The Imbalance of Circulating Follicular T Helper Cell Subsets in Primary Sjögren’s Syndrome Associates With Serological Alterations and Abnormal B-Cell Distribution

Krisztina Szabo¹*, Ilona Jámbor¹†, Antónia Szántó¹, Ildikó Fanny Horváth¹, Tünde Tarr¹, Britt Nakken², Peter Szodoray² and Gábor Papp¹

¹ Division of Clinical Immunology, Faculty of Medicine, University of Debrecen, Debrecen, Hungary, ² Department of Immunology, Oslo University Hospital, Rikshospitalet, Oslo, Norway

Since B-cell hyperactivity and pathologic antibody response are key features in the immunopathogenesis of primary Sjögren’s syndrome (pSS), the role of follicular T helper (TFH) cells as efficient helpers in the survival and differentiation of B cells has emerged. Our aim was to investigate whether a change in the balance of circulating (c)TFH subsets and follicular regulatory T (TFR) cells could affect the distribution of B cells in pSS. Peripheral blood of 38 pSS patients and 27 healthy controls was assessed for the frequencies of cTFH cell subsets, TFR cells, and certain B cell subpopulations by multicolor flow cytometry. Serological parameters, including anti-SSA, anti-SSB autoantibodies, immunoglobulin, and immune complex titers were determined as part of the routine diagnostic evaluation. Patients with pSS showed a significant increase in activated cTFH cell proportions, which was associated with serological results. Frequencies of cTFH subsets were unchanged in pSS patients compared to healthy controls. The percentages and number of cTFR cells exhibited a significant increase in autoantibody positive patients compared to patients with seronegative pSS. The proportions of transitional and naïve B cells were significantly increased, whereas subsets of memory B cells were significantly decreased and correlated with autoantibody production. Functional analysis revealed that the simultaneous blockade of cTFH and B cell interaction with anti-IL-21 and anti-CD40 antibodies decreased the production of IgM and IgG. Imbalance in TFR subsets and TFR cells indicates an ongoing over-activated humoral immune response, which contributes to the characteristic serological manifestations and the pathogenesis of pSS.

Keywords: follicular T helper cell, follicular regulatory T cell, B cell, chemokine receptors, primary Sjögren’s syndrome
INTRODUCTION

Primary Sjögren’s syndrome (pSS) is a chronic autoimmune disease displaying slow progression and affecting primarily middle-aged women nine-times more frequently than men. Histologically, the hallmark feature of pSS is the periductal cellular infiltration of lacrimal and salivary glands resulting in the damage and dysfunction of glandular tissue. Clinically, patients suffer from mouth dryness (xerostomia) and eye dryness (xerophthalmia) predominantly; however, various exocrinopathic symptoms are also frequently present. The clinical manifestations in pSS are not homogenous, besides glandular symptoms; the disease can involve any organ system and present with a wide variety of extraglandular manifestations affecting skin, genitourinary tract, kidneys, cardiovascular, nervous or musculoskeletal systems (1).

The development of pSS is still unclear; however, recent lines of evidence support a complex network between salivary gland epithelial cells and the innate and acquired immune systems. Epithelial cells, besides being the target of extrinsic and intrinsic factors that cause disruption of glandular tissue integrity, are also active participants in the initiation and perpetuation of the inflammatory process, through the expression of activation markers and toll-like receptors. The upregulation of adhesion and chemokine molecules, along with metabolic stress induced apoptosis initiates the recruitment of innate immune cells and triggers an interferon-related inflammatory cascade (1). Type I interferon production by plasmacytoid dendritic cells induces infiltrating leukocytes and epithelial cells to secrete B-cell activating factor (BAFF), which promotes B-cell survival and not only contributes to B-cell hyperactivity, but also acts as a mediator between innate and adaptive immunity (2, 3). Local accumulation and sustained survival of autoreactive B cells promote the development of ectopic germinal centers (GCs), which establish the perfect niche for autoantibody production or promote the development of ectopic germinal centers (GCs), accumulation and sustained survival of autoreactive B cells

B-cell selection, development, and activation. A subset of CD4+ T cells is involved in both the first stage of their differentiation during GC response (6). The ability to differentiate into GC B cells and GC T FH cells depends on antigen-specific contact between these cells and the successful delivery of mutual signals from the very first stage of their interaction (7). T-cell-dependent B-cell response proceeds in three spatiotemporal phases in the lymphoid tissues, including extrafollicular, follicular, and GC reaction phase, which demands the expression of several molecules. B cells in GCs receive help through CD40 ligation (CD40L), inducible T cell costimulator (ICOS) expression as well as via cytokine signals, especially interleukin-(IL)-21, which promotes B-cell differentiation. Stable connection between T FH and B cells is essential to prolong cell–cell contact and the effective delivery of help, which is established by signaling lymphocyte activation molecule associated adaptor protein (SAP) (8, 9). In the GCs, B cells undergo repetitive cycles of somatic hypermutation, clonal proliferation, and selection with the specialized aid from T FH cells to ensure the development of memory B cells and high-affinity antibody-secreting plasma cells (10). Whereas the evolution of GCs provides potent immune protection against foreign antigens, the tight regulation of T FH and B-cell interaction is essential to maintain immunological self-tolerance to prevent autoreactivity (11). Regarding the origin of circulating (c) T FH cells, strong evidence is supporting that T FH cells can be transported to neighboring GCs and migrate to the periphery. Activated T cells with decreased expression of characteristic T FH cell markers appear in the blood, and they have the potential to form memory cells (9). These circulating CD4+ C-X-C chemokine receptor 5 (CXCR5)+ memory T FH cells share functional properties with GC T FH cells and express ICOS and programmed cell death protein 1 (PD-1) molecules, however, in a tempered level. Human blood memory T FH cells form a heterogenous group and include several subsets with different phenotypes and functions according to the presence of ICOS, PD-1, C-C chemokine receptor type 7 (CCR7), CD62L, and chemokine receptors CXCR3, CCR6 (12, 13). The first set of parameters distinguishes CXCR5−CCR7−CD62L− effector memory and CXCR5+CCR7−CD62L− central memory cT FH cells. The expression of the two molecules is probably promoting the migration of the cells to secondary lymphoid tissues. Memory cell types can be further divided into ICOS−PD-1+CCR7int and ICOS−PD-1−CCR7hi quiescent and ICOS+PD-1−CCR7hi+activated memory cT FH cells. The second set of parameters defines CXCR3+CCR6−Th1-like cells (cT FH1), CXCR3−CCR6+Th2-like (cT FH2) and CXCR3+CCR6+Th17-like (cT FH17) cells (14). Concerning their distinct functional roles, cT FH2 and cT FH17 cells are declared as efficient helpers since they induce naïve B cell differentiation into plasma cells and produce immunoglobulin (Ig)A, IgG, and its subclasses. On the other hand, T FH1 cells are not competent naïve B-cell helpers; the only exception is when this subset is in an activated state (ICOS+PD-1+), but their helper capacity is limited to memory B cells (15–18). Recently, a subpopulation of regulatory T cells expressing Foxp3 and CXCR5, called follicular regulatory T cells (T FR), was identified, which have a capability to modify GC responses in several ways: direct regulation of T FH-cell proliferation via the interruption of costimulation or due to the modulation of metabolic pathways in T FH and B cells (19). Consequently, investigations on T FR and B cell interactions in pSS not only can shed light on disease pathogenesis, but can also show the potential use of T FH cells as a biomarker or therapeutic target in autoimmune diseases as an attractive possibility.

