

# HYPEREMESIS GRAVIDARUM (HG) AND THIAMIN



amplify, educate, advocate

## WHAT IS THIAMIN?

Thiamin (or thiamine) is a water-soluble B vitamin, also known as vitamin B1. Thiamin has many contributing roles including being an antioxidant, maintaining the myelin sheath, ATP production, branched-chain amino acid production, carbohydrate and lipid metabolism, and glucose-derived neurotransmitter synthesis (Pacei, 2020; Sechi, 2007; Thompson, 2002).

**Sources:** Thiamine is naturally present in some foods but also available from supplements. The body cannot make significant amounts on its own. Other B vitamins and electrolytes are important in proper absorption of B1 which occurs mainly in the upper jejunum, then it's transported for liver storage. The body has four forms of thiamin: free thiamine and three phosphorylated forms, including mono-(ThMP), di-(ThDP) and triphosphate (ThTP), with 80% of our thiamin being ThDP, which is metabolically the active form (Pacei, 2020).

**Usage:** Multiple enzyme complexes require the essential cofactor, ThDP, also known as thiamine pyrophosphate (TPP). Thiamin is delivered by erythrocytes to high use organs including the brain, heart, muscles, liver and pancreas. Without thiamin, rapid reductions in ATP will occur with oxidative stress and eventually cell death (Pacei, 2020).

## HG AND THIAMIN

Hyperemesis Gravidarum (HG) is defined as a potentially life-threatening pregnancy disease that may cause weight loss, malnutrition, dehydration, and debility due to severe nausea and/or vomiting and may cause long-

term health issues for mother and baby(ies).

- » **Storage:** The body's 25-30 mg of thiamine storage is depleted after about 2-3 weeks of restricted intake regardless of BMI, and more is needed during pregnancy (Chiossi, 2006; Manto, 2014; Veeprapaneni, 2014). Just two weeks of significant nausea and/or vomiting or weight loss puts patients at high risk of thiamin deficiency (TD).
- » **Intake:** The recommended thiamin intake of 1.4-1.5 mg during pregnancy is inadequate for pregnancies with multiple gestations or HG (MacGibbon, 2015).
- » **Absorption:** Studies report absorption by normal patients with oral dosing of thiamin at 300 mg/day will actually be reduced to 1/3rd or less of the dose (Thompson, 2002), suggesting those with HG and thus impaired absorption will need substantially higher oral doses. Further, vomiting and malnutrition may reduce absorption of thiamin by up to 70% (Thompson, 2002).
- » **Requirements:** HG patients also require high-dose B1 due to their high carbohydrate diet, coexisting deficiencies, limited food variety, impaired absorption, and reduced muscle mass for storage (Bellad, 2015; Ortega, 2004; Tanasescu, 2012; Xiong, 2015).
- » The half-life of thiamin is short, just 1-12 hours (Pacei, 2020), necessitating more frequent administration, especially in those with symptomatic thiamin deficiency.
- » **Deficiency & HG:** TD may make nausea and vomiting refractory to treatment.
- » **Complications:** Severe TD may trigger Wernicke's encephalopathy, while less severe or prolonged deficiency may lead to tachycardia, mental status changes, weight loss, peripheral neuropathy, and fetal loss or neurological damage in the child (Lonsdale, 2017; Sechi, 2007).



### Oral Thiamin Derivatives:

(Lonsdale, 2017 p203-204)

Thiamin hydrochloride is readily available but not as absorbable as derivatives such as TTFD which provide a higher blood concentration of B1.

1. Thiamine tetrahydrofurfuryl disulfide (TTFD), a synthetic derivative, is preferred because it penetrates into cells without a thiamine transporter and crosses the blood-brain barrier. (Fursultiamine, Lipothiamine)
2. S-Benzoylthiamine monophosphate (Benfotiamine) may cross the blood-brain barrier.

- » Women with HG report to the HER Foundation that they have varying degrees of memory and mild cognitive impairment after HG and symptoms of TD, however, they were not diagnosed with or treated for TD or Wernicke's encephalopathy (WE). This suggests chronic TD during HG may pose risks of long-term neurological deficits even if WE does not fully develop.
- » TD increases the risk of fetal loss, Sudden Infant Death Syndrome, and neurological disorders in children of mothers with HG (Lonsdale, 2017).

