Improving outcomes of refractory celiac disease –
current and emerging treatment strategies

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Abstract: Intestinal inflammation and symptoms of celiac disease (CD) usually respond well to gluten withdrawal, but rare cases are refractory to diet. Two types of refractory CD are discriminated on the basis of the presence or absence of an atypical population of mucosal lymphocytes that may progress to enteropathy-associated T-cell lymphoma. Challenges remain in the secure diagnosis of both types of refractory disease, and evidence on which to base treatment recommendations is flawed by the small numbers of reported patients and the use of different diagnostic strategies. Recent advances in our understanding of the mechanisms of the condition in conjunction with the development of immunomodulatory agents for managing other inflammatory diseases are helping to shape future approaches to targeted therapy. Progress will depend on collaboration and recruitment to trials. In the meantime, there is evidence to suggest that earlier diagnosis and better follow-up and management of CD may prevent the development of refractoriness.

Keywords: celiac disease, gluten, small intestine, lymphoma, lymphocytes

Overview
Celiac disease (CD) is a reversible, proximal immune enteropathy resulting from the interaction of dietary gluten with the intestinal immune system.1 Both innate and adaptive immune mechanisms are implicated in the response2 – the latter predominating as evidenced by the dependence on the presence of specific human leukocyte antigen (HLA) class II molecules for gliadin-derived antigen presentation. In keeping with this, dietary exclusion of gluten leads to resolution of the chronic inflammation, restoration of the intestinal mucosal architecture, and resolution of downstream effects of malabsorption.3–5 In a small proportion of cases and for reasons that are poorly understood, intestinal mucosal damage persists despite the removal of the environmental antigenic trigger – hence “celiac disease refractory to gluten withdrawal” or simply “refractory celiac disease” (RCD).6,7 This condition is now recognized as a premalignant overture to enteropathy-associated T-cell lymphoma (EATL), which carries a grave prognosis.8,9

Defining RCD
Understanding any condition relies fundamentally on stringent definition, without which data on prevalence are unreliable, clinical trials of treatment are impossible or uninterpretable, and it becomes difficult to correlate data from different institutions. Unfortunately the application of existing diagnostic criteria in RCD permits significant potential for variable interpretation.
RCD is currently defined as “the recurrence or persistence of malabsorptive symptoms and signs with villous atrophy despite a strict gluten-free diet for >12 months”.

A clinician will face a number of significant difficulties in applying this definition in practice:

1. Only a minority of patients experience the “classical” symptoms of weight loss and diarrhea, and the majority present with a wide array of different symptoms, including fatigue or headache, coincidental findings of various deficiencies on blood tests (such as iron, folic acid, vitamin D, and vitamin B12), and low bone density, or frequently with no reported symptoms at all (for instance, those identified through case finding in first-degree relatives). It may seem incongruous that the diagnosis of RCD should require the presence of malabsorptive signs and symptoms that are unlikely to have been present in the initial active (gluten-induced) lesion – it is possible (and common) to have severe villous atrophy in the absence of symptoms.

2. Malabsorptive symptoms and signs remain uncharacterized in this definition that requires both for the diagnosis of RCD. A common presentation of CD with bloating and irritable bowel syndrome-type symptoms is undoubtedly caused by proximal malabsorption and distal substrate fermentation. Whether this would be a sufficient attribution to define RCD is unclear; however, the coexistence of responsive CD and ongoing irritable bowel syndrome as a cause of persisting symptoms is overwhelmingly more likely. Similarly, diarrhea persisting despite gluten withdrawal may be due to causes that include pancreatic exocrine deficiency or microscopic colitis. Some malabsorptive symptoms and signs are likely to be multifactorial – for instance, associated with iron deficiency – and their presence as a defining feature of RCD may lead to overdiagnosis in a patient with ongoing villous atrophy on intestinal biopsy.

3. A strict gluten-free diet is exceptionally difficult to follow due to the environmental ubiquity of gluten. Furthermore, there are no definitive surrogate markers of ongoing gluten ingestion – we and others have recently shown that persisting IgA antitissue transglutaminase titers are not a reliable indicator of ongoing dietary gluten ingestion, and this has now been adopted in the UK national guidance. Interview of patients by experienced dietitians and the use of food diaries are equally likely to miss sources of gluten ingestion due to recall bias. Patients who are most likely to experience inadvertent gluten exposure are those who are least likely to prepare their own food and are therefore unable to relate the ingredients or precautions taken to prevent contamination during preparation. It is estimated that up to 45% of patients with ongoing symptoms after starting a gluten-free diet have inadvertent gluten exposure in the diet.

