Frequency and distribution of CD4+CXCR5+ follicular B helper T cells within involved tissues in IgG4-related ophthalmic disease

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Abstract. Immunoglobulin G4-related ophthalmic disease (IgG4-ROD) is a IgG4-RD and exhibits two main pathologically; and frequent formation of germinal centers (GCs). Follicular B helper T (Tfh) cells are now recognized as the true helper cells for B cells in antibody responses. Tfh cells were first described in the year 2000 when CD4+CXCR5+ Tfh cells were observed in patients with IgG4-ROD and IgG4-RLF. CD4+CXCR5+ Tfh cells were randomly distributed in areas demonstrating type 1 AIP. IL-21 expression was positively correlated with IL-21-mediated pathogenetic role in IgG4-ROD and type 1 AIP. Tfh cells may serve a direct role in the IL-21-mediated pathogenetic role in IgG4-ROD.

Introduction

Immunoglobulin G4-related disease (IgG4-RD) is a chronic disorder of an unknown etiology with multiple organ involvement. It was first observed in the pancreas and is now known as type 1 autoimmune pancreatitis (AIP)/IgG4-related pancreatitis (1-3). This condition has been observed to affect virtually every organ: Orbital contents; bile ducts; salivary glands; retroperitoneum; pachymeninges; kidneys; lungs; lymph nodes; aorta; arteries; breast; prostate; thyroid; pericardium; and skin (3,4). Clinicopathological features demonstrate striking similarities among the involved organs, including a tendency to form tumefactive lesions, a dense lymphoplasmacytic infiltrate rich in IgG4-positive plasma cells, storiform fibrosis, frequent elevations of serum IgG4 levels and good response to glucocorticoids (5,6). However, as a member of a broad spectrum of IgG4-RD diseases, IgG4-R ophthalmic (O)D (affects lacrimal glands, extraocular muscles, orbital nerve and eyelid) differs from the other members of the IgG4-RD group by two aspects: Fibrosis is not necessarily marked histopathologically, and germinal centers (GCs) are frequently observed (3,7). GCs are distinct structures within the B cell follicles in the secondary lymphoid tissues and are major sites of somatic hypermutation, class switching-associated recombination and affinity maturation of activated B cells, and the production of plasma cells and memory B cells (8,9). GCs depend on cluster of differentiation 4-positive (CD4+) T cells and it is now recognized that a subset of CD4+ T cells, termed follicular B helper T (Tfh) cells, are the true helper cells for B cells in antibody (Ab) responses (10,11).

Tfh cells were first described in the year 2000 when several groups reported that a significant proportion of C-X-C chemokine receptor type 5 (CXCR5)+CD4+ T cells involved...
in B cell help for Ab responses in tonsils was situated in GCs or follicles. These cells were subsequently termed Tfh cells (12,13). Following 17 years of research, there is now an improved understanding of the cellular and molecular mechanisms of helper functions of Tfh cells. The CXCR5 serves as a surface marker for Tfh cells, CD4+ T cells expressing CXCR5 can migrate in response to the follicular chemokine B lymphocyte chemoattractant (also known as CXCL13), and relocate to the border of T and B cell zone (T-B junction) and to the follicles in secondary lymphoid organs (14,15). B cell lymphoma 6 (Bcl-6) was strongly expressed in Tfh cells but not in T helper (Th)1 or 2 cells (16). Bcl-6 has been identified as a master regulator of Tfh cell differentiation in 2009 (17-19), in the absence of Bcl-6, Tfh cell differentiation ceases in vivo (17-19) while other CD4+ Th cell subsets are relatively unaffected (18,19). Tfh cells also express high levels of inducible T-cell costimulator (ICOS) and interleukin (IL)-21 (13,16,20). IL-21 is a Tfh cell-expressed helper cytokine (10) and the upregulation of ICOS is essential for the initiation and maintenance of Tfh differentiation (21,22).

