Special Article

Systematic review of the concentrations of oligosaccharides in human milk

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Context: *Oligosaccharides are the third largest solid component in human milk. These diverse compounds are thought to have numerous beneficial functions in infants, including protection against infectious diseases. The structures of more than 100 oligosaccharides in human milk have been elucidated so far.* ***Objective:*** *The aim of this review was to identify the main factors that affect the concentrations of oligosaccharides in human milk and to determine whether it is possible to calculate representative and reliable mean concentrations.* ***Data Sources:*** *A comprehensive literature search on oligosaccharide concentrations in human milk was performed in 6 electronic databases: BIOSIS, Current Contents Search, Embase, Lancet Titles, MEDLINE and PubMed.* ***Study Selection:*** *The initial search resulted in 1363 hits. After the elimination of duplicates, the literature was screened. The application of strict inclusion criteria resulted in 21 articles selected.* ***Data Extraction:*** *Oligosaccharide concentrations, both mean values and single values, reported in the literature were sorted by gestational age, secretor status of mothers, and defined lactation periods.* ***Results:*** *Mean concentrations, including confidence limits, of 33 neutral and acidic oligosaccharides reported could be calculated. Concentrations of oligosaccharides in human milk show variations that are dependent on both the secretor type of the mother and the lactation period as examined by analyses of variance. In addition, large interlaboratory variations in the data were observed.* ***Conclusions:*** *Worldwide interlaboratory quantitative analyses of identical milk samples would be required to identify the most reliable methods of determining concentrations of oligosaccharides in human milk. The data presented here contribute to the current knowledge about the composition and quantities of oligosaccharides in human milk and may foster greater understanding of the biological functions of these compounds.*

INTRODUCTION

Human milk oligosaccharides (HMOS) represent approximately 20% of the total carbohydrate content of human milk and are the third largest solid component, present at concentrations of up to 20 g/L or more in colostrum.[1] A wide variety of oligosaccharides are synthesized in the mammary gland by the action of specific

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glycosyltransferases that sequentially add *N*-acetylglucosamine, galactose, fucose, and *N*-acetylneuraminic acid to the basic acceptor molecule, lactose.[2] Currently, approximately 150 oligosaccharide structures in human milk have been elucidated,[3–12] and many more are present, at least in small quantities.[13] The presence of glycosyltransferases in the mammary gland is genetically determined. With the exception of fucosyltransferases, glycosyltransferases are common to all mothers. Fucosyltransferase 2 (corresponding to the secretor enzyme) transfers fucose residues as α1,2 linkages to acceptor molecules,[14] whereas fucosyltransferase 3 (ie, the Lewis enzyme) transfers fucose residues predominantly as α1,4 linkages, leading to different patterns of fucosylated HMOS.[2,15,16]

Since the infant's intestine cannot digest HMOS,[17,18] the possible physiological role of HMOS was the subject of intensive research for more than 60 years. The prebiotic effect of HMOS was the first function discovered.[19] There is increasing evidence that HMOS have other functions as well, as indicated by the direct interaction of HMOS with bacterial or protozoan lectins as well as with epithelial or immune cell receptors.[20–23]

Historically, the focus of HMOS research has been the development of methods to analyze the structure of HMOS. Since the biological functions of HMOS depend not only on the specific structure of an HMOS but also on the quantity of the HMOS present, different methods of quantitative measurements have also been developed[24–30] to understand both the functional properties of HMOS and the role of HMOS in the development and well-being of the breastfed infant.

The wide variability in the pattern of HMOS present in different individuals is well known. There are several studies indicating that factors such as polymorphism of the Lewis and secretor genes[2,31–33] as well as the period of lactation[31,34,35] determine both the pattern and the quantities of HMOS present in milk. Since data on human milk are often used as the basis for establishing infant feeding regimens, this review aimed to determine whether standard concentrations of individual HMOS in human milk can be calculated and to identify which confounding factors must be considered. Such data would be important for calculating the physiological intakes of different HMOS, which have various biological functions. The findings presented here may serve as a basis for the application of oligosaccharides with structures identical to or different from those of HMOS.

METHODS

Literature search and literature selection

Comprehensive electronic literature searches were performed (August 2015) to find all relevant literature reporting quantitative data for HMOS. The databases BIOSIS, Current Contents Search, Embase, Lancet Titles, MEDLINE, and PubMed were screened by CM and GB. The following search strategy was applied: "(human OR breast) AND (milk) AND (oligosaccharide*) AND (quantification OR concentration* OR content*)." The only language limitation was that the titles and the abstracts had to be published in English.

The full texts of all articles that appeared to report quantitative data for HMOS were evaluated by applying the predetermined PICOS (Population, Intervention, Comparator, Outcome, and Study design) criteria described in Table 1.

Design of the systematic review

Concentrations of individual HMOS reported in the literature vary widely. Such variation may be attributable to the general biological variability as well as to the influence of gestational age, Lewis blood group, secretor status of the mother, lactation period, and analytical methods. Although the effect of gestational age on HMOS concentrations has not been clearly demonstrated in previous studies, an influence cannot be excluded. Gidrewicz and Fenton[36] reported significantly higher amounts of oligosaccharides in preterm milk than in term milk during days 4 to 7 postpartum. Furthermore, lacto-*N*-tetraose (LNT), a nonfucosylated core HMOS, was more abundant in the milk of women who delivered preterm.[37] Therefore, in the present review, documentation of the gestational age was an inclusion criterion, and data from term and preterm milks were analyzed and presented separately.

Four different human milk types, characterized by the presence or absence of specific fucosylated oligosaccharides, have been described. The presence or absence of fucosylated oligosaccharides depends on the Lewis blood type of the lactating woman.[2,16] Furthermore, the 4 milk types contain different amounts of HMOS; these amounts are not directly affected by either of the 2 fucosyltransferases, ie, the secretor enzyme and the Lewis enzyme.[31–33] There are only a few HMOS studies that report the Lewis blood group of the donors, and even fewer compare HMOS concentrations of the different milk types. However, several authors report the secretor status of the mothers without indicating the exact milk group (secretors produce milk belonging to groups 1 and 3, whereas nonsecretors produce milk belonging to groups 2 and 4). For the purpose of this systematic review, HMOS data was included if the secretor status of the mothers was reported, even if the exact milk group was not.

