



UvA-DARE (Digital Academic Repository)

Barriers and challenges in hyperemesis gravidarum research

Grooten, I.J.

Publication date

2017

Document Version

Other version

License

Other

[Link to publication](#)

Citation for published version (APA):

Grooten, I. J. (2017). *Barriers and challenges in hyperemesis gravidarum research*. [Thesis, fully internal, Universiteit van Amsterdam].

General rights

It is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), other than for strictly personal, individual use, unless the work is under an open content license (like Creative Commons).

Disclaimer/Complaints regulations

If you believe that digital publication of certain material infringes any of your rights or (privacy) interests, please let the Library know, stating your reasons. In case of a legitimate complaint, the Library will make the material inaccessible and/or remove it from the website. Please Ask the Library: <https://uba.uva.nl/en/contact>, or a letter to: Library of the University of Amsterdam, Secretariat, P.O. Box 19185, 1000 GD Amsterdam, The Netherlands. You will be contacted as soon as possible.

1.1

Barriers and challenges in hyperemesis gravidarum research

IJ Grooten, TJ Roseboom, RC Painter

Nutrition and Metabolic Insights, 2015;8:33–39

SUMMARY

Nausea and occasional vomiting in early pregnancy (NVP) are common. When vomiting is severe or protracted it is referred to as hyperemesis gravidarum (HG). HG affects up to 3% of pregnancies and is characterised by weight loss, dehydration, electrolyte imbalance and the need of hospital admission. HG has significant consequences for maternal wellbeing, is associated with adverse birth outcomes and leads to major health care costs. Treatment options are symptomatic, hampered by the lack of evidence-based options including studies on nutritional interventions. One of the reasons for this lack of evidence is the use of a broad range of definitions and outcome measures. An internationally accepted definition and the formulation of core outcomes would facilitate meta-analysis of trial results and implementation of evidence in guidelines, to ultimately improve patient care.

INTRODUCTION

Progress in the understanding of the etiology and treatment of hyperemesis gravidarum (HG) has been slow in recent decades.¹⁻³ This might be due to difficulty in pinpointing a cause and presumed psychogenic aetiology, the latter remaining unsupported by evidence.⁴ Clinicians may find it challenging to differentiate HG from 'physiological' nausea and vomiting in pregnancy (NVP) and consider HG a self-limiting, non-life-threatening affliction, lacking curative treatment options. 'Therapeutic nihilism' among caregivers may lead to poor care for HG patients.⁵ Research is limited to single small study groups with no or minimal funding, often yielding underpowered studies. Meta-analysis has the ability to increase power and is considered to provide the highest level of evidence.⁶ It is therefore unfortunate that attempts to perform meta-analyses on HG etiology, diagnosis and treatment options are hampered by the lack of consensus on the exact definition of HG and most relevant outcomes.^{7,8}

In addition to the forthcoming Cochrane review on the treatment of HG,⁹ several systematic reviews on HG have been published, most recently in 2014.¹ Here, we therefore will not attempt to present an extensive overview of all evidence, but will summarise briefly where HG research stands today and suggest ways to bring HG research forward.

What is hyperemesis?

Nausea and occasional vomiting in early pregnancy (NVP) is common, affecting 50%-80% of pregnant women in the first half of gestation,^{10,11} with major impact on maternal wellbeing and quality of life.¹² When vomiting is severe or protracted, it is often referred to as hyperemesis gravidarum (HG).¹³ HG is much less common than NVP, affecting 0.2-3.6% of pregnant women.^{14,15} Despite this relatively low incidence, HG is the single most frequent reason for hospital admission in the first half of pregnancy,¹⁶ with substantial health care costs. In 2012, the costs per HG patient in the USA were estimated at \$47,351.¹⁷

Little is known about the pathophysiology of HG. Proposed underlying causes are related to maternal endocrine function, placental growth and function and gastrointestinal conditions. Early pregnancy rise of human chorionic gonadotrophin (hCG) coincides with the occurrence of HG and several conditions associated with increased hCG are more prevalent in HG patients.¹⁸ It is thought that rising hCG levels may cause nausea via several pathways, including through rises of other hormones such as thyroid hormone and estradiol. Although the majority of studies on hCG and HG describe a positive association, heterogenic use of definitions and outcome measures limit the possibility of meta-analysis and findings remain inconclusive.⁷

