

A new diagnosis of monoclonal B-cell lymphocytosis with cytoplasmic inclusions in a patient with COVID-19

Lorella Lanza¹ | Brisejda Korovesi² | Francesca Barducchi¹ | Angela Lorenzo³ |
Ezio Venturino¹ | Enrico Cappelli² | Flavia Lillo² | Barbara J. Bain⁴ 

¹Anatomical Pathology, ASL2 Liguria, Santa Corona Hospital, Pietra Ligure, Italy

²Laboratory of Clinical Pathology, ASL2 Liguria, S. Paolo Hospital, Savona, Italy

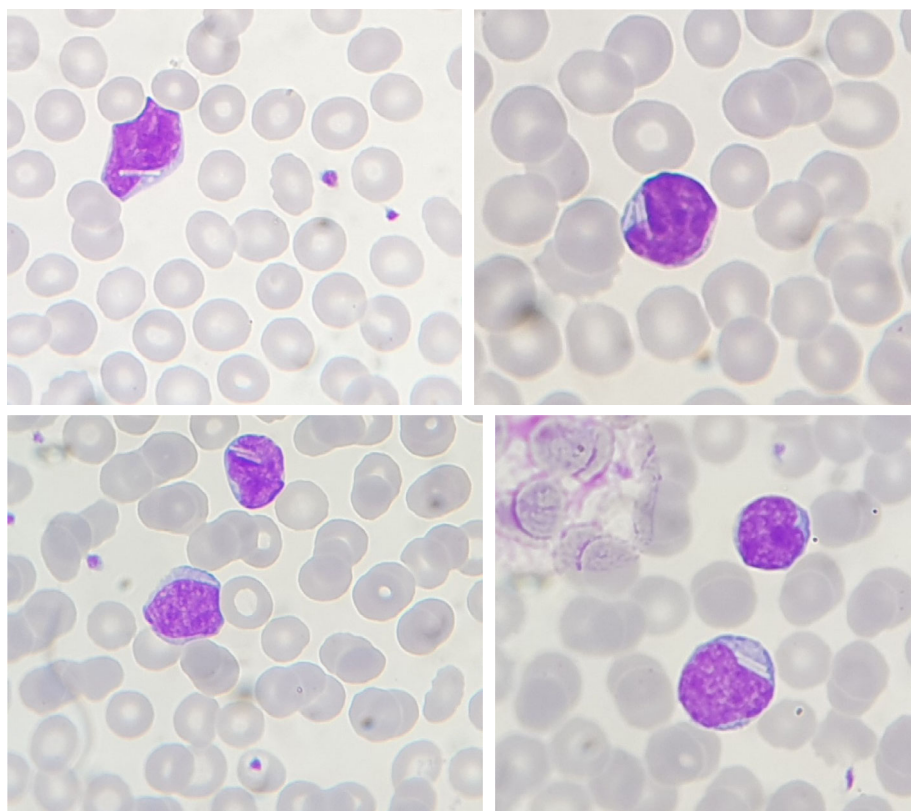
³Laboratory of Clinical Pathology, ASL2 Liguria, S. Corona Hospital, Savona, Italy

⁴Centre for Haematology, St Mary's Hospital Campus of Imperial College Faculty of Medicine, St Mary's Hospital, London, UK

Correspondence

Barbara J. Bain, Blood Sciences, St Mary's Hospital, Praed Street, London, W2 1NY, UK.

Email: b.bain@imperial.ac.uk



A 75-year-old man with a history of chronic ischemic heart disease with a previously normal blood count, presented to the emergency room with fever and tachycardia. There was no hepatosplenomegaly or lymphadenopathy. An electrocardiogram showed a left bundle branch block. Oxygen saturation was 95%. Because of the fever, the patient was tested for SARS-CoV-2; the

reverse transcriptase-polymerase chain reaction was positive. A diagnosis of COVID-19 was made.

The patient's blood count showed a white blood cell count (WBC) of $10.46 \times 10^9/L$, lymphocytes $4.51 \times 10^9/L$, hemoglobin 129 g/L, and platelet count $233 \times 10^9/L$. D-dimer was 659 $\mu\text{g/L}$ (normal range < 500) and interleukin 6 was 76.3 pg/ml (normal range < 6.4).

A computed tomography scan of the chest showed bilateral interstitial infiltrates associated with multiple enlarged mediastinal lymph nodes. Following a rapid and unexpected increase of the WBC to $17.49 \times 10^9/L$ and of the lymphocyte count to $8.37 \times 10^9/L$ over the next 48 h, a blood film and immunophenotyping were performed. The film showed small and medium-sized lymphocytes, with a variable N:C ratio and moderately basophilic cytoplasm. Smear cells were present. A quarter of the lymphocytes showed the negative images of one to three rod-shaped crystals (average two per cell). There were some immature monocytes and some neutrophils showed mild toxic granulation or abnormal nuclear shapes, consistent with COVID-19. Flow cytometric immunophenotyping showed an increased number of circulating B cells (93% of lymphocytes, $7.78 \times 10^9/L$) with lambda light chain restriction and expressing CD19, CD5, CD23, weak CD20, CD43, and CD200; the cells were negative for CD10, CD79b, CD81, FMC7, and CD38. The matutes score was 5/5. At this stage, the condition could not be distinguished from chronic lymphocytic leukemia (CLL).

Two months later the WBC and lymphocyte count had returned to normal and repeated immunophenotyping showed only $0.63 \times 10^9/L$ CD5-positive clonal B cells. However, lymphocytes with cytoplasmic crystals were still present. A diagnosis of monoclonal B-cell lymphocytosis (MBCL) was made. Patients with CLL in whom COVID-19 led to a marked but transient increase in the lymphocyte

count have been reported. In this case, COVID-19 in a patient with MBCL led to an increase in the lymphocyte count to a level simulating CLL with follow-up indicating the correct diagnosis. In addition, we report here the observation of the negative images of crystals, attributable to crystallization of immunoglobulin, in MBCL. This phenomenon has previously been reported in CLL.

CONFLICT OF INTEREST

The authors declare no conflict of interest.

DATA AVAILABILITY STATEMENT

Data sharing is not applicable to this article as no new data were created or analyzed in this study.

ORCID

Barbara J. Bain  <https://orcid.org/0000-0003-3077-4579>

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