

Abstract graphic on the left side of the slide, featuring overlapping curved shapes in various shades of blue and light blue.

National Cancer Action Team
Part of the National Cancer Programme

“What does a good
child, teenage and
young adult cancer
service look like?”

Louise Hooker

Purpose of workshop session

- To get delegates input into the process for developing:
 - Clinical Lines of Enquiry to support and enhance Peer Review of CYP services.
 - Metrics for measuring what a ‘good quality’ Children’s service and TYA service would look like.
 - To help develop service profiles, inform service specifications, commissioning and service provision.

Why?

- Access/Equity
- Safety
- Efficacy
- Productivity

To reach a expert consensus on which data will enable:

- Benchmarking of services
- Informed discussion – clinical practice, service provision
- Reviewing/understanding variation
- Commissioning of services

Who are the stakeholders?

- Commissioners – NHS Commissioning Board, SCG Clusters, Primary Care.
- Cancer Networks, CYP Cancer Network Coordinating Groups
- NHS Provider Organisations
- Clinical Teams
- External Bodies – Quality Assurance, regulators and professions
- Patients, families, the general public, taxpayers
- Partners – 3rd Sector, others.

Data from different sources – needs to be

- Relevant, meaningful
- Understandable
- Useable
- Accurate
- Available, collectable
- Collected/presented at correct level to fulfil purpose - MDT, Trust, Primary Care, CN, CYPCN, SCG etc

Which data?

- Reflect stages of patient pathways
- Diagnosis
- Treatment – different modalities
- Follow up care
- Survivorship issues
- Palliative/EOLC
- Non-cancer specific – generic CYP service quality metrics/measures?