Genetic and phenotypic characterization of indolent T-cell lymphoproliferative disorders of the gastrointestinal tract

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ABSTRACT

Indolent T-cell lymphoproliferative disorders of the gastrointestinal tract are rare clonal T-cell diseases that more commonly occur in the intestines and have a protracted clinical course. Different immunophenotypic subsets have been described, but the molecular pathogenesis and cell of origin of these lymphocytic proliferations is poorly understood. Hence, we performed targeted next-generation sequencing and comprehensive immunophenotypic analysis of ten indolent T-cell lymphoproliferative disorders of the gastrointestinal tract, which comprised CD4⁺ (n=4), CD8⁺ (n=4), CD4⁺/CD8⁺ (n=1) and CD4⁻/CD8⁻ (n=1) cases. Genetic alterations, including recurrent mutations and novel rearrangements, were identified in 8/10 (80%) of these lymphoproliferative disorders. The CD4⁺, CD4⁺/CD8⁺, and CD4⁻/CD8⁻ cases harbored frequent alterations of JAK-STAT pathway genes (5/6, 83%); STAT3 mutations (n=3), SOCS1 deletion (n=1) and STAT3-JAK2 rearrangement (n=1), and 4/6 (67%) had concomitant mutations in epigenetic modifier genes (TET2, DNMT3A, KMT2D). Conversely, 2/4 (50%) of the CD8⁺ cases exhibited structural alterations involving the 3' untranslated region of the IL2 gene. Longitudinal genetic analysis revealed stable mutational profiles in 4/5 (80%) cases and acquisition of mutations in one case was a harbinger of disease transformation. The CD4⁺ and CD4⁺/CD8⁺ lymphoproliferative disorders displayed heterogeneous Th1 (T-bet⁺), Th2 (GATA3⁺) or hybrid Th1/Th2 (T-bet⁺/GATA3⁺) profiles, while the majority of CD8⁺ disorders and the CD4⁻/CD8⁻ disease showed a type-2 polarized (GATA3⁺) effector T-cell (Tc2) phenotype. Additionally, CD103 expression was noted in 2/4 CD8⁺ cases. Our findings provide insights into the pathogenetic bases of indolent T-cell lymphoproliferative disorders of the gastrointestinal tract and confirm the heterogeneous nature of these diseases. Detection of shared and distinct genetic alterations of the JAK-STAT pathway in certain immunophenotypic subsets warrants further mechanistic studies to determine whether therapeutic targeting of this signaling cascade is efficacious for a proportion of patients with these recalcitrant diseases.

Introduction

Non-Hodgkin lymphomas frequently occur in the gastrointestinal (GI) tract, with the majority representing B-cell neoplasms.¹⁻³ T-cell lymphomas account for 10–20% of all primary GI lymphomas.¹⁻³ Aggressive lymphomas, including
enteropathy-associated T-cell lymphoma (EATL) and monomorphic epitheliotrophic intestinal T-cell lymphoma (MEITL), are among the more common types of primary GI T-cell lymphomas, which are associated with high morbidity and mortality. In recent years, there has been a growing awareness of indolent T- and natural killer (NK)-cell lymphoproliferative disorders, which can also arise within the GI tract and involve a variety of GI organs. The pathogenesis of indolent NK-cell disorders is unclear and it is not yet known if they constitute neoplastic proliferations of NK cells. Indolent T-cell lymphoproliferative disorders (ITLPD) of the GI tract, which constitute an immunophenotypically diverse group of clonal T-cell diseases, have been better characterized and hence included as provisional entities in the revised 4th edition of the World Health Organization (WHO) classification of lymphoid neoplasms. The clinical, morphological, and immunophenotypic features of ITLPD of the GI tract differ from those of other types of primary GI T-cell lymphomas and their cellular derivation, although not well established, is also considered to be distinct. Overlapping genomic and genetic alterations have been reported in EATL and MEITL. Limited data suggest a different spectrum of genomic aberrations in ITLPD of the GI tract, and until recently, no recurrent genetic abnormality had been identified in these disorders. However, the mutational landscape and molecular pathways underlying the initiation and progression of ITLPD of the GI tract are unknown and the cell of origin of the different immunophenotypic subsets has not been defined. To gain further insights into the biology of these rare diseases, we performed comprehensive immunohistochemical, molecular and targeted next-generation sequencing analyses of a series of ten cases.

Methods

Case selection
The pathology department databases of multiple institutions were searched for primary GI T-cell lymphomas, over a 23-year period (1996-2018), to identify cases fulfilling histopathological and clinical criteria of ITLPD as defined in the revised WHO classification. Clinical data, including therapy and outcomes, were obtained from the treating physicians or electronic medical records. The study was performed in accordance with the principles of the Declaration of Helsinki and protocols approved by the Institutional Review Boards of the participating institutions.

Morphology and immunophenotypic analysis
Hematoxylin and eosin-stained formalin-fixed, paraffin-embedded (FFPE) biopsy sections were reviewed to assess cyto-architectural features. Immunohistochemical staining was performed using a comprehensive panel of antibodies, including those directed against T-cell antigens, lineage-associated transcription factors, immune checkpoint molecules, histone modifications and cytokine signaling molecules (Online Supplementary Methods). The percentage of cells expressing nuclear T-bet and GATA3 was assessed in areas of dense lymphocytic infiltration determined by CD4 and CD8 staining. Cases with >50% cellular staining by both markers were deemed to co-express T-bet and GATA3. For pSTAT3 and pSTAT5, >10% nuclear staining was considered positive. Flow cytometry was performed on cell suspensions prepared from tissue samples (Online Supplementary Methods).

