Influence of prematurity and birth weight on the concentration of α-tocopherol in colostrum milk

Influência da prematuridade e do peso ao nascer sobre a concentração de α-tocoferol no leite colostro

Influencia de la prematuridad y del peso al nacer sobre la concentración de α-tocoferol en la leche calostro

Evellyn Câmara Grilo¹, Larissa Queiroz de Lira¹, Roberto Dimenstein¹, Karla Danielly da S. Ribeiro¹

ABSTRACT

Objective: To assess vitamin E levels in the breast milk, analyzing the prematurity and the birth weight influence in α-tocopherol concentration of colostrum milk.

Methods: Cross-sectional study, in which the colostrum was collected from 93 nursing mothers in a public maternity of Natal, Rio Grande do Norte, Northeast Brazil. The newborns were classified based on gestational age and birth weight. The analysis of α-tocopherol in the milk was carried out by high performance liquid chromatography.

Results: The α-tocopherol concentration in the colostrum of lactating women whose children were born at term was $1,093.6 \pm 532.4 \mu g/dL$; for preterm infants, the concentration was $1,321.6 \pm 708.5 \mu g/dL$ ($p=0.109$). In the preterm group, the α-tocopherol concentration in the colostrum of lactating women whose children were born with low and normal birth weight was $1,316.0 \pm 790.7$ and $1,327.2 \pm 655.0 \mu g/dL$, respectively ($p=0.971$). In the term group, the α-tocopherol levels were higher in mothers of children with birth weight >4000g, being $1,821.0 \pm 575.4 \mu g/dL$, compared to $869.5 \pm 532.1 \mu g/dL$ and $1,039.6 \pm 477.5 \mu g/dL$ with low and adequate birth weight, respectively ($p<0.05$).

Conclusions: Prematurity did not influence α-tocopherol levels in the colostrum milk. Mothers who had macrossomic term neonates presented increased α-tocopherol levels. These results indicate that birth weight can influence α-tocopherol levels in the colostrum milk.

Key-words: alpha-tocopherol; colostrum; infant, newborn; infant, premature; birth weight.

RESUMO

Objetivo: Avaliar os níveis de vitamina E no leite materno, analisando-se a influência da prematuridade e do peso ao nascer na concentração de α-tocoferol no colostro.

Métodos: Estudo transversal, com coleta de leite colostro de 93 mulheres atendidas em uma maternidade pública do Rio Grande do Norte. Os recém-nascidos foram classificados com base na idade gestacional e no peso ao nascer. O α-tocoferol no leite foi determinado por cromatografia líquida de alto desempenho.

Resultados: A concentração de α-tocoferol no colostro de lactantes cujos filhos nasceram a termo foi de $1,093,6 \pm 552,4 \mu g/dL$; no caso de recém-nascidos pré-termo, a concentração foi de $1,321,6 \pm 708,5 \mu g/dL$ ($p=0,109$). No grupo pré-termo, as mulheres com neonatos de baixo peso e de peso adequado apresentaram valores de α-tocoferol de $1,316,0 \pm 790,7$ e $1,327,2 \pm 655,0 \mu g/dL$, respectivamente ($p=0,971$), respectivamente. No grupo a termo, houve valores maiores de α-tocoferol em mulheres com crianças de peso ao nascer >4000g ($1,821,0 \pm 575,4 \mu g/dL$), em comparação a $869,5 \pm 532,1 \mu g/dL$ e $1,039,6 \pm 477,5 \mu g/dL$ com baixo peso e peso adequado, respectivamente ($p<0,05$).

Conclusões: Apesar de apresentar tendência de aumento em relação ao grupo a termo, a prematuridade não influenciou o α-tocoferol no colostro. Lactantes que tiveram recém-nascidos a termo com macrossomia apresentaram níveis aumentados de α-tocoferol. Esses resultados indicam que o peso ao nascer pode influenciar o α-tocoferol do colostro.