Several studies have demonstrated the influence of the lactation period on HMOS concentrations.[31,34,35,38,39] There is no international consensus about how to define

Table 1 **PICOS criteria for inclusion and exclusion of studies**

Parameter	Inclusion criteria	Exclusion criteria
Population	Milk samples from individual healthy mothers Documented duration of pregnancy Documented lactation days Individuals with defined secretor status in the case of neutral HMOS	Animal studies Pooled milk samples Milk samples at lactation periods not fitting the lactation periods defined in this review
Intervention	None	None
Comparator	None	None
Outcomes	Absolute concentrations of a single HMOS Mean values Single HMOS concentration values with n ≥ 2 at given gestational age, secretor status, and lactation period (ie, values from at least 2 mothers were required)	Relative concentrations Concentrations of mixtures of HMOS Median values Mass spectrometry data based on universal calibration
Study design	Original articles from peer-reviewed journals	Abstracts, monographs, review articles Studies with data already reported

Abbreviations: HMOS, human milk oligosaccharides.

lactation periods. The following 6 lactation periods were defined on the basis of the most commonly used definition in the articles selected for the present review: 0 to 4 days postpartum (colostrum), 5 to 10 days postpartum (transitional milk), 11 to 30 days postpartum, 31 to 60 days postpartum, 61 to 100 days postpartum, and more than 100 days postpartum. The effects of the lactation periods were examined in conjunction with the documented gestational age and with secretor status, respectively.

The above-mentioned variability in concentrations of different HMOS is probably also attributable to the various methods used to quantify free oligosaccharides in human milk. Although details about quantification methods were recorded for this review, the effect of these methods could not be tested because nearly every study used a unique methodology with regard to the preparation, derivatization, and separation of samples and the detection and quantification of HMOS.

Data extraction and data processing

From the studies selected, absolute concentrations of HMOS were extracted if the HMOS structures could be identified on the basis of several recent reports.[6,8–10]

The nomenclature according to Urashima et al.[8] served as a guideline for the designation of oligosaccharide structures. The structures of 3 HMOS reported in 1 study were identified after the main author was contacted.[34]

Most reports selected indicated the secretor status of the mothers as determined either by serological tests or by interpretation of the HMOS patterns found in human milk. The secretor status of the mothers in 2 studies[40,41] could be determined from the individual HMOS patterns reported. The secretor-positive status of the donors in 1 study[42] could be identified by combining the HMOS patterns and the concentration ranges reported. Erney et al.[35] did not mention the secretor status of the mothers. Nevertheless, the concentrations of α1,2 fucosylated HMOS were included because the authors indicated that zero values were excluded from the calculations of mean values. This was interpreted to mean that only secretor mothers were taken into account. Furthermore, data of other HMOS analyzed in milks from mothers from Latin America were also included in the present review because the HMOS patterns reported showed that these mothers were almost exclusively secretor donors.[35]

In most cases, concentrations of the HMOS were reported as grams per liter or milligrams per liter. Concentrations reported as millimoles per liter were converted to grams per liter by multiplying by the average molecular mass of the corresponding HMOS. Data presented in non-numerical form in graphs were measured manually with rulers, using multiple enlarged figures that were translated into numbers.[43–46] Two mean values of the acidic HMOS LST a were reported as "not determined."[34] During data processing, these expressions were transduced to zero values.

Usually, authors reported mean values as well as standard deviations, number of samples, and number of mothers. When concentrations from single milk samples were reported, mean values were calculated, with data sorted according to gestational age, secretor group, and lactation period as defined in the present review.[40,41,47,48] When authors reported several mean values within a certain lactation period at different days postpartum, the corresponding mean of the entire period was calculated. As a result, the different numbers of mothers or samples were not taken into account. Similarly, means reported for milk groups 1 and 3 as well as for milk groups 2 and 4 were averaged to yield the corresponding data for secretors and nonsecretors, respectively.[33,41] In addition, means reported from different world regions at given lactation times were also averaged.[35] As such, each study was represented at a given lactation period with 1 mean value.

Prior to the analyses of variance, outlier diagnostic tests were conducted using a cutoff value of 3.0. The rare studies that did not meet this criterion were excluded from the data processing. Furthermore, the prerequisites for the analysis of variance—homoscedasticity (Brown-Forsythe test) and homogeneity of variance (Shapiro–Wilk test)—were checked. These requirements were fulfilled in all cases. Data processing was performed with the software SAS, version 9.3, applying the procedures GLM (general linear models), MIXED (mixed linear models), and ROBUSTREG (robust regression) (using option outlier diagnostics).[49]

A one-way analysis of variance followed by a Tukey test of the means was applied to the factor "lactation period" as defined in this section. Tukey tests were carried out on a level of significance of $\alpha = 5\%$. In addition, 95% confidence limits were calculated for all mean values. All analyses were performed separately for term and preterm milks from secretor mothers. A two-factor analysis of variance with the factors "lactation period" and "secretor status" was performed when authors provided data for defined lactation periods from secretor as well as from nonsecretor mothers. Because these two-factor analyses were based on unequal numbers of observations, adjusted means (so-called least squares means) had to be calculated. Out of 24 HMOS tested, only 2 (3′-FL in term milks and DF-LNH II in preterm milks) showed significant interactions between the factors "lactation period" and "secretor status" detected. Hence, both factors were largely independent.

RESULTS

The application of a relatively simple search strategy using the keywords "human milk oligosaccharides" in combination with "quantification" in the most common medical and life science databases yielded 1363 hits (Figure 1). After duplicates were removed, the titles and abstracts of the articles were screened. Reports with irrelevant topics as well as monographs, meeting abstracts, and review articles were excluded, resulting in 129 articles. The corresponding full-text articles were examined intensively. These included all relevant articles already known to the authors, with 1 exception.[39] Forty-eight research studies reported quantitative data on single HMOS compounds. Of these, 27 had to be excluded because 1 or several inclusion criteria had not been fulfilled[24,25,27–29,37,38,50–69] (see Table S1 in the Supporting Information online). Several articles could not be included because the secretor status of the mothers could not be determined.[24,29,38,58,61,67] Finally, 21 studies meeting the predefined inclusion criteria were selected.[31–35,39–48,70–75] Table 2[31–35,39,40–48,70–75]

lists the main characteristics of these studies, the countries of the mothers, the secretor status, the gestational stage, and the quantification methods applied. The secretor status was either reported or could be deduced from 15 of the 21 studies selected, as shown in Table 2. The remaining 6 studies, which investigated acidic HMOS, lacked information on the secretor status of the mothers but were included since the effect of secretor status on the concentrations of acidic compounds is of minor importance, as noted in the next section. From all these reports, absolute concentrations of 33 HMOS (Figure 2) were extracted. The 33 oligosaccharides were classified according to their degree of fucosylation and sialylation: 22 were neutral, nonsialylated HMOS and 11 were acidic HMOS, half of which also contained fucose residues.

