

National Lung Cancer Audit Report 2013

Report for the audit period 2012



Prepared in partnership with:



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National Lung Cancer Audit Report 2013

Report for the audit period 2012

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Purpose

The purpose of this document, the ninth Annual Report of the National Lung Cancer Audit, is to summarise the key findings of the audit for patients diagnosed with lung cancer who were first seen in 2012. The history, purpose and methodology of the audit have been extensively documented and further details can be obtained from the HSCIC website (www.hscic.gov.uk/lung).

Based on the comments of service users we have updated this short report highlighting key issues. More extensive analyses on the 2012 data, including case-mix adjusted data in an electronic spreadsheet format will be available from the [HSCIC website](http://www.hscic.gov.uk) in due course.

Every Trust or Health Board in England and Wales and Scotland have participated in the audit, although because of differences in reporting schedules, standards and targets the Scottish data are tabulated separately. Guernsey has also participated in the audit. Unfortunately, the Northern Ireland data was not available in time to be included in this report and therefore will be published electronically at a later date. Details of care provided by individual organisations in this report are based on "place first seen" in secondary care. Place first seen is chosen since in the vast majority of cases it represents the location of the Multi-Disciplinary Team that co-ordinates the investigation and treatment of the individual patient. As a result some tertiary centres may appear to have little input into the care of lung cancer and to submit little data to the audit, however, on the contrary, they usually provide the most complex care for the most difficult patients and submit treatment data on behalf of other Trusts. Information about the number and types of treatment provided by these Trusts is provided in [Figure 10](#).

The data collected in this report is for patients first seen in the calendar year 2012. At this time the Cancer Networks were still in place and responsible for helping to facilitate cancer services in the areas they served therefore the audit has decided to report by Network for this year. Moving forward into 2013 the report will be modified to reflect the new commissioning structures. The NLCA Project Team would like to take this opportunity to thank colleagues who were employed by Cancer Networks for their invaluable support of the audit, often over many years. This has undoubtedly contributed hugely to the progress that the audit has made in improving lung cancer care.

Note that all data presented in this report refers to cases submitted to the National Lung Cancer Audit unless otherwise stated.

Changes to the Report Format

A major change for this year is that the data completeness reports for key items are available in online format only. Data completeness reporting was key in the early years of the audit because achieving a high enough standard of data to allow the reporting of clinical outcomes was the main focus. As the standard of data is so much higher than in recent years, this report can now focus more on clinical outcomes and uses of the data. Data completeness and quality is still key to the ongoing success of the NLCA and we would encourage audit participants to view their data at www.hscic.gov.uk/lung

A further change for this year is that we have excluded mesothelioma from the main report. Information about patients with mesothelioma will be published in PDF format at a later date. Where appropriate, at the bottom of the Key Outcome Tables, headline comparisons with last years data are provided both with mesothelioma cases included and excluded.

The report also, for the first time, includes some focus features about specific areas of cancer care and shows trends in care outcomes over time.

Key Messages

- The audit has collected data on 40,216 patients in Great Britain for this audit period, representing approximately 98 per cent¹ of the expected number of new lung cancer cases. This is thought to represent all cases of lung cancer presenting to secondary care.
- Overall measures of the standards of care have been sustained and have marginally improved compared to those seen last year, with small rises in the proportion of patients having surgery, and anti-cancer treatment. In many cases the measures of treatment now approach those seen in other western healthcare systems. Despite these improvements, there remains marked variation across Trusts and Networks and differences in case-mix do not appear to explain the whole of this variation. Trusts are encouraged to critically appraise their own results and perform reviews of lung cancer pathways and/or clinical cases where treatment rates are below the national average.

Key Outcomes ('Headline Indicators') for the NLCA for 2012 for England and Wales, Scotland and Guernsey)

Note; data for Scotland includes mesothelioma.
 Note; data for Guernsey for 2008 and 2009 was not recorded.
 Note; Scotland did not report on CT scan prior to bronchoscopy for 2008

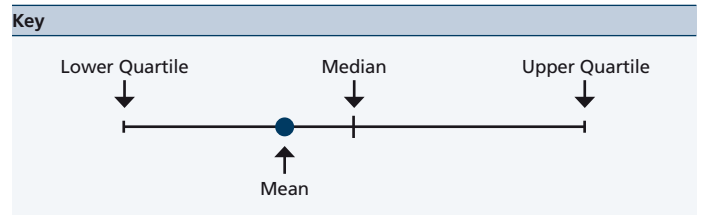
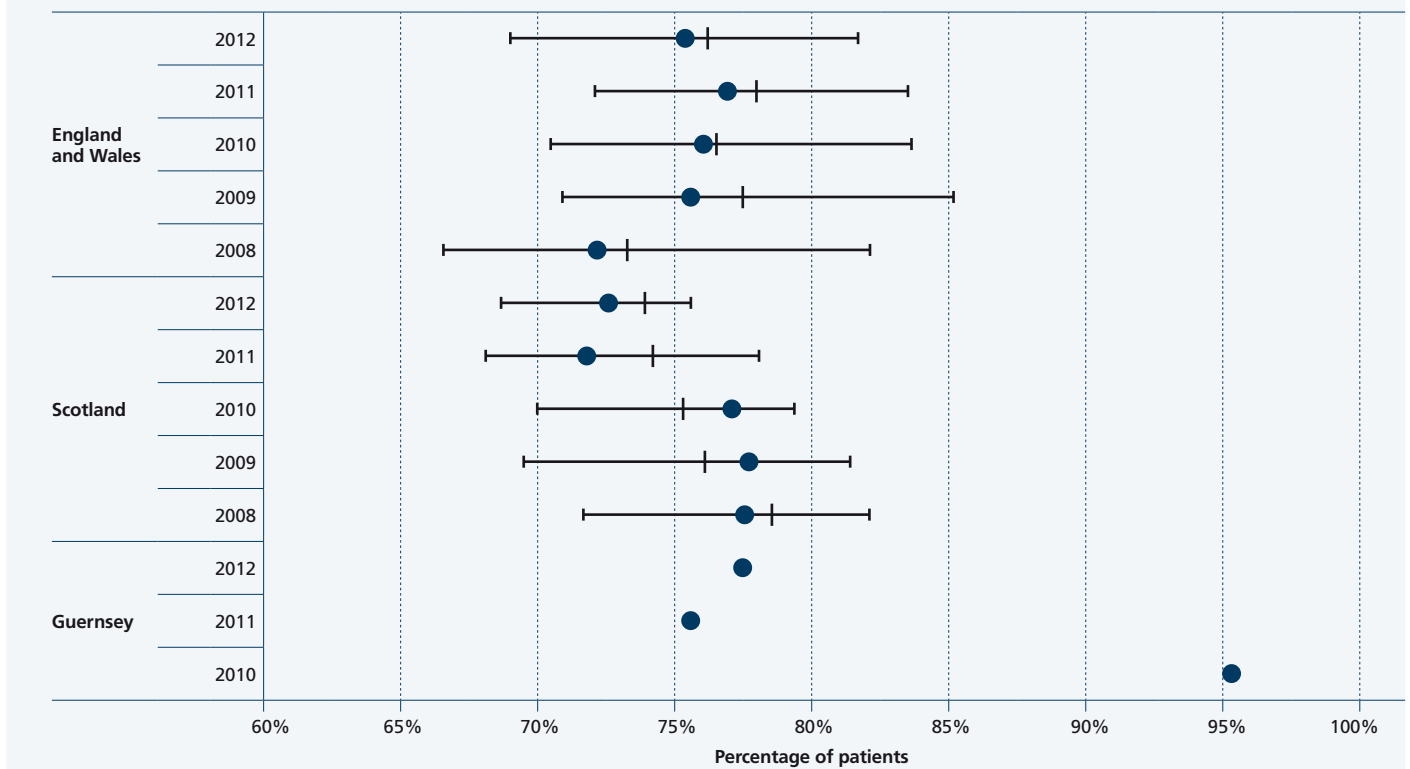
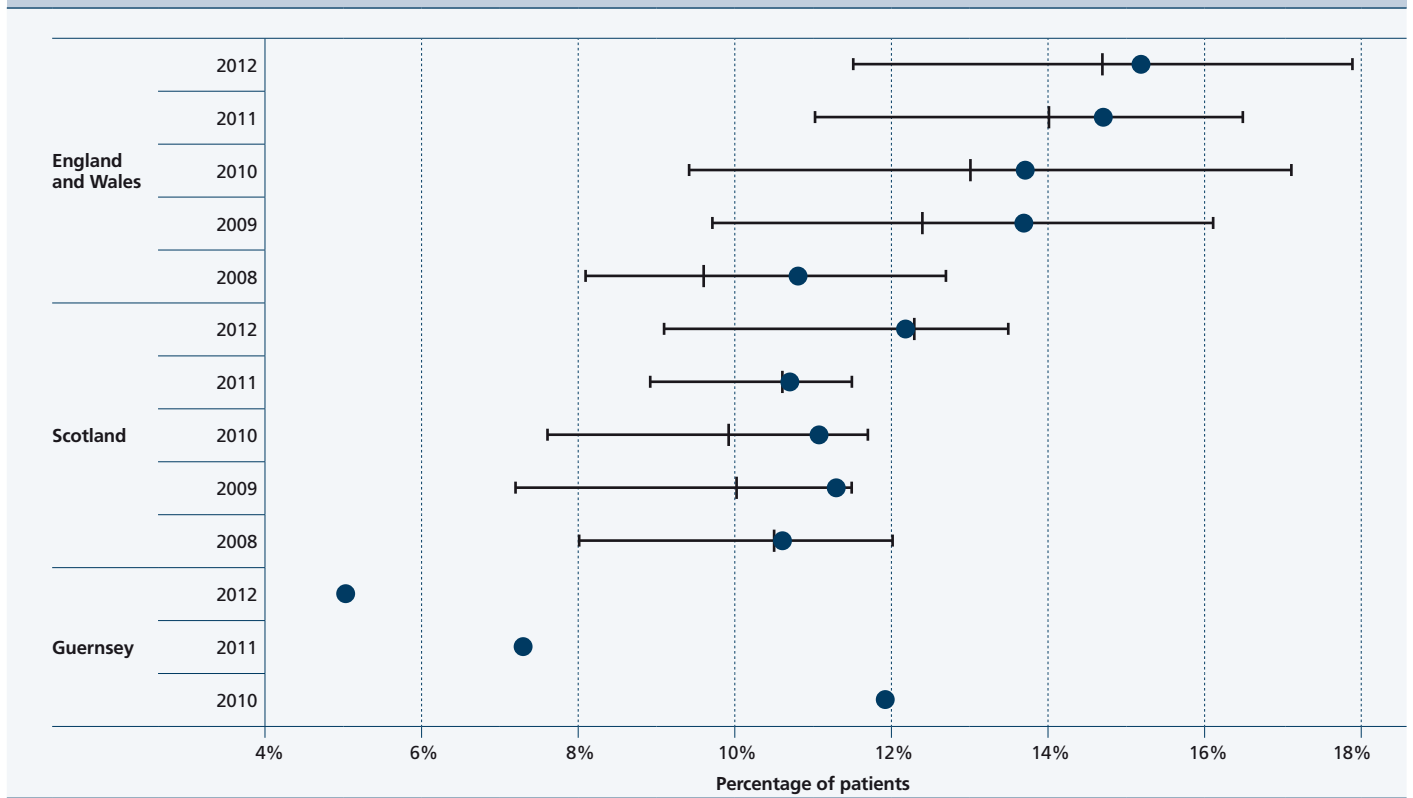


Figure 1
 Percentage of patients receiving a histological / cytological diagnosis



¹Based on Cancer Research UK (CRUK) data 2009.

Figure 2
Percentage of patients receiving an operation*



* Proportion of patients with histologically confirmed NSCLC receiving surgical resection is shown in Tables 3 and 4

Figure 3
Percentage of patients receiving any anti-cancer treatment

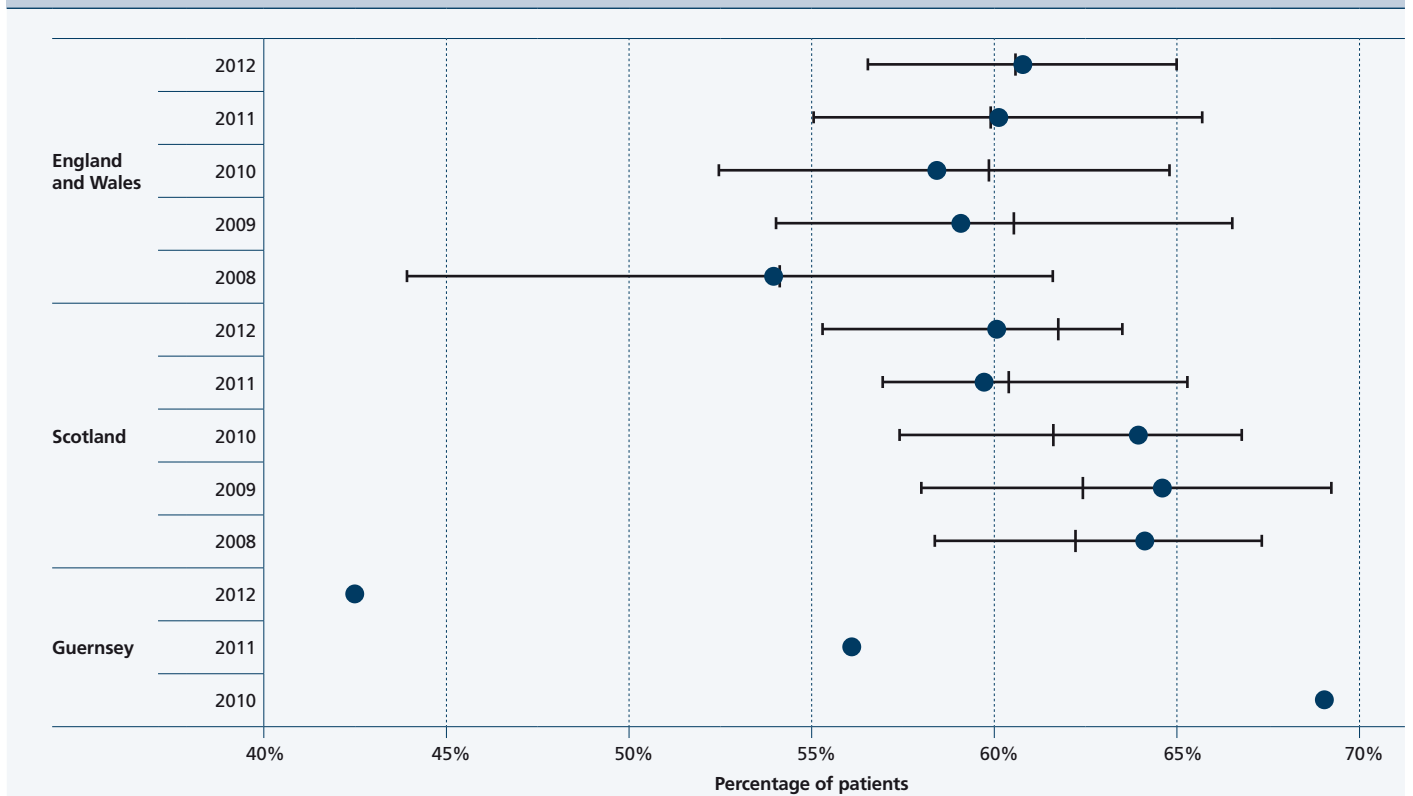


Figure 4
Percentage of patients receiving a CT scan before bronchoscopy

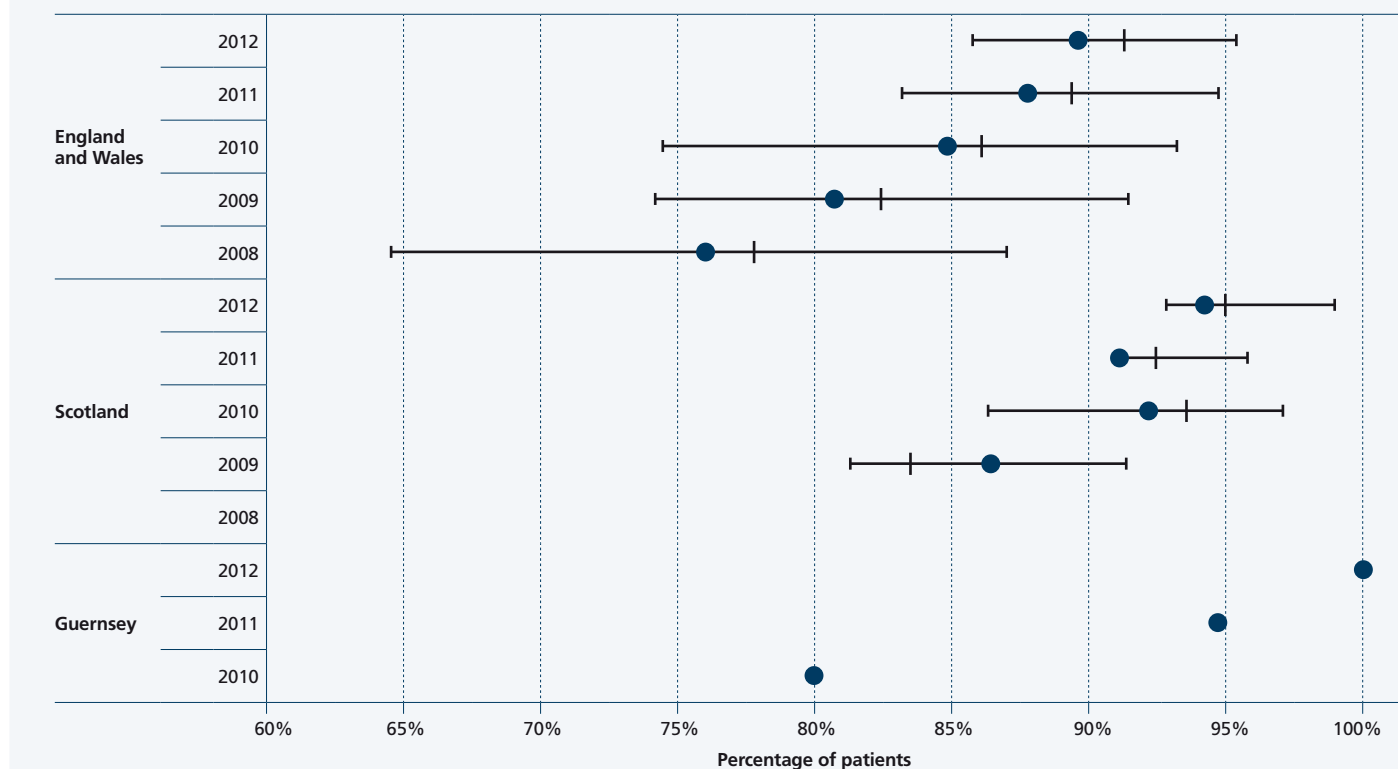
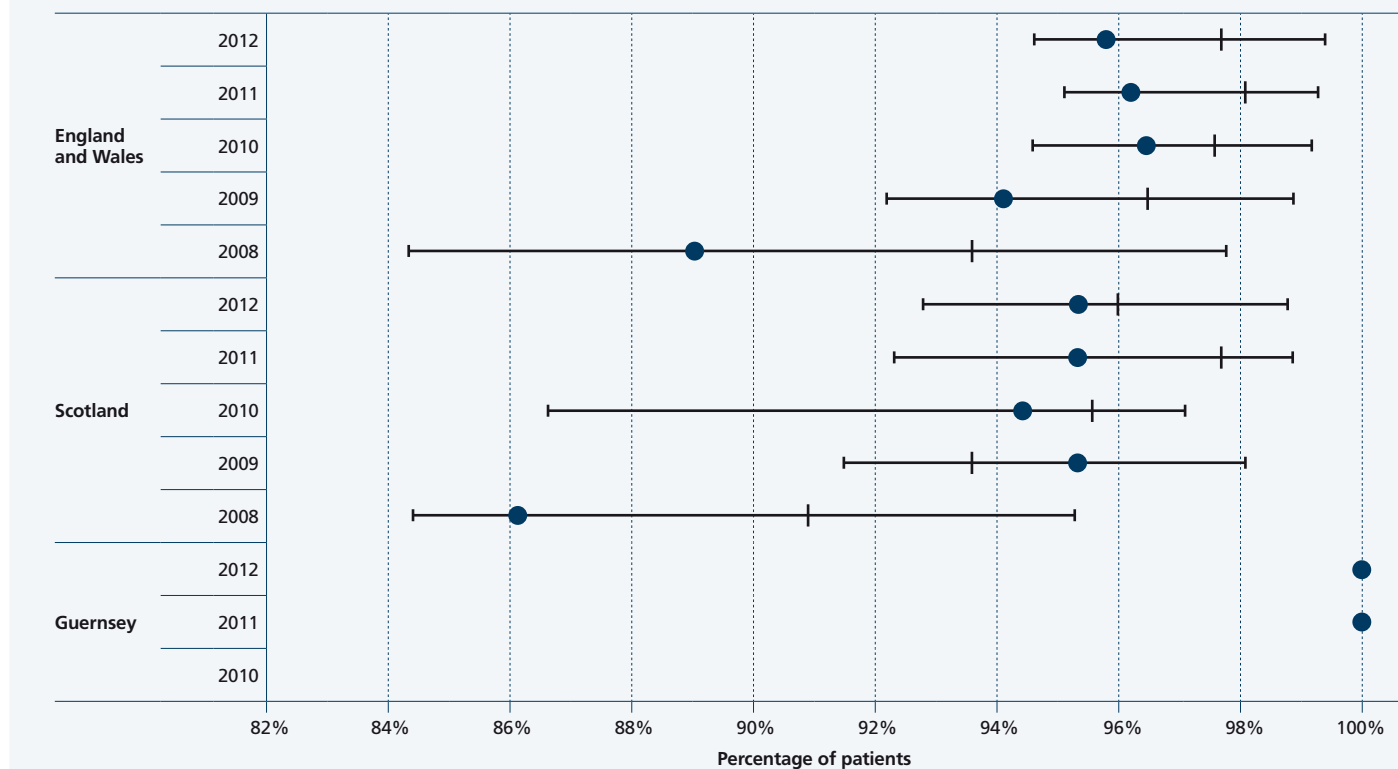


Figure 5
Percentage of patients discussed at MDT



Recommendations (England and Wales)

1. All Hospitals, Trusts and Health Boards should participate in this national audit, should submit data on all patients presenting to secondary care diagnosed with either lung cancer or mesothelioma, and should complete all relevant data fields for each individual patient.
2. Data completeness for key fields should exceed 85 per cent and for MDT completeness should exceed 95 per cent (See [Appendix 2](#) Local Action Plan).
3. Data completeness for the co-morbidity field should exceed 85 per cent, and for patients with Stage I-II and PS 0-1, completeness for FEV1 and FEV1% should exceed 75 per cent.
4. Maintain the level of 95 per cent of patients submitted to the audit being discussed at a Multi-Disciplinary Team (MDT) Meeting.
5. Histological/cytological confirmation rates below 75 per cent should be reviewed to determine whether best practice is being followed and whether patients have access to the whole range of biopsy techniques.
6. Non-Small Cell Lung Cancer, not otherwise specified (NSCLC NOS) rate of more than 20 per cent should be reviewed to ensure that best practice histological diagnostic techniques including immunohistochemistry are being followed, in order that patients receive appropriate chemotherapy regimens.
7. At least 80 per cent of patients are seen by a Lung Cancer Nurse Specialist (LCNS); at least 80 per cent of patients should have a Lung Cancer Nurse Specialist present at the time of diagnosis (note that these data are not available for Wales).
8. For patients undergoing bronchoscopy at least 95 per cent should have a CT scan prior to the procedure.
9. Surgical resection rates for NSCLC below the England and Wales average of 16 per cent should be reviewed. Furthermore, for early stage disease with good Performance Status (I and II), rates below 52 per cent should be reviewed to ensure that patients on the margins of operability/resectability are being offered access to specialist thoracic surgical expertise (including second opinions).
10. Active anti-cancer treatment rates below the England and Wales average of 61 per cent should be reviewed.