Palavras-chave: alfa-tocoferol; colostro; recém-nascido; prematuro; peso ao nasc.
RESUMEN

Objetivo: Evaluar los niveles de vitamina E en la leche materna, una vez que los lactantes son considerados grupo de riesgo para la deficiencia de esa vitamina, analizando la influencia de la prematuridad y del peso al nacer en la concentración de α-tocoferol en la leche calostro.

Métodos: Estudio transversal, con colecta de leche calostro de 93 mujeres atendidas en una maternidad pública de Rio Grande do Norte (Brasil). Los recién nacidos fueron clasificados con base en la edad gestacional y en el peso al nacer. El α-tocoferol fue determinado por cromatografía líquida de alta eficiencia.

Resultados: La concentración de α-tocoferol en el calostro de lactantes cuyos hijos nacieron a término fue de 1.093,6±532,4µg/dL; en el caso de recién nacidos pretérmino, la concentración fue de 1.321,6±708,5µg/dL (p=0,109). En el grupo pretérmino, las mujeres con neonatos de bajo peso y de peso adecuado presentaron valores de α-tocoferol de 1.316,0±790,7 y 1.327,2±655,0µg/dL (p=0,971), respectivamente. En el grupo a término, hubo valores superiores de α-tocoferol en mujeres con niños de peso al nacer >4.000g, siendo 1.821,0±575,4µg/dL en comparación a 869,5±532,1µg/dL y 1.039,6±477,5µg/dL con bajo peso y peso adecuado, respectivamente (p>0,05).

Conclusiones: A pesar de presentar tendencia de aumento respecto al grupo a término, la prematuridad no influyó el α-tocoferol en el calostro. Sin embargo, lactantes que tuvieron recién nacidos a término con macrosomía presentaron niveles aumentados de α-tocoferol. Esos resultados indican que el peso al nacer puede influenciar el α-tocoferol de la leche, sugiriendo que neonatos con bajo peso pueden ingerir menos vitamina E cuando amamantados.

Palabras clave: alfa-tocoferol; calostro; recién nacido; prematuro; peso al nacer.

Introduction

Vitamin E is a generic term used to designate eight different molecules: α-, β-, γ- and δ-tocopherol and α-, β-, γ- and δ-tocotrienol. Of these, α-tocopherol is the only isomer related to the vitamin E nutritional requirement. The naturally occurring form of the vitamin is RRR-α-tocopherol. Vitamin E is considered one of the best biological antioxidants because of the protection it offers to plasmatic membranes and low density lipoproteins against oxidation and lipid peroxidation reactions.

Oxidative stress can cause excessive production of reactive oxygen species and/or lead to deficiencies in antioxidant protection systems. This situation is considered to be a contributing factor to the pathophysiologic condition of many disorders. Free radicals are produced by normal metabolic processes and it is believed that their concentrations increase during periods of accentuated metabolic activity, such as pregnancy and birth.

Towards the end of pregnancy, the vitamin E concentration in fetal tissues increases as adipose tissues are built up, since they store around 90% of the vitamin. Newborn infants born preterm have scant adipose tissue and so they have limited vitamin E reserves, making them more susceptible to deficiencies of this micronutrient compared with full term newborns. According to Debier et al., newborn infants are more susceptible to oxidative damage than adults. Breastmilk plays an important role in protecting against oxidative stress, because it contains antioxidant molecules, including the tocopherols, and it is especially important for preterms since it can prevent retinopathy.

A study conducted in Spain found that very low weight neonates had higher hydroperoxide levels than a control group. This observation is linked to the increased degree of oxidative damage suffered by these babies, since hydroperoxides indicate the degree of oxidative stress suffered by erythrocytes. Furthermore, Baydas et al. demonstrated that premature newborn infants and full-term low birth weight infants had lower serum α-tocopherol than a control group.

