



Standing For Your Health

BADANAMU SICKLE CELL FOUNDATION

STRATEGIC PLAN FRAMEWORK

Badanamu Sickle Cell Foundation Strategic Plan Development Working Group Charter

Create a strategic plan draft process and utilize such process in engaging key stakeholders to contribute to the strategic plan development of the Badanamu Sickle Cell Foundation

Create and disseminate survey/s or similar strategies to ensure robust inputs are received from the global inherited blood disorders community in regard to the draft skeleton

Utilize the Badanamu Sickle Cell Foundation Vision, Mission and Objectives to develop its Goal areas

Finalize and disseminate the strategic plan to key stakeholders within specified timeline



Set Strategic Context	Solicit Collaborative Input	Define Strategic Priorities	Validate Plan with Membership	Approval Final Strategic plan
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A strong and united sickle cell and other inherited blood disorders patient and health care provider communities equipped to effectively advance Disease-specific and cross-disease priorities that improve outcomes for affected families



Through data-driven advocacy, collaboration, and capacity building, we empower our member organizations and advance cross-disease initiatives to improve outcomes for people living with sickle cell & other inherited blood disorders.



OBJECTIVES

- 1. Advance disease-specific and cross-disease initiatives that meet the immediate needs of people**
living with sickle cell disease & other inherited blood disorders
2. Equip inherited blood disorder patient organizations to shape and advance efforts that ensure the promise of new innovations can be fully realized for all.
3. Act as a medium for capacity building, training, and mentorship, and for translating knowledge and materials, compiling and disseminating information
4. Strengthen and empower sickle cell & other inherited blood patient organizations through the creation of regional networks, especially in areas of the highest unmet needs
5. Support cross-disease patient-centered education initiatives to strengthen care infrastructure, improve training/education of HCPs, and expand HCP access to resources
- 6. Encourage the formation and development of national and regional Inherited blood disorder organizations throughout the world, and assist and stimulate** such organizations through disease- **specific and cross-disease exchange of skills and knowledge.**
- 7. Enable advancement of shared priorities within the inherited blood disorders community through unified and coordinated data-driven advocacy**

Objective

Equip inherited blood disorder patient organizations to shape and advance efforts that ensure the promise of new innovations can be fully realized for all

Survey
Feedback

Advance disease-specific and cross-disease initiatives that meet the immediate needs of people living with sickle cell disease & other inherited blood disorders

Strategic
Priority

- | | |
|---|--|
| <ul style="list-style-type: none">• Enhanced collaboration and partnerships with HCPs• Funding for research, awareness campaigns, and patient support programs | <ul style="list-style-type: none">• Push campaigns to hard-to-reach areas and rural areas• Data sharing to enable connections between diseases• Education and awareness programs |
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Advance disease-specific and cross-disease initiatives in countries where collaboration is feasible.

Identify unmet needs

Create collaboration opportunities

Promote collaborative programs

Objective

Equip inherited blood disorder patient organizations to shape and advance efforts that ensure the promise of new innovations can be fully realized for all

Survey
Feedback

- | | |
|--|---|
| <ul style="list-style-type: none">• Integration of the patient voice• Collaboration and support in the form of funding, expertise, capacity building, or advocacy efforts | <ul style="list-style-type: none">• Coordination and effective, ongoing communication• Training of trainers for disseminating knowledge on inherited blood disorders |
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Strategic
Priority

Align Badanamu Sickle Cell Foundation initiatives with participating patient organizations' missions and activities

Program committees with
wide representation

Programs based on top
priorities

Programme evaluations

Objective

Act as a medium for capacity building, training, and mentorship, and for translating knowledge and materials, compiling and disseminating information

Survey
Feedback

- | | |
|--|--|
| <ul style="list-style-type: none">• Empowering patients and caregivers• Information compilation, knowledge acquisition, dissemination and application | <ul style="list-style-type: none">• Evidence generation and evidence dissemination• Curriculum development and research mentoring |
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Strategic
Priority

Identify trainings and workshops that increase knowledge where there are gaps and unmet needs

Core Competency and specialized trainings

Training sessions based on Surveying and feedback

Data Collection Activities

Objective

Enable advancement of shared priorities within the inherited blood disorder community through and unified and condinated data during advocacy.

Survey
Feedback

- | | |
|---|--|
| <ul style="list-style-type: none">• Important area of focus for any organization interested in blood disorders• Fostering a network of organizations | <ul style="list-style-type: none">• Participate in international conferencess and advocacy efforts• Challenging but highly needed |
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Strategic
Priority

Promotes sharing best practices to assist in the development of inherited blood disorder patient support orgnisations.

Capacity building programs

Support for emerging
orgnisations

Sharing best practices
among disorder
groups



STRATEGIC PRIORITIES

1. Advance disease-specific and cross-disease initiatives in countries where collaboration is feasible
2. Align Badanamu Sickle Cell Foundation initiatives with participating patient organizations' missions and activities
3. Identify trainings and workshops that increase knowledge where there are gaps and unmet needs
4. Develop regional patient networks which will help build capacity in countries with unmet needs
5. Support collaborative advocacy initiatives between patients and HCPs to improve care and treatment infrastructure
6. Promote sharing best practices to assist in the development of inherited blood disorder patient support organizations.