Concentrations of oligosaccharides in human milk

Table 3[31–35,42,43,45,46,48,71,72,74] shows the concentrations of neutral oligosaccharides in milk samples of secretor mothers, as most data reported were for HMOS from secretor milks. In addition, the reference numbers of the studies, the numbers of mothers and samples, and the corresponding lactation periods are listed. Mean values and confidence intervals of the HMOS concentrations are shown separately for term and preterm milk samples. Four studies reported the HMOS concentrations determined in preterm milks,[33,41,46,75] whereas 18 reported data for term milks. The total mean concentrations of 20 neutral HMOS from term milk and 17 neutral HMOS from preterm milk were 14.8 g/L and 11.6 g/L, respectively. The difference in mean values can be explained by the fact that 7 of the HMOS quantified were different and by an unusually high concentration of TF-LNH in term milks.[32,34] In most cases, the amounts of individual HMOS seemed to reach similar levels, although mothers with term infants tended to exhibit higher concentrations of LNnH, LNFP III, LNFP V, LNDFH II, and TF-LNH than mothers with preterm infants. However, owing to the particularly high variation between the 4 studies reporting preterm milk data, analyses of variance could not be performed to confirm this. In order to systematically examine the effect of gestational age, additional studies that determine HMOS concentrations both in term and in preterm milks are required so that the confounding effects of the factor method of quantification are eliminated. Only 1 study met this requirement, but it does not specify the secretor status of the milk donors.[75] Nine neutral HMOS from term milks, usually small-sized structures, were quantified in 5 or more studies. Mean concentrations of these compounds ranged between 0.14 g/L and 2.74 g/L, with the α1,2 fucosylated HMOS 2′-FL,
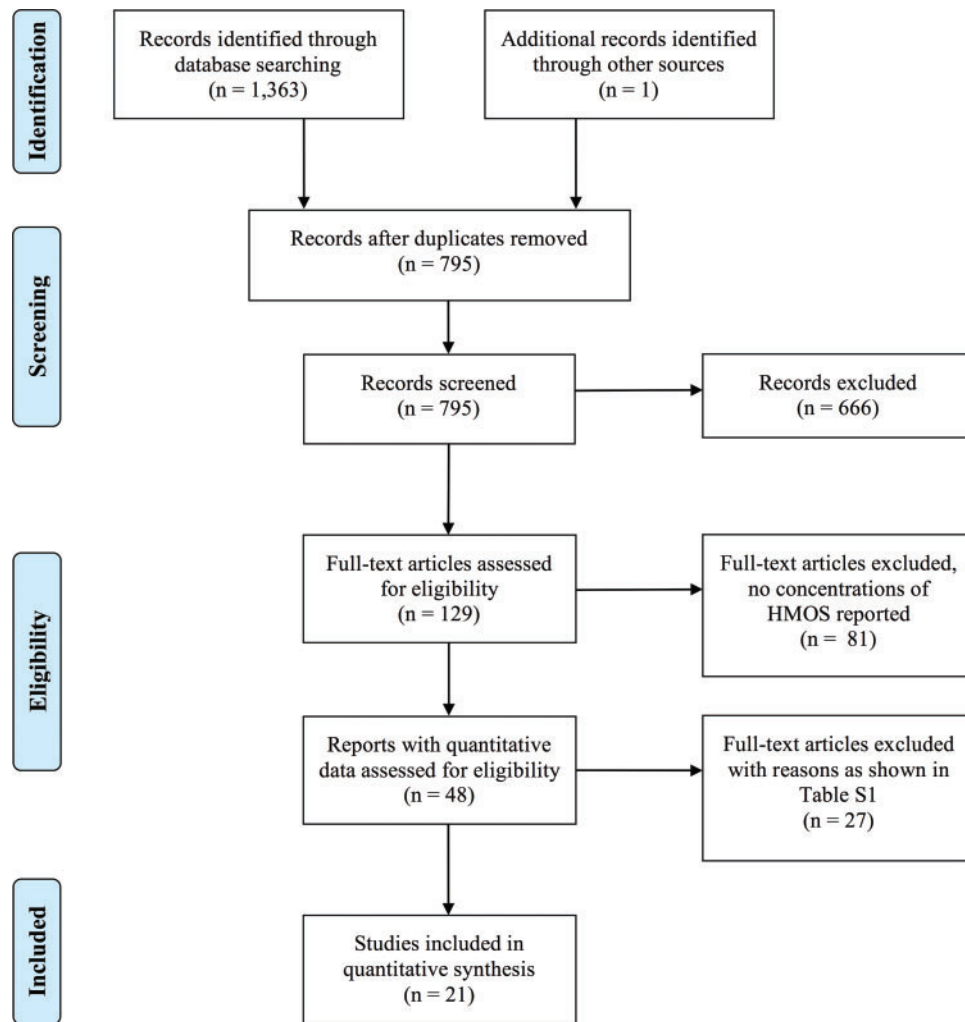
Figure 1 **Flow diagram of the literature search process.**

LNFP I, and LNDFH I as well as the nonfucosylated HMOS LNT and LNnT being the dominant oligosaccharides in secretor milks. In addition, the confidence intervals were narrower than those for the corresponding HMOS from preterm milks. The high concentrations of the larger neutral HMOS DF-LNH II and TF-LNH in term milks were reported in only 1 or 2 studies each.[32,34]

Table 4[31,33,34,40,45,46,71,72] lists the mean concentrations plus confidence intervals of 7 acidic HMOS from secretor mothers, reported in 1 to 6 studies. The sum of all mean concentrations of 7 acidic HMOS was 2.1 g/L for term milks (corresponding to the first 100 days postpartum) and 3.3 g/L for preterm milks (corresponding to the first 30 days postpartum). Generally, concentrations of acidic HMOS, particularly those of LST a and LST c, seem to be higher in preterm milks than in term milks. As was the case with neutral HMOS, this could not be confirmed by analyses of variance. In term milks, 6′-SL and DS-LNT were the dominant acidic HMOS.

