

**Appendix 2 – Informed Consent Form (Version 2.0, 26<sup>th</sup> July 2013)**

Patient Identification Number:

Name of Researcher: **[insert local principal investigator name]**

Title of the study: **Isotoxic Intensity Modulated Radiotherapy (IMRT) in Non-Small Cell Lung Cancer – A Feasibility Study**

**CONSENT FORM**

*Please initial in boxes*

1. I confirm that I have read and understood the information sheet Version 2, dated 26<sup>th</sup> July 2013 for the above study and have had the opportunity to ask questions.
2. I understand that my participation is voluntary and that I am free to withdraw at any time, without giving any reason, without my medical care or legal rights being affected.
3. I understand that sections of any of my medical notes may be looked at by responsible individuals from **[insert institution name]** and its authorised agents, by the sponsor for monitoring and audit or from Regulatory Authorities where it is relevant to my taking part in research. I give permission for these individuals to have access to my records.
4. I also understand that if I withdraw from the study early, the data collected whilst I was on the study will be retained to ensure the trial has been run in accordance with all applicable rules.
5. I understand that I will not benefit financially if this research leads to the development of a new treatment or medical test
6. I understand that my General Practitioner will be informed about my participation in this study.
7. I agree to take part in the above study.

\_\_\_\_\_  
Name of Patient

\_\_\_\_\_  
Date

\_\_\_\_\_  
Signature

\_\_\_\_\_  
Name of Person taking consent  
(If different from researcher)

\_\_\_\_\_  
Date

\_\_\_\_\_  
Signature

\_\_\_\_\_  
Researcher

\_\_\_\_\_  
Date

\_\_\_\_\_  
Signature

1 for patient; 1 for researcher; 1 for general practitioner, 1 to be kept with hospital notes

***Thank you for taking part in this research study***