As part of our commitment to improving the lives of people living with rare diseases Alexion, AstraZeneca's Rare Disease supports quality, independent Continuing Medical Education (CME) designed to enhance patient care and health outcomes.

This call for grant applications provides public notice of availability of funds to address areas related to the multidisciplinary care of patients with PNH.

Deadline for	
Submission	
Decision Notification	
Primary Area of	Rare Disease
Focus	
Therapeutic Area	Hematology – Paroxysmal Nocturnal Hemoglobinuria (PNH)
Geographic Focus	Global
CGA Code	AX012
Intended Audience	PNH Physicians and Care Teams
Budget	\$300,000
Educational Need	Given the diversifying landscape of therapies for managing PNH (e.g.,
	biosimilars, proximal inhibitor monotherapies and/or add-on as dual therapy, etc.) adult and pediatric hematologists would benefit from education on the pathophysiology, clinical manifestations, and appropriate treatment algorithms for managing PNH. It is critical for HCPs to understand the underlying pathophysiology and that life threatening consequences of PNH are associated with chronic, uncontrolled terminal complement activation leading to IVH, thrombosis, organ damage and premature mortality. It is a disease that affects RBCs, WBCs, and platelets.
Educational Design and Focus	<ul> <li>Alexion funding is intended to support multi-modal programs (i.e. with live/virtual and/or enduring components) including but not limited to: <ul> <li>Interactive self-directed programs designed for impactful learner engagement using proven distribution channels</li> <li>Symposium (face-to-face or virtual) that will be developed into a virtual enduring program. Slots should be secured by grant recipient.</li> <li>American Society of Hematology 2024</li> </ul> </li> </ul>
Application	Proposal must be independently developed and include the following:
Requirements	<ul> <li>Needs Assessment/Gaps/Barriers: Include a comprehensive, well-referenced needs assessment that provides a detailed description of the educational / practice gaps and barriers of the target audiences. The needs assessment must be independently developed and validated by the educational provider.</li> <li>Audience Generation: Describe methods for reaching the target audience(s) and any unique recruitment methods that will be utilized.</li> <li>Educational Strategy: Provide clearly defined and measurable learning objectives that are clearly designed to address the identified gaps and barriers. The proposal should demonstrate an understanding of instructional design issues as they relate to the gaps in the knowledge, competence, or performance of the targeted audience.</li> <li>Program Evaluation and Outcomes: Provide a description of the outcomes methodology that will be employed to measure the impact of the educational program and how these results will be presented, published, or disseminated. Additionally, describe the methods that will be used to determine the extent to which activity has served to close the identified healthcare gap.</li> </ul>

Programs should include an outcomes plan of at least Moore's level 4.
• <b>Budget:</b> Include a detailed budget with rationale, including breakdown of costs for content per activity, out-of-pocket cost per activity and management cost per activity.
• Accreditation: Programs must be accredited and fully compliant with all ACCME Criteria and Standards for Commercial Support <sup>™</sup> .

**Program Requirements:** The Program must be planned and executed as 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 nonpromotional 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.