T-cell receptor gene rearrangement analysis
Polymerase chain reaction (PCR) analysis to determine clonal T-cell receptor beta (TRB) and/or gamma (TRG) gene rearrangement was performed using the ‘Biomed-2’ primers on DNA extracted from fresh or FFPE GI biopsies, lymph nodes, peripheral blood, and bone marrow mononuclear cells, as previously described.

Next-generation sequencing
Targeted next-generation sequencing of lesional and matched normal (control) tissue samples was performed using a custom panel of 465 cancer-associated genes, as previously described. Variant calling required a variant allelic fraction of at least 5% and at least ten variant reads. Variants with an allele prevalence >0.01% in gnomAD, those reported as benign or likely benign in ClinVar, and germline variants present in the normal samples or inferred from variant allelic fractions were excluded from the analysis. Non-synonymous variants that were not known driver mutations were analyzed by PolyPhen-2, SIFT, REVEL, and MetaSVM algorithms. Copy number changes were determined based on read depths using fragments per kilobase per million mapped reads normalized to a pool of sex-matched control samples. The Fusion and Chromosomal Translocation Enumeration and Recovery Algorithm (FACTERA) was used to detect structural chromosomal alterations, which were confirmed by PCR using breakpoint-specific primers and Sanger sequencing of the PCR products (Online Supplementary Methods).

Fluorescent in-situ hybridization analysis
Fluorescent in-situ hybridization (FISH) analysis was performed to assess for SETD2 and JAK2 alterations on FFPE tissue sections using custom designed hybridization probes and dual-color break-apart probes (Oxford Gene Technologies Inc, Tarrytown, NY, USA), respectively, as previously described. Hybridization patterns of at least 100 tumor nuclei were reviewed for each probe. Cases were considered to have SETD2 deletion if the percentage of nuclei with SETD2 locus deletion exceeded the cut-off value of 11.2%, and JAK2 rearrangement if the frequency of split-signals exceeded the cut-off value of 5.0%.

Results

Clinical characteristics and patients’ outcomes
Ten patients (male:female = 8:2) with ITLPD of the GI tract were identified at the contributing centers (cases 1, 2, and 4 were reported previously). The clinical features are summarized in Table 1. The median age at diagnosis was 45 years (range, 37-64 years). The ethnicity of eight patients for whom this information was available was: White (n=5), Hispanic (n=2), and Asian (n=1). The most common signs and symptoms were diarrhea (70%), weight loss (60%), and abdominal pain (50%), with durations ranging from 2 to 16 years prior to diagnosis. Two patients lacked GI symptoms, with disease detected incidentally during routine colonoscopy and workup for inguinal lymphadenopathy. One patient had peptic ulcer disease, H. pylori infection and was serologically positive for hepatitis B and C viruses (case 9) and one patient (case 10) had a history of Crohn disease. Eight patients had been previously misdiagnosed as having celiac disease, seronegative and refractory to a gluten-free diet, and/or other types of lymphomas. The endoscopic findings included mucosal nodularity (70%), scalloping (40%), erythema (40%), decreased duodenal folds (50%), and polyps (20%). Common radiographic findings included abdomi-
Table 1. Clinical characteristics of patients with gastrointestinal indolent T-cell lymphoproliferative disorders.