Breastmilk is the only source of vitamin E that newborns on exclusive breastfeeding have. Research has concluded that the concentration of this micronutrient in breastmilk may be influenced by several different variables, including stage of lactation and gestational age. Studies that have compared the concentration of α-tocopherol in colostrum from mothers of full term and premature neonates have reported conflicting results. Some found that mothers recently-delivered of full term infants had significantly higher α-tocopherol levels in colostrum, but others found no association between gestational age and these levels. A previous study conducted at our center found no significant relationship between α-tocopherol in colostrum and birth weight.

There is evidence that preterm and/or very low weight neonates suffer increased oxidative stress and have lower levels of antioxidants such as vitamin E. In view of this, the relationships between gestational age and birth weight and colostrum α-tocopherol concentration needs to
be investigated, in order to verify whether there are certain
characteristics that can be identified as being linked with
risk of vitamin E deficiency in newborn infants. This study
investigated the influence of neonates’ gestational age and
birth weight on the concentration α-tocopherol in their
mothers’ colostrum milk.

Methods

This was a cross-sectional study that collected samples
of colostrum milk from 93 women seen at the Januário
Cicco teaching maternity hospital in Natal, Rio Grande do
Norte, Brazil, after prior approval by the Research Ethics
Committee at the Universidade Federal do Rio Grande do
Norte. Newborn infants were classified by gestational age
and birth weight. Their mothers were enrolled while still in
the maternity ward if they did not meet any of the following
exclusion criteria: maternal complications (diabetes, hyper-
tension, cancer, liver disease, infectious diseases, gastroin-
testinal tract diseases, heart disease, syphilis, HIV infection,
among others); fetal malformation; multiple births; vitamin
supplementation during pregnancy or postpartum vitamin A “megadose” supplementation, because this comprises
200,000UI of retinyl palmitate combined with 49.4mg of
all-rac-α-tocopherol.

Sample size was calculated using Statcalc (Epi-Info, ver-
sion 3.5.3). According to public health statistics for Natal,
the city has a 7.9% rate of premature live births(18). Since
the maternity hospital at which the study was conducted
hosts an average of 200 births per month, a sample of 90
recently-delivered mothers was needed to ensure a 95%
confidence level. Samples were collected between January
and September of 2010.

The study sample comprised 93 recently-delivered moth-
ers who volunteered after explanation of the study objectives,
signing free and informed consent forms.

Data on infant characteristics were taken from medi-
cal records. The sample of newborn infants was classified
by gestational age, with those born at less than 37 weeks
defined as preterm and those born at 37 to 42 weeks as full
term(19). Newborn infants were also classified by birth weight
as follows: low birth weight (LBW): <2500g; normal birth
weight: ≥2500g and ≤4000g; and macrosomia: >4000g(20).

On the first and second days after delivery, 2mL of colos-
trum was taken in the morning after an overnight fast of 8 to
12 hours. The samples were extracted by manual expression
of just one breast at the start and end of a feed.

The colostrum samples were pooled to avoid major
variations in vitamin levels. The samples were stored at
-20°C in a nitrogen atmosphere until lipid extraction and
α-tocopherol analysis.

The α-tocopherol extraction technique was adapted from
a method described by Ortega et al(21). For one 500μL aliquot
of colostrum, 500μL of 95% ethanol (Merck®, Rahway,
NJ, USA), was used to precipitate proteins and then 2mL
of hexane (Merck®) was used to extract lipids. Next the
samples were mixed for 1 minute and then centrifuged for
10 minutes, after which the hexane extract was transferred
to a new tube. This process was conducted twice, making
4mL of extract, from which a 2mL aliquot was drawn and
evaporated in a nitrogen atmosphere in a 37°C water bath.
For analysis, the dry extract was dissolved in 250μL of HPLC
grade absolute ethanol (Merck®, Rahway, NJ, USA) and
20μL taken for analysis in the High Performance Liquid
Chromatograph.

