine). Both commercially available infant formulas and bovine milk contain choline and choline-containing compounds [27]. Plasma choline concentration varies in response to diet [28] and can rise as much as two-fold after a two-egg meal. Fasting plasma choline concentrations vary from 7 to 20  $\mu$ M, with most subjects having concentrations of 10  $\mu$ M. Individuals that have starved for up to seven days have diminished plasma choline, but levels never drop below approximately 50% of normal [29].

**Table 1.** Recommended Adequate Intakes (AI) for Choline [1]

Population	Age	AI
AI for Infants	0 to 6 months	125 mg/day, 18 mg/kg
	6 to 12 months	150 mg/day
AI for Children	1 through 3 years	200 mg/day
	4 through 8 years	250 mg/day
	9 through 13 years	375 mg/day
AI for Males	14 through 18 years	550 mg/day
	19 and older	550 mg/day
AI for Females	14 through 18 years	400 mg/day
	19 years and older	425 mg/day
AI for Pregnancy	All ages	450 mg/day
AI for Lactation	All ages	550 mg/day

Pregnancy may be a time when dietary supplies of choline are especially limiting. Though female rats are resistant to choline deficiency, pregnant rats are as vulnerable to deficiency as are males [30]. During pregnancy, large amounts of choline are delivered to the fetus across the placenta [31], and this depletes maternal stores of choline [32]. At birth, humans and other mammals have plasma choline concentrations that are much higher than those in adults [33]. The need for choline is likely to be increased during lactation because so much must be secreted into milk (human milk contains 1.5 to 2 mM choline moiety per liter [27]). Lactating rats are more sensitive to choline deficiency than are non-lactating rats [30].

Ensured availability of choline appears to be important to infants because organ growth - extremely rapid in the neonate - requires large amounts of choline for membrane biosynthesis [34,35]. During development, there is a progressive decline in blood choline concentration that begins *in utero*. In fact, plasma or serum choline concentrations are sevenfold higher in the fetus and neonate than they are in the adult [33,34]. This decline in serum choline concentration occurs during the first weeks of life in the rat and the human [36]. High levels of choline circulating in the fetus presumably ensure enhanced availability of choline to tissues. Neonatal rat brain efficiently extracts choline from blood [37,38]. Supplementing choline during the perinatal period further increases blood and brain choline metabolite concentrations [39]

An interesting effect of dietary choline deficiency in rats has never been studied in humans. Choline availability during embryogenesis and perinatal development of the rat and mouse may be especially important. There are two sensitive periods in rodent brain development during which treatment with choline (about 1 mmol/day) produces long-lasting enhancement of spatial memory that is lifelong [40-49]. The first occurs during embryonic days 12 to 17 (rats give birth on day 21), and the second during postnatal days 16 to 30. Choline supplementation during these critical periods elicits a major improvement in memory performance at all stages of training on a 12-arm radial maze. Though all animals were cross-fostered with untreated mothers, even when the rats are old these memory changes persist and can be used to easily identify which rats got more or less choline during the perinatal period; the normal age-associated decrement that occurs in rat memory seems to be delayed in the choline supplemented group ([44], Warren Meck, personal communication). Pups from mothers fed a choline deficient diet during this same period of pregnancy have diminished memory function [44,47]. During embryogenesis, progenitors of neurons and glia divide, many migrate to new locations, and unnecessary cells die by apoptosis [50-52]. These early developmental events determine the future structure and function of the brain. The choline-induced spatial memory facilitation correlates with changes in the birth, death and migration of cells in the hippocampus during fetal brain development [53-56], with altered distribution and morphology of neurons involved in memory storage within the brain [41,46], with biochemical changes in the hippocampus [57] and with electrophysiological changes in the hippocampus [45]. Are these findings in rats likely to be true of humans? We do not know. Human and rat brains mature at different rates; the rat brain is comparatively more mature at birth than is the human brain, but in humans hippocampal development may continue for months or years after birth [58]. A controlled human study is needed, but until then it seems prudent to ensure that dietary intake of choline is adequate during pregnancy. Two eggs per day contain approximately the dietary requirement for choline, and until more foods are analyzed for choline content, pregnant women might want to include eggs in their diets.