| Case | Age | Sex | Eth | Presenting signs & symptoms | Duration of symptoms prior to diagnosis (years) | Prior conditions | Endoscopic findings | Radiographic findings | Sites of involvement | Ann Arbor stage (at diagnosis) | Treatment | Outcome (cause of death) |
|------|-----|-----|-----|-------------------------------|-----------------------------------------------|-----------------|---------------------|---------------------|---------------------|---------------------------|------------|------------------------|
| 1*   | 53  | M   | W   | Diarrhea, weight loss, night sweats | 16                                             | None            | Celiac disease      | Mucosal nodularity, scalloping, decreased duodenal folds, erythema | Mild mesenteric LAD, mild FDG activity | Duodenum, jejunum, ileum | IEB        | Bud                  | AWD, 9 years |
| 2*   | 50  | F   | W   | Diarrhea, weight loss, abd pain, fatigue | 3                                              | None            | Celiac disease      | Mucosal nodularity, scalloping | SB wall thickening and dilation | Duodenum, ileum, appendix, colon, stomach, BM | IEB        | Pral, Romi, Bud       | AWD, 7 years |
| 3    | 64  | F   | NA  | No GI symptoms | 0                                              | NA              | None                | Sesile polyp in colon | NA                  | Colon                  | NA         | NA                    | NA         |
| 4**  | 37  | M   | W   | Diarrhea, weight loss | 2                                              | None            | Celiac disease      | Mucosal nodularity, scalloping | Normal               | Duodenum, ileum, colon, stomach | IEB        | Bud, Pred, Aza        | D, 11 years (Large cell trans) |
| 5    | 62  | M   | H   | Diarrhea, weight loss | NA                                             | NA              | Celiac disease, EATL | Mucosal nodularity, scalloping, mosaic pattern, increased vascularity, ulcer | SB and LB dilation | Duodenum, jejunum, inguinal LN | IEB        | CP, Dox, VCR, Pred   | D, 1 year (SB perf) |
| 6    | 41  | M   | NA  | No GI symptoms | 0                                              | MG              | None                | Polypoidal ileal lesions | Mesenteric and iliac LAD | Ileum, colon, stomach, inguinal LN, BM | IVE        | None                  | AWD, 1 year |
| 7    | 38  | M   | W   | Diarrhea, abd pain, vomiting | 5                                              | Lyme disease   | EATL                | Mucosal nodularity, decreased duodenal folds, gastric erythema | SB wall thickening, intuss, mesenteric LAD | Duodenum, jejunum, ileum, colon | IE         | None                  | AWD, 21 years |
| 8    | 38  | M   | H   | Diarrhea, weight loss, abd pain | 5                                              | CHD            | MEITL               | Mucosal nodularity, erythema, friability | Mesenteric and retroperitoneal LAD, incr FDG activity | Duodenum, ileum, colon | IEB        | CP, Dox, VCR, Bud, Pred, Aza | AWD, 7 years |
| 9†   | 41  | M   | A   | Abd pain | 3                                               | PUD, H. pylori, viral hep (B & C) | Atyp lymphoid infiltrate, favor MZL | Mucosal nodularity, decreased duodenal folds | Abd LAD, mild FDG activity, splenomegaly | Duodenum, stomach, BM | IE         | IFN, CP, Dox, VCR, Pred, Gem | D, 27 years (Large cell trans) |
| 10   | 49  | M   | W   | Diarrhea, weight loss, abd pain | 5                                              | Crohn disease | Celiac disease, EATL | Flattened SB mucosa, gastric erythema | Mild SB wall thickening and dilation, partial SB obstruction | Duodenum, jejunum | IEB        | CP, Dox, VCR, Pred, Mes, Aza | AWD, 19 years |

A: Asian; abd: abdominal; AGS67E: anti-CD37 monoclonal antibody AGS67E; AWD: alive with disease; Aza: azathioprine; BM: bone marrow; Bud: budesonide; CHD: congenital heart disease; CP: cyclophosphamide; D: dead; Dox: doxorubicin; EATL: enteropathy associated T-cell lymphoma; Eth: ethnicity; Etop: etoposide; F: female; FDG: fluorodeoxyglucose; Gen: gemcitabine; GI: gastrointestinal; H: Hispanic; hep: hepatitis; IFN: interferon; incr: increased; intuss: intussusception; LAD: lymphadenopathy; LB: large bowel; LN: lymph node; M: male; MEITL: monomorphic epitheliotropic intestinal T-cell lymphoma; Mes: mesalamine; MG: monoclonal gammopathy; MZL: marginal zone lymphoma; NA: not available; perf: perforation; PUD: peptic ulcer disease; Pral: pralatrexate; Pred: prednisone; Romi: romidepsin; SB: small bowel; trans: transformation; VCR: vincristine; W: White. *Previously published cases. Findings prior to large cell transformation. †Bone marrow involvement was detected by cytogenetic analysis; there was no morphological or immunophenotypic evidence of disease and TCRβ polyclonal chain reaction showed polyclonal products. §Biopsies diagnosed as Crohn disease were not reviewed by authors.
nine (66%) patients are alive with persistent disease and three (33%) have died; one (case 5) due to septicemia and multorgan failure following chemotherapy-induced intestinal perforation 1 year after diagnosis and two (cases 4 and 9) due to disease transformation 11 and 27 years after diagnosis.

**Morphological features**

All cases with involvement of the small intestines displayed a dense diffuse or nodular infiltrate of small-sized lymphocytes in the lamina propria (Figures 1A and 2A), with extension into the submucosa noted in a subset. Villous atrophy was observed in three of the nine cases of ITLPD (cases 2, 4, 9) (Figure 1B), however the villi were expanded (blunted appearance) (Figure 2B) in many cases, and all except one (case 10) showed crypt hyperplasia. The lymphocytes had round, ovoid or mildly irregular nuclei, variable fine or coarse chromatin, indistinct or small nucleoli, and scant or moderate cytoplasm (Figures 1C and 2C). No significant increase in intraepithelial lymphocytes was identified (Figures 1B and 2B), although focal lymphocytic infiltration of the epithelium was present in four of nine cases of ITLPD (cases 1, 2, 4, and 7). Scattered lymphoid aggregates were seen in all except one ITLPD (case 5). Sparse, patchy mucosal infiltrates were noted in the seven cases with gastric and/or colonic involvement. Mitotic figures and apoptotic cells were inconspicuous. No angiocentricity, angiodestruction, ulceration, or necrosis was observed. The histopathological findings of the small intestinal biopsy from one patient with large cell transformation, available for review (case 4), were reported previously.11

