We report an autopsy case of acute myocarditis, in which the mediastinal lymph nodes exhibited unique findings. A 15-year-old Japanese boy was diagnosed with the secondary onset of acute myocarditis. No viruses were identified. Autopsy confirmed acute lymphocytic myocarditis. Lymphadenopathy was observed, especially in pulmonary hilar/mediastinal areas. Microscopically, interfollicular areas were uniformly filled with medium-sized, round cells that resembled lymphocytes. They were immunohistochemically CD3⁻ CD5⁻ CD19⁻ CD20⁻ Pax-5⁻ CD138⁺ MUM1⁻ LMP1⁻ EBNA2⁻ cytoplasmic IgG⁺ IgA⁻ and IgM⁻. No monotypia was observed for kappa and lambda light chains, and multiplex polymerase chain reaction analyses of immunoglobulin heavy chain variable region diversity demonstrated oligoclonal peaks, suggesting reactive change. IgG⁺ or VS38c⁺ cells frequently co-expressed Ki-67 (up to 80%). We considered these cells abundantly present in lymph nodes to be reactive plasmablasts because they were early plasma cells with proliferative activity.

Keywords: plasmablasts, reactive lymphadenopathy, double-labeling immunohistochemistry
2) no increase in germinal centers, and 3) vascular transformation of lymphatic sinuses. The basic architecture of lymph nodes was preserved. Retroperitoneal or mesenteric lymph nodes demonstrated no notable microscopic changes (Fig. 1B). Higher magnification in pulmonary hilar nodes revealed that interfollicular areas were filled with diffuse proliferation of round cells with scant amphophilic cytoplasm, which were larger (up to two-times) than small lymphocytes (Fig. 1C, D, Fig. S3). They occasionally had one or two prominent nucleoli. These cells were later identified as plasmablasts for the reasons described below. Typical plasma cells were hardly observed. Neutrophils were sporadically observed among round cells. The above morphological findings suggested T cell responses (i.e., paracortical hyperplasia) caused by infection of unidentified organisms. However, the immunohistochemical results were unexpected: most round cells in the interfollicular areas were CD3− CD5− CD19− CD20− CD79a− Pax-5− CD138+ (up to 50%), MUM1+, LMP1−, EBNA2−, and cytoplasmic IgG+ IgA−, and IgM− (Fig. 2A-D, Fig. S4A-D). Kappa and lambda light chain staining did not demonstrate monotypia (Fig. 2E, F). As Ki-67+ cells were abundant in this area, we performed double staining. First, we confirmed that neither CD3+ T nor CD20+ B cells in the expanded interfollicular area correspond to Ki-67+ cells (Fig. 3A, Fig. S5A). In sharp contrast, immunofluorescence staining revealed that both cytoplasmic IgG+ cells and VS38c+ cells frequently co-expressed Ki-67 (78% among IgG+ cells) (Fig. 3B, C). Monoclonal antibody VS38c, which recognizes rough endoplasmic reticulum, is a marker of plasma cell differentiation among lymphoid cells.3 Based on the staining results, the cytoplasmic IgG-positive round cells were considered to have significant proliferative activity. These cells corresponded to plasmablasts based on the definition used mainly in the immunology field because plasmablasts are defined as early plasma cells that have proliferative and/or migratory activity.4-6

To exclude the possibility of neoplastic changes, genomic DNA was extracted from the formalin-fixed, paraffin-embedded sections of pulmonary lymph nodes and junctional diversity of the immunoglobulin heavy chain variable regions was analyzed according to the standardized multiplex PCR protocol.7 Amplification with the FR3-JH primer set yielded several distinct peaks embedded within a polyclonal background (Fig. S6A), whereas no monoclonal peak was observed with any of the used primer sets, including DH1-6-JH (Fig. S6B). This suggested that the plasmablast expansion was oligoclonal in nature.