## THIAMIN DEFICIENCY (TD) SYMPTOMS

Thiamin deficiency (TD) can manifest after just 1 week as tachycardia at rest, peripheral neuropathy, weakness, and decreased reflexes (Pancei, 2020). Continued deficiency develops into dry beriberi (neurological symptoms) or wet beriberi (cardiac symptoms), Wernicke's encephalopathy, or Central Pontine Myelinolysis (Manzo, 2014) that require immediate and aggressive intervention to prevent serious morbidity and mortality.



Early signs of TD mimic and exacerbate HG symptoms.

Distinguishing symptoms from HG and the side-effects of some antiemetics that cause sleepiness, movement disorders, depression, dizziness and instability is problematic.

Assume all patients with HG have TD and prescribe IV and/or oral thiamin. Symptomatic patients require close assessment and IV infusion of thiamin.

- » Weight Loss
- » Confusion
- » Memory loss
- » Muscle weakness
- » Cardiovascular symptoms (tachycardia, bradycardia, enlarged heart)
- » GI Symptoms (vomiting, diarrhea, colitis)
- » Peripheral neuropathy
- » Fatigue
- » Lack of appetite
- » Vision, speaking, or thinking changes
- » Irritability
- » Pain in abdomen or head
- » Mental changes, such as apathy or depression

## MATERNAL COMPLICATIONS

(Said, 2010)

- » Peripheral nerve damage
- » Degeneration of thalamus and cerebellum
- » Reduction of blood flow
- » Vascular resistance
- » Cardiac/respiratory failure
- » Wernicke-Korsakoff Syndrome

## CHILD COMPLICATIONS

85.2% of babies born to TD mothers are also deficient. (Xiong, 2015) If maternal TD while breastfeeding, infants develop TD within 3-4 weeks and have greater incidence of (Butterworth 2001; Lonsdale, 2017; Ortega, 2004):

- » SIDS
- » Behavioral changes
- » Autism
- » Delayed language development, and
- » Decreased visual alertness.

## RECOMMENDATIONS/ TREATMENT

- » **Administer** B1 proactively and routinely during pregnancy, including all HG patients, to prevent potentially life-threatening morbidity and mortality.
- » Slower IV administration results in higher absorption and less renal elimination (Drewe, 2003).
- » **Dilute** B1 in 100 cc of IV fluid and infuse over 30+ minutes.
- » **Monitor electrolytes** (esp. Mg, Phos, Na) and methodically correct along with B vitamin cofactors (B2, B3, B6, B9).
- » **Screen** at each visit for signs of TD and Wernicke's encephalopathy.

### Safety

High dose thiamin has not been associated with any adverse effects and anaphylaxis from IV thiamin is extremely rare (Juel, 2013).