In our study, we investigated the disproportion of T FR and cT FH cell subsets and assessed their correlations with disease activity and with various circulating B cell subpopulations in patients with pSS and healthy individuals. Moreover, we evaluated the role of cT FH cells in helping B cells by an in vitro functional assay.
MATERIALS AND METHODS

Subjects
Based on our clinical practice, patients underwent evaluation according to the American-European Consensus Group (AECG), and the diagnosis of pSS was based on it (20). The diagnosis of patients was also revised in agreement with the 2016 American College of Rheumatology (ACR)/European League Against Rheumatism (EULAR) criteria for pSS (21). A total of 38 patients with pSS (37 female and one male; median age: 54 (range 37–74) years) fulfilling both classification criteria were included in the present study. Patients were diagnosed at the Division of Clinical Immunology, University of Debrecen, where they received regular follow-up treatment. The median disease duration was 12 (range 1–25) years. Since immunosuppressive and immunomodulatory drugs fundamentally affect the functions and numbers of the investigated lymphocytes, pSS patients with ongoing immunomodulatory treatments were excluded from the study. Systemic disease activity in pSS was assessed with the EULAR Sjögren’s Syndrome Disease Activity Index (ESSDAI) at the time of the laboratory examinations (22). The mean ESSDAI was 2.44 (+ 0.51, range 1–4), which indicated currently a low disease activity without the necessity of ongoing immunosuppressive drug administration. During the total disease course, 23 patients suffered from systemic manifestations, such as vasculitis, Raynaud’s phenomenon, polyarthralgia, and polyneuropathy, while 15 patients had common glandular involvements. Based on routine laboratory results, 24 patients were positive for anti-Ro/SSA autoantibody (SSA >10 U/ml); moreover, 14 of them showed anti-La/SSB seropositivity as well. Twenty-nine age and sex-matched healthy volunteers (27 female and two male; median age: 46 (36–57) years) served as controls. The characteristics of pSS patients and healthy participants are presented in Table 1. The study participants diagnosed with viral or bacterial infections were excluded. Informed written consent was obtained from all patients. The study has been approved by the Ethics Committee of our University (protocol number: 4879-2017) and the Policy Administration Services of Public Health of the Government Office (protocol number: 1660-4/2018). All data stored and all experiments carried out were in compliance with the Declaration of Helsinki.

Cell Surface Staining and Flow Cytometric Analysis of TFH and B Cell Subsets in Peripheral Blood Mononuclear Cells

Immunophenotyping of lymphocytes was performed by multicolor flow cytometry. Venous blood was collected in vacutainer heparin tubes, and peripheral blood mononuclear cells (PBMCs) were isolated by Ficoll-Histopaque (Sigma-Aldrich, St Louis, MO, USA) density-gradient centrifugation. Harvested cells were washed twice and stained with anti-IgD-fluorescein isothiocyanate (FITC) (clone: IADB6, Beckman Coulter Inc, Fullerton, CA, USA), anti-CD27-phycocerythrin (PE) (clone: 1A4CD27, Beckman Coulter), anti-CD19-phycocerythrin–Cyanine dye 5 (PE-Cy5) (clone: J3-119, Beckman Coulter), anti-CD38-FITC (clone: H7T2, BioLegend, San Diego, CA, USA), anti-CD244-PE (clone: ML5, BioLegend), anti-CXCR5-Alexa Fluor 488 (clone: RF8B2, BD Biosciences, San Diego, CA, USA), anti-ICOS-PE (clone: DX29, BD Biosciences), anti-CD44-PE-Cy5 (clone: RPA-T4, BioLegend) monoclonal antibodies. Cell staining was carried out for 20 min at 4°C in the dark, then washed twice, and prepared for measurements. In case of B cells, at least 10,000 CD19+ events were analyzed, while TFH cell data were collected from at least 75,000 CD4+ events per sample within the whole lymphocyte population.

As described by Morita et al., three TFH subsets could be identified according to the expression of CXCR3 and CCR6 chemokines (15). Based on this study, Th1-like TFH1 (CXCR3+CCR6−), TFH1/17 (CXCR3+CCR6+), Th2-like TFH2 (CXCR3−CCR6−) or Th17-like TFH17 (CXCR3+CCR6+) cells were defined in CD4+CXCR5+ blood T cells with the help of anti-CXCR3-PE (clone: G025H7, BioLegend) and anti-CCR6-PerCP-Cy5.5 (clone: G034E3, BioLegend) monoclonal antibodies. For the determination of TFH cells we used anti-CD127-PE (clone: R34.34) and anti-CD25-PE-Cy5 (clone: B1.49.9) (both from Beckman Coulter) monoclonal antibodies besides CD4 and CXCR5. Fluorescence Minus One (FMO) controls were used to determine gate settings in multicolor panels. The flow cytometric examination was performed with a FACS Calibur instrument (Becton Dickinson, Franklin Lakes, NJ, USA). Data were analyzed using FlowJo Software (Treestar, Ashland, OR, USA). Fig. 1A demonstrates the gating strategy and the division of TFH cell subpopulations. Cell viability was assessed in a number of fresh samples using 7-Aminoactinomycin D (7-AAD) staining. The percentages of non-viable cells were 3.69 ± 0.73 in total PBMCs, 0.13 ± 0.10 in CD4+ lymphocytes, and 0.03 ± 0.02 in case of CD19+ lymphocytes (Supplementary Figure 1C).

