method of treating peripheral early stage (T1, T2) NSCLC. It is considered as a valid option for treatment when surgery is high risk or declined. Many clinicians now question whether SABR can be considered as a genuine alternative to surgery for peripheral stage I NSCLC.

One concern about any comparison with surgery is that those patients who receive SABR may miss out on adjuvant chemotherapy due to lack of pathological nodal staging that would be done post operatively.

Our aim was to determine what proportion of cases this might represent.

Methods We carried out a retrospective review of surgical resections for lung cancer between 2008 and 2011 at a teaching hospital in Yorkshire. We identified those patients who were staged preoperatively as T1 or T2 with no nodal or distant metastases.

We also identified a subgroup of patients with peripheral tumours no greater than 5cm in diameter (i.e. those potentially suitable for SABR).

We recorded post operative staging and whether they received adjuvant chemotherapy.

Results We identified 162 cases in total (81 female) of which 13.7% had nodal involvement on post operative staging. Of these, 72.7% received adjuvant chemotherapy. Within the subgroup with peripheral tumours <5cm there were 116 cases of which 7.0% had nodal involvement. Of these, 62.5% received adjuvant chemotherapy.

Conclusions Our findings show that a significant number of patients who are preoperatively staged as N0 do have hilar and/or mediastinal nodal involvement at the time of surgery. However, many of these patients have central tumours not suitable for SABR. In the subgroup of patients with small peripheral tumours, that would potentially be suitable for SABR, this proportion is smaller.

It is thought that on direct comparison with surgery those who receive SABR might be more likely to develop recurrent disease due to nodal involvement outside of the radiotherapy field in some patients. Our results suggest that this proportion is small. Any effect on overall survival would be weighed against peri-operative morbidity and mortality.

S95

SOCCAR: INTERNATIONALLY RESONANT RESULTS FROM A RANDOMISED TRIAL BASED ON UK CLINICAL PRACTISE

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Concurrent chemoradiation is now recognised as a standard of care for patients with inoperable Stage III NSCLC and good Performance Status. Two year survival rates in published trials ranges from 29% to 43%, with median survival in the range of 14 to 17 months. An ongoing multi-centre study performed in the United States has recently reported median survival of 21 months for this group of patients, with no advantage from dose escalation from 60 to 74 Gy in conventional fractionation. In SOCCAR, we employed a standard UK radical lung cancer radiotherapy treatment schedule; 130 patients were randomised to receive 55Gy in 20 fractions over 4 weeks, given either concurrently or sequentially with cisplatin and vinorelbine. Trial entry criteria were: pathologically confirmed stage III NSCLC, judged in-operable by a thoracic surgeon and treatable within a radical radiotherapy volume with V20 <30% and <12 cm oesophagus in PTV. Patients were required to have PS 0 or 1, FEV1 >= 1L and TLCO >= to 50%. Patients with weight loss and N3 disease were included, with no upper age limit. Characteristics were well balanced between the two arms.

Results Median survival is 27.4 months in the concurrent arm and 18.6 months in the sequential arm. Two year survival rates are 54% and 42% respectively. Toxicity was acceptable and similar in both arms. Documented local relapse was 10% in the concurrent arm and 22% in the sequential arm. Quality of Life was measured prospectively in all patients, there being no significant difference in either arm. In comparison with 16 published RCTs, comprising 1733 patients treated with concurrent chemoradiation, the concurrent treatment arm in SOCCAR ranks top in survival with comparable toxicity. These results are recognised as being of major relevance internationally and are the basis of a proposed multi-centre international study.

Clinical studies in interstitial lung disease

S96

A COMBINATION SCORE OF RAISED SACE, LYMPHOPENIA AND HYPERGAMMAGLOBULINAEMIA CORRELATES WITH ACTIVE DISEASE ON THORACIC CT SCAN OF PATIENTS WITH PULMONARY SARCOIDOSIS

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Introduction Clinical decisions about treatment in pulmonary sarcoidosis, and phenotyping research studies are hampered by inability to objectively measure disease activity. We reviewed 120 consecutive patients from our Sarcoidosis Clinic and observed that raised sACE, immunoglobulin and/or presence of lymphopenia were associated with clinical decisions to change treatment. We question if these markers could be combined to form an objective measure of disease activity in pulmonary sarcoidosis.

Methods As there is no 'gold standard' for disease activity in sarcoidosis, we used thoracic CT scan to reflect disease activity on the basis that ground glass opacity, nodularity, consolidation, interlobular septal thickening and conglomeration reflect T cell alveolitis, cellular infiltrate and granulomatous deposits¹⁻². Using Fleischner Society definition of terms³, we designed a scoring system ("CT activity score; CTAS") to quantify this, and examined if a composite clinical score ("clinical activity score; CAS"; IgG > 13 g/l = 1, Lymphocytes < 1x10⁹/l = 1, sACE < 55 U/l = 0, 56-100 U/l = 1, > 100 U/l = 2) correlated with changes in CTAS. An enhanced CAS incorporating extent of defined CXR abnormalities was also examined. We collected data from 100 consecutive patients diagnosed according to WASOG/ATS criteria (with histological confirmation), who had thoracic CT scan, CXR and blood parameters within three months of each other. Two radiologists scored the CT scans blindly. We present results for the first 20 cases.

