National Consent Model Workshop – Programme

Date	**Thursday, 21 April 2016**
Time	**10:00 to 16:00**
Venue	**Lift, 45 White Lion Street, London, N1 9PW**

1000 - 1030	**Arrival, tea & coffee**	
	Welcome & aims for the day	
1030 - 1045	▪ Chair’s opening remarks	**Richard Stephens** use MY data patient advocate, Chair, NCRI Consumer Forum
	Session 1 – The National Data Guardian’s review of data security & consent	
1045 - 1130	▪ Getting the right balance between sharing data and privacy concerns	**Peter Knight** Deputy Director, Head of Research Contracting, Information Intelligence & Stakeholder Engagement, DH
1130 - 1145	**Refreshment break**	
	Session 2 – Cancer registration, analysis & research: how changes to consent will alter the use of patient data	
1145 - 1215	▪ Why the use of cancer data is so important, and why the current model (no consent) is so valuable	**Professor Michel Coleman** London School of Hygiene & Tropical Medicine
1215 - 1300	▪ Cancer registration – how will it change if consent is needed? ▪ Routes to Diagnosis reimagined	**Chris Carrigan** Head, National Cancer Registration and Analysis Service **Sam Johnson** Cancer Information Analyst, National Cancer Registration and Analysis Service
1300 - 1330	**Lunch**	
	Session 3 – Consent around the UK	
1330 - 1430	▪ The devolved nations and England – contrasting approaches to patient consent	**Dr Anna Gavin** Director, N. Ireland Cancer Registry
1430 - 1445	**Refreshment break**	
	Session 4 – Improving the way patients learn about cancer registration	
1445 - 1545	▪ How patients are currently informed ▪ Using patients’ experiences to define how new patients could/should be informed	**Julie Flynn** Strategic Data and Influencing Lead, Macmillan Cancer Support **Table discussions**
	Closing summary	
1545 -1600	▪ Chair’s closing remarks	**Richard Stephens** use MY data patient advocate, Chair, NCRI Consumer Forum

* use MY data