Since the effect of secretor status on concentrations of acidic HMOS has not been examined in the literature[31,33,71,72] and seems to be less important than the effect on neutral HMOS, data for HMOS without defined secretor status are presented in Table 5.[31,33,34,39,40,44–47,70,72,73,75] In addition to data for the 7 acidic HMOS from term milk shown in Table 4, data for 4 fucosylated acidic HMOS were available. The major acidic HMOS were 6′-SL (10 studies) and DS-LNT (7 studies). To allow adequate evaluation of the mean concentration data, the number of studies that reported data for each HMOS is shown. In most cases, the corresponding mean values seem to be similar, independent of the secretor status of the donors. However, in term milks, the average concentrations of 6′-SL and LST b seem to be lower when 10 instead of 6 studies and when 6 instead of 3 studies reported data, respectively. These contradictory results can be explained by outliers detected by the software when a larger number of studies was included. Thus, 6′-SL concentrations reported by Thurl et al.[31]

Table 2 **Studies included in the systematic review**

Reference	Mothers				Quantification methods	Type of HMOS quantified
	Site	Gestation	Secretor status	N		
Asakuma et al. (2007)[70]	Japan	Term	Unknown	20	HPLC-UV	Acidic
Asakuma et al. (2008)[42]	Japan	Term	+	12	HPLC-UV	Neutral
Bao et al. (2007)[47]	USA	Term	Unknown	8	CE-UV	Acidic
Bao et al. (2013)[48]	USA	Term	+	4	LC-MS	Neutral
Chaturvedi et al. (2001)[43]	Mexico	Term	+	11	HPLC-UV	Neutral
Coppa et al. (1999)[34]	Italy	Term	+	18	HPAEC-PAD	Neutral, acidic
Coppa et al. (2011)[32]	Italy	Term	+/−	16/23	HPAEC-PAD	Neutral
Erney et al. (2000)[35]	Various world regions	Term	+	197	HPAEC-PAD	Neutral
Gabrielli et al. (2011)[33]	Italy	27.9 wk	+/−	42/21	HPAEC-PAD	Neutral, acidic
Goehring et al. (2014)[71]	USA	Term	+/−	13/4	LC-MS	Neutral, acidic
Hong et al. (2014)[72]	USA	Term	+/−	10/10	LC-MS	Neutral, acidic
Kunz et al. (1999)[40]	Germany	Term	+/−	2/2	HPAEC-PAD	Neutral, acidic
Leo et al. (2010)[73]	Samoa	Term	Unknown	16	HPLC-UV	Neutral, acidic
Martin-Sosa et al. (2003)[44]	Spain	Term	Unknown	12	HPLC-UV	Acidic
Nakhla et al. (1999)[41]	USA	29.5 wk	+	12	HPAEC-PAD	Acidic
Olivares et al. (2015)[74]	The Netherlands	Term	+/−	7/5	CE-FL	Neutral
Smilowitz et al. (2013)[45]	USA	Term	+/−	40/12	NMR	Neutral
Spevacek et al. (2015)[75]	USA	Term/preterm	Unknown	15/13	NMR	Neutral, acidic
Sumiyoshi et al. (2003)[39]	Japan	Term	Unknown	20	HPLC-UV	Acidic
Thurl et al. (2010)[31]	Germany	Term	+	21	HPAEC-PAD	Neutral, acidic
Van Niekerk et al. (2014)[46]	South Africa	500–1250 g[a]	+/−	20/21	HPLC-UV	Neutral, acidic

Abbreviations: CE-FL, capillary electrophoresis with fluorescence detection; CE-UV, capillary electrophoresis with UV detection; HPAEC-PAD, high-pH anion-exchange chromatography with pulsed amperometric detection; HPLC-UV, high-performance liquid chromatography with UV detection; LC-MS, liquid chromatography with mass spectrometric detection; NMR, nuclear magnetic resonance spectrometry; +, positive; −, negative.
[a]Birth weight of the premature infants.

and Goehring et al.[71] and LST b values reported by Coppa et al.[34] were eliminated.

Effect of secretor status

Secretor mothers regularly produced HMOS with fucose residues bound in α1,2 linkages to terminal galactose units of various oligosaccharides. In contrast, nonsecretor women usually did not produce these HMOS. However, Hong et al.[72] and van Niekerk et al.[46] reported very low concentrations of the α1,2 fucosylated compound LNFP I in milk samples from nonsecretor mothers. In addition, Hong et al.[72] found a relatively high mean value of 2′-FL (0.45 g/L) for 10 nonsecretor mothers. They attributed this extraordinarily high content to 2 nonsecretor mothers who produced amounts of the α1,2 fucosylated structure 2′-FL that were similar to amounts produced by secretor donors.[72] In their study, secretor status was determined on the basis of serological tests. Since the Le[b] epitope was shown to be weakly expressed at birth,[76] it is possible that the secretor status was incorrectly assigned in some cases.[76,77]

The effect of the secretor status was tested for all HMOS except the above-mentioned α1,2 fucose-containing structures that usually are absent in milks of nonsecretor mothers. To avoid a possible influence of the quantification methods, only data from studies reporting HMOS concentrations from both secretor and nonsecretor mothers in a given lactation period were used for the analyses of variance. Statistical comparisons, usually based on 1 to 3 studies, were possible with 4 neutral HMOS from term milks and 12 neutral HMOS from preterm milks (Figure 3). Concentrations of 3-FL in term milks and of LNT, LNFP II, LNDFH II, F-LNH II, and DF-para-LNH in preterm milks were significantly higher in milks from nonsecretor mothers than in milks from secretor mothers. In addition, concentrations of the acidic HMOS 6′-SL in term and preterm milks and of 6 other acidic HMOS from preterm milks could be compared (Figure 4). Concentrations of 6′-SL, LST a, and FS-LNnH I in preterm milks were significantly different between secretor and nonsecretor mothers, but no general tendency toward higher concentrations of acidic HMOS in milks from nonsecretor mothers was detectable.

