

RESEARCH ARTICLE

Maternal pre-pregnancy overweight/obesity and gestational diabetes interaction on delayed breastfeeding initiation

Tanara Vogel Pinheiro^{1*}, Marcelo Zubaran Goldani^{1,2}, IVAPSA group^{1,2}

1 Department of pediatrics, Universidade Federal do Rio Grande do Sul, Porto Alegre, Rio Grande do Sul, Brazil, **2** Department of pediatrics, Hospital de Clínicas de Porto Alegre, Porto Alegre, Rio Grande do Sul, Brazil

* tanaravogel@gmail.com



Abstract

Background

Cumulative evidence indicates an association between maternal overweight and gestational diabetes with delayed breastfeeding initiation; however, the presence of both conditions simultaneously has been little explored. This study aims to investigate the interaction between maternal overweight/obesity and gestational diabetes on breastfeeding initiation.

Methods

This study comprises data from the IVAPSA Birth Cohort, a prospective follow-up of mothers and their newborns. Two of the five groups from IVAPSA were evaluated, considering women with and without gestational diabetes. These women were further categorized according to their pre-pregnancy body mass index as normal weight or overweight/obese.

Results

219 women were evaluated, 53.4% of them had pre-pregnancy overweight/obesity and 32% had gestational diabetes. Most women were able to initiate breastfeeding within 12 hours from delivery (92.7%) and only eight (3.7%) women had not breastfed in the first 24 hours postpartum. Of these, seven were overweight/obese (77.8%) and five had gestational diabetes (66.7%), with four of them having overweight/obesity and gestational diabetes concomitantly. Women with both adverse conditions had an adjusted relative risk of delayed breastfeeding initiation of 1.072 (95% CI 1.006; 1.141), $p = 0.032$.

Conclusions

The results indicate an additive interaction between maternal pre-pregnancy overweight/obesity and gestational diabetes on delayed breastfeeding initiation.

OPEN ACCESS

Citation: Pinheiro TV, Goldani MZ, IVAPSA group (2018) Maternal pre-pregnancy overweight/obesity and gestational diabetes interaction on delayed breastfeeding initiation. PLoS ONE 13(6): e0194879. <https://doi.org/10.1371/journal.pone.0194879>

Editor: Cassandra Nichole Spracklen, University of North Carolina at Chapel Hill, UNITED STATES

Received: December 5, 2017

Accepted: March 12, 2018

Published: June 18, 2018

Copyright: © 2018 Pinheiro et al. This is an open access article distributed under the terms of the [Creative Commons Attribution License](https://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Data Availability Statement: All relevant data are within the paper and its Supporting Information files.

Funding: The research was supported by FAPERGS/CNPq 10/0018.3 to MZG (Conselho Nacional de Desenvolvimento Científico e Tecnológico), PRONEX 2009 to MZG (Programa de Apoio a Núcleos de Excelência), FIPE/HCPA 110097 to MZG (Fundo de Incentivo à Pesquisa e Eventos do Hospital de Clínicas de Porto Alegre)

and CAPES (Comissão de Aperfeiçoamento de Pessoal do Nível Superior)—Brazil to TVP.

Competing interests: The authors have declared that no competing interests exist.

Abbreviations: aRR, Adjusted Relative Risk; BF, Breastfeeding; BMI, body mass index; GDM, gestational diabetes; IVAPSA, Impact of Perinatal Different Intrauterine Environments on Child Growth and Development in the First Six Months of Life Study; SD, Standard Deviation; SPSS, Statistical Package for the Social Sciences; WHO, World Health Organization.

Introduction

The prevalence of overweight and obesity among women of reproductive age has risen all over the world during the last decades [1, 2] and has been associated with a greater risk of pregnancy, obstetric and neonatal complications [3], including greater difficulties to initiate breastfeeding (BF) after delivery compared with healthy weight women [4–6]

The increase of maternal body mass index (BMI) is strongly associated with a higher risk of developing gestational diabetes (GDM) during pregnancy [7–9], which in turn is also a risk factor for BF failure [10, 11]. GDM has been associated with delayed onset of lactogenesis II (onset of copious milk secretion in the first few days postpartum), lower rates of BF initiation and earlier weaning [12].

The health benefits of BF are of particular importance for mothers with overweight and/or gestational diabetes, since BF is associated with a decreased probability of type 2 diabetes [13] and a potential decrease of postpartum weight retention [14, 15]. Children born from those women are also benefited by BF since increasing evidence indicates a protective effect against obesity and diabetes for children as well [16–18].

Considering that women with overweight and gestational diabetes are at increased risk of failure to initiate BF and that the timing of BF initiation is critical to BF success [19, 20], this study aims to investigate the interaction between maternal overweight/obesity and gestational diabetes on the timing of BF initiation.

Materials and methods

Participants

The subjects in this study are part of the “Impact of Perinatal Different Intrauterine Environments on Child Growth and Development in the First Six Months of Life—IVAPSA study”, a prospective follow-up of mothers and their newborns that aims to investigate health outcomes after exposure to adverse intrauterine environments. The IVAPSA study has a convenience sample of 400 mother-offspring pairs, recruited in three public hospitals of Porto Alegre, Brazil. The hospitals are Baby-Friendly, meaning that they follow the Ten Steps to Successful Breastfeeding and have written policies for breastfeeding protection, promotion and support. A detailed description of the IVAPSA recruitment and interviews has been previously published elsewhere [21].

This paper comprises only two of the five groups that integrate the IVAPSA study: women with gestational diabetes (GDM) and their offspring and a control group of mother-offspring with none of the adverse conditions investigated by IVAPSA. Women with positive test for HIV, multiple pregnancy or preterm delivery (<37 weeks) and infants with acute diseases or congenital birth defects were not included in the study. All women and babies were together in the maternity ward of the hospitals in the moment of recruitment, meaning that mother and child were healthy and with no greater health risks.