4. Sensitivity to dietary gluten may also vary significantly between individuals with CD as a large number of different immunomodulatory genes have been implicated to date in its pathogenesis. It is likely that many patients with ongoing symptoms and villous atrophy 1 year after the initiation of treatment will still have gluten in the diet at a level that leads to toxic effects in that individual rather than a diagnosis of RCD. The reported prevalence of RCD is therefore related to the inverse function of the reliability of the individual or health professionals to be able to exclude sufficient gluten from the diet. It should also be borne in mind that the enforcement of national legislation, involvement of patient support groups, socioeconomic status, and cultural and ethnic dietary factors will impact on successful dietary gluten exclusion and therefore on the persistence of symptoms and villous atrophy at 1 year. This may account for the majority of the large difference between the reported prevalences of “RCD” from different cultures.

5. A degree of villous atrophy is present in ~40% of patients who are rebiopsied at 1 year despite good dietary compliance. As there is an inevitable learning curve with the gluten-free diet, it is not possible to state for certain whether this is due to delayed initiation of adequate gluten exclusion or a fixed length of time required to heal the mucosal lesion. Most studies to date that purport to indicate the length of time required for mucosal healing were not designed to do so. However, while we have found no difference in the healing rates between patients rebiopsied at 1 year and after 1 year, a significant recent study has demonstrated that the likelihood of ongoing villous atrophy at 1 year is related to the severity of the initial mucosal lesion, and this is the first reliable evidence that supports a time to mucosal healing after gluten exclusion of >1 year in certain cases. There is a significant observer variability between histopathologists in the reporting of duodenal biopsies, and therefore, minor degrees of ongoing intestinal inflammation need to be treated with caution and not necessarily considered to represent refractory disease.
6. Many patients with severe malabsorption due to (primarily unresponsive) RCD become clearly apparent long before the 12 months required by this definition due to the severity of the symptoms and the mucosal lesion despite rigorous exclusion of gluten with artificial enteral or parenteral feeds.

7. There is a need to exclude pathologies that may masquerade as RCD, such as olmesartan enteropathy, or chronic norovirus infection associated with immunodeficiency – this is particularly relevant as many RCD patients are seronegative for antitissue transglutaminase or endomyosal antibodies. While the presence of an appropriate HLA is a necessary prerequisite, positivity does not prove the case due to the high background prevalence of HLA-DQ2 or HLA-DQ8 in Caucasian populations.

Therefore, as a result of this working definition, RCD is most likely to be overdiagnosed due to the likelihood of ongoing gluten exposure in the majority of cases and the presence of villous atrophy in a high proportion of normally responsive cases at 1 year. However, the requirement for the presence of unspecified symptoms in the definition (which is not necessary for the diagnosis of CD itself) is likely to underestimate those asymptomatic or paucisymptomatic cases with severe refractory mucosal inflammation (histologically refractory) that may be at risk of late complication.

Classification of RCD
Clarification of the diagnosis of RCD came with the finding of an aberrant population of intestinal intraepithelial lymphocytes (IELs) in some patients with RCD that lack the usual expression of CD3 and CD8 on the cell surface. These cells often share a clonal T-cell receptor (TCR) rearrangement and show immune-phenotypic similarity to the malignant cells in EATL, lending credence to a continuum between RCD and EATL. Patients with an excess of such IELs are described as having type 2 RCD in contrast to type 1 where they are absent (Figure 1 and Table 1). RCD2 carries a far greater risk of transformation to EATL (Figure 2) than RCD1 and a correspondingly poor prognosis and would therefore warrant a more aggressive therapeutic approach if it were shown that this could reduce the risk of malignant transformation or resolution of symptoms. However, such an approach mandates reliable discrimination of type 1 from type 2 RCD, but unfortunately, despite a clearly identifiable cellular phenotype, the diagnosis of RCD2 is not always clear. The presence of clonal TCR rearrangements is problematic as the background IEL population is oligoclonal under normal circumstances, and a clonal rearrangement can predominate in patients who are newly diagnosed or continue to ingest gluten. The presence of clonality can be transient and is not therefore definitive evidence of RCD2.