### TD Prevention

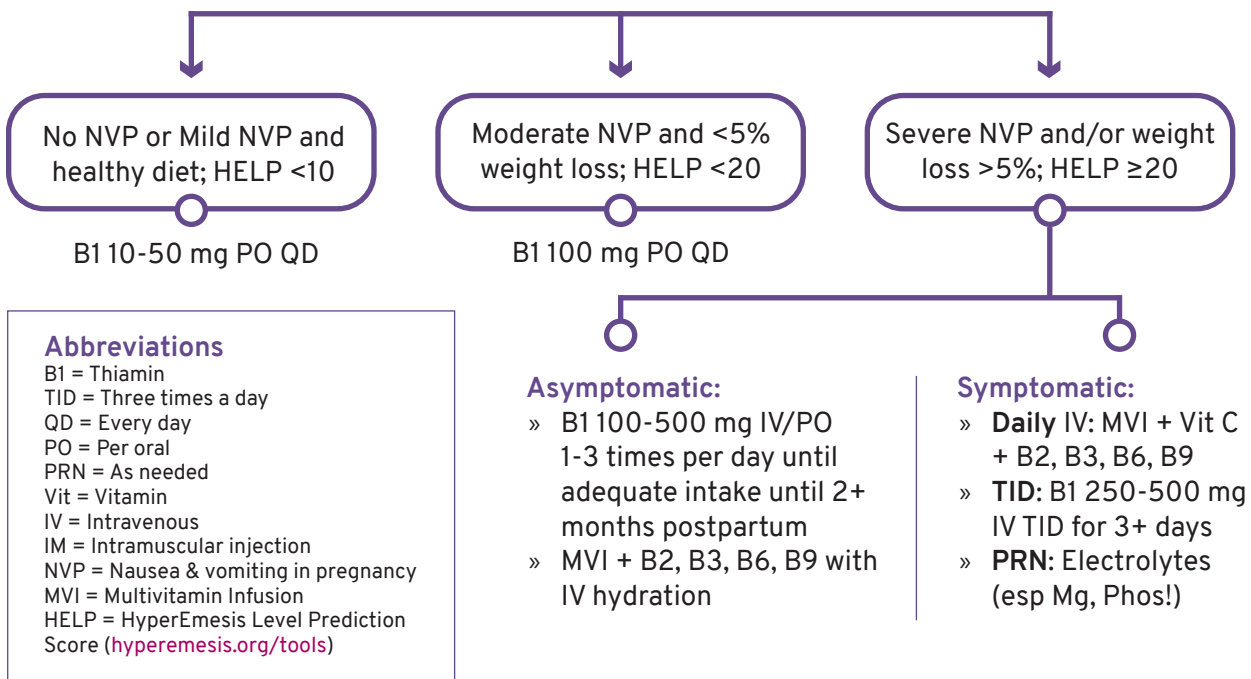
- » Give B1 100-500 mg IV in every IV bag.
- » Proactively administer 200 mg of IV thiamin with every bag of IV dextrose.
- » Give B1 100-500 mg sublingually, orally or IV 1-3 times per day until NVP resolves.
- » Consult dietary to encourage high B1 foods.
- » Monitor diet for high carbohydrate intake and B1 rich foods using the HG Care app food diary ([hyperemesis.org/HGapp](http://hyperemesis.org/HGapp)).

### TD Treatment

- » B1 250-500 mg IV 3 times daily for 3-5 days, or until symptoms resolve. Continue with B1 100-500 mg TID orally until 2+ months postpartum & eating a healthy diet.
- » Continue B1 2 months after breastfeeding.

**For more info:** See the HER website ([hyperemesis.org/neuro](http://hyperemesis.org/neuro)) for the Wernicke's Encephalopathy Fact Sheet, algorithms and protocols ([hyperemesis.org/tools](http://hyperemesis.org/tools)).

## THIAMIN (B1) DEFICIENCY PREVENTION & TREATMENT



## MEDICATIONS THAT IMPACT THIAMIN LEVELS

- » Antacids
- » Anticonvulsants
- » Diuretics
- » Hormonal therapy
- » Parenteral nutrition
- » Antibiotics
- » IV dextrose

## FOOD SOURCES WITH THIAMIN

Heat and processing reduce thiamin levels in foods.

- » Pork
- » Poultry
- » Eggs
- » Fish (trout)

- » Legumes (navy beans, lentils)
- » Nuts (macadamia)
- » Whole grains & cereals
- » Seeds

## CLINICAL KEYS

- » MVI has only 6 mg B1.
- » IV dextrose can cause life-threatening complications in thiamin deficient patients.
- » Always give 200 mg B1 IV with dextrose and 100 mg TID IV/PO with a high carbohydrate diet.
- » Methodically correct electrolytes.
- » B1 Lab testing is unreliable.
- » Avoid IM B1 (atrophy, pain, hematoma).
- » High doses of B1 TID increase absorption.
- » Oral absorption decreased by vomiting and malnutrition (up to 70%) (Thomson, 2002).

