



Addressograph

**Homocystinuria New Diagnosis Care Plan**

<b>PROBLEM:</b> Knowledge deficit due to new diagnosis.	S/N Sig:	Date: _____ Planned by: _____	Problem no: <b>45</b>
<b>GOAL:</b> 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			
<b>NURSING CARE</b>	<b>SELF/ FAMILY CARE</b>	<b>DATE AND SIGN / COUNTERSIGN ANY CHANGES</b>	
<ol style="list-style-type: none"> <li>1. Record vital signs (Temperature, Pulse and Respiratory rate) daily or more frequently as condition indicates.</li> <li>2. _____ will continue on existing feeding regime until baseline blood specimens have been obtained and pyridoxine responsiveness has been established.</li> <li>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"/>, B<sub>12</sub> &amp; Folate <input type="checkbox"/>, CBS Enzyme (EDTA) <input type="checkbox"/>.</li> <li>4. Liaise with Metabolic Consultant regarding trial of Pyridoxine (to confirm / out rule Pyridoxine responsiveness).</li> <li>5. If pyridoxine non-responsive, establish dietary and medication management as prescribed by Metabolic Consultant.</li> <li>6. Take blood for Total &amp; 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></li> <li>7. Liaise with Metabolic Team and Dietetic Team re. commencement of special diet.</li> <li>8. Administer Vit B<sub>12</sub> and Folate and B6 as prescribed.</li> <li>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.</li> <li>10. Assess parent's level of understanding of the condition/management throughout _____'s admission and prior to discharge.</li> <li>11. Monitor fluid and calorie intake.</li> <li>12. Weigh _____ (insert frequency)</li> <li>13. Involve Psychology and Social Work teams in ongoing care and establishing support networks for discharge.</li> <li>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 _____.</li> <li>15. Liaise with PHN and provide him/her with information re HCU.</li> <li>16. Establish, if possible, the facility to take bloods at local hospital.</li> </ol> <p>Reference: <u>Management Protocol for Patients diagnosed with homocystinuria (HCU) due to Cystathionine β-syntase deficiency.</u></p>			