The presence of immunophenotypically aberrant IELs, regardless of whether or not they are clonal, should indicate RCD as type 2. However, even the immunophenotype of the aberrant IELs is not unique to RCD2 and can occur in normal individuals (Figure 1A) – it is the relative proportion of the surface CD3-CD8+ cells that is key to the diagnosis. How these cells are detected is also critical – immunohistochemistry using dual color immunostaining for CD3 and CD8 helps localization, but is most often carried out singly, leading to very rough approximation of the proportion of CD8-CD3+ cells (immunohistochemistry cannot differentiate between surface and intracellular CD3). Flow cytometry has additional benefits of multiple marker identification on cells but is affected by variable cell yields, the requirement for fresh biopsies, and the lack of localization of the cells isolated (CD4+ lymphocytes predominate in the lamina propria compared to CD8+ cells in the epithelium). Different cutoffs of the proportion of aberrant IELs are used to define RCD type 2 depending on the techniques used to quantify them. Therefore, given the lack of a clear defining test, continual monitoring of aberrant IEL numbers over time in sequential biopsies may be preferable.

Epidemiology of RCD
A further prerequisite for optimizing the management of any condition is to understand its natural history. The main groups reporting RCD have collected cases over many years or even decades, and this reflects the rarity of the condition. However, its reported prevalence ranges from 0.3% to 10% of CD cases – an extraordinary 30-fold discrepancy. This span most likely reflects the loose definition and problems in ruling out gluten exposure in the diagnosis of type 1 RCD as the ratio of RCD1 to RCD2 is higher in those with the highest prevalence of RCD (11:1) compared to those with the lowest prevalence of RCD (3.4:1). However, even when allowing for this, there is still a sevenfold difference in the reported prevalence of RCD2. This may be due to ascertainment bias within tertiary centers; however, it is likely that the use of different diagnostic tests – and in particular the overreliance on TCR clonality rather than flow cytometry – contributes significantly to the variability that inevitably impacts on the reported outcomes of interventions.
Younger patients have been described, but the peak age of onset of RCD is in the sixth decade with a predominance in females – in contrast to EATL that is more frequently identified in males.

The prognosis of RCD2 is poor with a 5-year survival of only 44%–58%, largely accounted for by progression to EATL in 33%–67% of cases. In contrast, while RCD1 can progress to EATL, this is rare and the corresponding 5-year survival is >90%. Transformation to EATL carries a very poor prognosis with the majority dying within 8 months of diagnosis and a 5-year survival of only 11%. Whether RCD1 progresses to RCD2 is contentious – only one study has retrospectively analyzed sequential intestinal biopsies taken prior to the diagnosis of RCD and demonstrated a progressive increase in aberrant IELs averaging 1.8% per month to above the diagnostic threshold for RCD2 in tandem with the development of TCR clonality.

Pathogenesis of RCD

The age at presentation and the finding of de novo RCD or even EATL (without a prior diagnosis of CD) in a significant number of cases implicate chronic antigen exposure, driving the transformation from inflammation to neoplasia. RCD1 is clinically and histologically indistinguishable from active CD and could represent an increased sensitivity to background gluten or a delayed response to gluten withdrawal. This would be in keeping with the findings of resolution of RCD1 over time with the use of strict antigen exclusion (elemental diet), careful and sustained dietary follow-up, or immunosuppressive agents. No genetic markers of unre-
Table 1  Characteristics, outcomes, and possible future discriminatory tests in CD and RCD

