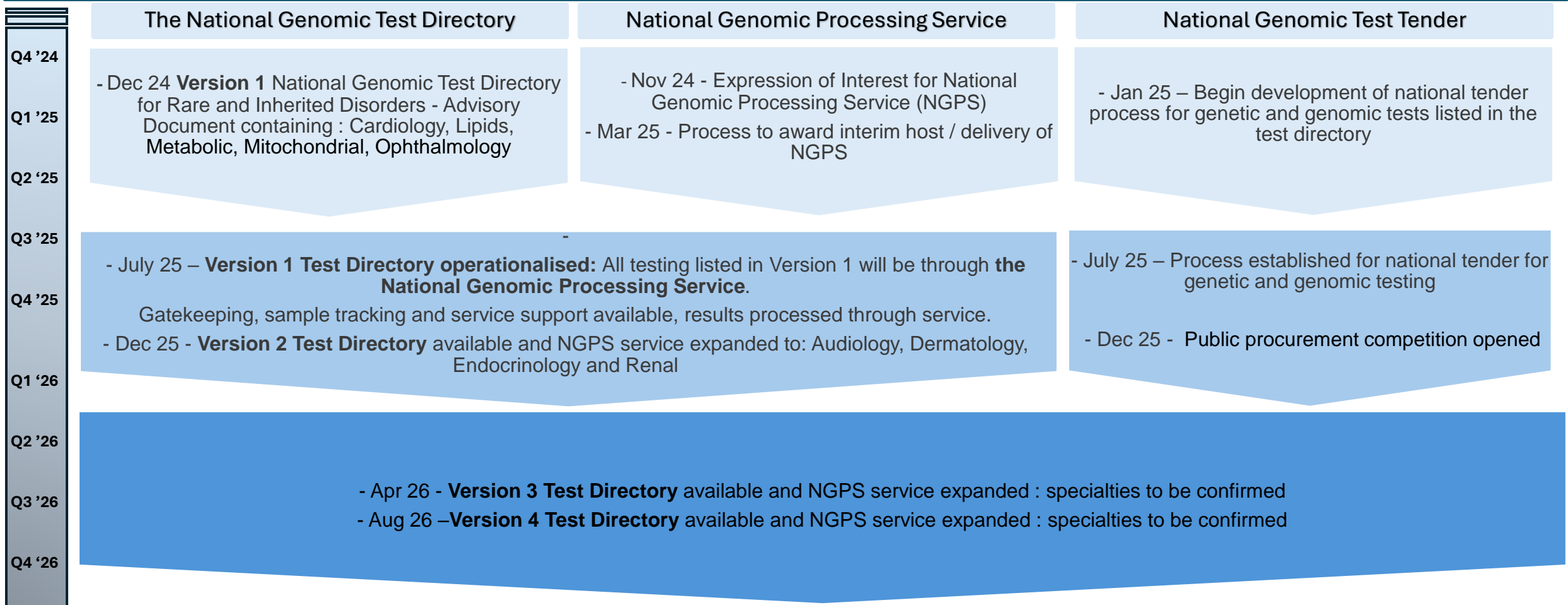


National Genomic Test Directory for Rare and Inherited Disease and National Genomic Processing Service Project Timeline



Q1 '27

Final Outcomes 2027

- ✓ **Version 5 National Genomic Test Directory for Rare and Inherited Disease completed for all parts** : Acutely Unwell Children, Audiology, Cardiology, Clinical Genetics, Dermatology, Developmental Disorders, Endocrinology, Fetal including NIPD, Gastro-hepatology, Haematology, Immunology, Inherited Cancer, Lipids, Metabolic, Mitochondrial, Mosaic & Structural Chromosomal Abnormalities, Multi-Purpose Tests, Musculoskeletal, Neurology, Ophthalmology, Renal, Respiratory, Ultrarare and atypical disorders. All changes to the Test Directory processed through Test Directory review structure.
- ✓ **Fully operational National Genomic Processing Service**: Service provider laboratories as dictated by national tender
- ✓ Compliance of service user and service providers achieved through **national tender**