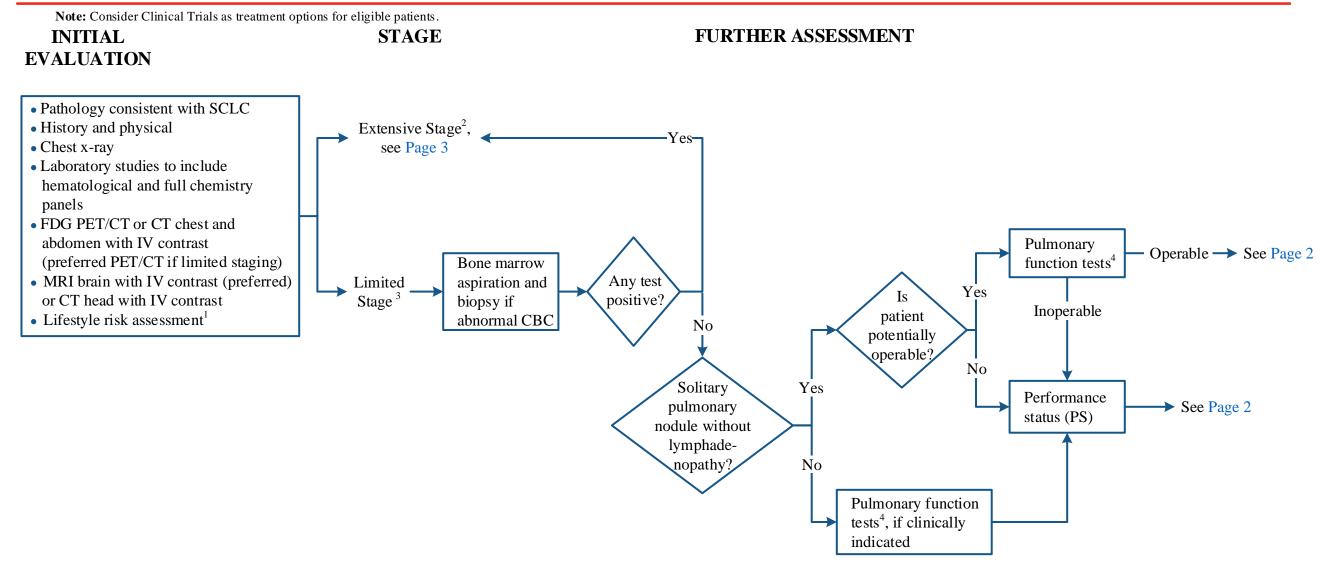
MDAnderson Small Cell Lung Cancer (SCLC)

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EBUS = endobronchial ultrasound

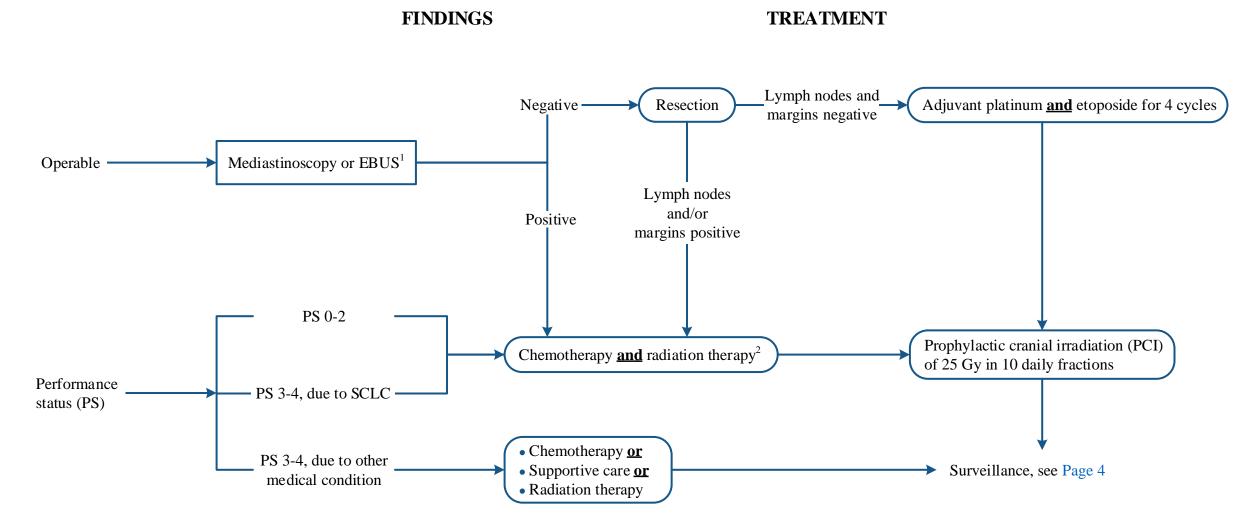
- ¹See Physical Activity, Nutrition, and Tobacco Cessation algorithms; ongoing reassessment of lifestyle risks should be a part of routine clinical practice
- ² Extensive stage: disease beyond ipsilateral hemithorax or malignant pleural effusion or obvious metastatic disease
- ³Limited stage: disease confined to the ipsilateral hemithorax within a single radiation port
- ⁴Pulmonary function tests include: spirometry pre-and-post-bronchodilators, xenon if clinically indicated, exercise oxygen consumption testing if clinically indicated