Influence of lactation periods

In order to examine the effect of the time of lactation on HMOS concentrations, the time postpartum was classified into 6 lactation periods, as described above (see *Design of the systematic review* in Methods). The large volume of data on concentrations of HMOS is provided in the Supporting Information online (see Tables S2-A through S2-F and Tables S3, S4, and S5). Hereby, the concentrations of HMOS are listed
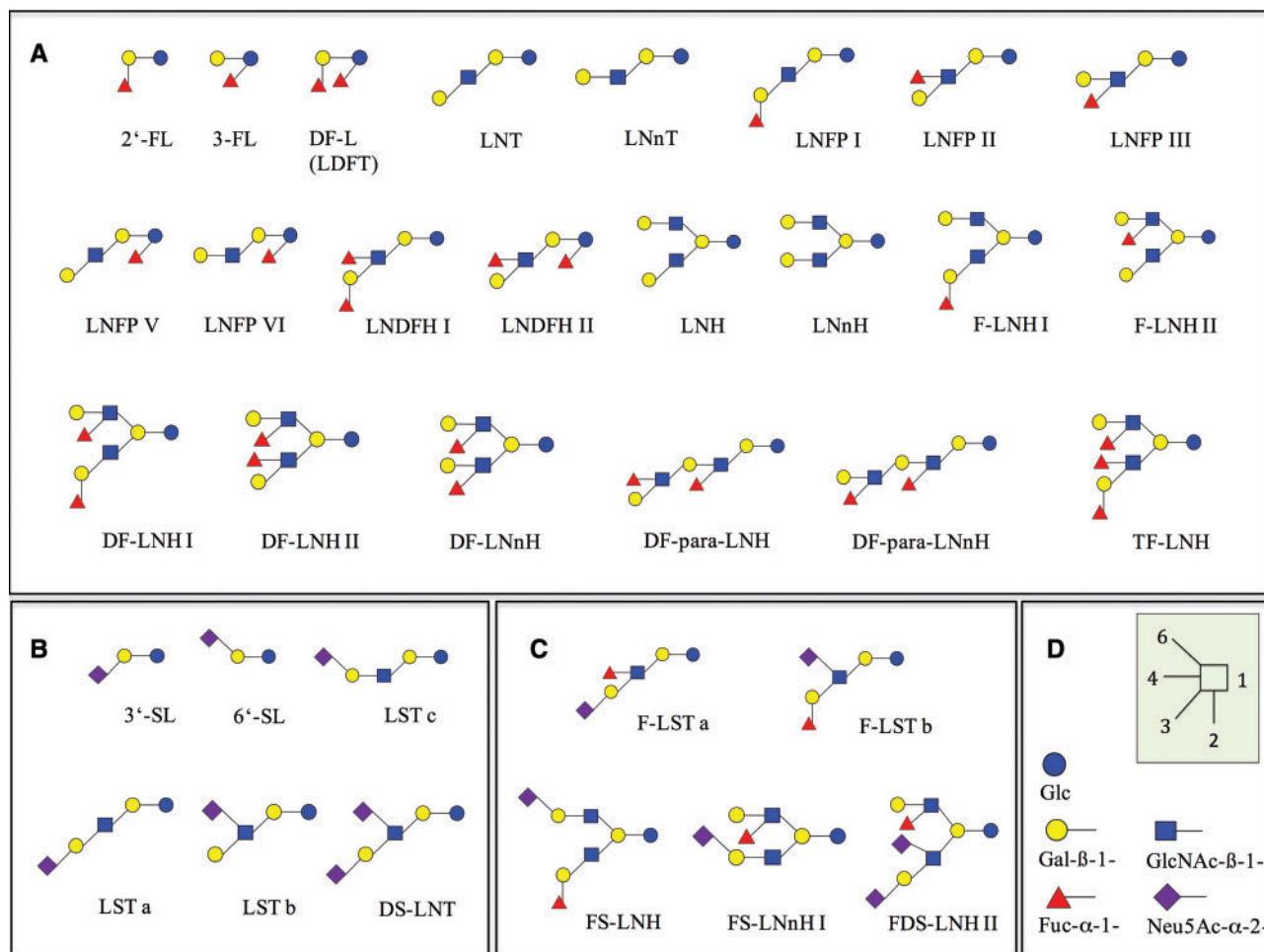
Figure 2 **Structures of the HMOS examined in this review.** (**A**) Neutral HMOS, (**B**) acidic nonfucosylated HMOS, (**C**) acidic fucosylated HMOS, (**D**) monomers and linkages.

separately for mothers with term and preterm infants, considering the lactation periods. The influence of the lactation period was tested in secretor mothers because these data on concentrations were readily available. No significant effect of the lactation period on any of the 13 HMOS from preterm milks was detected. Twenty-two HMOS from term milks could be tested. Concentrations of the neutral, ie, nonsialylated, HMOS 3-FL increased significantly (at least 2-fold) during the first 3 and 4 months postpartum (see Table S2-A in the Supporting Information online). Similarly, the concentration of LNFP III rose, but not significantly (see Table S3 in the Supporting Information online). During the course of lactation, the neutral α1,2 fucosylated HMOS LNFP I decreased approximately 2-fold (see Table S2-B in the Supporting Information online), and the acidic nonfucosylated HMOS LST c decreased approximately 4-fold (see Table S2-D in the Supporting Information online). Further significant differences were detected with the neutral HMOS LNDFH I (see Table S2-C in the Supporting Information online). In addition,

concentrations of the neutral HMOS 2′-FL, DF-L, LNFP II, F-LNH II, and DF-LNH II and the acidic HMOS 3′-SL, 6′-SL, and LST a in term milks decreased during the course of lactation, although not significantly (see Tables S3 and S4 in the Supporting Information online). Concentrations of the 2 acidic HMOS 6′-SL and LST a declined significantly during the first 100 days postpartum when the secretor status of mothers was not considered (see Tables S2-E and S2-F in the Supporting Information online). This outcome could be explained by the additional studies that were taken into account in these cases (Tables 4 and 5).

DISCUSSION

During the 1980s and early 1990s, many quantitative data on oligosaccharide fractions in human milk were published. These data were obtained mainly by use of the separation techniques gel permeation chromatography and low-resolution liquid chromatography.[78–82] The application of more advanced chromatographic

Table 3 Concentrations of neutral human milk oligosaccharides (HMOS) from secretor mothers[a]