# ACKNOWLEDGMENTS

Supported by the National Institutes of Health (AG09525). Support for this work was also provided by grants from the NIH to the UNC Clinical Nutrition Research Unit (DK56350).

# REFERENCES

- Institute of Medicine and National Academy of Sciences USA: 'Dietary Reference Intakes for Folate, Thiamin, Riboflavin, Niacin, Vitamin B12, Panthothenic Acid, Biotin, and Choline.' Washington DC: National Academy Press, 1998.
- Zeisel SH, Blusztajn JK: Choline and human nutrition. Ann Rev Nutr 14:269–296, 1994.
- 3. Kuksis A, Mookerjea S: Choline. Nutr Rev 36:201-207, 1978.
- Zeisel SH, daCosta KA, Franklin PD, Alexander EA, Lamont JT, Sheard NF, Beiser A: Choline, an essential nutrient for humans. FASEB J 5:2093–2098, 1991.
- Hershey JM, Soskin S: Substitution of 'lecithin' for raw pancreas in a diet of depancreatized dog. Am J Physiol 93:657–658, 1931.
- 6. Hoffbauer FW, Zaki FG: Choline deficiency in the baboon and rat compared. Arch Path 79:364–369, 1965.
- daCosta K, Cochary EF, Blusztajn JK, Garner SC, Zeisel SH: Accumulation of 1,2-sn-diradylglycerol with increased membraneassociated protein kinase C may be the mechanism for spontaneous hepatocarcinogenesis in choline deficient rats. J Biol Chem 268: 2100–2105, 1993.
- Newberne PM, Rogers AE: Labile methyl groups and the promotion of cancer. Ann Rev Nutr 6:407–432, 1986.
- Lieber CS, Robins SJ, Li J, DeCarliL M, Mak KM, Fasulo JM, Leo MA: Phosphatidylcholine protects against fibrosis and cirrhosis in the baboon. Gastroenterology 106:152–159, 1994.
- Buchman AL, Dubin M, Jenden D, Moukarzel A, Roch MH, Rice K, Gornbein J, Ament ME, Eckhert CD: Lecithin increases plasma free choline and decreases hepatic steatosis in long-term total parenteral nutrition patients. Gastroenterology 102:1363–1370, 1992.
- 11. Buchman A, Dubin M, Moukarzel A, Jenden D, Roch M, Rice K, Gornbein J, Ament M: Choline deficiency: a cause of hepatic steatosis during parenteral nutrition that can be reversed with

