Glucose tolerance and insulin resistance in Indian children: relationship to infant feeding pattern

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Abstract

Aims/hypothesis Our objective was to examine whether longer duration of breast-feeding and later introduction of complementary foods are associated with lower glucose concentrations and insulin resistance (IR-HOMA) in Indian children.

Methods Breast-feeding duration (six categories from <3 to ≥18 months) and age at introduction of complementary foods (four categories from <4 to ≥6 months) were recorded at 1, 2 and 3 year follow-up of 568 children from a birth cohort in Mysore, India. At 5 and 9.5 years of age, 518 children were assessed for glucose tolerance and IR-HOMA.

Results All the children were initially breast-fed; 90% were breast-fed for ≥6 months and 56.7% started complementary foods at or before the age of 4 months. Each category increase in breast-feeding duration was associated with lower fasting insulin concentration ($\beta = -0.05$ pmol/l [95% CI $-0.10, -0.004$]; $p = 0.03$) and IR-HOMA ($\beta = -0.05$ [95% CI $-0.10, -0.001$]; $p = 0.046$) at 5 years, adjusted for the child's sex, age, current BMI, socioeconomic status, parent's education, rural/urban residence, birthweight and maternal gestational diabetes status. Longer duration of

breastfeeding was associated with higher 120-min glucose concentration at 5 years ($\beta = 0.08$ mmol/l [95% CI 0.001, 0.15; $p = 0.03$]) but lower 120-min glucose concentration at 9.5 years ($\beta = -0.09$ [95% CI $-0.16, -0.03$]; $p = 0.006$). Age at starting complementary foods was unrelated to the children's glucose tolerance and IR-HOMA.

Conclusions/interpretation Within this cohort, in which prolonged breast-feeding was the norm, there was evidence of a protective effect of longer duration of breast-feeding against glucose intolerance at 9.5 years. At 5 years longer duration of breast-feeding was associated with lower IR-HOMA.

Keywords Breast-feeding · Children · Complementary foods · Glucose tolerance · India · Insulin resistance

Abbreviations

GDM Gestational diabetes mellitus
IR-HOMA Insulin resistance HOMA

Introduction

Children and adults who were breast-fed rather than bottle-fed, and breast-fed for a longer duration during infancy have lower rates of type 2 diabetes and lower insulin resistance (IR-HOMA) [1, 2]. It has been suggested that differences between breast-fed and bottle-fed babies in the nutrient quality of the milk, patterns of infant weight gain, or in learned feeding behaviour influence later diabetes risk [1]. Few studies have examined diabetes risk in relation to the age at starting complementary foods in infancy.

In the Mysore Parthenon birth cohort study [3], data collected on infant feeding practices, and subsequent

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measurements of plasma glucose and insulin concentrations in the children enabled us to examine whether longer duration of breast-feeding and later introduction of complementary foods are associated with lower glucose concentrations and IR-HOMA in Indian children, and whether these associations change with age.

Methods

As described previously [3], 830 pregnant women underwent an OGTT at 30±2 weeks of gestation (49 had gestational diabetes [GDM]). Of these, 663 delivered live normal babies at the Holdsworth Memorial Hospital, Mysore. The children

had detailed anthropometry at birth, annually until the age of 5 years and every 6 months thereafter.

Infant feeding data were collected at 1, 2 and 3 years of age by asking mothers the following questions: How was the baby fed from birth (breast, bottle, breast+bottle or other)? If breast-fed, was the baby still being breast-fed? If not, at what age (months) was breast-feeding stopped? At 1 year, mothers were asked the age in months at which their baby started taking solid foods regularly.

Five hundred and eighty-five children (93% of survivors) were studied at 5 years and 539 (86%) at 9.5 years. Weight (Salter, Tonbridge, UK) and height (Microtoise, CMS Instruments, London, UK) were measured using standardised methods. After an overnight fast, blood