The concentration of α-tocopherol in samples was then
determined using a chromatograph (Shimadzu Corporation®,
Kyoto, Japan). The system comprises an LC-20 AT Shimadzu
pump connected to a SPD-20A Shimadzu UV-VIS Detector,
with a Shim-pack CLC-ODS (M) 4.6mm x 15cm column
and a computer running the LC solution program (Shimadzu
Corporation®, Kyoto, Japan) to process the data. The mobile
phase used for α-tocopherol analysis was methanol (chro-
matography grade) and water MilliQ® (97:3), in an isocratic
system with a 1.5mL/min flow rate. The wavelength chosen
to monitor absorption was 292nm, and retention time of 8.2
minutes was obtained.

Alpha-tocopherol was identified and quantified in samples
by comparing the area of the chromatographic profile with
the area for standard α-tocopherol (Sigma®, St. Louis, MO,
USA). The concentration of the standard was confirmed
against the specific extinction coefficient in absolute ethanol
for α-tocopherol (ε 1%, 1cm=75.8 to 292nm)(22).

Statistical analysis was performed using the open
source statistical software R, version 2.15. The results for
α-tocopherol concentrations in colostrum were expressed as
means with standard deviations. Numerical variables were
shown not to be normal, by the Kolmogorov-Smirnov test,
characterizing the samples as parametric.

The preterm and full term subsets were compared for the
response variable α-tocopherol concentration in colostrum
using analysis of variance (ANOVA), with Tukey’s post hoc
test. This statistical treatment evaluated both the main ef-
ffects of gestational age and birth weight on α-tocopherol

Evellyn Câmara Grilo et al

Rev Paul Pediatr 2013;31(4):473-9.

475
concentration in colostrum and the interaction between the two variables. Differences were considered significant when $p<0.05$.

### Results

Twenty-two neonates were defined as preterm and 71 were defined as full term. Half (50%) of the preterm neonates also had low birth weight and none of them had macrosomia. Just 7.0% of the full term neonates had low birth weight and 8.5% had macrosomia (Table 1). Overall, 16 of the 93 newborn infants had low birth weight, 71 had normal birth weights and six were abnormally heavy. Mean birth weight for the preterm subset was 2560±710g and mean full term birth weight was 3260±500g ($p<0.001$).

Mean $\alpha$-tocopherol concentration in colostrum for the whole sample ($n=93$) was 1,147.6±582.9µg/dL. Mean concentration of $\alpha$-tocopherol in colostrum from breastfeeding mothers whose children were born full term was 1,093.6±532.4µg/dL; and the mean for mothers of preterm newborn infants was 1,321.6±708.5µg/dL ($p>0.05$) (Figure 1).

When $\alpha$-tocopherol concentration was further broken down by birth weight, it was found that mean $\alpha$-tocopherol concentration in the colostrum of breastfeeding mothers whose children were born preterm and with low weight was 1,316.0±790.7µg/dL; and that mothers whose children were preterm but born with normal weight had a mean concentration of 1327.2±655.0µg/dL, with no significant difference between them ($p>0.05$) (Figure 2). The colostrum of women whose children were in the full term group and were born with macrosomia had a mean $\alpha$-tocopherol concentration of 1,821.0±575.4µg/dL, which was higher than for women with full term children born at low weight, with a concentration of 869.5±532.1µg/dL ($p<0.05$), and higher than for mothers of full term children born at normal weight, whose concentration was 1,039.6±477.5 ($p<0.05$) (Figure 2).

Thus, ANOVA indicated a main effect from the variable birth weight ($F=5.81; p<0.05$). However, gestational age did

### Table 1 - Characteristics of the sample of newborn infants from the Januário Cicco teaching maternity hospital, by birth weight and gestational age, 2010

| Birth weight                  | Preterm (n=22) | Full term (n=71) |
|------------------------------|----------------|------------------|
| Low weight (n=16)            | 11 (50.0)      | 5 (7.0)          |
| Normal weight (n=71)         | 11 (50.0)      | 60 (84.5)        |
| Macrosomia (n=6)             | 0 (0)          | 6 (8.5)          |

$\alpha$-tocopherol concentration in colostrum of breastfeeding mothers, by gestational age of infants. There was no statistically significant difference between the full term and preterm groups according to Student’s $t$ test ($p>0.05$)

$\alpha$-tocopherol concentration in colostrum of breastfeeding mothers, by birth weight and gestational age. *Statistically significant difference between subset with macrosomia and other full term subsets (low birth weight and normal birth weight) according to ANOVA with Tukey’s post hoc test $p<0.05$
not exhibit a main effect (F=2.87; p>0.05), and there was no statistically significant interaction between gestational age and birth weight (F=0.21; p>0.05).

