## APPENDIX V: Treatment algorithms for NVP and HG in primary care (Vai and ii), ambulatory care (Vb), emergency department (Vc) and inpatient care (Vd)

## Vai. Summary for General Practitioners

## Why is the active management of nausea and vomiting of pregnancy (NVP)/ hyperemesis gravidarum (HG) important?

- NVP/ HG is associated with serious health consequences for both mother and baby
- Patients with NVP/HG often present to primary care as onset of symptoms occur prior to their pregnancy being booked by a midwife
- Patients are likely to have tried non-pharmacological options prior to presenting thus they
  may have severe disease at first presentation to primary care

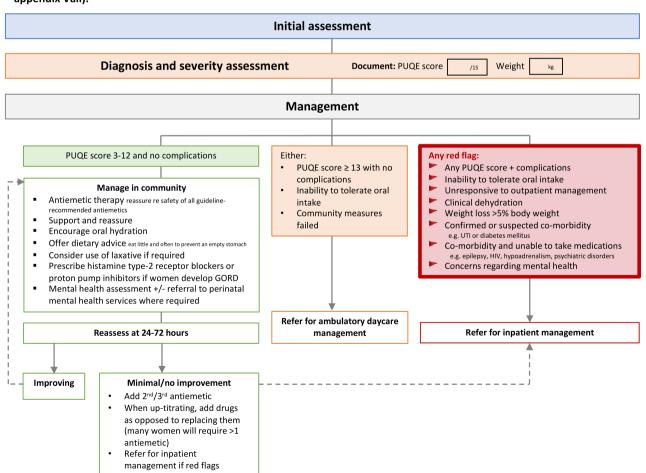




## Practice points for general practitioners:

- · Validate patients' symptoms
- There are safety and efficacy data for first line antiemetic therapy including anti (H1) histamines, phenothiazines and doxylamine/pyridoxine and they should be prescribed when required for the management of NVP/HG
- In patients with severe disease multiple antiemetics prescribed together will be required
- · Ketonuria is not an indicator of dehydration and should not be used to assess severity of NVP/HG
- Guidance for referral to secondary care is included in the algorithm below
- NVP/HG is likely to recur in subsequent pregnancies and pre-emptive use of medication can reduce severity of disease future pregnancies
- An assessment of mental as well as physical is important

Recommended simplified management algorithm for management of NVP/HG in primary care (for detailed algorithm see appendix Vaii):



	Vaii. Management of Nausea and Vomiting of Pregr	nancy (NVP)/ Hyperer	nesis Gravidarum (HG) in General Practico	
	History:  Previous history of NVP/HG	e	Royal College of Obstetricians & GPCPC  Investigations:  Urine dipstick +/- MSU  intrites may indicate urinary tract infection  188. Ketones are not a marker of dehydration  Urea and electrolytes  to assess for hypo/hyperkalaemia, hyponatraemia, kidney injury  Full blood count  infection, raised the or Mct may indicate dehydration  Blood glucose to assess for diabetes	
	NVP:  onset of nausea and/or vomiting in early pregnancy with no other cause is identified  HG:  Nausea and vomiting (one of which is severe)  Onset <16 weeks' gestation  Inability to eat and drink normally	Document: PUQE s  E-24 scoring system: e last 24 hours: ong have you felt nauseated or your stomach for? nany times have you vomited? nany times have you had ng or dry heaves?	Not at ≤1h 2-3hrs 4-6hrs >6hrs all [1] r[2] [3] [4] [5]  0x [1] 1-2x 3-4x 5-6x ≥7x [2] [3] [4] [5]  0x [1] 1-2x 3-4x 5-6x ≥7x [2] [3] [4] [5]	
	PUQE score 3-12 and no complications  Manage in community  Antiemetic therapy reasure re safety of all guideline- recommended antiemetics  Support and reassure  Encourage oral hydration  Offer dietary advice eat little and often to prevent an empty stomach  Mental health assessment +/- referral to perinatal mental health services where required	Either:  PUQE score ≥ 13 with no complications Inability to tolerate oral intake Community measures failed	Any red flag:  Any PUQE score + complications Inability to tolerate oral intake Unresponsive to outpatient management Clinical dehydration Weight loss >5% body weight Confirmed or suspected co-morbidity e.g. U1 or diabetes mellitus Co-morbidity and unable to take medications e.g. epilepsy, diabetes mellitus, HIV, hypoadrenalism and psychiatric disorders Concerns regarding mental health	
  -  -	Reassess at 24-72 hours    Minimal/no improvement : Add 2 <sup>/3-4</sup> antismetic Refer for impatient management if red flags	Refer for ambulatory daycare management	Refer for inpatient management	
Prochlorperazine 5–10 mg 6–8 hourly PO (or 3 mg buccal); 12.5 mg 8 hourly IM/IV; 25 mg PR daily Promethazine 12.5–25 mg 4–8 hourly PO, IM or IV Chlorpromazine 10–25 mg 4–6 hourly PO, IM or IV  2nd line Metoclopramide 5–10 mg 8 hourly PO, IV/IM/SC Democridate 10 mg 8 hourly PO, IV/IM/SC Democridate 10 mg 8 hourly PO; 30 mg 13 hourly PP  For all patients consider:		Up titration of antiemetics: Initially select a 1st line antiemetic Use combinations of drugs in women who do not respond to a single antiemetic When up titrating add drugs as opposed to replacing them Different classes of drugs may have synergistic effects and some women will require a combination of 3+ antiemetics to control symptoms  For all patients consider:		
	Ondersetron 4 mg 8 hourly PO; 30 mg 12 hourly PR Ondansetron 4 mg 8 hourly or 8 mg 12 hourly PO; 8 mg over 16 mg daily PR Women taking ondansetron may require laxatives if constipation develops  3rd line Prednisolone 40–50 mg daily PO, with the dose gradua maintenance dose that controls the symptoms is reached Controcteriodis should be reserved for cases where standard therapies have falled; when initials addition to previously started antiemetics. Women taking them should have their BP monitored	ally tapered until lowest	Histamine type-2 receptor blockers or proton pump inhibitors if women develop GORD Both safe in pregnancy     Thiamine supplementation in those with severely reduced dietary intake     Laxatives if required for constipation     VTE risk assessment (see RCOG risk assessment tool)	
	Post-partum care, planning for future pregnancy and signposting  Patients with severe HG are risk of PTSD if required they should be referred to appropriate services In future pregnancy early use of lifestyle modifications should be used  HER Foundation			

- Patients with severe HG are risk of PTSD if required they should be referred to appropriate services
- In future pregnancy early use of lifestyle modifications should be used
- Pre-emptive use of doxylamine and pyridoxine can be used to reduce severity of disease in subsequent pregnancy 20/20mg PO at night to be started on confirmation of positive pregnancy test and up titrated when required



(1)

- Pregnancy Sickness Support
- HER Foundation
- UK Teratology Information Service