11. Chemotherapy rates for small cell lung cancer below the England and Wales average of 68 per cent should be reviewed.
12. Chemotherapy rates for good Performance Status (0-1) Stage III B / IV NSCLC below the England and Wales average of 57 per cent should be reviewed.

A Local Action Planning toolkit (LAP) is provided in [Appendix 2](#) to assist organisations in benchmarking against these quality measures. All organisations are encouraged to use the audit data to drive their service development in order to improve the standard of care for lung cancer patients. Trusts whose results in 2012 meet these recommendations should work to maintain their high standards and exceed them where appropriate.

Performance against some of these recommendations is highlighted by a system of colour coding in the data [Tables 1, 2, 4 and 5](#).

Scotland

The above recommendations do not apply to Scotland; therefore the data in the Tables are not colour coded. NHS Quality Improvement Scotland published National Lung Cancer Standards in March 2008. NHS Boards in all Scottish Networks participate in comparing 2012 results measured against these Standards, and where variance is shown action plans can be developed by Networks and NHS Boards and monitored by Regional Cancer Advisory Groups.

As part of the Scottish Government's National Cancer Quality Programme new Quality Performance Indicators (QPIs) for Lung Cancer were implemented for all patients diagnosed on, or after, 1st July 2013. Performance against these QPIs will be monitored following one year of implementation and will be subject to a robust governance process through Regional Cancer Networks, the Scottish Government and Healthcare Improvement Scotland.

It is important to stress that these quality measures are not targets, since in some cases there will be valid reasons for variation, such as case-mix and patient choice. Where applicable, organisations should take the case-mix adjusted results (published separately) into consideration in the evaluation of their service, although it is noted that in general case-mix does not explain the whole of the variation in practice across organisations.

Summary Details of Key Findings

How Many People Were Diagnosed With Lung Cancer?

In 2012 there were 35,366 patient records submitted from England and Wales (see [Figure 6](#)), 4,810 submitted from Scotland ([Figure 7](#)), and 40 submitted from Guernsey ([Figure 8](#)). Combined, this is approximately 98 per cent of the expected annual incidence and probably almost all of those cases presenting to secondary care

(some cases are diagnosed and treated in primary care, or are diagnosed at a post-mortem). Of these records, 367 were not suitable for further analysis (mainly from the English submissions) and predominantly due to no 'date first seen' being recorded, meaning that it was not possible to be certain that these were cases from 2012. [Figures 6, 7, and 8](#) show the incidence by cancer type.

Figure 6
Number of patient records submitted to the NLCA – England and Wales

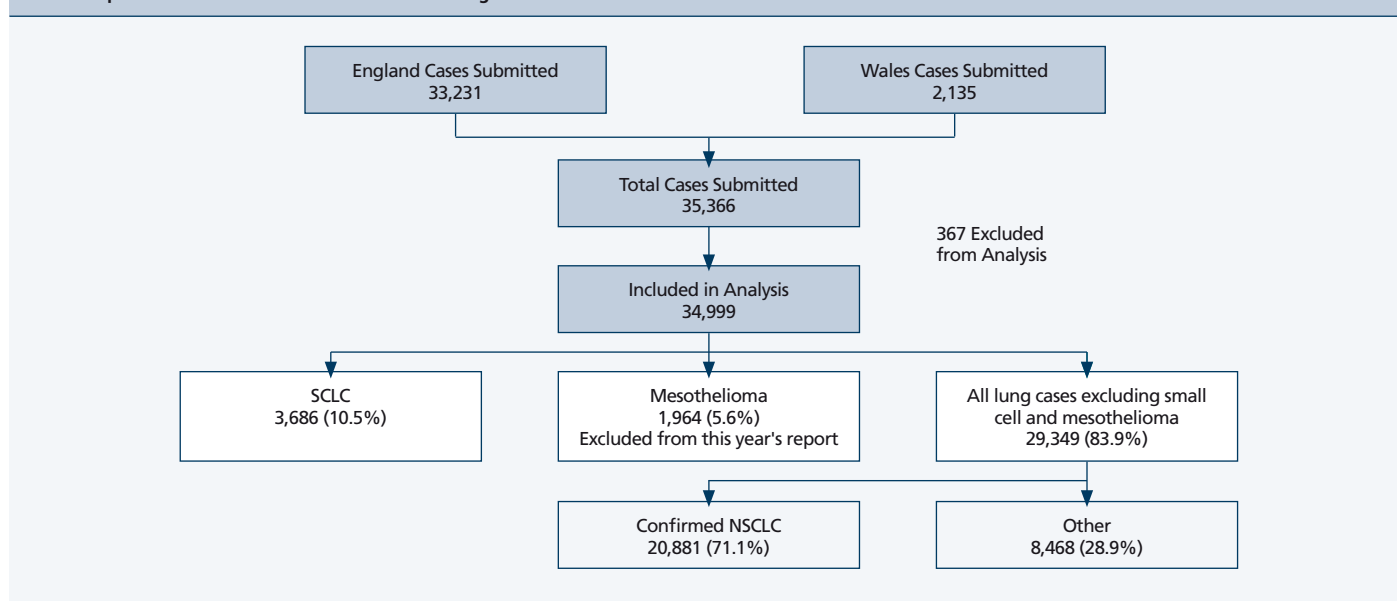


Figure 7
Number of patient records submitted to the NLCA – Scotland

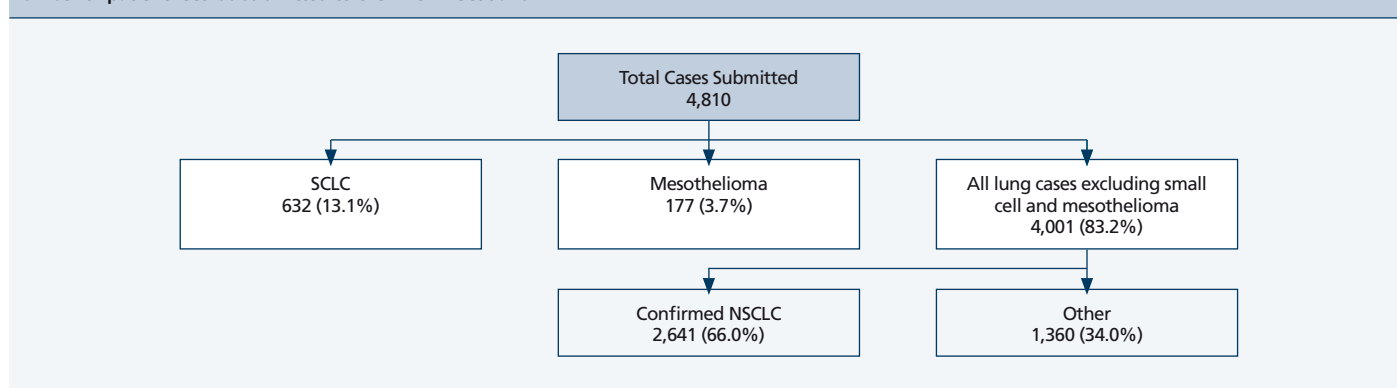
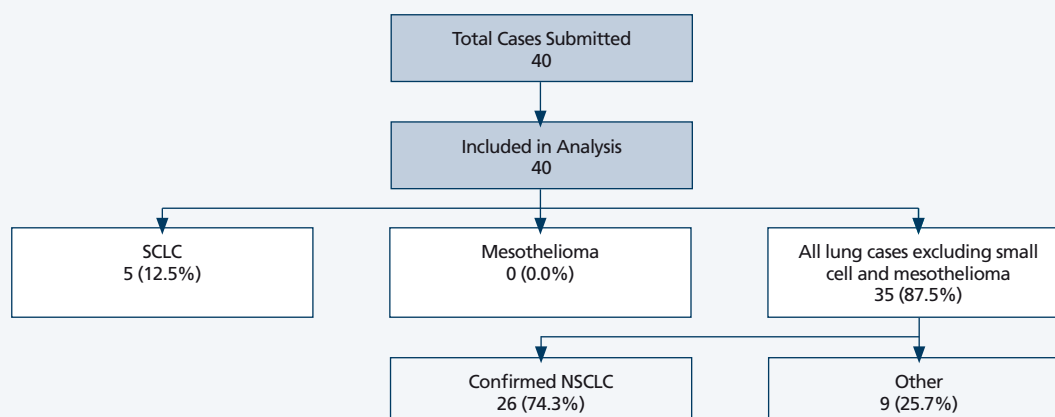


Figure 8
Number of patient records submitted to the NLCA – Guernsey



How Accurate are the Data in this Report?

Data submitted to the National Lung Cancer Audit need to be as complete as possible in terms of healthcare organisation participation, population coverage and data field completeness both to ensure the representative nature of the information and to make case-mix adjustment possible. Please refer to previous versions of the Annual Report for a full explanation of this issue.

Healthcare Organisation Participation

Every Trust or Health Board in England and Wales, and every Health Board in Scotland has participated in the audit. Princess Elizabeth Hospital, Guernsey has also participated in the audit.

Population Coverage

Figures 6-8 show that the audit has captured approximately 98 per cent of the expected number of cases nationally and almost all of those patients presenting to secondary care.

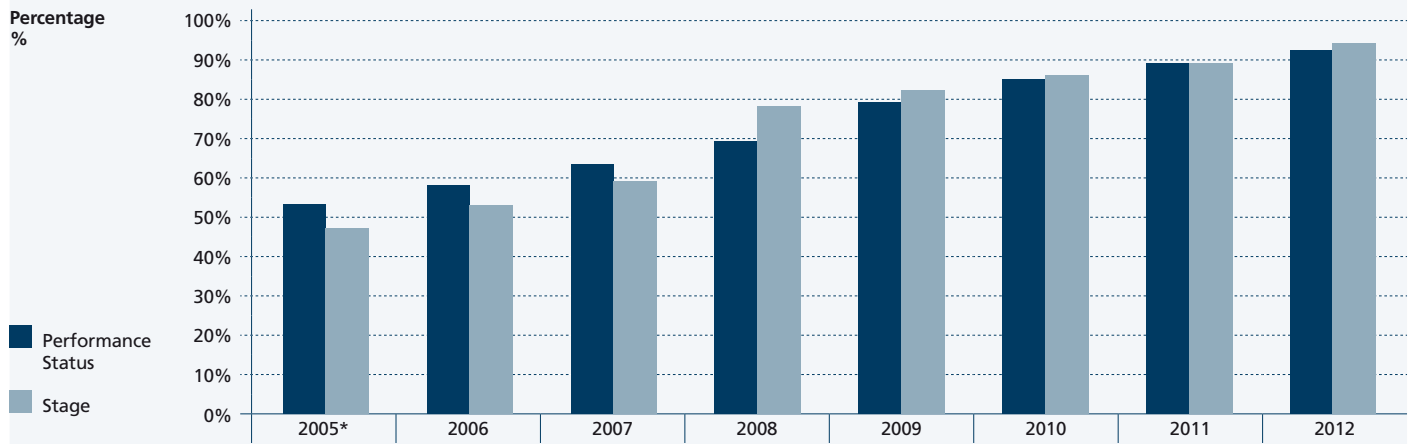
The “Data Completeness” section (formerly shown in Tables 1a-c) in previous years is now included in the online information sheet (published on www.hscic.gov.uk/lung).

The colour coding in the Tables reflects the targets set in the 2011 Local Action Plan (LAP). Trusts with a high tertiary workload or where the targets are known to not be applicable for other reasons are shown in blue throughout. Many of the Trusts in this category fully participate in the audit by submitting treatment data for other Trusts. However, their full contribution to the audit process may not be reflected by the way these audit results are presented. The treatment data entered by these Trusts are shown in Figure 10 on page 35.

Data Field Completeness

As previously stated data completeness results are available online only this year. Figure 9 illustrates that once again the data completeness of the key fields of Stage and Performance Status (PS) have improved for another year.

Figure 9
Data Completeness - England and Wales (2012)



* England Only

What is the Standard of Care Given to Patients?

Table 1 lists "Headline Indicators" (Process and Clinical Outcomes for England and Wales 2012) by Network and by Trust (key to codes given in Appendix 1) for all lung cancer cases across England and Wales. These indicators have been chosen to reflect the overall standard of care provided to patients. In interpreting these figures data completeness must be borne in mind, furthermore, the results as presented do not take into account the case-mix of patients. Adjustments to the results to account for case-mix will be available from the HSCIC website in due course. Where applicable, organisations should take the case-mix adjusted results into consideration in the evaluation of their service since although case-mix does not explain the whole of the variation in practice across organisations, it may show a particular result to be, or not to be, a statistical outlier. The colour coding in the Tables reflects the targets set in the 2011 Local Action Plan (LAP).

Data for Guernsey are shown in Table 2.

Similar data for Scotland are shown in Table 3. LAP targets do not apply to Scotland; hence the data are not colour coded. National Lung Cancer Standards published by NHS Quality Improvement Scotland in 2008 include Standards for rate of histological confirmation (minimum 75 per cent) and percentage of SCLC having chemotherapy (minimum 60 per cent) however these do not specify rates of resection or anti-cancer treatment.

Improvements in Care

Further details of the changes in the key outputs of the audit are shown in the Tables in the "Key Messages" section on page 9. For England and Wales, the proportion of patients receiving a histological/cytological diagnosis shows a marginal decrease to 75.3 per cent, the proportion of patients discussed at an MDT remains stable at 95.8 per cent, and the proportion of patients who receive a CT scan prior to a bronchoscopy procedure remains stable at 89.6 per cent. The anti-cancer treatment rate and the overall surgical treatment rate have both increased by almost one percentage point this year to 60.8 per cent and 15.2 per cent respectively.

In interpreting these results it should be noted that mesothelioma has been excluded from the denominator for England and Wales. Comparison to last year's results with mesothelioma removed from the denominator reveals exactly the same data pattern with almost no change to the relative increase or decrease in results.

It is clear from the key messages that there remains a marked variation in the outputs that the audit measures across organisations. This is apparent both at Network and even more markedly at Hospital Trust level. In the latter case, some of the more extreme variation is explained by low numbers of cases, or poor quality data, so a useful way of reporting the variation is the "interquartile range" (IQR), describing the range of values in the middle 50 per cent. In England and Wales, the IQR for histological/cytological confirmation is 69.0 - 81.7 per cent, and for surgical treatment it is 11.5 -17.9 per cent, the variation in range is almost identical to last year for these two measures. For anti-cancer treatment the IQR has reduced by over 2 percentage points and this year is 56.5 - 65.0 per cent. Similar variation is apparent for Scotland and Guernsey.

Converting Data into Service Improvement

Collecting data is only part of the audit process and it is important that the data is used to improve the services provided to patients and the outcomes of their treatment. There are numerous examples of local organisations doing just this. Furthermore, national organisations such as the National Institute for Health and Clinical Excellence, the British Thoracic Society and the National Cancer Peer Review Programme have all utilised data from the audit in their work programmes for lung cancer. Examples of some of the uses of the audit data are described in the list on page 17.

National Cancer Peer Review Programme (part of the National Cancer Action Team)	To provide data for the 'Clinical Lines of Enquiry' – outcome measures for the assessment of Lung Cancer Multi-Disciplinary Teams in England
Nottingham University – 'LUCADA Fellowship'. Funded by the Royal College of Physicians	Academic MD fellowships based on the use and interpretation of data from the NLCA has resulted in 6 peer-reviewed publications to date
European Respiratory Society Thoracic Oncology Assembly : 'European Initiative for the Quality Management of Lung Cancer'	Underpinning the long term goal of a pan-European comparative audit of lung cancer performance and outcomes
LungPATH – a National Audit and Service Improvement programme in lung cancer pathology in collaboration with Guy's & St Thomas' Hospital and King's College London	A programme, based on the elements of the NLCA. This national audit (funded by an unrestricted educational grant from the pharmaceutical industry) demonstrated marked variation in the quality of the process of the pathological diagnosis of lung cancer and in 2013 published a report including recommendations to address this http://www.hqip.org.uk/assets/NCAPOP-Library/NCAPOP-2012-13/LungPath-Phase-2-Report.pdf
Society of Cardiothoracic Surgeons, The National Cancer Intelligence Network and Nottingham University	Examining the detail underpinning the variation in surgical resection rates and surgical outcomes for lung cancer patients across the UK
The Health Foundation	Improving Lung Cancer Outcomes Project (ILCOP) - described separately
The Department of Health and Cancer Research UK's International Cancer Benchmarking Partnership and the UK Cancer Registries: the collection of staging data on lung cancer	Data on the stage of cancers is essential for the interpretation of variations in cancer survival both within the UK and across national boundaries. The collection of staging data for lung cancer in the NLCA has improved the proportion of patients with stage recorded in the Cancer Registries having significant impact such initiatives as the International Cancer Benchmarking Partnership
Oxford University Department of Biomedical Engineering	Ph.D. project on clinical decision support and machine learning. The output of the work will be in the form of a clinical decision support platform, intended to act as a software tool to assist the clinicians in coming to informed, timely, safe and effective decisions in lung cancer care.
Scottish Government Health Directorate: Detect Cancer Early Programme	From 2012 routine quarterly staging data is being supplied by Health Boards from audit within the three Scottish Cancer Networks to measure the target to increase the number of Scots diagnosed with Stage 1 cancer by 25 per cent for three cancer types including lung cancer.

Focus on Surgical Resection



Focus on Surgical Resection

Surgery is the treatment of choice for patients with Stage I-II NSCLC, offering the best chance of a cure, although there is increasing evidence that new radiotherapy techniques may offer an alternative for selected patients. Careful preoperative assessment of the patient's overall medical condition, especially the patient's pulmonary reserve and cardiac status, is critical in considering the benefits of surgery.

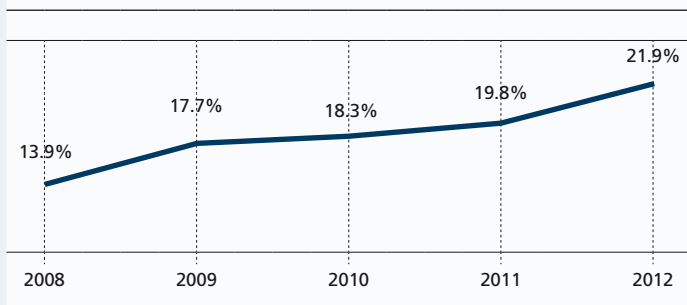
A low number of surgical resections has been considered as part of the explanation for poorer survival of lung cancer patients in the UK. Measurement of the proportion of patients undergoing surgical resection is an important measure of performance of lung cancer teams. This proportion can be calculated against a variety of denominators (all patients, all patients excluding SCLC and mesothelioma, all histologically confirmed NSCLC). Data presented here uses the latter, but the other measures, including calculation of "odds ratio" adjusted for clinical features will be included in the online audit data reports.

Focus on Surgical Resection

For patients first seen in 2012, the mean resection rate in histologically confirmed NSCLC for England and Wales was 21.9 per cent, and this figure has been rising steadily over the past few years, although as indicated in [Focus: Chart 2](#), a wide geographic variation persists.

The equivalent figure for 2012 for Scotland is 20.5 per cent.

Focus: Chart 1
Resection rate in confirmed NSCLC by year (England and Wales)



Focus: Chart 2
Surgical resection rates in 2012

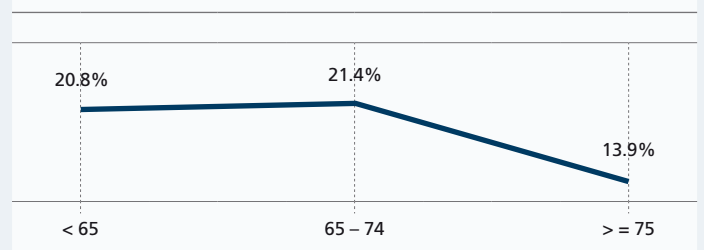
Network	2012 Resection	Trend
N01	16.1%	
N02	27.0%	
N03	30.8%	
N06	26.0%	
N07	23.8%	
N08	22.4%	
N11	21.6%	
N12	23.8%	
N20	18.0%	
N21	21.0%	
N22	19.6%	
N23	15.9%	
N24	15.8%	
N25	15.1%	
N26	19.0%	
N27	19.8%	
N28	25.6%	
N29	17.4%	
N30	27.8%	
N31	22.2%	
N32	18.4%	
N33	15.2%	
N34	19.5%	
N35	26.1%	
N36	20.9%	
N37	17.8%	
N38	18.9%	
N39	27.0%	
NWW	18.4%	
SWCN	15.8%	

Making Sense of the Data

The Table ([Focus: Chart 2](#)) shows the resection rate in all histologically confirmed NSCLC for Cancer Networks in England and Wales, as well as the five year trend compared to the national average. Each result is made up of the results from individual hospitals in that region. The "optimum" resection rate in this group of patients is not known. For younger patients with good performance status and no co-morbidities, surgery is without doubt the treatment of choice, but as age increases and general health declines, the risks of a surgical procedure may be considered too great by the patient or Surgeon. This approach to risk may explain some of the geographic variation in resection rates even when case-mix is taken into account.

The graph ([Focus: Chart 3](#)) demonstrates the impact age has on resection rates and organisations looking to increase their rate of surgery can look in detail at their approach to older patients as a means of improvement. Results for individual hospitals are available in the Tables in this report as well as in the online audit reports (the latter will include case-mix adjusted odds ratios which take account of variation in patient clinical features).

Focus: Chart 3
Resection rate by age group, 2008-2012



Why Increase Resection Rate?

Compared with some European countries, England has low lung cancer survival and low use of surgical resection. A recent paper extracted data on 77,349 NSCLC patients diagnosed between 2004 and 2006 from the English National Cancer Repository Dataset. The researchers divided the patients into five groups, according to the frequency of surgical resection in their Primary Care Trust (PCT) area, and looked at mortality rates in each group. They found large geographical variation in the surgical resection rate for NSCLC (3-18 per cent). A low frequency of resection in a PCT area was strongly associated with overall mortality.