## SOURCES

1. Bellad, A., Shrinivas, B., Arif, M., Suhas., 2015. A Rare Case of Wernicke's encephalopathy due to hyperemesis gravidarum. Online J Health Allied Scs, 14(1), p.3.
2. Butterworth, R., 2001. Maternal thiamin deficiency: still a problem in some world communities. Am J Clin Nutr, 74(6), pp.712-713. 72.
3. Chiossi, G., Neri, I., Cavazzuti, M., Basso, G., Facchinetti, F., 2006. Hyperemesis gravidarum complicated by Wernicke encephalopathy: background, case report, and review of the literature. Obstet Gynecol Surv, 61(4), pp.255- 268.
4. Drewe, J., Delco, F., Kissel, T. and Beglinger, C., 2003. Effect of intravenous infusions of thiamine on the disposition kinetics of thiamine and its pyrophosphate. Journal of Clinical Pharmacy and Therapeutics, 28(1), pp.47-51.
5. HER Foundation. (n.d.) Common Medications. Retrieved from <https://www.hyperemesis.org/about-hyperemesis-gravidarum/treatment/medications>.
6. Juel, J., Pareek, M., Langfrits, C. and Jensen, S., 2013. Anaphylactic shock and cardiac arrest caused by thiamine infusion. Case Reports, 2013(jul12 1), pp.bcr2013009648-bcr2013009648.
7. Lonsdale, D. and Marrs, C., 2017. Thiamine Deficiency Disease, Dysautonomia, And High Calorie Malnutrition. San Diego: Academic Press. pp 172.
8. MacGibbon KW, Fejzo MS, Mullin PM, 2015. Mortality Secondary to Hyperemesis Gravidarum: A Case Report. Womens Health Gynecol 1(2).
9. Manzo, G., De Gennaro, A., Cozzolino, A., Serino, A., Fenza, G., Manto, A., 2014. MR Imaging Findings in Alcoholic and Nonalcoholic Acute Wernicke's Encephalopathy: A Review. BioMed Res Int, 2014, pp.1-12.
10. Ortega, R., Martínez, R., Andrés, P., Marín-Arias, L., López-Sobaler, A., 2004. Thiamin status during the third trimester of pregnancy and its influence on thiamin concentrations in transition and mature breast milk. BJN, 92(01), p.129.
11. Oudman, E., Wijnia, J., Oey, M., van Dam, M., Painter, R. and Postma, A., 2019. Wernicke's encephalopathy in hyperemesis gravidarum: A systematic review. European Journal of Obstetrics & Gynecology and Reproductive Biology, 236, pp.84-93.
12. Pacei, F., Tesone, A., Laudi, N., Laudi, E., Cretti, A., Pnini, S., Varesco, F. and Colombo, C., 2020. The Relevance of Thiamine Evaluation in a Practical Setting. Nutrients, 12(9), p.2810.
13. Said, H., 2010. Thiamin. In: Coates, P., Betz, J., Blackman, M., et al., eds. Encyclopedia of Dietary Supplements. 2nd ed. London and New York: Informa Healthcare, pp.748-53.
14. Sechi, G., Serra, A., 2007. Wernicke's encephalopathy: new clinical settings and recent advances in diagnosis and management. The Lancet Neurology, 6(5), pp.442-455.
15. Tanasescu, R., Dumitrescu, L., Dragos, C., Luca, D., Oprisan, A., Coclitu, C., et al. 2012. Wernicke's Encephalopathy, Miscellanea on Encephalopathies - A Second Look, Dr. Radu Tanasescu (Ed.), ISBN: 978-953-51-0558-9, InTech, DOI: 10.5772/27988.
16. Thomson, A., 2002. The Royal College of Physicians report on alcohol: guidelines for managing Wernicke's encephalopathy in the accident and emergency department. Alcohol and Alcoholism, 37(6), pp.513-521.
17. Veerapaneni, K., Brown, T., Dooley, D., 2014. Combined deficiencies of vitamins B1 and C in well-nourished patients. Primary Care Companion CNS Disord, 16(4).
18. Xiong, G., & Bienenfeld, D., 2015. Wernicke-Korsakoff Syndrome Treatment & Management: Approach Considerations, Diet and Activity, Referral and Follow-Up Care. Emedicine.medscape.com.