HMOS (ref. nos. in superscript)	Lactation (days)	No. of mothers/no. of samples	Mean concentration (g/L)	95%CL (g/L)
Term studies				
2'-FL [31–35,42,43,45,71,72]	0 to > 100	353/556	2.74	2.43–3.04
3-FL [a31,32,34,42,43,45,48]	0 to > 100	122/365	0.44	0.31–0.58
DF-L [31,35,42,43,45,74]	0 to > 100	288/455	0.42	0.32–0.51
LNT [31,34,40,42,45,48,72,74]	0 to 100	114/308	0.79	0.59–0.98
LNnT [31,34,35,42,48]	0 to 100	184/372	0.74	0.36–1.12
LNFP I [a31,32,34,35,40,42,43,45,48,72]	0 to > 100	331/580	1.31	1.08–1.53
LNFP II [31,32,34,35,45,48,74]	0 to 100	229/389	0.28	0.21–0.34
LNFP III [31,35,48]	0 to 100	154/246	0.33	0.24–0.42
LNFP V [35,48]	0 to 30	133/137	0.06	−0.15 to 0.26
LNFP VI [48]	0 to 30	4/8	0.01	0.00–0.02
LNDFH I [a31,34,43,48,74]	0 to > 100	61/214	0.80	0.66–0.94
LNDFH II [31,32,34,42,43,48]	0 to > 100	76/319	0.14	0.10–0.18
LNH [31,34,72]	0 to 100	49/209	0.09	0.06–0.13
LNnH [34]	0 to 100	18/90	0.16	0.06–0.25
F-LNH I [31]	0 to 100	21/109	0.20	0.08–0.33
F-LNH II [31,32,34]	0 to 100	53/176	0.27	0.14–0.40
DF-LNH I [31]	0 to 100	21/109	0.31	0.19–0.43
DF-LNH II [32,34]	0 to 100	28/100	2.31	1.93–2.68
DF-LNnH [34]	0 to 100	18/90	0.54	0.10–0.98
TF-LNH [32,34]	0 to 100	28/100	2.84	2.60–3.07
Sum of means			14.78	
Preterm studies				
2'-FL [33,41,46]	0–60	74/230	2.77	0.76–4.78
3-FL [33,41,46]	0–60	75/230	0.32	0.17–0.48
DF-L [33,41]	0–60	54/190	0.41	0.17–0.65
LNT [33,41,46]	0–60	75/356	1.04	0.39–1.68
LNnT [33,41,46]	0–30	75/227	0.66	0.04–1.28
LNFP I [33,41,46]	0–60	74/230	1.09	0.39–1.78
LNFP II [33,41,46]	0–60	68/198	0.27	0.14–0.40
LNFP III [33,41,46]	0–60	75/230	0.16	0.04–0.28
LNFP V [41]	0–60	12/22	0.02	0.00–0.04
LNDFH II [33,41]	0–60	47/168	0.07	0.00–0.14
LNH [33]	0–30	42/168	0.06	0.04–0.08
LNnH [33]	0–30	42/168	0.08	0.06–0.11
F-LNH II [33]	0–30	41/168	0.33	0.26–0.41
DF-LNH II [33]	0–30	35/140	2.70	2.54–2.86
DF-para-LNH [33]	0–30	35/140	0.44	0.10–0.79
DF-para-LNnH [33]	0–30	42/168	0.50	−0.09 to 1.09
TF-LNH [33]	0–30	35/140	0.65	0.21–1.08
Sum of means			11.57	

Abbreviations: CL, confidence limit; ref. nos., reference numbers.
[a]HMOS exhibiting significant differences between the means of the lactation periods (see Tables S2-A, S2-B, and S2-C in the Supporting Information online).

and electrophoretic techniques like high-performance liquid chromatography–UV, high-pH anion-exchange chromatography with pulsed amperometric detection, capillary electrophoresis with UV detection, and capillary gel electrophoresis with laser-induced fluorescence detection after 1995 allowed the quantification of individual oligosaccharides in their native form or after derivatization.[24–29] Recently, a variety of promising mass spectrometric methods, along with nuclear magnetic resonance techniques, have emerged.[13,30,45,48,63,72,83–85] The growing information on oligosaccharide structures, quantities, and biological functions has been examined in numerous review articles.[2,6,7,20,21,23,86–89] In recent years, several reviews have also described the analytical methods applied for quantification and structural

analysis of HMOS.[1,90–93] Some of the reviews mentioned above, as well as some original studies, summarize data on HMOS concentrations without providing a comprehensive overview.[1,47,48,54,89] To the best of knowledge, this is the first systematic review of quantitative data for individual HMOS. Data on concentrations are reported, and the influence of 2 biological parameters—secretor status and lactation period—is examined.

Variations in HMOS data

Mean values and confidence intervals of a variety of neutral and acidic HMOS could be calculated from the studies selected. On the basis of the available literature

Table 4 **Concentrations of acidic human milk oligosaccharides (HMOS) from secretor mothers[a]**

HMOS (ref. nos. in superscript)	Lactation (days)	No. of mothers/no. of samples	Mean concentration (g/L)	95%CL (g/L)
Term studies				
3′-SL[31,34,40,72]	0–100	51/217	0.19	0.14–0.24
6′-SL[31,34,40,45,71,72]	0–100	104/270	0.64	0.38–0.91
LST a[31,34]	0–30	39/199	0.06	0.02–0.11
LST b[31,34,40]	0–100	41/203	0.13	0.08–0.18
LST c[31,34,72]	0–100	49/191	0.25	0.13–0.38
DS-LNT[31,34]	0–100	39/199	0.50	0.34–0.66
FS-LNnH I[34]	0–100	18/90	0.36	0.21–0.51
Sum of means			2.13	
Preterm studies				
3′-SL[33,46]	0–30	62/208	0.29	0.21–0.36
6′-SL[33]	0–30	42/168	0.66	0.25–1.08
LST a[33]	0–30	42/168	0.29	0.13–0.44
LST b[33,46]	0–30	62/208	0.13	0.06–0.19
LST c[33,46]	0–30	62/208	0.71	0.24–1.17
DS-LNT[33,46]	0–30	62/208	0.77	0.22–1.32
FS-LNnH I[33]	0–30	42/168	0.44	0.27–0.62
Sum of means			3.29	

Abbreviations: CL, confidence limit; ref. nos., reference numbers.
[a]HMOS exhibiting significant differences between the means of the lactation periods (see Table S2-D in the Supporting Information online).

Table *5* **Concentrations of acidic human milk oligosaccharides (HMOS) from lactating women, regardless of secretor status[a]**

HMOS (ref. nos. in superscript)	Lactation (days)	No. of mothers/no. of samples	Mean concentration (g/L)	95%CL (g/L)
Term studies				
3′-SL[31,34,39,40,44,45,47,70,72,73,75]	0–100	200/509	0.16	0.12–0.19
6′-SL[a34,39,40,44,45,47,70,72,73,75]	0–100	179/400	0.35	0.29–0.42
LST a[a31,34,39,70,73]	0–100	91/363	0.07	0.04–0.10
LST b[31,39,40,47,70,73]	0–100	83/285	0.06	0.05–0.07
LST c[31,34,39,47,70,72,73]	0–100	119/373	0.26	0.16–0.36
DS-LNT[31,34,39,44,47,70,73]	0–100	111/395	0.54	0.37–0.71
F-LST a[47]	11–30	3/3	0.02	–
F-LST b[70]	0–4	20/60	0.08	–
FS-LNH[47]	0–30	8/8	0.12	−0.08 to 0.33
FS-LNnH I[34,47]	0–100	26/98	0.29	0.14–0.44
FDS-LNH II[47]	0–100	3/3	0.12	–
Sum of means			2.07	
Preterm studies				
3′-SL[33,46,75]	0–30	114/350	0.24	0.20–0.28
6′-SL[33,75]	0–30	73/268	0.60	0.40–0.80
LST a[33]	0–30	63/252	0.34	0.17–0.51
LST b[33,46]	0–30	104/334	0.14	0.10–0.17
LST c[33,46]	0–30	104/334	0.65	0.19–1.12
DS-LNT[33,46]	0–30	104/334	0.92	0.26–1.58
FS-LNnH I[33]	0–30	63/252	0.51	0.26–0.75
Sum of means			3.40	