High hCG levels in the second trimester may be caused by abnormal placentation. This is in line with the study of Bolin *et al.* who found placental dysfunction disorders to be associated with HG.¹⁹ Colonisation with the gastric bacterium *Helicobacter pylori* (*H pylori*) is associated with an increased risk of HG, but the strength and size of these associations vary between different populations and countries.²⁰ Despite many causes for HG have been proposed, no single mechanism has yet been identified, making a multifactorial origin more likely.¹

Large registry studies have identified several risk factors. The recurrence rate of HG after a first pregnancy complicated by HG ranges from 15% to 81% and may vary with HG definition based on symptoms or hospital admission and study methodology.^{21, 22} In addition to having had HG in a previous pregnancy, having a mother or sister who had HG are important risk factors. This suggests yet-unidentified genetic factors are involved in HG aetiology.^{21, 23} Furthermore, young age, nulliparity, non-Western ethnicity, low socioeconomic status, diabetes, hypertensive disorders, psychopathology, assisted reproductive techniques, female fetus and multiple gestation are all associated with HG.¹⁵ In the past, psychopathology has been suggested to play a causative role,²⁴ but convincing evidence is lacking.⁴

How do we diagnose HG?

HG is a clinical diagnosis, made after other causes of nausea and vomiting have been excluded. Commonly used criteria include nausea and vomiting before 20 weeks gestation, the need for hospitalisation, weight loss, dehydration and electrolyte imbalances.^{13, 25} Patients generally undergo a diagnostic work-up including laboratory testing and ultrasound to rule out molar pregnancy or multiple gestation. However, such a work-up is not based on reliable data from the literature. A recent review showed that, despite its widespread use as a major diagnostic criterion, there is little evidence to support the use of ketonuria in HG diagnosis due to its lack of association with symptom severity, admission duration or readmission rates.⁷ There is no international consensus on the exact definition of HG.

Do we have treatment options?

Management of care

Hospital admission, intravenous rehydration and antiemetic medication are often applied.¹³ The only randomised controlled trial investigating rehydration for HG did not show any benefits after 24 hours of treatment when 5% dextrose was added to 0.9% saline solution on ketonuria, nausea and vomiting symptoms.²⁶ Early thiamine (vitamin B1) supplementation before the administration of dextrose is mandatory to

prevent the development of Wernicke encephalopathy.¹³ It could be debated whether HG should always be treated in hospital. McCarthy *et al.* recently showed that day care management of HG reduced total days of inpatient stay without affecting patient satisfaction.²⁷ Another trial on the effectiveness of outpatient management of HG in reducing symptoms is still ongoing (ISRCTN24659467).

Pharmacologic interventions

According to the most recent Cochrane review on interventions for NVP, evidence on the effectiveness of pyridoxine (vitamin B6), ginger and antiemetic medication is inconsistent.³ Pyridoxine mainly reduces nausea but not vomiting. However, pyridoxine in combination with antihistamines (H1-receptor blockers including doxylamine and meclizine) significantly reduces nausea and vomiting symptoms and is not associated with teratogenic effects.²⁸ Phenothiazines (e.g. phenergan) and dopamine-antagonists (e.g. metoclopramide) may be used as a second-line treatment followed by 5-hydroxytryptamine₃-receptor antagonists (e.g. ondansetron). They are all described to reduce nausea and vomiting symptoms but might have maternal side-effects, and potential teratogenic effects are less studied.¹³ The use of corticosteroids (CCS) has been advocated.²⁹ CCS are often used for chemotherapy induced nausea and vomiting, but strong evidence for their effectiveness in the management of HG is missing.³⁰ Several case studies have reported that *H pylori* eradication can effectively relieve symptoms in women with persistent HG who are unresponsive to conventional treatment, although a randomised controlled trial has yet to be conducted.³¹⁻³³

Non-pharmacologic interventions

Mindfulness-based cognitive therapy (MBCT) just proved to effectively reduce symptoms in women with moderate NVP,³⁴ but has not yet been studied in women with HG. Dietary advice based on anecdotal evidence include frequent small meals and avoidance of spicy, fatty and odorous foods.³⁵ Nutritional interventions for HG have never been studied in clinical trials. This is surprising because poor nutritional intake^{36, 37} and maternal weight loss^{37, 38} are important features of HG, while substantial evidence indicates that poor maternal nutrition in early gestation has permanent detrimental effects on short- and long-term offspring health.³⁹ Small case series indicated that enteral tube feeding alleviated symptoms and was well tolerated when continued in a home setting.^{40, 41} Furthermore, a retrospective cohort study by Stokke *et al.* suggested that enteral tube feeding improved maternal weight gain during pregnancy in women with severe HG and substantial early pregnancy weight loss.⁴² Currently, the first randomized controlled trial (RCT) on enteral tube feeding in HG patients is being performed (NTR4197; www.studies-obsgyn.nl/mother), to investigate

the extent to which enteral tube feeding can affect birth outcomes, as well as reducing nausea and vomiting symptoms or time in hospital.

