



This is a repository copy of *Radical radiotherapy for non-small cell lung cancer (NSCLC): real world outcomes for two accelerated fractionation schedules.*

White Rose Research Online URL for this paper:  
<http://eprints.whiterose.ac.uk/128942/>

Version: Supplemental Material

---

**Proceedings Paper:**

Robinson, S.D., Absalom, K.A., Lankathilake, A. et al. (5 more authors) (2018) Radical radiotherapy for non-small cell lung cancer (NSCLC): real world outcomes for two accelerated fractionation schedules. In: Lung Cancer. The 16th Annual BTOG, 24-26 Jan 2018, Dublin. Elsevier , S71-S71.

[https://doi.org/10.1016/S0169-5002\(18\)30189-2](https://doi.org/10.1016/S0169-5002(18)30189-2)

---

**Reuse**

Items deposited in White Rose Research Online are protected by copyright, with all rights reserved unless indicated otherwise. They may be downloaded and/or printed for private study, or other acts as permitted by national copyright laws. The publisher or other rights holders may allow further reproduction and re-use of the full text version. This is indicated by the licence information on the White Rose Research Online record for the item.

**Takedown**

If you consider content in White Rose Research Online to be in breach of UK law, please notify us by emailing [eprints@whiterose.ac.uk](mailto:eprints@whiterose.ac.uk) including the URL of the record and the reason for the withdrawal request.



[eprints@whiterose.ac.uk](mailto:eprints@whiterose.ac.uk)  
<https://eprints.whiterose.ac.uk/>

**INTRODUCTION**

Numerous radiotherapy regimes are used for inoperable NSCLC who are not suitable for stereotactic ablative radiotherapy. Our centre has used continuous hyperfractionated accelerated radiotherapy (CHART, 54Gy in 36 fractions over 12 days) and accelerated hypofractionated radiotherapy (55Gy in 20 fractions over 4 weeks) with selection largely down to patient choice (in-patient vs out-patient treatment).

This audit reviews patients treated with radical radiotherapy between 2010 - 2015.

**METHODS**

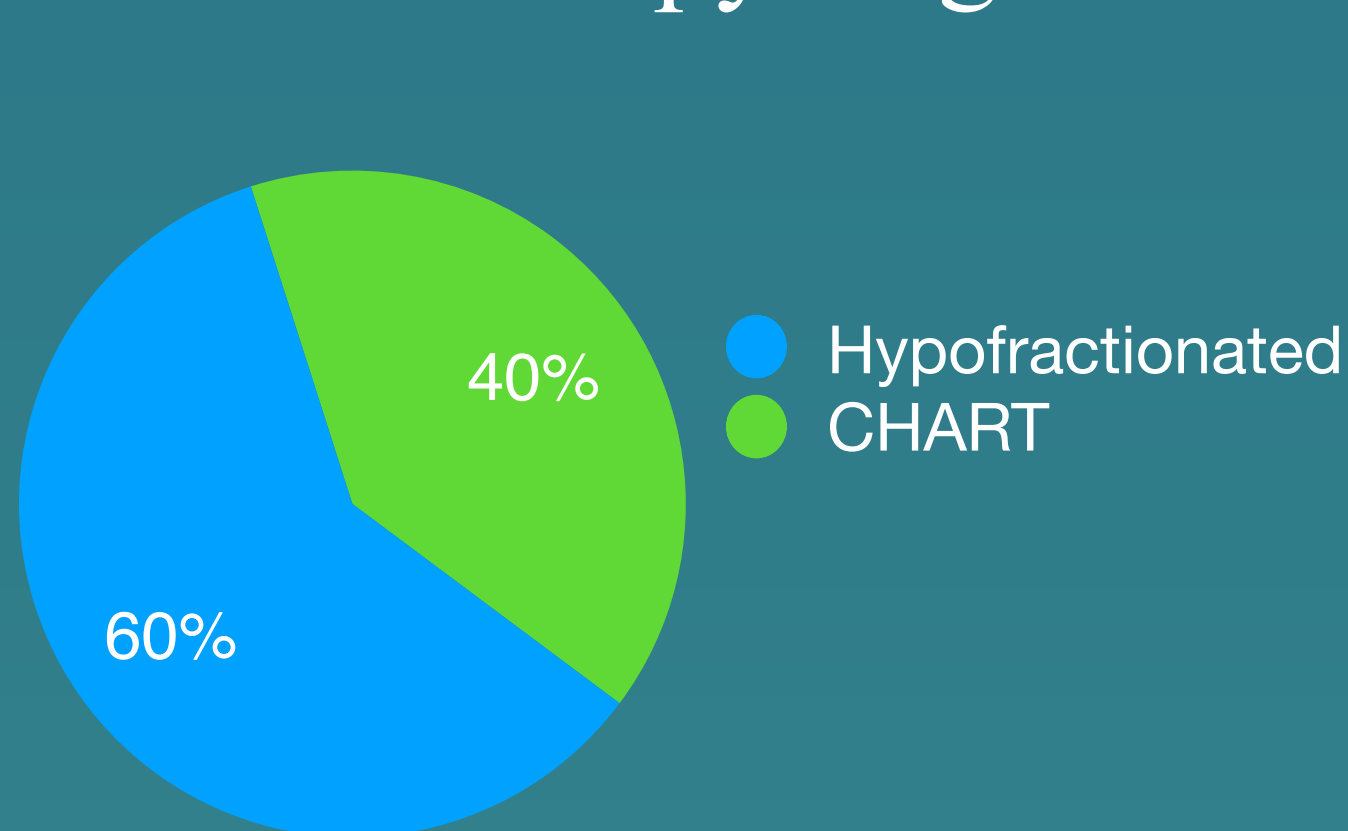
Case notes and radiotherapy records for all patients receiving radical radiotherapy were retrospectively reviewed. Basic patient demographics, tumour characteristics, radiotherapy and survival data were collected. Descriptive statistical analysis and Cox regression analysis was performed using SPSS.