The pathogenesis of IgG4-RD remains to be elucidated. Researchers in the field of immunology identified certain abnormal immunological mechanisms involved in the pathogenesis of IgG4-RD. The results suggested that Th2 cells and regulatory T cells (Tregs) as well as associated cytokines, including IL-4, IL-10, IL-13 and transforming growth factor β1, may serve key roles in the development of Mikulicz disease, IgG4-related pancreatitis and cholangitis (23,24). More recently, studies on the role of Th2 cytokines in IgG4-RD indicated that IL-18 and interferon-γ are implicated in the pathogenesis of IgG4-related dacyoadenitis and sialoadenitis (25,26). Currently, increasing attention is paid to Tfh cells; a higher frequency of GCs in patients with IgG4-RD indicates an increased number of activated B cells, which may reflect the increase in Tfh cell number.

An association between Tfh cells and autoimmunity has been suggested in many autoimmune diseases, including systemic lupus erythematosus (27,28), autoimmune thyroid disease (29) and myasthenia gravis (30,31). Since the major function of Tfh cells is to aid B cell activation and to stimulate Ab responses, patients with IgG4-RD display elevated IgG4 concentrations in sera and infiltrated IgG4-positive cells in associated tissues, suggesting the involvement of Tfh cells in the pathogenesis. In the studies performed with peripheral blood samples, Tfh cells were related to the pathogenic process of IgG4-RD (32,33). The profile and distribution of Tfh cells in involved tissue, which demonstrates the association between Tfh cells and IgG4-RD directly, has not been examined among patients with IgG4-RD and IgG4-RD patients demonstrating GC formation.

In the following study, the expression of ICOS and Bcl-6 was increased in tissues of patients with IgG4-RD compared with IgG4-negative controls. It was also demonstrated that CD4+CXCR5+ Tfh cells were more abundant in patients with IgG4-RD compared with the IgG4-negative controls. Among patients with IgG4-RD, CD4+CXCR5+ Tfh cells were present in and outside GCs in patients with IgG4-RD and IgG4-related lymphadenopathy (IgG4-RL) while staining for CD4 and CXCR5 in pancreatic lesions appeared as a randomly distributed pattern. Fewer CD4+CXCR5+ Tfh cells were observed in patients with type 1 AIP compared with patients with IgG4-ROD and IgG4-RL. The expression of IL-21 was coincidental with dual immunofluorescence of CD4 and CXCR5, and increased expression of IL-21 was observed in patients with IgG4-RD and IgG4-RL. IL-21 expression in patients with IgG4-RD was positively correlated with the IgG4/IgG ratio in immunohistochemically (IHC)-positive cells.

Materials and methods

Patients and samples. A total of 7 patients with IgG4-RD (4 men and 3 women; mean age ± standard deviation age 60.9±7.5 years) referred to the Department of Ophthalmology at the Shanghai Changzheng Hospital (Shanghai, China) between January 2013 and June 2016 were included in the present study. A total of 7 patients with type 1 AIP or IgG4-RL (6 men and 1 woman; mean age ± standard deviation age 56.0±7.0 Years) who were referred to Renji Hospital (Shanghai, China) between July 2012 and June 2016 were studied as disease positive controls. Cases with normal serum IgG4 and no IgG4-positive cells detected in tissue specimens (n=7) were designated as IgG4-negative controls. IgG4-RD was diagnosed using the following criteria (5,34): i) Clinical examination reveals diffuse/localized swelling or masses in single or multiple organs; ii) tissue biopsy meets at least two out of three major histopathologic features of the disease: A lymphoplasmacytic infiltrate, fibrosis (typically storiform); and oblitative phlebitis; and iii) infiltration of IgG4-positive plasma cells: Ratio of IgG4-positive cells to IgG-positive cells of >40% and >10 IgG4-positive cells/high power field (magnification, x400).

The lesion specimens were collected from all patients with IgG4-RD from whom written informed consent was obtained, and tissues were used only for research purposes. The study was approved by the Ethics Committee of Renji Hospital.

IHC staining. The primary antibodies used for the IHC were as follows: Anti IgG4 (cat. no. ab109493; Abcam, Shanghai, China; at 1/500 dilution); IgG (cat. no. ab109489; Abcam, Shanghai, China; at 1/300 dilution); Bcl-6 (cat. no. ab18485; Abcam, Shanghai, China; at 1/50 dilution); ICOS (cat. no. ab105227; Abcam, Shanghai, China; at 1/50 dilution); IL-21 (cat. no. 118510; Abcam, Shanghai, China; at 1/200 dilution).