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Note: Consider Clinical Trials as treatment options for eligible patients.



EBUS = endobronchial ultrasound

¹Consider EBUS for patients treated with radiation therapy also

²Start radiation therapy within the first 2 cycles of chemotherapy

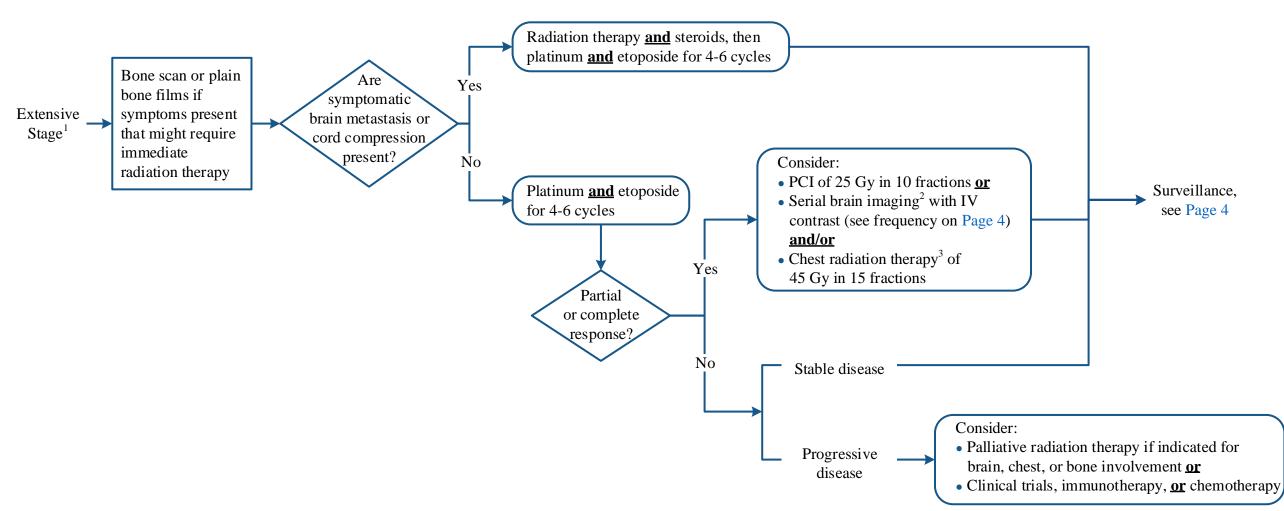
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TREATMENT

Note: Consider Clinical Trials as treatment options for eligible patients.

STAGE FURTHER WORKUP



¹Extensive stage: disease beyond ipsilateral hemithorax or malignant pleural effusion or obvious metastatic disease

²MRI brain preferred over CT as it is more sensitive in identifying brain metastases

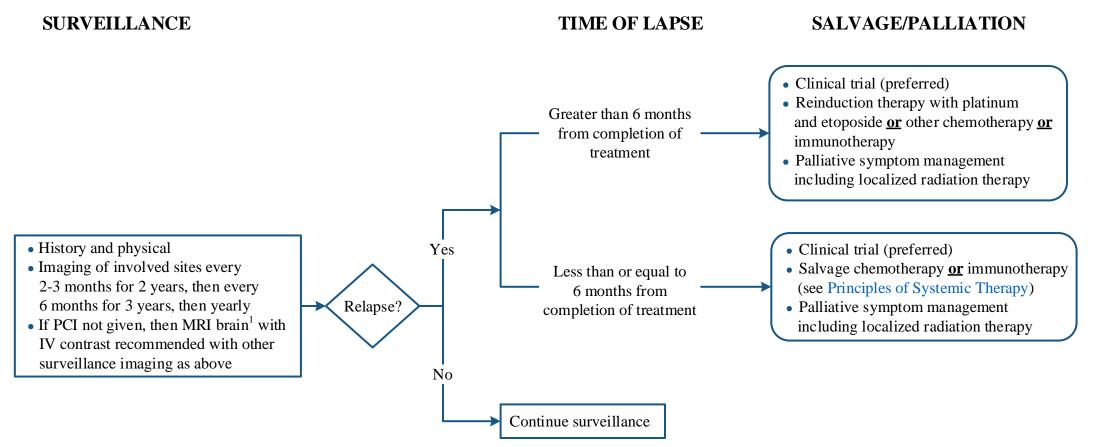
³ For selected patients with residual thoracic disease and low-bulk extrathoracic metastatic disease that has responded to systemic therapy