**Immunophenotypic features**

The immunophenotypic profiles of all cases are summarized in Table 2. Four of ten (40%) ITLPD were CD4+ (Figure 1D), four (40%) were CD8+ (Figure 2D) and one each (10%) was CD4+/CD8+ (“double-positive”) and CD4/CD8- (“double-negative”). All cases analyzed expressed CD2 (Figure 1E) and CD3 (Figures 1F and 2E). Other T-cell antigens were expressed by the majority (Figure 1G, H); variable downregulation or loss of CD5 and/or CD7 was seen in four of ten cases (2/4 CD4+, 1/4 CD8+, and 1/1 double-negative). All except one CD8+ case and the CD4+/CD8- case displayed a cytotoxic immunophenotype, with TIA-1 expression (Figure 2F) noted in three of four cases and variable granzyme B expression (Figure 2G) observed in two of four CD8+ cases.
cases and in the CD4+/CD8+ case. CD103 expression was detected in two of four CD8+ cases (Figure 2H), with one also showing partial CD56 expression (case 8) (Figure 2I). The CD4+/CD8+ and the CD4-/CD8- cases expressed PD-1. CD20 highlighted mucosal lymphoid follicles, but the neoplastic cells were CD20- in all ITLPD. Surface TCRαβ expression was observed in all cases evaluated by flow cytometry and none expressed TCRγδ. All analyzed cases were negative for BCL6, CD10, FoxP3, MATK, PD-L1 or CD30, however CD30 expression (and acquisition of cytotoxic proteins) was observed, and previously reported, upon large cell transformation (case 4). The Ki-67 proliferation index was low (<5%) in all ITLPD evaluated (Figures 1J and 2J).

**Determination of the cell of origin**

Since a good correlation between the transcriptional profiles and immunohistochemistry for T-bet and GATA3 has been reported in T-cell lymphomas, we assessed T-bet and GATA3 expression by immunohistochemistry to determine the cell of origin of ITLPD (Table 2, Online Supplementary Table S1, Online Supplementary Figure S1). The CD4+ cases showed heterogeneity with regards to T-bet and GATA3 expression: one case each was T-bet+ and GATA3+, suggesting T-helper type 1 (Th1) and type 2 (Th2) lineage, respectively and two cases showed T-bet and GATA3 co-expression - hybrid Th1/Th2 profile (Figure 1K, L). The CD4+/CD8+ ITLPD also co-expressed T-bet and GATA3. The CD4+/CD8- case and three of the four (75%) CD8+ cases were GATA3+, implying a type-2-polarized effector T-cell (Tc2) phenotype and one CD8+ case showed T-bet and GATA3 co-expression (Figure 2K, L). Sequential analysis of one CD4+ ITLPD (case 2) showed a shift from a Th1/Th2 (T-bet and GATA3 co-expression) to Th2 (GATA3) phenotype over the course of disease. Double staining for T-bet and GATA3, performed in a subset (cases 2, 7, and 8), confirmed distinct T-bet and GATA3 as well as T-bet and GATA3 co-expressing lymphocytes (data not shown).

**T-cell receptor gene rearrangement analysis**

Clonal TRB and/or TRG rearrangement products were detected in all ITLPD. In patients in whom longitudinal testing was performed, similar sized peaks were observed in all samples, confirming persistence of the same lymphocytic clone.

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**Figure 2. Morphological and immunophenotypic features of CD8+ indolent T-cell lymphoproliferative disorders of the gastrointestinal tract.** (A) An ileal biopsy (case 8) shows a dense mucosal lymphocytic infiltrate expanding the lamina propria and widening the villi; no villous atrophy is present but the crypts are hyperplastic. (B) Small clusters of lymphocytes are seen within the villus epithelium along the lateral edges. There is no increase in intraepithelial lymphocytes. (C) The lymphocytes are small and have round or oval nuclei, condensed chromatin, indistinct nucleoli, and scant to moderate clear or pale pink cytoplasm. The lymphocytes express (D) CD8 and (E) CD3. Most of the cells express the cytotoxic marker (F) TIA1 and (G) granzyme B is expressed by a subset. (H) The lymphocytes are CD103+ and a subset expresses (I) CD56. (J) The Ki-67 proliferation index is low (<5%). The majority of cells express (K) GATA3, but 60% also show (L) T-bet expression.
Table 2. Immunophenotypic characteristics of gastrointestinal indolent T-cell lymphoproliferative disorders.