Data collection

Information presented in this paper was collected during two interviews: the first conducted between 24 and 48 hours postpartum, in the maternity ward of the hospitals, and the second between 30 and 45 days after delivery, either in the participants’ house or in clinics located in one of the hospitals.

In addition to the information provided by the mothers during the interviews, we also collected data from the mother’s prenatal care booklet, which contains medical records of the

entire gestation period, and from the hospital's medical registries for detailed information on hospital admission, delivery, exams and hospital discharge.

Maternal pre-pregnancy body mass index

Maternal pre-pregnancy weight was collected from the prenatal care booklet or, in the absence of this information, was reported by the mother in the first interview in response to the question "Just before you got pregnant with your new baby, how much did you weight?". Maternal height was measured in three occasions: before delivery (by hospital staff), at 6–8 days and at 30–45 days after delivery (by trained researchers). We preferably used the last measure (30 days) because there may be a postural adjustment during pregnancy and in the first days after delivery [22]. Pre-pregnancy body mass index (BMI) was calculated by the equation "Pre-pregnancy weight (kg)/height squared (m^2)" and categorized as Normal weight: 18.5–24.9 kg/m^2 ; Overweight: 25.0–29.9 kg/m^2 ; or Obese: $\geq 30.0 kg/m^2$. Women under 20 years were evaluated using the WHO AnthroPlus software BMI-for-age Z-score. We applied the BMI-for-age cut-offs suggested by WHO: Normal weight: Z-score between -2SD and +1SD; Overweight: Z-score $> +1SD$; and Obese: Z-score $> +2SD$.

Gestational diabetes

The Brazilian Society of Diabetes defines gestational diabetes as any level of glucose intolerance with onset or diagnosis during gestation [23]. Threshold values for diagnosis of GDM recommended by the Brazilian Society of Diabetes are: fasting plasma glucose ≥ 92 mg/dL, 1-h plasma glucose after a 75 g oral glucose load ≥ 180 mg/dL or 2-h plasma glucose after a 75 g oral glucose load ≥ 153 mg/dL. The diagnosis of gestational diabetes was identified by review of hospital medical records and later confirmed through detailed questions about diabetes during the in-hospital interview with the mother.

Breastfeeding initiation

During the first interview, conducted between 24 and 48 hours postpartum, the mothers were asked if they had breastfed their child any time until that moment. If the answer was positive, they were asked to indicate how long (in minutes or hours) after delivery they had breastfed for the first time. The answer was categorized in more or less than 24 hours after delivery and it is presented as the primary outcome in the analysis.

Ethics

Approval to conduct this study has been granted by the Human Research Ethics Committees of the Hospital de Clínicas de Porto Alegre (reference n° 110097) and of the Grupo Hospitalar Conceição (reference n° 11027).

Statistical analysis

Characteristics of the study sample according to pre-pregnancy BMI and gestational diabetes were compared using Pearson's Chi Square test, Student T-test, and Fischer's Exact test, as appropriate. The frequency of BF initiation in the first 24 hours after delivery was compared according to pre-pregnancy BMI and gestational diabetes diagnosis using Pearson's Chi Square test with Bonferroni correction. Poisson regression with robust error variance was used to estimate the adjusted relative risk of failure to initiate BF in the first 24 hours after delivery, with normal weight women without gestational diabetes as the reference group. All statistical tests were two-sided and $p < 0.05$ was considered statistically significant. Underweight women

($n = 5$) and women with type 1 diabetes ($n = 2$) and type 2 diabetes ($n = 2$) were not included in the analyses due to their reduced prevalence in the sample. All statistical analyses were performed using SPSS software (version 20.0; SPSS Inc, Chicago, IL).

Results

Within 295 eligible women, 46 refused to participate (15.6%). Underweight women and women with type 1 or 2 diabetes were not included in the analyses due to their reduced frequency in the sample. Complete information on maternal BMI and BF initiation were available for 219 participants (Fig 1).

Women in the included sample had no statistically significant differences compared to those who refused to participate regarding race/ethnicity ($p = 0.856$), schooling ($p = 0.628$), parity ($p = 0.718$) and type of delivery ($p = 0.476$), however the participants were significantly younger than those who refused to participate [mean (SD): 26.75 (6.8) vs. 29.10 (7.0) years, $p = 0.031$].

Characteristics of the study sample according to pre-pregnancy BMI and gestational diabetes (GDM) are described in Table 1. Pre-pregnancy BMI was higher in those with GDM compared to those without GDM in both categories of BMI. The frequency of multiparous women among overweight/obese was greater than among their healthy weight counterparts. Additionally, having GDM raised the frequency of cesarean delivery, but only among overweight/obese women.