**Discussion**

Some studies have found that macrosomic, premature and/or low weight newborn infants have lower vitamin E concentrations in plasma or the umbilical cord\(^{(17,23-25)}\). It is therefore essential that infants in these groups that could be at risk of deficiency receive their intake of this micronutrient in breastfeeding in order to protect them from oxygen toxicity and to stimulate immune system development\(^{(26)}\).

The mechanisms by which \(\alpha\)-tocopherol is transported to the mammary gland are not well understood, but there is consensus that the colostrum-secretion cells increase \(\alpha\)-tocopherol uptake at the end of pregnancy, during labor and at the start of lactation. Schweigert\(^{(27)}\) studied transport of liposoluble vitamins and lipids in cows and suggested that cholesterol and vitamin E may be transferred to colostrum by a secretory cell transport system that is specific to LDL.

Considering that colostrum is an exclusively breastfed infant's sole source of nutrients, it is important to determine the concentration of vitamin E in breastmilk. The mean \(\alpha\)-tocopherol concentration in the colostrum of the breastfeeding mothers studied here was in line with the results of studies of similar populations, such as a study by Dimenstein *et al*\(^{(28)}\) who reported a figure of 1,155±811 µg/dL and a study by Garcia *et al*\(^{(29)}\) who reported an \(\alpha\)-tocopherol concentration of 1,206±859 µg/dL.

When results were broken down by gestational age, it was found that the \(\alpha\)-tocopherol concentration in the colostrum of recently-delivered mothers whose babies were born prematurely was similar to the results of similar studies undertaken in Germany\((1,450 µg/dL)\)\(^{(13)}\) and Spain (from 1,292.1 to 1,722.8 µg/dL)\(^{(12)}\). However, our result is higher than the mean concentration observed in a study in China \((777.84 µg/dL)\)\(^{(14)}\). The mean \(\alpha\)-tocopherol concentration in colostrum from mothers of full term infants observed here was similar to the result from a German population \((1,140 µg/dL)\)\(^{(13)}\), higher than the concentration found in a Chinese population \((741.67 µg/dL)\)\(^{(14)}\) and lower than the concentration observed in colostrum from a Spanish population (from 2,153.5 to 2,584.2 µg/dL)\(^{(12)}\).

When \(\alpha\)-tocopherol concentration in colostrum from mothers of full term infants was compared with the concentration in colostrum from mothers of preterms there was no statistical difference. Similar results have been published by Haug *et al*\(^{(15)}\) and Zheng *et al*\(^{(14)}\). Only the Spanish study found a significant difference between their full term and preterm groups, with higher \(\alpha\)-tocopherol concentrations in colostrum from mothers whose children were born full term (2,154 to 2,584 µg/dL) compared with mothers of preterm infants (1,292 to 1,723 µg/dL). It should be pointed out, however, that only 30 women were enrolled on the Spanish study, 15 with full term and 15 with preterm infants\(^{(12)}\).

It has been shown that the plasma lipoprotein profile changes during colostrum formation, with the observation that the LDL fraction transports around 20% of lipids 4 weeks before birth but just 4% at the time of birth\(^{(27)}\). On the basis of this evidence, it has been hypothesized that the imminence of birth causes biochemical changes to lipoproteins irrespective of the duration of gestation.