| Case | CD4  | CD8  | CD2  | CD3  | CD5  | TIA-1 | GrzB | Perf | CD103 | CD56 | BCL6 | CD10 | PD-1 | PD-L1 | FoxP3 | MATK | TCRαβ | TCRγδ | CD30 | K67 (%) | T-bet | GATA3 |
|------|------|------|------|------|------|-------|------|------|-------|------|------|------|------|------|-------|------|-------|-------|-------|--------|-------|-------|
| 1*   | +    | -    | +    | +    | +/-  | -     | -    | -    | -     | -    | -    | +    | -    | -    | <5    | +    | -     |       |       |        |       |       |
| 2*   | +    | -    | NA   | +    | +    | -     | -    | -    | NA    | NA   | -    | -    | -    | -    |       |       | -     |       |       |        |       |       |
| 3    | +    | -    | NA   | +    | +    | -     | -    | -    | NA    | NA   | -    | -    | -    | -    |       |       | -     |       |       |        |       |       |
| 4**  | +    | -    | +    | +    | +/-  | -     | -    | -    | -     | -    | -    | +    | -    | -    |       |       | +     |       |       |        |       |       |
| 5    | +    | +    | +    | +    | +/-  | -     | -    | -    | +     | -    | +    | -    | -    | -    |       |       |       |       |       |        |       |       |
| 6    | -    | -    | +    | +    | +    | +/-  | NA   | -    | -     | NA   | -    | NA   | +    | -    |       |       |       |       |       |        |       |       |
| 7    | -    | +    | +    | +    | +    | +/-  | -    | +    | +/-   | -    | -    | -    | -    | -    | +     |       | -     |       |       |        |       |       |
| 8**  | -    | +    | +    | +    | +    | +/-  | -    | -    | -     | -    | -    | +    | -    | -    |       |       | +     |       |       |        |       |       |
| 9†   | -    | +    | +    | +    | +/-  | +     | -    | -    | -     | -    | -    | +    | -    | -    |       |       | -     |       |       |        |       |       |
| 10   | -    | +    | +    | +    | +    | +/-  | NA   | +    | -     | NA   | -    | -    |       |       |       |       |       |       |        |       |       |       |
| TOTAL| 5/10 | 5/10 | 9/9 | 10/10| 8/10 | 6/10 | 2/10 | 2/10 | 1/10 | 0/9  | 2/9  | 0/9  | 0/9  | 0/9  | 0/8  | 0/10 | 0/10 | 0/10 | 5/10 | 5/10  |       |       |

CD4+: positive; -: negative; DN: double-negative; DP: double-positive; GrzB: granzyme B; Perf: perforin; TCRαβ: T-cell receptor alpha-beta; TCRγδ: T-cell receptor gamma-delta. *Previously published cases. †Findings at diagnosis prior to large cell transformation.
Next-generation sequencing analysis

Targeted sequencing of 20 ITLPD biopsies from ten patients and seven matched normal samples (cases 1, 2, 4, 7-10) revealed 86 genetic variants, including 29 nonsynonymous single nucleotide variants, one small indel, and six structural variants. The average on-target coverage was 1059x (range 809x - 1639x). Twenty-three of the 36 alterations were predicted to be pathogenic based on the published literature or prediction algorithms; the remaining 13 mutations were classified as variants of uncertain significance (Online Supplementary Table S2).

The genetic alterations and their expected functional consequences are summarized in Table 3. Pathogenic or potentially pathogenic changes were identified in eight of ten (80%) ITLPD. Three of four (75%) CD4+ cases and the CD4+/CD8+ and CD4-/CD8- cases harbored alterations of JAK-STAT signaling pathway genes. STAT3 SH2 domain hotspot mutations (D661Y and S614R) were noted in three cases and one case each had a SOCS1 deletion and a STAT3-JAK2 rearrangement. Of note, conventional cytogenetic analysis had previously revealed a balanced translocation t(9;17)(p24;q21) in the latter case, the breakpoints corresponding to the JAK2 and STAT3 loci, and JAK2 rearrangement was confirmed by FISH analysis. Concomitant mutations in epigenetic modifier genes (TET2, DNMT3A, and KMT2D) were observed in four cases.

Table 3. Genetic alterations in gastrointestinal indolent T-cell lymphoproliferative disorders.

| Case | Phenotype | Time point (years following diagnosis) | Genetic alterations | Predicted functional consequence |
|------|-----------|--------------------------------------|--------------------|----------------------------------|
| 1    | CD4+      | 2.5                                  | STAT3 (c.1981G>T, p. D661Y) | Activation of JAK-STAT pathway |
|      |           |                                      | TET2 (c.2457T>G, p. Y819*)  | Altered DNA methylation         |
| 2    | CD4+      | 0                                    | STAT3 (c.1981G>T, p. D661Y) | Activation of JAK-STAT pathway |
|      |           |                                      | TET2 (c.2457T>G, p. Y819*)  | Altered DNA methylation         |
| 3    | CD4+      | 0                                    | SOCS1 deletion         | Activation of JAK-STAT pathway  |
| 4    | CD4+      | 0.5                                  | KMT2D (c.1108G>del, p.L369Gfs) | Altered histone modification    |
|      |           | 7.4                                  | KMT2D (c.1108G>del, p.L369Gfs) | Altered RNA processing and decay |
|      |           |                                      | DIS3 (c.1115T>C, p.L372P)  | Altered histone modification    |
|      |           |                                      |                  | Altered RNA processing and decay |
| 5    | CD4+/CD8+ | 0                                    | STAT3 (c.1842G>C, p.S614R) | Activation of JAK-STAT pathway |
|      |           |                                      | DNMT3A (c.2116G>A, p.G706W) | Altered DNA methylation         |
|      |           |                                      | CDKN2A (c.2080G>A, p.I694F) | Altered DNA damage              |
| 6    | CD4+/CD8+ | 0                                    | STAT3 (c.1842G>C, p.S614R) | Activation of JAK-STAT pathway |
|      |           |                                      | KMT2D (c.9415G>C, p.P3139A) | Altered RNA processing and decay |
|      |           |                                      |                  | Altered histone modification    |
| 7    | CD8+      | 0                                    | IL2-RHOH rearrangement | Unknown                         |
|      |           |                                      |                  | Unknown                         |
| 8    | CD8+      | 0                                    | IL2 3' UTR deletion, IL2-TNIP3 rearrangement | Unknown |
|      |           |                                      |                  | Cell cycle dysregulation        |
|      |           |                                      |                  | Unknown                         |
|      |           |                                      |                  | Cell cycle dysregulation        |
| 9    | CD8+      | 14                                   | None identified    | NA                              |
| 10   | CD8+      | 10.8                                 | None identified    | NA                              |

NA: not applicable. †Large cell transformation. ‡Confirmed by breakpoint-specific polymerase chain reaction and Sanger sequencing.
cases. A missense mutation in the cell cycle regulatory gene $CDKN2A$ and a nonsense mutation in $TNFAIP3$ were detected in one case each.