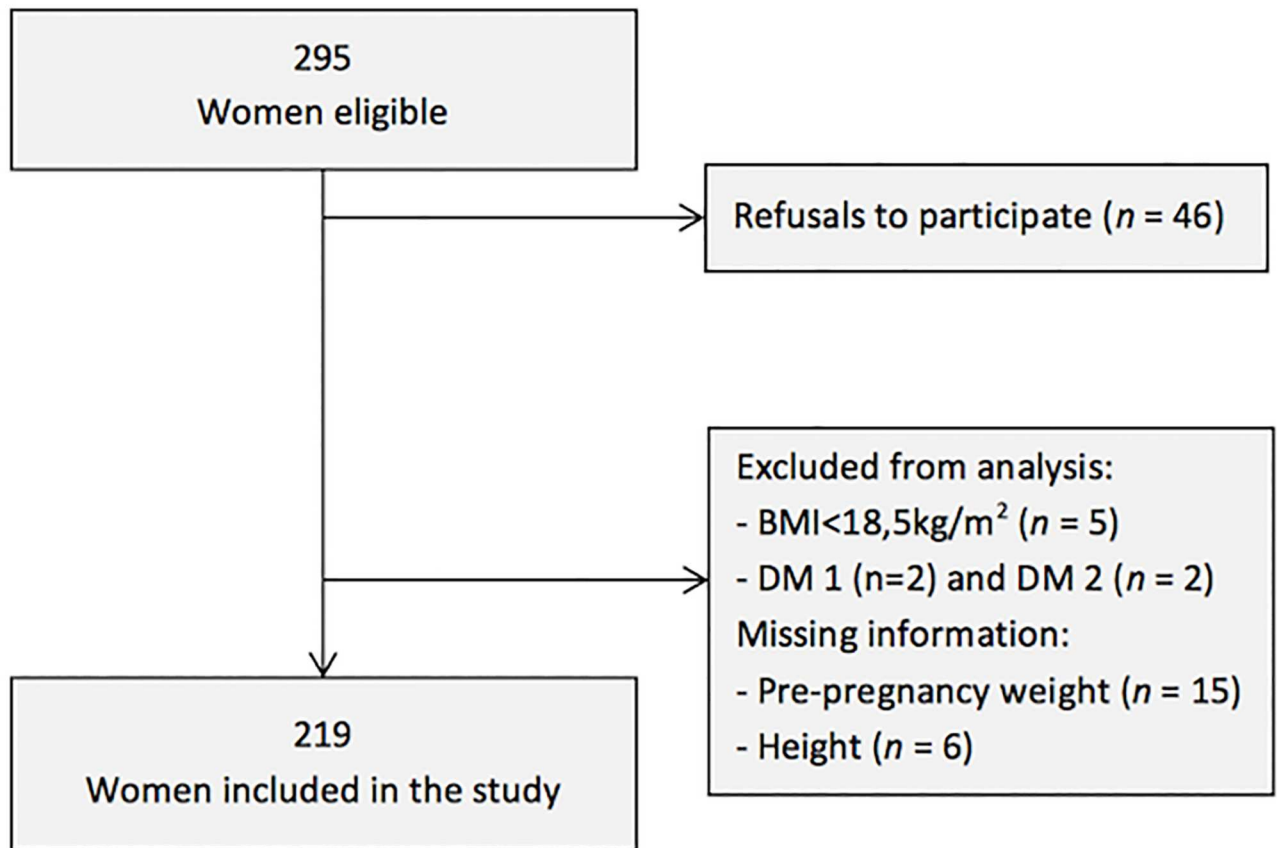


Fig 1. Progress of participants through the study.

<https://doi.org/10.1371/journal.pone.0194879.g001>

Table 1. Characteristics of the study sample according to pre-pregnancy body mass index and gestational diabetes, collected at 24–48 hours after delivery.

	Normal Weight (n = 102)			Overweight/Obese (n = 117)			P ³
	No GDM (n = 85)	DGM (n = 17)	P ¹	No GDM (n = 64)	DGM (n = 53)	P ²	
Age (years) ^a	25.3 (7.3)	28.4 (5.8)	0.109	27.0 (6.5)	28.1 (6.2)	0.360	0.083
Pre-pregnancy BMI (kg/m ²) ^a	21.7 (1.8)	22.7 (1.7)	0.026	29.4 (3.9)	31.2 (5.5)	0.036	<0.001
Schooling ^b			1.000			0.096	0.268
< 8 years	14 (16.5)	2 (11.8)		10 (15.6)	15 (28.3)		
≥ 8 years	71 (83.5)	15 (88.2)		54 (84.4)	38 (71.7)		
Parity ^b			0.425			0.222	0.014
None	44 (51.8)	7 (41.2)		25 (39.1)	15 (28.3)		
≥1	41 (48.2)	10 (58.8)		39 (60.9)	38 (71.7)		
Type of delivery ^b			0.260			0.021	0.623
Vaginal	57 (67.1)	14 (82.4)		48 (75.0)	29 (54.7)		
Cesarean	28 (32.9)	3 (17.6)		16 (25.0)	24 (45.3)		

Chi Square and Fisher's exact test used to compare categorical variables; t test used to compare continuous variables.

^a continuous variables, presented as mean (SD);

^b categorical variables, presented as n (%)

¹ P-value for comparison of women with and without GDM among the Normal weight group;

² P-value for comparison of women with and without GDM among the Overweight/obese group;

³ P-value for comparison of women Normal weight and women Overweight/obese.

<https://doi.org/10.1371/journal.pone.0194879.t001>

Most women were able to initiate BF within 12 hours from delivery (92.7%). Only eight (3.7%) women did not breastfeed in the first 24 hours postpartum. Of these, seven were overweight/obese (77.8%) and five had gestational diabetes (66.7%), with four of them having both conditions.

The percentage of mothers that did not breastfeed in the first day postpartum, according to pre-pregnancy BMI and GDM is presented in Fig 2. All women with a healthy pre-pregnancy weight without GDM managed to breastfeed in the first 24 hours postpartum. Among women with GDM and healthy weight 5.9% did not breastfeed in this period, while among overweight/obese women without GDM 4.8% had delayed initiation of BF. When both conditions were present, the frequency of delayed initiation raised to 7.5% ($p = 0.021$).

Table 2 indicates Poisson Regression analysis for the risk of not BF during the first 24h after delivery according to pre-pregnancy BMI and gestational diabetes, adjusted for cesarean delivery. Women who had both overweight/obesity and gestational diabetes presented the highest risk for delayed BF initiation among all groups.