Table 1 General characteristics of the study cohort

Infant feeding	Boys (<i>n</i> = 246)	Girls (<i>n</i> =272)	<i>p</i> value
Breast-feeding duration categories, <i>n</i> (%)			
<3 months	7 (2.9)	7 (2.6)	0.6
3–5 months	23 (9.4)	19 (7.0)	
6–8 months	26 (10.6)	23 (8.5)	
9–11 months	34 (13.8)	45 (16.5)	
12–17 months	105 (42.7)	131 (48.2)	
≥18 months	51 (20.7)	47 (17.3)	
Age at starting regular solids, <i>n</i> (%)			
<4 months	43 (18.7)	51 (20.2)	0.5
4 months	81 (35.2)	99 (39.1)	
5 months	70 (30.4)	61 (24.1)	
≥6 months	36 (15.7)	42 (16.6)	
Child characteristics at 5 years (mean±SD)			
Age (years)	5.0±0.04	5.0±0.03	0.09
Height (cm)	106.3±4.3	104.9±4.3	0.0004
BMI (kg/m ²)	13.6±1.0	13.5±1.2	0.6
Fasting glucose (mmol/l)	4.9±0.5	4.8±0.3	0.3
30 min glucose (mmol/l)	7.2±1.5	7.4±1.3	0.1
120 min glucose (mmol/l)	5.9±1.0	5.9±0.9	0.5
Fasting insulin (pmol/l) ^a	18.6 (11.4, 28.6)	22.7 (15.0, 22.7)	<0.0001
30 min insulin (pmol/l) ^a	120.1 (75.8, 211.1)	168.4 (108.0, 242.0)	0.0006
Insulin resistance (IR-HOMA) ^a	0.6 (0.4, 1.0)	0.8 (0.5, 1.2)	0.0001
Insulin increment ^a	14.3 (9.0, 24.7)	19.5(12.8, 27.0)	0.02
Child characteristics at 9.5 years (mean±SD)			
Age (years)	9.4±0.1	9.4±0.1	0.3
Height (cm)	131.3±5.5	130.3±5.9	0.07
BMI (kg/m ²)	14.6±1.8	14.7±2.0	0.9
Fasting glucose (mmol/l)	4.7±0.4	4.7±0.4	0.1
30 min glucose (mmol/l)	6.7±1.3	6.9±1.2	0.1
120 min glucose (mmol/l)	5.0±1.0	5.2±0.8	0.067
Fasting insulin (pmol/l) ^a	19.2 (12.0, 28.8)	26.4 (18.6, 36.0)	<0.0001
30 min insulin (pmol/l) ^a	208.2 (124.8, 342.0)	270.0 (166.8, 408.0)	<0.0001
Insulin resistance (IR-HOMA) ^a	0.7 (0.4, 1.0)	0.9 (0.6, 1.3)	<0.0001
Insulin increment ^a	26.5 (16.8, 45.5)	35.3 (21.6, 49.7)	0.005

p values for the differences between boys and girls derived using Student's *t* test or χ^2 test

^a Log-transformed variable; values are median and interquartile range

samples were collected (fasting, and 30 and 120 min after an oral glucose load [1.75 g/kg body weight]). Insulin was analysed by a time-resolved, fluoroimmunoassay (DEL-FIA) and plasma glucose concentrations by standard enzymatic method (Alcyon 3000, Abbott Laboratories, Abbott Park, IL, USA). Inter-assay coefficients of variations for insulin and glucose were <10% and <5%, respectively. IR-HOMA was estimated using the HOMA equation [4]. Insulin increment was calculated ([30-min insulin–fasting insulin]/30-min glucose) [5].

The hospital's ethics committee approved the study; informed consent/assent was obtained from parents and children.

Statistical methods Insulin (fasting, 30 and 120 min), IR-HOMA and insulin increment were log transformed to normality. From a public health perspective (for example, having a bin for <3 months) and also from a statistical perspective (i.e. having enough participants in each bin), to reduce the effect of outliers and maintain ordering, total duration of breast-feeding in months was split into six

categories (<3, 3–5, 6–8, 9–11, 12–17 and ≥ 18 months). Age at starting complementary foods was split into four categories (<4, 4, 5, ≥ 6 months). The associations of breast-feeding duration and age at starting complementary foods with outcomes and potential confounders were examined by multiple linear regression using Stata v10 (Stata, College Station, TX, USA).

Results

A total of 518 children had complete feeding data and outcomes at both 5 and 9.5 years (electronic supplementary material [ESM] Fig. 1). All were initially breast-fed, 90% were breast-fed for ≥ 6 months and 64% for ≥ 12 months (Table 1). More than half (56.7%) of the children had started complementary foods by 4 months. At both timepoints girls had higher insulin concentrations (fasting and 30 min), insulin increment and IR-HOMA than boys (Table 1).