Are there adverse maternal outcomes?

Wernicke encephalopathy is a very rare but potentially life-threatening complication of HG, caused by thiamine deficiency.¹³ Patients are also at risk of developing refeeding syndrome when nutrition is not introduced carefully after a period of starvation.⁴³ A much more common consequence of HG with great impact is the psychological distress experienced by patients. This might result in pregnancy termination⁴⁴ and increases the risk of depression, anxiety and posttraumatic stress disorder.^{45, 46} No studies so far have yet addressed whether potential treatments have any beneficial effects on maternal wellbeing.

Are there adverse offspring outcomes?

HG is repeatedly associated with adverse birth outcomes. A systematic review and meta-analysis on offspring consequences of HG showed increased risk of low birth weight, small for gestational age and prematurity.⁴⁷ However, not all studies found adverse outcomes as a consequence of HG,⁴⁸ which might be due to heterogeneous patient populations. Maternal characteristics partly determine pregnancy outcome in HG patients¹⁵ but it is also likely that other factors associated with HG, such as insufficient weight gain during pregnancy,⁴² maternal stress responses during pregnancy⁴⁹ or *H pylori* infection involving the placenta⁵⁰ could each contribute to adverse birth outcomes associated with HG.

Thus far, reports on possible long-term consequences of HG beyond the perinatal period are limited. One study reported the association of HG and testicular cancer,⁵¹ but a large registry study found no increased cancer risk in HG offspring.⁵² Two studies reported on neurodevelopment in children with conflicting findings,^{53, 54} although there are some indications that risk of psychological and behavioural problems in adulthood is increased.⁵⁵ Lastly, Ayyavoo *et al.* recently showed that insulin sensitivity in children of hyperemetic mothers is decreased,⁵⁶ and others found that early pregnancy weight loss was associated with increased blood pressure in offspring,⁵⁷ suggesting that risk for cardiometabolic disease in later life might be increased.

Can we improve HG research?

HG severely impacts the lives of many women and their families and better understanding of causes, consequences and treatment options is needed. The lack of guidelines for the treatment of HG might have its origin in the inability to synthesize

results across studies in systematic reviews. It is clear that the lack of consensus on a definition, diagnostic criteria and classification of HG can hamper progress in the context of both clinical trials and clinical practice. Primary outcomes for HG are numerous, often differentially reported and performed in small trials.

Tables 1 and 2 illustrate the variations in HG-defining characteristics and outcome measures in eight clinical trials investigating the effectiveness of metoclopramide and promethazine.⁵⁸⁻⁶⁵ We chose these studies because they were specifically designed to study the effectiveness of antiemetic medication in patients with HG (and not NVP). Although there are similarities, only the trials by Abas⁵⁸ and Tan⁶⁰ used the same set of defining characteristics and outcome measures (Table 1). Both trials were conducted by the same research group. Gestational age^{58-62, 64, 65} and ketonuria^{58-61, 63-65} were included in the definition by all but one trial. Sullivan⁶³ did not include gestational age, Neri⁶² did not include ketonuria. Weight loss^{59, 61-65}, vomiting episodes^{59, 61, 62, 64, 65} and hospitalisation for HG^{58, 60, 61, 64} were used by at least four studies but these did not completely overlap. Also the definition of each criterion varied widely. Regarding trial outcomes (Table 2), five studies reported vomiting episodes and nausea severity,^{58-60, 62, 65} of which four^{58, 60, 62, 65} also included some, but not identical, measure of wellbeing. Interestingly, only one study included birth outcomes.⁶²

Not only varying outcome measures, but also the diversity of interventions studied limit the possibility of meta-analysis. Communicating study protocols might overcome this problem. Furthermore, the patient perspective is hardly recognized to play a role in HG management⁶⁶ while patient involvement could help prioritize the research agenda. In other fields of medicine, this problem has long been acknowledged and methods to reach consensus on definition and core outcome sets (COS) have been developed.^{67, 68} The Delphi process is a method that can be used to reach consensus among relevant stakeholders, including patients, to formulate a definition and COS that should be reported in clinical trials⁶⁹ to ultimately improve implementation of research findings in clinical practice. Currently a Delphi process to formulate a definition and COS for HG is in preparation (www.comet-initiative.org).