Table 1. Demographic characteristic of patients with pSS and healthy controls.

|                   | pSS (n = 38) | Control (n = 29) |
|-------------------|-------------|-----------------|
| Age, years (range) | 54 (37–74)  | 46 (36–57)      |
| Female sex, n (%)  | 37 (97.4)   | 27 (93.1)       |
| Systemic involvement, n (%) | 23 (60.5) | NA              |
| ESSDAI, mean [SD (min-max)] | 2.44 ± 0.51 [1–4] | 2.71 ± 0.98 [1.2–5.5] |
| Anti-Ro/SSA positivity, n (%) | 24 (63.2) | NA              |
| Anti-Ro/SSA titer, U/ml | 100.0 (99.7–100.0) | 99.0 (98.5–100.5) |
| Anti-La/SSB positivity, n (%) | 14 (36.8) | NA              |
| Anti-La/SSB titer, U/ml | 89.9 (86.2–98.4) | 95.5 (92.0–101.4) |
| RF, n (%)          | 29 (76.3)   | NA              |
| RF titer, IU/ml    | 24 (16.8–71.5) | NA              |
| IC titer, extinction | 81.5 (56.0–150.9) | 82.0 (73.5–160.4) |
| IgG titer, g/L     | 16.2 ± 8.4  | NA              |
| IgA titer, g/L     | 2.80 ± 1.56 | NA              |
| IgM titer, g/L     | 1.33 ± 0.91 | NA              |
| Lymphocyte count, ×10^9/L | 1.52 (1.23–1.84) | 1.86 (1.39–2.41) |

Data are displayed as mean ± SD or median (25th–75th percentiles). NA, no data available; pSS, primary Sjögren’s syndrome; ESSDAI, EULAR Sjögren’s Syndrome Disease Activity Index; RF, rheumatoid factor; IC, immune complex; Ig, immunoglobulin.
Cell Separation, Cell Culture and Functional Analysis

B cells were magnetically isolated from the PBMCs using CD19+ B cell isolation kit (Miltenyi Biotec; Bergisch Gladbach, Germany). CD4+ T cells were enriched from PBMCs using CD4+ T cell isolation kit, then CXCR5+ and CXCR5− cell populations were separated within these cells with the application of CD185 (CXCR5)-biotin antibody and anti-biotin microbead (all from Miltenyi Biotec) according to the manufacturer’s instructions (healthy controls, n = 7; pSS patients, n = 5). Cell purity was assessed by flow cytometry: it was greater than 98% for B cells, while the purity was above 80% for CD4+ T cells. B cells were pre-activated by cross-linking of B-cell receptor (BCR) with 2.5 µg/ml affinity purified F(ab′)2 fragment of goat anti-human IgG + IgM (H + L) (Jackson ImmunoResearch, Baltimore, PA, USA) in the presence of recombinant human IL-2 and IL-10 proteins (both 50 ng/ml) (PeproTech, Cranbury, NJ, USA) for 24 h at 37°C in 5% CO2 milieu in Roswell Park Memorial Institute (RPMI) 1640 medium supplemented with 10% heat-inactivated fetal bovine serum in 96-well flat-bottom plates. For in vitro co-culture assay, 5 × 10^4 autologous B cells were cultured with 5 × 10^4 CD4+CXCR5+ T cells or 5 × 10^4 CD4+CXCR5− cTFH cell in the presence of staphylococcal enterotoxin B (SEB) (Sigma-Aldrich; Steinheim, Germany) (1 µg/ml) in 96-well U-bottom plates. Blocking of T–B cell interaction was performed using 3 µg/ml anti-human IL-21 neutralizing antibody (clone MT216G/21.3m, MabTech AB) and 5 µg/ml anti-human CD40/TNFRSF5 (clone 82102; R&D Systems, MN, USA). At day 7, supernatants were collected and stored at −70°C until further measurements, while cells were harvested then stained with the combination of anti-CD38-FITC, anti-CD27-PE, anti-CD4-APC monoclonal antibodies and 7-AAD viability solution. Plasmablasts were assessed using FACS Calibur flow cytometer and further analyzed with FlowJo software.

Enzyme Linked Immunosorbent Assay

IgG and IgM concentrations were measured in the supernatant of the co-culture by enzyme-linked immunosorbent assay (ELISA) kits according to the manufacturer’s protocols (MabTech AB, Nacka Strand, Sweden).

Assessment of Routine Laboratory Parameters

All patients were followed up on a routine basis, and their records contain detailed information on symptoms, clinical conditions, laboratory, and other findings of each visit. Routine laboratory measurements were carried out at the Department of Laboratory Medicine, Faculty of Medicine, University of Debrecen. Blood cell counts including total lymphocyte counts were analyzed with ADVIA 2120i hematology system (Siemens, Munich, Germany). Autoantibody levels were determined by enzyme-linked immunosorbent assay (ELISA) with AUTOSTAT II kits (Hycor Bio-medical, Indianapolis, IN, USA), according to the manufacturer’s instructions (anti-Ro/SSA; normal: ≤10 U/ml and anti-La/SSB; normal: ≤10 U/ml). Serum Ig concentrations were analyzed by turbidimetric method (Dialab GmbH, Wiener Neudorf, Austria). Rheumatoid factor (RF; normal: ≤20 IU/ml) was measured by quantitative nephelometry (Dialab). Circulating immune complexes (ICs) were detected by polyethylene glycol precipitation method. Laboratory reference range was an extinction of 0 to 170 for IC. The results were given according to the analytical measurement range stated in the manufacturers’ instructions.