|                          | CD                                                                 | Histologically RCD                                                                 | RCD1                                                                                           | RCD2                                                                                           |
|--------------------------|--------------------------------------------------------------------|-------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------------------|
| Response to gluten       | Symptoms usually respond rapidly (2–8 weeks)                      | Symptoms respond as for CD                                                          | By definition, symptomatic                                                                     | As for RCD1                                                                                      |
| withdrawal               |                                                                    | Ongoing villous atrophy recovers 60%–70% within 1 year                              | Symptoms may respond initially and return at a later stage without obvious dietary indiscretion |                                                                                                  |
|                          |                                                                    | Villous atrophy present at symptom presentation or representation                 |                                                   |                                                                                                |
| Intestinal TCR b/g       | Identical clone may persist transiently after diagnosis           | TCR clonality may be present transiently or persist or develop over time            | Transient clonality may occur as with CD                                                        | Clonality usually present but may be below limit of detection (10%)                             |
| or g clonality           |                                                                    | As for CD (usually symptomatic when aberrant lymphocytes present)                   |                                                   | Surface CD3 CD8+ cells express intracellular CD3 >20% by flow cytometry                         |
| IEL phenotype            | Increased gd-TCR+, reduced CD3–CD7+ may persist after treatment    | Surface CD3-CD8+ cells express intracellular CD3 <20% by flow cytometry             | As for CD                                                                                        | CD3+CD8+ IELs >40% by immunostaining                                                           |
|                          |                                                                    | CD3-CD8+ IELs <40% by immunostaining                                               |                                                   |                                                                                                |
| Future test              | As for RCD1                                                         | Cytokine assays or unique antibody profiles may differentiate from responsive CD   | Better definition of lymphocyte subsets on flow cytometry may improve targeted therapy         |                                                                                                |
| possibilities            |                                                                    | Urine and stool gliadin peptide assays may identify dietary indiscretion           |                                                   |                                                                                                |
| Prognosis                | Excellent                                                          | 93% 5-year survival                                                                 | 44%–58% 5-year survival                                                                         | 33%–67% 5-year progression to EATL                                                            |
|                          |                                                                    | 14% 3-year progression to EATL                                                      |                                                   |                                                                                                |

Note: The requirement for symptoms in the definition of RCD results in the entity of “histologically RCD” without symptoms or signs of malabsorption that may go undetected unless routine follow-up biopsies are performed after initiating diet.

Abbreviations: CD, celiac disease; RCD, refractory celiac disease; TCR, T-cell receptor; IEL, intraepithelial lymphocyte; EATL, enteropathy-associated T-cell lymphoma.

In contrast, the aberrant IEL phenotype in RCD2 appears to be expanded in vitro and in vivo by the mucosal expression of interleukin (IL)-15, which is stimulated by innate immune pathways, including interferon-a, a nonimmunogenic gliadin epitope, and viral RNA pattern recognition via TLR3. The aberrant IELs demonstrate direct cell-mediated cytotoxicity against enterocytes, which is activated by the interaction of a stress-derived major histocompatibility complex class I-like molecule (MICA) on the surface of enterocytes with NKG2D expressed on the IELs. This direct epithelial cytotoxicity may account for the severity of the mucosal lesion in RCD2. Cytogenetic alterations have been identified in RCD2 IELs similar to those in EATL, including partial trisomy.
of 1q, which is common in other forms of peripheral T-cell lymphoma. An interesting cytogenetic finding in EATL that has not to date been replicated in RCD2 IELs is a chromosomal gain at q93–34, a region that encompasses both Notch (required for T-cell differentiation of CD127-IELs in the abovementioned study) and the oncogene ABL. 

**Prevention of RCD**

Despite our incomplete understanding of the etiology of RCD, it is likely that chronic antigenic exposure is the driving force. In most western populations, there is a seroprevalence of −1%, but only a small proportion of these patients with CD are diagnosed. Indeed, EATL or osteoporosis often arises in patients previously not known to have the condition. It is possible that early detection and good subsequent management of CD could lead to a reduction in the later development of such complications, including RCD. Such data could be gleaned from comparative studies of different health care economies where case finding and management vary significantly – for instance in Finland, there is a high detection rate of CD and a very low reported incidence of RCD in comparison to the USA with a high reported incidence of RCD and a low detection rate of CD. Quality of adherence to a gluten-free diet in preventing the development of RCD is further supported by the increase in lymphoproliferative disease in patients with incomplete mucosal healing on follow-up biopsies. 

**Treatment of the mucosal lesion in RCD**

Following establishment of the diagnosis, treatment of the patient with RCD requires a careful search for evidence of underlying EATL using cross-sectional imaging, fluorodeoxyglucose-positron emission tomography, enteroscopy, and video capsule studies. Nutritional deficiencies usually require parenteral replacement. Associated osteoporosis should not be overlooked. Treatment is aimed at the mucosal lesion for two reasons: to improve absorption and symptoms and to reduce the risk of progression to EATL.