**RESULTS - Demographics**

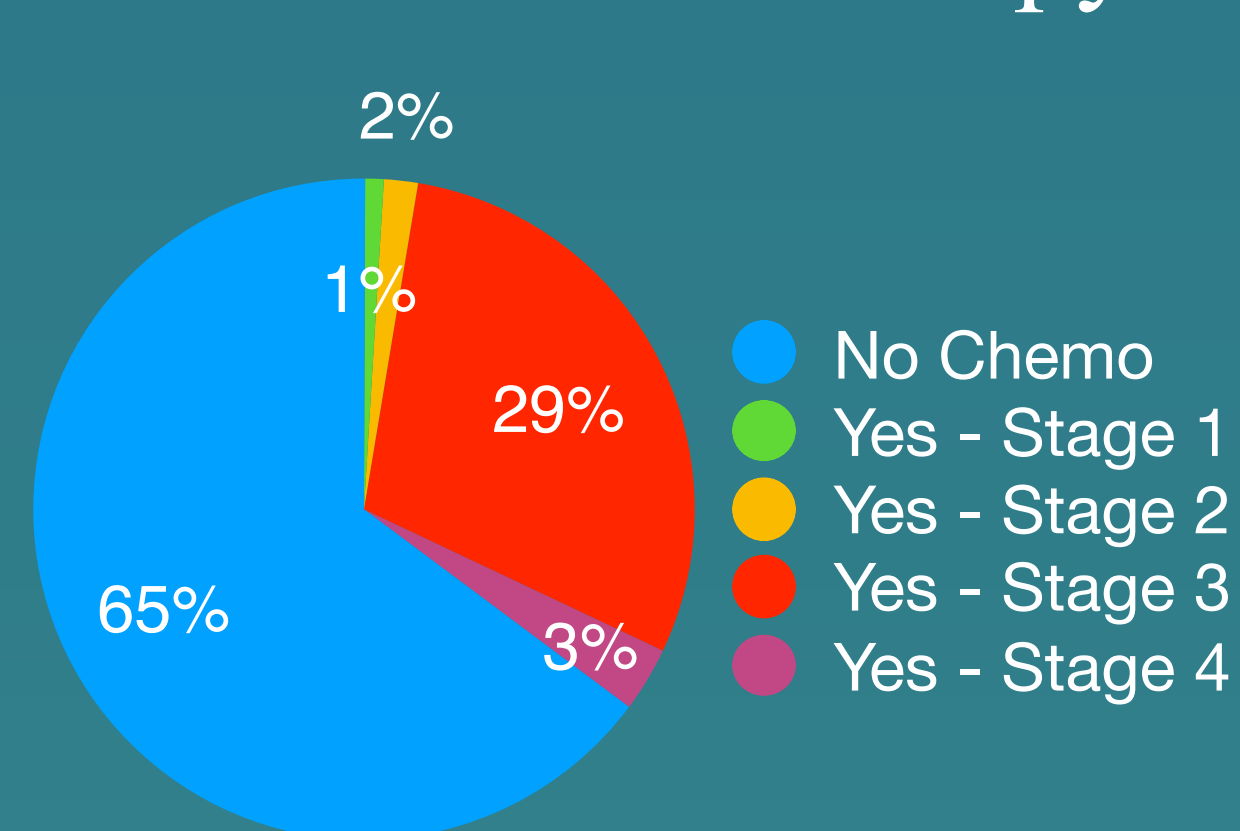
563 patients received radical radiotherapy between 2010-15.

