Results: One-hundred-eighty-seven patients with EGPA were identified, from which 89 patients with documented asthma assessment at baseline and after 3 years from diagnosis were included. Severe/uncontrolled asthma was observed in 42.7% of patients at diagnosis and was associated with a previous history of respiratory allergy (p<0.01), higher serum total IgE levels (p<0.05), and increased use of high dose inhaled corticosteroids (ICS) (p<0.05) and OCS (p<0.001) for respiratory symptoms the year before the diagnosis of EGPA. Improvement or worsening of asthma during follow-up was experienced by 22.4% patients, with no discrimination of baseline features that allowed their distinction. Severe/uncontrolled asthma was present in 40.5% of patients at 3 years, and was associated with increased airway resistance on pulmonary function testing (p<0.05). Long-term PFT did not improve during long-term follow-up regardless of ICS and OCS therapy (both FEV1/FVC and FEV1, p>0.05). Using multivariate binary logistic regressions, severe rhinosinusitis, pulmonary infiltrates, over-weight and severe/uncontrolled asthma at vasculitis diagnosis independently predicted severe/uncontrolled asthma at the 3-year endpoint (Table below).

<table>
<thead>
<tr>
<th>Variables</th>
<th>Odds Ratio</th>
<th>95% Confidence Interval</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Severe/uncontrolled asthma</td>
<td>10.64</td>
<td>3.43-39.08</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Pulmonary infiltrates</td>
<td>1.16</td>
<td>1.43-21.85</td>
<td>0.011</td>
</tr>
<tr>
<td>Severe rhinosinusitis</td>
<td>3.57</td>
<td>1.07-13.55</td>
<td>0.038</td>
</tr>
<tr>
<td>Overweight (BMI&gt;25 kg/m²)</td>
<td>3.41</td>
<td>1.05-12.90</td>
<td>0.041</td>
</tr>
</tbody>
</table>

Conclusion: In this large cohort of patients with EGPA, long-term severe/uncontrolled asthma is frequent, and patients affected display a higher proportion of bronchial obstruction and have more severe air-way resistance. Overall, pulmonary function does not improve during the follow-up regardless of ICS and OCS therapy. Finally, long-term severe/uncontrolled asthma is associated with baseline pulmonary and ENT manifestations, but not with vasculitic features.

REFERENCES


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SAT0218 PATIENTS’ JOURNEY THROUGH GIANT CELL ARTERITIS: A QUALITATIVE STUDY

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Background: Giant cell arteritis (GCA) is the most common form of vasculitis. Diagnosis is difficult due to multiple presenting symptoms: headache, jaw and limb claudication, myalgia and visual impairment. Access to care may be delayed because often multiple providers are involved in the diagnosis of GCA and disease management. Treatment with high-dose glucocorticoids (GCs) can relieve symptoms and prevent vision loss, but GC-related adverse events are common; GCA often relapses once GCs are tapered.

Objective: To understand the GCA patient care pathway and unmet needs in GCA through in-depth patient interviews.

Methods: US patients with GCA were recruited through outreach to physicians and The Vasculitis Foundation, which used email newsletters and social media to recruit 50% of participating patients. Extensive individual interviews with patients were conducted by qualitative researchers by phone or in person and explored patients’ perspectives and experiences from the onset of GCA symptoms to diagnosis and disease management. Patients were asked open-ended questions and encouraged to share their stories to provide additional insights into the patient journey and their individual perspectives. The qualitative data collected were analyzed using human-centered design methodology, including patient typologies (personas: optimist, fearful, stoic or despondent), forced temporal zoom (journeys), forced semantic zoom (stakeholder system mapping) and affinity mapping for pattern recognition of unmet needs.

Results: A total of 28 patients were interviewed; 23 (82%) were women and mean age was 69 years (Table). The number of patients in each persona category is shown in the Table. Stoic and optimist personas had medium to high levels of self-advocacy and a positive/engaged attitude toward their condition. Patients often ascribed their milder GCA symptoms to causes such as stress and did not consult a physician until they developed moderate to severe symptoms, such as persistent headache, jaw pain or visual disturbances. Patients with existing inflammatory disorders were less likely to share symptoms of GCA unless the symptoms substantially worsened. In most cases, physicians diagnosed GCA based on abnormal erythrocyte sedimentation rate, often followed by a temporal artery biopsy. After diagnosis of GCA, all patients received GCs with little information on the chronicity of GCA. Few patients were offered treatment alternatives to GCs. Overall, patients managed their GCA independently, with moderate support from friends or family, and sought to balance relief of GCA symptoms with the adverse effects of GCs. Patients concentrated on tapering and discontinuing GCs, with less concern about relapse. Furthermore, patients who were most uncomfortable with the adverse effects of GCs often waited until their GCA symptoms became debilitating before telling their physician. Almost all patients reported searching for a support group after the diagnosis.

Conclusion: Patients with GCA experience adverse effects from GCs and remain focused on reducing their GC dose. For those with inflammatory comorbidities, diagnosis of GCA is another burden in a debilitating journey that results in a sense of disempowerment and resignation toward their condition and paucity of therapeutic options. Patients with GCA want a clearer understanding of treatment options and access to support groups. Patients’ attitudes and self-advocacy vary depending on their personas; recognizing these persons may help HCPs coordinate patient care. Increased awareness of GCA among patients and HCPs may accelerate the path to diagnosis and treatment, and emerging therapies may help reduce GC burden.