*Abbreviation*: CL, confidence limit; ref. nos., reference numbers.
[a]HMOS exhibiting significant differences between the means of the lactation periods (see Tables S2-E and S2-F in the Supporting Information online).

and because of the strict study and data selection criteria applied in this review, only data from secretor mothers with term infants were available from several different studies. For this review, HMOS for which data were available from 5 or more studies, eg, 2′-FL, 6′-SL, LNDFH II, were considered particularly relevant. The high concentrations of the relatively large structures DF-LNH II and TF-LNH were reported by 2 studies, both performed by the same working group that applied a similar quantification method.[32,34] These high concentrations, which strongly influenced the sum values of HMOS, need to be confirmed by other research groups. Data processing revealed wide variations in HMOS concentrations, which is reflected in the relatively wide limits of confidence shown in Tables 3, 4, and 5. In particular, results obtained by different
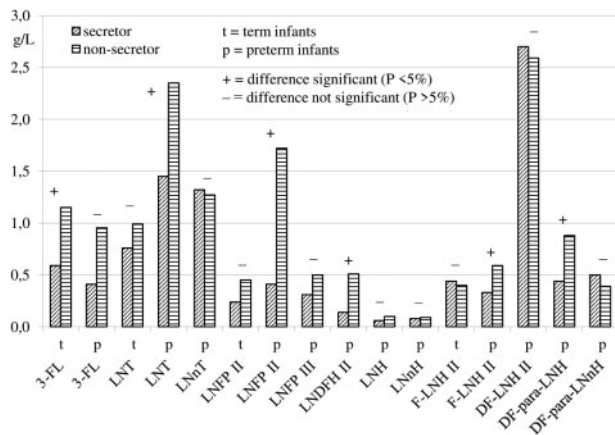
Figure 3 **Comparison of neutral HMOS concentrations in milks from secretor and nonsecretor mothers.**
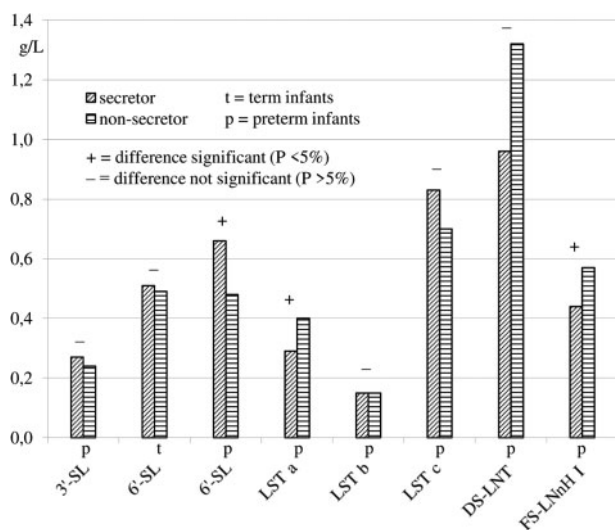
Figure 4 **Comparison of acidic HMOS concentrations in milks from secretor and nonsecretor mothers.**

working groups analyzing HMOS varied considerably. For example, the concentration of LNnT exhibiting a type 2 structure reported by Coppa et al.[32,34] exceeded that detected by Thurl et al.[31] by at least 4-fold. As a consequence, although the results of this review show concentrations of type 1 structures to be approximately twice that of type 2 structures (Tables 3 and 4), the clear dominance of type 1 structures reported[1,31] could not be confirmed. It is hypothesized that the predominance of type 1 structures coevolved with the dominance of bifidobacteria in infants' guts, as exoenzymes produced by bifidobacteria preferentially degrade oligosaccharide type 1 structures.[94] In the case of the acidic compound 6′-SL, unusually high concentrations reported in 1 study[31] were detected as outliers when compared with concentrations reported in most other studies.[34,39,40,47,70] This finding has biological implications,

since acidic HMOS, including 6′-SL, for example, are thought to modulate immunologic responses in the infants.[22]

The variation in HMOS concentrations was also attributed to biological parameters like gestational age, secretor status, lactation period, or general biological variability. The influence of gestational age on the composition of human milk, eg, on lipids and proteins, has been discussed in pediatrics for decades.[36] Although the data analyses with term and preterm milks were conducted separately in this review, no clear effects of gestational age on HMOS concentrations were found. The different sums of total neutral HMOS might be attributable to the different compounds quantified and by an unusually high concentration of TF-LNH in term milks.[32,34] Concentrations of acidic HMOS may have been higher in preterm milks than in term milks because preterm samples were obtained during the early lactation period of 30 days, while term samples were obtained during the period of 100 days. Some data in this review as well as some from published studies show that concentrations of acidic HMOS, in particular, decrease significantly during the first 3 months postpartum.[31,34,39]

This overview confirms the general finding that secretor mothers produce high amounts of α1,2 fucosylated HMOS compared with nonsecretor mothers,[2,15,16,32,33] whose milk either contain nondetectable or very low concentrations.[46,72] The prevalence of α1,2 fucosylated HMOS in secretor milks likely has biological consequences for infants. These compounds significantly reduced the incidence of diarrhea associated with enterotoxigenic *Escherichia coli*, *Campylobacter jejuni*, or caliciviruses,[60,62] thus possibly conferring an evolutionary advantage for infants of secretor mothers. The indirect effects of the secretor status on HMOS lacking α1,2 fucoses were also examined. An influence of the secretor status on HMOS concentrations was found in 9 of 24 comparative tests (1 test with term, 8 tests with preterm milks). All 6 tests of neutral HMOS that had significant results showed that nonsecretor mothers produced higher concentrations than secretor mothers, which is in accordance with findings reported in the literature.[31,33,45] Probably owing to the lack of the α1,2 fucose-transferring secretor enzyme, milks from nonsecretor mothers contain higher amounts of the remaining nonfucosylated as well as the α1,3-, and α1,4-fucosylated HMOS. Since very few studies were available reporting HMOS data from both secretor and nonsecretor mothers, all results—whether significant or not—should be evaluated with care. Different results cannot be ruled out when more studies become available.