It is therefore assumed that recently-delivered mothers whose gestations lasted less than 37 weeks and/or who gave birth to low weight infants have a similar biochemical pattern of \(\alpha\)-tocopherol transport to the mammary glands to that of mothers whose children were born full-term and/or with normal birth weight, which may be explained by an accelerated physiological adaptation, in response to the imminence of birth. This would explain the absence of significant differences in \(\alpha\)-tocopherol concentrations between subsets divided by gestational age and also after further subdivision by birth weight and gestational age (into low birth weight and normal birth weight subsets).

However, \(\alpha\)-tocopherol concentration was significantly higher in the colostrum of breastfeeding mothers whose children were born full-term and macrosomic than in the normal and low birth weight subsets of the full-term group (Figure 2), which contrasts with the results of a study conducted in the same city as this one (Natal), in which birth weight had no influence on vitamin levels\(^{(15)}\).

Studies of the kinetics and metabolism of \(\text{RRR-}\alpha\)-tocopherol have found that 99% of the average quantity of this compound that is estimated to exist in the human body is contained in adipose tissues. High rates of \(\alpha\)-tocopherol transfer between adipose tissue and plasma lipoproteins were also observed\(^{(30)}\). On this basis, it is supposed that when the body’s \(\alpha\)-tocopherol stocks are high there will be a higher concentrations of vitamins in lipoproteins, which in turn...
in erythrocytes and oxidative stress in erythrocytes - Coenzyme Q concentration and total antioxidant capacity of

The fact that both this metabolic condition and excess weight gain either before or during pregnancy are linked with macrosomia\(^{(32)}\), suggests that the increased \(\alpha\)-tocopherol concentrations in the colostrum of recently-delivered mothers whose children were born macrosomic is the result of their greater accumulation of adipose tissue which in turn leads to increased maternal stocks of the vitamin and, as a consequence, to greater availability to the mammary glands. Further studies should be conducted to test for an association between gestational body mass index and the concentrations of \(\alpha\)-tocopherol in colostrum and breastmilk and for a relationship between gestational weight gain and vitamin E concentrations.

In view of the findings of this study, it is believed that the colostrum received by these newborn infants may be more beneficial in terms of nutritional vitamin E intake. This micronutrient is essential for macrosomic newborns who may be more susceptible to deficiency of the vitamin because of lower plasma \(\alpha\)-tocopherol concentrations and impaired antioxidant capacity, considering the enzymatic mechanisms involved\(^{(22,32)}\). The authors of a study of newborn infants with macrosomia that was conducted in Algeria concluded that excess weight is potentially a factor that increases oxidative stress\(^{(32)}\).

The vitamin E content of human milk can affect the biochemical status of this vitamin in breastfed babies\(^{(33)}\). Newborn infants with macrosomia therefore benefit more in terms of vitamin E transfer when fed on their own mothers’ milk, since they are potentially at risk of increased oxidative stress\(^{(32)}\).

Certain elements of this study could be considered limitations, including the lack of an assessment of maternal nutritional status, by means of serum \(\alpha\)-tocopherol assay, and the lack of a dietary assessment of the breastfeeding mothers enrolled on the study.

Studies such as this one that investigate the influence of neonatal characteristics on \(\alpha\)-tocopherol in colostrum are important to delineate subsets at risk of vitamin E deficiency, considering the vitamin supply provided to newborn infants in breastmilk, and to widen understanding of the influence that factors such as gestational age have on the adaptive capacity of mechanisms that transfer \(\alpha\)-tocopherol to the mammary glands.

