











Key Objectives



Advocate for political recognition and dedicated policy frameworks for RDs

Campaign for budget allocations to address the unique needs of individuals with RDs.

Review national plans to shorten diagnosis time and improve intervention efficiency.



Ensure availability, affordability, and coverage of RD tests and medications.

Improve coordination of care across fragmented healthcare systems. Address geographical and cultural factors impacting RD care and access to services.



Data Collection and Sharing

Explore strategies for systematic, standardized data collection and sharing.

Utilize technology infrastructure and telemedicine to enhance RD care.



Patient Empowerment and Support

Enhance social and cultural acceptance, equality, and inclusion of individuals with RDs.

Empower patients and families to self-care and advocate for their needs.

Promote prevention, screening, and early intervention in RD management.



Research and Innovation

Discuss incentives for innovation in the RD field.

Promote research advancements and collaboration to improve RD care.