IHC analyses. Paraffin sections (4 µm-thick) were deparaffinized by immersion in xylene, followed by a hydration in ethanol. An antigen retrieval was performed with EDTA pH 8.0 microwave repair method. Endogeneous peroxide activity was quenched by 3% hydrogen peroxide in water. Then the sections were incubated for 30 min at room temperature with 3% bovine serum albumin (BSA) (cat. no. A8020; Beijing Solarbio Science and Technology Co., Ltd., Beijing, China) and incubated overnight at 4°C with primary antibody. Thereafter, the sections were incubated with a horseradish peroxidase-conjugated secondary antibody (cat. no. K5007; Dako; Agilent Technologies, Inc., Santa Clara, CA, USA; at 1/1 dilution) for 50 min at room temperature. The stain was prepared using diaminobenzidine substrate (cat. no. K5007; Dako; Agilent Technologies, Inc.). Sections were counterstained 3 min at room temperature with Harris hematoxylin and mounted prior to analysis. Between each
step of the analysis, sections were rinsed with PBS three times, 5 min each. IHC images were captured using a light microscope equipped with a digital camera (Olympus Corporation, Tokyo, Japan). Stained IgG4- and IgG-positive cells were counted in three high-power fields with the highest density of positive cells and an average was calculated (5). Semiquantification of IL-21 was performed as previously described with a slight modification in the grade classification of staining intensity and percentage of positive cells (35). Depending on the proportion of positive stained cells (PP): <5% was scored 0, 6-25% was scored 1, 26-50% was scored 2, 51-75% was scored 3 and >75% was scored 4. According to the staining intensity (SI): No color was scored 0, light yellow was scored 1, light brown was scored 2, brown was scored 3. PP was multiplied by SI to obtain a total IHC score for any given sample.

**Table I. Clinical and serological characteristics of patients with immunoglobulin G4-related ophthalmic disease.**

| Patient no. | Sex | Age at onset, years | Onset time | Exophthalmos (mm) Right eye (orbital distance) left eye | Onset signs and symptoms | IgG (g/l) | IgG4 (g/l) | IgE IU/ml |
|-------------|-----|---------------------|------------|--------------------------------------------------------|--------------------------|----------|-----------|-----------|
| 1           | M   | 74                  | 1 year     | 15 (105) 14 | Diplopia, proptosis, swelling | Bil | 33.7 | 38.5 | 368 |
| 2           | M   | 57                  | 8 years    | 24.5 (113) 31 | Proptosis, swelling | Bil | 31.3 | 43.2 | 415 |
| 3           | F   | 68                  | 3 months   | 20 (103) 24 | Proptosis | Bil | 9.77 | 3.26 | 415 |
| 4           | F   | 59                  | 6 months   | 13 (110) 13 | Swelling | Bil | 22.6 | 20.2 | 1140 |
| 5           | M   | 59                  | 5 years    | 14 (110) 20 | Swelling | Bil | 18.9 | 12.3 | 5 |
| 6           | F   | 52                  | 5 months   | 14 (107) 15 | Swelling, ptosis | Left | 16.2 | 8.48 | 495 |
| 7           | M   | 57                  | 7 years    | 26 (99) 28 | Proptosis, swelling | Bil | 25.8 | 17.5 | 319 |

M, male; F, female; Bil, bilateral; IgG, immunoglobin G.

**Table II. Clinical and serological characteristics of patients with type 1 AIP or IgG4-RL and IgG4-negative controls.**

A. Patients with type 1 AIP or IgG4-RL

| Patient no. | Sex | Age at onset, years | Affected organ | IgG4 (g/l) | Onset time | Onset signs and symptoms |
|-------------|-----|---------------------|----------------|-----------|------------|--------------------------|
| 8           | M   | 57                  | Pancreas, lymph node | N          | 0.5 months | Asymptomatic, occupying |
| 9           | M   | 67                  | Pancreas       | N          | 10 days    | Abdominal pain, swelling |
| 10          | M   | 62                  | Pancreas       | N          | 2 months   | Abdominal pain, swelling |
| 11          | M   | 49                  | Pancreas, lymph node | 10.1       | 2 weeks    | Asymptomatic, occupying |
| 12          | M   | 54                  | Lymph node    | 28.1       | 1 years    | Swelling |
| 13          | M   | 56                  | Lymph node    | N          | 0.5 months | Swelling |
| 14          | F   | 47                  | Lymph node    | N          | 4 years    | Swelling |