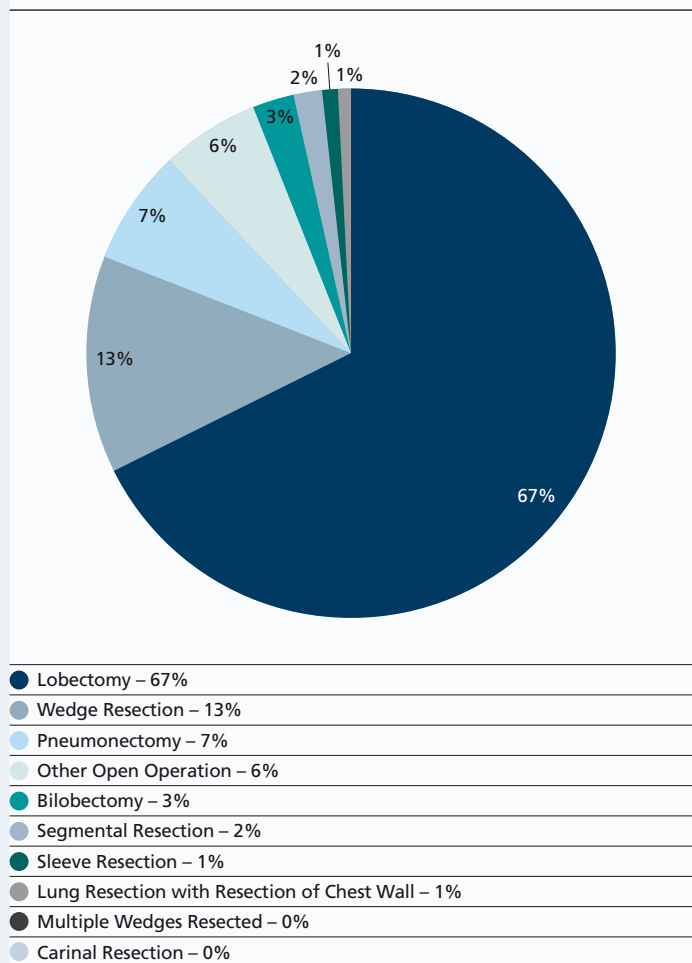
Focus on Surgical Resection

The researchers concluded that if compared all PCT areas had the same resection rate as the highest resecting PCT, then 5420 deaths could be delayed in the overall NSCLC group, whereas about 146 more deaths could be expected amongst the resected patients (due to the complications of the increased surgery).¹¹

17,628 Operations (2008 to 2012)

The surgical procedures carried out are recorded as Office of Population Censuses and Surveys (OPCS) codes. The vast majority of surgical procedures are lobectomies, with smaller numbers of larger resections (pneumonectomy) and smaller resections (wedge and segmental). A small number of sleeve resections and chest wall resections are also recorded (See [Focus: Chart 4](#)).

Focus: Chart 4
Operations, 2008-2012



¹¹Variation in surgical resection for lung cancer in relation to survival: population-based study in England 2004-2006. *Eur J Cancer*. 2012 Jan;48(1):54-60. doi: 10.1016/j.ejca.2011.07.012. Epub 2011 Aug 24. Riaz SP, Lüchtenborg M, Jack RH, Coupland VH, Linklater KM, Peake MD, Møller H. <http://www.ncbi.nlm.nih.gov/pubmed/21871792>

Focus on Patient Pathway



Focus on Patient Pathway

Regardless of the lung cancer type and stage, all patients should be investigated quickly and efficiently, and be supported through the diagnostic pathway with high quality and personalised information. All patients should have their case discussed in a Multi-Disciplinary Team (MDT) meeting. Here a panel of specialists (including Surgeon and Oncologist) will recommend the most appropriate form of treatment as well as any other treatment option that the patient can consider.

Whilst the diagnostic pathway for lung cancer patients is often complex, national guidance makes clear recommendations on ways to streamline investigations, for example by ensuring that CT scans are carried out before more invasive tests, and choosing a mode of biopsy that offers information on stage as well as diagnosis.

Relevant measures collected by the audit include:

- Proportion of patients discussed in an MDT
- Proportion of patients seen by a Lung Cancer Nurse Specialist (LCNS)
- Proportion of patients where the Nurse Specialist is present at the time of diagnosis
- Proportion of patients where a CT scan was carried out before a bronchoscopy

Whilst the audit collects data on the timing of different elements of the patient pathway, these are not routinely reported upon, since the NHS already collects and analyses data on "Cancer Waiting Times".

Focus on Patient Pathway

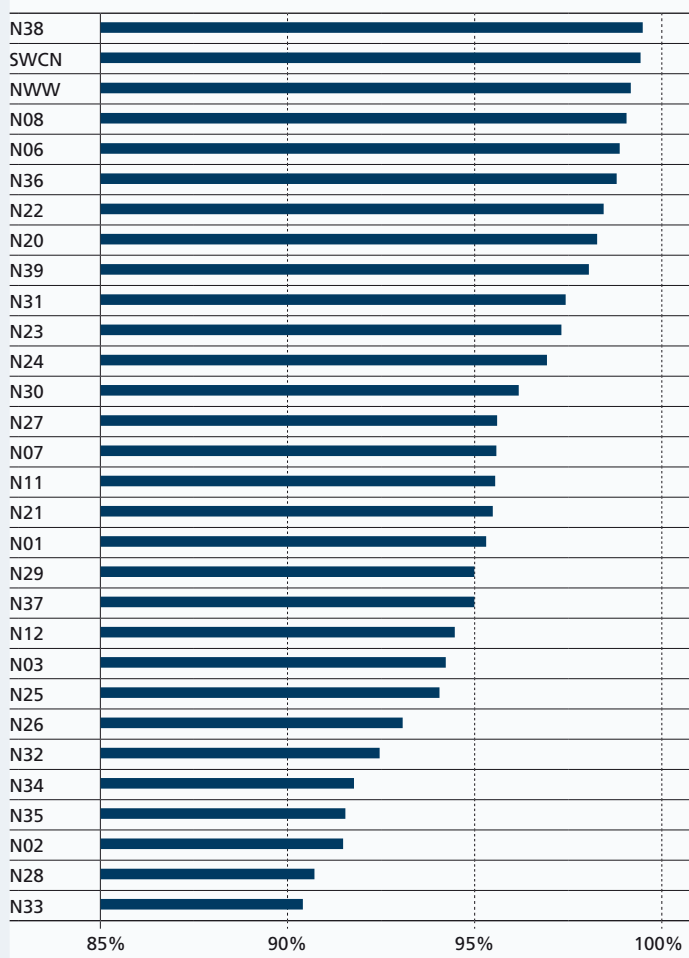
Discussed at MDT

For patients first seen in England and Wales in 2012, 95.8 per cent are discussed at an MDT.

The graph (Focus: Chart 1) shows the results for individual Networks and data on individual Trusts is available in the data Tables.

Since some patients will unfortunately have died before there is an opportunity to discuss their case, a figure of 100 per cent will rarely be achieved, but where the figure falls below 95 per cent it is advisable that organisations review their mechanisms and policies to ensure that all suitable patients benefit from the multi-disciplinary expertise.

Focus: Chart 1
Patients discussed in an MDT, 2012



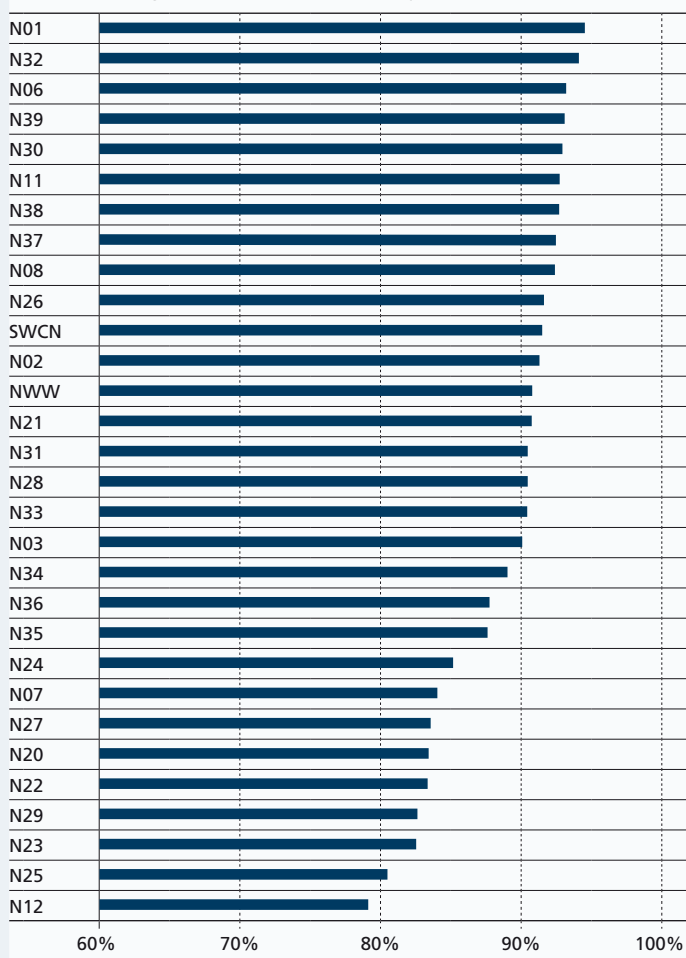
CT Scan before Bronchoscopy

It is now recognised that it is rarely appropriate for a bronchoscopy to be carried out unless a CT scan has been done first. The two main drivers of this are the frequent need to carry out a repeat bronchoscopy to obtain staging information from mediastinal lymph nodes, and the finding of more appropriate biopsy sites once CT information is available.

For patients first seen in England and Wales in 2012, 89.6 per cent have a CT scan before any bronchoscopy procedure.

The graphs (Focus: Chart 1 and 2) show the results for individual Networks and data on individual Trusts is available in the data Tables.

Focus: Chart 2
Patients receiving CT Scan before bronchoscopy, 2012



Focus on Patient Pathway

Lung Cancer Nurse Specialist

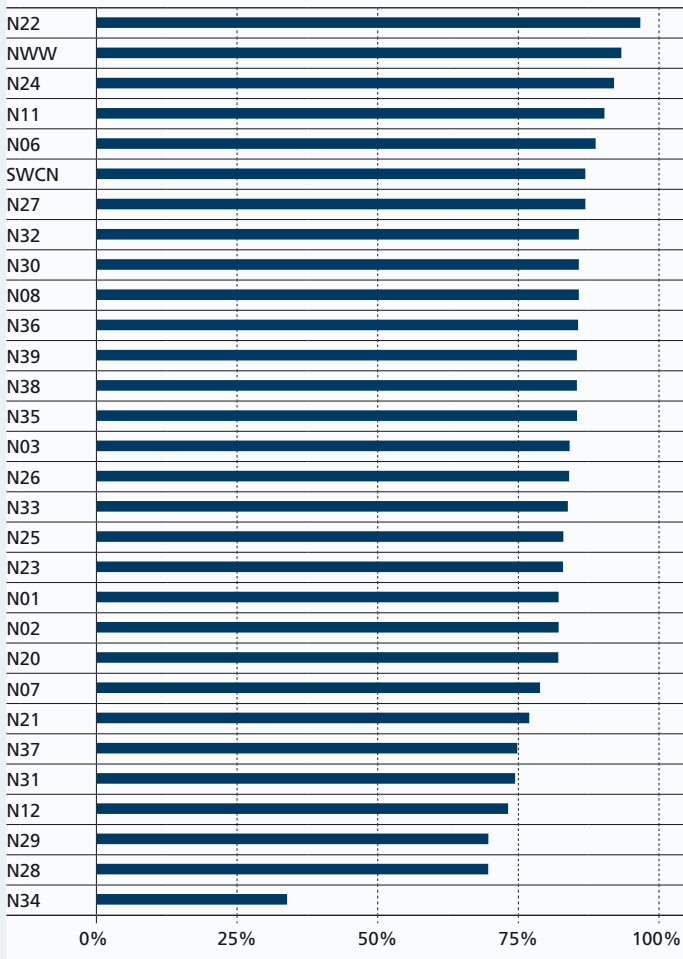
The role of Clinical Nurse Specialist was introduced in 1995, and they play a vital role in the delivery of high quality care and treatment to patients with lung cancer. Surveys of patients and carer experience repeatedly demonstrate the value placed upon the skills of these nurses in such areas as information provision, holistic assessment, symptom management, psychological support, co-ordination of care and patient advocacy. It is recommended that all patients should have equitable access to a LCNS at the time of diagnosis to guarantee that their physical, social and emotional needs, and their treatment options are appropriately assessed and discussed from the beginning of their cancer journey.

For patients first seen in England and Wales in 2012, 82.3 per cent are seen by a Lung Cancer Nurse Specialist (LCNS), and 61.2 per cent have the nurse present at the time of diagnosis. These results are improvements on previous years (79.5 per cent and 58.6 per cent in 2011).

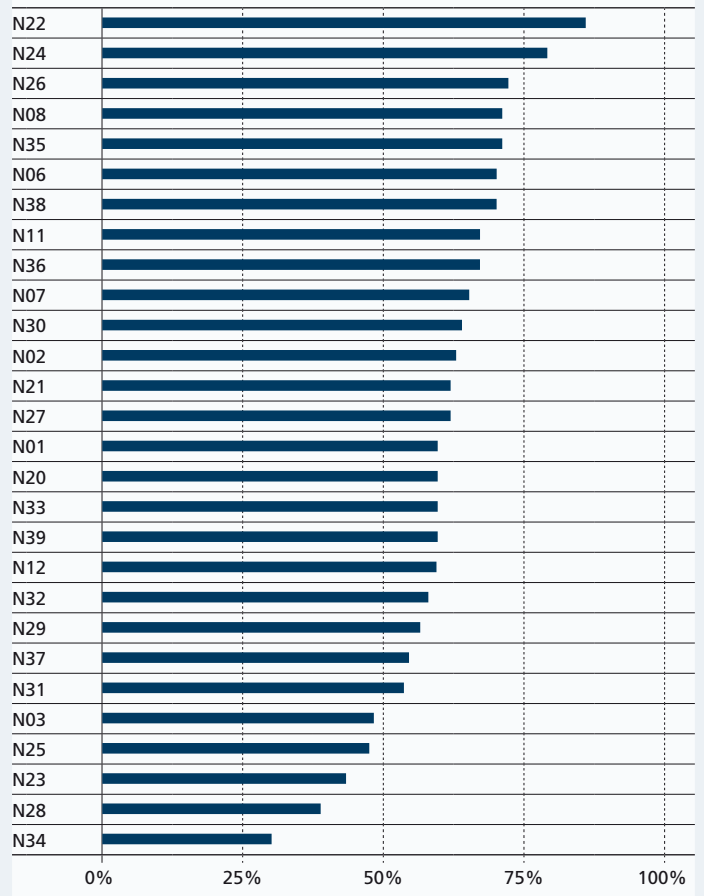
The graphs (Focus: Chart 3 and Focus: Chart 4) show the results for individual Networks. Data on individual Trusts is available in the data Tables. Note that Welsh Trusts do not collect data on whether the patient sees the LCNS at diagnosis.

The Roy Castle Lung Cancer Foundation produced a report in January 2013 entitled "Understanding the value of lung cancer nurse specialists". This document, available from their website (www.roycastle.org) examines the role of LCNSs, with the aim of helping people understand the vital contribution they make to the delivery of high quality care and to improved outcomes for patients with lung cancer. It includes 11 recommendations designed to ensure that LCNS posts are protected and, where necessary, increased in number, so that all patients with lung cancer have equitable access to LCNS input in their lung cancer pathway, and demonstrates the importance of fully supporting LCNSs with sufficient resources to carry out their work to the highest possible standards.

Focus: Chart 3
Seen by LCNS



Focus: Chart 4
LCNS present at diagnosis



Focus on Survival



Focus on Survival

Treatment of lung cancer aims to prolong survival and improve quality of life by improving symptoms. At this time, measuring patient quality of life is not routinely carried out, so measurement of survival offers the best way of assessing effectiveness of treatment. Median survival (the time taken for 50 per cent of the patients to die from their cancer) is one way of measuring survival of the whole cohort of patients in England and Wales from 2012.

Lung cancer has one of the lowest survival outcomes of any cancer because around 50 per cent of patients are diagnosed at a late stage when curative treatment is not possible. Similarly, the median age

at diagnosis of 72 years, coupled with high smoking prevalence means that there is a high incidence of co-morbidities in these patients that impacts upon the ability to deliver curative treatments. Even for those patients who have “curative” treatment with surgery or radical radiotherapy, lung cancer may recur some months or years later.

However, for patients with incurable lung cancer, active anti-cancer treatment and specialist palliative care still offer the potential to extend life and can be measured by six-month or one-year survival.

Focus on Survival

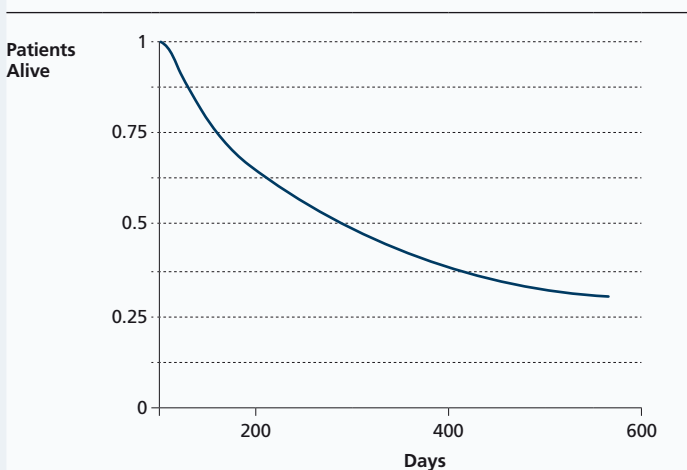
Overall Survival

The graph (Focus: Chart 1) is a Kaplan-Meier plot of survival of the whole cohort of patients from 2012. For these patients, the median survival is 221 days (7.4 months), with a six-month survival of 55 per cent and a one-year survival of 39 per cent (based on the whole cohort).

These short-term survival figures appear to have improved compared to 2008 when median survival was 191 days (6.4 months) and one-year survival was 3 per cent.

The audit does not hold individual patient data for other countries and so is unable to calculate survival.

Focus: Chart 1
Overall survival



Median Survival

The Table (Focus: Chart 2) shows the median survival for Cancer Networks in England and Wales. Each result is made up of the results from individual hospitals in that region.

Results for individual hospitals will be available in the online audit reports and will include statistically adjusted "odds ratios" to take account of differing clinical features of patients (such as age, Stage and Performance Status).

Survival data has to be interpreted with caution, to avoid making inappropriate judgements. Survival time has been calculated from the date first seen in secondary care rather than date of diagnosis since the latter is not always recorded accurately in the audit. If patients in one geographical area tend to be seen in secondary care (or are diagnosed) earlier, then the patient's survival time is increased. This increase in survival time makes it seem as though those patients are living longer when that may not be happening. This is called lead-time bias.

Focus: Chart 2
Median survival of all lung cancer patients, 2012

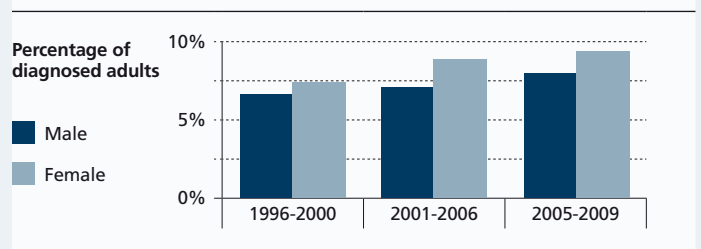
Network	Number of Days	Total
N01	210	210
N02	224	224
N03	240	240
N06	244	244
N07	224	224
N08	215	215
N11	211	211
N12	223	223
N20	152	152
N21	235	235
N22	216	216
N23	227	227
N24	213	213
N25	260	260
N26	236	236
N27	280	280
N28	241	241
N29	216	216
N30	251	251
N31	206	206
N32	242	242
N33	184	184
N34	198	198
N35	205	205
N36	218	218
N37	272	272
N38	209	209
N39	224	224
NWW	190	190
SWCN	179	179

Making Sense of the Data

The NLCA has been collecting data for eight years and it is not yet possible to use the data to measure improvements in long-term survival. Cancer Registries on the other hand, have collected survival data over many years and this data does indicate a slowly improving one-year and five-year survival over the past 20 years (see Focus: Chart 3, data from www.cancerresearchuk.org).

Improving longer-term survival depends upon finding lung cancers at an earlier stage and offering curative treatment to as many of these patients as possible.

Focus: Chart 3
Five-year survival for lung cancer, England, 1996-2009



Case-Mix Adjustment

A typical explanation for different audit results from different organisations (Hospital Trusts or Cancer Networks) is that there is a different "case-mix". For example, a Hospital with a low treatment rate might argue that the patients they treat are older, more socially deprived, have more advanced disease, or poorer fitness (Performance Status).

The NLCA collects data that allows such factors to be taken into account. Taking anti-cancer treatment as an example, a statistical technique known as "logistic regression" calculates the likelihood of a patient in an organisation getting treatment compared to a baseline (typically the largest organisation) assuming that patients are matched for their case-mix.

This measure of likelihood of treatment is called an "odds ratio". The baseline organisation will always have an odds ratio of 1.0. If Hospital X has an odds ratio of 0.9, we can say that patients in that Hospital are 10 per cent less likely to have treatment (1.0 minus 0.9, converted to a percentage). Odds ratios have a further benefit, in that they provide so-called "confidence intervals", indicating how confident we can be that the observed differences are statistically important.

Improvements in data collection mean that Stage and Performance Status are now recorded in around 90 per cent of cases. In order to further refine the statistical analyses, it is important in future that organisations improve recording of co-morbidity and lung function. As mentioned in "Key Recommendations", we have suggested that data completeness for the co-morbidity field should exceed 85 per cent and for patients with Stage I-II and PS 0-1, completeness for FEV1 and FEV1% should exceed 75 per cent.

Case-mix adjusted data in an electronic spreadsheet format will be available from the HSCIC website in due course.

Improving Lung Cancer Outcomes Project (ILCOP)

The Improving Lung Cancer Outcomes Project funded by the Health Foundation and hosted by the Royal College of Physicians aimed to reduce variation in outcomes and improve patient experience via reciprocal peer to peer review visits and facilitated quality improvement. During the intervention phase, the thirty participating Trusts provided many examples of local improvement projects which can be found on the RCP ILCOP website: www.rcplondon.ac.uk/projects/improving-lung-cancer-outcomes-project-ilcop

Evaluation of NLCA Key Indicators in 2012 showed that, compared to 2009, there was a small but statistically significant increase in active treatment rates in the intervention group compared to the controls and non-participants. In the intervention group overall patient experience scores did not change significantly during the study however the mean total score for the five Trusts with the worst baseline scores did show significant improvement.

The Project Team are currently exploring whether this form of supportive peer review for lung cancer quality improvement can be incorporated into a comprehensive national programme.

Focus on Chemotherapy



Focus on Chemotherapy

Chemotherapy is used widely in the treatment of lung cancer. It is the primary mode of treatment for small cell tumours, where used alone or in combination with radiation, it may be curative. In NSCLC, it is most often used as part of a palliative treatment regime in locally-advanced or metastatic disease where it prolongs survival and improves quality of life. It is also used before surgery (neo-adjuvant) and after surgery (adjuvant) to reduce the chance of later relapse.

The audit focusses on two areas of chemotherapy usage that are recommended in National Guidelines such as the NICE Quality Standards:

- Chemotherapy in SCLC
- Chemotherapy in IIIB/IV NSCLC

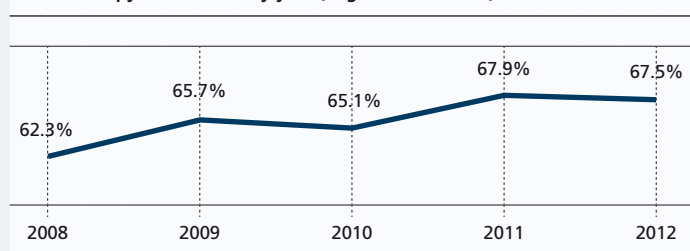
Focus on Chemotherapy

Chemotherapy in SCLC

For patients first seen in 2012, the mean chemotherapy rate in SCLC for England and Wales was 67.5 per cent, a small increase over previous years (See [Focus: Chart 1](#)).

The equivalent figure for 2012 for Scotland is 70.6 per cent.

