



Addressograph

Homocystinuria New Diagnosis Care Plan

PROBLEM: Knowledge deficit due to new diagnosis.	S/N Sig:	Date: _____ Planned by: _____	Problem no: 45
GOAL: 1) To establish recommended dietary regime for management of Homocystinuria (HCU) 2) To educate parent (s) on HCU, management, complications of non-treatment / non-compliance with recommended management			
NURSING CARE	SELF/ FAMILY CARE	DATE AND SIGN / COUNTERSIGN ANY CHANGES	
<ol style="list-style-type: none"> 1. Record vital signs (Temperature, Pulse and Respiratory rate) daily or more frequently as condition indicates. 2. _____ will continue on existing feeding regime until baseline blood specimens have been obtained and pyridoxine responsiveness has been established. 3. Assist in obtaining baseline bloods for: Quantitative Amino Acids <input type="checkbox"/>, Total and Free Homocysteine <input type="checkbox"/>, DNA mutation <input type="checkbox"/>, B₁₂ & Folate <input type="checkbox"/>, CBS Enzyme (EDTA) <input type="checkbox"/>. 4. Liaise with Metabolic Consultant regarding trial of Pyridoxine (to confirm / out rule Pyridoxine responsiveness). 5. If pyridoxine non-responsive, establish dietary and medication management as prescribed by Metabolic Consultant. 6. Take blood for Total & Free Homocysteine sample (deproteinised amino acids) daily / alternate days / or other _____ (insert frequency) as instructed by Metabolic Consultant. <i>Ensure rapid delivery of amino acids specimens to Metabolic Laboratory (require deproteinising within 10 mins of collection).</i> 7. Liaise with Metabolic Team and Dietetic Team re. commencement of special diet. 8. Administer Vit B₁₂ and Folate and B6 as prescribed. 9. Commence parental education programme re. condition, diet, implications of non-compliance and non-treatment in liaison with Metabolic Nurse Specialist. Give HCU Handbook to parents to provide reference material and learning resource. Encourage to read and ask questions. 10. Assess parent's level of understanding of the condition/management throughout _____'s admission and prior to discharge. 11. Monitor fluid and calorie intake. 12. Weigh _____ (insert frequency) 13. Involve Psychology and Social Work teams in ongoing care and establishing support networks for discharge. 14. Liaise with Metabolic team re. scheduling of consults to other disciplines and screening of siblings. Cardiology date _____, Ophthalmology date _____, Screening of Sibling(s) date _____. 15. Liaise with PHN and provide him/her with information re HCU. 16. Establish, if possible, the facility to take bloods at local hospital. <p>Reference: <u>Management Protocol for Patients diagnosed with homocystinuria (HCU) due to Cystathionine β-syntase deficiency.</u></p>			