activity. While no obvious correlation was seen in Th1, Th2, Th17 and NK cells. Then the results showed there was a statistically significant decrease in the secretion of IL-10 in the BD patients ($P = 0.004$), not for IFN-$
$y, IL-4, IL-17 and IL-6.

To evaluate the effects of IL-2 on lymphocytes in vivo, we examined 39 inpatients who received daily low-dose IL-2 at the dosage of 50 WIU for 5 days. It showed that, besides NK cells, total T cells, B cells, CD4+ T cells, CD8+ T cells, Th1 cells, Th2 cells, and Th17 cells were all increased after IL-2 treatment. But only Treg cells were amplified more dramatically, with the four-fold increase. Accordingly, the ratio of Th17/Treg was decreased significantly in patients with IL-2 treatment, tended to balance and had no difference with healthy individuals. At the same time, we found that the symptom were mitigated obviously and disease activity including ESR and CRP were both decreased distinctly without observed side effects.

**Conclusion:** Absolute decrease of PB Tregs in patients with BD was associated with disease activity which might be the major reason for imbalance of Th17/Tregs. It is speculated that BD is an autoimmune disease triggered by the defect of immunotolerance. More importantly, low-dose IL-2 proposes a selective biological treatment strategy by restoring immune tolerance and promoting rapidly remission.

**References:**


Objectives: We aimed to test the hypothesis on aggregation of increased longevity in families of PMR/GCA patients because the family members of long-lived subjects have a survival advantage.

Methods: After questioning our patients we compared age of death of 358 parents of 179 PMR and GCA patients with corresponding data retrieved from 506 parents of 235 randomly collected age and sex matched controls.

Results: We found the number of nonagenarian (≥90 year old) mothers of PMR/GCA patients significantly higher vs controls. Both nonagenarian parents were found in 6 patients (3.35%) and in none of the controls. Decreased number of nonagenarian fathers of our patients remains unexplained.

Conclusion: Confirming our findings in a wider studies would imply a need of including some genetic or behavioural factors to explain PMR/GCA survival advantage.

References:

Acknowledgments: To professor Maciej Markiewski for revision of the manuscript

Table 1. Number of nonagenarians (≥90 years old) in parents of PMR/GCA patients

<table>
<thead>
<tr>
<th></th>
<th>Parents age</th>
<th>PMR/GCA (N=179)</th>
<th>Controls (N=253)</th>
<th>OR</th>
<th>95%CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mothers</td>
<td>&lt;90</td>
<td>148 (82.6%)</td>
<td>243 (96.05%)</td>
<td>0.24</td>
<td>0.11-0.5</td>
</tr>
<tr>
<td></td>
<td>≥90</td>
<td>31 (17.35%)</td>
<td>10 (3.95%)</td>
<td>2.34</td>
<td>1.11-9.15</td>
</tr>
<tr>
<td>Fathers</td>
<td>&lt;90</td>
<td>161 (89.94%)</td>
<td>203 (80.24%)</td>
<td>0.45</td>
<td>0.24-0.83</td>
</tr>
<tr>
<td></td>
<td>≥90</td>
<td>37 (10.06%)</td>
<td>50 (19.76%)</td>
<td>0.9412</td>
<td></td>
</tr>
<tr>
<td>One of parents</td>
<td>&lt;90</td>
<td>136 (75.8%)</td>
<td>193 (76.28%)</td>
<td>1.02</td>
<td>0.63-1.63</td>
</tr>
<tr>
<td></td>
<td>≥90</td>
<td>43 (24.20%)</td>
<td>60 (23.72%)</td>
<td>0.003</td>
<td></td>
</tr>
<tr>
<td>Both parents</td>
<td>&lt;90</td>
<td>173 (96.65%)</td>
<td>253 (100.00%)</td>
<td>0.003</td>
<td></td>
</tr>
<tr>
<td></td>
<td>≥90</td>
<td>6 (3.35%)</td>
<td>0 (0%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mothers of female patients</td>
<td>&lt;90</td>
<td>105 (84%)</td>
<td>149 (97%)</td>
<td>0.0015</td>
<td>0.22-7.29</td>
</tr>
<tr>
<td></td>
<td>≥90</td>
<td>20 (16%)</td>
<td>4 (3%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mothers of male patients</td>
<td>&lt;90</td>
<td>43 (80)</td>
<td>94 (94%)</td>
<td>0.006</td>
<td>4.01-12.13</td>
</tr>
<tr>
<td></td>
<td>≥90</td>
<td>11 (20%)</td>
<td>6 (6%)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