**Steroids and immunosuppressant agents**

Corticosteroids are used as the initial treatment for both types of RCD. Subjective (symptomatic) improvement is common, but the extent of mucosal response is unclear. Interestingly, corticosteroids were used in the treatment of CD prior to Dicke’s thesis on the association with cereals, and brought about a significant and rapid histological and symptomatic response. The extent of the response in RCD is less clear. An observational study from Paris demonstrated good clinical response (defined as a 50% reduction in symptoms and/or a 50% regain of lost weight prior to treatment); however, histological response was only 40% in RCD1 and 33% in RCD2. Only two of eleven patients treated with steroids alone achieved histological recovery in the Mayo clinic report, and a small study in Argentina reported no benefit from oral steroids.

There is a substantial risk of steroid dependence when used in this setting, in patients already at risk of complications such as osteoporosis, and two strategies have been employed to reduce this risk – topical steroids (modified release budesonide) and purine analog antimetabolites as “steroid-sparing agents”.

While mostly inactivated by first-pass metabolism, modified release budesonide has a systemic bioavailability of 12%, but this may be higher with mucosal disease. Only 30% of the delivered dose is released in the proximal small intestine. Higher doses may therefore work by the systemic route, and it is possible that in CD, where the systemic bioavailability is unknown, the patient may be exposed to a similar magnitude of long-term risks as with oral prednisolone therapy. Subjective clinical measures have improved in both RCD1 and RCD2 with modified release budesonide but without any sign of histological recovery.

Azathioprine is initially combined with steroids due to the delay in onset in effect (when used in inflammatory bowel disease). Early studies showed promising subjective and histological responses in both RCD1 and RCD2, but a worrying trend to increasing progression to EATL in RCD2. Good responses have been reported in RCD1 but not subsequently in RCD2. Limiting its use to RCD1, the azathioprine metabolite, thioguanine (tioguanine), achieved an 83% clinical response and 78% histological response, but there was still a considerable degree of corticosteroid dependency.

One center reporting exceptionally good responses to azathioprine in RCD2 reported no incidence of EATL in their patients >15 years. This discrepancy demonstrates the difficulty in diagnosing RCD2 by over-reliance on T-cell clonality as these patients were most likely RCD1 and the responses to Azathioprine therefore in keeping with previous reports.

An alternative approach in RCD2 has been to use the adenosine nucleoside analog cladribine. Promising
results have been demonstrated with an 81% response and histological recovery in 47% patients – some with complete mucosal recovery.\textsuperscript{76,77} However, the proportion of aberrant IELs, while reduced by the treatment, still remained within the diagnostic criteria for RCD2; hence, pending longer term follow-up for these patients, they cannot be considered to have been “cured”.

A variety of other immunosuppressant agents are reported to have been used in small numbers, including infliximab,\textsuperscript{88}\textsuperscript{–}82 campath (anti-CD52),\textsuperscript{83,84} methotrexate,\textsuperscript{85} cyclosporin A,\textsuperscript{85} and a trial of immunomodulation with recombinant IL-10.\textsuperscript{86} Patient numbers are too small and outcomes too variable to consider the routine use of these agents. However, the number of case reports of benefit with infliximab coinciding with the recent findings of TNF-\( \alpha \) upregulation in RCD might support a reevaluation of the use of anti-TNF-\( \alpha \) monoclonal therapy in RCD1 following failure of steroids and azathioprine.