Rural and GDM mothers breast-fed for longer than urban ($p=0.001$) and non-GDM mothers ($p=0.05$), respectively.

Table 2 Associations of duration of breast-feeding and age at introduction of complementary foods with glucose/insulin concentrations at 5 and 9.5 years: multiple linear regression analysis

	Age							
	5 years				9.5 years			
	β (95% CI)	p value ^b	p value ^c	p value ^d	β (95% CI)	p value ^b	p value ^c	p value ^d
Association of breast-feeding duration with:								
Glucose								
Fasting glucose (mmol/l)	0.01 (−0.02, 0.05)	0.4	0.6	0.4	0.002 (−0.02, 0.03)	0.9	0.9	0.9
120 min glucose (mmol/l)	0.08 (0.02, 0.15)	0.02	0.04	0.03	−0.06 (−0.13, 0.0003)	0.051	0.02	0.02
Insulin								
Fasting insulin (pmol/l) ^a	−0.06 (−0.11, −0.02)	0.006	0.01	0.03	0.001 (−0.04, 0.04)	0.9	0.7	0.9
30 min insulin (pmol/l) ^a	−0.04 (−0.10, 0.01)	0.1	0.1	0.3	0.02 (−0.03, 0.07)	0.4	0.5	0.4
Insulin resistance ^a	−0.06 (−0.11, −0.01)	0.01	0.02	0.046	0.001 (−0.04, 0.04)	0.9	0.7	0.9
Insulin increment ^a	−0.01 (−0.07, 0.05)	0.8	0.8	0.9	0.01 (−0.04, 0.06)	0.6	0.6	0.4
Association of age at introduction of complementary foods with:								
Glucose								
Fasting glucose (mmol/l)	−0.02 (−0.06, 0.03)	0.5	0.7	0.7	0.01 (−0.02, 0.05)	0.5	0.5	0.4
120 min glucose (mmol/l)	0.05 (−0.04, 0.14)	0.3	0.2	0.1	−0.01 (−0.09, 0.07)	0.8	0.9	1.0
Insulin								
Fasting insulin (pmol/l) ^a	0.01 (−0.05, 0.07)	0.7	0.7	0.6	0.01 (−0.04, 0.07)	0.6	0.4	0.1
30 min insulin (pmol/l) ^a	0.04 (−0.03, 0.11)	0.3	0.1	0.09	−0.01 (−0.08, 0.05)	0.7	0.8	0.5
Insulin resistance ^a	0.01 (−0.06, 0.07)	0.8	0.8	0.6	0.02 (−0.04, 0.07)	0.5	0.3	0.1
Insulin increment ^a	0.01 (−0.07, 0.09)	0.8	0.7	0.6	0.01 (−0.08, 0.06)	0.8	0.6	0.4

^a Log transformed variable; regression co-efficient (β) is the effect size per category increase in breast-feeding duration (<3, 3–5, 6–8, 9–11, 12–17 and ≥ 18 months) or age at starting complementary foods (<4, 4, 5 and ≥ 6 months)

^b p value adjusted for sex and child's current age

^c p value further adjusted for confounders (socioeconomic status, parents' education, rural/urban residence maternal GDM status and birthweight)

^d p value further adjusted for child's current BMI

There were no significant associations between breast-feeding duration and parental education or socioeconomic status.

At 5 years, 120-min glucose concentration increased, whereas fasting insulin concentration and IR-HOMA decreased with increasing duration of breast-feeding, independent of potential confounders (Table 2). The 30-min insulin concentration tended to fall with increasing breast-feeding duration, although this was non-significant. At 9.5 years, 120-min glucose concentration decreased with increasing duration of breast-feeding. IR-HOMA was unrelated to breast-feeding duration (Table 2).

Breast-feeding duration correlated with the age at starting complementary foods (Spearman $r=0.1$; $p=0.02$). Earlier introduction of complementary foods was associated with higher BMI at 9.5 years ($p=0.04$) and maternal education ($p=0.051$). There were no significant associations, at either 5 or 9.5 years, of age at starting complementary foods with glucose tolerance or IR-HOMA (Table 2).

The findings were unchanged when we re-analysed our data using breast-feeding duration or age at starting complementary foods as continuous or normalised variables, or if the categories were changed; when breast-feeding duration and age at starting complementary foods were included simultaneously in the regression; when preterm ($n=34$) and offspring of GDM mothers ($n=33$) were excluded and when weight at 1 year was included with all the other confounders (data not shown). The associations were similar in boys and girls and there were no non-linear associations.