Discussion

The results of this study indicate an additive interaction between maternal pre-pregnancy overweight/obesity and gestational diabetes on delayed BF initiation after 24 hours postpartum. This result added a new insight to other studies that also found worse outcomes for BF initiation in overweight and obese women [4, 5, 17, 24] and in women with gestational diabetes [10, 11, 25].

There is an international consensus on the importance of early BF initiation, since it has been strongly associated with reduced neonatal and early infant mortality [26–28]. Smith et al., conducted a large systematic review and meta-analysis and found that mortality risk for infants who were breastfed for the first time after 24 hours was more than 100% higher compared

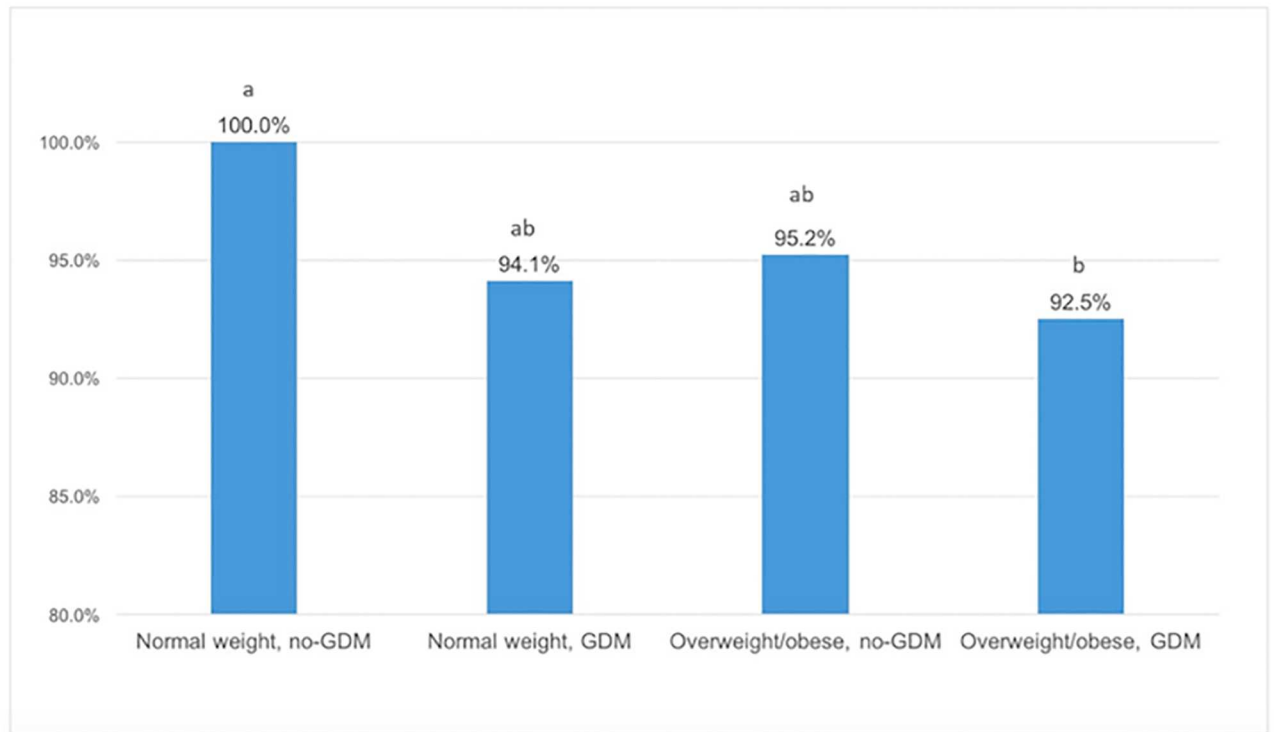


Fig 2. Percentage of mothers that initiated BF in the first 24 hours after delivery, according to pre-pregnancy body mass index and gestational diabetes. Chi Square test with Bonferroni correction. Distinct superscript letters represent statistical difference ($p < 0.05$).

<https://doi.org/10.1371/journal.pone.0194879.g002>

with infants that were breastfed within the first hour of life [28]. Also, previous studies indicate that women who did not breastfeed in the first days after delivery were less able to sustain any or exclusive BF 4 weeks postpartum [19], and had earlier weaning [29].

Poor lactation performance in overweight and obese women may be explained by several different aspects, such as: abnormal development of mammary glands before, during and after pregnancy [30, 31]; complications of pregnancy, including higher rates of cesarean section [9, 32], preterm birth [9, 32], and babies with macrosomia [9, 33]; psychosocial, economic and cultural factors, such as postpartum depression [34, 35], body image concerns [36, 37], education and income [38]; maternal confidence and behavior beliefs [38]; latching and positioning difficulties [39–41]; less support from family, friends and hospital staff [38, 42, 43]; and hormonal and metabolic alterations [44, 45].

Table 2. Adjusted relative risk for failure to initiate BF during the first 24 hours after delivery.

	aRR (95% confidence interval)	<i>p</i>
Normal weight, no-GDM	1 (reference)	-
Normal weight, GDM	1.063 (0.956; 1.182)	0.257
Overweight/Obese, no-GDM	1.050 (0.998; 1.105)	0.060
Overweight/Obese, GDM	1.072 (1.006; 1.141)	0.032

aRR, Adjusted Relative Risk; GDM, Gestational Diabetes. Poisson regression with robust error variance, adjusted for type of delivery

<https://doi.org/10.1371/journal.pone.0194879.t002>

Gestational diabetes, on the other hand, may negatively impact BF initiation with an effect on mammary development and milk production. Emerging evidence connect different levels of glucose intolerance with poor development of mammary glands during pregnancy [46, 47], delayed onset of lactogenesis [48, 49], and reduced milk production in early and mature lactation [50, 51]. This delayed milk production is thought to be a result of protein tyrosine phosphatase, receptor type F overexpression in the mammary gland caused by decreased insulin sensitivity [50].