B. IgG4-negative controls

| Patient no. | Sex | Age at onset, years | Affected organ | IgG4 (g/l) | Onset time | Onset signs and symptoms |
|-------------|-----|---------------------|----------------|-----------|------------|--------------------------|
| 15          | F   | 25                  | Labial gland   | 0.85      | 1.5 years  | Dry mouth, dry skin |
| 16          | F   | 52                  | Orbital tissue | 0.35      | 10 years   | Swelling |
| 17          | F   | 44                  | Peripancreatic lymph node | 0.90      | 1 years    | Abdominal pain, swelling |
| 18          | M   | 59                  | Submandibular gland | 1.27      | 1 month   | Swelling |
| 19          | M   | 43                  | Pancreas       | 0.71      | 1 year     | Abdominal pain |
| 20          | F   | 55                  | Orbital tissue | 1.25      | 7 months   | Orbital pain |
| 21          | F   | 63                  | Bile duct     | 0.74      | 2 months   | Yellowing and pruritus of the skin |

M, male; F, female; N, not measured; type 1 AIP, autoimmune pancreatitis; IgG4-RL, immunoglobin G4-related lymphadenopathy.

**Double-labeled immunofluorescence for CD4, CXCR5 and IL-21.** Sections were de-waxed by xylene, then hydrated through...
a graded ethanol series, and washed with distilled water. Antigen retrieval and serum block were performed using a similar method to the IHC staining. The primary antibodies used for the analysis of co-expression of CD4 and CXCR5 were anti-CD4 (cat. no. GB13064; Goodbio technology, Wuhan, Hubei, China; at 1/50 dilution), anti-CXCR5 (cat. no. GTX100351; GeneTex, Inc., Irvine, CA, USA; at 1/100 dilution). The anti-IL-21 goat polyclonal antibody was used to analyze the co-expression of IL-21 and CXCR5 (cat. no. GTX82914; GeneTex, Inc.; at 1/200 dilution while anti-CXCR5 at 1/200 dilution). Cy3-conjugated goat anti-mouse IgG (cat. no. GB21301; Goodbio technology; at 1/300 dilution), Alexa 488-conjugated goat anti-rabbit IgG (cat no. GB25303; Goodbio technology; at 1/400 dilution), Cy3-conjugated donkey anti-goat IgG (cat no. GB254031; Goodbio technology; at 1/300 dilution), Alexa 488-conjugated donkey anti-rabbit IgG (cat no. GB254031; Goodbio technology; at 1/400 dilution) were used as fluorochrome-conjugated secondary antibodies. Then the sections were sequentially incubated with primary antibodies overnight at 4°C, corresponding second antibodies for 50 min at room temperature and DAPI for 10 min at room temperature. Between each step, sections were rinsed with PBS three times, 5 min each. Fluorescence images were captured using a fluorescence microscope (Nikon Corporation).

Statistical analysis. Differences in the IHC staining scores between more than two groups were determined using the Kruskal-Wallis test and differences between two groups were determined using the Mann-Whitney U test (two-tailed). Spearman’s rank correlation coefficient was used to assess the correlation between IL-21 expression and the ratio of IgG-positive cells to IgG-negative cells in IgG4-ROD samples. All statistical analyses were performed using SPSS software (version 20; IBM Corp., Armonk, USA). P<0.05 was considered to indicate a statistically significant difference.