Demographic		Number of patients / Median	Percentage of patients (%) / Range
Gender	Male	316	56.1
	Female	247	43.9
Age	Median; Range	71	36-93
Performance Status	0	94	16.7
	1	203	36.1
	2	123	21.8
	3	9	1.6
	Unknown	134	23.8
FEV1, L	Median; Range	1.6	0.6-3.67
	Unknown	269	47.8
Site of Primary	Central	4	0.7
	Right	308	54.7
	Left	247	43.9
	Unknown	4	0.7
Histology	Squamous Cell	264	46.9
	Adenocarcinoma	144	25.6
	Other histology	38	6.7
	No histology	117	20.8
Stage (TNM v7)	1	171	30.4
	2	77	13.7
	3	281	49.9
	4	33	5.9
	Unknown	1	0.2
PET performed?	Yes	532	94.5
	No	31	5.5

**Radiotherapy Regimen**



**Prior Chemotherapy**



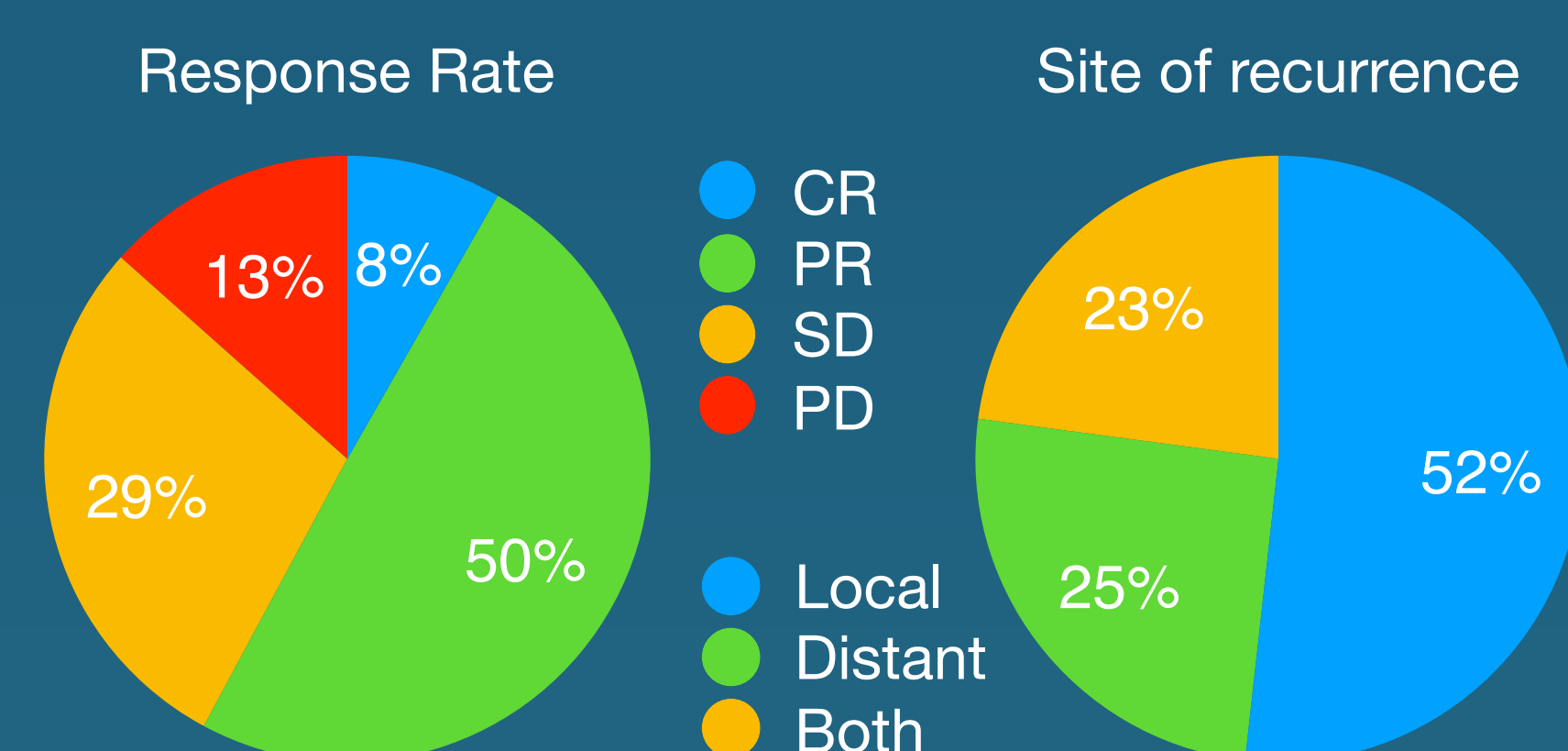
99.1% completed their prescribed radiotherapy treatment.