Results CTAS score (maximum possible 81) in our patients ranged from 0 to 40; CAS from 0-4 (maximum 5); when including CXR score - from 1-12 (maximum 12). We found a strong correlation between CAS and CTAS (Figure 1). No correlation was observed between the CTAS and FVC, TLC or KCO, supporting the premise that severe irreversible functional loss can be inactive.

Conclusions A combination score incorporating lymphopenia, sACE levels, and hypergammaglobulinemia is strongly correlated with CT quantification of disease activity in pulmonary sarcoidosis. Addition of CXR scoring improved the correlation. This score could be used as an indicator of disease activity to aid clinical decisions on treatment, and paves the way for collation of larger numbers and longitudinal studies to further validate the tool.

References

1. Nishimura K et al. Radiology 1993.
2. Oberstein A et al. Sarc Vasc Diffuse Lung Dis 1997.
3. Hansell DM et al. Radiology 2008.

Abstract S96 Graph 1

S97 IS AN INCREASED TENDENCY TO CLOT A RISK FACTOR FOR DEVELOPING IDIOPATHIC PULMONARY FIBROSIS?

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Background The aetiology of idiopathic pulmonary fibrosis (IPF) remains poorly understood. Recent animal studies and epidemiological data have suggested that activation of the coagulation cascade in the lung may have an important role in the pathogenesis of IPF.

Methods We recruited incident cases of physician diagnosed IPF from five teaching hospitals and eight district general hospitals in England and Wales. Age and sex matched general population controls were recruited from the same region. Participants were asked for details of lifetime occupational history, current or previous illnesses, medication and smoking. Each case and control then had a venous blood sample taken for a thrombophilia screen, including inherited and acquired clotting defects. We also collected high resolution computed tomography (HRCT) scans for all cases, which were reviewed by two experienced thoracic radiologists to confirm

the diagnosis of IPF. After excluding individuals on warfarin, logistic regression was used to estimate odds ratios for the association between a prothrombotic state and IPF, adjusting for age, sex and highly sensitive C Reactive Protein (hsCRP). Individuals were defined as being prothrombotic if they had at least one clotting defect present. The analysis was then repeated, stratifying cases by radiological diagnosis.

Results Our study included 306 incident cases of IPF (mean age at diagnosis 72.6 years, 70.6% male) and 256 controls (see Table 1). We found an increased tendency to clot among our cases (Odds Ratio [OR] 4.67; 95% Confidence Interval [CI] 3.00 to 7.23) compared to controls. After stratifying by radiological diagnosis, this association was stronger in those with definite usual interstitial pneumonia (UIP) (OR 5.86; 95% CI 3.08 to 11.15) compared to probable UIP (OR 3.47; 95% CI: 1.92 to 6.29). There was no effect modification by age, sex or hsCRP.

Conclusion An increased tendency to clot appears to be an independent risk factor for developing IPF. A clinical trial using one of the new, safer anticoagulants may be warranted.

S98 EARLY CLINICAL EXPERIENCE WITH PIRFENIDONE FOR IDIOPATHIC PULMONARY FIBROSIS (IPF) IN THE UK: INTERIM RESULTS FROM A UK COHORT

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Introduction and Objectives In March 2011, the novel anti-fibrotic, pirfenidone (Esbriet®), became the first drug to be licenced in Europe for the treatment of IPF. Since September 2011 pirfenidone has been available in the UK through a named patient programme (NPP). We present initial findings from a real-world study describing clinical experience with pirfenidone in routine UK clinical practise.

Methods A multi-centre, retrospective, cohort review was undertaken across 4 NHS Trusts. Data (through to July 2012) were collected from the clinical records of individuals receiving pirfenidone for IPF through the NPP.

Results Data was available from 68 patients (72% Male). Mean ($\pm$ S.D.) age at diagnosis was 67.3 $\pm$ 8.1 years. At initiation of pirfenidone FVC was 69.4 $\pm$ 21.5% predicted and DLco 39.8 $\pm$ 15.3% predicted. Domiciliary oxygen was being administered to 38.2% (26/68).

Abstract S97 Table 1

Table 1: Demographic features of incident cases of IPF and matched general population controls

		Cases (n=306)	Controls (n=256)
Mean age (standard deviation)		72.6 (8.5)	70.9 (9.0)
Number of men (%)		216 (70.6)	189 (73.8)
Smoking status (%)	Never smoked	85 (27.8)	113 (44.1)
	Ex-smokers	194 (63.4)	119 (46.5)
	Current smokers	27 (8.8)	24 (9.4)
Previous venous thromboembolic event (%)		34 (11.1)	15 (5.9)
Anticoagulant status (%)	Never been on warfarin	258 (84.3)	239 (93.4)
	Previously been on warfarin	21 (6.9)	11 (4.3)
	Currently on warfarin	27 (8.8)	6 (2.3)