Results

Patient profiles. The clinical and serological characteristics of patients with IgG4-ROD are presented in Table I. Patients with IgG4-ROD were predominantly middle-aged to elderly males. IgG4-ROD affected the lacrimal gland in 6 patients the orbital soft tissue in 2 patients, including the orbital apex. Computed tomography or magnetic resonance imaging identified space-occupying lesions of lacrimal gland or orbital cavity. Serum IgG and IgG4 were elevated in all patients with a median level of 22.6 g/l (P25-P75: 16.2-31.3 g/l) and 17.5 g/l (P25-P75: 8.5-38.5 g/l), respectively. With the exception of 1 patient, the serum IgE value for all patients was elevated above the normal limit (<165 IU/ml). Case 2 exhibited para-nasal sinusitis and case 5 exhibited bronchial asthma. Clinical and serological characteristics of patients with type 1 AIP or IgG4-RL and IgG4-negative controls are summarized in Table II. Immunohistochemistry results demonstrated that IgG4-positive cells mainly infiltrated the interfollicular areas in lymph nodes (Fig. 1A), areas around the acinar and ductal cells in pancreas (Fig. 1B) and diffusely in lacrimal glands (Fig. 1C).

Expression of Bcl-6 and ICOS. The expression of Bcl-6, a master regulator transcription factor in Tfh cells and ICOS, a surface marker of Tfh cells, was examined (Fig. 2). In patients with IgG4-ROD, the nuclear expression of Bcl-6 and expression of ICOS were predominantly detected in GCs, and to a lesser extent in infiltrating lymphocytes outside the GCs (Fig. 2A and B). A small proportion of Bcl-6- and ICOS-positive cells were randomly distributed within the pancreatic lesions (Fig. 2E and F). In lymph nodes from patients with IgG4-RL positive staining for Bcl-6 and ICOS was only observed in GCs (Fig. 2C and D). By contrast, the nuclear expression of Bcl-6 was detected in relatively few lymphocytes in all IgG4-negative controls and ICOS-positive cells were rarely observed (Fig. 2G and H).

Increased number of CD4+CXCR5+ cells in involved tissues of patients with IgG4-ROD. The above results indicated that expression of Bcl-6 and ICOS, which are the markers for Tfh cells, were increased in patients with IgG4-ROD compared with patients with IgG4-negative. CD4+CXCR5+ cells were examined using dual immunofluorescence staining for CD4 and CXCR5 using paraffin-embedded specimens from patients...
with IgG4-RD and IgG4-negative patients (Figs. 3 and 4). In the IgG4-RD group, CD4+CXCR5+ cells were randomly distributed in areas of type 1 AIP (Figs. 3 and 4E) and were predominantly detected in GCs and T-B junctions in lymph nodes from IgG4-RL (Fig. 4C). CD4+CXCR5+ cells were scattered and distributed in GCs within the orbital lesions in patients with IgG4-ROD (Figs. 3 and 4A). In IgG4-negative controls, numerous CD4-positive cells were observed; however, few CD4+CXCR5+ cells could be identified (Fig. 3). The number of CD4+CXCR5+ cells among involved tissues from patients with IgG4-RD demonstrated significantly decreased CD4+CXCR5+ cells in patients with type 1 AIP compared with patients with IgG4-ROD and IgG4-RL (P=0.024 and 0.032, respectively), there was no significant difference between IgG4-ROD and IgG4-RL (Fig. 5).

Expression of IL-21 in involved tissues of patients with IgG4-ROD. IL-21 is the main effector cytokine produced by Tfh cells (10) and the present study, using immunohistochemical analyses, determined whether IL-21 is expressed in samples from patients with IgG4-RD. All 16 IgG4-RD samples expressed IL-21 (Fig. 6A). IL-21 was prominent in GCs and less evident in infiltrating lymphocytes outside the GCs in orbital lesions and lymph nodes. Staining for IL-21 in pancreatic lesions appeared as a scattered pattern, primarily in infiltrating lymphocytes around the acinar and slightly detected in ductal cells. Fluorescence microscopy revealed that the IL-21 expression in involved tissues from patients with IgG4-RD coincided with their CXCR5 expression (Fig. 6B). While the scores from the IHC staining exhibit decreased expression of patients with IL-21 in type 1 AIP compared with patients with IgG4-RD and IgG4-RL (P=0.042 and 0.016, respectively), the scores were comparable between IgG4-RD and IgG4-RL patients (Fig. 6C).

Association between IgG4 production and expression of IL-21 in patients with IgG4-ROD. The association between IgG4 production and the levels of IL-21 IHC scores in orbital lesions was examined. Spearman’s rank correlation analysis demonstrated that the IL-21 IHC scores in orbital lesions were positively correlated with the ratio of IgG4-positive cells to IgG-positive cells in patients with IgG4-ROD (Fig. 6D).