Statistical Analysis

Data and statistical analyses were performed with GraphPad Prism 8 software (GraphPad Software, San Diego, USA). Descriptive data was represented in box plots of interquartile range (IQR) with a line in the middle as median and “+” sign as the mean value. Tukey whiskers show the 25th percentile minus 1.5IQR and 75th percentile plus 1.5IQR. Functional assay data displayed in scattered dot plots and bars with mean ± standard deviation. Kolmogorov–Smirnov and Shapiro–Wilks normality tests were used to assess normal distribution. In cases of normal distribution, one-way analysis of variance (ANOVA) with Tukey’s post hoc test for pairwise multiple comparisons was carried out. When data sets deviated from normality, Kruskal–Wallis test followed by Dunn’s post hoc test for pairwise multiple comparison was used. Two-way ANOVA with posttest Tukey’s test or Bonferroni’s test for multiple comparisons was used to assess data in in vitro experiments. Statistics and visualization of the correlation matrices were done with the R 3.4.0 software, using the Hmisc and Corrplot packages (23–25). Spearman’s correlation analysis was used for the correlations between the percentages of cell subsets and serological parameters, and R-values were obtained using Corrplot function. Differences were considered statistically significant at p ≤ 0.05.

RESULTS

Changes in the Proportions of Circulating T_{FH} Cell Subsets and T_{FR} Cells in Patients With pSS

We previously described elevated percentages of circulating T_{FH} cells in pSS (26). In order to confirm our former observation, we determined cTFH cell frequencies in a significantly larger patient cohort with well characterized B-cell abnormalities. Additionally, we further characterized blood T_{FH} cell subsets in these patients. Lymphocytes were gated based on a forward and side scatter plot followed by a CD4+ cell gate. The percentages of T_{FH} cell subsets, including T_{FH1}, T_{FH2}, T_{FH17}, and T_{FH1/17} were quantified within the CD4+CXCR5+ lymphocytes of peripheral blood (Figure 1A). Several studies have shown that cTFH cells belong to memory CD45RA+CD4+ T cells (15). To confirm the credibility of our protocol, we evaluated the relative fluorescence expression of CD45RA on circulating CD4+CXCR5+ as well as CD4+CXCR5+PD-1+ T cells and reinforced that both subsets barely expressed CD45RA compared to CXCR5− T cells; thus they resembled memory cells (Supplementary Figure 2B). To assess whether cTFH subsets were altered in pSS, CXCR3+CCR6− T_{FH1},
CXCR3⁺CCR6⁻ TFH1/17, CXCR3⁻CCR6⁺ TFH2, and CXCR3⁻CCR6⁺ TFH17 cells were identified and quantified. There was no detectable difference in percentages of each TFH subsets in pSS patients in comparison to controls or in either designated group within pSS (Figures 1B–E). Interestingly, the entire group of pSS patients (p = 0.0248) and patients with anti-Ro/SSA positivity (p = 0.0047) showed a significantly increased frequency of activated ICOS⁺PD-1⁺ cTFH cells, compared to healthy individuals (Figure 1F).

Blood TFR cells were identified as CD127⁻CD25bright cells within the CD4⁺CXCR5⁺ T cell population (Figure 1G).

**FIGURE 1** | The distribution of cTFH subsets and TFR cells in patients with pSS and their association with laboratory serological markers. PBMCs were isolated from 38 pSS patients and 29 healthy controls (HC) then were stained with fluorochrome-labeled antibodies as described previously. Peripheral circulating (c)TFH subsets were quantified as their percentage within CD4⁺CXCR5⁺ lymphocyte population. (A) Representative dot plots show the gating strategy and the division of TFH cell subpopulations. Percentages of cTFH1 (B), cTFH1/17 (C), cTFH2 (D), cTFH17 (E) in pSS patients (n = 38), pSS without anti-SSA(−) (n = 14), pSS with anti-SSA(+) (n = 24) and HCs (n = 29). (F) Representative dot plots show the gating of ICOS⁺PD-1⁺ activated cTFH cells in pSS and HC. Percentages of activated cTFH cells. (G) Representative dot plots display the gating of TFR cells in pSS and HC. Frequencies of TFR cells. (H) Correlation analysis of the percentages of TFH cell subsets and TFR cells with different serological parameters in patients with pSS. One-way ANOVA with Tukey’s multiple comparisons test (B, D, E) or Kruskal–Wallis test with Dunn’s multiple comparisons test (C, F, G) was used. Correlation analysis was performed using Spearman’s test (H). Box plots represent the interquartile range (IQR) with a line in the middle as median and ‘+’ sign as the mean value. Statistically significant differences are indicated by *p < 0.05; **p < 0.01.
Although the percentages of T_{FR} cells in the whole population of pSS patients were similar to that of healthy controls, when patients were divided into two subgroups based on the presence or absence of autoantibodies, we found that the proportion of T_{FR} cells in patients having anti-Ro/SSA antibodies was significantly higher than those measured in autoantibody-negative patients (p = 0.0363, Figure 1G). Regarding the absolute number of circulating T_{FR} cells, pSS patients had significantly decreased values compared to healthy individuals. This observation presumably stems from the fact that pSS patients have lower lymphocyte counts, which tends to bias the comparative analysis of absolute number of T cell subsets between pSS patients and healthy individuals (27, 28). Nevertheless, among patients, a significant elevation in absolute numbers of T_{FR} cell was detected in anti-Ro/SSA positive individuals, compared to autoantibody-negative patients (p = 0.0157) (Supplementary Figure 3F).

**Altered Circulating T_{FH} Subset and T_{FR} Cell Frequencies Are Associated With Serological Findings**

We investigated whether the alterations in activated cT_{FH} cells as well as T_{FH} cell subsets were associated with the disease etiology and other serological findings in pSS. Interestingly, the percentages of activated cT_{FH} cells showed a significant positive correlation with the levels of anti-La/SSB autoantibody (R = 0.3504; p = 0.0310) and with serum IgA titer (R = 0.3530; p = 0.0297) (Figure 1H). In addition, we found that the proportion of T_{FH}1 (R = 0.3368; p = 0.0387) and T_{FH}1/17 cells (R = 0.3734; p = 0.0209), but not of T_{FH}2 cells, was significantly positively correlated with the levels of ICs (Figure 1H). The frequency of T_{FH}1 cells also showed a significant positive correlation with the levels of serum IgG (R = 0.3977; p = 0.0134) and anti-La/SSB autoantibody (R = 0.3948; p = 0.0142) (Figure 1H). Interestingly, significant positive associations were found between the ratio of T_{FR} cells and levels of anti-La/SSB autoantibody (R = 0.5158, p = 0.0140; Figure 1H). Associations between the percentages of T_{FR} cells and other T_{FH} cell subsets are shown in the supplement, but it is worth mentioning that there was a positive correlation between the frequency of T_{FR} cells and activated T_{FH}1 cells (R = 0.5054, p = 0.0164) (Supplementary Figure 4A).