CONCLUSION

If clinicians do not know how to approach and treat HG patients, this is likely to impact patient care. Although an internationally accepted definition and the formulation of a core outcome set is no panacea, we believe researchers, clinicians and patients would benefit greatly. Bringing stakeholders together could furthermore increase collaborative research on HG and enable prioritising research questions that are not just relevant to researchers and clinicians but also relevant to patients.

TABLE 1. HG-defining characteristics in published randomised controlled trials on metoclopramide and/or promethazine for HG

Trial	Comparison	Defining characteristics							
		Hospitalisation for HG ^a	Gestational age ^b	Vomiting episodes ^c	Dehydration ^d	Ketonuria ^e	Electrolyte disturbances ^f	Weight loss ^g	Antiemetics ^d
Abas ^{58*} (2014)	metoclopramide vs ondansetron	X	X		X	X			
Kashifard ⁵⁹ (2013)	metoclopramide vs ondansetron		X	X		X		X	
Tan ^{60*} (2010)	metoclopramide vs promethazine	X	X		X	X			
Bondok ⁶¹ (2006)	metoclopramide vs hydrocortisone	X	X	X		X		X	
Neri ⁶² (2005)	metoclopramide vs acupuncture		X	X	X			X	
Sullivan ⁶³ (1996)	promethazine vs ondansetron				X	X	X	X	X
Safari ⁶⁴ (1998)	promethazine vs methylprednisolone	X	X	X		X		X	
Ziaei ⁶⁵ (2004)	promethazine vs prednisolone		X	X		X		X	

^a Abas⁵⁸ Tan⁶⁰ Sullivan⁶³ Any hospitalisation for HG; Bondok⁶¹ ICU admission for HG; Safari⁶⁴ Hospitalisation or readmission for HG

^b Ziaei⁶⁵ GA 6-12 weeks; Neri⁶² GA <12 weeks; Kashifard⁵⁹ GA <16 weeks; Abas⁵⁸ Tan⁶⁰ Bondok⁶¹ Safari⁶⁴ GA ≤16 weeks

^c Kashifard⁵⁹ vomiting 3 times per day; Ziaei⁶⁵ vomiting >3 times per day during last 72 hrs; Bondok⁶¹ Safari⁶⁴ persistent vomiting; Neri⁶² any nausea and vomiting

^d Sullivan⁶³ and/or >2 ER visits resulting in i.v. rehydration or promethazine suppositories

^e Abas⁵⁸ ketonuria ≥2+; Sullivan⁶³ ketonuria >80 mg/dl; Ziaei⁶⁵ ketonuria not responding to dietary manipulation; Kashifard⁵⁹ Tan⁶⁰ Bondok⁶¹ Safari⁶⁴ any ketonuria

^f Sullivan⁶³ potassium <3.0 mEq/dl or sodium <134 mEq/dl and/or positive serum acetone

^g Sullivan⁶³ weight loss ≥2.25kg; Kashifard⁵⁹ weight loss >3 kg; Bondok⁶¹ Neri⁶² weight loss >5% pre-pregnancy weight; Safari⁶⁴ Ziaei⁶⁵ any weight loss

* Same research group

TABLE 2. Outcome measures in published randomised controlled trials on metoclopramide and/or promethazine for HG

Trial	Comparison	Outcome measures								
		Vomiting episodes ^a	Nausea severity ^b	Ketonuria ^c	Wellbeing ^d	Length of hospital stay ^e	Readmission ^f	Weight gain ^g	Side-effects ^h	Other ⁱ
Abas ^{58*} (2014)	metoclopramide vs ondansetron	X	X	X	X	X				X
Kashifard ⁵⁹ (2013)	metoclopramide vs ondansetron	X	X							
Tan ^{60*} (2010)	metoclopramide vs promethazine	X	X	X	X	X			X	X
Bondok ⁶¹ (2006)	metoclopramide vs hydrocortisone	X					X			X
Neri ⁶² (2005)	metoclopramide vs acupuncture	X	X		X				X	X
Sullivan ⁶³ (1996)	promethazine vs ondansetron		X	X		X		X	X	X
Safari ⁶⁴ (1998)	promethazine vs methylprednisolone	X			X		X			X
Ziaei ⁶⁵ (2004)	promethazine vs prednisolone	X	X		X					

