Policy Priorities for Improving FH Care

1. Greater Awareness
   Familial hypercholesterolemia impacts 34 million people, 90% undiagnosed.

2. Continued Advocacy
   Advocacy organizations around the world provide patient support & education.

3. Screening, Testing & Diagnosis
   FH screening, through genetic or clinical testing, should follow country-specific conditions and guidelines.

4. Accessible Treatment
   Patient-centered FH treatment should be accessible, begin in childhood and continue over the course of life.

5. Guidelines for Severe, Homozygous FH
   Severe and rare, homozygous FH needs separate guidelines because patients with this form of FH require unique resources.

6. Family-Based Care
   FH is a lifelong condition, and care should be organized and accessible throughout all life stages.

7. Registries
   Government-funded FH registries can provide valuable country-specific information on awareness, treatment & outcomes.

8. Research
   Robustly funded research can advance the scientific understanding of FH and help identify new treatments.

9. Cost
   FH care must be affordable and provide value to impacted families and society at large.

Learn More
Adapted from Reducing the Clinical and Public Health Burden of Familial Hypercholesterolemia: A Global Call to Action