**RESULTS - Outcome**

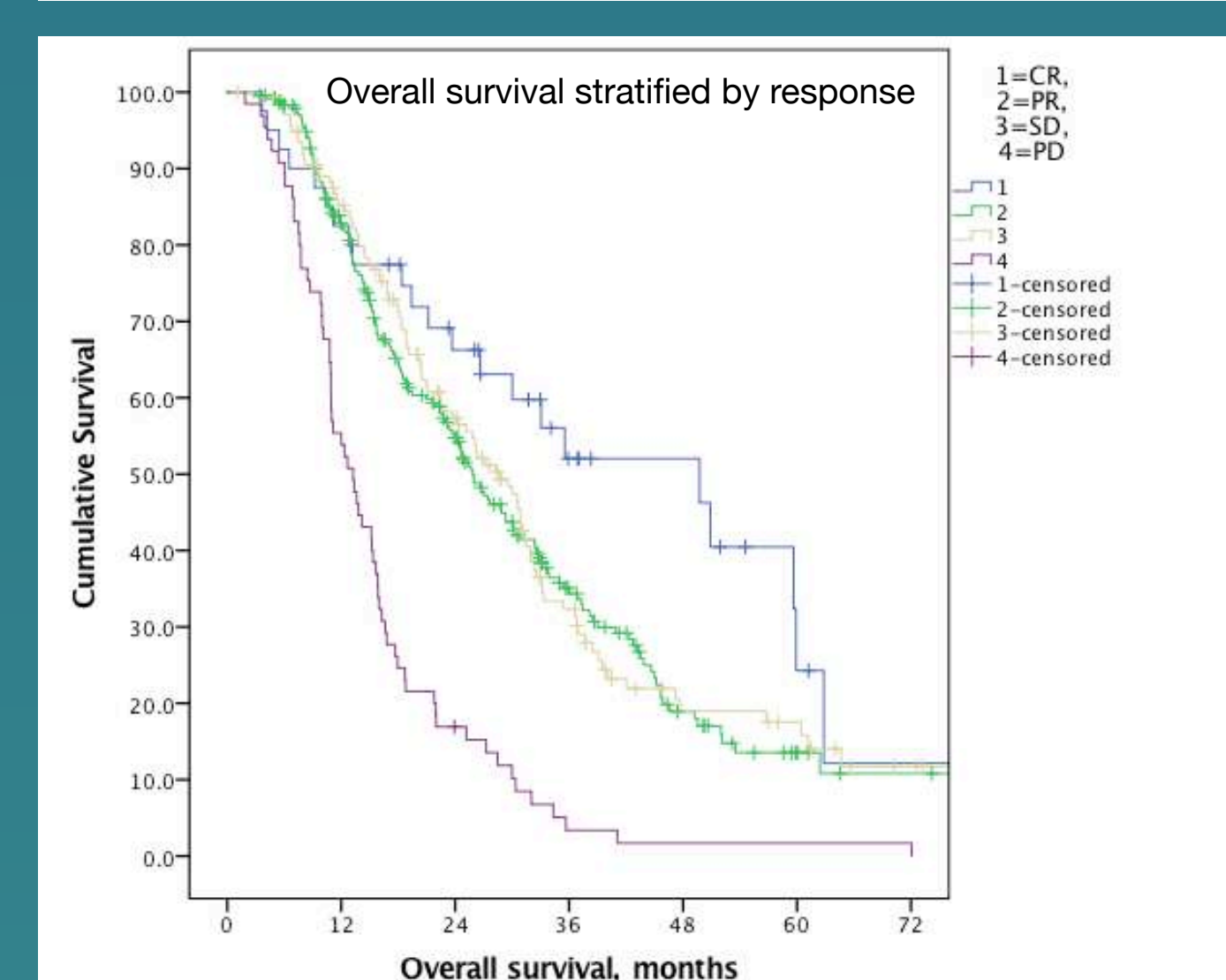