Most studies analyzed only the effect of maternal overweight or the effect of gestational diabetes over BF initiation. Matias et al., however, assessed the impact of maternal obesity evaluating GDM women from the Study of Women, Infant Feeding and Type 2 Diabetes After GDM Pregnancy (SWIFT), and found that pre-pregnancy obesity was independently associated with delayed lactogenesis in women with GDM [52]. The authors, however, were not able to identify whether the outcome was caused only by maternal obesity or if obesity and GDM were interacting to produce this effect, since there were no control women without diabetes in the study.

Given all the evidence of worse BF outcome in overweight women, a few intervention studies have been conducted to improve BF initiation rates for this group. Rasmussen et al. conducted two low-intensity interventions with obese women [53]: the first one design to give telephone support to mothers on BF and the second one to provide manual or electric pumps for milk supply. Neither of them were successful in enhancing BF rates. Another study provided intensive peer counseling to overweight/obese mothers, including prenatal visits, daily in-hospital visits, postpartum home visits and phone calls [54]. This intervention also had no impact on BF initiation and on exclusive BF rates at 1, 3, or 6 months postpartum.

Antenatal breast expression is lately being applied, especially to diabetic women, in order to store colostrum and handle neonatal hypoglycemia and as a form of hastening milk production and preventing delayed initiation of BF after delivery [55]. The efficacy and safety of this practice, however, is yet to be proven, with some small studies indicating that this procedure could induce preterm delivery and increase the frequency of NICU admissions [56]. Forster et al. conducted a big randomized control trial and found no harm in advising diabetic women with low risk of complications to express breast milk from 36 weeks' gestation, however the authors enunciated their concerns with the extrapolation of this results to other high risk groups without further research [57].

Our results were both strengthened and limited by the design of the IVAPSA Study. To our knowledge this is the first study to present women with and without GDM and women with and without overweight/obesity to investigate the interaction of these adverse conditions over BF initiation. Greater statistical power, with the increase in the number of participants in each group, however, would be substantial to carry out more complex statistical analysis. We can not refute the possibility that the aggravation of obesity for women with both overweight and GDM could, in part, be responsible for the increased relative risk for delayed BF in this group. Another strength of this study was the quality of the prospective data collection, especially on BF and maternal anthropometry.

Conclusion

This study indicates an additive interaction between maternal pre-pregnancy overweight/obesity and gestational diabetes on delayed BF initiation. Intervention studies that aim to improve BF initiation in these target groups are essential given the increasing rates of overweight and diabetes in women of reproductive age worldwide and the beneficial effect of BF for these women and their offspring.

Supporting information

S1 File. MDS_plos: Minimal data set for the analysis presented in this paper.
(SAV)

S2 File. MDS labels: Labels for the minimal data set.
(DOCX)

Acknowledgments

We would like to thank the researchers Clécio Homrich da Silva, Isabel Cristina Werlang, Juliana Rombaldi Bernardi and Vera Lucia Bossa, from the IVAPSA group, who were fully involved in the rationale and design of the IVAPSA project and contributed to data collection and analysis in all studies involving this project.

Author Contributions

Conceptualization: Tanara Vogel Pinheiro, Marcelo Zubaran Goldani.

Formal analysis: Tanara Vogel Pinheiro, Marcelo Zubaran Goldani.

Investigation: Tanara Vogel Pinheiro.

Methodology: Tanara Vogel Pinheiro, Marcelo Zubaran Goldani.

Resources: Marcelo Zubaran Goldani.

Writing – original draft: Tanara Vogel Pinheiro.

Writing – review & editing: Tanara Vogel Pinheiro, Marcelo Zubaran Goldani.

References

1. Williams EP, Mesidor M, Winters K, Dubbert PM, Wyatt SB. Overweight and Obesity: Prevalence, Consequences, and Causes of a Growing Public Health Problem. *Curr Obes Rep*. 2015; 4(3):363–70. <https://doi.org/10.1007/s13679-015-0169-4> PMID: 26627494
2. Zera C, McGirr S, Oken E. Screening for obesity in reproductive-aged women. *Prev Chronic Dis*. 2011; 8(6):A125. PMID: 22005618
3. Marchi J, Berg M, Dencker A, Olander EK, Begley C. Risks associated with obesity in pregnancy, for the mother and baby: a systematic review of reviews. *Obes Rev*. 2015; 16(8):621–38. <https://doi.org/10.1111/obr.12288> PMID: 26016557
4. Rasmussen KM. Association of maternal obesity before conception with poor lactation performance. *Annu Rev Nutr*. 2007; 27:103–21. <https://doi.org/10.1146/annurev.nutr.27.061406.093738> PMID: 17341160
5. Turcksin R, Bel S, Galjaard S, Devlieger R. Maternal obesity and breastfeeding intention, initiation, intensity and duration: a systematic review. *Matern Child Nutr*. 2014; 10(2):166–83. <https://doi.org/10.1111/j.1740-8709.2012.00439.x> PMID: 22905677
6. Bever Babendure J, Reifsnider E, Mendias E, Moramarco MW, Davila YR. Reduced breastfeeding rates among obese mothers: a review of contributing factors, clinical considerations and future directions. *Int Breastfeed J*. 2015; 10:21. <https://doi.org/10.1186/s13006-015-0046-5> PMID: 26140049
7. Torloni MR, Betran AP, Horta BL, Nakamura MU, Atallah AN, Moron AF, et al. Prepregnancy BMI and the risk of gestational diabetes: a systematic review of the literature with meta-analysis. *Obes Rev*. 2009; 10(2):194–203. <https://doi.org/10.1111/j.1467-789X.2008.00541.x> PMID: 19055539
8. Chu SY, Callaghan WM, Kim SY, Schmid CH, Lau J, England LJ, et al. Maternal obesity and risk of gestational diabetes mellitus. *Diabetes Care*. 2007; 30(8):2070–6. <https://doi.org/10.2337/dc06-2559a> PMID: 17416786
9. Poston L, Caleyachetty R, Cnattingius S, Corvalan C, Uauy R, Herring S, et al. Preconceptional and maternal obesity: epidemiology and health consequences. *Lancet Diabetes Endocrinol*. 2016; 4(12):1025–36. [https://doi.org/10.1016/S2213-8587\(16\)30217-0](https://doi.org/10.1016/S2213-8587(16)30217-0) PMID: 27743975