Focus: Chart 1
Chemotherapy rate in SCLC by year (England and Wales)



The Table ([Focus: Chart 2](#)) shows the proportion of patients diagnosed with SCLC who receive chemotherapy in each Cancer Network in England and Wales, as well as the five year trend, compared to the national average. SCLC often progresses rapidly from the time of presentation to secondary care, and unless diagnosis and treatment are carried out expeditiously, the patient may well deteriorate during the diagnostic pathway to the point where they can no longer be offered treatment due to their poor general condition. NICE Quality Statement 13 states that "People with small cell lung cancer have treatment initiated within two weeks of the pathological diagnosis". Therefore, a low rate of treatment may represent a slow pathway, or alternatively a generally nihilistic approach to treatment. On the other hand, a very high rate of treatment may mean that some patients may receive chemotherapy inappropriately, leading to side effects, deterioration in quality of life and reduced survival. The results for individual Trusts are shown in the later data Tables, with suggestions for service improvement contained within the Local Action Plan Template. ([Appendix 2](#))

Focus: Chart 2
Patients diagnosed with SCLC receiving chemotherapy in England and Wales

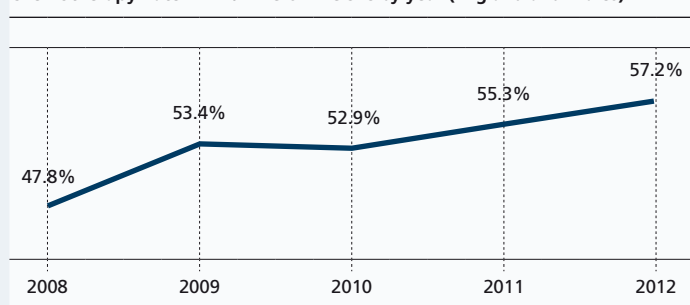
Network	2012 SCLC Chemo	Trend
N01	80.4%	
N02	69.4%	
N03	68.4%	
N06	72.6%	
N07	69.9%	
N08	63.8%	
N11	63.6%	
N12	56.7%	
N20	49.3%	
N21	65.8%	
N22	63.8%	
N23	63.0%	
N24	71.1%	
N25	67.3%	
N26	73.5%	
N27	66.7%	
N28	72.0%	
N29	60.7%	
N30	69.6%	
N31	72.5%	
N32	53.7%	
N33	63.8%	
N34	63.8%	
N35	69.6%	
N36	66.5%	
N37	70.1%	
N38	67.8%	
N39	69.6%	
NWWW	63.5%	
SWCN	63.1%	

Focus on Chemotherapy

Chemotherapy in Stage IIIB/IV PS 0-1 NSCLC

For patients first seen in 2012, the mean chemotherapy rate for good Performance Status patients with locally advanced or metastatic NSCLC for England and Wales was 57.2 per cent, a small increase over previous years (See [Focus: Chart 3](#)).

Focus: Chart 3
Chemotherapy Rate in IIIB/IV PS 0-1 NSCLC by year (England and Wales)



Offering systemic therapy (most often chemotherapy) to patients with people with Stage IIIB or IV non-small cell lung cancer and eligible Performance Status is recommended in NICE Quality Statement 12. Patients with this type of cancer often question whether it is worth them having treatment, as they will perceive treatment as likely to impair their quality of life. However, published research confirms that the majority of patients having palliative chemotherapy will actually enjoy improved quality of life as well as modest improvements in survival. Communicating these issues clearly and sensitively to patients may be the key to improving treatment rates. The Table ([Focus: Chart 4](#)) shows the chemotherapy rates and five year trend compared to the national average for these patients by Cancer Network, with results for individual Trusts available in the later data Tables.

Focus: Chart 4
NSCLC Chemotherapy rates, 2012

Network	2012 NSCLC Chemo	Trend
N01	70.9%	
N02	51.1%	
N03	63.3%	
N06	67.6%	
N07	55.2%	
N08	63.8%	
N11	58.5%	
N12	47.9%	
N20	48.1%	
N21	51.9%	
N22	63.0%	
N23	64.2%	
N24	62.6%	
N25	59.1%	
N26	49.0%	
N27	51.0%	
N28	66.2%	
N29	44.9%	
N30	52.1%	
N31	48.7%	
N32	62.4%	
N33	45.2%	
N34	43.6%	
N35	58.2%	
N36	62.2%	
N37	56.9%	
N38	58.4%	
N39	51.8%	
NWWW	63.5%	
SWCN	58.9%	

Linking Data

Radiotherapy

There have been fields in the National Lung Cancer Audit dataset since its inception and the 'fact of radiotherapy' being given has been included as part of the 'Active Treatment Rate' Headline Indicator since the first Annual Report. Although there is a field for RT treatment intent in the LUCADA dataset, palliative or curative intent has never been utilized in the reporting because its interpretation is uncertain. Since the NLCA's inception, the NHS developed a Radiotherapy Dataset (RTDS) which was mandated for England in 2009 and has been managed by the National Clinical Analysis and Specialised Applications Team (NatCanSat) in Liverpool. The RTDS draws data on dose, fractionation and scheduling directly from linear accelerators and there are other data fields which are completed manually in RT Centres. There have been delays in gaining access to data in a form that allows the NLCA to examine RT by dose and treatment intent. These issues are now resolved and the audit plans to use these data to report more extensively on the utilization of radiotherapy.

Chemotherapy

As with radiotherapy, there have been data fields in the NLCA dataset on chemotherapy from the outset, recording whether or not a patient has received chemotherapy rather than details of drug regimen or dose. This field robustly records the 'fact of chemotherapy' occurring with regards first line treatment, but the LUCADA database was not designed to record second line treatment. These chemotherapy data contribute to the reporting of 'Active Treatment Rate' and to chemotherapy rates for small cell lung cancer and the treatment of patients with Stages IIIB & IV non-small cell lung cancer of good Performance Status. The National Cancer Intelligence Network (NCIN) developed the Systemic Anti-Cancer Therapy (SACT) dataset and that became mandatory in England from April 2012. Most of the data is now being drawn from e-prescribing systems and, as of autumn 2013, the level of coverage of English chemotherapy units and the quality of the data coming from them has exceeded expectations. The SACT dataset includes fields on drug regimen, dosage, number of cycles administered and treatment intent. It also covers first and all subsequent lines of treatment. The NLCA intends to link data from the NLCA to these data as soon as is feasible and hope to be able to enhance the analyses of chemotherapy and other anti-cancer drug therapies in future reports.

Trust and Health Board Performance

Handling of Low Case Numbers

It should be noted that Trusts submitting very low numbers of cases with high levels of data completeness have been omitted from all Tables to ensure that no details about specific patients can be identified in this report. Because of this Network totals may not equal the sum of the composite Trusts. For example, in a Trust with only two submitted cases of lung cancer, with 100 per cent data completeness and a resection rate of 100 per cent, it would be possible to know the details of treatment of all lung cancer patients seen at that Trust. However, in most cases, each reported value is composed of multiple variables so it is impossible to surmise information about specific individuals from this report.

Data Groupings

The data has been divided into the following groups for analysis:

- NSCLC – non-small cell lung cancer or, more correctly, this should be considered NOT small cell lung cancer. This group includes all lung cancers including those that are clinically diagnosed, but excludes pathological diagnoses of small cell lung cancer and clinical/pathological diagnoses of mesothelioma.
- Histologically confirmed non-small cell lung cancer – all cases of non-small cell lung cancer that are confirmed by a histological or cytological specimen.
- Small cell lung cancer – all cases of lung cancer that are confirmed to be of small cell type by a histological or cytological specimen.

A separate report on mesothelioma will follow in due course.

Focus on Histological Diagnosis



Focus on Histological Diagnosis

In recent years, a full pathological (histological) diagnosis of lung cancer has become increasingly important, with the realisation that therapies have different activity dependent upon the histological subtype and the presence or absence of molecular markers. Consequently, measures of the adequacy of histological diagnosis are considered markers of the quality of care.

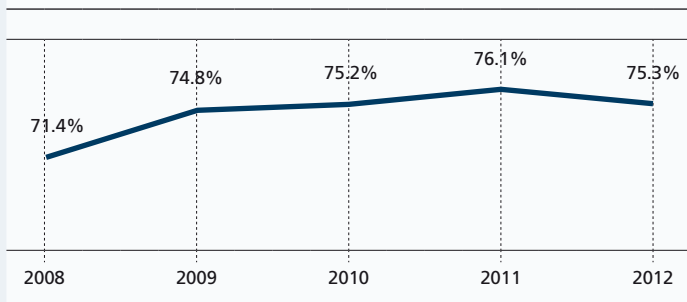
Histological Confirmation Rate (HCR) is the proportion of patients that have a diagnosis of lung cancer confirmed on a histological (tissue) or cytological (cellular) specimen rather than based on clinical or radiological features only. This includes patients who only have their cancer histologically confirmed at the time of their surgical treatment. The optimum HCR is unknown and will only rarely be 100 per cent since for some patients, the risks or burden of a biopsy cannot be justified. Evaluating the HCR in subgroups of patients or as an "odds ratio" adjusted for clinical features may therefore provide more useful information and will be included in the online audit data reports.

Focus on Histological Diagnosis

For patients first seen in 2012, the mean HCR for England and Wales was 75.3 per cent, similar to the previous year, but significantly higher than earlier years, reflecting the increasing importance of histological subtyping and the availability of low toxicity targeted therapies (See [Focus: Chart 1](#)).

The equivalent figure for 2012 for Scotland is 72.6 per cent.

Focus: Chart 1
Histocytological Confirmation Rate by year (England and Wales)



Focus: Chart 2
HCR for Cancer Networks in England and Wales

Network	2012 HCR	Trend
N01	71.2%	
N02	68.9%	
N03	72.6%	
N06	73.3%	
N07	66.3%	
N08	70.9%	
N11	76.2%	
N12	76.6%	
N20	71.6%	
N21	76.8%	
N22	83.1%	
N23	81.0%	
N24	84.1%	
N25	85.7%	
N26	70.9%	
N27	70.4%	
N28	74.4%	
N29	76.8%	
N30	83.7%	
N31	77.9%	
N32	83.7%	
N33	76.2%	
N34	77.1%	
N35	75.0%	
N36	74.4%	
N37	78.5%	
N38	85.1%	
N39	75.3%	
NWW	75.0%	
SWCN	71.5%	

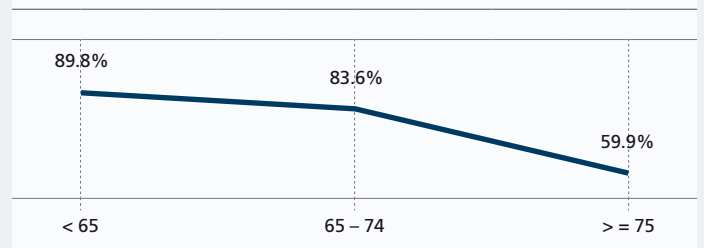
Making Sense of the Data

The Table ([Focus: Chart 2](#)) shows the overall HCR for Cancer Networks in England and Wales. Each result is made up of the results from individual hospitals in that region. A high HCR may indicate a more thorough approach to diagnosis of a lung cancer, although some patients may decline to have a biopsy, and for some patients a biopsy cannot be justified, so too high an HCR may also require further evaluation. Results for individual hospitals are available in the online audit reports.

The Table also demonstrates the trend in HCR rates for these Networks over the past five years, indicating whether the situation is improving or worsening compared to the national average results.

Clinical features of the patients in hospital will have an impact upon the HCR, since for older and less fit patients, the utility of a biopsy may be less. The graph ([Focus: Chart 3](#)) demonstrates the impact age has on HCR rates. Organisations looking to improve their HCR should look in detail at their approach to older patients as a means of improvement.

Focus: Chart 3
HCR by age group (England and Wales 2012)



Focus on Histological Diagnosis

LungPATH Project

The LungPATH Project examined the diagnostic and pathology processes of 19 hospitals. Each hospital reported the availability of investigations, waiting times and Pathologist workload. Data was collected on lung cancer patients seen over a six month period. Pathology reports for each invasive investigation were used to detect histological subtyping, and if immunohistochemistry or EGFR testing was used. Wide disparity in the availability of CT scanning, PET scanning and EBUS was found. There was a link between a tests availability and how likely it was to be used; patients were many times more likely to receive a PET scan or EBUS in some centres than others. The Project recommended maximum waiting times for CT scans of one week and two weeks for PET-CT and EBUS.

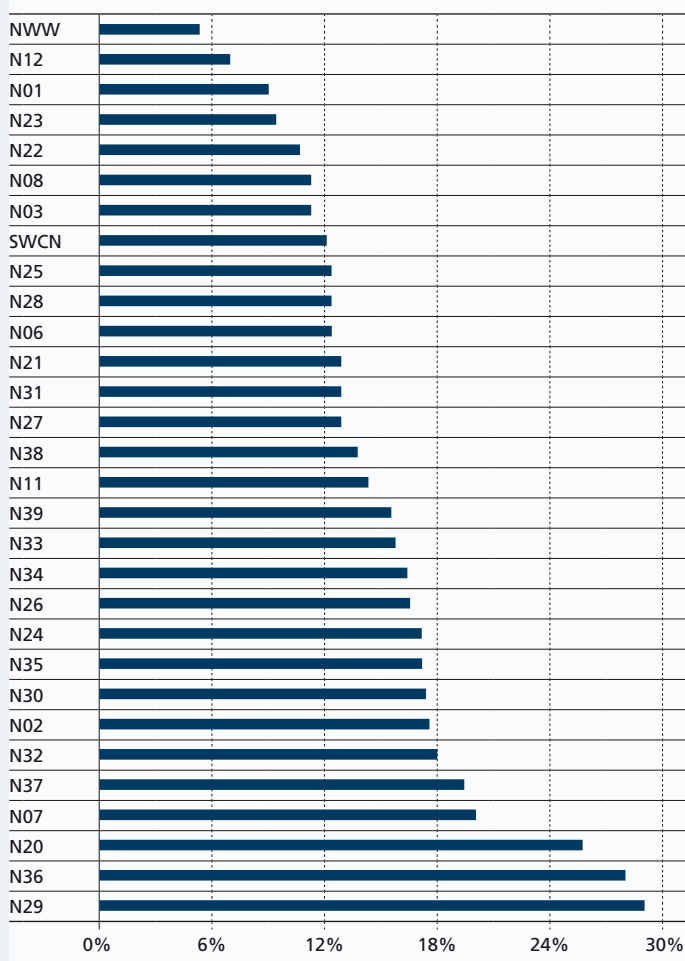
There was variation in the lung cancer workload Pathologists had in different centres, (1-14 hours per week). There was overuse of immunohistochemistry causing concerns around misuse of the tumour present, supported by the high EGFR test failure rate in some centres. The project recommended sub-specialisation of Pathologists within centres to enable concentration of workload, development of expertise and identified a need for education on immunohistochemistry use in lung cancer samples.

The full Project Report can be seen at:
www.hqip.org.uk/ncapop-library

NSCLC NOS

Most cases of non-small cell lung cancer can be further classified to adenocarcinoma, squamous carcinoma or other subtype using a range of techniques. This subtyping has an important impact upon the choice of chemotherapy, and so those patients whose tumours cannot be sub-classified (referred to as "Not Otherwise Specified") may not receive the most appropriate chemotherapy. Keeping this "NOS Rate" low will help to deliver best treatment for patients (See [Focus: Chart 4](#)).

Focus: Chart 4
NSCLC NOS



Tertiary Trusts

Most activity relating to lung cancer initial diagnosis occurs in the secondary care Trusts which range from small district general hospitals, to large teaching hospitals. Subsequent treatment often take place in the same Trust, or for some smaller Trusts, the patient may be transferred to another secondary care organisation. Activity in these organisations is well represented by the audit since the analysis of cases by "place first seen" allocates patients to the decision making Multi-Disciplinary Team.

However, there are several Tertiary Trusts which do not provide diagnostic services and which are therefore only rarely the "place first seen". These Trusts do provide a very important treatment service for patients both in their local area, and also on a regional/national basis, and for this reason we have chosen to record their activity separately, as shown in the Table below (Figure 10). Due to the absence of a common denominator, it is not possible to compare outcomes in these organisations at the present time.

Figure 10
Tertiary Trusts

Trust Code	Trust Name	Surgery (n)	Chemotherapy (n)	Radiotherapy including Brachytherapy (n)	Any (n)
RBV	The Christie NHS Foundation Trust	0	435	793	991
REN	The Clatterbridge Cancer Centre NHS Foundation Trust	0	557	610	872
RGM	Papworth Hospital NHS Foundation Trust	99	0	0	99
RM2	University Hospital of South Manchester NHS Foundation Trust	394	305	2	585
RPY	The Royal Marsden NHS Foundation Trust	0	167	195	327
RT3	Royal Brompton and Harefield NHS Foundation Trust	331	0	0	331

Data Completeness 2012 – Key Data Items

Data completeness reports for 2012 are available online at www.hscic.gov.uk/lung

Process and Outcome

Date first seen = 2012 (all). Date of extract = 08/07/2013 (England) & 28/06/2013 (Wales)

Table 1
Process and Clinical Outcomes for England and Wales 2012

Code	Actual number	Discussed at MDT (%)	Histological diagnosis (%)	Patient seen by Nurse Specialist (%)	Nurse Specialist present at diagnosis (%)	% Having active treatment	% of patients receiving CT before bronchoscopy	% receiving surgery all cases	% receiving radiotherapy
N01 Total	1,086 ●	95.3 ●	71.2 ▲	82.3 ●	60.1 ▲	61.2 ●	94.5 ▲	9.9	26.2
RTX	251 ●	96.8 ●	62.2 ▲	61.4 ▲	4.0 ▲	65.7 ●	93.4 ▲	10.0	29.1
RXL	276 ●	98.9 ●	77.2 ●	92.0 ●	85.1 ●	65.2 ●	94.3 ▲	6.2	19.2
RXN	244 ●	90.6 ▲	79.1 ●	84.4 ●	55.3 ▲	58.6 ▲	93.1 ▲	11.9	34.4
RXR	315 ■	94.6 ▲	67.0 ▲	88.9 ●	86.7 ●	56.2 ▲	96.2 ●	11.7	23.5
N02 Total	2,302 ●	91.5 ▲	68.9 ▲	82.3 ●	63.0 ▲	60.9 ●	91.3 ▲	17.0	33.1
RBT	122 ●	78.7 ▲	65.6 ▲	75.4 ▲	14.8 ▲	63.1 ●	71.2 ▲	21.3	35.2
RJN	135 ●	100.0 ●	67.4 ▲	98.5 ●	91.1 ●	61.5 ●	95.4 ●	14.8	35.6
RM2	255 ◆	90.2 ◆	80.8 ◆	80.0 ◆	27.1 ◆	66.7 ◆	97.9 ◆	24.3 ◆	25.5 ◆
RM3	196 ●	96.4 ●	75.0 ●	86.2 ●	71.4 ▲	61.7 ●	93.4 ▲	20.4	36.2
RMC	217 ●	86.6 ▲	53.9 ▲	81.1 ●	62.2 ▲	59.0 ▲	95.5 ●	16.1	35.0
RMP	161 ●	95.0 ●	67.7 ▲	83.9 ●	51.6 ▲	62.1 ●	96.4 ●	13.7	38.5
RRF	253 ●	94.9 ▲	71.9 ▲	80.2 ●	77.5 ▲	63.2 ●	87.5 ▲	19.0	35.2
RW3	226 ■	93.8 ▲	80.1 ●	88.9 ●	70.4 ▲	57.1 ▲	89.7 ▲	17.7	29.2
RW6	534 ●	86.9 ▲	62.9 ▲	74.9 ▲	71.2 ▲	59.0 ▲	88.9 ▲	13.9	31.5
RWJ	202 ●	98.0 ●	67.3 ▲	90.1 ●	72.8 ▲	59.4 ▲	90.3 ▲	11.9	36.1
N03 Total	1,849 ●	94.2 ▲	72.6 ▲	84.2 ●	48.4 ▲	61.0 ●	90.0 ▲	20.6	30.8
LLCU	420 ●	96.7 ●	81.9 ●	96.0 ●	77.6 ▲	73.1 ●	97.1 ●	27.4	36.7
RBL	315 ●	99.0 ●	67.6 ▲	90.5 ●	61.0 ▲	61.3 ●	89.2 ▲	19.4	34.9
RBN	249 ●	96.0 ●	75.9 ●	76.7 ▲	23.7 ▲	52.2 ▲	85.7 ▲	19.3	18.1
REM	327 ●	95.1 ●	72.2 ▲	74.0 ▲	3.7 ▲	64.5 ●	92.0 ▲	23.2	34.6
RJR	196 ●	87.8 ▲	62.8 ▲	71.4 ▲	36.7 ▲	51.5 ▲	75.3 ▲	14.8	25.0
RVY	156 ●	86.5 ▲	69.9 ▲	90.4 ●	71.2 ▲	59.0 ▲	95.4 ●	17.9	31.4
RWW	181 ●	91.2 ▲	68.5 ▲	84.0 ●	66.3 ▲	49.7 ▲	88.6 ▲	11.0	27.1
N06 Total	2,015 ●	98.9 ●	73.3 ▲	88.9 ●	70.2 ▲	63.9 ●	93.1 ▲	17.1	36.8
RAE	266 ●	99.2 ●	60.5 ▲	89.8 ●	70.7 ▲	60.9 ●	98.1 ●	17.7	35.3
RCB	194 ●	99.0 ●	72.2 ▲	89.2 ●	89.2 ●	57.7 ▲	95.9 ●	16.5	33.5
RCD	95 ●	98.9 ●	77.9 ●	88.4 ●	77.9 ▲	68.4 ●	100.0 ●	14.7	27.4
RCF	145 ●	100.0 ●	71.7 ▲	93.8 ●	81.4 ●	57.9 ▲	100.0 ●	15.2	25.5
RR8	608 ●	99.0 ●	75.2 ●	86.2 ●	53.5 ▲	69.2 ●	88.3 ▲	17.3	44.9
RWY	251 ●	100.0 ●	78.5 ●	87.6 ●	61.4 ▲	69.7 ●	92.0 ▲	16.7	36.7
RXF	456 ●	97.4 ●	75.4 ●	91.0 ●	84.0 ●	58.8 ▲	93.5 ▲	18.0	33.8
N07 Total	790 ●	95.6 ●	66.3 ▲	78.9 ▲	65.2 ▲	59.4 ▲	84.0 ▲	16.5	23.8
RCC	108 ●	93.5 ▲	55.6 ▲	88.9 ●	74.1 ▲	50.9 ▲	78.1 ▲	9.3	25.9
RJL	272 ●	95.2 ●	65.8 ▲	86.8 ●	75.4 ▲	58.1 ▲	83.8 ▲	16.5	26.8
RV9	10	80.0 ▲	60.0 ▲	100.0 ●	70.0 ▲	80.0 ●	100.0 ●	0.0	60.0
RWA	400 ●	96.8 ●	69.8 ▲	70.3 ▲	55.8 ▲	62.0 ●	84.9 ▲	18.8	20.3
N08 Total	1,285 ●	99.1 ●	70.9 ▲	85.8 ●	71.3 ▲	57.6 ▲	92.4 ▲	14.9	19.0
RFF	163 ●	98.8 ●	77.3 ●	95.7 ●	81.6 ●	64.4 ●	93.8 ▲	14.1	7.4
RFR	192 ●	95.3 ●	68.8 ▲	94.3 ●	77.6 ▲	53.1 ▲	100.0 ●	12.5	9.9
RFS	172 ●	100.0 ●	69.8 ▲	84.9 ●	71.5 ▲	45.9 ▲	86.2 ▲	10.5	8.1
RHQ	423 ●	100.0 ●	68.8 ▲	79.9 ▲	70.4 ▲	56.5 ▲	94.5 ▲	19.6	23.2
RP5	335 ●	99.7 ●	72.2 ▲	83.9 ●	63.6 ▲	64.2 ●	88.8 ▲	13.1	30.1

