**ADELE - ADaptive dEtaiLed comparEd to standard imaging follow up after lung cancer surgery**

**SITE SURVEY FORM**

The ADELE trial is planning to be conducted across the UK, and we are currently undertaking a site survey on management of these patients at your site – we would be grateful if you could review the below study description, complete the questionnaire and return to: Gemma Nanson at [g.simpson@liv.ac.uk](mailto:g.simpson@liv.ac.uk) or via fax 0151 794 8930

**Study description**

The study will be undertaken in ~14 investigational sites in the UK. This is an unblinded two-arm parallel randomised design. Patients will be randomised to either adaptive intensive or standard follow up.

|  |  |  |
| --- | --- | --- |
| **Time since surgery** | **Adaptive detailed** | **Standard** |
| 3 month | CT | CXR |
| 6 month | PET/CT | CXR |
| 9 month | CT | CXR |
| 12 month | CT | CXR |
| 18 month | PET/CT | CXR |
| 24 months | CT | CXR |
| 36 months | CT | CXR |
| 48 months | CT | CXR |
| 60 months | CT | CXR |

CT and PET/CT imaging from the adaptive intensive arm will be reported locally for clinical management and centrally for the study. Patients with identified abnormalities on the CT, PET/CT or CXR will be managed according to local policies.

Patients who are receiving adjuvant chemotherapy will be randomised after the completion of the chemotherapy to join the study at the 6 month interval.

|  |  |
| --- | --- |
| **Site details** | |
| **Name:** |  |
| **Site name & NHS Trust:** |  |
| **Email:** |  |
| **Are you interested in participating in this trial?** | Yes  No  Comments: |
| **Are you taking part or planning to take part in any study that would conflict with the recruitment to this trial? If so please state”.** | Yes  No  Comments: |
| **Do you have any conflicts of interest for this study?** | Yes  No  Comments: |
|  | |
| **Recruitment** | |
| **How many patients with non-small lung cancer in your department received surgery during the last calendar year?** | *Please base this on actual figures rather than subjective estimate* |
| **Please review the inclusion and exclusion criteria carefully. How many of the patients detailed above do you think would have been eligible for the ADELE study?** |  |
| **Please indicate your site’s routine standard of care for follow up of patients with non-small cell lung cancer following surgery** | |  |  |  |  | | --- | --- | --- | --- | | Time point | Please tick method | | | | 3 month | Chest X-Ray | CT | PET- CT | | 6 month | Chest X-Ray | CT | PET- CT | | 9 month | Chest X-Ray | CT | PET- CT | | 12 month | Chest X-Ray | CT | PET- CT | | 18 month | Chest X-Ray | CT | PET- CT | | 24 months | Chest X-Ray | CT | PET- CT | | 36 months | Chest X-Ray | CT | PET- CT | | 48 months | Chest X-Ray | CT | PET- CT | | 60 months | Chest X-Ray | CT | PET- CT | |
| **The current 5 year survival rate following surgery for non-small lung cancer is estimated at 56% What increase in the survival rate would persuade you change your current clinical practice for follow up to a more intensive regimen e.g. from 56% to 65%** |  |