THE UNIVERSITY OF TEXAS MDAnderson Small Cell Lung Cancer (SCLC)

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PRINCIPLES OF SYSTEMIC THERAPY

First-line therapy

- Acceptable regimens for limited stage disease (maximum of 4-6 cycles) include:
 - $_{\odot}$ Cisplatin 60 mg/m² IV on Day 1 and etoposide 120 mg/m² IV on Days 1, 2, 3
 - $_{\odot}$ Cisplatin 80 mg/m² IV on Day 1 and etoposide 100 mg/m² IV on Days 1, 2, 3
 - $_{\odot}$ Carboplatin AUC 5-6 IV on Day 1 and etoposide 100 mg/m² IV on Days 1, 2, 3
 - During systemic therapy plus radiation therapy, cisplatin/etoposide is recommended (category 1)
 - The use of myeloid growth factors is not recommended during concurrent systemic therapy plus radiation therapy (category 1 or not using GM-CSF)
- Acceptable regimens for extensive stage disease (maximum of 4-6 cycles) include:
 - \circ Carboplatin AUC 5-6 IV on Day 1 and etoposide 100 mg/m² IV on Days 1, 2, 3
 - \circ Cisplatin 75 mg/m² IV on Day 1 and etoposide 100 mg/m² IV on Days 1, 2, 3
 - Cisplatin 80 mg/m² IV on Day 1 and etoposide 80 mg/m² IV on Days 1, 2, 3
 - \circ Cisplatin 25 mg/m² IV on Day 1, 2, 3 and etoposide 100 mg/m² IV on Days 1, 2, 3

Second-line therapy

- Clinical trial (preferred)
- If relapse occurs less than or equal to 6 months and performance status 0-2:
 - Topotecan PO or IV
 - Irinotecan
 - Paclitaxel
 - Docetaxel
 - Temozolomide PO
 - Nivolumab plus ipilimumab
 - Vinorelbine
 - Etoposide PO
 - Gemcitabine
- If relapse occurs greater than 6 months after completion of first-line therapy: original regimen
- Consider dose reduction or growth factor support for patients with performance status of 2 or age greater than or equal to 70 years

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Note: Consider Clinical Trials as treatment options for eligible patients.

PRINCIPLES OF RADIATION THERAPY

Radiation therapy for Limited Stage disease

- Radiation therapy should be given 1.5 Gy twice a day (with at least 6 hours between fractions) to a total dose of 45 Gy. In circumstances where twice daily fractionation is not feasible, an acceptable alternate schedule is 1.8 2.0 Gy/day to a dose of 60 70 Gy.
- Radiation therapy should be administered concurrently with chemotherapy, ideally beginning during cycle 1 of chemotherapy
- Radiation therapy should be delivered to original tumor volume unless there is marked risk of radiation pneumonitis; decrease field as tumor shrinks
- Appropriate schedule for prophylactic cranial irradiation (PCI) is 25 Gy in 10 fractions
- In patients receiving radiation therapy or chemoradiation with curative intent, treatment interruptions or dose reductions for temporary and manageable toxicities, such as esophagitis and myelosuppression, should be avoided. Careful patient monitoring and aggressive supportive care are preferable to treatment breaks in potentially curable patients. Patients should be evaluated at least once per every 5 fractions to monitor weight changes and toxicity.
- 45 Gy in 30 fractions over 3 weeks would not be recommended with concurrent chemotherapy on Day 1, if the DVH shows V20 more than 35% of target lesion. If the GTV is too large to meet dose volume constraints, give one cycle of chemotherapy or go daily fraction of radiation and cone down of the GTV after re-simulation after 2-3 weeks treatment. This will apply for patients who have FEV1 or DLCO less than 30% of predicted value.
- Elective nodal radiation therapy is not recommended

DVH = dose volume histogram GTV = gross tumor volume

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DEVELOPMENT CREDITS

This practice algorithm is based on majority expert opinion of the Thoracic Oncology Center Faculty at the University of Texas MD Anderson Cancer Center. It was developed using a multidisciplinary approach that included input from the following:

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