Two of the CD8$^+$ ITLPD exhibited structural chromosome alterations involving the interleukin-2 (IL2) gene. One case demonstrated an IL2-RHOH (Ras homolog family member H) rearrangement, representing an inversion of chromosome 4, with breakpoints occurring in the 3' untranslated region (3' UTR) of both IL2 (chr4:123372363, c.*44) (Figure 3A) and RHOH (chr4:40246032, c.*449) genes. This rearrangement did not affect the coding sequence, but resulted in the deletion of a portion of the 3' UTR of IL2, including five of the six AU-rich regulatory elements (ARE, AUUUA). The “reciprocal” RHOH-IL2 rearrangement had breakpoints in the 3' UTR of RHOH (chr4:40246006, c.*424) and intron 3 of IL2 (chr4:123373085, c.352-67). Another CD8$^+$ case demonstrated a 1.2 Mb deletion on chromosome 4q, beginning 5 base pairs downstream of the IL2 stop codon (chr4:123372908, c.*5) (Figure 3D) and ending 6 kilobases upstream of the TNFAIP3 interacting protein 3 (TNIP3) gene (chr4:122154953), deleting all regulatory elements from the IL2 3' UTR. In addition, an inversion, with breakpoints in exon 4 of IL2 (chr4:123372912, c.457) and intron 2 of TNIP3 (chr4:122128556, c.89-9014) was identified (Figure 3D). A missense mutation in the minichromosome maintenance complex component 5 (MCM5) gene was also identified in this case. The chromosome breakpoints were confirmed in all ITLPD samples with structural IL2 alterations via PCR amplification and Sanger sequencing.

Figure 3. Structural chromosome alterations of the IL2 gene in CD8$^+$ indolent T-cell lymphoproliferative disorders. In case 7, (A) two chromosome breaks were detected as a consequence of a rearrangement involving the 3' untranslated region (UTR) of IL2 and 3' UTR of RHOH (“IL2-RHOH”) and a reciprocal rearrangement involving intron 3 of IL2 and the 3' UTR of RHOH (“RHOH-IL2”). (B) Pile-up of a subset of reads mapping to the IL2-RHOH rearrangement. (C) Sanger sequencing validation of the fusion breakpoints. In case 8, (D) two chromosome breaks were observed due to a 1.2 Mb deletion spanning the majority of the 3' UTR of IL2 and a portion of the intergenic region between IL2 and TNIP3 (“IL2 3' UTR del”) and an inversion involving exon 4 of IL2 and intron 2 of TNIP3 (“IL2-TNIP3”). (E) Pile-up of a subset of reads mapping to the IL2 3' UTR deletion. (F) Sanger sequencing validation of the deletion breakpoints. *Chromosome position based on assembly GRCh37.p13.
No pathogenic mutations or structural abnormalities were observed in two CD8+ ITLPD (cases 9 and 10), although a variant of uncertain significance was observed in one case (Online Supplementary Table S2).

Longitudinal analysis of five ITLPD (cases 1, 2, 4, 7, 8) revealed stable mutational profiles in four ITLPD. Accrual of mutations over time was noted in one CD4+ ITLPD (case 4). Only a KMT2D frameshift mutation was detected in the first biopsy, obtained shortly after diagnosis. Additional mutations were identified at later time points, including a missense TP53 mutation prior to disease transformation. Of interest, biopsies at the first, second, and fourth time-points had shown different chromosome copy number changes (reported previously),11 but none of the altered regions corresponded to the loci of mutated genes.

Evaluation of the SETD2-H3K36me3 axis

No SETD2 mutations were observed by next-generation sequencing analysis and FISH did not detect any SETD2 deletions in the cases analyzed. Additionally, no loss of SETD2 protein or H3K36me3 was detected by immunohistochemistry and H3K36me2 expression was observed in all analyzed cases (Figure 4A-C, Online Supplementary Table S3).

Evaluation of JAK-STAT pathway activation

Due to the presence of frequent and recurrent genetic alterations targeting the JAK-STAT pathway and IL2 genes, we evaluated pSTAT3-Y705 and pSTAT5-Y694 expression by immunohistochemistry to assess activation of the JAK-STAT signaling pathway. All nine tested cases only showed single scattered or small clusters of nuclear pSTAT3-Y705 and pSTAT5-Y694 positive cells (<10%) in all biopsies (Figure 4E, F, Online Supplementary Table S3).