Surprisingly, the influence of the time of lactation on HMOS concentrations was significant for only 4 HMOS out of 22 tested in term milks from secretor mothers, although this had already been reported in several studies.[31,34,35,39,43] This result can be explained by the wide variations in findings within the studies and the even greater variations between the studies. As reported previously,[31,34] LNFP I concentrations decreased significantly during the first months of lactation. However, a decrease in the major HMOS 2′-FL and a general tendency that α1,2 fucosylation and the activity of the secretor enzyme decline during the course of lactation could not be convincingly demonstrated, in contrast to findings in other studies.[31,34,43] It is hypothesized that the large interlaboratory differences in findings prevented the detection of this effect. In contrast, the concentration of 3-FL was shown to increase during lactation. This outcome, already reported in some studies,[31,35,43] can be explained by differing activities of the Lewis enzyme and other fucosyltransferases during the course of lactation.[31] Two of the major HMOS, 3-FL and LNFP II, were shown to specifically block a fucose-binding lectin of the human pathogen *Pseudomonas aeruginosa*, thus possibly protecting newborns against infection.[95]

Future HMOS studies

A variety of biological factors influence or could influence HMOS concentrations. Thus, future HMOS studies should precisely define the milk sampling procedures, such as time postpartum, Lewis blood groups of the donors (at least the secretor status), gestational age, ethnicity, and techniques of milk sampling. In addition, there is an initial report that sialylated HMOS might be influenced by the nutritional status of the mother.[96] Further studies on the effect of the maternal diet on HMOS are needed.

A variety of chromatographic, electrophoretic, and spectrometric methods of quantification were used in the studies analyzed in this review, which likely influenced the concentrations reported. In recent years, mass spectrometry and nuclear magnetic resonance methods emerged and were judged by several researchers to be superior to the established methods.[63,92] However, because of the complexity of overlapping signals of the same monosaccharide residue in very similar chemical environments, quantitative data obtained by nuclear magnetic resonance should be evaluated with care.[93,97] Data on absolute HMOS concentrations obtained with mass spectrometry or nuclear magnetic resonance are rare. In most cases, methods were described, but the number of samples analyzed was small and was not well defined biologically.[28,30,53,63,65] Currently, there are no studies that compare the performance of the different quantification techniques. In the case of chromatographic methods, the application of internal standards is recommended. With mass spectrometric methods, the use of the corresponding standard substances should be obligatory, and universal calibration should not be applied. Hong et al.[72] reported some differences in results when the 2 methods of quantification were compared. Ideally, interlaboratory tests should be conducted with several standardized human milk samples, with homogenized pooled human milk preparations from secretor and Lewis-positive donors representing the most complex matrix. Several laboratories should participate, all applying the major techniques: high-performance liquid chromatography, high-pH anion-exchange chromatography, capillary electrophoresis, capillary gel electrophoresis, mass spectrometry, and nuclear magnetic resonance. Moreover, several laboratories should examine the human milk samples in parallel, using different methods. As was recently shown for a large set of glycoprotein glycans,[98] an adapted approach and effort could also be applied to the quantification of HMOS.

This review focused on the concentrations of HMOS. However, the biological effects of HMOS are related to the daily amounts ingested by infants during the entire period of breastfeeding. Therefore, the concentration values would have to be multiplied by the milk volume or milk mass consumed. Average milk intakes reported in the literature could be used for purposes of estimation. It is generally assumed that, after the first month postpartum, infants drink about 700 mL of milk per day.[99,100] Data on HMOS concentrations and the daily amounts consumed are a prerequisite for evaluating the possible multiple biological functions of HMOS or oligosaccharide subgroups, eg, antiadhesive or prebiotic effects.

Detailed knowledge about HMOS structures and quantities is of more than just academic interest. For about 15 years, certain oligosaccharides, particularly mixtures of short-chain galactooligosaccharides and long-chain fructooligosaccharides (9:1) that mimic the complexity of size and the biological effects of HMOS, have been added to infant formulas and are currently recommended if exclusive breastfeeding is not possible. More recently, studies in newborn infants showed that the HMOS 2′-FL, whether combined with short-chain galactooligosaccharides or not, is safe and tolerated as a formula supplement.[101,102] Moreover, infants fed 2′-FL-fortified formula showed immune biomarkers similar to those detected in breastfed infants.[103]

Most recent studies

New studies reporting concentrations of HMOS are continuously being published. An update of the

literature search from August 27, 2015, corresponding to the deadline of this review, until December 14, 2016, yielded 231 records. After duplicate publications were eliminated, 135 reports were screened. Ten promising studies were examined in detail,[104–113] but 9 did not meet the strict inclusion criteria applied in this review (see Table S6 in the Supporting Information online). One article[113] reported concentrations of 3 neutral HMOS.

CONCLUSION

Data on the concentrations of oligosaccharides in human milk, as presented in this review, are a prerequisite for understanding the biological functions of HMOS and should help guide further developments in infant and maternal nutrition. Nevertheless, more detailed information on the composition of human milk is needed. Further studies with well-defined human milk samples (ie, documentation of lactation period, Lewis blood group, gestational age of the mothers, etc) using state-of-the art quantification methods will likely reveal valuable information about the composition and quantity of HMOS.

Acknowledgments

Supporting Information

The following Supporting Information is available through the online version of this article at the publisher's website.

Table S1 **Excluded studies reporting quantitative HMOS data**

Table S2-A **Concentration of 3-FL during the course of lactation (secretor mothers with term infants)**

Table S2-B **Concentration of LNFP I during the course of lactation (secretor mothers with term infants)**

Table S2-C **Concentration of LNDFH I during the course of lactation (secretor mothers with term infants)**

Table S2-D **Concentration of LST c during the course of lactation (secretor mothers with term infants)**

Table S2-E **Concentration of 6′-SL during the course of lactation (term infants)**

Table S2-F **Concentration of LST a during the course of lactation (term infants)**

Table S3 **Concentrations of neutral HMOS from secretor mothers during the course of lactation**

Table S4 **Concentrations of acidic HMOS from secretor mothers during the course of lactation**

Table S5 **Concentrations of acidic HMOS during the course of lactation regardless of secretor status of the mothers**

Table S6 **Most recent studies reporting quantitative HMOS data**

Table S7 **PRISMA 2009 checklist**

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