**Altered Distribution of Peripheral B Cell Subpopulations Is a Characteristic Feature of pSS**

In parallel with the assessment of circulating T_{FH} subsets in pSS, we measured the proportions of different B cell subpopulations, in order to get a better view on T_{FH}–B cell axis in pSS pathogenesis. Lymphocytes were first gated on their characteristic scatter pattern and then gated on the CD19^{+} population. Memory B cell subsets and naïve or mature-naïve B cells were identified according to IgD, CD27, CD38, and CD24 expression (Figure 2A). Proportions of CD19^{+}IgD^{-}CD27^{+} naïve B cells were significantly increased in pSS patients compared to healthy individuals (p = 0.0009, Figure 2B). As expected, patients with anti-Ro/SSA seropositivity had significantly higher percentages than controls (p < 0.0001, Figure 2B). On the other hand, the frequency of CD19^{+}IgD^{-}CD27^{-} un-switched memory B cells was significantly diminished in the total pSS population (p = 0.0010, Figure 2C), while the difference was more pronounced in patients with anti-Ro/SSA antibodies (p < 0.0001, Figure 2C) compared to controls. Interestingly, the proportion of un-switched memory B cells in autoantibody-negative patients was similar to control results (Figure 2C). Peripheral CD19^{+}IgD^{-}CD27^{-} switched memory B cell percentages closely followed the above described tendency, since they were significantly reduced in the whole pSS patient group compared to controls (p = 0.0016, Figure 2D). Moreover, the anti-Ro/SSA positive group showed a significant diminished ratio compared to healthy controls (p = 0.0007, Figure 2D), as well. The percentages of CD19^{+}CD38^{+}CD24^{hi}CD27^{+} transitional B cells were significantly elevated in the total patient group (p = 0.0029), along with anti-Ro/SSA antibody positive patients (p = 0.0002) (Figure 3E) compared to control values. On the other hand, we found that the proportion of transitional B cells had a tendency to be increased in patients having autoantibodies compared to the autoantibody-negative patient population (p = 0.0662, Figure 2E). The ratio of CD19^{+}CD38^{int}CD24^{int} mature-naïve B-cell subset followed a similar tendency as naïve B cells (HC vs. Total: p = 0.0064; HC vs. aSSA+: p = 0.0038, Figure 2F). The percentages of CD19^{+}CD38^{+}CD24^{hi}CD27^{+} primary memory B cells were significantly diminished in the total pSS patients (p = 0.0005) as well as in the anti-Ro/SSA subgroup (p < 0.0001) (Figure 2G) in comparison with healthy subjects.

Upon assessing the associations between B cell subpopulations and other measured parameters, we identified a significant negative correlation between the levels of serum anti-Ro/SSA and the frequency of switched memory B (R = -0.4047; p = 0.0117), un-switched memory B (R = -0.4046; p = 0.0118) and primarily memory B cells (R = -0.4311; p = 0.0069), while they showed positive correlation with naïve B cell percentages (R = 0.3957; p = 0.0139) (Figure 2H). The presence of serum anti-La/SSB showed a significant negative correlation with switched memory B (R = -0.3232; p = 0.0478) and primarily memory B cells (R = -0.3888; p = 0.0159), whereas their serum concentrations revealed a significant positive association with the frequencies of naïve B (R = 0.3532; p = 0.0296) as well as transitional B cells (R = 0.4641; p = 0.0033) (Figure 2H). The levels of IgG and IgA showed a similar correlation as autoantibodies, since they correlated negatively with the proportion of primarily memory B cells (IgG: R = -0.3258; p = 0.0459, IgA: R = -0.3233; p = 0.0477), while IgG also displayed a significant positive association with the percentages of naïve B cells (R = 0.3490; p = 0.0318) and a significant negative correlation with the frequency of un-switched memory B (R = -0.3570; p = 0.0278) (Figure 2H). Interestingly, we found a positive association between the percentages of transitional B cells and serum levels of IgG (R = 0.5861; p = 0.0001) and IgA (R = 0.5882; p = 0.0001) (Figure 2H). Levels of serum IgA also showed a significant positive correlation with the percentages of plasmablasts (R = 0.3821; p = 0.0179) while IgM exhibited a positive correlation with primarily memory B cells (R = 0.3237; p = 0.0474) (Figure 2H). Moreover, the presence of serum ICs correlated positively with naïve (R = 0.3794; p = 0.0188) and transitional B cells (R =
0.4049; \( p = 0.0117 \), and negatively with switched memory B cells (\( R = -0.3302; \ p = 0.0429 \)) (Figure 2H).

We also assessed the possible associations between peripheral T_{FH} cell subsets and B cell subpopulations. The frequency of activated cT_{FH} cells within CD4^{+}CXCR5^{+} cells revealed a significant positive correlation with the percentages of transitional B cells (\( R = 0.5520; \ p = 0.0003 \)) and plasmablasts (\( R = 0.3960; \ p = 0.0139 \)), whereas it showed a significant negative correlation with un-switched memory B cell proportions (\( R = -0.3345; \ p = 0.0401 \)) (Figure 2H). Interestingly, only the frequency of T_{FH2} subset showed any association with B cell subsets. A significant positive correlation was found with every memory B cell subpopulation (switched B: \( R = 0.3640; \ p = 0.0246 \), un-switched B: \( R = 0.4301; \ p = 0.0070 \), DN B: \( R = 0.3826; \ p = 0.0177 \), primarily memory B: \( R = 0.3986; \ p = 0.0132 \)), while a significant negative correlation was observed with the frequency of naïve B cells (\( R = -0.4645; \ p = 0.0033 \)) and mature-naïve B cells (\( R = -0.3967; \ p = 0.0137 \)) (Figure 2H). Of note, significant positive associations were found between the proportion of T_{FR} cells and transitional B cells (\( R = 0.5438, \ p = 0.0089 \)) (Figure 2H). In summary, the present results reinforced our previously published observations on the distribution of B cell subsets in pSS (29). Moreover, our current findings on the proportions of certain B cell subsets along with the high percentages of activated cT_{FH} cells and T_{FR} cells, point at the crucial role of T_{FR}-T_{FH}-B cell axis in the development of aberrant humoral immunity in pSS.