10. Chamberlain CR, Wilson AN, Amir LH, O'Dea K, Campbell S, Leonard D, et al. Low rates of predominant breastfeeding in hospital after gestational diabetes, particularly among Indigenous women in Australia. *Aust N Z J Public Health*. 2017; 41(2):144–50. <https://doi.org/10.1111/1753-6405.12629> PMID: 28110518
11. Oza-Frank R, Moreland JJ, McNamara K, Geraghty SR, Keim SA. Early Lactation and Infant Feeding Practices Differ by Maternal Gestational Diabetes History. *J Hum Lact*. 2016; 32(4):658–65. <https://doi.org/10.1177/0890334416663196> PMID: 27550377
12. Nommsen-Rivers LA. Does Insulin Explain the Relation between Maternal Obesity and Poor Lactation Outcomes? An Overview of the Literature. *Adv Nutr*. 2016; 7(2):407–14. <https://doi.org/10.3945/an.115.011007> PMID: 26980825
13. Schwarz EB, Brown JS, Creasman JM, Stuebe A, McClure CK, Van Den Eeden SK, et al. Lactation and maternal risk of type 2 diabetes: a population-based study. *Am J Med*. 2010; 123(9):863.e1–6.
14. Ostbye T, Krause KM, Swamy GK, Lovelady CA. Effect of breastfeeding on weight retention from one pregnancy to the next: results from the North Carolina WIC program. *Prev Med*. 2010; 51(5):368–72. <https://doi.org/10.1016/j.ypmed.2010.07.017> PMID: 20655944
15. Krause KM, Lovelady CA, Peterson BL, Chowdhury N, Ostbye T. Effect of breast-feeding on weight retention at 3 and 6 months postpartum: data from the North Carolina WIC Programme. *Public Health Nutr*. 2010; 13(12):2019–26. <https://doi.org/10.1017/S1368980010001503> PMID: 20519049
16. Mayer-Davis EJ, Rifas-Shiman SL, Zhou L, Hu FB, Colditz GA, Gillman MW. Breast-feeding and risk for childhood obesity: does maternal diabetes or obesity status matter? *Diabetes Care*. 2006; 29(10):2231–7. <https://doi.org/10.2337/dc06-0974> PMID: 17003298
17. Bider-Canfield Z, Martinez MP, Wang X, Yu W, Bautista MP, Brookey J, et al. Maternal obesity, gestational diabetes, breastfeeding and childhood overweight at age 2 years. *Pediatr Obes*. 2017; 12(2):171–8. <https://doi.org/10.1111/ijpo.12125> PMID: 26956226
18. Dugas C, Perron J, Kearney M, Mercier R, Tchernof A, Marc I, et al. Postnatal Prevention of Childhood Obesity in Offspring Prenatally Exposed to Gestational Diabetes mellitus: Where Are We Now? *Obes Facts*. 2017; 10(4):396–406. <https://doi.org/10.1159/000477407> PMID: 28848122
19. Brownell E, Howard CR, Lawrence RA, Dozier AM. Does Delayed Onset Lactogenesis II Predict the Cessation of Any or Exclusive Breastfeeding? *J Pediatr*. 2012; 161(4):608–14. <https://doi.org/10.1016/j.jpeds.2012.03.035> PMID: 22575242
20. Zhu P, Hao J, Jiang X, Huang K, Tao F. New insight into onset of lactation: mediating the negative effect of multiple perinatal biopsychosocial stress on breastfeeding duration. *Breastfeed Med*. 2013; 8:151–8. <https://doi.org/10.1089/bfm.2012.0010> PMID: 23057642
21. Bernardi JR, Ferreira CF, Nunes M, da Silva CH, Bosa VL, Silveira PP, et al. Impact of Perinatal Different Intrauterine Environments on Child Growth and Development in the First Six Months of Life—IVAPSA Birth Cohort: rationale, design, and methods. *BMC Pregnancy Childbirth*. 2012; 12:25. <https://doi.org/10.1186/1471-2393-12-25> PMID: 22471837
22. Rodacki CL, Fowler NE, Rodacki AL, Birch K. Stature loss and recovery in pregnant women with and without low back pain. *Arch Phys Med Rehabil*. 2003; 84(4):507–12. <https://doi.org/10.1053/apmr.2003.50119> PMID: 12690588
23. Milech A, Angelucci AP, Golbert A, Matheus A, Carrilho AJF, Ramalho AC, et al. *Diretrizes da Sociedade Brasileira de Diabetes (2015–2016)*. São Paulo: A C Farmacêutica; 2016. 348 p.
24. Amir LH, Donath S. A systematic review of maternal obesity and breastfeeding intention, initiation and duration. *BMC Pregnancy Childbirth*. 2007; 7:9. <https://doi.org/10.1186/1471-2393-7-9> PMID: 17608952
25. Verd S, de Sotto D, Fernandez C, Gutierrez A. The Effects of Mild Gestational Hyperglycemia on Exclusive Breastfeeding Cessation. *Nutrients*. 2016; 8(11).
26. Fawzi WW. Timing of initiation, patterns of breastfeeding, and infant survival: prospective analysis of pooled data from three randomised trials. 2016.
27. Debes AK, Kohli A, Walker N, Edmond K, Mullany LC. Time to initiation of breastfeeding and neonatal mortality and morbidity: a systematic review. *BMC Public Health*. 2013; 13 Suppl 3:S19.
28. Smith ER, Hurt L, Chowdhury R, Sinha B, Fawzi W, Edmond KM. Delayed breastfeeding initiation and infant survival: A systematic review and meta-analysis. *PLoS One*. 2017; 12(7):e0180722. <https://doi.org/10.1371/journal.pone.0180722> PMID: 28746353
29. Hossain MM, Department of Community Medicine FoMHS, UAE University P.O. Box 17666, Al Ain, United Arab Emirates, Reves RR, Denver Disease Control Service and University of Colorado Health Sciences Center Denver C, USA, Radwan MM, Epidemiology Study Center Bilbeis Sharqiya E, and Center for Infectious Diseases, University of Texas, Medical School and School of Public

