

# National Health Council

## Domains & Values:

### PCORI Reauthorization



Domains	Values
<b>Continue Focus on Patient Involvement</b>	<ul style="list-style-type: none"> <li>• Ensure that patients have at least the same level of representation on Board and other governance bodies</li> <li>• Continue requiring patient partnership in research</li> <li>• Continue to hold workshops and engagement events aimed at gathering patient stakeholder priorities and input</li> </ul>
<b>Include Preamble or Report Language Directing Incremental Changes to Enhance Impact and Relevance to Patients</b>	<ul style="list-style-type: none"> <li>• Encourage research that assesses treatment value from the patient and family perspective (e.g., studies that consider benefits and costs beyond clinical outcomes such as direct/indirect costs, impact on families, productivity, travel, etc.)</li> <li>• Increase patient input and transparency in the selection of national priority topics and research questions</li> <li>• Expand funding mechanisms that support patient-driven research (e.g., led by patients and patient groups)</li> <li>• Increase emphasis on building patient capacity to engage in PCOR activities</li> </ul>
<b>Avoid Substantial Changes that Would Negatively Impact Patients or Decrease Likelihood of Reauthorization</b>	<ul style="list-style-type: none"> <li>• Continue ban on conducting cost-effectiveness research (e.g., cost-per QALY)</li> <li>• Continue ban on using research to drive Medicare national coverage decisions</li> </ul>