References

1. Traber MG. Vitamin E. In: Bowman BA, Russell RM, editors. Present knowledge in nutrition. 9th ed. Washington: ILSI Press; 2006. p. 211-9.
2. Batista ES, Costa AG, Pinheiro-Sant’Ana HM. Adding vitamin E to foods: implications for the foods and for human health. Rev Nutr 2007;20:525-35.
3. Sánchez-Pérez A, Delgado-Zamarreño MM, Bustamante-Rangel M, Hernández-Méndez J. Automated analysis of vitamin E isomers in vegetable oils by continuous membrane extraction and liquid chromatography-electrochemical detection. J Chromatogr A 2000;881:229-41.
4. Erdem M, Harma M, Harma IM, Arikan I, Barut A. Comparative study of oxidative stress in maternal blood with that of cord blood and maternal milk. Arch Gynecol Obstet 2012:285:371-5.
5. Wey M. Vitamina E no plasma de recém-nascidos de pré-termo de muito baixo peso no primeiro mês de vida. Relação com a vitamina E recebida [tese de doutorado]. Botucatu (SP): Unesp; 2008.
6. Debier C, Pottier J, Goffe CH, Larondelle Y. Present knowledge and unexpected behaviours of vitamins A and E in colostrum and milk. Livest Prod Sci 2005;98:135-47.
7. De Azeredo VB, Trugo NM. Retinol, carotenoids, and tocopherols in the milk of lactating adolescents and relationships with plasma concentrations. Nutrition 2008;24:133-9.
8. Duda G, Nogala-Kalucka M, Karwowska W, Kupczyk B, Lampart-Szczapa E. Influence of the lactating women diet on the concentration of the lipophilic vitamins in human milk. Pak J Nutr 2009;8:629-34.
9. Ochoa JJ, Contreras-Chova F, Muñoz S, Araujo-Nepomuceno E, Bonillo A, Molina-Carbollo A et al. Fluidity and oxidative stress in erythrocytes from very low birth weight infants during their first 7 days of life. Free Radic Res 2007;41:1035-40.
10. Baydas G, Karatas F, Gursu MF, Bozkurt HA, Ilhan N, Yasar A et al. Antioxidant vitamin levels in term and preterm infants and their relation to maternal vitamin status. Arch Med Res 2002;33:276-80.
11. Macias C, Schweigert FJ. Changes in the concentration of carotenoids, vitamin A, alpha-tocopherol and total lipids in human milk throughout early lactation. Ann Nutr Metab 2001;45:82-5.
12. Quiles JL, Ochoa JJ, Ramirez-Tortosa MC, Linde J, Bompard S, Battino M et al. Coenzyme Q concentration and total antioxidant capacity of human milk at different stages of lactation in mothers of preterm and full-term infants. Free Radic Res 2006;40:199-206.
13. Haug M, Laubach C, Burke M, Harzer G. Vitamin E in human milk from mothers of preterm and term infants. J Pediatr Gastro Nutr 1987;6:605-9.
14. Zheng MC, Zhang GF, Zhou LS, Guo XG, Quan YF. Alpha-tocopherol concentrations in human milk from mothers of preterm and full-term infants in China. Biomed Environ Sci 1993;6:259-64.
15. Garcia LR. Avaliação da suplementação materna com megadose de vitamina A sobre os níveis de retinol e alfa-tocoferol no colostro [tese de mestrado]. Natal (RN): UFRN; 2009.
16. Kositamongkol S, Suthutvoravut U, Chongviriyaphan N, Feungpean B, Nuntnarumit P. Vitamin A and E status in very low birth weight infants during their first 72 h of life. Free Radic Res 2003;37:317-22.
17. Brasil - Ministério da Saúde - DATASUS [homepage on the Internet]. In view of the findings of this study, it is believed that the colostrum received by these newborn infants may be more beneficial in terms of nutritional vitamin E intake. This micronutrient is essential for macrosomic newborns who may be more susceptible to deficiency of the vitamin because of lower plasma \(\alpha\)-tocopherol concentrations and impaired antioxidant capacity, considering the enzymatic mechanisms involved\(^{(22,32)}\). The authors of a study of newborn infants with macrosomia that was conducted in Algeria concluded that excess weight is potentially a factor that increases oxidative stress\(^{(32)}\).

The vitamin E content of human milk can affect the biochemical status of this vitamin in breastfed babies\(^{(33)}\). Newborn infants with macrosomia therefore benefit more in terms of vitamin E transfer when fed on their own mothers’ milk, since they are potentially at risk of increased oxidative stress\(^{(32)}\).