Discussion

We have shown in a cohort of Indian children, among whom prolonged breast-feeding was the norm across all socioeconomic groups, that longer breast-feeding duration was associated with lower fasting insulin concentrations and IR-HOMA at 5 years, but not at 9.5 years. Longer breast-feeding duration was associated with higher 120-min glucose concentration at 5 years, but lower 120-min glucose at 9.5 years. These associations were independent of all potential confounding factors measured. There were no significant associations of age at starting complementary foods with glucose/insulin concentrations.

A strength of the study was that breast-feeding data were obtained prospectively. Limitations were a lack of information on the exclusivity and frequency of breast-feeding, the nutritional quality of the breast milk or the type of milk used after stopping breast-feeding, the type and nutritional quality of complementary foods, and current diet and physical activity.

Longer breast-feeding duration was associated with lower fasting insulin concentrations and IR-HOMA at 5 but not 9.5 years, suggesting that this association changes with age. There are no previous reports of serial measures of insulin in relation to breast-feeding. A meta-analysis of studies in developed countries (four in adults and two in children) [2] showed no differences in fasting insulin concentrations among those who were breast-fed or not. Similarly, two studies in adults [1, 2] and three in children [1, 2, 6] reported no association of exclusive breast-feeding and/or its duration with IR-HOMA [1, 2]. However, a recent study reported an inverse association between breast-feeding duration and IR-HOMA in adult men [7].

A notable finding in our study was that the association between breast-feeding duration and 120-min glucose concentrations changed in direction between 5 and 9.5 years. Children who stopped breast-feeding earlier had lower 120-min glucose concentrations at 5 years, but higher 120-min glucose at 9.5 years. In addition to having higher fasting insulin concentrations, they tended to have higher 30-min insulin concentrations. This may be analogous to the phenomenon observed in animal studies, in which offspring of rat dams exposed to a nutritional insult (high-carbohydrate or low-protein diets) during pregnancy have fewer and smaller pancreatic islets, long-term changes in the structure and function of insulin-sensitive tissues (liver, adipocytes and muscle) and reduced insulin signalling protein production. These changes are associated with hyperinsulinaemia, enhanced glucose tolerance or reactive hypoglycaemia in the young animal and to glucose intolerance later [8]. Such a progression has not previously been reported in humans, although few have serial data like ours through childhood.

Consistent with findings from many studies including a meta-analysis, mainly from high-income countries, our data at 9.5 years suggest that longer breast-feeding is associated with better glucose tolerance, and possibly lower diabetes risk, in later life. Whether these associations are causal is unknown, but animal studies have demonstrated that the high long-chain polyunsaturated fatty acid content of breast milk may suppress proinflammatory cytokine production, regulate neurotransmitter function, enhance insulin receptor numbers in the brain and other tissues, and decrease IR-HOMA [9]. Whether these changes could persist up to the age of 5 or 9.5 years is unknown. Breast-fed babies are thought to develop better satiety sensing than formula-fed babies and gain less weight during infancy [1], both of which may protect against later obesity and thus diabetes.

Few studies have examined the association of age at starting complementary foods with later IR-HOMA and glucose tolerance. Consistent with our findings, a recent study found no association among young adults [10]. As growth in infancy influences later health, timely introduction

and nutritional quality of complementary foods are important for promoting optimal growth and lifelong health.

To conclude, in this cohort of healthy Indian children, there was evidence of a possible protective effect of longer breast-feeding duration against later diabetes, suggesting that promoting WHO guidelines (exclusive breast-feeding for 6 months, introduction of nutritious complementary foods from 6 months and continued breast-feeding up to 2 years) [11] may contribute to reducing the escalating incidence of diabetes in developing countries. This improved glucose tolerance may be associated with a transient period of lower IR-HOMA and higher glucose concentrations in early childhood.

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Contribution statement SRV: conception and design, analysis and interpretation of data, drafting the article and approval of final version. GVK, JCH, SCK: conception and design, revising the manuscript critically for intellectual content and approval of final version. AKW: analysis and interpretation of data, revising the manuscript critically for intellectual content and approval of final version. CHDF: conception and design, interpretation of data and revising the manuscript critically for intellectual content and approval of final version.

Duality of interest The authors declare that there is no duality of interest associated with this manuscript.

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