Table 1 (continued)
Process and Clinical Outcomes for England and Wales 2012

Code	Actual number	Discussed at MDT (%)	Histological diagnosis (%)	Patient seen by Nurse Specialist (%)	Nurse Specialist present at diagnosis (%)	% Having active treatment	% of patients receiving CT before bronchoscopy	% receiving surgery all cases	% receiving radiotherapy
N11 Total	1,124 ●	95.6 ●	76.2 ●	90.3 ●	67.1 ▲	54.4 ▲	92.7 ▲	15.3	18.7
RBK	153 ●	98.0 ●	83.0 ●	93.5 ●	77.8 ▲	58.2 ▲	96.1 ●	13.7	19.6
RR1	455 ●	91.9 ▲	66.4 ▲	90.3 ●	66.4 ▲	45.9 ▲	85.7 ▲	13.2	9.7
RRK	265 ●	97.0 ●	85.7 ●	91.3 ●	66.8 ▲	60.4 ●	91.4 ▲	20.4	24.2
RXK	251 ●	99.2 ●	79.7 ●	87.3 ●	62.2 ▲	61.0 ●	91.5 ▲	14.7	28.7
N12 Total	505 ●	94.5 ▲	76.6 ●	73.3 ▲	59.4 ▲	57.8 ▲	79.1 ▲	17.2	18.4
RJC	90 ●	88.9 ▲	72.2 ▲	83.3 ●	71.1 ▲	48.9 ▲	78.3 ▲	12.2	12.2
RKB	209 ●	97.1 ●	75.6 ●	83.7 ●	72.2 ▲	60.8 ●	82.4 ▲	18.7	23.4
RLT	126 ●	92.9 ▲	78.6 ●	82.5 ●	63.5 ▲	57.1 ▲	87.1 ▲	14.3	16.7
RWPO0	80 ●	96.3 ●	81.3 ●	20.0 ▲	6.3 ▲	61.3 ●	60.0 ▲	23.8	15.0
N20 Total	585 ●	98.3 ●	71.6 ▲	81.5 ●	60.0 ▲	47.7 ▲	83.4 ▲	12.8	18.3
RC9	213 ●	99.1 ●	71.4 ▲	77.5 ▲	47.9 ▲	46.5 ▲	77.9 ▲	17.4	12.7
RWG	193 ●	97.4 ●	71.0 ▲	88.6 ●	67.9 ▲	38.3 ▲	81.7 ▲	8.3	16.6
RWH	179 ●	98.3 ●	72.6 ▲	78.8 ▲	65.9 ▲	59.2 ▲	91.4 ▲	12.3	26.8
N21 Total	797 ●	95.5 ●	76.8 ●	77.0 ▲	62.1 ▲	60.5 ●	90.7 ▲	15.4	29.9
RAS	131 ●	87.0 ▲	61.1 ▲	86.3 ●	81.7 ●	46.6 ▲	84.2 ▲	12.2	22.9
RC3	67 ●	100.0 ●	55.2 ▲	83.6 ●	80.6 ●	55.2 ▲	100.0 ●	10.4	25.4
RFW	124 ●	97.6 ●	64.5 ▲	51.6 ▲	24.2 ▲	50.0 ▲	92.7 ▲	10.5	33.9
RQM	70 ●	94.3 ▲	87.1 ●	72.9 ▲	42.9 ▲	61.4 ●	100.0 ●	14.3	28.6
RT3	22 ◆	86.4 ◆	90.9 ◆	90.9 ◆	31.8 ◆	86.4 ◆	16.7 ◆	63.6 ◆	22.7 ◆
RV8	106 ●	99.1 ●	81.1 ●	91.5 ●	74.5 ▲	54.7 ▲	88.1 ▲	21.7	7.5
RYJ	277 ●	97.1 ●	89.5 ●	76.9 ▲	67.9 ▲	72.9 ●	94.4 ▲	14.4	41.9
N22 Total	781 ●	98.5 ●	83.1 ●	96.8 ●	86.0 ●	67.2 ●	83.3 ▲	15.7	33.5
RAL	88 ●	97.7 ●	95.5 ●	100.0 ●	98.9 ●	65.9 ●	89.5 ▲	23.9	25.0
RAP	101 ●	99.0 ●	80.2 ●	98.0 ●	94.1 ●	62.4 ●	95.8 ●	9.9	40.6
RKE	87 ●	97.7 ●	71.3 ▲	93.1 ●	74.7 ▲	64.4 ●	71.4 ▲	11.5	37.9
RQW	143 ●	98.6 ●	86.7 ●	98.6 ●	91.6 ●	69.2 ●	86.8 ▲	15.4	42.7
RRV	112 ●	100.0 ●	91.1 ●	92.9 ●	55.4 ▲	88.4 ●	90.9 ▲	19.6	39.3
RVL	247 ●	98.4 ●	78.1 ●	97.6 ●	93.9 ●	59.5 ▲	71.7 ▲	15.0	24.7
N23 Total	709 ●	97.3 ●	81.0 ●	82.9 ●	43.3 ▲	56.1 ▲	82.5 ▲	12.3	20.5
R1HKH	102 ●	99.0 ●	87.3 ●	65.7 ▲	48.0 ▲	59.8 ▲	95.0 ●	19.6	13.7
R1HMO	112 ●	100.0 ●	88.4 ●	98.2 ●	90.2 ●	63.4 ●	94.9 ▲	20.5	20.5
R1HNNH	86 ■	96.5 ●	68.6 ▲	87.2 ●	67.4 ▲	37.2 ▲	78.6 ▲	10.5	7.0
RF4	311 ●	95.2 ●	83.3 ●	83.6 ●	11.6 ▲	62.1 ●	69.9 ▲	7.4	30.9
RQX	98 ●	100.0 ●	69.4 ▲	77.6 ▲	64.3 ▲	41.8 ▲	95.6 ●	12.2	6.1
N24 Total	719 ●	96.9 ●	84.1 ●	92.1 ●	79.3 ▲	62.4 ●	85.1 ▲	12.7	31.0
RJ1	125 ▲	97.6 ●	95.2 ●	90.4 ●	73.6 ▲	81.6 ●	90.7 ▲	20.8	43.2
RJ2	104 ●	97.1 ●	81.7 ●	89.4 ●	46.2 ▲	59.6 ▲	68.2 ▲	13.5	23.1
RJZ	118 ●	100.0 ●	87.3 ●	78.8 ▲	78.8 ▲	63.6 ●	89.3 ▲	13.6	34.7
RYQ	372 ●	95.7 ●	80.1 ●	97.6 ●	90.6 ●	56.5 ▲	88.8 ▲	9.4	28.0
N25 Total	588 ■	94.0 ▲	85.7 ●	83.3 ●	47.3 ▲	62.4 ●	80.5 ▲	12.1	28.2
RAX	115 ■	96.5 ●	85.2 ●	87.8 ●	59.1 ▲	63.5 ●	96.9 ●	14.8	25.2
RJ6	133 ●	94.7 ▲	83.5 ●	96.2 ●	95.5 ●	54.9 ▲	94.7 ▲	9.0	27.8
RJ7	141 ■	96.5 ●	87.2 ●	79.4 ▲	4.3 ▲	72.3 ●	73.9 ▲	17.7	29.8
RVR	198 ●	90.4 ▲	86.4 ●	74.7 ▲	38.9 ▲	59.6 ▲	61.1 ▲	8.6	28.8

Table 1 (continued)
Process and Clinical Outcomes for England and Wales 2012

Code	Actual number	Discussed at MDT (%)	Histological diagnosis (%)	Patient seen by Nurse Specialist (%)	Nurse Specialist present at diagnosis (%)	% Having active treatment	% of patients receiving CT before bronchoscopy	% receiving surgery all cases	% receiving radiotherapy
N26 Total	1,056 ●	93.1 ▲	70.9 ▲	84.2 ●	72.1 ▲	61.9 ●	91.5 ▲	13.4	36.6
RA9	186 ●	94.1 ▲	76.3 ●	82.8 ●	64.5 ▲	64.5 ●	92.4 ▲	16.7	39.2
RBZ	96 ●	93.8 ▲	57.3 ▲	93.8 ●	61.5 ▲	39.6 ▲	91.7 ▲	10.4	6.3
REF	247 ●	96.8 ●	75.7 ●	94.7 ●	84.6 ●	70.0 ●	100.0 ●	13.0	45.3
RH8	221 ●	99.5 ●	85.5 ●	91.4 ●	83.7 ●	71.5 ●	86.5 ▲	16.7	47.1
RK9	306 ●	84.6 ▲	57.5 ▲	68.3 ▲	61.4 ▲	53.9 ▲	88.0 ▲	10.5	29.7
N27 Total	476 ●	95.6 ●	70.4 ▲	86.6 ●	61.8 ▲	55.3 ▲	83.5 ▲	13.4	23.3
RBD	122 ●	98.4 ●	68.9 ▲	91.0 ●	36.9 ▲	50.8 ▲	83.3 ▲	11.5	18.9
RD3	148 ●	92.6 ▲	76.4 ●	81.1 ●	46.6 ▲	64.2 ●	73.6 ▲	18.2	31.1
RDZ	206 ●	96.1 ●	67.0 ▲	87.9 ●	87.4 ●	51.5 ▲	90.1 ▲	11.2	20.4
N28 Total	938 ●	90.7 ▲	74.4 ▲	69.5 ▲	38.8 ▲	62.8 ●	90.4 ▲	18.0	28.5
RA3	89 ●	91.0 ▲	88.8 ●	73.0 ▲	47.2 ▲	66.3 ●	97.8 ●	19.1	22.5
RA4	92 ●	88.0 ▲	62.0 ▲	77.2 ▲	66.3 ▲	62.0 ●	85.7 ▲	17.4	15.2
RA7	144 ●	96.5 ●	74.3 ▲	73.6 ▲	24.3 ▲	62.5 ●	87.7 ▲	16.7	30.6
RBA	196 ●	94.4 ▲	63.8 ▲	70.9 ▲	51.5 ▲	54.6 ▲	84.4 ▲	15.8	28.1
RD1	202 ●	82.7 ▲	74.8 ▲	76.7 ▲	30.7 ▲	61.4 ●	95.2 ●	19.3	31.2
RVJ	215 ●	92.1 ▲	83.3 ●	54.0 ▲	29.3 ▲	70.7 ●	87.2 ▲	19.5	33.0
N29 Total	538 ●	95.0 ●	76.8 ●	69.9 ▲	56.5 ▲	57.8 ▲	82.6 ▲	12.5	31.0
RLQ	107 ●	100.0 ●	75.7 ●	69.2 ▲	58.9 ▲	57.0 ▲	100.0 ●	15.0	30.8
RTE	298 ●	95.0 ●	76.2 ●	77.9 ▲	57.7 ▲	58.4 ▲	80.1 ▲	13.8	32.2
RWV50	133 ●	91.0 ▲	78.9 ●	52.6 ▲	51.9 ▲	57.1 ▲	68.3 ▲	7.5	28.6
N30 Total	1,075 ●	96.2 ●	83.7 ●	85.9 ●	64.0 ▲	66.7 ●	92.9 ▲	21.4	27.8
RD7	151 ●	89.4 ▲	82.8 ●	68.9 ▲	29.1 ▲	44.4 ▲	81.6 ▲	12.6	16.6
RD8	103 ●	84.5 ▲	89.3 ●	76.7 ▲	49.5 ▲	76.7 ●	87.8 ▲	30.1	39.8
RHW	182 ●	97.3 ●	74.2 ▲	86.3 ●	44.5 ▲	63.7 ●	98.4 ●	10.4	46.7
RN3	158 ●	100.0 ●	77.8 ●	91.8 ●	74.1 ▲	57.6 ▲	95.7 ●	17.1	15.8
RTH	306 ●	99.3 ●	88.6 ●	92.8 ●	85.0 ●	77.1 ●	95.0 ●	30.7	21.6
RXQ	172 ●	99.4 ●	88.4 ●	88.4 ●	77.9 ▲	72.7 ●	93.7 ▲	23.3	31.4
N31 Total	1,061 ●	97.5 ●	77.9 ●	74.5 ▲	53.7 ▲	59.9 ▲	90.6 ▲	15.7	27.2
R1F	101 ●	99.0 ●	78.2 ●	94.1 ●	93.1 ●	61.4 ●	87.0 ▲	11.9	42.6
RHM	214 ●	98.6 ●	72.9 ▲	50.5 ▲	22.9 ▲	74.3 ●	87.5 ▲	18.7	45.8
RHU	332 ●	97.3 ●	80.1 ●	71.4 ▲	68.7 ▲	45.2 ▲	96.5 ●	17.2	5.7
RN506	66 ●	92.4 ▲	86.4 ●	74.2 ▲	31.8 ▲	57.6 ▲	68.0 ▲	19.7	24.2
RN541	90 ●	97.8 ●	78.9 ●	88.9 ●	54.4 ▲	66.7 ●	96.2 ●	7.8	32.2
RNZ	107 ●	99.1 ●	81.3 ●	89.7 ●	45.8 ▲	69.2 ●	95.3 ●	17.8	32.7
RYR16	151 ●	96.0 ●	73.5 ▲	82.8 ●	53.0 ▲	61.6 ●	89.5 ▲	12.6	32.5
N32 Total	588 ●	92.5 ▲	83.7 ●	85.9 ●	57.7 ▲	57.8 ▲	94.1 ▲	15.0	20.7
RA2	98 ●	82.7 ▲	94.9 ●	76.5 ▲	23.5 ▲	59.2 ▲	94.7 ▲	10.2	28.6
RDU	176 ●	84.7 ▲	79.5 ●	89.2 ●	51.1 ▲	59.1 ▲	96.1 ●	18.2	27.8
RTK	135 ●	100.0 ●	78.5 ●	87.4 ●	75.6 ▲	45.9 ▲	86.2 ▲	14.1	8.1
RTP	179 ●	100.0 ●	85.5 ●	86.6 ●	69.3 ▲	64.8 ●	95.3 ●	15.1	19.0
N33 Total	698 ●	90.4 ▲	76.2 ●	83.7 ●	59.6 ▲	56.6 ▲	90.4 ▲	10.0	32.1
RXC	256 ●	98.0 ●	77.3 ●	85.5 ●	62.9 ▲	57.4 ▲	90.7 ▲	8.6	31.6
RXH	263 ●	82.1 ▲	76.0 ●	77.9 ▲	51.0 ▲	62.0 ●	98.0 ●	13.3	36.9
RYR18	179 ●	91.6 ▲	74.9 ▲	89.4 ●	67.6 ▲	47.5 ▲	80.5 ▲	7.3	25.7

Table 1 (continued)
Process and Clinical Outcomes for England and Wales 2012

Code	Actual number	Discussed at MDT (%)	Histological diagnosis (%)	Patient seen by Nurse Specialist (%)	Nurse Specialist present at diagnosis (%)	% Having active treatment	% of patients receiving CT before bronchoscopy	% receiving surgery all cases	% receiving radiotherapy
N34 Total	948 ●	91.8 ▲	77.1 ●	34.0 ▲	30.2 ▲	59.6 ▲	89.0 ▲	13.7	35.1
RN7	113 ●	100.0 ●	92.9 ●	98.2 ●	83.2 ●	75.2 ●	90.5 ▲	21.2	33.6
RPA	177 ●	98.3 ●	76.8 ●	8.5 ▲	8.5 ▲	55.9 ▲	93.7 ▲	15.3	34.5
RVV*	467 ●	83.9 ▲	69.0 ▲	4.3 ▲	4.1 ▲	51.8 ▲	88.0 ▲	8.8	34.5
RWF	191 ●	100.0 ●	88.0 ●	92.1 ●	82.7 ●	72.8 ●	82.1 ▲	19.9	38.2
N35 Total	1,182 ●	91.5 ▲	75.0 ●	85.3 ●	71.2 ▲	60.2 ●	87.6 ▲	18.5	30.1
RJD	135 ●	96.3 ●	81.5 ●	89.6 ●	65.9 ▲	66.7 ●	79.7 ▲	19.3	33.3
RJE	323 ●	86.1 ▲	68.1 ▲	80.5 ●	53.3 ▲	58.8 ▲	91.3 ▲	13.3	38.7
RL4	228 ●	100.0 ●	70.6 ▲	96.5 ●	95.6 ●	57.0 ▲	91.5 ▲	23.2	18.9
RNA	207 ●	84.1 ▲	80.2 ●	88.9 ●	74.4 ▲	59.4 ▲	90.4 ▲	18.8	20.8
RWP31	44 ●	81.8 ▲	90.9 ●	54.5 ▲	52.3 ▲	68.2 ●	78.3 ▲	20.5	29.5
RXW	245 ●	96.3 ●	77.1 ●	81.2 ●	75.9 ▲	60.4 ●	85.8 ▲	20.0	35.5
N36 Total	2,676 ●	98.8 ●	74.4 ▲	85.7 ●	66.9 ▲	61.4 ●	87.7 ▲	13.6	33.1
RE9	148 ●	98.6 ●	66.9 ▲	98.6 ●	90.5 ●	58.1 ▲	98.2 ●	14.2	29.7
RLN	296 ●	98.0 ●	81.1 ●	87.2 ●	64.9 ▲	65.9 ●	80.1 ▲	16.9	41.2
RNL	236 ●	98.7 ●	83.5 ●	82.6 ●	58.5 ▲	62.7 ●	90.1 ▲	17.8	34.7
RR7	208 ●	100.0 ●	69.7 ▲	96.2 ●	72.1 ▲	54.8 ▲	93.3 ▲	12.5	23.6
RTD	308 ●	98.1 ●	76.6 ●	84.4 ●	69.5 ▲	66.2 ●	84.7 ▲	15.9	33.8
RTF	348 ●	98.3 ●	72.7 ▲	86.8 ●	46.6 ▲	55.7 ▲	77.9 ▲	9.8	19.0
RTR	384 ●	100.0 ●	74.5 ▲	90.4 ●	87.8 ●	62.5 ●	94.0 ▲	12.2	35.4
RVW	307 ●	96.7 ●	73.0 ▲	88.6 ●	60.9 ▲	67.4 ●	93.6 ▲	12.4	47.6
RXP	441 ●	100.0 ●	70.5 ▲	70.7 ▲	62.4 ▲	58.0 ▲	88.9 ▲	13.2	30.8
N37 Total	1,552 ●	95.0 ●	78.5 ●	74.9 ▲	54.6 ▲	67.3 ●	92.4 ▲	12.8	39.7
RC1	111 ●	99.1 ●	85.6 ●	74.8 ▲	49.5 ▲	72.1 ●	86.2 ▲	15.3	36.9
RCX	153 ●	98.0 ●	88.2 ●	92.8 ●	66.7 ▲	71.9 ●	91.9 ▲	15.7	34.6
RGM	13 ◆	100.0 ◆	100.0 ◆	92.3 ◆	76.9 ◆	84.6 ◆	0.0 ◆	69.2 ◆	15.4 ◆
RGN	182 ●	100.0 ●	84.6 ●	88.5 ●	49.5 ▲	77.5 ●	91.7 ▲	15.4	55.5
RGP	163 ●	88.3 ▲	66.3 ▲	65.0 ▲	35.0 ▲	61.3 ●	97.1 ●	8.0	31.9
RGQ	162 ●	96.3 ●	77.2 ●	86.4 ●	63.0 ▲	80.2 ●	81.8 ▲	11.1	64.8
RGR	147 ●	94.6 ▲	79.6 ●	74.1 ▲	54.4 ▲	55.8 ▲	96.6 ●	10.2	25.2
RGT	233 ●	99.6 ●	90.6 ●	90.1 ●	89.7 ●	68.7 ●	100.0 ●	10.7	44.2
RM1	322 ●	87.9 ▲	64.9 ▲	59.6 ▲	43.5 ▲	59.0 ▲	97.7 ●	11.8	30.1
RQQ	66 ●	98.5 ●	78.8 ●	10.6 ▲	4.5 ▲	60.6 ●	90.0 ▲	16.7	37.9
N38 Total	827 ●	99.5 ●	85.1 ●	85.4 ●	69.5 ▲	60.2 ●	92.7 ▲	14.5	25.8
RAJ	219 ●	98.2 ●	80.4 ●	82.6 ●	71.7 ▲	56.6 ▲	93.3 ▲	17.8	16.4
RDD	203 ●	100.0 ●	89.7 ●	84.7 ●	66.5 ▲	60.6 ●	89.9 ▲	11.8	18.7
RDE	238 ●	100.0 ●	82.4 ●	84.0 ●	59.7 ▲	73.1 ●	100.0 ●	14.7	50.8
RQ8	167 ●	100.0 ●	89.8 ●	91.6 ●	84.4 ●	46.1 ▲	84.1 ▲	13.2	10.8
N39 Total	2,253 ●	98.0 ●	75.3 ●	85.4 ●	59.6 ▲	64.0 ●	93.1 ▲	18.4	30.2
RJF	137 ●	97.1 ●	78.8 ●	95.6 ●	86.1 ●	59.1 ▲	97.8 ●	19.7	19.0
RK5	207 ●	95.2 ●	81.6 ●	100.0 ●	97.6 ●	64.7 ●	88.4 ▲	15.0	26.6
RNQ	201 ●	93.5 ▲	66.7 ▲	90.5 ●	78.1 ▲	59.7 ▲	96.6 ●	15.4	25.9
RNS	177 ●	98.3 ●	63.3 ▲	92.1 ●	45.2 ▲	56.5 ▲	84.5 ▲	16.4	29.4
RTG	289 ●	98.6 ●	81.3 ●	81.0 ●	59.2 ▲	71.3 ●	97.6 ●	21.1	34.9
RWD	334 ●	99.7 ●	72.5 ▲	88.0 ●	36.5 ▲	59.3 ▲	88.6 ▲	18.6	24.3
RWE	474 ●	99.2 ●	67.1 ▲	81.2 ●	57.6 ▲	65.0 ●	95.3 ●	18.4	35.2
RX1	430 ●	98.8 ●	87.2 ●	75.8 ▲	50.7 ▲	67.7 ●	94.3 ▲	19.8	33.5
England Total	31,003 ●	95.6 ●	75.5 ●	81.9 ●	61.2 ▲	61.0 ●	89.5 ▲	15.5	29.6

* East Kent Hospitals University Foundation Trust (RVV) has had local issues with their system uploader, resulting in inaccuracies to their reported data.