- intravenous choline supplementation. Hepatology 22:1399–1403, 1995.
- Buchman AL, Moukarzel A, Jenden DJ, Roch M, Rice K, Ament ME: Low plasma free choline is prevalent in patients receiving long term parenteral nutrition and is associated with hepatic aminotransferase abnormalities. Clin Nutr 12:33–37, 1993.
- 13. Chawla RK, Wolf DC, Kutner MH, Bonkovsky HL: Choline may be an essential nutrient in malnourished patients with cirrhosis. Gastroenterology 97:1514–1520, 1989.
- Shapira G, Chawla RK, Berry CJ, Williams PJ, Roy RGB, Rudman
   Cysteine, tyrosine, choline and carnitine supplementation of patients on total parenteral nutrition. Nutr Internat 2:334–339, 1986.
- Sheard NF, Tayek JA, Bistrian BR, Blackburn GL, Zeisel SH: Plasma choline concentration in humans fed parenterally. Am J Clin Nutr 43:219–224, 1986.
- Vance JE: Secretion of VLDL, but not HDL, by rat hepatocytes is inhibited by the ethanolamine analogue N-monomethylethanolamine. J Lipid Res 32:1971–1982, 1991.
- Fast D, Vance D: Nascent VLDL phospholipid composition is altered when phosphatidylcholine biosynthesis is inhibited: evidence for a novel mechanism that regulates VLDL secretion. Biochim Biophys Acta 1258:159–168, 1995.
- Handler P, Bernheim F: Choline deficiency in the hamster. Proc Soc Exptl Med 72:569, 1949.
- Fairbanks BW, Krider JL: Significance of B vitamins in swine nutrition. N Am Vet 26: 18–23, 1945.
- Eagle H: The minimum vitamin requirements of the L and Hela cells in tissue culture, the production of specific vitamin deficiencies, and their cure. J Exptl Med 102:595–600, 1955.
- Albright CD, Lui R, Bethea TC, daCosta KA, Salganik RI, Zeisel SH: Choline efficiency induces apoptosis in SV40-immortalized CWSV-1 rat hepatocytes in culture. FASEB J 10:510–516, 1996.
- Shin OH, Mar MH, Albright CD, Citarella MT, daCosta KA, Zeisel SH: Methyl-group donors cannot prevent apoptotic death of rat hepatocytes induced by choline-deficiency. J Cell Biochem 64:196–208, 1997.
- James S, Miller B, Basnakian A, Pogribny I, Pogribna M, Muskhelishvili L: Apoptosis and proliferation under conditions of deoxynucleotide pool imbalance in liver of folate/methyl deficient rats. Carcinogenesis 18:287–293, 1997.
- Bremer J, Greenberg D: Methyl transfering enzyme system of microsomes in the biosynthesis of lecithin (phosphatidylcholine). Biochim Biophys Acta 46:205–216, 1961.
- Varela-Moreiras G, Selhub J, daCosta K, Zeisel SH: Effect of chronic choline deficiency in rats on liver folate content and distribution. J Nutr Biochem 3:519–522, 1992.
- 26. Zeisel SH, Zola T, daCosta K, Pomfret EA: Effect of choline deficiency on S-adenosylmethionine and methionine concentrations in rat liver. Biochem J 259:725–729, 1989.
- Holmes-McNary M, Cheng WL, Mar MH, Fussell S, Zeisel SH: Choline and choline esters in human and rat milk and infant formulas. Am J Clin Nutr 64:572–576, 1996.
- Zeisel SH, Growdon JH, Wurtman RJ, Magil SG, Logue M: Normal plasma choline responses to ingested lecithin. Neurology 30:1226–1229, 1980.
- 29. Savendahl L, Mar MH, Underwood L, Zeisel S: Prolonged fasting

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- results in diminished plasma choline concentration but does not cause liver dysfunction. Am J Clin Nutr 66:622–625, 1997.
- Zeisel SH, Mar MH, Zhou ZW, daCosta KA: Pregnancy and lactation are associated with diminished concentrations of choline and its metabolites in rat liver. J Nutr 125:3049–3054, 1995.
- Welsch F: Studies on accumulation and metabolic fate of (N-Me<sub>3</sub>H)choline in human term placenta fragments. Biochem. Pharmacol 25:1021–1030, 1976.
- Gwee MC, Sim MK: Free choline concentration and cephalin-N-methyltransferase activity in the maternal and foetal liver and placenta of pregnant rats. Clin Exper Pharmacol Physiol 5:649–653, 1978.
- Zeisel SH, Epstein MF, Wurtman RJ: Elevated choline concentration in neonatal plasma. Life Sci 26:1827–1831, 1980.
- Zeisel SH, Wurtman RJ: Developmental changes in rat blood choline concentration. Biochem J 198:565–570, 1981.
- Zeisel SH: Choline: an important nutrient in brain development, liver function and carcinogenesis. J Am Coll Nutr 11:473–481, 1992.
- McMahon KE, Farrell PM: Measurement of free choline concentrations in maternal and neonatal blood by micropyrolysis gas chromatography. Clin Chim Acta 149:1–12, 1985.
- Braun LD, Cornford EM, Oldendorf WH: Newborn rabbit bloodbrain barrier is selectively permeable and differs substantially from the adult. J Neurochem 34:147–152, 1980.
- Cornford EM, Braun LD, Oldendorf WH: Developmental modulations of blood-brain barrier permeability as an indicator of changing nutritional requirements in the brain. Pediatr Res 16:324–328, 1982.
- Garner SC, Mar MH, Zeisel SH: Choline distribution and metabolism in pregnant rats and fetuses are influenced by the choline content of the maternal diet. J Nutr 125:2851–2858, 1995.
- Cermak JM, Blusztajn JK, Meck WH, Williams CL, Fitzgerald CM, RoseneDL, LoyR: Prenatal availability of choline alters the development of acetylcholinesterase in the rat hippocampus. Dev Neurosci 21:94–104, 1999.
- Loy R, Heyer D, Williams CL, Meek WH: Choline-induced spatial memory facilitation correlates with altered distribution and morphology of septal neurons. Adv Exp Med Biol 295:373–382, 1991.
- Meck W, Williams C: Perinatal choline supplementation increases the threshold for chunking in spatial memory. Neuroreport 8:3053– 3059, 1997.
- Meck W, Williams C: Characterization of the facilitative effects of perinatal choline supplementation on timing and temporal memory. Neuroreport 8:2831–2835, 1997.
- Meck W, Williams C: Simultaneous temporal processing is sensitive to prenatal choline availability in mature and aged rats. Neuroreport 8:3045