Discussion

The present study analyzed the expression of the key Tfh cell immunological proteins ICOS, Bcl-6, CXCR5 and IL-21 in the affected tissues of patients with IgG4-RD, using IHC and dual immunofluorescence. The results demonstrated increased expression of ICOS and Bcl-6 in the involved tissues of patients with IgG4-RD compared with the IgG4-negative control group. The CD4+CXCR5+ Tfh cells were more prevalent in IgG4-RD and scarce in the IgG4-negative control group. Among patients with IgG4-RD, CD4+CXCR5+ Tfh cells were detected inside and outside GCs in patients with IgG4-ROD and IgG4-RL, whereas staining for CD4 and CXCR5 in pancreatic lesions appeared as a randomly distributed pattern. A decreased number of CD4+CXCR5+ Tfh cells was observed in patients with type 1 AIP compared with IgG4-ROD and IgG4-RL patients. The expression of IL-21 was almost coincidental with the dual immunofluorescence of CD4 and CXCR5 and elevated expression of IL-21 was observed in patients with IgG4-ROD and IgG4-RL. IL-21 expression was positively correlated with the IgG4/IgG ratio in IHC positive cells.

Naive CD4+ T cells may differentiate into one of several lineages of Th cells and exert a variety of immunological functions. These include clearing viruses, helminths and fungi by Th1, Th2 and Th17 cells, respectively, and suppressing immune responses by induced Treg cells (36). Tfh cells are a subset of CD4+ T cells that provide help for B cells during the GC reaction (10). A large GC is frequently formed within involved organs in individuals with IgG4-RD (7,37) and is presumably the source of IgG4-positive plasma cells. Based on this observation, the present study targeted IgG4-ROD and investigated the Tfh cell abundance in affected tissues.
Tfh cells depend on the expression of the master regulator transcription factor Bcl-6 and distinguishing features of Tfh cells are the expression of ICOS, CXCR5, IL-21, programmed cell death protein 1 and SLAM-associated protein (10). The analysis of the expression of Bcl-6 and ICOS in patients with IgG4-RD and IgG4-negative controls, revealed elevated expression of ICOS and Bcl-6 in the involved tissues of patients with IgG4-RD. Tfh cells require upregulation of Bcl-6 to develop and perform their function in B cell maturation in GCs (38). Bcl-6 binding is associated with the control of Tfh cell migration and repression of alternative cell fates (39). The upregulation of ICOS is essential for the initiation and maintenance of Tfh differentiation, and CD4+CXCR5+ cells were scarce in mice with B cell-specific deletion of ICOS ligand (11,21). Increased expression of both Bcl-6 and ICOS in involved tissues of patients with IgG4-RD indicated the presence of Tfh cells.

Dual immunofluorescence of paraffin-embedded specimens revealed increased numbers of CD4+CXCR5+ Tfh cells
in all cases of IgG4-RD irrespective of the organs involved, and an enhanced expression of Bcl-6 and ICOS, which histopathologically indicated that this cell population was involved in the pathogenesis of IgG4-RD. The frequency and distribution of CD4+CXCR5+ Tfh cells differed in patients with IgG4-ROD and type 1 AIP. CD4+CXCR5+ Tfh cells were observed inside and outside GCs in patients with IgG4-ROD, however they were randomly distributed and fewer in areas of type 1 AIP compared with IgG4-ROD. The distribution of CD4+CXCR5+ Tfh cells corresponded with the expression of Bcl-6 and ICOS. CXCR5 is the canonical Tfh marker and in the present study, CD4+CXCR5+ cells which localized in GCs were considered as GC Tfh cells and predecessors of those cells with Tfh-like characteristics (localized in extrafollicles) were termed Pre-Tfh cells (10,11,40). Human autoimmune diseases predominantly affect nonlymphoid tissues, which frequently contain large numbers of infiltrating, activated lymphocytes, as well as lymphoid-like tissues with a GC. Tfh cells have been identified in nonlymphoid tissues, particularly in autoimmune diseases (41). High frequency of GC formation and an increased number of Tfh cells in IgG4-ROD may suggest that IgG4-ROD is an autoimmune disease and may also indicate that Tfh cells serve a different role in IgG4-ROD and type 1 AIP, since GCs are not commonly observed in type 1 AIP (42). Previous studies considered blood CD4+CXCR5+ cells as equivalents of Tfh cells. Akiyama et al (32) demonstrated that the proportion of circulating Tfh cells was significantly increased in patients with IgG4-RD compared with patients with allergic rhinitis or with healthy controls, and these results are consistent with those obtained in the present study (11,15).