Table 1 (continued)

Process and Clinical Outcomes for England and Wales 2012

Code	Actual number	Discussed at MDT (%)	Histological diagnosis (%)	Patient seen by Nurse Specialist (%)	Nurse Specialist present at diagnosis (%)	% Having active treatment	% of patients receiving CT before bronchoscopy	% receiving surgery all cases	% receiving radiotherapy
NWW Total	476 ●	99.2 ●	75.0 ●	93.3 ●	N/A	60.9 ●	90.8 ▲	12.0	38.0
7A1A1	193 ●	97.9 ●	79.8 ●	97.4 ●	N/A	63.7 ●	88.2 ▲	12.4	41.5
7A1A4	155 ●	100.0 ●	69.0 ▲	91.0 ●	N/A	57.4 ▲	92.3 ▲	16.1	34.2
7A1AU	128 ●	100.0 ●	75.0 ●	89.8 ●	N/A	60.9 ●	92.4 ▲	6.3	37.5
SWCN Total	1,556 ●	99.5 ●	71.5 ▲	86.9 ●	N/A	57.7 ▲	91.5 ▲	9.4	34.8
7A2AG	29 ▲	100.0 ●	65.5 ▲	51.7 ▲	N/A	48.3 ▲	66.7 ▲	3.4	20.7
7A2AJ	39 ●	100.0 ●	82.1 ●	12.8 ▲	N/A	61.5 ●	100.0 ●	5.1	23.1
7A2AL	117 ●	100.0 ●	86.3 ●	87.2 ●	N/A	73.5 ●	97.6 ●	13.7	23.1
7A2BL	90 ●	94.4 ▲	72.2 ▲	98.9 ●	N/A	51.1 ▲	90.0 ▲	15.6	18.9
7A3B7	99 ●	100.0 ●	63.6 ▲	80.8 ●	N/A	65.7 ●	64.9 ▲	7.1	49.5
7A3C4	90 ●	100.0 ●	73.3 ▲	92.2 ●	N/A	54.4 ▲	100.0 ●	7.8	33.3
7A3C7	89 ■	100.0 ●	76.4 ●	89.9 ●	N/A	60.7 ●	100.0 ●	6.7	32.6
7A3CJ	83 ●	100.0 ●	74.7 ▲	88.0 ●	N/A	61.4 ●	69.2 ▲	7.2	39.8
7A4C1	312 ●	100.0 ●	67.0 ▲	91.7 ●	N/A	52.6 ▲	97.6 ●	9.3	36.9
7A5B1	136 ●	98.5 ●	78.7 ●	84.6 ●	N/A	60.3 ●	93.5 ▲	12.5	41.9
7A5B3	114 ●	100.0 ●	75.4 ●	94.7 ●	N/A	75.4 ●	94.9 ▲	8.8	56.1
7A6AM	123 ●	100.0 ●	61.8 ▲	94.3 ●	N/A	49.6 ▲	93.3 ▲	8.1	33.3
7A6AR	235 ●	99.6 ●	67.2 ▲	85.1 ●	N/A	49.4 ▲	96.1 ●	8.9	27.7
Wales Total	2,032 ●	99.4 ●	72.3 ▲	88.4 ●	N/A	58.5 ▲	91.3 ▲	10.0	35.6
England and Wales Total	33,035 ●	95.8 ●	75.3 ●	82.3 ●	N/A	60.8 ●	89.6 ▲	15.2	30.0

Range Network

Min		90.4	66.3	34.0	30.2	47.7	79.1	9.4	18.3
LQ		94.1	71.9	79.6	56.0	57.8	84.3	12.7	24.3
Median		95.6	75.8	84.2	61.0	60.4	90.5	14.7	30.0
UQ		98.2	78.4	85.9	67.7	62.3	92.4	16.9	33.1
Max		99.5	85.7	96.8	86.0	67.3	94.5	21.4	39.7

Range Trust

Min		78.7	53.9	4.3	3.7	37.2	60.0	0.0	5.7
LQ		94.6	69.0	77.6	50.7	56.5	85.8	11.5	23.1
Median		97.7	76.2	86.8	65.9	60.6	91.3	14.7	30.1
UQ		99.4	81.7	91.0	77.5	65.0	95.4	17.9	35.3
Max		100.0	95.5	100.0	98.9	88.4	100.0	30.7	64.8

Last year's results (including mesothelioma)

2011 Total	33,374 ●	96.2 ●	77.0 ●	79.7 ▲	58.8 ▲	60.2 ●	87.8 ▲	14.7	29.3
Difference	-339	-0.4	-1.7	2.6	2.4	0.6	1.8	0.5	0.7

Last year's results (excluding mesothelioma)

2011 Total	31,557 ●	96.3 ●	76.1 ●	79.5 ▲	58.6 ▲	59.8 ▲	87.9 ▲	13.6	29.3
Difference	1,478	-0.5	-0.8	2.8	2.6	1.0	1.7	1.3	0.7

Counts aggregated by place first seen Trust.

Indicator	Definition	▲	■	●
Actual number	Number of cases (excluding mesothelioma) with date first seen in year specified	<50%	50-75%	≥75%
Discussed at MDT (%)	Complete when MDT discussion indicator = Y (denominator = all cases excluding mesothelioma)	<95%		≥95%
Histological diagnosis (%)	Complete when histology is present or basis of diagnosis equals 5, 6 or 7 (denominator = all cases excluding mesothelioma)	<75%		≥75%
Patient seen by Nurse Specialist (%)	Complete when patient assessed by a Lung Cancer Nurse Specialist = Y (denominator = all cases excluding mesothelioma)	<80%		≥80%
Nurse Specialist present at diagnosis (%)	Complete when Lung Cancer Nurse Specialist present when received diagnosis = Y (denominator = all English cases excluding mesothelioma)	<80%		≥80%
% having active treatment	Complete when date present for brachytherapy, anti-cancer drug regimen, surgery or teletherapy (denominator = all cases excluding mesothelioma)	<60%		≥60%
% of patients receiving CT scan before bronchoscopy	Complete when CT scan date before or equal to bronchoscopy date (denominator = cases excluding mesothelioma with bronchoscopy date present)	<95%		≥95%
% receiving surgery all cases	Complete when surgery procedure date is present (denominator = all cases excluding mesothelioma)			
% receiving radiotherapy	Complete when either teletherapy treatment course start date or brachytherapy therapy treatment course start date is present (denominator = all cases excluding mesothelioma)			
◆ – Tertiary Trust standards do not apply				

Table 2
Process and Clinical Outcomes for Guernsey 2012

Code	Actual number	Discussed at MDT (%)	Histological diagnosis (%)	Patient seen by Nurse Specialist (%)	Nurse Specialist present at diagnosis (%)	% Having active treatment	% of patients receiving CT before bronchoscopy	% receiving surgery all cases	% receiving radiotherapy
2012 Total	40 ●	100.0 ●	77.5 ●	N/A	40.0 ▲	42.5 ▲	100.0 ●	5.0	2.5
2011 Total	41 ●	100.0 ●	75.6 ●	N/A	65.9 ▲	56.1 ▲	94.7 ▲	7.3	19.5
Difference	-1	0	1.9	N/A	-25.9	-13.6	5.3	-2.3	-17

Table 3
Process and Clinical Outcomes for Scotland 2012

Health Board	Actual number (Total)	% of expected	Discussed at MDT (%)	Histological diagnosis (%)	Patient seen by Nurse Specialist (%)	% having active treatment	% of patients receiving CT before bronchoscopy	% receiving surgery all cases	% receiving radiotherapy	Number of histologically confirmed NSCLC	% histologically confirmed NSCLC having surgery	Number of patients small cell lung cancer	% small cell receiving chemotherapy
SCAN	1,219	90	96.6	67.0	81.9	55.9	96.7	12.1	35.3	628	21.0	147	70.1
Borders	86	86	100.0	75.6	94.2	60.5	100.0	9.3	36.0	53	11.3	10	90.0
Dumfries and Galloway	123	85	92.7	65.0	65.9	55.3	100.0	8.1	37.4	58	17.2	19	78.9
Fife	329	98	100.0	66.0	84.2	55.0	99.2	11.6	34.0	173	20.8	35	62.9
Lothian	681	88	95.3	66.8	82.1	55.8	93.7	13.5	35.4	344	23.3	83	68.7
WoSCAN	2,585	96	95.9	74.3	86.8	60.2	89.8	13.8	35.3	1,417	23.5	363	70.0
Ayrshire and Arran	371	106	97.8	76.0	79.2	63.3	90.3	12.7	44.7	201	20.9	58	70.7
Clyde	391	99	92.8	72.6	93.1	49.9	76.5	12.3	25.1	212	21.2	56	53.6
Forth Valley	215	84	98.6	74.0	94.4	68.4	97.0	13.5	40.5	119	24.4	34	64.7
Lanarkshire	495	90	98.8	79.4	90.9	64.4	87.8	16.4	31.9	298	24.2	69	71.0
North Glasgow	707	100	96.0	74.1	82.7	63.5	95.2	13.9	39.0	391	23.5	85	80.0
South Glasgow	406	91	92.1	68.7	86.0	52.0	92.8	13.5	31.5	196	27.0	61	72.1
NoSCAN	1,006	91	93.6	74.9	83.3	64.6	95.5	8.3	44.3	596	12.9	122	73.0
Grampian	357	84	86.1	74.3	71.0	69.2	94.0	8.8	51.5	218	14.2	42	69.0
Orkney	0	0											
Shetland	16	229											
Highland	220	101	95.8	78.9	82.4	61.7	99.0	6.1	37.9	160	8.8	36	83.3
Argyll and Clyde (H)	27	79											
Western Isles	14	117											
Tayside	372	92	99.7	72.6	96.2	62.1	95.0	9.1	41.7	218	14.7	44	68.2
Total	4,810	93.2	95.6	72.6	84.8	60.0	92.4	12.2	37.2	2,641	20.5	632	70.6
2011 Total	4,655	91.7	95.3	71.8	81.2	59.7	91.1	10.7	37.9	2,531	18.5	654	67.6
Difference	155	1.5	0.3	0.7	3.6	0.3	1.3	1.6	-0.7	110	2.0	-22	3.0

Range Health Board

Min		0	86.1	65.0	65.9	49.9	76.5	6.1	25.1		8.8		53.6
LQ		85	92.8	68.7	82.1	55.3	92.8	9.1	34.0		14.7		68.2
Median		91	96.0	74.0	84.2	61.7	95.0	12.3	37.4		20.9		70.7
UQ		100	98.8	75.6	93.1	63.5	99.0	13.5	40.5		23.5		78.9
Max		229	100.0	79.4	96.2	69.2	100.0	16.4	51.5		27.0		90.0

Table 4
Further Process and Clinical Outcomes for England and Wales 2012

Code	Actual number	NSCLC Stage IA, IB, IIA or IIB	% of NSCLC Stage IA, IB, IIA or IIB having surgery	Number of PS 0-1 NSCLC Stage IIIB or IV	% PS 0-1 Stage IIIB or IV NSCLC having chemotherapy	Number of histologically confirmed NSCLC	% histologically confirmed NSCLC having surgery	Number of pre-treatment NSCLC	% pre-treatment NSCLC histology NOS	Number of patients small cell lung cancer	% small cell receiving chemotherapy
N01 Total	1,086 ●	218	38.5 ▲	251	70.9 ●	627	16.1	624	9.1 ●	138	80.4 ●
RTX	251 ●	54	27.8 ▲	67	65.7 ●	125	17.6	125	12.8 ●	28	92.9 ●
RXL	276 ●	48	35.4 ▲	66	72.7 ●	176	9.7	174	8.6 ●	36	88.9 ●
RXN	244 ●	50	48.0 ▲	50	80.0 ●	157	17.2	157	6.4 ●	34	82.4 ●
RXR	315 ■	66	42.4 ▲	68	67.6 ●	169	20.7	168	9.5 ●	40	62.5 ▲
N02 Total	2,302 ●	571	45.5 ▲	440	51.1 ▲	1,320	27.0	1,284	17.6 ●	248	69.4 ●
RBT	122 ●	25	56.0 ●	13	53.8 ▲	69	34.8	70	18.6 ●	10	60.0 ▲
RJN	135 ●	35	37.1 ▲	29	72.4 ●	80	25.0	80	20.0 ●	11	90.9 ●
RM2	255 ◆	77 ◆	57.1 ◆	51 ◆	52.9 ◆	168 ◆	33.9 ◆	165 ◆	7.9 ◆	35 ◆	71.4 ◆
RM3	196 ●	47	55.3 ●	33	30.3 ▲	118	30.5	113	17.7 ●	27	63.0 ▲
RMC	217 ●	57	42.1 ▲	28	50.0 ▲	98	32.7	92	6.5 ●	17	70.6 ●
RMP	161 ●	28	42.9 ▲	28	64.3 ●	85	20.0	85	20.0 ●	21	81.0 ●
RRF	253 ●	59	54.2 ●	56	44.6 ▲	151	30.5	142	12.7 ●	29	62.1 ▲
RW3	226 ■	68	45.6 ▲	42	45.2 ▲	155	23.9	154	15.6 ●	25	64.0 ▲
RW6	534 ●	120	38.3 ▲	96	59.4 ●	276	24.3	276	26.8 ▲	58	72.4 ●
RWJ	202 ●	55	32.7 ▲	64	42.2 ▲	119	17.6	106	23.6 ▲	15	60.0 ▲
N03 Total	1,849 ●	480	60.2 ●	264	63.3 ●	1,112	30.8	1,104	11.4 ●	206	68.4 ●
LLCU	420 ●	136	69.9 ●	54	66.7 ●	282	37.9	280	6.1 ●	58	63.8 ▲
RBL	315 ●	77	61.0 ●	42	69.0 ●	175	30.3	171	12.9 ●	33	66.7 ●
RBN	249 ●	58	69.0 ●	27	59.3 ●	153	28.1	155	9.0 ●	33	78.8 ●
REM	327 ●	94	53.2 ●	55	58.2 ●	195	32.8	192	17.7 ●	35	74.3 ●
RJR	196 ●	50	46.0 ▲	30	46.7 ▲	103	28.2	103	15.5 ●	20	75.0 ●
RVY	156 ●	31	61.3 ●	25	76.0 ●	94	25.5	94	8.5 ●	9	77.8 ●
RWW	181 ●	33	42.4 ▲	30	70.0 ●	106	18.9	105	13.3 ●	18	44.4 ▲
N06 Total	2,015 ●	509	46.4 ▲	370	67.6 ●	1,206	26.0	1,158	12.5 ●	252	72.6 ●
RAE	266 ●	81	38.3 ▲	57	59.6 ●	133	32.3	132	11.4 ●	26	84.6 ●
RCB	194 ●	50	44.0 ▲	26	73.1 ●	124	25.0	123	16.3 ●	15	66.7 ●
RCD	95 ●	19	42.1 ▲	17	76.5 ●	62	19.4	54	13.0 ●	11	72.7 ●
RCF	145 ●	33	39.4 ▲	30	70.0 ●	83	22.9	81	12.3 ●	18	72.2 ●
RR8	608 ●	175	41.1 ▲	101	69.3 ●	376	25.3	376	10.4 ●	74	74.3 ●
RWY	251 ●	51	62.7 ●	68	73.5 ●	158	24.1	154	12.3 ●	38	81.6 ●
RXF	456 ●	100	58.0 ●	71	60.6 ●	270	28.1	238	14.7 ●	70	62.9 ▲
N07 Total	790 ●	160	60.6 ●	183	55.2 ●	450	23.8	359	20.1 ▲	73	69.9 ●
RCC	108 ●	15	46.7 ▲	15	66.7 ●	58	13.8	13	15.4 ●	2	0.0 ▲
RJL	272 ●	46	69.6 ●	73	53.4 ▲	156	28.2	136	11.8 ●	22	81.8 ●
RV9	10	0	0.0 ▲	1	100.0 ●	5	0.0	0	0.0 ●	1	100.0 ●
RWA	400 ●	99	58.6 ●	94	54.3 ▲	231	23.8	210	25.7 ▲	48	66.7 ●
N08 Total	1,285 ●	276	52.9 ●	271	63.8 ●	741	22.4	684	11.4 ●	160	63.8 ▲
RF	163 ●	26	57.7 ●	37	70.3 ●	98	20.4	86	20.9 ▲	28	67.9 ●
RF	192 ●	38	50.0 ▲	31	61.3 ●	92	13.0	85	9.4 ●	39	61.5 ▲
RFS	172 ●	37	35.1 ▲	43	48.8 ▲	93	16.1	80	15.0 ●	25	56.0 ▲
RHQ	423 ●	102	58.8 ●	73	79.5 ●	250	30.4	251	9.6 ●	34	67.6 ●
RP5	335 ●	73	53.4 ●	87	56.3 ●	208	20.7	182	8.8 ●	34	64.7 ▲
N11 Total	1,124 ●	227	52.9 ●	258	58.5 ●	728	21.6	717	14.5 ●	118	63.6 ▲
RBK	153 ●	21	52.4 ●	38	65.8 ●	102	17.6	99	19.2 ●	23	78.3 ●
RR1	455 ●	90	47.8 ▲	85	60.0 ●	260	21.2	252	9.9 ●	40	60.0 ▲
RRK	265 ●	68	60.3 ●	62	64.5 ●	195	25.1	195	15.9 ●	28	53.6 ▲
RXK	251 ●	48	52.1 ●	73	47.9 ▲	171	20.5	171	17.0 ●	27	66.7 ●

Table 4 (continued)
Further Process and Clinical Outcomes for England and Wales 2012

Code	Actual number	NSCLC Stage IA, IB, IIA or IIB	% of NSCLC Stage IA, IB, IIA or IIB having surgery	Number of PS 0-1 NSCLC Stage IIIB or IV	% PS 0-1 Stage IIIB or IV NSCLC having chemotherapy	Number of histologically confirmed NSCLC	% histologically confirmed NSCLC having surgery	Number of pre-treatment NSCLC	% pre-treatment NSCLC histology NOS	Number of patients small cell lung cancer	% small cell receiving chemotherapy
N12 Total	505 ●	100	59.0 ●	94	47.9 ▲	323	23.8	266	7.1 ●	60	56.7 ▲
RJC	90 ●	13	53.8 ●	16	56.3 ●	59	18.6	51	19.6 ●	6	66.7 ●
RKB	209 ●	37	67.6 ●	47	42.6 ▲	132	24.2	90	4.4 ●	24	50.0 ▲
RLT	126 ●	33	48.5 ▲	19	63.2 ●	81	21.0	74	4.1 ●	17	58.8 ▲
RWV00	80 ●	17	64.7 ●	12	33.3 ▲	51	33.3	51	3.9 ●	13	61.5 ▲
N20 Total	585 ●	82	35.4 ▲	104	48.1 ▲	344	18.0	332	25.9 ▲	69	49.3 ▲
RC9	213 ●	39	33.3 ▲	33	48.5 ▲	127	22.8	118	35.6 ▲	24	20.8 ▲
RWG	193 ●	15	20.0 ▲	30	23.3 ▲	113	12.4	110	9.1 ●	24	54.2 ▲
RWH	179 ●	28	46.4 ▲	41	65.9 ●	104	18.3	104	32.7 ▲	21	76.2 ●
N21 Total	797 ●	145	58.6 ●	206	51.9 ▲	525	21.0	461	13.0 ●	76	65.8 ●
RAS	131 ●	13	46.2 ▲	28	39.3 ▲	67	23.9	37	27.0 ▲	13	46.2 ▲
RC3	67 ●	15	33.3 ▲	20	40.0 ▲	35	20.0	34	5.9 ●	2	100.0 ●
RFW	124 ●	20	50.0 ▲	24	45.8 ▲	65	15.4	58	15.5 ●	12	66.7 ●
RQM	70 ●	15	60.0 ●	27	59.3 ●	56	17.9	54	7.4 ●	4	100.0 ●
RT3	22 ◆	12 ◆	75.0 ◆	1 ◆	100.0 ◆	17 ◆	76.5 ◆	8 ◆	12.5 ◆	2 ◆	100.0 ◆
RV8	106 ●	21	76.2 ●	24	54.2 ▲	73	26.0	66	13.6 ●	13	69.2 ●
RYJ	277 ●	49	61.2 ●	82	57.3 ●	212	16.5	204	12.3 ●	30	63.3 ▲
N22 Total	781 ●	131	59.5 ●	192	63.0 ●	556	19.6	524	10.7 ●	80	63.8 ▲
RAL	88 ●	14	85.7 ●	21	66.7 ●	71	21.1	65	6.2 ●	9	66.7 ●
RAP	101 ●	15	33.3 ▲	23	60.9 ●	72	13.9	66	27.3 ▲	9	44.4 ▲
RKE	87 ●	12	66.7 ●	26	50.0 ▲	49	18.4	49	4.1 ●	12	75.0 ●
RQW	143 ●	28	60.7 ●	13	69.2 ●	102	19.6	95	12.6 ●	20	55.0 ▲
RRV	112 ●	24	45.8 ▲	39	74.4 ●	90	20.0	89	9.0 ●	8	100.0 ●
RVL	247 ●	38	65.8 ●	68	58.8 ●	169	21.3	158	7.6 ●	22	59.1 ▲
N23 Total	709 ●	138	41.3 ▲	148	64.2 ●	490	15.9	461	9.5 ●	73	63.0 ▲
R1HKH	102 ●	19	36.8 ▲	21	52.4 ▲	74	21.6	68	7.4 ●	11	81.8 ●
R1HM0	112 ●	25	68.0 ●	15	73.3 ●	84	25.0	76	11.8 ●	13	76.9 ●
R1HNN	86 ■	14	42.9 ▲	15	40.0 ▲	53	15.1	38	5.3 ●	6	33.3 ▲
RF4	311 ●	55	34.5 ▲	75	70.7 ●	223	10.3	223	4.0 ●	33	60.6 ▲
RQX	98 ●	25	32.0 ▲	22	63.6 ●	56	17.9	56	33.9 ▲	10	50.0 ▲
N24 Total	719 ●	126	44.4 ▲	171	62.6 ●	525	15.8	504	17.3 ●	76	71.1 ●
RJ1	125 ▲	30	53.3 ●	34	82.4 ●	103	23.3	99	25.3 ▲	16	75.0 ●
RJ2	104 ●	13	61.5 ●	26	46.2 ▲	74	16.2	65	26.2 ▲	11	81.8 ●
RJZ	118 ●	20	60.0 ●	35	51.4 ▲	88	15.9	87	8.0 ●	12	58.3 ▲
RYQ	372 ●	63	31.7 ▲	76	64.5 ●	260	12.7	253	15.0 ●	37	70.3 ●
N25 Total	588 ■	112	47.3 ▲	137	59.1 ●	451	15.1	420	12.4 ●	49	67.3 ●
RAX	115 ■	23	60.9 ●	22	50.0 ▲	85	20.0	83	12.0 ●	13	69.2 ●
RJ6	133 ●	25	36.0 ▲	26	53.8 ▲	99	12.1	94	8.5 ●	12	66.7 ●
RJ7	141 ■	30	60.0 ●	40	65.0 ●	107	22.4	106	12.3 ●	13	69.2 ●
RVR	198 ●	33	36.4 ▲	49	61.2 ●	159	9.4	136	14.7 ●	11	63.6 ▲
N26 Total	1,056 ●	229	46.3 ▲	249	49.0 ▲	615	19.0	594	16.7 ●	113	73.5 ●
RA9	186 ●	40	50.0 ▲	44	52.3 ▲	117	20.5	107	3.7 ●	20	70.0 ●
RBZ	96 ●	19	36.8 ▲	26	42.3 ▲	49	16.3	44	54.5 ▲	6	66.7 ●
REF	247 ●	46	56.5 ●	63	36.5 ▲	153	18.3	153	16.3 ●	28	67.9 ●
RH8	221 ●	50	56.0 ●	55	56.4 ●	156	19.9	149	6.0 ●	29	86.2 ●
RK9	306 ●	74	33.8 ▲	61	55.7 ●	140	18.6	141	26.2 ▲	30	70.0 ●

Table 4 (continued)
Further Process and Clinical Outcomes for England and Wales 2012