  –3051, 1997.

- Pyapali G, Turner D, Williams C, Meck W, Swartzwelder HS: Prenatal choline supplementation decreases the threshold for induction of long-term potentiation in young adult rats. J Neurophysiol 79:1790–1796, 1998.
- 46. Williams C, Meck W, Heyer D, Loy R: Hypertrophy of basal forebrain neurons and enhanced visuospatial memory in perinatally choline-supplemented rats. Brain Res 794: 225–238, 1998.
- Meck WH, Williams CL: Choline supplementation during prenatal development reduces proactive interference in spatial memory. Dev Brain Res 118:51–59, 1999.
- 48. Tees RC: The influences of rearing environment and neonatal choline dietary supplementation on spatial learning and memory in adult rats. Behav Brain Res 105:173–188, 1999.
- Tees RC: The influences of sex, rearing environment, and neonatal choline dietary supplementation on spatial and nonspatial learning and memory in adult rats. Dev Psychobiol 35:328–342, 1999.
- Oppenheim RW: Cell death during the development of the nervous system. Ann Rev Neurosci 14:453–501, 1991.
- Thompson CB: Apoptosis in the pathogenesis and treatment of disease. Science 267: 1456–1462, 1995.
- 52. Merry D, Korsmeyer S: Bcl-2 gene family in the nervous system. Annu Rev Neurosci 20: 245–267, 1997.
- Albright CD, Tsai AY, Friedrich CB, Mar MH, Zeisel SH: Choline availability alters embryonic development of the hippocampus and septum in the rat. Dev Brain Res 113:13–20, 1999.
- 54. Albright CD, Tsai AY, Mar MH, Zeisel SH: Choline availability modulates the expression of TGFβ1 and cytoskeletal proteins in the hippocampus of developing rat brain. Neurochem Res 23:751– 758, 1997.
- Albright CD, Zeisel SH: Choline deficiency causes increased localization of TGFb1 signaling proteins and apoptosis in rat liver. Pathobiology 65:264–270, 1997.
- Holmes-McNary MQ, Loy R, Mar MH, Albright CD, Zeisel SH: Apoptosis is induced by choline deficiency in fetal brain and in PC12 cells. Devel Brain Res 101:9–16, 1997.
- Cermak J, Holler T, Jackson D, Blusztajn J: Prenatal availability of choline modifies the development of the hippocampal cholinergic system. FASEB J 12:349–357, 1998.
- Eriksson, P.S., Perfilieva E., Bjork-Eriksson T., Alborn A.M., Nordborg C., Peterson D.A. and Gage F.H. Neurogenesis in the adult human hippocampus. Nat Med 4:1313–1317, 1998.

Received June 2000.