

**AstraZeneca – Call For Grant Applications**  
**CGA Rheumatology SLE State Chapter Meetings 2021-01**

<b>Submission Deadline</b>	Rolling review/support decisions, with submission deadline of October 6 <sup>th</sup> at 9 am EST
<b>Primary Area of Focus</b>	Rheumatology
<b>Therapeutic Area</b>	Systemic Lupus Erythematosus (SLE)
<b>Burden to the Patient</b>	<p>Patients with SLE who have been properly diagnosed have many burdens/challenges, including:</p> <ul style="list-style-type: none"> <li>• Daily symptom management to maximize quality-of-life</li> <li>• Distress over timing and severity of next flare</li> <li>• Toxicities associated with conventional immunosuppressants, including glucocorticoids</li> <li>• Advocating for individualized treatments for better symptom management and quality-of-life</li> </ul>
<b>Healthcare Burden</b>	<p>SLE is a chronic, multisystem autoimmune disorder that impacts approximately 204,000 Americans, primarily women of child-bearing age.</p> <p>Rheumatologists face substantial challenges in the diagnosis and management of patients with SLE, owing to the:</p> <ul style="list-style-type: none"> <li>• Diverse symptomatology due to the involvement of different organ systems</li> <li>• Diverse patient populations, with its prevalence being 2-3 times greater among women of color</li> <li>• Complexity of the pathophysiology of SLE and targets being explored in clinical trials</li> <li>• Limited awareness of current clinical trials and results</li> <li>• Limited experience with biologic therapy in the management of patients with moderate-to-severe SLE</li> <li>• Underappreciation of the impact of conventional therapies on the accumulation of organ damage and SLE-related comorbidities</li> </ul>
<b>Educational Program</b>	Consisting of, but not limited to, live in-person or virtual educational sessions at state rheumatology chapter meetings and enduring programming.
<b>Budget</b>	≤ \$250,000
<b>Successful submission</b>	<ul style="list-style-type: none"> <li>• Independently-developed application, providing rationale/basis for and detailed description of the goals, learning objectives, educational strategy, format, and execution and measurement/analysis of program outcomes</li> <li>• Key findings of analyses conducted to determine root cause(s) of identified educational needs and practice gaps (in contrast to a disease state background)</li> <li>• Expertise executing high-quality sessions at state rheumatology chapter meetings</li> <li>• Outcomes from previously executed program</li> <li>• Outcome methodology that will analyze for regional differences</li> </ul>
<b>CGA Code</b>	CGA_Rheumatology_SLE_State Chapter Meetings_ 2021-01
<b>Website URL</b>	<a href="https://www.astrazenecagrants.com/us-grants.html">https://www.astrazenecagrants.com/us-grants.html</a>

### **Program Requirements:**

The Program must be planned and executed as if an accredited activity and fully compliant with the criteria and/or standards of commercial support for ACCME, AAFP, AOA, ACPE, ANCC, AANP, or NCCPA. Furthermore, the program will be educational and non-promotional in nature and will be planned, designed and implemented in accordance with the U.S. Food and Drug Administration's Guidance on Industry-Supported Scientific and Educational Activities ("Policy Statement").

The Policy Statement and the ACCME Standards require, among other things, that (i) Institution conduct the Program independently and without control or influence by AstraZeneca over the Program's planning, content (including the selection of speakers or moderators), or execution; (ii) the Program be free of commercial bias for or against any product; (iii) Institution make meaningful disclosure of AstraZeneca support of the Program and any prior relationship between Institution and AstraZeneca, and the relationship, if any, between AstraZeneca and the speakers selected by Institution; and (iv) AstraZeneca not engage in, and Institution not permit any other sponsor to engage in, promotional activities in or near the Program room or advertise its products in any materials disseminated as part of the Program.

In addition, Institution is required by the Policy Statement and, if applicable, accreditation standards to ensure that any product discussions at the Program be accurate, objective, balanced and scientifically rigorous. This includes a balanced discussion of each product and of treatment alternatives, that limitations on data be disclosed, that unapproved uses be identified as such, and that for live presentations there be opportunities for questioning or debate.