**FIGURE 2** | Abnormal distribution of different B cell subsets reflects the pathophysiology of pSS. Peripheral blood B cell subsets were quantified as their percentage in CD19^{+} lymphocyte population. (A) Representative dot plots indicate the gating strategy and determination of different B cell subsets. Percentages of IgD^{-}CD27^{-} naïve B cells (B), IgD^{+}CD27^{+} un-switched memory B cells (C), IgD^{+}CD27^{-} switched memory B cells (D), CD38^{+}CD24^{+}CD27^{-} transitional B cells (E), CD38^{-}CD24^{-} mature-naïve B cells (F), CD38^{+}CD24^{+}DN B cells (G) in pSS patients (n = 38), pSS without anti-SSA(-) (n = 14) and pSS with anti-SSA(+) (n = 24) and healthy controls (HC; n = 29). (H) Frequencies of certain B cell subsets of pSS patients were acquired, and correlation analysis with serological parameters as well as with the percentages of cT_{FH} subsets was carried out. Data analyses were performed using one-way ANOVA with Tukey’s multiple comparisons test or Kruskal–Wallis test with Dunn’s multiple comparisons test. Correlation analysis was carried out with Spearman’s test. Box plots represent the interquartile range (IQR) with a line in the middle as median and “+” sign as the mean value. Statistically significant differences are indicated by * \( p < 0.05 \); ** \( p < 0.01 \); *** \( p < 0.001 \); **** \( p < 0.0001 \).
**TFH Cells Help B Cells in the Peripheral Blood of pSS Patients**

It is known that TFH cells play an important role in B-cell proliferation and high-affinity antibody production by elevated expression of costimulatory signals as well as secreting stimulating cytokines, including IL-21. Our aforementioned results suggested a direct association between activated TFH cells and B cells in the early differentiation state or with antibody production. To further investigate this issue, we cocultured B cells with isolated T helper or circulating TFH cells *in vitro* in the presence of SEB superantigen and investigated the competence of certain T cell subsets to induce B cell differentiation into CD27+CD38hi plasmablasts (Figure 3A).

In order to reinforce the key function of TFH cells in pSS, we investigated the neutralization of T and B-cell interaction in co-culture, blocking antibodies against IL-21, and CD40/TNFRSF5 was added in T–B co-cultures. As expected, cTFH cells supported B-cell differentiation and plasmablast development more efficiently than CD4+CXCR5− T cells in both controls and pSS patients (HC untreated: *p* = 0.0314 and pSS untreated: *p* = 0.0432, Figure 3B), but we failed to detect any difference after neutralization. The results of pSS patients and controls in IgM and IgG production showed similar patterns, but a tendency towards lower production of IgM in healthy individuals after simultaneous treatment with anti-IL-21 and anti-CD40 was evident (*p* = 0.0539, Figure 3C). When we measured the fold

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**FIGURE 3** The effects of anti-human CD40/TNFRSF5 and anti-IL-21 on B-cell differentiation and antibody production. (A) Graphical representation of B-T_H/cTFH co-culture assay. Magnetically isolated CD19+ B cells were pre-activated with Fab′2 fragment in the presence of recombinant IL-2 and IL-10 for 24 h. Isolated CD4+CXCR5+ cTFH or CD4+CXCR5− TH cells were co-cultured for 7 days with autologous B cells under stimulation by SEB and in the presence or absence of anti-human CD40/TNFRSF5 and anti-human IL-21 antibodies. (B) Representative dot plots show the neutralization effect of anti-human CD40 and IL-21 on plasmablast differentiation. Percentages of 7-AAD−CD4−CD27+CD38hi viable plasmablasts were measured by flow cytometry in healthy controls (HC; n = 6) and pSS patients (n = 3). Two-way ANOVA followed by Bonferroni’s posttest was used. Bars represent mean with SD. (C–F) Concentration and fold decrease of IgM and IgG were measured by ELISA in patients with pSS (n = 5) and HCs (n = 7). Bars show the mean with SD and each data point represents an individual subject. Data analysis performed with two-way ANOVA followed by Tukey’s multiple comparison tests. Statistically significant differences are defined as *p* < 0.05.
decrease in the production of IgG and IgM antibodies, patients with pSS appeared to be more affected by both neutralizations than healthy individuals. There was a 31-fold decrease in IgM secretion by the treatment with both CD40 and IL-21 blockade in patients with pSS \( (p = 0.0125, \text{Figure 3D}) \) compared to a 12-fold decrease in healthy subjects. In case of IgG production, we found a 15-fold decrease after anti-CD40 and anti-IL-21 treatment in pSS patients \( (p = 0.0105, \text{Figure 3F}) \) in comparison with a sixfold decrease in controls. These data reinforce the importance of the \( \text{T}_{\text{FH}}-\text{B cell} \) axis in the development of aberrant humoral immunity in pSS.

**DISCUSSION**

Disturbance of B-cell homeostasis and autoantibody production are key hallmarks of pSS \( (30) \). We have previously shown the important role of \( \text{T}_{\text{FH}} \) cells and their IL-21 secretion in autoreactive B cell activation and autoantibody production in pSS \( (26) \). Recently, we also demonstrated decreased percentages of peripheral IL-10\(^+\) regulatory B cells in pSS and confirmed the strong relationship between IL-21 production of blood \( \text{T}_{\text{FH}} \) cells and the characteristic aberrant distribution of circulating B-cell subsets in the disease \( (29) \). In the present study, we confirmed the increased frequency of activated \( \text{cT}_{\text{FH}} \) cells in an independent cohort of pSS patients and that their altered distribution is associated with anti-Ro/SSA seropositivity. Previous studies have examined circulating ICOS\(^+\)PD-1\(^+\) \( \text{T}_{\text{FH}} \) cells in pSS patients and have found a correlation with ESSDAI or with the presence of ectopic lymphoid structures \( (31-33) \). Other studies focused on CCR7\(^+\)PD-1\(^-\) or PD-1\(^-\) \( \text{cT}_{\text{FH}} \) cells and found a positive association with the degree of focal lymphocytic infiltration in the salivary gland and also with ESSDAI \( (34, 35) \). An important difference between our study and former research is that we included only patients with low disease activity without the necessity of ongoing immunosuppressive treatment. We investigated the distribution of recently described subtypes of \( \text{cT}_{\text{FH}} \) cells in a relatively large number of pSS patients. Although the ratio of these cells did not differ significantly in pSS patients compared to healthy controls, the percentages of \( \text{cT}_{\text{FH}1} \) cells tended to increase within the CD4\(^+\)CXCR5\(^+\) \( \text{cT}_{\text{FH}} \) pool. However, the alteration in the distribution of \( \text{T}_{\text{FH}} \) subsets is still a matter of debate. The previously published data of Li et al. described that frequencies of Th17-like cells within CXCR5\(^+\)CD4\(^+\) T cells were significantly higher in pSS patients than in healthy controls \( (36) \).