**Hemopoeitic stem cell transplantation**

Given that a high proportion of patients with RCD2 progress to EATL (32%–67%),\textsuperscript{36–38} which in turn carries a dire survival of \(-15\%\) at 2 years,\textsuperscript{8,9} it is worth considering the possibility of more radical therapies. Autologous hemopoietic stem cell transplantation (aHSCT) has been used successfully in this setting.\textsuperscript{87} In the largest report to date, 13 patients received autologous marrow precursors following melphalan and fludarabine conditioning.\textsuperscript{88} Three died subsequently – one as a result of EATL and the other two most likely as a result of the procedure itself, giving a procedural mortality rate of 15%. Of the 18 patients initially selected on the basis of nonresponse to cladribine, two failed stem cell harvest, possibly as a result of the prior use of cladribine itself and three developed EATL prior to the possibility of transplantation. Overall, excluding those who developed EATL prior to commencement, the intention to treat survival was ten out of 15 (66%), but this is based on a minimum follow-up of 10 months (median \( >2 \) years). All the survivors at the time of reporting had an excellent World Health Organization performance status. The possibility of a lead-time bias should be borne in mind as two of the patients who underwent successful transplantation had effectively normal intestinal biopsies at the time (Marsh grades 0 and 1 effectively), which makes it unclear when or whether they would have ever progressed to EATL. Against the apparent benefits of aHSCT must be balanced the subsequent risk of secondary malignancy (including acute myeloid leukemia), which may be as high as 5% at 5 years and plateau at 10% after 10 years, and the risk of solid tumors, which appears to continue beyond 20 years.\textsuperscript{89} This is clearly dependent on the type of conditioning regimen used, and with milder treatments (that then risk failure of engrafting), there is a lower risk of complications.

Unfortunately, none of the patients undergoing autologous transplantation obliterated their aberrant IEL population and thus remain at risk of further EATL after transplantation. It is well known that IELs are very resistant to chemotherapy and radiotherapy, and therefore, depletion of these cells will require much more intensive conditioning therapy that will significantly increase the risks in the patient. Patients with RCD2 therefore face an uncertain 50% 5-year survival rate with standard therapies, including cladribine, or an equally uncertain 5-year survival rate of \(<66\%\) following aHSCT, but still with ongoing risks of EATL and additional complications of toxicity of the procedure, including secondary malignancies.

**Potential new therapeutic approaches**

With greater understanding of the molecular mechanisms of the development of RCD, it is likely that newer immunomodulatory treatments being developed for other inflammatory conditions will become available for use in RCD. However, given the small number of patients with this condition, trials will have to be focused on the most appropriate agent in order to avoid incomplete evidence based only on case reports. There is experimental evidence for the use of anti-IL-15 monoclonal antibody treatment,\textsuperscript{90} and data from the use of this agent in patients are awaited. However, recent evidence suggesting that certain aberrant IELs in RCD differ with regard to IL-15 receptor expression might also suggest that any effect would be limited.\textsuperscript{91} Therefore, it is likely that downstream regulation of intracellular signaling pathways in the aberrant IELs will be subjected to suppression in an attempt to change the natural history of the condition. One such agent already in use for rheumatoid arthritis and ulcerative colitis is the JAK3 inhibitor tofacitinib\textsuperscript{92} that inhibits downstream signaling from the common g-chain receptor component of IL-2, IL-4, IL-7, IL-9, IL-15, and IL-21, and it may find value in the setting of RCD2 in combination with other agents.\textsuperscript{93}

**Current and emerging strategies in the management of RCD1**

There is currently no way of knowing whether patients with RCD1 have ongoing gluten ingestion, a high immunological
sensitivity to gluten, or slow response to withdrawal or have already entered a different biological phase of refractoriness. Differential cytokine expression in CD and RCD\textsuperscript{52} and different epitope recognition patterns by anticalreticulin antibodies in CD and RCD\textsuperscript{94} hint, however, at a true biological entity, and novel diagnostic tests may become available as a result. Newer methods of detecting gluten ingestion – in stool\textsuperscript{95} or \(\alpha\) measures, a trial of anti-TNF-\(\alpha\) optimize the dosage. If the diagnosis is not in doubt and the 6-thioguanine and shunt pathway metabolites in order to prompt a reevaluation of the diagnosis of RCD\textsuperscript{1}, and fail-

In keeping with other conditions where chronic inflammation and antigen exposure lead to the development of lymphoproliferative disease,\textsuperscript{97} all patients with RCD must ensure strict dietary gluten exclusion, even beyond normal measures. Given the current lack of defining characteristics for RCD\textsuperscript{1}, patients should undergo rigorous reappraisal of the original diagnosis and a search for alternative causes of symptoms or histological damage.