Discussion

Despite an increasing awareness of ITLPD of the GI tract, deciphering their molecular pathogenesis and cellular origins has been challenging, in part due to the rarity of these disorders. In this study, comprising one of the largest series of cases evaluated, we delineate novel genetic alterations, including recurrent mutations and rearrangements, suggest cellular origins, and expand the immunophenotypic spectrum of these diseases.

The clinical presentations and disease course of our patients were largely congruent with previous descriptions.6,8-16 Of interest, the ITLPD were detected incidentally in two asymptomatic patients, which has rarely been documented.10 A history of Crohn disease has been reported in some patients with CD8+ ITLPD,12,13 which was also the case for one patient in our series. Prior erroneous diagnoses of seronegative, refractory celiac disease in a high proportion (50%) of patients were deemed to be the consequence of misinterpretation of the histopathological changes and incomplete laboratory testing. Due to the relatively recent recognition of these disorders, it is not surprising that 40% of the ITLPD in the current study had been previously misdiagnosed as aggressive intestinal T-cell lymphomas (EATL and MEITL). Extra-GI disease was observed more frequently (40%) in our series than in previously reported series, and transformation to aggressive lymphoma, which is considered rare,8,11,15,28 occurred in two patients, including one with a CD8+ ITLPD. These findings emphasize the need for comprehensive clinical and laboratory evaluation and long-term follow-up of individuals with these disorders.

Next-generation sequencing of the ITLPD revealed genetic alterations in 80% of the cases, including mutations in JAK-STAT signaling pathway genes, observed in

Figure 4. Analysis of the SETD2-H3K36me3 axis and JAK-STAT pathway activation. Immunohistochemical analysis of a CD4+ indolent T-cell lymphoproliferative disorder with STAT3-JAK2 rearrangement (case 2) shows preserved (A) SETD2, (B) H3K36me2, and (C) H3K36me3 protein expression. The lymphocytes express (D) CD4. Only a few scattered (E) pSTAT3-Y705+ and (F) pSTAT5-Y694+ cells are noted (comprising <10% of the neoplastic lymphocytes).
75% of the CD4+ cases and in the CD4+/CD8− and CD4+/CD8+ cases. The STAT3 D661Y and S614R mutations are well-characterized hotspot mutations that impart greater hydrophobicity to the SH2 dimerization surface and promote STAT3 nuclear localization and activation.24 These mutations have been described in a myriad of lymphoid neoplasms and are quite frequent in T-large granular lymphocytic leukemia.25 Perry et al. did not detect STAT3 SH2 domain hotspot mutations in five cases analyzed by Sanger sequencing, although all tested cases were CD8+.12 Deletion of SOCS1, a negative regulator of the JAK family proteins,6 which was seen in a colonic CD4+/ITLPD, is a recurrent abnormality in a variety of T-cell lymphomas and more commonly reported in mycosis fungoides.31 We confirm that STAT3-JAK2 rearrangement is a recurrent event in CD4+ ITLPD although this alteration was only observed in one (25%) of our cases compared to four of five (80%) cases in the series reported by Sharma et al.13

Loss-of-function mutations in epigenetic modifier genes (TET2, DNMT3A, KMT2D) represented the next most commonly altered gene class, identified in 40% of cases and restricted to CD4+, CD4+/CD8−, and CD4+/CD8+ cases. Mutations in epigenetic modifiers, which are believed to be early events in lymphomagenesis9,32,33 and known to cooperate with other mutations in fostering neoplastic transformation,34,35 have also been reported in diverse T-cell malignancies.36,37 However, in contrast to other T-cell lymphomas,19 IDH1/2 mutations were not observed in any ITLPD. Although not recurrent, mutations in CDKN2A and TNAIP3 suggest roles of cell cycle deregulation17 and NF-κB activation38 in the pathogenesis of at least some ITLPD.

Structural chromosome alterations recurrently targeting the 3′ UTR of the IL2 gene, which were identified in 50% of the CD8− ITLPD, have not been described before. The rearrangements and deletions led to the loss of most or all of the regulatory ARE involved in mRNA stability. Studies in mitogen-stimulated Jurkat cells have shown that deletion of these regulatory elements, which act as binding sites for components of the mRNA degradation machinery,9,39 results in a longer half-life of IL2 mRNA.40 Whether these alterations lead to changes in the cellular localization of the IL2 transcript or affect the assembly or composition of protein complexes that modulate activities beyond its 3′ UTR-independent functions has not been investigated. An IL2-TNFRSF17 rearrangement,41 resulting from t(4;16)(q26;p13),41 was previously reported in a CD4+ ITLPD. However, in contrast to our cases, the breakpoints in that case mapped to intron 3 of IL2 and exon 1 of the B-cell maturation antigen (BCMA) gene, also known as tumor necrosis factor receptor superfamily member 17 (TNFRSF17).42 The authors detected chimeric IL2-TNFRSF17 mRNA, but no fusion protein was identified. The functional significance of the prior and current IL2 genetic alterations remains unknown.

Despite the frequent JAK-STAT pathway gene mutations and structural alteration of the IL2 gene, which encodes a key T-cell cytokine that signals via the JAK-STAT pathway,42 none of the ITLPD analyzed showed high-level pSTAT3 or pSTAT5 expression. Our findings are similar to those of Perry et al. who also did not observe significant pSTAT3 expression,12 but contrast with those of Sharma et al. who reported pSTAT5 expression in three of four cases with STAT3-JAK2 rearrangements.43 The reasons for these discrepant findings are unclear. It is plausible that the mutations simply augment the sensitivity of ITLPD to cytokine stimulation, enhancing ligand-mediated signaling as described in other T-cell lymphomas,44 and aberrant proliferations of intraepithelial lymphocytes in refractory celiac disease type 24 that harbor STAT3 mutations.