Certain elements of this study could be considered limitations, including the lack of an assessment of maternal nutritional status, by means of serum \(\alpha\)-tocopherol assay, and the lack of a dietary assessment of the breastfeeding mothers enrolled on the study.

Studies such as this one that investigate the influence of neonatal characteristics on \(\alpha\)-tocopherol in colostrum are important to delineate subsets at risk of vitamin E deficiency, considering the vitamin supply provided to newborn infants in breastmilk, and to widen understanding of the influence that factors such as gestational age have on the adaptive capacity of mechanisms that transfer \(\alpha\)-tocopherol to the mammary glands.
19. World Health Organization. Neonatal and perinatal mortality: country, regional and global estimates. Geneva: WHO; 2006.
20. Strutz KL, Richardson LJ, Hussey JM. Preconception health trajectories and birth weight in a national prospective cohort. J Adolesc Health 2012;51:629-36.
21. Ortega RM, López-Sobaier AM, Martínez RM, Andrés P, Quintas ME. Influence of smoking on vitamin E status during the third trimester of pregnancy and on breast-milk tocopherol concentrations in Spanish women. Am J Clin Nutr 1998;68:662-7.
22. Nierenberg DW, Nann SL. A method for determining concentrations of retinol, tocopherol, and five carotenoids in human plasma and tissue samples. Am J Clin Nutr 1992;56:417-26.
23. Grissa O, Atègbo JM, Yessoufou A, Tabka Z, Miled A, Jerbi M et al. Antioxidant status and circulating lipids are altered in human gestational diabetes and macrosomia. Transl Res 2007;150:164-71.
24. Kumar A, Ranjan R, Basu S, Khanna HD, Bhargava V. Antioxidant levels in cord blood of low birth weight newborns. Indian Pediatr 2008;45:583-5.
25. Saker M, Mokhtar NS, Merzouk SA, Merzouk H, Belarbi B, Narce M. Oxidant and antioxidant status in mothers and their newborns according to birthweight. Eur J Obstet Gynecol Reprod Biol 2008;141:95-9.
26. Debier C, Larondele Y. Vitamins A and E: metabolism, roles and transfer to offspring. Br J Nutr 2005;93:153-74.
27. Schweigert FJ. Effect of gestation and lactation on lipoprotein pattern and composition in dairy cows. J Anim Physiol Anim Nutr (Berl) 1990;63:75-83.
28. Dimenstein R, Lira L, Medeiros AC, Cunha LR, Stamford TL. Efeito da suplementação com vitamina E sobre a concentração de alfa-tocoferol no colostro humano. Rev Panam Salud Publica 2011;29:399-403.
29. García L, Ribeiro K, Araújo K, Pires J, Azevedo G, Dimenstein R. Alpha-tocopherol concentration in the colostrum of nursing women supplemented with retinyl palmitate and alpha-tocopherol. J Hum Nutr Diet 2010;23:529-34.
30. Novotny JA, Fadel JG, Holstege DM, Furr HC, Clifford AJ. This kinetic, bioavailability, and metabolism study of RRR-α-tocopherol in healthy adults suggests lower intake requirements than previous estimates. J Nutr 2012;142:2105-11.
31. Amorim MM, Leite DF, Gadelha TG, Muniz AG, Melo AS, Rocha AM. Risk factors for macrosomia in newborns at a school-maternity in Northeast of Brazil. Rev Bras Ginecol Obstet 2009;31:241-8.
32. Haddouche M, Aribi M, Moullessehoul S, Smahi MC, Lammani M, Benyoucef M. Alteration of antioxidant defense status precedes humoral immune response abnormalities in macrosomia. Med Sci Monit 2011;17:CR650-6.
33. Romeu-Nadal M, Morera-Pons S, Castellote AI, López-Sabater MC. Determination of gamma- and alpha-tocopherols in human milk by a direct high-performance liquid chromatographic method with UV-vis detection and comparison with evaporative light scattering detection. J Chromatogr A 2006;1114:132-7.