In contrast, another study showed that the frequency of circulating CXCR5\(^+\) Th17 cells is decreased in pSS patients, as a result of their extensive infiltration towards the site of the inflammation related to the local expression of CCL20 ligand \( (37) \). In support of our data, Kim et al. recently reported similar distributions of \( \text{T}_{\text{FH}} \) cell subsets in pSS compared to healthy subjects \( (34) \). Interestingly, patients with IgG4-related disease (RD) are also in the center of interest concerning \( \text{T}_{\text{FH}} \) research, but the results are more in agreement with the original work by Morita et al. \( (15) \) stating that CXCR5\(^+\) Th2 and Th17 cells are more proficient B cell helpers than the Th1 phenotype. In line with this, Akiyama et al. described that only \( \text{cT}_{\text{FH}2} \) cells induce the differentiation of naive B cells, enhance the production of IgG4, and associate with disease activity in patients with untreated IgG4-RD \( (38) \). Based on the similarity of organ involvement, pSS patients were enrolled as controls in that study, but did not show any difference in \( \text{T}_{\text{FH}} \) subsets compared to healthy individuals \( (38) \). In another investigation, IgG4-RD was also characterized by a shift toward Th2/TFH2 and Th17/TFH17 polarization, but patients with pSS failed to show any abnormality in the \( \text{T}_{\text{FH}} \) subsets \( (39) \). Our data are in support of no specific abnormality within \( \text{T}_{\text{FH}} \) subsets, but rather an enhanced activation status.

Since their discovery, \( \text{cT}_{\text{FH}} \) cell subsets were investigated extensively in several autoimmune diseases, including systemic lupus erythematosus \( (40-44) \), rheumatoid arthritis \( (45-48) \), myasthenia gravis \( (49) \), multiple sclerosis \( (50) \), and IgA vasculitis \( (51) \) with varying results. Considering the contradictions as well as limited amount of data in pSS, we attempted to resolve this discrepancy and clarify the distribution of blood \( \text{T}_{\text{FH}} \) subpopulations. Since the assessment of peripheral B cells is lacking in several previous studies \( (33, 52) \), we measured a wide spectrum of B-cell subsets to assess how their characteristic distribution correlates with the proportions of \( \text{T}_{\text{FH}} \) subsets.

We correlated the expression of activation markers ICOS and PD-1 with the percentages of \( \text{cT}_{\text{FH}1} \) subsets and discovered that mainly \( \text{cT}_{\text{FH}1} \) cells were activated while \( \text{cT}_{\text{FH}17} \) and \( \text{cT}_{\text{FH}2} \) cells were probably in a quiescent state in the peripheral blood of patients with pSS. Correlation analysis with serological markers reinforced our previous findings that activated \( \text{cT}_{\text{FH}} \) cells are in relation with serum IgA and anti-La/SSB but regarding the cell subsets, only \( \text{cT}_{\text{FH}1} \) and partially \( \text{cT}_{\text{FH}17} \) cells seem to play a role in the pathogenesis of pSS. In order to correlate the ratio of \( \text{cT}_{\text{FH}} \) cells and B cells, we measured a wide variety of B-cell subpopulations. Our results on B cells confirmed previous observations \( (29) \); moreover, the present pSS patient group showed a more characteristic imbalance regarding memory B cells and transitional B-cell distributions. Hence, according to our data, significant correlation with B cells was restricted only to activated \( \text{cT}_{\text{FH}} \) and \( \text{cT}_{\text{FH}2} \) cells. Activated \( \text{cT}_{\text{FH}} \) showed negative correlation with un-switched memory B cells and correlated positively with transitional B cells and plasmablasts, while, \( \text{cT}_{\text{FH}2} \) cells correlated positively with memory B cell subtypes and demonstrated negative association with naïve and mature-naïve B cells. Inevitably, these data raise the question of how the counterintuitive decrease in memory B cell and altered \( \text{cT}_{\text{FH}} \) cell frequencies could contribute to humoral autoimmunity in pSS. Central and peripheral tolerance checkpoints are known to be defective in patients with autoimmune diseases, leading to the expansion of transitional and naïve autoreactive B cells in peripheral blood. A recent study has demonstrated that activated naïve B cells could generate highly polyclonal antibody-secreting cells and could be responsible for serum autoantibody repertoire in lupus \( (53) \). From several studies it has become clear that the decreased ratio of peripheral memory B cells may be interpreted by the upregulation of CXCR4 and CXCR3 chemokines which arrange their recruitment into the inflamed tissues of affected organs \( (3, 54-57) \). Ectopic expression of chemokines, including CXCL13, CCL21, and CXCL12 not
only induces the migration and survival of memory B cells and long-lived plasma cells, but supports lymphoid neogenesis in the pSS salivary glands (58). Upregulated expression of CXCL10 and CCL20 in the salivary gland tissue of pSS raises the possibility that an altered T FH subset distribution at the periphery may be a consequence of their migration towards the affected organs (37).