Code	Actual number	NSCLC Stage IA, IB, IIA or IIB	% of NSCLC Stage IA, IB, IIA or IIB having surgery	Number of PS 0-1 NSCLC Stage IIIB or IV	% PS 0-1 Stage IIIB or IV NSCLC having chemotherapy	Number of histologically confirmed NSCLC	% histologically confirmed NSCLC having surgery	Number of pre-treatment NSCLC	% pre-treatment NSCLC histology NOS	Number of patients small cell lung cancer	% small cell receiving chemotherapy
N27 Total	476 ●	96	46.9 ▲	151	51.0 ▲	288	19.8	275	13.1 ●	42	66.7 ●
RBD	122 ●	29	31.0 ▲	31	61.3 ●	75	16.0	70	15.7 ●	7	57.1 ▲
RD3	148 ●	24	70.8 ●	49	53.1 ▲	99	24.2	94	7.4 ●	12	75.0 ●
RDZ	206 ●	43	44.2 ▲	71	45.1 ▲	114	18.4	111	16.2 ●	23	65.2 ●
N28 Total	938 ●	187	52.4 ●	198	66.2 ●	602	25.6	601	12.5 ●	82	72.0 ●
RA3	89 ●	12	41.7 ▲	19	78.9 ●	59	25.4	59	28.8 ▲	18	66.7 ●
RA4	92 ●	19	42.1 ▲	21	66.7 ●	51	29.4	51	27.5 ▲	4	100.0 ●
RA7	144 ●	32	46.9 ▲	24	87.5 ●	89	27.0	89	14.6 ●	18	66.7 ●
RBA	196 ●	36	50.0 ▲	38	63.2 ●	113	25.7	113	7.1 ●	10	70.0 ●
RD1	202 ●	43	48.8 ▲	50	52.0 ▲	125	28.0	125	4.0 ●	22	72.7 ●
RVJ	215 ●	45	68.9 ●	46	67.4 ●	165	21.8	164	11.0 ●	10	80.0 ●
N29 Total	538 ●	101	44.6 ▲	118	44.9 ▲	351	17.4	346	29.2 ▲	56	60.7 ▲
RLQ	107 ●	23	43.5 ▲	25	52.0 ▲	68	22.1	66	22.7 ▲	11	63.6 ▲
RTE	298 ●	60	46.7 ▲	71	36.6 ▲	193	19.2	190	40.0 ▲	30	50.0 ▲
RWP50	133 ●	18	38.9 ▲	22	63.6 ●	90	10.0	90	11.1 ●	15	80.0 ●
N30 Total	1,075 ●	211	62.1 ●	242	52.1 ▲	763	27.8	724	17.5 ●	125	69.6 ●
RD7	151 ●	16	50.0 ▲	25	20.0 ▲	114	16.7	82	30.5 ▲	11	27.3 ▲
RD8	103 ●	13	84.6 ●	14	71.4 ●	84	35.7	83	14.5 ●	8	62.5 ▲
RHW	182 ●	31	38.7 ▲	49	34.7 ▲	121	14.9	118	24.6 ▲	13	46.2 ▲
RN3	158 ●	31	58.1 ●	30	56.7 ●	98	23.5	98	13.3 ●	23	78.3 ●
RTH	306 ●	76	72.4 ●	80	61.3 ●	210	39.0	209	11.5 ●	52	78.8 ●
RXQ	172 ●	42	64.3 ●	44	63.6 ●	134	29.9	133	18.0 ●	18	77.8 ●
N31 Total	1,061 ●	191	60.2 ●	298	48.7 ▲	702	22.2	681	13.1 ●	120	72.5 ●
R1F	101 ●	9	66.7 ●	26	46.2 ▲	69	15.9	65	24.6 ▲	10	90.0 ●
RHM	214 ●	49	55.1 ●	46	54.3 ▲	131	28.2	130	6.9 ●	25	88.0 ●
RHU	332 ●	62	66.1 ●	97	39.2 ▲	230	24.3	226	13.3 ●	36	63.9 ▲
RN506	66 ●	15	60.0 ●	19	31.6 ▲	50	26.0	50	16.0 ●	7	42.9 ▲
RN541	90 ●	12	41.7 ▲	25	68.0 ●	59	10.2	58	15.5 ●	12	91.7 ●
RNZ	107 ●	20	70.0 ●	34	55.9 ●	73	23.3	67	9.0 ●	12	58.3 ▲
RYR16	151 ●	24	54.2 ●	51	54.9 ▲	90	17.8	85	12.9 ●	18	66.7 ●
N32 Total	588 ●	108	50.9 ▲	125	62.4 ●	418	18.4	415	18.1 ●	67	53.7 ▲
RA2	98 ●	13	46.2 ▲	3	66.7 ●	73	11.0	73	27.4 ▲	16	50.0 ▲
RDU	176 ●	36	61.1 ●	35	65.7 ●	114	25.4	114	8.8 ●	25	52.0 ▲
RTK	135 ●	31	41.9 ▲	42	50.0 ▲	99	13.1	96	3.1 ●	5	40.0 ▲
RTP	179 ●	28	50.0 ▲	45	71.1 ●	132	20.5	132	31.8 ▲	21	61.9 ▲
N33 Total	698 ●	115	44.3 ▲	157	45.2 ▲	433	15.2	428	15.9 ●	94	63.8 ▲
RXC	256 ●	30	46.7 ▲	58	55.2 ●	154	13.6	150	12.0 ●	42	59.5 ▲
RXH	263 ●	52	51.9 ▲	43	41.9 ▲	163	19.6	163	14.1 ●	34	76.5 ●
RYR18	179 ●	33	30.3 ▲	56	37.5 ▲	116	11.2	115	23.5 ▲	18	50.0 ▲
N34 Total	948 ●	204	41.2 ▲	188	43.6 ▲	636	19.5	598	16.4 ●	94	63.8 ▲
RN7	113 ●	26	57.7 ●	25	68.0 ●	92	26.1	91	3.3 ●	13	76.9 ●
RPA	177 ●	39	59.0 ●	44	34.1 ▲	121	21.5	116	6.0 ●	14	64.3 ▲
RVV*	467 ●	100	20.0 ▲	69	29.0 ▲	275	14.2	244	19.3 ●	47	59.6 ▲
RWF	191 ●	39	66.7 ●	50	60.0 ●	148	23.6	147	27.9 ▲	20	65.0 ●

* East Kent Hospitals University Foundation Trust (RVV) has had local issues with their system uploader, resulting in inaccuracies to their reported data.

Table 4 (continued)
Further Process and Clinical Outcomes for England and Wales 2012

Code	Actual number	NSCLC Stage IA, IB, IIA or IIB	% of NSCLC Stage IA, IB, IIA or IIB having surgery	Number of PS 0-1 NSCLC Stage IIIB or IV	% PS 0-1 Stage IIIB or IV NSCLC having chemotherapy	Number of histologically confirmed NSCLC	% histologically confirmed NSCLC having surgery	Number of pre-treatment NSCLC	% pre-treatment NSCLC histology NOS	Number of patients small cell lung cancer	% small cell receiving chemotherapy
N35 Total	1,182 ●	237	59.5 ●	239	58.2 ●	762	26.1	761	17.3 ●	112	69.6 ●
RJD	135 ●	26	65.4 ●	36	63.9 ●	99	25.3	98	12.2 ●	10	70.0 ●
RJE	323 ●	25	60.0 ●	42	78.6 ●	182	22.5	182	19.8 ●	36	80.6 ●
RL4	228 ●	67	58.2 ●	46	58.7 ●	141	36.2	141	13.5 ●	19	68.4 ●
RNA	207 ●	50	62.0 ●	45	66.7 ●	149	21.5	149	11.4 ●	15	60.0 ▲
RWP31	44 ●	9	77.8 ●	12	66.7 ●	37	24.3	37	29.7 ▲	3	100.0 ●
RXW	245 ●	60	53.3 ●	58	31.0 ▲	154	26.6	154	24.0 ▲	29	58.6 ▲
N36 Total	2,676 ●	556	49.1 ▲	590	62.2 ●	1,664	20.9	1,594	28.1 ▲	313	66.5 ●
RE9	148 ●	25	68.0 ●	24	62.5 ●	73	27.4	73	9.6 ●	26	69.2 ●
RLN	296 ●	63	58.7 ●	56	62.5 ●	201	23.4	201	46.8 ▲	35	82.9 ●
RNL	236 ●	36	69.4 ●	71	43.7 ▲	171	24.6	168	23.8 ▲	25	52.0 ▲
RR7	208 ●	59	39.0 ▲	39	53.8 ▲	134	18.7	116	31.0 ▲	10	90.0 ●
RTD	308 ●	70	62.9 ●	64	67.2 ●	198	23.7	174	19.0 ●	38	81.6 ●
RTF	348 ●	55	45.5 ▲	97	64.9 ●	206	14.6	199	24.1 ▲	44	61.4 ▲
RTR	384 ●	84	46.4 ▲	59	86.4 ●	230	20.0	230	13.9 ●	56	53.6 ▲
RVW	307 ●	67	41.8 ▲	68	66.2 ●	190	19.5	190	25.3 ▲	33	66.7 ●
RXP	441 ●	97	36.1 ▲	112	56.3 ●	261	20.3	243	45.3 ▲	46	63.0 ▲
N37 Total	1,552 ●	294	47.6 ▲	355	56.9 ●	1,051	17.8	1,008	19.4 ●	157	70.1 ●
RC1	111 ●	17	58.8 ●	7	57.1 ●	85	17.6	87	16.1 ●	9	55.6 ▲
RCX	153 ●	30	46.7 ▲	46	76.1 ●	111	20.7	111	19.8 ●	23	78.3 ●
RGM	13 ◆	11 ◆	81.8 ◆	0 ◆	0.0 ◆	13 ◆	69.2 ◆	7 ◆	14.3 ◆	0 ◆	0.0 ◆
RGN	182 ●	35	57.1 ●	53	58.5 ●	137	19.7	133	39.1 ▲	17	58.8 ▲
RGP	163 ●	33	27.3 ▲	53	43.4 ▲	96	11.5	88	13.6 ●	11	81.8 ●
RGQ	162 ●	27	51.9 ▲	31	51.6 ▲	113	14.2	113	15.9 ●	10	80.0 ●
RGR	147 ●	18	50.0 ▲	30	60.0 ●	97	13.4	97	11.3 ●	17	76.5 ●
RGT	233 ●	41	51.2 ▲	74	55.4 ●	185	13.5	171	15.2 ●	26	65.4 ●
RM1	322 ●	61	39.3 ▲	57	56.1 ●	173	21.4	170	17.1 ●	33	72.7 ●
RQQ	66 ●	21	47.6 ▲	4	50.0 ▲	41	26.8	31	35.5 ▲	11	54.5 ▲
N38 Total	827 ●	157	57.3 ●	219	58.4 ●	576	18.9	563	13.9 ●	115	67.8 ●
RAJ	219 ●	44	63.6 ●	46	58.7 ●	146	25.3	136	11.0 ●	28	71.4 ●
RDD	203 ●	38	47.4 ▲	50	68.0 ●	143	14.7	143	9.1 ●	36	69.4 ●
RDE	238 ●	38	68.4 ●	80	61.3 ●	158	18.4	158	13.3 ●	33	72.7 ●
RQ8	167 ●	37	48.6 ▲	43	41.9 ▲	129	17.1	126	23.0 ▲	18	50.0 ▲
N39 Total	2,253 ●	439	59.0 ●	517	51.8 ▲	1,416	27.0	1,297	15.6 ●	260	69.6 ●
RJF	137 ●	28	71.4 ●	38	65.8 ●	91	28.6	76	14.5 ●	16	68.8 ●
RK5	207 ●	42	52.4 ●	51	52.9 ▲	138	21.7	133	18.8 ●	31	83.9 ●
RNQ	201 ●	43	55.8 ●	39	61.5 ●	108	28.7	105	38.1 ▲	26	69.2 ●
RNS	177 ●	33	45.5 ▲	42	47.6 ▲	91	27.5	87	18.4 ●	20	60.0 ▲
RTG	289 ●	60	65.0 ●	82	69.5 ●	203	28.1	178	9.6 ●	31	64.5 ▲
RWD	334 ●	48	54.2 ●	61	57.4 ●	200	27.5	187	13.9 ●	36	75.0 ●
RWE	474 ●	97	56.7 ●	105	41.0 ▲	270	28.9	221	13.6 ●	41	63.4 ▲
RX1	430 ●	88	65.9 ●	99	37.4 ▲	314	25.8	309	12.0 ●	57	68.4 ●
England Total	31,003 ●	6,400	51.3 ▲	6,735	57.0 ●	19,675	22.3	18,783	16.2 ●	3,428	67.9 ●
NWW Total	476 ●	103	47.6 ▲	115	63.5 ●	293	18.4	291	5.5 ●	63	63.5 ▲
7A1A1	193 ●	39	48.7 ▲	39	53.8 ▲	126	18.3	126	7.1 ●	27	74.1 ●
7A1A4	155 ●	47	48.9 ▲	34	76.5 ●	84	28.6	83	4.8 ●	23	60.9 ▲
7A1AU	128 ●	17	41.2 ▲	42	61.9 ●	83	8.4	82	3.7 ●	13	46.2 ▲

Table 4 (continued)
Further Process and Clinical Outcomes for England and Wales 2012

Code	Actual number	NSCLC Stage IA, IB, IIA or IIB	% of NSCLC Stage IA, IB, IIA or IIB having surgery	Number of PS 0-1 NSCLC Stage IIIB or IV	% PS 0-1 Stage IIIB or IV NSCLC having chemotherapy	Number of histologically confirmed NSCLC	% histologically confirmed NSCLC having surgery	Number of pre-treatment NSCLC	% pre-treatment NSCLC histology NOS	Number of patients small cell lung cancer	% small cell receiving chemotherapy
SWCN Total	1,556 ●	346	35.5 ▲	353	58.9 ●	913	15.8	903	12.2 ●	195	63.1 ▲
7A2AG	29 ▲	6	16.7 ▲	6	83.3 ●	18	5.6	18	5.6 ●	1	100.0 ●
7A2AJ	39 ●	7	28.6 ▲	11	54.5 ▲	27	7.4	27	18.5 ●	5	60.0 ▲
7A2AL	117 ●	27	55.6 ●	36	77.8 ●	79	19.0	80	2.5 ●	19	89.5 ●
7A2BL	90 ●	26	50.0 ▲	28	46.4 ▲	62	22.6	60	3.3 ●	3	66.7 ●
7A3B7	99 ●	21	33.3 ▲	23	65.2 ●	49	14.3	48	10.4 ●	14	64.3 ▲
7A3C4	90 ●	17	35.3 ▲	24	54.2 ▲	54	13.0	53	7.5 ●	12	50.0 ▲
7A3C7	89 ■	12	25.0 ▲	28	60.7 ●	58	10.3	58	12.1 ●	10	70.0 ●
7A3CJ	83 ●	16	31.3 ▲	31	51.6 ▲	52	9.6	52	17.3 ●	9	77.8 ●
7A4C1	312 ●	79	31.6 ▲	41	63.4 ●	177	16.4	176	18.8 ●	32	68.8 ●
7A5B1	136 ●	28	53.6 ●	40	40.0 ▲	91	18.7	90	15.6 ●	16	31.3 ▲
7A5B3	114 ●	29	31.0 ▲	30	70.0 ●	71	14.1	69	10.1 ●	15	60.0 ▲
7A6AM	123 ●	27	18.5 ▲	11	54.5 ▲	54	18.5	53	9.4 ●	22	63.6 ▲
7A6AR	235 ●	51	33.3 ▲	44	59.1 ●	121	17.4	119	13.4 ●	37	56.8 ▲
Wales Total	2,032 ●	449	38.3 ▲	468	60.0 ●	1,206	16.4	1,194	10.6 ●	258	63.2 ▲
England and Wales Total	33,035 ●	6,849	50.4 ▲	7,203	57.2 ●	20,881	21.9	19,977	15.8 ●	3,686	67.5 ●

Last year's results (including mesothelioma)

2011 Total	33,374 ●	5,821	50.3 ▲	6,682	55.3 ●	20,035	19.8	19,110	19.2 ●	3,748	67.9 ●
Difference	-339	1,028	0.1	521	1.9	846	2.1	867	-3.4	-62	-0.4

Last year's results (excluding mesothelioma)

2011 Total	31,557 ●	5,821	50.3 ▲	6,682	55.3 ●	20,035	19.8	19,108	19.2 ●	3,748	67.9 ●
Difference	1,478	1,028	0.1	521	1.9	846	2.1	869	-3.4	-62	-0.4

Counts aggregated by place first seen Trust.

Range Network

Min			35.4		43.6		15.1		5.5		49.3
LQ			44.8		51.0		17.9		12.3		63.7
Median			48.4		58.3		19.7		14.2		67.0
UQ			58.9		62.9		23.8		17.5		69.8
Max			62.1		70.9		30.8		29.2		80.4

Range Trust

Min			0.0		20.0		0.0		0.0		0.0
LQ			41.2		50.0		16.5		9.1		60.0
Median			50.0		59.1		20.5		13.6		66.7
UQ			60.0		66.7		25.3		19.3		76.5
Max			85.7		100.0		39.0		54.5		100.0

Indicator	Definition	▲	■	●
Actual number	Number of cases (excluding mesothelioma) with date first seen in year specified	<50%	50-75%	≥75%
NSCLC Stage IA, IB, IIA or IIB	Number of NSCLC cases with TNM Stage IA, IB, IIA or IIB			
% of NSCLC Stage IA, IB, IIA or IIB having surgery	Complete when surgery procedure date is present (excluding where primary procedure (OPCS) = E59.5) (denominator = NSCLC cases with TNM Stage IA, IB, IIA or IIB)	<52%		≥52%
Number of PS 0-1 NSCLC Stage IA, IB, IIA or IIB	Number of NSCLC cases with Performance Status 0 or 1 and TNM Stage IA, IB, IIA or IIB			
% PS 0-1 Stage IA, IB, IIA or IIB NSCLC having FEV1 absolute and % predicted	Complete when both FEV1 % and FEV1 absolute amount are present (denominator = NSCLC cases with Performance Status 0 or 1 and TNM Stage IA, IB, IIA or IIB)	<85%		≥85%
Number of PS 0-1 NSCLC Stage IIIB or IV	Number of NSCLC cases with Performance Status 0 or 1 and TNM Stage IIIB or IV			
% PS 0-1 Stage IIIB or IV NSCLC having chemotherapy	Complete when chemotherapy start date is present (denominator = NSCLC cases with Performance Status 0 or 1 and TNM Stage IIIB or IV)	<55%		≥55%
Number of histologically confirmed NSCLC	Number of histologically confirmed NSCLC cases			
% histologically confirmed NSCLC having surgery	Complete when surgery procedure date is present (excluding where primary procedure (OPCS) = E59.5) (denominator = histologically confirmed NSCLC cases)			
Number of pre-treatment NSCLC	Number of pre-treatment NSCLC cases			
% pre-treatment NSCLC histology NOS	Percentage of pre-treatment NSCLC cases with histology NOS (M8046/3) (denominator = pre-treatment NSCLC cases)	<20%		≥20%
Number of patients small cell lung cancer	Number of SCLC cases			
% small cell receiving chemotherapy	Complete when chemotherapy start date is present (denominator = SCLC cases)	<65%		≥65%

◆ – Tertiary Trust standards do not apply

Table 5
Further Process and Clinical Outcomes for Guernsey 2012

Code	Actual number	Number of NSCLC	% of NSCLC having surgery	NSCLC Stage IA, IB, IIA or IIB	% of NSCLC Stage IA, IB, IIA or IIB having surgery	PS 0-1 NSCLC Stage IA, IB, IIA or IIB	% PS 0-1 NSCLC Stage IA, IB, IIA or IIB having FEV1 absolute and % predicted	Number of PS 0-1 NSCLC Stage IIIB or IV	% PS 0-1 Stage IIIB or IV NSCLC having chemotherapy	Number of histologically confirmed NSCLC	% histologically confirmed NSCLC having surgery	Number of pre-treatment NSCLC	% pre-treatment NSCLC histology NOS	Number of patients small cell lung cancer	% small cell receiving chemotherapy
2012 Total	40 ●	35	5.7 ▲	3	66.7 ●	3	66.7 ▲	9	77.8 ●	26	7.7	23	4.3 ●	5	60.0 ▲
2011 Total	41 ●	33	9.1 ▲	6	50.0 ▲	5	100.0 ●	9	55.6 ●	23	13.0	23	13.0 ●	7	100.0 ●
Difference	-1	2	-3.4	-3	16.7	-2	-33.3	0	22.2	1	-5.3	0	-8.7	-2	-40.0

Appendix 1: Trust and Health Board Identification for England and Wales

N01	Lancashire and South Cumbria
RTX	University Hospitals of Morecambe Bay NHS Foundation Trust
RXL	Blackpool Teaching Hospitals NHS Foundation Trust
RXN	Lancashire Teaching Hospitals NHS Foundation Trust
RXR	East Lancashire Hospitals NHS Trust

N02	Greater Manchester and Cheshire
RBT	Mid Cheshire Hospitals NHS Foundation Trust
RBV	The Christie NHS Foundation Trust
RJN	East Cheshire NHS Trust
RM2	University Hospital of South Manchester NHS Foundation Trust
RM3	Salford Royal NHS Foundation Trust
RMC	Bolton NHS Foundation Trust
RMP	Tameside Hospital NHS Foundation Trust
RRF	Wrightington, Wigan and Leigh NHS Foundation Trust
RW3	Central Manchester University Hospitals NHS Foundation Trust
RW6	Pennine Acute Hospitals NHS Trust
RWJ	Stockport NHS Foundation Trust

N03	Merseyside and Cheshire
LLCU*	Liverpool Lung Cancer Unit
RBL	Wirral University Teaching Hospital NHS Foundation Trust
RBN	St Helens and Knowsley Hospitals NHS Trust
REM	Aintree University Hospitals NHS Foundation Trust
REN	The Clatterbridge Cancer Centre NHS Foundation Trust
RJR	Countess of Chester Hospital NHS Foundation Trust
RVY	Southport and Ormskirk Hospital NHS Trust
RWW	Warrington and Halton Hospitals NHS Foundation Trust

N06	Yorkshire Cancer Network
RAE	Bradford Teaching Hospitals NHS Foundation Trust
RCB	York Teaching Hospital NHS Foundation Trust
RCD	Harrogate and District NHS Foundation Trust
RCF	Airedale NHS Foundation Trust
RR8	Leeds Teaching Hospitals NHS Trust
RWY	Calderdale and Huddersfield NHS Foundation Trust
RXF	Mid Yorkshire Hospitals NHS Trust

N07	Humber and Yorkshire Coast Cancer Network
RCC	Scarborough and North East Yorkshire Health Care NHS Trust
RJL	Northern Lincolnshire and Goole Hospitals NHS Foundation Trust
RV9	Humber NHS Foundation Trust
RWA	Hull and East Yorkshire Hospitals NHS Trust

N08	North Trent
RFF	Barnsley Hospital NHS Foundation Trust
RFR	The Rotherham NHS Foundation Trust
RFS	Chesterfield Royal Hospital NHS Foundation Trust
RHQ	Sheffield Teaching Hospitals NHS Foundation Trust
RP5	Doncaster and Bassetlaw Hospitals NHS Foundation Trust

N11	Pan Birmingham
RBK	Walsall Healthcare NHS Trust
RR1	Heart of England NHS Foundation Trust
RRK	University Hospitals Birmingham NHS Foundation Trust
RXK	Sandwell and West Birmingham Hospitals NHS Trust