- HealthHouston, Texas, USA, et al. The Timing of Breastfeeding Initiation and its Correlates in a Cohort of Rural Egyptian Infants. *Journal of Tropical Pediatrics*. 2017; 41(6):354–9.
30. Sejrnsen K, Purup S, Vestergaard M, Foldager J. High body weight gain and reduced bovine mammary growth: physiological basis and implications for milk yield potential. *Domest Anim Endocrinol*. 2000; 19(2):93–104. PMID: [11025189](#)
 31. Thorn SR, Purup S, Cohick WS, Vestergaard M, Sejrnsen K, Boisclair YR. Leptin does not act directly on mammary epithelial cells in prepubertal dairy heifers. *J Dairy Sci*. 2006; 89(5):1467–77. [https://doi.org/10.3168/jds.S0022-0302\(06\)72214-7](https://doi.org/10.3168/jds.S0022-0302(06)72214-7) PMID: [16606717](#)
 32. Ma RC, Schmidt MI, Tam WH, McIntyre HD, Catalano PM. Clinical management of pregnancy in the obese mother: before conception, during pregnancy, and post partum. *Lancet Diabetes Endocrinol*. 2016; 4(12):1037–49. [https://doi.org/10.1016/S2213-8587\(16\)30278-9](https://doi.org/10.1016/S2213-8587(16)30278-9) PMID: [27743977](#)
 33. Liu P, Xu L, Wang Y, Zhang Y, Du Y, Sun Y, et al. Association between perinatal outcomes and maternal pre-pregnancy body mass index. *Obes Rev*. 2016; 17(11):1091–102. <https://doi.org/10.1111/obr.12455> PMID: [27536879](#)
 34. Mannan M, Mamun A, Doi S, Clavarino A. Is there a bi-directional relationship between depression and obesity among adult men and women? Systematic review and bias-adjusted meta analysis. *Asian J Psychiatr*. 2016; 21:51–66. <https://doi.org/10.1016/j.ajp.2015.12.008> PMID: [27208458](#)
 35. Mehta UJ, Siega-Riz AM, Herring AH, Adair LS, Bentley ME. Maternal obesity, psychological factors, and breastfeeding initiation. *Breastfeed Med*. 2011; 6(6):369–76. <https://doi.org/10.1089/bfm.2010.0052> PMID: [21492019](#)
 36. Hauff LE, Demerath EW. Body image concerns and reduced breastfeeding duration in primiparous overweight and obese women. *Am J Hum Biol*. 2012; 24(3):339–49. <https://doi.org/10.1002/ajhb.22238> PMID: [22308116](#)
 37. Gavin AR, Simon GE, Ludman EJ. The association between obesity, depression, and educational attainment in women: the mediating role of body image dissatisfaction. *J Psychosom Res*. 2010; 69(6):573–81. <https://doi.org/10.1016/j.jpsychores.2010.05.001> PMID: [21109045](#)
 38. Hauff LE, Leonard SA, Rasmussen KM. Associations of maternal obesity and psychosocial factors with breastfeeding intention, initiation, and duration. *Am J Clin Nutr*. 2014; 99(3):524–34. <https://doi.org/10.3945/ajcn.113.071191> PMID: [24401717](#)
 39. Katz KA, Nilsson I, Rasmussen KM. Danish health care providers' perception of breastfeeding difficulty experienced by women who are obese, have large breasts, or both. *J Hum Lact*. 2010; 26(2):138–47. <https://doi.org/10.1177/0890334409349805> PMID: [19910520](#)
 40. Mok E, Multon C, Piguell L, Barroso E, Goua V, Christin P, et al. Decreased full breastfeeding, altered practices, perceptions, and infant weight change of prepregnant obese women: a need for extra support. *Pediatrics*. 2008; 121(5):e1319–24. <https://doi.org/10.1542/peds.2007-2747> PMID: [18450874](#)
 41. Garner CD, McKenzie SA, Devine CM, Thornburg LL, Rasmussen KM. Obese women experience multiple challenges with breastfeeding that are either unique or exacerbated by their obesity: discoveries from a longitudinal, qualitative study. *Matern Child Nutr*. 2016.
 42. Kair LR, Colaizy TT. Obese Mothers have Lower Odds of Experiencing Pro-breastfeeding Hospital Practices than Mothers of Normal Weight: CDC Pregnancy Risk Assessment Monitoring System (PRAMS), 2004–2008. *Matern Child Health J*. 2016; 20(3):593–601. <https://doi.org/10.1007/s10995-015-1858-z> PMID: [26515471](#)
 43. Garner CD, Ratcliff SL, Devine CM, Thornburg LL, Rasmussen KM. Health Professionals' Experiences Providing Breastfeeding-Related Care for Obese Women. *Breastfeed Med* [Internet]. 2014; 9(10): [503–9 pp.]. Available from: <http://dx.doi.org/10.1089/bfm.2014.0104>.
 44. Rasmussen KM, Kjolhede CL. Prepregnant overweight and obesity diminish the prolactin response to suckling in the first week postpartum. *Pediatrics*. 2004; 113(5):e465–71. PMID: [15121990](#)
 45. Moynihan AT, Hehir MP, Glavey SV, Smith TJ, Morrison JJ. Inhibitory effect of leptin on human uterine contractility in vitro. *Am J Obstet Gynecol*. 2006; 195(2):504–9. <https://doi.org/10.1016/j.ajog.2006.01.106> PMID: [16647683](#)
 46. Neville MC, Webb P, Ramanathan P, Mannino MP, Pecorini C, Monks J, et al. The insulin receptor plays an important role in secretory differentiation in the mammary gland. *Am J Physiol Endocrinol Metab*. 2013; 305(9):E1103–14. <https://doi.org/10.1152/ajpendo.00337.2013> PMID: [23982156](#)
 47. Vanky E, Nordskar JJ, Leithe H, Hjorth-Hansen AK, Martinussen M, Carlsen SM. Breast size increment during pregnancy and breastfeeding in mothers with polycystic ovary syndrome: a follow-up study of a randomised controlled trial on metformin versus placebo. *Bjog*. 2012; 119(11):1403–9. <https://doi.org/10.1111/j.1471-0528.2012.03449.x> PMID: [22827167](#)