Lymph nodes in other regions of the body (i.e.,
retroperitoneal or mesenteric lymph nodes) were not abundant in IgG+ Ki-67+ cells (Fig. S5B), highlighting the peculiarity of lymph nodes in the pulmonary hilar and mediastinal regions. Plasmablasts were not detected in the liver (1,685 g), spleen (210 g), or bone marrow, and peripheral blood analysis did not demonstrate a significant number of plasmablasts. A true paracortex (T cell zone) with abundant CD3+ T cells was located only beneath the surface area of the pulmonary hilar lymph nodes. Of note, some T cells in the paracortex co-expressed Ki-67 (Fig. S5C), suggesting the presence of antigenic stimuli. Negative controls did not exhibit specific reactivity (Fig. S5D). No other inflammatory diseases or findings of viral infection were observed in other organs or tissues.

Fig. 2. Immunohistochemistry for round cells (plasmablasts) in the extrafollicular areas of the pulmonary hilar nodes, labeled by antibodies against A) CD19, B) CD20, C) IgG, D) IgM, E) kappa chain, and F) lambda chain. Plasmablasts are CD19+ CD20- IgG+ IgM-. Sparse CD20+ cells were judged to be reactive B cells. No monotypia was observed for kappa and lambda chains. Scales: 20 μm (A-D).

Fig. 3. Multi-labeling immunohistochemistry for phenotypic characterization of plasmablasts in the pulmonary hilar lymph nodes. A) Chromogenic method. Ki-67+ cells (brown) did not correspond to CD20+ (red) or CD3+ (green) cells. B), C) Immunofluorescence method. Merged figures showing that IgG+ cells (B, green) and VS38c+ cells (C, green) were frequently double-positive for Ki-67 (red). Note that Ki-67 is positive in the nucleus (located in the center). Round cells in the extrafollicular areas were finally identified as plasmablasts. Scales: 20 μm (A-C). Methods are described in the legends of Supplementary Figure 5.

**DISCUSSION**

We judged the pulmonary hilar and mediastinal lymph node pathology described above to be reactive lymphadenopathy with abundant plasmablasts. Such cells are early...
plasma cells that have proliferative and/or migratory activity. Of note, this usage of plasmablasts is different from the same term used for neoplasms (for example, plasmablastic lymphoma).\(^8\) We demonstrated that double-labeling immunohistochemistry for immunoglobulin and Ki-67, in addition to VS38c and Ki-67, was useful to identify plasmablasts in tissue sections. The tissue preservation of lymph nodes in the present case was fairly good for an autopsy case, probably because ECMO was used until the terminal stage, which enabled us to apply the above double-staining technique and PCR analyses.

The present patient was 15 years old. More active immune responses may occur in lymph nodes in young patients than in older adults. However, a previous report on reactive lymphadenopathy in children did not describe a similar change.\(^9\) Moreover, a report on lymphadenopathy associated with plasma cell responses did not describe cases similar to ours.\(^10\) To our knowledge, lymph node lesions similar to those in our case, including childhood and adult cases, have not been reported.

The lesion we observed may have been caused by reactions by memory B lymphocytes, resulting in the rapid accumulation of IgG\(^+\) (not IgM\(^+\)) plasmablasts in the medullary cord of lymph nodes without de novo formation of germinal centers,\(^11,12\) which was observed in our case. If the present lymphadenopathy was caused by a primary immune response, a mixture of IgM\(^+\) and IgG\(^+\) plasma cells, and proliferating B cells with the development of germinal centers,\(^11,12\) which was observed in our case. The cause of the observed lymph node pathology remains unclear because no viruses were identified. However, the spatial relationship between the heart and the location of lymphadenopathy (only observed in the mediastinal areas) suggests that the lymphadenopathy was related to myocarditis.

One of the authors (HO) previously reported that the base of ulcerative colitis is abundant 'proliferating plasma cells'.\(^13\) We (MM and HO) now consider the previously described 'proliferating plasma cells' to be plasmablasts for the following reasons: a) these round cells were immunohistochemically CD19\(^+\) CD20\(^-\) CD3\(^-\) CD138\(^+\) IgG\(^+\) or IgA\(^+\) (but IgM\(^-\)), and b) CD19\(^+\) or IgG\(^-\) cells were frequently double-positive for Ki-67 (Table S2). Immuno-electron microscopy revealed that CD19\(^+\) cells had well-developed rough endoplasmic reticulum, a feature consistent with antibody-secreting cells. In relation to this, flow cytometry revealed plasmablasts (i.e., early plasma cells with migratory potential) in the peripheral blood of patients with ulcerative colitis.\(^14\)

The present report described a case of plasmablast-rich, reactive lymphadenopathy. Further studies are required to clarify the significance of the lesion and the distribution of reactive plasmablasts in different inflammatory lesions. Our double-staining method for immunoglobulin and Ki-67 may be useful for this purpose.