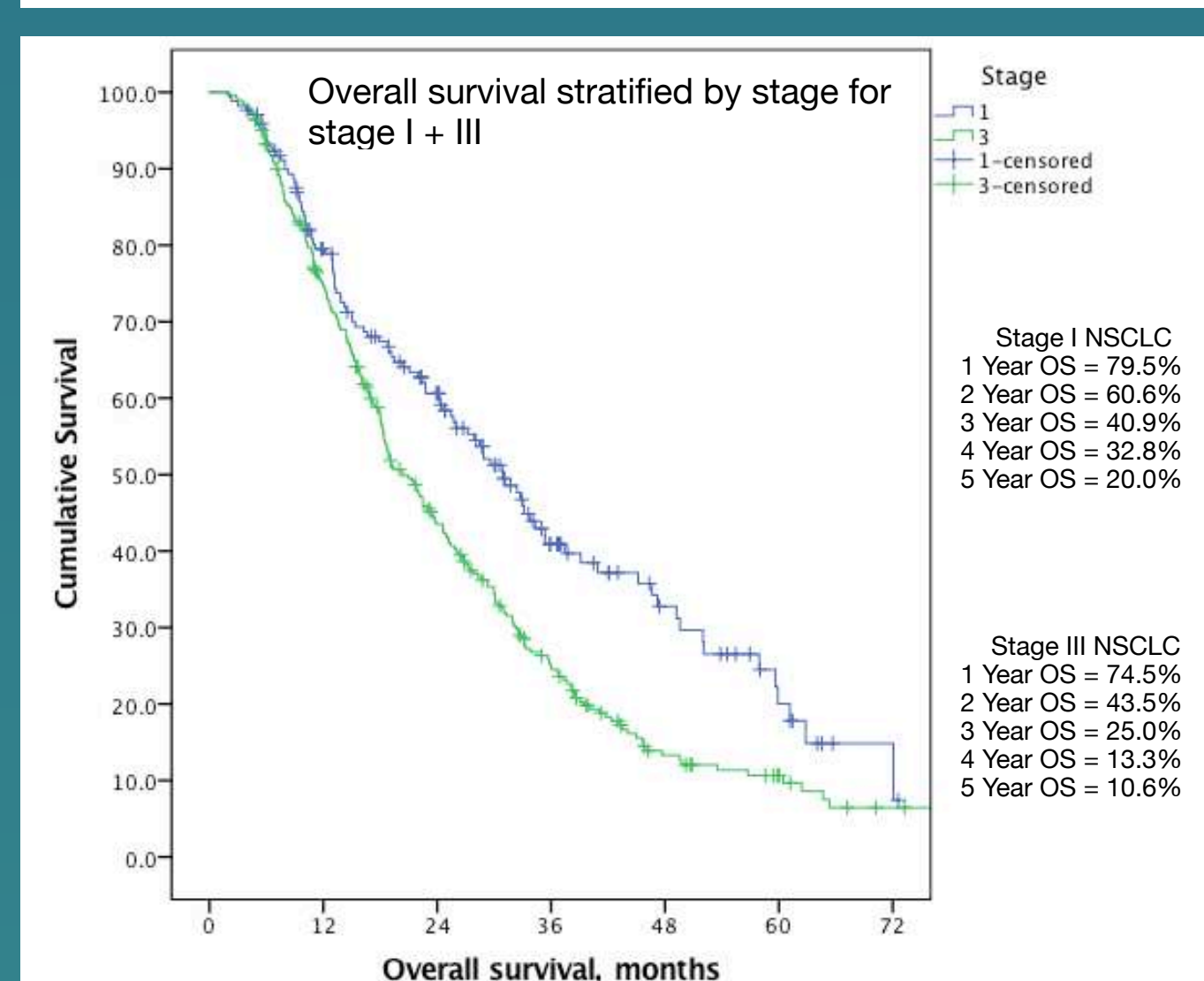
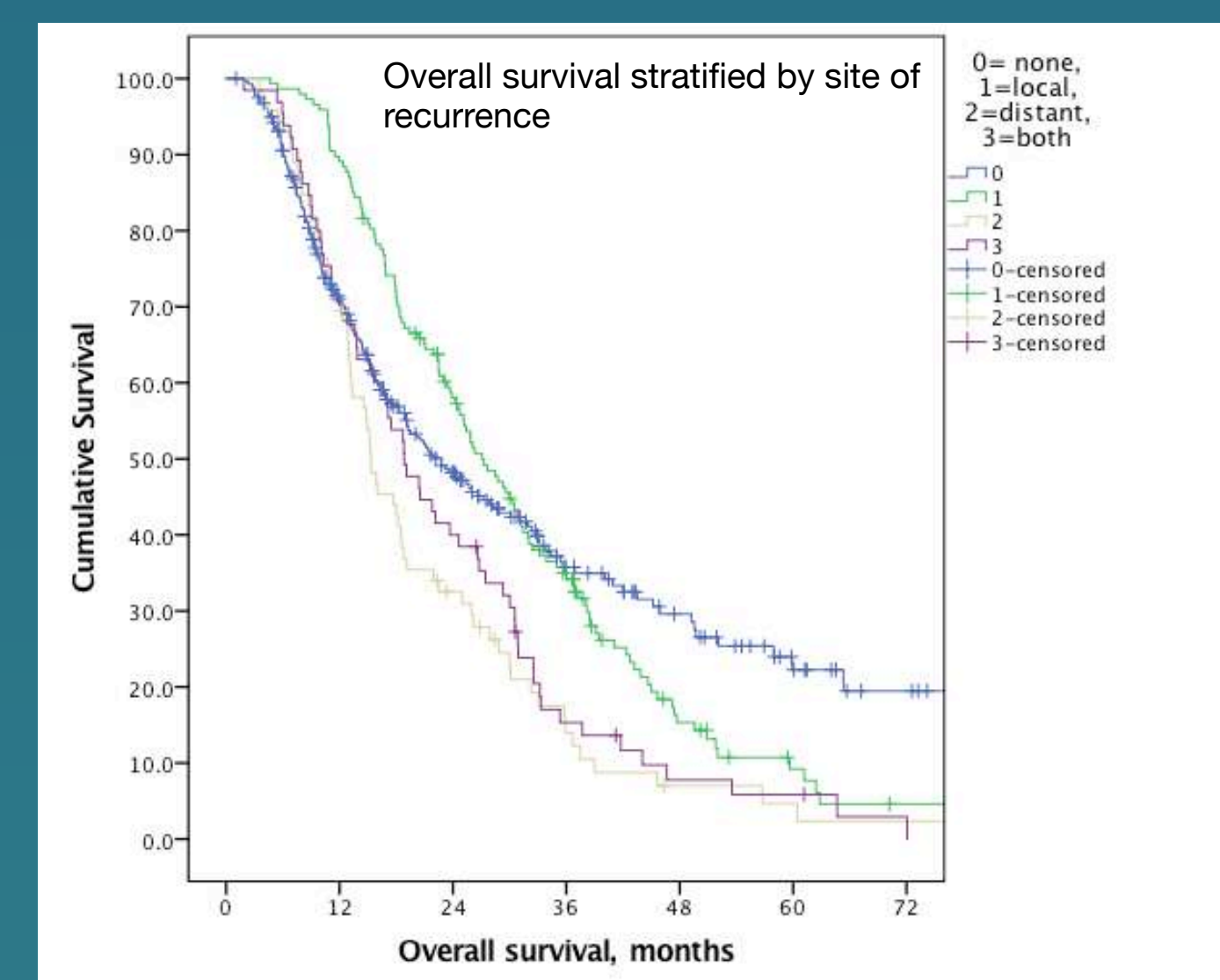