Besides revealing the alterations and associations of the distribution of cTFH cell subsets, the present study also explored the possible importance of T FR cells in pSS. Our study defined peripheral T FR cells as CD127lo/CD25hi within CD4+CXCR5+ cTFH cells and found that their frequency and absolute number were increased in pSS patients with seropositivity for anti-Ro/SSA autoantibody compared to seronegative patients. In addition, we investigated the relationship between T FR cells and the investigated T FH subsets and demonstrated a positive correlation with activated cTFH and cTFH1 cells, whereas a negative correlation was found with cTFH17 cells. Furthermore, the ratio of T FR cells also showed a positive association with the percentages of transitional B cells as well as the serum levels of anti-La/SSB and IgM. Our observations on T FR ratios are in accordance with the recent findings of Kim et al. and Fonseca et al., who also reported such increased percentages of T FR cells in pSS (34, 59). Additionally, when the latter workgroup compared the suppressive capacity of T FR cells and CXCR5+ Treg cells, they found that T FR cells do not have specialized humoral regulatory capacity, but these cells were able to up-regulate CXCR5 upon in vitro activation and migrate towards CXCL13 gradient, therefore circulating T FR cells could be recruited into subsequent GC responses (59). It is assumed, that blood T FR cells form before the full differentiation into follicle resident T FR cells and retain the expression of CD25 and CXCR5, but lack BCL6 expression as well as the humoral suppression capacity of the mature phenotype (60). Therefore, the increase in cTFR ratio in the peripheral blood could indicate an ongoing humoral response and not a measure of suppressor activity (31, 59). Possibly, the elevated IL-21 production during the follicular and GC phase inhibits T FR commitment through the down-regulation of Foxp3 and CD25 (61, 62), thus the elevation of pre-T FR cells and their traffic to the periphery could be a part of an increased counter-regulatory reaction.

Hence, the negative correlation of T FR/T FH17 in the peripheral blood may be not the outcome of a T FR-mediated suppression, but it is rather derived from the imbalance of the differentiation process in secondary lymphoid tissue.

Costimulatory and cytokine-mediated signals are needed to support T cell-dependent B-cell responses (8). Since CD40 ligation is necessary for the survival and differentiation of GC B cells, blockade of CD40L signaling or disruption of CD40L–CD40 interaction could be beneficial for preventing autoantibody production and B cell abnormality (63). The role of this interaction has been implicated by others, as an increased expression of CD40 has been observed in the salivary gland of pSS and in mice (64, 65). IL–21 is a pleiotropic cytokine, mainly secreted by T FH cells, and it signals through IL-21 receptor which is broadly expressed by different types of lymphoid and myeloid cells. This cytokine has a crucial role not only in the development of T FH cells but also in the differentiation of B cells and the generation of humoral immune responses as well (66). Of note, in vitro analyses on T FH mediated B cell responses with blocking CD40L–CD40 interaction and neutralizing IL-21 effects in pSS were previously performed only in animal models (67, 68). To further clarify the role of costimulation in our in vitro assay, anti-human CD40/TNFRSF5 and anti-human IL-21 antibodies were added together or by itself to inhibit T FH-B cell interaction. To examine their functions, CXCR5+ T helper and CXCR5+ cTFH cells were isolated from CD4+ lymphocytes. As expected, blood T FH cells offered more efficient B-cell help than conventional T helper cells, resulting in the generation of more CD19+CD27+CD38hi plasmablasts in the T–B co-culture in both patients and controls. Nevertheless, B cells co-cultured with CD4+CXCR5+ T cells were still able to differentiate into plasmablast. Of note, a recent study defined a population of PD-1hi expressing CD4+CXCR5+ peripheral helper T (T FH) cells that effectively induce the formation of ectopic lymphoid structures by recruiting B cells via CXCL13 secretion and support their survival and maturation by the production of IL-21 (69). There was a clear difference in the expression of factors associated with B-cell help between the PD-1+ and PD-1hi populations within the CD4+CXCR5+ T cells for the benefit of the latter; presumably this subset interfered with our results. Blockade of CD40 decreased the production of both IgM and IgG in the in vitro assay, suggesting a CD40L–CD40 dependent interaction between CD4+ T cells and B cells. Neutralization of human IL-21 bioactivity affected IgG and IgM production similarly, but simultaneous treatment with both substances proved to be more effective. When we analyzed the fold-decrease of Ig production, we were able to demonstrate a difference between patients with pSS and healthy individuals, since the neutralization effect was more pronounced in pSS. The limitation of our study is the use of total CD4+CXCR5+ T and CD19+ B cells in our functional analysis, as the B-cell population consists of an increased proportion of naïve B cells in patients with pSS; Furthermore cTFR cells probably occur within CD4+CXCR5+ T cells. Since Fonseca et al. demonstrated in a functional assay that blood T FR cells suppress T FH cell and naïve B cell proliferation, thus it could have influenced the outcome of the assay in pSS patients compared to controls (59). Other limitations are the inclusion of patients with a broad range of disease duration and small sample size in the functional analysis.

Of note, the blockade of the CD40–CD40L co-stimulatory pathway is a promising therapeutic approach in pSS. A phase II study on subcutaneous or intravenous administration of an inhibitory monoclonal antibody against CD40 (iscalimab/CFZ533) revealed a decrease in circulating CXCL13 levels and B-cell activation (70). Another study demonstrated that there was no significant difference in ESSDAI score between subcutaneous icosalimab and placebo; therefore therapeutic potential for CD40 blockade in pSS should be further investigated (71).

In conclusion, our findings suggest that there may be no detectable imbalance in the distribution of cTFH subsets in the periphery of pSS. However, our study points out that increased ratios of activated cTFH cells and T FR cells are involved in the pathogenesis of pSS by providing help to B cells. Neutralization of CD40 and/or IL-21 reduced antibody production more
effectively in pSS than in healthy controls in vitro; thus, the interruption of T and B-cell collaboration may be a potential therapeutic approach for down-regulating pathological humoral immunity in pSS.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/Supplementary Material. Further inquiries can be directed to the corresponding author.

**ETHICS STATEMENT**

The studies involving human participants were reviewed and approved by the Ethics Committee of our University (protocol number: 4879-2017) and the Policy Administration Services of Public Health of the Government Of Hungary. All data stored and all experiments carried out were in compliance with the Declaration of Helsinki. The patients/participants provided their written informed consent to participate in this study.

**AUTHOR CONTRIBUTIONS**

KS designed and performed the experiments, analyzed data, prepared the figures, and wrote the main manuscript text. GP designed and collected human samples. TT, BN, and PS provided conceptual advices and edited the manuscript. GP designed and financed the study and contributed to the final version of the manuscript. All authors contributed to the article and approved the submitted version.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fimmu.2021.639975/full#supplementary-material
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**Conflict of Interest:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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