^a Abas,⁵⁸ Tan,⁶⁰ Ziaei⁶⁵ number of episodes; Kashifard⁵⁹ VAS-score; Neri⁶² vomiting episodes improvement; Safari⁶⁴ symptom improvement based on vomiting episodes, ability to tolerate fluids and wellbeing

^b Abas,⁵⁸ Kashifard,⁵⁹ Tan,⁶⁰ Sullivan,⁶³ Ziaei⁶⁵ VAS-score; Neri⁶² nausea severity improvement

^c Sullivan⁶³ daily ketonuria

^d Abas,⁵⁸ Tan,⁶⁰ VAS-score; Neri⁶² daily functioning; Ziaei⁶⁵ sickness

^e Length of hospital stay following the intervention

^f Bondok⁶¹ ICU readmission within 2 weeks of study completion; Safari⁶⁴ hospital readmission within 2 weeks after start intervention

^g Sullivan⁶³ daily weight

^h Abas,⁵⁸ Tan,⁶⁰ Sullivan⁶³ recording of symptoms

ⁱ Abas,⁵⁸ Tan,⁶⁰ Sullivan⁶³ treatment failure; Abas,⁵⁸ Tan,⁶⁰ Sullivan⁶³ use of antiemetics; Tan⁶⁰ duration of i.v. rehydration; Bondok⁶¹ serum albumin; Neri⁶² food intake improvement; Neri⁶² birth outcomes

* Same research group

REFERENCES

1. McCarthy FP, Lutomski JE, Greene RA. Hyperemesis gravidarum: current perspectives. *International journal of women's health*. 2014;6:719-25.
2. Festin M. Nausea and vomiting in early pregnancy. *BMJ clinical evidence*. 2014;2014.
3. Matthews A, Haas DM, O'Mathuna DP, Dowswell T. Interventions for nausea and vomiting in early pregnancy. *Cochrane Database Syst Rev*. 2015;9:CD007575.
4. Munch S. Chicken or the egg? The biological-psychological controversy surrounding hyperemesis gravidarum. *Soc Sci Med*. 2002;55(7):1267-78.
5. Power Z, Campbell M, Kilcoyne P, Kitchener H, Waterman H. The Hyperemesis Impact of Symptoms Questionnaire: Development and validation of a clinical tool. *Int J Nurs Stud*. 2009;47(1):67-77.
6. Sackett DL, Rosenberg WMC, Gray JAM, Haynes RB, Richardson WS. Evidence based medicine: what it is and what it isn't. *BMJ*. 1996;312(7023):71-2.
7. Niemeijer MN, Grooten IJ, Vos N, Bais JM, van der Post JA, Mol BW, et al. Diagnostic markers for hyperemesis gravidarum: a systematic review and metaanalysis. *Am J Obstet Gynecol*. 2014;211(2):150 e1-15.
8. Matthews A, Dowswell T, Haas DM, Doyle M, O'Mathuna DP. Interventions for nausea and vomiting in early pregnancy. *Cochrane database of systematic reviews (Online)*. 2010(9):CD007575-CD.
9. Boelig Rupsa C, Berghella V, Kelly Anthony J, Barton Samantha J, Edwards Steve J. Interventions for treating hyperemesis gravidarum. *Cochrane Database of Systematic Reviews [Internet]*. 2013; (6). Available from: <http://onlinelibrary.wiley.com/doi/10.1002/14651858.CD010607/abstract>.
10. Gadsby R, Barnie-Adshead AM, Jagger C. A prospective study of nausea and vomiting during pregnancy. *Br J Gen Pract*. 1993;43(371):245-8.
11. Lacroix R, Eason E, Melzack R. Nausea and vomiting during pregnancy: A prospective study of its frequency, intensity, and patterns of change. *Am J Obstet Gynecol*. 2000;182(4):931-7.
12. Lacasse A, Rey E, Ferreira E, Morin C, Bérard A. Nausea and vomiting of pregnancy: what about quality of life? *BJOG*. 2008;115(12):1484-93.
13. Niebyl JR. Clinical practice. Nausea and vomiting in pregnancy. *N Engl J Med*. 2010;363(16):1544-50.
14. Matsuo K, Ushioda N, Nagamatsu M, Kimura T. Hyperemesis gravidarum in Eastern Asian population. *Gynecol Obstet Invest*. 2007;64(4):213-6.
15. Roseboom TJ, Ravelli ACJ, van der Post JA, Painter RC. Maternal characteristics largely explain poor pregnancy outcome after hyperemesis gravidarum. *Eur J Obstet Gynecol Reprod Biol*. 2011;156(1):56-9.
16. Adams MM, Harlass FE, Sarno AP, Read JA, Rawlings JS. Antenatal hospitalization among enlisted servicewomen, 1987-1990. *Obstet Gynecol*. 1994;84(1):35-9.
17. Piwko C, Koren G, Babashov V, Vicente C, Einarson TR. Economic burden of nausea and vomiting of pregnancy in the USA. *J Popul Ther Clin Pharmacol*. 2013;20(2):e149-60.
18. Verberg MFG, Gillott DJ, Al-Fardan N, Grudzinskas JG. Hyperemesis gravidarum, a literature review. *Hum Reprod Update*. 2005;11(5):527-39.
19. Bolin M, Åkerud H, Cnattingius S, Stephansson O, Wikström aK. Hyperemesis gravidarum and risks of placental dysfunction disorders: a population-based cohort study. *BJOG*. 2013;120(5):541-7.

