



Disseminated Intravascular Large B-Cell Lymphoma Presenting as Chronic Pancreatitis and Ischemic Colitis

Kaila Buckley, Nyla A Heerema and Weiqiang Zhao*

Department of Pathology, The Ohio State University Wexner Medical Center, USA

Clinical Image

A 56-year-old Caucasian male with no past medical history presented to a local hospital with fevers of unknown origin, recurrent pancreatitis (2 episodes were 3 months apart, and 3 months prior to this diagnosis). He quickly deteriorated with anemia, thrombocytopenia, coagulopathy, renal failure, as well as several embolic events, including splenic infarcts and colitis. A bone marrow was performed to rule out immune thrombocytopenia purpura which showed hypercellularity (80%) with increased medullary compartments and sinusoidal closure with large atypical cells but difficult to assess based on H&E stain (Figure 1A). Immunohistochemistry (IHC) for CD20 on the bone marrow biopsy highlighted monotonous large lymphoid cells located intravascular and intra-sinusoidal spaces (Figure 1B). The patient expired after his code status was switched to “Do Not Resuscitate”. The post-mortem examination showed ischemic colitis with mucosal and focal transmural necrosis and spotted microthrombi by monotonous intravascular large lymphoma cells (Figure 1C) which were uniformly positive for CD20 (Figure 1D).

Intravascular lymphoma is a systemic disorder that usually affects the elderly in their sixth decade [1]. Although it is an extremely rare lymphoma, IVLBCL should be included in the differential diagnosis when recurrent pancreatitis and nonspecific abdominal manifestations are the major complaints due to ischemic events caused by microthrombi and lymphoma aggregates [2,3]. A high-dose chemotherapy followed by ASCT can sometimes achieve remission. Therefore, a prompt bone marrow biopsy or tissue biopsy should be done. The morphological examination sometimes might be difficult but atypical large lymphoid cells, increased bone marrow cellularity, and positivity for B cell markers by Immunohistochemistry should not be missed. There is no defined characteristic karyo type and molecular insights are still underexplored and might provide new diagnostic tool as well as therapeutic opportunities for this deadly disease. To our knowledge this is the first case published in IVLBCL presenting with recurrent pancreatitis [1-3].

OPEN ACCESS

*Correspondence:

Weiqiang Zhao, Department of Pathology, The Ohio State University Wexner Medical Center, USA, E-mail: Weiqiang.Zhao@osumc.edu

Received Date: 10 Aug 2019

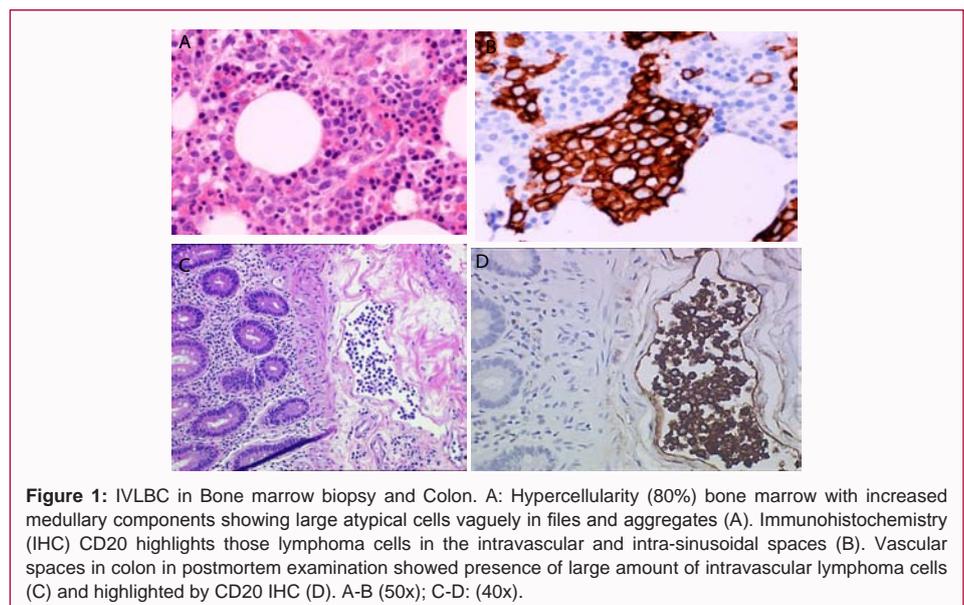
Accepted Date: 28 Aug 2019

Published Date: 04 Sep 2019

Citation:

Buckley K, Heerema NA, Zhao W. Disseminated Intravascular Large B-Cell Lymphoma Presenting as Chronic Pancreatitis and Ischemic Colitis. *Am J Leuk Res.* 2019; 3(1): 1018.

Copyright © 2019 Weiqiang Zhao. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.



References

1. Swerdlow SH, Campo E, Pileri SA, Harris NL, Stein H, Siebert R, et al. The 2016 revision of the World

- Health Organization classification of lymphoid neoplasms. *Blood*. 2016;127(20):2375-90.
2. Testoni PA. Acute recurrent pancreatitis: Etiopathogenesis, diagnosis and treatment. *World J Gastroenterol*. 2014;20(45):16891-01.
 3. Opie EL. The etiology of acute hemorrhagic pancreatitis. *Bull Johns Hopkins Hosp*. 1901;12:182-8.