Tfh cells are a predominant source of IL-21 (43). In the present study, IL-21 was abundant in GCs in orbital lesions and lymph nodes, whereas staining for IL-21 in pancreatic lesions appeared as a scattered pattern, primarily in infiltrating lymphocytes around the acinar and ductal cells. Expression of IL-21 was weaker in Type 1 AIP patients compared with IgG4-ROD and IgG4-RL patients. The IL-21 expression in involved tissues from patients with IgG4-RD was coincidental with their CXCR5 expression. Concurrent expression of IL-21 and CXCR5, together with the distribution of IL-21-positive cells indicates that the IL-21 in affected tissues of IgG4-RD is mainly produced by Tfh cells and the high expression of IL-21 in IgG4-RD is due to the increased number of Tfh cells. IgG4 production was positively correlated with the IL-21 IHC scores in patients with IgG4-RD. IL-21 can promote CD40L-mediated GC B cell proliferation and drive human B cell differentiation.
into IgG4-related ophthalmic disease (IgG4-ROD) controls the maintenance and optimal affinity maturation of the GC reaction by maintaining the Bcl-6 expression in GC B cells in vivo (46,47). IL-21 can also contribute to IgG4 production in Mikulicz’s disease (48) and the data obtained in the present study demonstrates that Tfh-derived IL-21 can promote B cell differentiation into IgG4-secreting cells in patients with IgG4-ROD.

Based on the available data (32,33,41), it can be hypothesized that chronic stimulation of orbital tissues by an unknown antigen induces production of CXCL13, which in turn recruits circulating Tfh and B cells. Interaction between Tfh and B cells triggers GC formation and B cell differentiation into IgG4-producing cells through secretion of IL-21 in IgG4-ROD.

Steroids represent an effective short-term treatment of IgG4-RD, their effect typically becomes evident within weeks; however, 40% of patients relapse within the 1st year (49). Rituximab is a promising medication but resistant cases have been reported (50). The aforementioned results of the present study suggest that the use of T cell activation inhibitors for Tfh cells, including abatacept, could represent an effective treatment of IgG4-ROD. Yamamoto et al (51) recently reported a rituximab-resistant IgG4-ROD patient who showed a good response to treatment with abatacept. Since abatacept may affect Tfh cells in the GCs (51), it may exert greater effect on patients with IgG4-ROD. Understanding of the pathogenetic relevance of IL-17 in rheumatoid arthritis and collagen-induced arthritis led to the development of a semi-specific immunotherapy based on cytokine antagonism for the treatment of these disorders (52,53). Identification of IL-21 as a potentially key cytokine in IgG4-ROD may enable the use of specific IL-21 antagonists in the treatment of IgG4-ROD.

However, the significance of the present study is limited by the number of patients analyzed. As clinicians are becoming increasingly aware of IgG4-ROD and the administration of glucocorticoid therapy is more prevalent, invasive medical procedures are less prevalent, decreasing the availability of samples from IgG4-ROD patients, especially those with IgG4-ROD. Larger cohort studies performed in the future may further elucidate the underlying mechanism of Tfh cell action in affected tissues.

In summary, in the present study CD4+CXCR5+ Tfh cells were highly prevalent in IgG4-ROD, and the frequency and distribution of CD4+CXCR5+ Tfh cells differed between patients with IgG4-ROD and type 1 AIP, which may suggest different roles for Tfh cells in IgG4-ROD and type 1 AIP. IL-21 was highly expressed in patients with IgG4-ROD and Tfh-derived IL-21 can promote B cell to differentiate into IgG4-secreting cells in IgG4-ROD, suggesting that Tfh cells may have a direct role in the pathogenesis of IgG4-ROD. These results make Tfh cells and IL-21 an important focus of potential therapeutic methods to treat IgG4-ROD.

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