20. Li L, Li L, Zhou X, Xiao S, Gu H, Zhang G. *Helicobacter pylori* infection is associated with an increased risk of hyperemesis gravidarum: a meta-analysis. *Gastroenterol Res Pract*. 2015;2015:278905-.
21. Vikanes A, Skjaerven R, Grjibovski AM, Gunnes N, Vangen S, Magnus P. Recurrence of hyperemesis gravidarum across generations: population based cohort study. *BMJ*. 2010;340:c2050.
22. Fejzo MS, Macgibbon KW, Romero R, Goodwin TM, Mullin PM. Recurrence risk of hyperemesis gravidarum. *Journal of midwifery & women's health*. 2011;56(2):132-6.
23. Zhang Y, Cantor RM, MacGibbon K, Romero R, Goodwin TM, Mullin PM, et al. Familial aggregation of hyperemesis gravidarum. *Am J Obstet Gynecol*. 2011;204(3):230 e1-7.
24. Buckwalter JG, Simpson SW. Psychological factors in the etiology and treatment of severe nausea and vomiting in pregnancy. *Am J Obstet Gynecol*. 2002;186(5, Supplement 2):S210-S4.
25. Fairweather DV. Nausea and vomiting in pregnancy. *Am J Obstet Gynecol*. 1968;102(1):135-75.
26. Tan PC, Norazilah MJ, Omar SZ. Dextrose saline compared with normal saline rehydration of hyperemesis gravidarum: a randomized controlled trial. *Obstet Gynecol*. 2013;121(2 Pt 1):291-8.
27. McCarthy FP, Murphy A, Khashan AS, McElroy B, Spillane N, Marchocki Z, et al. Day care compared with inpatient management of nausea and vomiting of pregnancy: a randomized controlled trial. *Obstet Gynecol*. 2014;124(4):743-8.
28. Nausea and vomiting of pregnancy. Practice Bulletin No. 153. American College of Obstetricians and Gynecologists. *Obstet Gynecol*. 2015;126:e12-24.
29. Jarvis S, Nelson-Piercy C. Management of nausea and vomiting in pregnancy. *BMJ*. 2011;342:d3606.
30. Grooten IJ, Vinke ME, Roseboom TJ, Painter RC. A systematic review and meta-analysis of the utility of corticosteroids in the treatment of hyperemesis gravidarum. *Nutr Metab Insights*. 2015;8(S1):1-10.
31. Jacoby EB, Porter KB. *Helicobacter pylori* infection and persistent hyperemesis gravidarum. *Am J Perinatol*. 1999;16(2):85-8.
32. Strachan BK, Jokhi RP, Filshie GM. Persistent hyperemesis gravidarum and *Helicobacter pylori*. *J Obstet Gynaecol*. 2000;20(4):427.
33. Mansour GM, Nashaat EH. *Helicobacter pylori* and hyperemesis gravidarum. *International Journal of Gynecology & Obstetrics*. 2009;106(1):63-4.
34. Faramarzi M, Yazdani S, Barat S. A RCT of psychotherapy in women with nausea and vomiting of pregnancy. *Hum Reprod*. 2015;30(12):2764-73.
35. Bischoff SC, Renzer C. Nausea and nutrition. *Auton Neurosci*. 2006;129(1-2):22-7.
36. van Stuijvenberg ME, Schabert I, Labadarios D, Nel JT. The nutritional status and treatment of patients with hyperemesis gravidarum. *Am J Obstet Gynecol*. 1995;172(5):1585-91.
37. Birkeland E, Stokke G, Tangvik RJ, Torkildsen EA, Boateng J, Wollen AL, et al. Norwegian PUQE (Pregnancy-Unique Quantification of Emesis and nausea) identifies patients with hyperemesis gravidarum and poor nutritional intake: a prospective cohort validation study. *PLoS One*. 2015;10(4):e0119962.