On analysis of serial samples, acquisition of additional mutations, including those targeting genes involved in the DNA damage response (TP53, POLE) were only identified in an ITLPD that transformed to aggressive lymphoma. It is possible that ineffective DNA repair mechanisms fueled acquisition of additional mutations and complex chromosome changes in this case.31 It is unclear if prolonged azathioprine therapy played a role in genomic evolution. Nonetheless, this and other cases in our series as well as those published previously highlight the futility of genotoxic chemotherapeutic agents for treating ITLPD of the GI tract. The prognostic relevance of periodic genetic analysis needs to be assessed in future larger studies.

Our findings indicate that ITLPD of the GI tract share certain pathogenic mechanisms with other intestinal T-cell lymphomas. As in our cohort, mutations in JAK1-JAK2 pathway genes represent the most frequent alterations in EATL, MEITL, and intestinal T-cell lymphoma, not otherwise specified.17-21 Similarly, loss-of-function mutations in epigenetic modifier genes and DNA damage repair genes have also been reported in aggressive intestinal T-cell lymphomas.15,18 In contrast to EATL and MEITL, however, SETD2 mutations or deletions were not seen in any ITLPD and the burden of pathogenic alterations in ITLPD appears lower.17,18

ITLPD of the GI tract are immunophenotypically heterogeneous diseases. Our study revealed a few unique features that are worth highlighting. In addition to CD4+, CD8−, and CD4+/CD8− ITLPD, we describe a CD4+/CD8+ (double-positive) case. Two ITLPD with a similar phenotype were recently reported from the USA and China.45 Two of our CD8+ ITLPD expressed CD103, which has not been documented before. Prior sporadic cases of CD103+ ITLPD have all been of CD4 T-cell lineage.11,13 These ITLPD could arise from αε integrin-expressing lamina propria T cells,46 but the possibility of activation-induced upregulation of CD103 cannot be excluded.47-49 Of note, one CD103+ CD8+ ITLPD also showed focal CD86 expression. Distinguishing such cases from MEITL can be challenging; however, in addition to the clinical presentation and course, the presence of small lymphocytes with bland cytomorphology confined to the lamina propria, absent MATK expression, and a low Ki-67 index, can help establish a diagnosis of ITLPD. Evaluation of SETD2 and H3K36me3 expression can also aid in differentiating ITLPD from MEITL, which frequently show loss of SETD2 and H3K36 trimethylation.47

ITLPD of the GI tract are thought to originate from mucosal T cells, but the cell of origin of different disease subsets has not been clarified. Absence of FoxP3 and T-follicular helper (THF) cell markers in the current and previously reported CD4+ ITLPD11,13 argues against their derivation from regulatory T cells or TFH cells. Based on expression of T-bet and GATA3, which are transcription factors regulating CD4+ Th1 vs. Th2 cell fate decisions, the CD4+ and CD4+/CD8+ ITLPD in our series displayed Th1, Th2, or hybrid Th1/2 profiles. It is not known whether ITLPD with the latter profile develop directly
from naive T cells into bifunctional mucosal Th1/2 cells, similar to those described in primary immune responses against parasites, which help dampen inflammation, or derive from Th1 or Th2 cells that have undergone cytokine-mediated reprogramming to acquire a Th1/Th2 phenotype, with concomitant production of Th1 and Th2 cytokines. The phenotypic shift from a Th1/Th2 to Th2 profile over time, observed in one CD4+ case, suggests lineage (and possibly functional) plasticity of at least a subset of ITLPD. The majority of the CD8+ cases and the CD4+/CD8+ ITLPD displayed a Tc2 phenotype. Besides orchestrating diverse functions in CD4+ T-helper cells, GATA3 also regulates the activation, homeostasis, and cytolytic activity of CD8+ T cells. The significance of T-bet/GATA3 co-expression in CD8+ ITLPD is unknown. It must be pointed out that despite the reported concordance between the transcriptional and protein expression profiles of T-bet and GATA3 in certain T-cell lymphomas, the definitive lineage (and function) of neoplastic T cells cannot be ascertained based on the expression of single lineage-associated transcription factors. Cytokine profiling and in vitro functional studies are awaited for confirmation of our observations. Contrary to observations in peripheral T-cell lymphoma, not otherwise specified, however, an inferior prognostic impact of GATA3 expression was not apparent in our series of ITLPD.

In conclusion, our study reveals considerable immunophenotypic and genetic heterogeneity of GI ITLPD. We describe recurrent and novel genetic abnormalities in different immunophenotypic subtypes of GI ITLPD which implicate deregulated cytokine signaling and epigenetic alterations in disease pathogenesis. It is hoped that future unbiased interrogation of ITLPD genomes and transcriptomes as well as mechanistic studies will help to clarify the cell of origin and the functional consequences of the underlying genetic aberrations in these rare disorders, opening the door for targeted, less toxic and more effective therapies.

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