

Sample Preservation



Tissue preservation methods can impact test results and therefore should be optimized to maintain sample quality.¹⁴

Decrease time to fixation



Immediate fixation (within **1 hour** of acquisition) is important for preserving tumor characteristics for diagnostic testing.¹⁵

Choose an appropriate fixative



Tissues should be preserved using appropriate solutions based on the testing that will be performed.³

10% NBF^{3,14}

- Most common
- **Recommended by guidelines**
- **Can be used for histologic and molecular analyses**

70% ethanol¹⁴

- Often used for cytology
- **Not suitable for most histologic analyses**

Optimize duration of fixation

Both underfixation and overfixation can affect sample quality and the accuracy of the biomarker test results.^{14,16}



Guidelines recommend fixation for **6 to 48 hours**, depending on the size of the sample.¹⁴

Proper sample preservation is critical for ensuring that the quality of the sample is appropriate for diagnostic testing.^{1,14}

A Focus on Guidelines:

Best Practices for Tissue Sample Collection in mNSCLC

In mNSCLC, tissue collection may be necessary at diagnosis and upon disease progression to test for biomarkers that can inform treatment decisions.¹

Guidelines recommend testing for multiple different biomarkers, including EGFR, ALK, BRAF, ROS1, and PD-L1; therefore, sufficient tissue must be collected and preserved appropriately to perform all biomarker tests.^{1,2}

According to the **AMERICAN THORACIC SOCIETY**¹

*“Potentially all treatment decisions, at the time of diagnosis and later, will be based on the information obtained from [the tumor sample]. Therefore, it is essential that the **specimen collection and processing procedures be optimized** to ensure that the quality and quantity of the specimen are adequate.”*

Tissue Collection



Multidisciplinary Team Collaboration



Collect sufficient sample

Multiple diagnostic tests are required to determine the most appropriate treatment options for a patient.^{1,3} To ensure that all necessary testing can be performed, guidelines agree that obtaining sufficient sample quantity is essential.^{1,2,4}

NCCN Clinical Practice Guidelines in Oncology (NCCN Guidelines®) ²	"[Biopsying physicians] should procure sufficient tissue to enable all appropriate testing."
American Thoracic Society ¹	"[Biopsying physicians] should adapt diagnostic procedures...to ensure that specimens are sufficient for molecular and immunohistochemical testing."
CHEST ⁴	"It is recommended that adequate tissue be obtained to accurately define the histologic type and to perform molecular analysis."

Coordinate rapid on-site evaluation (ROSE)

Guidelines recommend using ROSE, which allows a pathologist or technologist to assess sample quality and quantity on-site and thereby:

- Ensure that the sample is sufficient for biomarker testing^{4,5,8}
- Provide the biopsying physician with an opportunity to collect additional samples if necessary^{4,5,8}

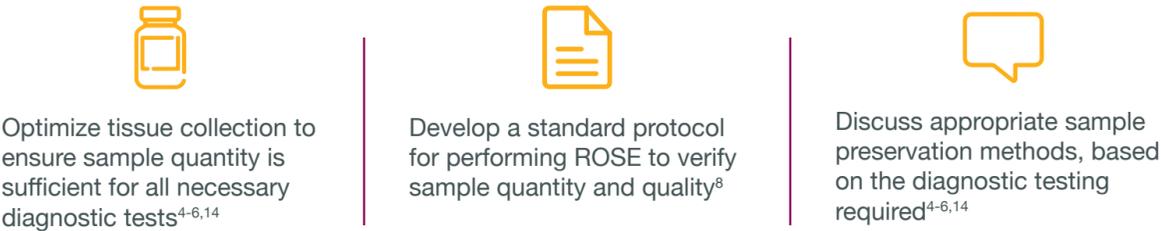
ROSE has been shown to:



Sufficient sample collection is critical for timely testing and informing treatment decisions at diagnosis and progression.^{1,4,8}

Because biomarker testing can inform treatment decisions, communicating with your multidisciplinary team to reduce turnaround time is critical.^{1,4,8} Multidisciplinary team collaboration has been shown to increase adherence to guidelines and improve patient outcomes, including survival.^{1,17}

Communicate with the medical oncologist and pathologist in your multidisciplinary team to:



Guidelines recommend that the biopsying physician play an active role in the multidisciplinary team to ensure accurate and timely biomarker test results and thereby inform treatment decisions.^{1,4,5,14}

For even more information about best practices in biomarker testing, visit AZOncologyID.com.

Choose an appropriate collection method

Different sample collection methods may yield different amounts of tissue and therefore must be chosen carefully.^{5,6} To maximize the amount of tissue available for diagnostic testing, guidelines recommend:

- Using image-guided procedures⁵
- Performing dedicated passes for ancillary studies⁵
- Obtaining concurrent CNB and FNAs if possible⁵

Both **CT-guided CNB** and **FNA** have been shown to yield sufficient tumor tissue for biomarker testing.⁷

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References: 1. Gaga M et al; ATS/ERS Task Force on the Role of the Pulmonologist in the Management of Lung Cancer. *Am J Respir Crit Care Med.* 2013;188(4):503-507. 2. Referenced with permission from the NCCN Clinical Practice Guidelines in Oncology (NCCN Guidelines®) for Non-Small Cell Lung Cancer V6.2018. © National Comprehensive Cancer Network, Inc. 2018. All rights reserved. Accessed October 9, 2018. To view the most recent and complete version of the guideline, go online to NCCN.org. 3. Lindeman NI et al. *Arch Pathol Lab Med.* 2018;142(3):321-346. 4. Rivera MP et al. *Chest.* 2013;143(5 suppl):e142S-e165S. 5. Roy-Chowdhuri S et al. *Arch Pathol Lab Med.* 2016;140(11):1267-1272. 6. Ofiara LM et al. *Front Oncol.* 2014;4:253. 7. Coley SM et al. *Cancer Cytopathol.* 2015;123(5):318-326. 8. Levy BP et al. *Oncologist.* 2015;20(10):1175-1181. 9. Trisolini R et al. *Chest.* 2015;148(6):1430-1437. 10. da Cunha Santos G et al. *Cancer Cytopathol.* 2013;121(1):4-8. 11. Choi SM et al. *Ann Thorac Surg.* 2016;101(2):444-450. 12. Chen C et al. *J Thorac Dis.* 2015;7(suppl 4):S238-S245. 13. Biliçeroğlu S. *Curr Opin Pulm Med.* 2017;23(3):247-253. 14. Lindeman NI et al. *Arch Pathol Lab Med.* 2013;137(6):828-860. 15. Hammond ME et al. *Arch Pathol Lab Med.* 2010;134(7):e48-e72. 16. Engel KB et al. *Arch Pathol Lab Med.* 2011;135(5):537-543. 17. Bilfinger TV et al. *Clin Lung Cancer.* 2018;19(4):346-351.