38. Fejzo MS, Poursharif B, Korst LM, Munch S, MacGibbon KW, Romero R, et al. Symptoms and pregnancy outcomes associated with extreme weight loss among women with hyperemesis gravidarum. *J Womens Health (Larchmt)*. 2009;18(12):1981-7.
39. Painter RC, Roseboom TJ, Bleker OP. Prenatal exposure to the Dutch famine and disease in later life: An overview. *Reprod Toxicol*. 2005;20(3):345-52.
40. Hsu JJ, Clark-Glena R, Nelson DK, Kim CH. Nasogastric enteral feeding in the management of hyperemesis gravidarum. *Obstet Gynecol*. 1996;88(3):343-6.
41. Pearce CB, Collett J, Goggin PM, Duncan HD. Enteral nutrition by nasojejunal tube in hyperemesis gravidarum. *Clin Nutr*. 2001;20(5):461-4.
42. Stokke G, Gjelsvik BL, Flaatten KT, Birkeland E, Flaatten H, Trovik J. Hyperemesis gravidarum, nutritional treatment by nasogastric tube feeding: a 10-year retrospective cohort study. *Acta Obstet Gynecol Scand*. 2015;94(4):359-67.
43. National Institute for Health and Clinical Excellence. Nutrition support in adults: NICE guideline CG32. London (UK): NICE Clinical Guidelines; 2006.
44. Mazzotta P, Stewart D, Atanackovic G, Koren G, Magee LA. Psychosocial morbidity among women with nausea and vomiting of pregnancy: prevalence and association with anti-emetic therapy. *J Psychosom Obstet Gynaecol*. 2000;21(3):129-36.
45. McCarthy FP, Khashan AS, North RA, Moss-Morris R, Baker PN, Dekker G, et al. A prospective cohort study investigating associations between hyperemesis gravidarum and cognitive, behavioural and emotional well-being in pregnancy. *PLoS One*. 2011;6(11):e27678.
46. Christodoulou-Smith J, Gold JI, Romero R, Goodwin TM, MacGibbon KW, Mullin PM, et al. Posttraumatic stress symptoms following pregnancy complicated by hyperemesis gravidarum. *J Matern Fetal Neonatal Med*. 2011;24(11):1307-11.
47. Veenendaal MVE, van Abeelen AFM, Painter RC, van der Post JAM, Roseboom TJ. Consequences of hyperemesis gravidarum for offspring: a systematic review and meta-analysis. *BJOG*. 2011;118(11):1302-13.
48. Vikanes AV, Stoer NC, Magnus P, Grjibovski AM. Hyperemesis gravidarum and pregnancy outcomes in the Norwegian Mother and Child Cohort - a cohort study. *BMC Pregnancy Childbirth*. 2013;13:169.
49. Grote N, Gavin A, Melville J, Iyengar S, Katon W, Bridge J. A meta-analysis of depression during pregnancy and the risk of preterm birth, low birth weight, and intrauterine growth restriction. *Arch Gen Psychiatry*. 2010;67(10):1012-24.
50. Franceschi F, Di Simone N, D'Ippolito S, Castellani R, Di Nicuolo F, Gasbarrini G, et al. Antibodies anti-CagA cross-react with trophoblast cells: a risk factor for pre-eclampsia? *Helicobacter*. 2012;17(6):426-34.
51. Henderson B, Jing J, Yu M, Pike M, Benton B. Risk factors for cancer of the testis in young men. *Int J Cancer*. 1979;23(5):598-602.
52. Vandraas KF, Vikanes AV, Stoer NC, Troisi R, Stephansson O, Sorensen HT, et al. Hyperemesis gravidarum and risk of cancer in offspring, a Scandinavian registry-based nested case-control study. *BMC Cancer*. 2015;15:398.
53. Roberts CJ. Developmental and neurological sequelae of the common complications of pregnancy and birth. *British Journal of Preventiv & Social Medicine*. 1970;24(1):33-8.

