Turk J Hematol 2020;37:207-219 LETTERS TO THE EDITOR

aspirate cytology. In pregnancy, management is performed with prednisolone, although immunosuppressive therapy has also been used [2]. In Table 1, the different modes of treatment used in cases of pregnancy-related PRCA by different authors are summarized [3,4,5,6,7]. PRCA in pregnancy has a better prognosis compared to pre-existing PRCA and aplastic anemia in pregnancy. In general, PRCA that develops during pregnancy spontaneously resolves postpartum [8]. Although it rarely recurs in subsequent pregnancy, recurrence was observed here and hence the permanence of PRCA is not known [3].

in pregnancy, Hypoproliferation, Erythroid Promegaloblast

Anahtar Sözcükler: PRCA, Edinsel saf kırmızı hücre aplazisi, anemi, Hipoproliferasyon, Eritroid Gebelikte öncüller. Promegaloblast

Informed Consent: Obtained.

Conflict of Interest: No conflict of interest was declared by the authors.

Keywords: PRCA, Acquired pure red cell aplasia, Anemia

received no financial support. Acknowledgements: Salem Polyclinic Pvt. Ltd. References

1. Sawada K, Fujishima N, Hirokawa M. Acquired pure red cell aplasia: updated review of treatment.Br J Haematol 2008;142:505-514.

Financial Disclosure: The authors declared that this study

- Means RT. Pure red cell aplasia. Hematol Am Soc Hematol Educ Program 2016;2016:51-56.
- Miller AC, Rashid RM. Three episodes of acquired pure red cell aplasia restricted to pregnancy. J Perinat Med 2008:36:270-271.
- Kashyap R, Pradhan M. Maternal and fetal outcome in pregnancy-associated pure red cell aplasia. J Obstet Gynaecol 2010;30:733-734.
- Ito S, Ikuta K, Yamamoto M, Okamura N, Ichiki K, Sugiyama J, Shindo M, Torimoto Y, Kohgo Y. Case report: A case of pure red cell aplasia associated with pregnancy. Nihon Naika Gakkai Zasshi 2012;101:2042-2044.
- Aggarwal S. Reversible pure red cell aplasia of pregnancy: a therapeutic challenge. J Obstet Gynaecol India 2013;63:138-139.
- Edahiro Y, Yasuda H, Ando K, Komatsu N. Self-limiting pregnancy-associated pure red cell aplasia developing in two consecutive pregnancies: case report and literature review. Int J Hematol 2020:111:579-584.
- Choudry M, Moffett B, Laber D. Pure red-cell aplasia secondary to pregnancy, characterization of a syndrome. Ann Hematol 2007;86:233-237.

©Copyright 2020 by Turkish Society of Hematology Turkish Journal of Hematology, Published by Galenos Publishing House



Address for Correspondence/Yazışma Adresi: Ashwin Rao, MD, Salem Polyclinic, Department of Obstetrics

and Gynecology, Tamil Nadu, India

Phone: +91-7708922999

E-mail: ashwinrao2404@gmail.com ORCID: orcid.org/0000-0002-9462-5321

Received/Geliş tarihi: April 14, 2020 Accepted/Kabul tarihi: May 11, 2020

DOI: 10.4274/tjh.galenos.2020.2020.0170

# A Case of Burkitt's Lymphoma Mimicking Peritonitis Carcinomatosa

## Peritonitis Karsinomatozayı Taklit Eden Bir Burkitt Lenfoma Olgusu

Deram Büyüktaş¹, Serdar Örnek², Tülay Tecimer³, Burhan Ferhanoğlu¹

<sup>1</sup>Koç University Faculty of Medicine, Department of Hematology, İstanbul, Turkey

<sup>2</sup>American Hospital, Department of Hematology, İstanbul, Turkey

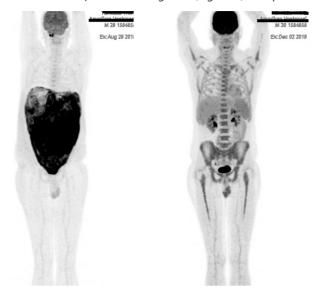
<sup>3</sup>Acıbadem University Faculty of Medicine, Department of Pathology, İstanbul, Turkey