**ACKNOWLEDGMENTS**

We are grateful to Dr. Roderick E. Mitcham, Kindai University Faculty of Medicine, Japan, for critical reading of the manuscript. Technical assistance by Mr. Kakeru Sannomaru was greatly appreciated.

**INFORMED CONSENT**

Written informed consent was received from the parents of the patient.

**AUTHOR CONTRIBUTIONS**

HO performed all pathological analyses. MM interpreted the pathological data from the immunological viewpoint. YN, TM, LL, and JS treated the patient and summarized the clinical records. HO drafted the manuscript, and all authors participated in the discussion and completion of the manuscript.

**CONFLICT OF INTEREST**

None declared.

**REFERENCES**

1. Bracamonte-Baran W, Čiháková D. Cardiac Autoimmunity: Myocarditis. Adv Exp Med Biol. 2017; 1003: 187-221.
2. Pollack A, Kontorovich AR, Fuster V, Dec GW. Viral myocarditis—diagnosis, treatment options, and current controversies. Nat Rev Cardiol. 2015; 12: 670-680.
3. Turley H, Jones M, Erber W, et al. VS38: a novel monoclonal antibody for detecting plasma cell differentiation in routine sections. J Clin Pathol. 1994; 47: 418-422.
4. Nutt SL, Hodgkin PD, Tarlinton DM, Corcoran LM. The generation of antibody-secreting plasma cells. Nat Rev Immunol. 2015; 15: 160-171.
5. Tellier J, Nutt SL. Plasma cells: The programming of an antibody-secreting machine. Eur J Immunol. 2019; 49: 30-37.
6. Tarlinton D. Plasma cell biology. In : Honjo T, Reth M, Radbruch A, et al. (eds) : Molecular Biology of B Cells, 2nd ed, Cambridge, Academic Press, 2015; pp. 232-243.
7. van Dongen JJM, Langerak AW, Brüggemann M, et al. Design and standardization of PCR primers and protocols for detection of clonal immunoglobulin and T-cell receptor gene recombination in suspect lymphoproliferations: Report of the BIOMED-2 Concerted Action BMH4-CT98-3936. Leukemia. 2003; 17: 2257-2317.
8. Campo E, Stein H, Harris NL. Plasmablastic lymphoma. In : Swerdlow HS, Campo E, Harris NL, et al. (eds) : WHO Classification of Tumours of Haematopoietic and Lymphoid Tissues, Revised 4th ed, Lyon, IARC. 2017; pp. 321-322.
9. Ramsay AD. Reactive lymph nodes in pediatric practice. Am J Clin Pathol. 2004; 122(suppl): S87-S97.
10. Xie Y, Vallangeon B, Liu X, Lagoo AS. Plasmacytic or lymphoplasmacytic infiltrate in lymph nodes: Diagnostic approach and
differential considerations. Indian J Pathol Microbiol. 2016; 59: 446-456.
11 Kurosaki T, Kometani K, Ise W. Memory B cells. Nat Rev Immunol. 2015; 15: 149-159.
12 Hoffman W, Lakkis FG, Chalasani G. B Cells, antibodies, and more. Clin J Am Soc Nephrol. 2016; 11: 137-154.
13 Jinno Y, Ohtani H, Nakamura S, et al. Infiltration of CD19+ plasma cells with frequent labeling of Ki-67 in corticosteroid-resistant active ulcerative colitis. Virchows Arch. 2006; 448: 412-421.
14 Tarlton NJ, Green CM, Lazarus NH, et al. Plasmablast frequency and trafficking receptor expression are altered in pediatric ulcerative colitis. Inflamm Bowel Dis. 2012; 18: 2381-2391.