54. Fejzo MS, Magtira A, Schoenberg FP, Macgibbon K, Mullin PM. Neurodevelopmental delay in children exposed in utero to hyperemesis gravidarum. *Eur J Obstet Gynecol Reprod Biol.* 2015;189:79-84.
55. Mullin PM, Bray A, Schoenberg F, MacGibbon KW, Romero R, Goodwin TM, et al. Prenatal exposure to hyperemesis gravidarum linked to increased risk of psychological and behavioral disorders in adulthood. *J Dev Orig Health Dis.* 2011;2(4):200-4.
56. Ayyavoo A, Derraik JG, Hofman PL, Biggs J, Bloomfield FH, Cormack BE, et al. Severe hyperemesis gravidarum is associated with reduced insulin sensitivity in the offspring in childhood. *The Journal of clinical endocrinology and metabolism.* 2013;98(8):3263-8.
57. Grooten IJ, Painter RC, Pontesilli M, van der Post JAM, Mol BWJ, van Eijsden M, et al. Weight loss in pregnancy and cardiometabolic profile in childhood: findings from a longitudinal birth cohort. *BJOG: An International Journal of Obstetrics & Gynaecology.* 2014:n/a-n/a.
58. Abas MN, Tan PC, Azmi N, Omar SZ. Ondansetron compared with metoclopramide for hyperemesis gravidarum: a randomized controlled trial. *Obstet Gynecol.* 2014;123(6):1272-9.
59. Kashifard M, Basirat Z, Kashifard M, Golsorkhtabar-Amiri M, Moghaddamnia A. Ondansetron or metoclopramide? Which is more effective in severe nausea and vomiting of pregnancy? A randomized trial double-blind study. *Clin Exp Obstet Gynecol.* 2013;40(1):127-30.
60. Tan PC, Khine PP, Vallikkannu N, Omar SZ. Promethazine compared with metoclopramide for hyperemesis gravidarum: a randomized controlled trial. *Obstet Gynecol.* 2010;115(5):975-81.
61. Bondok RS, El Sharnouby NM, Eid HE, Abd Elmaksoud AM. Pulsed steroid therapy is an effective treatment for intractable hyperemesis gravidarum. *Crit Care Med.* 2006;34(11):2781-3.
62. Neri I, Allais G, Schiapparelli P, Blasi I, Benedetto C, Facchinetti F. Acupuncture versus pharmacological approach to reduce Hyperemesis gravidarum discomfort. *Minerva Ginecol.* 2005;57(4):471-5.
63. Sullivan CA, Johnson CA, Roach H, Martin RW, Stewart DK, Morrison JC. A pilot study of intravenous ondansetron for hyperemesis gravidarum. *Am J Obstet Gynecol.* 1996;174(5):1565-8.
64. Safari HR, Fassett MJ, Souter IC, Alsulyman OM, Goodwin TM. The efficacy of methylprednisolone in the treatment of hyperemesis gravidarum: a randomized, double-blind, controlled study. *Am J Obstet Gynecol.* 1998;179(4):921-4.
65. Ziaei S, Hosseiney FS, Faghizadeh S. The efficacy low dose of prednisolone in the treatment of hyperemesis gravidarum. *Acta Obstet Gynecol Scand.* 2004;83(3):272-5.
66. Munch S, Schmitz MF. Hyperemesis gravidarum and patient satisfaction: a path model of patients' perceptions of the patient-physician relationship. *J Psychosom Obstet Gynaecol.* 2006;27(1):49-57.
67. Ferguson ND, Davis AM, Slutsky AS, Stewart TE. Development of a clinical definition for acute respiratory distress syndrome using the Delphi technique. *J Crit Care.* 2005;20(2):147-54.
68. Fearon K, Strasser F, Anker SD, Bosaeus I, Bruera E, Fainsinger RL, et al. Definition and classification of cancer cachexia: an international consensus. *Lancet Oncol.* 2011;12(5):489-95.