N12	Arden
RJC	South Warwickshire NHS Foundation Trust
RKB	University Hospitals Coventry and Warwickshire NHS Trust
RLT	George Eliot Hospital NHS Trust
RWP00	Worcestershire Acute Hospitals NHS Trust

N20	Mount Vernon Cancer Network
RC9	Luton and Dunstable University Hospital NHS Foundation Trust
RWG	West Hertfordshire Hospitals NHS Trust
RWH	East and North Hertfordshire NHS Trust

N21	West London Cancer Network
RAS	The Hillingdon Hospitals NHS Foundation Trust
RC3	Ealing Hospital NHS Trust
RFW	West Middlesex University Hospital NHS Trust
RQM	Chelsea and Westminster Hospital NHS Foundation Trust
RT3	Royal Brompton and Harefield NHS Foundation Trust
RV8	North West London Hospitals NHS Trust
RYJ	Imperial College Healthcare NHS Trust

N22	North London
RAL	Royal Free London NHS Foundation Trust
RAP	North Middlesex University Hospital NHS Trust
RKE	The Whittington Hospital NHS Trust
RQW	The Princess Alexandra Hospital NHS Trust
RRV	University College London Hospitals NHS Foundation Trust
RVL	Barnet and Chase Farm Hospitals NHS Trust

N23	North East London Cancer Network
R1HKH	Whipps Cross University Hospital (Barts Health NHS Trust)
R1HM0	Barts and the London NHS Trust
R1HNH	Newham General Hospital NHS Trust (Barts Health NHS Trust)
RF4	Barking, Havering and Redbridge University Hospitals NHS Trust
RQX	Homerton University Hospital NHS Foundation Trust

N24	South East London
RJ1	Guy's and St Thomas' NHS Foundation Trust
RJ2	Lewisham and Greenwich NHS Trust
RJZ	King's College Hospital NHS Foundation Trust
RYQ	South London Healthcare NHS Trust

N25	South West London
RAX	Kingston Hospital NHS Foundation Trust
RJ6	Croydon Health Services NHS Trust
RJ7	St George's Healthcare NHS Trust
RPY	The Royal Marsden NHS Foundation Trust
RVR	Epsom and St Helier University Hospitals NHS Trust

N26	Peninsula
RA9	South Devon Healthcare NHS Foundation Trust
RBZ	Northern Devon Healthcare NHS Trust
REF	Royal Cornwall Hospitals NHS Trust
RH8	Royal Devon and Exeter NHS Foundation Trust
RK9	Plymouth Hospitals NHS Trust

N27	Dorset Cancer Network
RBD	Dorset County Hospital NHS Foundation Trust
RD3	Poole Hospital NHS Foundation Trust
RDZ	The Royal Bournemouth and Christchurch Hospitals NHS Foundation Trust

N28	Avon Somerset and Wiltshire
RA3	Weston Area Health NHS Trust
RA4	Yeovil District Hospital NHS Foundation Trust
RA7	University Hospitals Bristol NHS Foundation Trust
RBA	Taunton and Somerset NHS Foundation Trust
RD1	Royal United Hospital Bath NHS Trust
RVJ	North Bristol NHS Trust

N29	3 Counties Cancer Network
RLQ	Wye Valley NHS Trust
RTE	Gloucestershire Hospitals NHS Foundation Trust
RWP50	Worcestershire Acute Hospitals NHS Trust

N30	Thames Valley
RD7	Heatherwood and Wexham Park Hospitals NHS Foundation Trust
RD8	Milton Keynes Hospital NHS Foundation Trust
RHW	Royal Berkshire NHS Foundation Trust
RN3	Great Western Hospitals NHS Foundation Trust
RTH	Oxford University Hospitals NHS Trust
RXQ	Buckinghamshire Healthcare NHS Trust

N31	Central South Coast
R1F	Isle of Wight NHS Trust
RHM	University Hospital Southampton NHS Foundation Trust
RHU	Portsmouth Hospitals NHS Trust
RN506	Basingstoke and North Hampshire Hospital (Hampshire Hospitals NHS Foundation Trust)
RN541	Royal Hampshire County Hospital (Hampshire Hospitals NHS Foundation Trust)
RNZ	Salisbury NHS Foundation Trust
RYR16	St Richard's Hospital (Western Sussex Hospitals NHS Trust)

N32	Surrey, West Sussex and Hampshire
RA2	Royal Surrey County Hospital NHS Foundation Trust
RDU	Frimley Park Hospital NHS Foundation Trust
RTK	Ashford and St Peter's Hospitals NHS Foundation Trust
RTP	Surrey and Sussex Healthcare NHS Trust

N33	Sussex
RXC	East Sussex Healthcare NHS Trust
RXH	Brighton and Sussex University Hospitals NHS Trust
RYR18	Worthington Hospital (Western Sussex Hospitals NHS Foundation Trust)

N34	Kent and Medway
RN7	Dartford and Gravesham NHS Trust
RPA	Medway NHS Foundation Trust
RVV	East Kent Hospitals University NHS Foundation Trust
RWF	Maidstone and Tunbridge Wells NHS Trust

N35	Greater Midlands
RJD	Mid Staffordshire NHS Foundation Trust
RJE	University Hospital of North Staffordshire NHS Trust
RL4	The Royal Wolverhampton Hospitals NHS Trust
RNA	The Dudley Group NHS Foundation Trust
RWP31	Kidderminster Hospital (Worcestershire Acute Hospitals NHS Trust)
RXW	Shrewsbury and Telford Hospital NHS Trust

N36	North of England Cancer Network
RE9	South Tyneside NHS Foundation Trust
RLN	City Hospitals Sunderland NHS Foundation Trust
RNL	North Cumbria University Hospitals NHS Trust
RR7	Gateshead Health NHS Foundation Trust
RTD	The Newcastle Upon Tyne Hospitals NHS Foundation Trust
RTF	Northumbria Healthcare NHS Foundation Trust
RTR	South Tees Hospitals NHS Foundation Trust
RVW	North Tees and Hartlepool NHS Foundation Trust
RXP	County Durham and Darlington NHS Foundation Trust

N37	Anglia Cancer Network
RC1	Bedford Hospital NHS Trust
RCX	The Queen Elizabeth Hospital King's Lynn NHS Foundation Trust
RGM	Papworth Hospital NHS Foundation Trust
RGN	Peterborough and Stamford Hospitals NHS Foundation Trust
RGP	James Paget University Hospitals NHS Foundation Trust
RGQ	Ipswich Hospital NHS Trust
RGR	West Suffolk NHS Foundation Trust
RGT	Cambridge University Hospitals NHS Foundation Trust
RM1	Norfolk and Norwich University Hospitals NHS Foundation Trust
RQQ	Hinchingbrooke Health Care NHS Trust

N38	Essex Cancer Network
RAJ	Southend University Hospital NHS Foundation Trust
RDD	Basildon and Thurrock University Hospitals NHS Foundation Trust
RDE	Colchester Hospital University NHS Foundation Trust
RQ8	Mid Essex Hospital Services NHS Trust

N39	East Midland Cancer Network
RJF	Burton Hospitals NHS Foundation Trust
RK5	Sherwood Forest Hospitals NHS Foundation Trust
RNQ	Kettering General Hospital NHS Foundation Trust
RNS	Northampton General Hospital NHS Trust
RTG	Derby Hospitals NHS Foundation Trust
RWD	United Lincolnshire Hospitals NHS Trust
RWE	University Hospitals of Leicester NHS Trust
RX1	Nottingham University Hospitals NHS Trust

	Welsh Cancer Network
7A1A1	Ysbyty Glan Clwyd
7A1A4	Ysbyty Maelor Wrexham
7A1AU	Ysbyty Gwynedd
7A2AG	West Wales General Hospital
7A2AJ	Bronglais General Hospital
7A2AL	Prince Philip Hospital Site
7A2BL	Withybush General Hospital
7A3B7	Princess Of Wales Hospital
7A3C4	Singleton Hospital
7A3C7	Morrison Hospital
7A3CJ	Neath Port Talbot Hospital
7A4C1	University Hospital Llandough
7A5B1	The Royal Glamorgan Hospital
7A5B3	Prince Charles Hospital
7A6AM	Nevill Hall Hospital
7A6AR	Royal Gwent Hospital

* LLCU is a partnership between Liverpool Heart and Chest NHS Foundation Trust and Liverpool and Broadgreen University Hospital NHS Trust for management of lung cancer.

Appendix 2: Local Action Plan

Recommendation	Achieved Y/N/P/NK	Planned Action	Suggested Actions	Suggested Responsibility	Date Plan Actioned	Date Issue Resolves
Data Completeness and Quality						
The organisation participates in this national audit.			Contact Clinical Audit Support Unit (nlca@hscic.gov.uk) Visit http://www.ic.nhs.uk/lung for information. Obtain, read and disseminate the National Lung Cancer Audit Annual Report.	Cancer Manager, Governance Lead		
Data on all patients diagnosed with either lung cancer or mesothelioma are submitted to the audit.			Use MDT meetings to capture all cases discussed, Try to record cases in real time or near real time. Liaise with Pathology Departments to correlate cases. Work with IT Department to set up CSV file upload facility if information is collected on a third party system or identify resources to input data directly.	MDT Chair		
All relevant data fields are completed for each patient.			Use proforma for data collection at MDT. Identify key person to quality assure data prior to submission. Ensure data inputters understand clinical implications of data. Map and allocate responsibility along patient pathway. Agree protocols and submission routes for patients that are treated across different organisations.	Data Co-ordinator, Cancer Manager		
Key data fields including Stage and Performance Status should be completed in at least 85 per cent and in at least 95 per cent with respect to the MDT field.			Refer to the documentation on the National Lung Cancer Audit website and ensure that key fields are completed for all relevant cases. MDT Chair assists Co-ordinator by ensuring that Stage, Performance Status and other key fields are discussed and recorded for each patient.	MDT Chair, Data Co-ordinator, Manager		
FEV1 absolute and FEV1% predicted for Stage I and II NSCLC patients with PS 0 or 1 should be recorded in at least 85 per cent.			Record data in real time at MDT where possible; foster links with Physiology Departments to obtain data on relevant patients; quality assure data prior to submission.	MDT Chair, Data Co-ordinator, Cancer Manager		
Process of Care						
Over 95 per cent of patients submitted to the audit are discussed at an MDT.			Liaise with cancer waiting times team to identify lung cancer referrals. Liaise with Radiology Department to identify all imaging suspicious of lung cancer or mesothelioma. Liaise with Pathology Department to identify cases.	MDT Chair, Lung Cancer Clinical Lead		
The Histological Confirmation Rate should be at least 75 per cent. To be reviewed in light of case-mix adjusted odds ratio.			This result should be interpreted in conjunction with the case-mix adjusted odds ratio, which might better reflect whether the organisation is an outlier. Ensure all histological diagnoses are submitted to the audit, including those confirmed only by resection. Liaise with Pathology Department to identify cases. Review clinical diagnoses and diagnostics protocols if HCR is below optimum.	MDT Chair, Lung Cancer Clinical Lead		
The proportion of patients receiving CT scan prior to bronchoscopy should exceed 95 per cent.			Ensure that all CT / bronchoscopy data is submitted to the audit. Review patient pathway and individual clinician practices.	MDT Chair, Lung Cancer Clinical Lead, Radiologists		
Over 80 per cent of patients are seen by a Lung Cancer Nurse Specialist.			Review the Specialist Nurse service, ensuring all nursing posts are staffed and that clear referral pathways exist.	MDT Chair, Lung Cancer Clinical Lead, Specialist Nurse		
Over 80 per cent of patients have a Lung Cancer Nurse Specialist present at the time of diagnosis.			Review the Specialist Nurse service, allocate extra nursing support alongside lung cancer clinics	MDT Chair, Lung Cancer Clinical Lead, Specialist Nurse		
Co-morbidity that prevents a patient receiving treatment of choice should be recorded for all relevant cases.			Ensure that all relevant co-morbidity data is discussed at MDT, and ensure that cases where co-morbidity prevents treatment of choice are submitted to the audit. It is important that the collected data adheres to the definitions in the LUCADA data manual.	MDT chair, Lung Cancer Clinical Lead, Specialist Nurse		
PET scan dates should be recorded for all relevant cases.			Ensure that all PET data is captured at MDT submitted to the audit.	MDT Chair, Lung Cancer Clinical Lead, Specialist Nurse		
NSCLC NOS rate of more than 20 per cent should be reviewed to ensure that best practice histological diagnostic techniques including immunohistochemistry are being followed, in order that patients receive appropriate chemotherapy regimens.			Ensure that Pathologist is an integral part of the lung MDT and understands the importance of tumour subtyping. Ensure that a locally approved panel of immunohistochemical markers are being used for subtyping and that locally approved appropriate mutation testing is being applied.	MDT Chair, Pathologist, Lung Cancer Clinical Lead, Specialist Nurse, MDT Co-ordinator		

Recommendation	Achieved Y/N/P/NK	Planned Action	Suggested Actions	Suggested Responsibility	Date Plan Actioned	Date Issue Resolves
Clinical Outcomes						
<p>Surgical resection rates below 14 per cent for all patients excluding small cell lung cancer or mesothelioma must be reviewed</p> <p>To be reviewed in light of case-mix adjusted odds ratio.</p>			<p>This result should be interpreted in conjunction with the case-mix adjusted odds ratio, which might better reflect whether the organisation is an outlier. Ensure that all surgical resections are submitted to the audit. If data is complete then review treatment policies for early stage lung cancer in patients with good Performance Status.</p> <p>Ensure that the Thoracic Surgeon attends MDT meetings. Consider offering a second opinion in borderline cases.</p>	MDT Chair, Lung Cancer Clinical Lead, Thoracic Surgeons		
<p>Surgical resection rates for patients for all patients excluding small cell lung cancer or mesothelioma with Stage I or II disease below 52 per cent must be reviewed.</p>			<p>This result should be interpreted in conjunction with the case-mix adjusted odds ratio, which might better reflect whether the organisation is an outlier. Ensure that all surgical resections are submitted to the audit. If data is complete then review treatment policies for early stage lung cancer in patients with good Performance Status.</p> <p>Ensure that the Thoracic Surgeon attends MDT meetings. Consider offering a second opinion in borderline cases.</p>	MDT Chair, Lung Cancer Clinical Lead, Thoracic Surgeons		
<p>Active anti-cancer treatment rates below 60 per cent should be reviewed.</p> <p>To be reviewed in light of case-mix adjusted odds ratio.</p>			<p>This result should be interpreted in conjunction with the case-mix adjusted odds ratio, which might better reflect whether the organisation is an outlier.</p> <p>Ensure that all treatments are submitted to the audit. Review treatment policies for small cell lung cancer patients. Review pathway from diagnosis to treatment to ensure it is as expeditious as possible.</p>	MDT Chair, Lung Cancer Clinical Lead, MDT members		
<p>Chemotherapy rates for small cell lung cancer below 65 per cent should be reviewed.</p> <p>To be reviewed in light of case mix adjusted odds ratio</p>			<p>This result should be interpreted in conjunction with the case-mix adjusted odds ratio, which might better reflect whether the organisation is an outlier.</p> <p>Ensure that all treatments are submitted to the audit. Review treatment policies for small cell lung cancer patients.</p>	MDT Chair, Lung Cancer Clinical Lead, MDT members		
<p>Chemotherapy rates for patients of PS 0-1 with advanced Stage NSCLC IIIB/IV below 55 per cent should be reviewed.</p> <p>To be reviewed in light of case-mix adjusted odds ratio.</p>			<p>This result should be interpreted in conjunction with the case-mix adjusted odds ratio, which might better reflect whether the organisation is an outlier.</p> <p>Ensure that all treatments are submitted to the audit. Review treatment policies for non-small cell lung cancer patients with advanced stage.</p>	MDT Chair, Lung Cancer Clinical Lead, MDT members		
<p>Low median survival, as demonstrated by a case-mix adjusted hazard ratio significantly below the baseline, should be investigated.</p>			<p>Ensure that all relevant data has been submitted to the audit, Identify areas where audit standards have not been met or where CMA demonstrates the Trust to be an outlier and review.</p>	MDT Chair, Lung Cancer Clinical Lead, MDT members		

Appendix 3: Glossary

Adjuvant treatment – An additional therapy (e.g. chemotherapy or radiotherapy) provided to improve the effectiveness of the primary treatment (e.g. surgery). This may aim to reduce the chance of local recurrence of the cancer or to improve the patient's overall chance of survival.

Adenocarcinoma – a type of cancer arising from glandular tissue.

Anti-cancer treatment (active treatment) – a term used to define treatments for lung cancer that have an effect on the tumour itself, not just on symptoms. In lung cancer patients these are most often surgery, chemotherapy, radiotherapy or a combination.

Benchmarking – a method of comparing processes and outcomes against standards.

Biopsy – removal and examination of tissue, usually microscopic, to establish a precise (histological) diagnosis.

Bronchoscopy – a procedure for examining the airways by inserting an instrument (bronchoscope) into the trachea and lungs, normally via the nose. Enables a bronchial biopsy to be taken.

Bronchial biopsy – removal of a small piece of lung tissue during a bronchoscopy in order to make a histological diagnosis.

Cancer Network – a system within the NHS to organise the integrated and care of cancer patients across a geographic region.

Cancer Registry/ies – organisations who systematically collect high level data about all cases of cancer in the UK. Cancer registries are unique in being able to provide historical trend and population-based data to monitor changes in cancer incidence or survival over long periods of time.

Case ascertainment – the number of cases of lung cancer actually recorded by an organisation as a proportion of the number expected. Gives assurance that organisations are submitting data on all relevant cases.

Case-mix – refers to the different characteristics of patients seen in different hospitals (for example age, sex, disease stage, social deprivation and general health). Knowledge of differing case-mix enables a more accurate method of comparing quality of care (case-mix adjustment).

Case-mix adjustment – a statistical method of comparing quality of care between organisations that takes into account important and measurable patient characteristics.

Chemotherapy – medicines used in the treatment of cancer that can be given by mouth or by injection.

Common denominator – (in a non-mathematical context) factors that link objects (e.g. hospitals) together.

Co-morbidity – medical conditions or disease processes that are additional to the disease under investigation (in this case lung cancer). In the NLCA this is recorded when a co-morbidity restricts the type of treatment that can be given for lung cancer.

CT scan – the abbreviated term for computed or computerised axial tomography. These are tests that produce detailed images of the body using X-rays images that are enhanced by a computer.

Cytological – refers to a pathological examination of cells outside the architecture of the actual tissue or organ they are taken from (as opposed to histological).

Data completeness – a measure of the standard of data submitted to the audit, both in terms of the numbers of cases submitted as well as the data on each individual case.

Diagnosis – confirming the presence of the disease.

Health Board – an organisation providing healthcare services in Scotland and Wales. A Health Board may manage one or several hospitals within a region.

Histological – refers to a pathological examination of cells within the architecture of a tissue or organ rather than just the cells themselves (as opposed to cytological).

Hospital Trust – an organisation providing secondary healthcare services in England. A Hospital Trust may be made up of one or several hospitals within a region.

Immunohistochemistry – is a special test used by pathologists to detect specific molecules on cells. It is used to help diagnose diseases, such as cancer, and to detect the presence of microorganisms.

Improving Lung Cancer Outcomes Project (ILCOP) – a project sponsored by the Health Foundation and managed by the Royal College of Physicians to look at ways to improve care offered to people diagnosed with lung cancer.

Interquartile range – the range of a particular variable excluding the highest quarter and lowest quarter of the values recorded. Can be useful to give a sense of the spread of a set of data without being affected by very high or very low results.

Lead Clinician – a healthcare professional in a hospital taking overall responsibility for the services provided for a specific disease area.

Lobectomy – an operation to remove a whole section (lobe) of lung tissue see also wedge resection. There are three lobes in the right lung and two lobes in the left lung.

LUCADA – National Lung Cancer Audit Dataset. (LUNG CANCER DATA)

Lung Cancer Nurse Specialist – a nurse specialising in care of people diagnosed with lung cancer or mesothelioma.

Lymph nodes – small, oval-shaped organs of the immune system, whose main job is to fight infection. Distributed widely throughout the body (including the neck, armpit, abdomen and thorax) they are a common place for cancers to spread.

MDT – Multi-Disciplinary Team, a group of healthcare professionals working in a co-ordinated manner for patient care.

Mediastinum/Mediastinal – refers to an area within the center of the thorax (chest) between the two lungs, where the heart, blood vessels and lymph nodes are found.

Mediastinotomy/oscopy – an operation that enables visualization and biopsy of the mediastinal lymph nodes. These procedures are often used to determine whether a cancer has spread to the lymph nodes, which affects the stage of the disease.

Mesothelioma – cancer of the lining of the lung caused by exposure to asbestos.

Metastasis – cancer that has spread from the place where it was formed to grow in another part of the body.

Network – see ‘Cancer Network’.

NLCA – National Lung Cancer Audit.

Nodule (lung nodule) – a small abnormality on the lung often found on chest X-rays or CT scans. Most lung nodules are non-cancerous (benign). However, some lung nodules may be cancerous - either early-stage lung cancer or metastatic cancer that has spread to the lungs from another site in the body.

Non-small cell carcinoma – a group of types of lung cancer sharing certain characteristics, that makes up 85-90 per cent of all lung cancers. Includes squamous carcinoma and adenocarcinoma. See also small cell carcinoma.

NOS – not otherwise specified. In the case of NSCLC histology, this implies that the histological diagnosis has not been sub-classified to a particular cell type e.g. squamous carcinoma, adenocarcinoma etc.

NSCLC – non-small cell lung cancer.

Operability – in the consideration of surgical treatment of a lung cancer, refers to the patients’ ability to cope with both the operation and the subsequent reduction of lung volume and function. See also resectability.

Oncologist – a doctor who specializes in treating people with cancer.

Oncology – the study and treatment of tumours.

Palliative – (of a medicine or medical care) relieving pain without dealing with the cause of the condition.

Performance Status – a systematic method of recording the ability of an individual to undertake the tasks of normal daily life compared with that of a normal person.

PET scan – an abbreviation for positron emission tomography. This is a computerised diagnostic technique that uses radioactive substances to examine structures of the body. Nowadays usually combined with a CT scan (PET-CT scan). It produces a three-dimensional image that reflects the metabolic and chemical activity of the body.

Pneumonectomy – surgical removal of a lung or part of a lung.

Radiologist – a doctor specialising in the use of imaging technologies, including radiation, to diagnose and treat disease.

Radiotherapy – the treatment of cancer using radiation, which is most often delivered by X-ray beams (external beam radiotherapy) but can be given internally (brachytherapy).

Resectability – in the consideration of surgical treatment of a lung cancer, refers to the ability of the Surgeon to remove the tumour taking into account its location and stage. See also operability.

RCP – abbreviation for The Royal College of Physicians, the professional body of doctors practicing general medicine and its sub-specialties.

SCLC – small cell lung cancer.

Secondary care – care provided by a hospital as opposed to that provided in the community by a general practitioner and allied staff (primary care).

Small cell lung cancer – a type of lung cancer making up around 10-15 per cent of all lung cancers. See also non-small cell carcinoma.

Squamous Carcinoma – a type of cancer arising from cells which line body cavities.

Staging/Stage – the anatomical extent of a cancer.

Surgical resection – an operation to remove abnormal tissues or organs.

Tertiary Centres – hospitals that specialise in diagnosis and treatment of specific conditions, often handling very complex cases. Other hospitals may refer patients to these centres for specialist treatment.

Thoracic Surgeon – specialist surgeon who operates on the chest and lungs.

Wedge resection – an operation to remove a section of lung tissue smaller than a lobe – see also lobectomy.

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