Children's Wisconsin

Co-Management Guidelines

To support collaborative care, we have developed guidelines for our community providers to utilize when referring to, and managing patients with, the pediatric specialists at Children's Wisconsin. These guidelines provide protocols for jointly managing patient cases between community providers and our pediatric specialists.

Thrombocytopenia A condition in which the patient has a low blood platelet count. Normal platelet count is 150,00 to 400,000 throughout childhood									
Diagnosis/symptom	Referring provider's initial evaluation and management:	When to initiate referral/ consider refer to Hematology Clinic:	What can referring provider send to Hematology Clinic?	Specialist's workup will likely include:	Model of Care				
 Signs and symptoms: Onset, Bleeding site, Severity Acute: petechiae, purpura, epistaxis, menorrhagia Chronic: easy bruising, epistaxis, prolonged and/or frequent menstrual bleeding 	 Initial evaluation and therapy: History and physical exam CBC, differential, Coagulation studies if bleeding (PT, PTT, consider von Willebrand testing) If ITP, some pediatricians may prefer to manage without referral. 	 Sudden onset of moderate- severe thrombocytopenia (<20,000 platelets) Chronic thrombocytopenia of unknown cause Thrombocytopenic patient with uncontrolled bleeding Thrombocytopenic patient to undergo surgical intervention IF SEVERE THROMBOCYTOPENIA AND BLEEDING PLEASE CALL OUR CLINIC 	 Internal Provider using Epic: Place Ambulatory Referral to Hematology. External Provider using EPIC: Please complete the external referral order to CHW HEMATOLOGY & ONCOLOGY CLINICS or or Fax to Hematology at (414) 266-2426 or Online ambulatory referral form Please include: History and physical exam including past medical history Recent laboratory studies especially CBC and differential 	 CBC, differential, and reticulocyte count and smear Consideration of coagulation studies Genetic analysis for chronic thrombocytopenia syndromes If more than one cell line is involved, consideration of bone marrow aspiration biopsy ANA, HIV studies for new onset ITP in an adolescent After referral to Hematology Clinic: For acute ITP: CBC to be followed at local lab or PMD office with phone contacts between primary physician's office and pediatric hematology For chronic thrombocytopenic syndromes: observation with eventual genetic testing to be 	Initial consultation with followup care by primary physician, perhaps with input or additional hematology followup If ITP, initial treatment observation vs steroids, possibly IVIG. Chronic treatment may involve immune suppression, thrombopoietin receptor mimetics, splenectomy. If congenital platelet problem, may need treatment for bleeding with desmopressin, antifibrinolytics, platelet transfusion.				



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			Office Number: (414) 266-2420	performed if thrombocytopenia persists for greater than one year and there is no pre-existing history of a normal platelet count					
Causes									
 ITP, most common cause of acute thrombocytopenia children in otherwise well child Consumptive coagulopathy (DIC, HUS, TTP) in moderate to severely ill children 									
 Chronic thrombocytopenic syndromes, likely genetic and etiology in a well-child with chronic thrombocytopenia 									
References									
ITP: new guidelines available at https://ashpublications.org/bloodadvances/article/3/23/3829/429213/American-Society-of-Hematology-2019-guidelines-for									
General thrombocytopenia: https://www.uptodate.com/contents/approach-to-the-child-with-unexplained-									
thrombocytopenia#:~:text=Thrombocytopenia%2C%20defined%20as%20a%20platelet,or%20bleeding%2C%20or%20mucosal%20hemorrhage.									
Isolated thrombocytopenia: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6056371/									

*Approved by Specialty Medical Leader, CSG Clinical Integration, CMG Clinical Guidelines Core Team

Medical Disclaimer

Medicine is a dynamic science; as research and clinical experience enhance and inform the practice of medicine, changes in treatment protocols and drug therapies are required. The authors have checked with sources believed to be reliable in their effort to provide information that is complete and generally in accord with standards accepted at the time of publication. However, because of the possibility of human error and changes in medical science, neither the authors nor Children's Hospital and Health System, Inc. nor any other party involved in the preparation of this work warrant that the information contained in this work is in every respect accurate or complete, and they are not responsible for any errors in, omissions from, or results obtained from the use of this information. Readers are encouraged to confirm the information contained in this work with other sources.



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Up to date citations 4/10/2024

https://www.uptodate.com/contents/immune-thrombocytopenia-itp-in-children-clinical-features-anddiagnosis?search=pediatric%20itp&source=search_result&selectedTitle=3%7E150&usage_type=default&display_rank=3

https://www.uptodate.com/contents/immune-thrombocytopenia-itp-in-children-initialmanagement?search=pediatric%20itp&source=search_result&selectedTitle=2%7E150&usage_type=default&display_rank=2

General thrombocytopenia

https://www.uptodate.com/contents/approach-to-the-child-with-unexplainedthrombocytopenia?search=pediatric%20thrombocytoepnia&source=search_result&selectedTitle=1%7E150&usage_type=default&display_rank=1



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