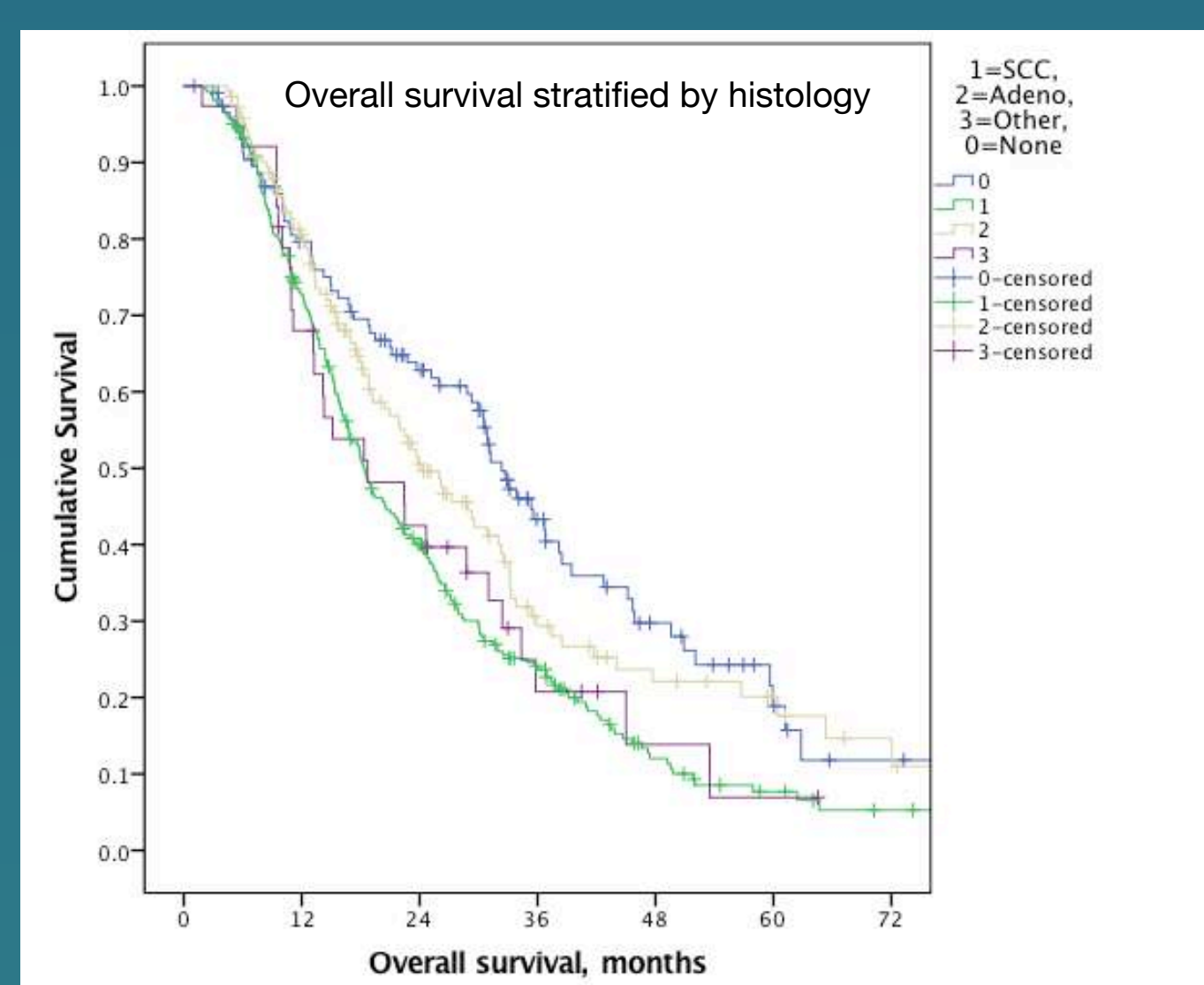