### To the Editor,

A 30-year-old man was admitted to the hospital with fatigue, fever, nausea, and abdominal distension in August 2019. Laboratory analyses were as follows: white blood cell count, 13,400/μL; absolute neutrophil count, 9,700/μL; absolute lymphocyte count, 2,100/μL; hemoglobin, 14.5 g/dL; platelets, 442,000/μL; C-reactive protein, 15.6 mg/L; lactate dehydrogenase, 186 U/L; ferritin, 927 ng/mL; alanine transaminase, 108 U/L; aspartate transaminase, 245 U/L. Abdominal ultrasound showed massive ascites. Cytospinning of the ascites revealed B-cell nonHodgkin's lymphoma. PET-CT showed increased FDG uptake of the whole peritoneum, omentum, and small intestine (Figure 1). Peritonitis carcinomatosa was considered in the differential diagnosis. The patient underwent tru-cut peritoneal biopsy; the findings were consistent with Burkitt's lymphoma. In immunohistochemical analysis, CD20, CD10, bcl6, and c-myc were positive; CD5, bcl2, CD23, MUM1, and TDT were negative. The Ki-67 index was 99%. FISH analysis for myc/IGH translocation was positive. Bone marrow was normocellular with no sign of lymphoma involvement and conventional cytogenetics showed a normal karyotype: 46, XY [20]. Cerebrospinal fluid cytospinning

LETTERS TO THE EDITOR Turk J Hematol 2020;37:207-219

was also negative for any atypical cells. He was treated with the GMALL protocol [1]. Interim PET was consistent with complete response after four cycles of the regimen (Figure 1). The patient



**Figure 1.** PET-CT before and after treatment.

PET-CT: Positron emission tomography-computed tomography

completed the rest of the regimen uneventfully and the final PET-CT did not show any residual disease or recurrence.

**Keywords:** Burkitt's lymphoma, Peritonitis carcinomatosa, PET-CT

**Anahtar Sözcükler:** Burkitt lenfoma, Peritonitis karsinomatoza, PET-BT

Informed Consent: Obtained.

**Conflict of Interest:** No conflict of interest was declared by the authors.

**Financial Disclosure:** The authors declared that this study received no financial support.

### References

 Pohlen M, Gerth HU, Liersch R, Koschmieder S, Mesters RM, Kessler T, Appelmann I, Müller-Tidow C, Berdel WE. Efficacy and toxicity of a rituximab and methotrexate based regimen (GMALL B-ALL/NHL 2002 protocol) in Burkitt's and primary mediastinal large B-cell lymphoma. Am J Hematol 2011;86:61-64.

©Copyright 2020 by Turkish Society of Hematology Turkish Journal of Hematology, Published by Galenos Publishing House



Address for Correspondence/Yazışma Adresi: Deram Büyüktaş, MD, Koç University Faculty of Medicine, Department of Hematology, İstanbul, Turkey

E-mail: derambuyuktas@yahoo.com ORCID: orcid.org/0000-0002-3623-2925

Received/Geliş tarihi: January 11, 2020 Accepted/Kabul tarihi: March 16, 2020

DOI: 10.4274/tjh.galenos.2020.2020.0015

# **CD4+CD8+ Double-Positive T-Lymphocytes: Pitfalls**

CD4+CD8+ Çift Pozitif T-Lenfositler: Tuzaklar

İrfan Yavaşoğlu

Aydın Adnan Menderes University Faculty of Medicine, Division of Hematology, Aydın, Turkey

#### To the Editor.

The article entitled "Percentages of CD4+CD8+ Double-positive T Lymphocytes in the Peripheral Blood of Adults from a Blood Bank in Bogotá, Colombia," written by Gonzalez-Mancera et al. [1] and published in a recent issue of your journal, was quite interesting. Herein, I wish to contribute to the article.

Nicotine has been reported to affect the cell-mediated immune system. In addition, nicotine exposure can lead to regulatory T-cell induction [2,3]. Therefore, I think that it is important to know the smoking status and also the number of

lymphocytes for the subjects in Gonzalez-Mancera et al.'s [1] study. Data have been published revealing that the prevalence of monoclonal B-cell lymphocytosis is higher than previously reported in blood donors [4]. Also, the large number of monoclonal B-cell lymphocytes determines the biological fate of cells transfused in recipients [4]. The use of CD45 during gating in flow cytometry could provide accurate identification. CD3+CD16/56 is important in determining natural killer T (NKT) cells and could have identified NKT cell contamination in the study.