Management of RCD\textsuperscript{1} is largely symptom driven with an emphasis on surveillance in case of development of characteristics defining features of RCD\textsuperscript{2}. Treatment should be initiated with oral prednisolone at 0.5–1 mg/kg/d with bone disease prophylaxis as necessary. Following an initial symptomatic response with steroids, azathioprine can be initiated at 2–2.5 mg/kg/d. Intestinal rebiopsy should be performed after 3 months on azathioprine. Failure to respond symptomatically should once again prompt a reevaluation of the diagnosis of RCD\textsuperscript{1}, and failure to respond to azathioprine warrants measurement of 6-thioguanine and shunt pathway metabolites in order to optimize the dosage. If the diagnosis is not in doubt and the patient remains symptomatic despite the abovementioned measures, a trial of anti-TNF-\(\alpha\) monoclonal therapy could be considered.

Patients who respond to treatment should undergo annual endoscopy and biopsy with quantification of aberrant IELs by flow cytometry or detailed immunohistochemistry and TCR clonality studies. Clinicians could consider withdrawing azathioprine after 2–3 years of complete response in order to confirm the diagnosis of RCD\textsuperscript{1} rather than a slow response to gluten withdrawal (Figure 3).

**Current and emerging strategies in the management of RCD\textsuperscript{2}**

Identification of an aberrant IEL phenotype is fundamental to the diagnosis of RCD\textsuperscript{2}. However, the mere presence of TCR clonality or an aberrant IEL phenotype is insufficient by itself and requires clinician integration of evidence from symptoms, histology, flow cytometry, and molecular studies of clonality. Patients mislabeled as RCD\textsuperscript{2} are liable to considerable harm as a result of Internet search-induced anxiety or toxic overtreatment.

Management of RCD\textsuperscript{2} is aimed at improving symptoms and removing the aberrant IEL clone – a goal that is at present difficult to achieve. The hope is that directed treatment at an early stage – when the patient is still well – could reduce the devastating risk of subsequent EATL. If this were proven, then potentially toxic therapies would be warranted; however, there is as yet no convincing evidence that the development of EATL can be prevented, and current data suggest that at best it may be delayed.

Following rigorous evaluation for the presence of EATL, initial treatment should be steroid based with subsequent biopsy reevaluation of the diagnosis regardless of the response. Reconfirmation of RCD\textsuperscript{2} – often with an increase in the number of aberrant IELs – warrants treatment intensification. Given the risk of enhancing progression to EATL with azathioprine, this agent should be avoided. Although based on limited experience, intravenous cladribine could be used at this stage at 0.1 mg/kg/d for 5 days.

Patients who continue to experience a symptomatic decline following cladribine therapy should undergo a reevaluation of their diagnosis, and at this stage, there are limited options available. The evidence in support of aHSCT is limited but remains the only available option for patients who are declining rapidly – patients who reach this stage of their disease should be managed in conjunction with a center collating data and preferably as part of a clinical trial. Better understanding of the nature of RCD\textsuperscript{2} as a diffuse EATL may lead to resetting thresholds for the use of high-dose chemotherapy in patients without a solid tumor, in such cases with aHSCT as salvage therapy. At present, there is no role for aHSCT in attempting to eradicate the aberrant IEL population in patients with RCD\textsuperscript{2} who are subjectively well (Figure 3).

**Summary**

The current state of the art in treating RCD depends on reliable diagnosis, and outcomes in this condition will only be improved when there is consensus and the development of better diagnostic tools that can adequately predict prognosis. Given the rarity of the condition, the development of national centers or databases and international collaborations will be required. Better case finding and management of CD (including follow-up biopsies on diet) may lead in the future to even lower rates of RCD.
Figure 3 A flow chart showing suggested pathways for diagnosis and management of RCD.

Notes: Dotted lines and italics show pathways that remain unclear or controversial. Arrow marked with * indicates that RCD can develop secondarily after initial response to gluten withdrawal.

Abbreviations: RCD, refractory celiac disease; CD, celiac disease; IBS, irritable bowel syndrome; PEI, pancreatic exocrine insufficiency; LC, lymphocytic colitis; SIBO, small intestinal bacterial overgrowth; LI, lactose intolerance; IEL, intraepithelial lymphocyte; TCR, T-cell receptor; EATL, enteropathy-associated T-cell lymphoma; TNF-α, tumor necrosis factor-α; Tx, transplantation.
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