N - number of all parents

Disclosures of Interests: Marcin Milchert Consultant of: Sanofi, Roche, Marek Brzozko; None declared DOI: 10.1136/annrheumdis-2020-eular.4266

SAT0267 ROLE OF AGGRESSIVE IMMUNOSUPPRESSION ON SUBGLOTTIC STENOSIS IN GRANULOMATOSIS WITH POLYANGIITIS: RETROSPECTIVE ANALYSIS OF A MONOCENTRIC COHORT

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Background: Subglottic stenosis (SGS) is defined as airway narrowing below the vocal cords and is a common and potentially life-threatening manifestation of Granulomatosis with Polyangiitis (GPA), with an estimated prevalence of 16-23% (1). Balloon catheter dilation is effective in GPA-related SBS, but relapses are frequent. Little is known about the role of immunosuppression in this setting.

Objectives: to analyse the clinical characteristics of a monocentric GPA cohort, describe phenotype differences among patients with and without SGS and investigate the role of surgical and medical treatments on relapse risk and general outcome.

Methods: Biopsy-proven patients with SGS were identified by review of medical charts among a cohort of patients with GPA, classified according to the algorithm of the European Medicine Agency (2). The clinical characteristics

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SAT0266 RISK FACTORS FOR INFECTIONOUS COMPLICATIONS FOLLOWING RITUXIMAB TREATMENT – MULTICENTER POLISH EXPERIENCE

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Background: Rituximab (RTX) is a B cell depleting monoclonal antibody with proven efficacy in the treatment of ANCA-associated vasculitides (AAV). The infectious complications occur in 15-25%.

Objectives: We aimed to assess the frequency and risk factors of infections in patients with AAV receiving RTX among Polish patients.

Methods: 7 tertiary referral centers experienced in the treatment of vasculitis completed a questionnaire regarding AAV patients treated with RTX.

Results: Among 49 patients included in the analysis (47 with GPA, 2 with MPA; 36/73% men; mean age at diagnosis 42,45±14,9 yrs., mean age on RTX initiation 46,14±14,7 yrs.), at least one infection occurred in 20 patients (40.82%) after mean time of 16,65±16,01 months since the administration of RTX. Patients were followed for a mean time of 26,88±21,94 months. There were no differences in the incidence of infectious complications by gender, age, BMI, smoking status, severity of the disease, activity of the disease (BVAS), time from diagnosis to RTX initiation, carriage of staphylococcus aureus in the upper respiratory tract, total dose of CYC before RTX treatment. We didn’t observe severe hypogammaglobulinemia or neutropenia after RTX treatment. 40% of the observed infections occurred during the first month, 35% between second and sixth month of follow-up, while 25% were observed between 6 and 12 months after the RTX initiation. Of the 20 patients who developed infection, 12 (24.5%) had further infections. Antibiotic prophylaxis with trimethoprim–sulfamethoxazole was administered in 40 out of 49 (81.63%). Upper respiratory tract infection was the most common infectious complication (n=11), followed by lower respiratory tract (n=4), soft tissues (n=4) and urinary tract infections (n=4), lacrimal gland abscess (n=2) and abdomen (n=1). In cases with a positive microbial result Staphylococcus aureus (n=4), Klebsiella pneumoniae (n=2), Pseudomonas aeruginosa (n=1), Candida (n=1) and others (n=6) were identified. No fatalities were recorded and only 3 patients had severe infection with the necessity of prolonged treatment.

Conclusion: Despite the high number of infections in our group treated with RTX, most of them were not severe. Upper respiratory tract was the most common site of infection.

Disclosure of Interests: None declared DOI: 10.1136/annrheumdis-2020-eular.1607

SAT0265 FAMILIAR AGGREGATION OF LONGEVITY IN GIANT CELL ARTERITIS AND POLYMYALGIA RHEUMATICA

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Background: The long-term mortality in giant cell arteritis (GCA) and polymyalgia rheumatica (PMR) is unexpectedly decreased (1,2,3,4) or at least not increased regardless increased mortality risk factors that these diseases share with other systemic inflammatory disorders.

Objectives: To investigate the role of surgical and medical treatments on relapse risk and general outcome.

Disclosures of Interests: None declared DOI: 10.1136/annrheumdis-2020-eular.1607