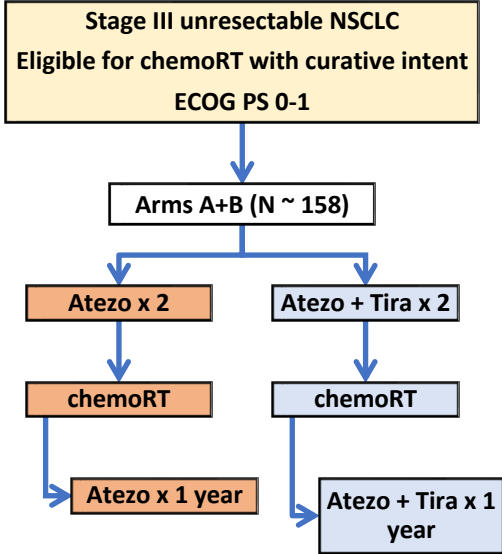
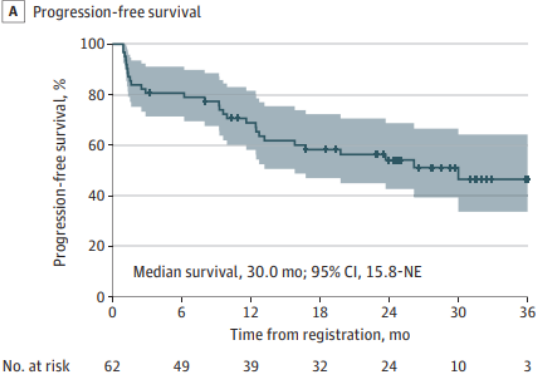
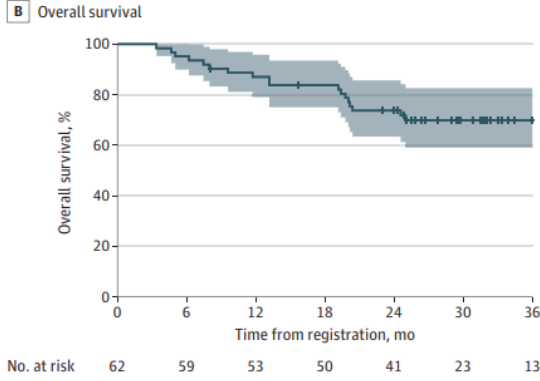


<b>AFT-57 / Randomized phase II trial of neoadjuvant and adjuvant atezolizumab with or without tiragolumab in conjunction with chemoradiotherapy for unresectable stage III NSCLC</b> ClinicalTrials.gov Identifier: NCT05798663	
<p>Schema</p>	<div style="display: flex; justify-content: space-between;"> <div style="width: 45%;">  </div> <div style="width: 50%;"> <p><b>Non-surgical Candidate considerations:</b></p> <ul style="list-style-type: none"> <li>• Multiple nodal stations</li> <li>• N3 nodes</li> <li>• Bulky nodes</li> </ul> <p><b>Key Eligibility Criteria:</b></p> <ul style="list-style-type: none"> <li>• Histologically confirmed stage III NSCLC per AJCC v8</li> <li>• ECOG PS 0-1</li> <li>• Adequate organ function</li> <li>• No active immunologic disorder</li> <li>• No active other malignancy</li> <li>• No contraindication to chemotherapy, immunotherapy, thoracic radiation</li> </ul> </div> </div>
<p>Study Status</p>	<ul style="list-style-type: none"> <li>• 21 selected sites out of 25</li> <li>• 5 sites active</li> <li>• 3 participants enrolled</li> <li>• First subject first visit: December, 2024</li> <li>• Expected last participant first visit: January, 2026</li> </ul>
<p>Objectives</p>	<p><i>Primary:</i> Progression-free survival (PFS)</p> <p><i>Secondary:</i> Overall survival (OS), overall response (ORR) and safety</p> <p><i>Principal Translational Science:</i> Assess immunologic markers including PD-L1 to develop biomarkers for efficacy and toxicity.</p>
<p>The predecessor AFT-16 trial of neoadjuvant atezolizumab in this setting provided proof of concept for safety of the neoadjuvant approach and for outcomes that compare favorably with standard of care therapy.</p>	<p>Progression-Free and Overall Survival and Exploratory Analysis of the Patients Who Completed Concurrent Chemoradiation Therapy (CRT)</p> <div style="display: flex; justify-content: space-around;"> <div style="width: 45%;"> <p><b>A</b> Progression-free survival</p>  </div> <div style="width: 45%;"> <p><b>B</b> Overall survival</p>  </div> </div> <p>No. at risk: 62, 49, 39, 32, 24, 10, 3 (for PFS); 62, 59, 53, 50, 41, 23, 13 (for OS)</p> <p>Shading indicates the 95% CI; tick marks, survival time censored. NE indicates not evaluable</p> <p>JAMA Oncol. 2024;10(9):1212-1219. doi:10.1001/jamaoncol.2024.1897 Published online July 25, 2024.</p>
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