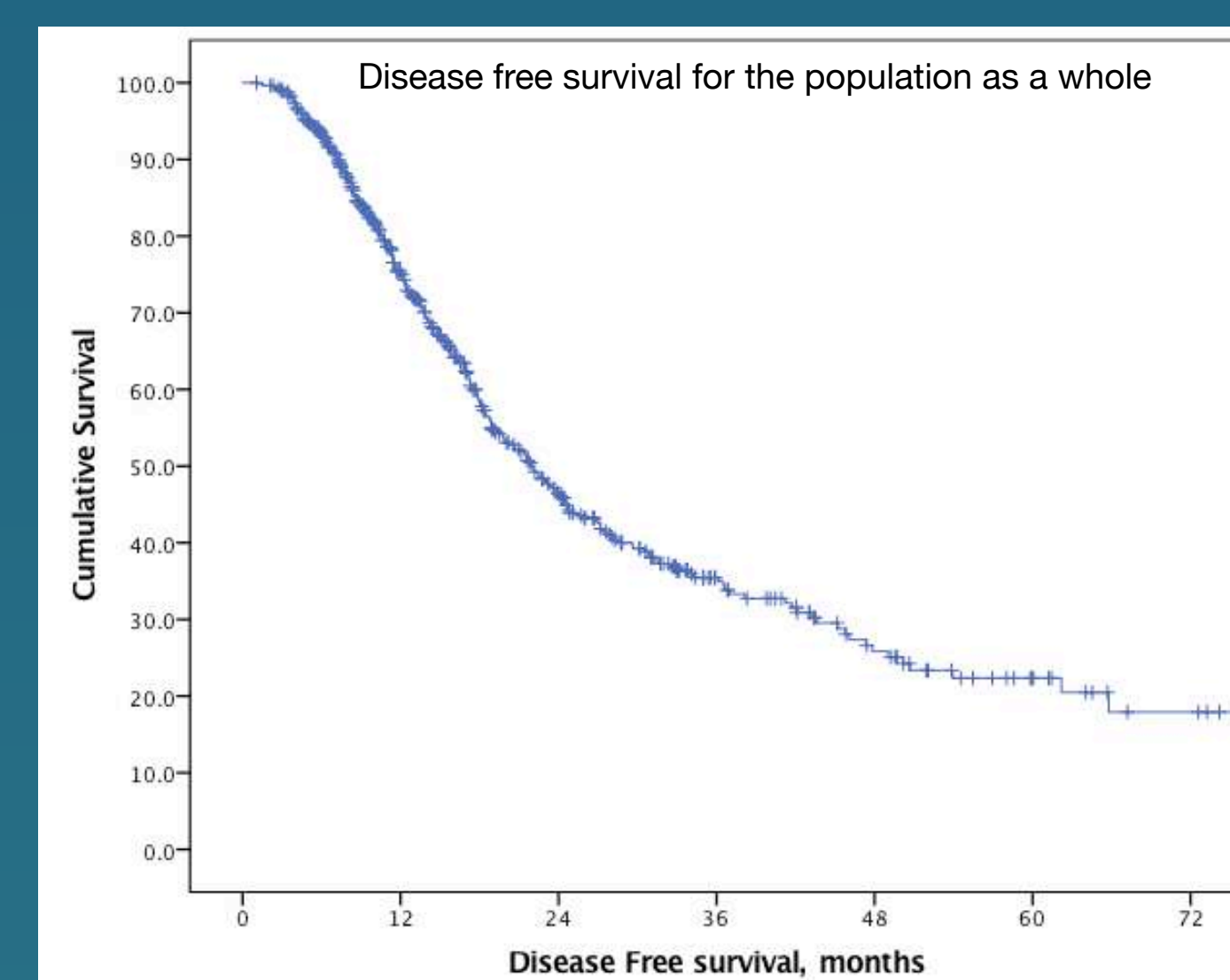
Median disease-free survival was 19 months.