48. Neubauer SH, Ferris AM, Chase CG, Fanelli J, Thompson CA, Lammi-Keefe CJ, et al. Delayed lactogenesis in women with insulin-dependent diabetes mellitus. *Am J Clin Nutr* [Internet]. 1993 Jul; 58(1): [54–60 pp.]. Available from: <http://dx.doi.org/10.1093/ajcn/58.1.54>
49. Nommsen-Rivers LA, Dolan LM, Huang B. Timing of Stage II Lactogenesis Is Predicted by Antenatal Metabolic Health in a Cohort of Primiparas. *Breastfeed Med* [Internet]. 2012; 7(1):[43–9 pp.]. Available from: <http://dx.doi.org/10.1089/bfm.2011.0007>.
50. Lemay DG, Ballard OA, Hughes MA, Morrow AL, Horseman ND, Nommsen-Rivers LA. RNA Sequencing of the Human Milk Fat Layer Transcriptome Reveals Distinct Gene Expression Profiles at Three Stages of Lactation. *PLoS One* [Internet]. 2013; 8(7). Available from: <http://dx.doi.org/10.1371/journal.pone.0067531>.
51. Riddle SW, Nommsen-Rivers LA. A Case Control Study of Diabetes During Pregnancy and Low Milk Supply. *Breastfeed Med*. 2016; 11(2):80–5. <https://doi.org/10.1089/bfm.2015.0120> PMID: 26859784
52. Matias SL, Dewey KG, Quesenberry CP, Gunderson EP. Maternal prepregnancy obesity and insulin treatment during pregnancy are independently associated with delayed lactogenesis in women with recent gestational diabetes mellitus. *Am J Clin Nutr* [Internet]. 2014; 99(1):[115–21 pp.]. Available from: <http://dx.doi.org/10.3945/ajcn.113.073049>.
53. Rasmussen KM, Dieterich CM, Zelek ST, Altabet JD, Kjolhede CL. Interventions to increase the duration of breastfeeding in obese mothers: the Bassett Improving Breastfeeding Study. *Breastfeed Med*. 2011; 6(2):69–75. <https://doi.org/10.1089/bfm.2010.0014> PMID: 20958105
54. Chapman DJ, Morel K, Bermúdez-Millán A, Young S, Damio G, Pérez-Escamilla R. Breastfeeding Education and Support Trial for Overweight and Obese Women: A Randomized Trial. *Pediatrics*. 131 2013. p. e162–70. <https://doi.org/10.1542/peds.2012-0688> PMID: 23209111
55. Lamba S, Chopra S, Negi M. Effect of Antenatal Breast Milk Expression at Term Pregnancy to Improve Post Natal Lactational Performance. *J Obstet Gynaecol India*. 2016; 66(1):30–4. <https://doi.org/10.1007/s13224-014-0648-7> PMID: 26924904
56. Chapman T, Pincombe J, Harris M. Antenatal breast expression: a critical review of the literature. *Midwifery*. 2013; 29(3):203–10. <https://doi.org/10.1016/j.midw.2011.12.013> PMID: 22342171
57. Forster DA, Moorhead AM, Jacobs SE, Davis PG, Walker SP, McEgan KM, et al. Advising women with diabetes in pregnancy to express breastmilk in late pregnancy (Diabetes and Antenatal Milk Expressing [DAME]): a multicentre, unblinded, randomised controlled trial. *Lancet*. 2017; 389(10085):2204–13. [https://doi.org/10.1016/S0140-6736\(17\)31373-9](https://doi.org/10.1016/S0140-6736(17)31373-9) PMID: 28589894