Median overall survival of 22.5 months, with a 6.5% 90-day mortality rate.

Median OS was 31 and 20 months respectively for stage I and III NSCLC.



**On Univariate Analysis**  
 Histology: p=0.000  
 Stage: p=0.001  
 Response: p=0.000  
 Recurrence: p=0.022  
 Site of recurrence: p=0.000  
 Gender: p=0.101  
 Performance status: p=0.512  
 Chemotherapy: p=0.762  
 Radiotherapy regimen: p=0.736  
**On Multivariate Analysis**  
 Performance status: p=0.008  
 Histology: p=0.003  
 Stage: p=0.002  
 Chemotherapy: p=0.002  
 Response: p=0.000  
 Recurrence: p=0.000  
 Gender: p=0.844  
 Age: p=0.304  
 Radiotherapy regimen: p=0.945



**CONCLUSIONS**

This represents a large unselected cohort of patients treated with radical radiotherapy for NSCLC. It demonstrates both schedules are deliverable and safe with no statistically significant difference in survival. Future dose escalation studies (eg ADSCAN [1]) are required to develop these techniques to match outcomes reported by recent concurrent chemo-radiation studies [2].

**REFERENCES:**

1. Lawless C, Boyd K, Faivre-finn C, Fenwick J, Haswell T, Landau D, et al ADSCAN: A randomised phase II study of accelerated, dose escalated, sequential chemo-radiotherapy in non-small cell lung cancer (NSCLC). NCRI Cancer Conference. 3 Nov; Liverpool: 2015.
2. Bradley JD, Paulus R, Komaki R, Masters G, Blumenschein G, Schild S, et al. Standard-dose versus high dose conformal radiotherapy with concurrent and consolidation carboplatin plus paclitaxel with or without cetuximab for patients with stage IIIA or IIIB non-small-cell lung cancer (NSCLC) RTOG 0617: a randomised, two-by-two factorial phase 3 study. Lancet Oncol. Feb 2015;16(2):187-199.

