IGH-BCL2 Fusion, t(14;18) by FISH

Indications for Ordering
Diagnosis of follicular lymphoma (FL) in conjunction with clinical, morphologic, and flow cytometric data

Test Methodology
- Fluorescence in situ hybridization (FISH)
- Dual color, dual fusion probes detect t(14;18) or IGH-BCL2 gene rearrangement

Tests to Consider

Typical Testing Strategy
FL can often be diagnosed using the combination of morphology, immunohistochemistry (IHC), immunophenotyping, and clinical presentation
- Lymph node biopsy with morphologic and IHC evaluation
- Leukemia/lymphoma phenotyping by flow cytometry
- CD5 negative and CD10 positive suggests FL
- Bone marrow (BM) evaluation for staging – marrow involvement is common

Chromosome analysis by FISH – may be useful when the above combination does not yield a diagnosis
- Presence of IGH-BCL2 fusion, t(14;18) supports a diagnosis of FL

Primary Test
IGH-BCL2 Fusion, t(14;18) by FISH 3001298
- Most sensitive method to detect IGH-BCL2 fusion in formalin-fixed, paraffin-embedded (FFPE) tissue specimens

Related Tests
Leukemia/Lymphoma Phenotyping Evaluation by Flow Cytometry 3001780
- Aids in diagnosis of hematopoietic neoplasms

Chromosome FISH, Interphase 2002298
- Specific FISH probe for t(14;18) must be requested

Chromosome Analysis, Bone Marrow 2002292
- Diagnosis, prognosis, and monitoring of lymphoma in BM

Chromosome Analysis, Solid Tumor 2002296
- May identify additional, useful cytogenetic abnormalities in tissues that are not targeted by FISH assays

Disease Overview
Incidence – most common non-Hodgkin B-cell lymphoma

Signs/symptoms
- Early – usually asymptomatic
- Advanced
  - Peripheral and central lymphadenopathy
  - Splenomegaly
  - BM involvement
  - B symptoms – fever, night sweats, weight loss

Diagnostic Criteria for Follicular Lymphoma
- Morphology
  - Lymph node – neoplastic mixture of centrocytes and centroblasts recapitulating germinal centers of secondary lymphoid follicles
  - BM – paratrabeicular lymphoid aggregates
- Flow cytometry
  - CD5 negative; CD10 positive monoclonal B cells

Genetics
Gene – IGH-BCL2

Structure/Function
- Translocation juxtaposes the immunoglobulin enhancer region (IGH, 14q32) with the BCL-2 oncogene (18q21)
- Translocation causes overexpression of BCL-2, an anti-apoptotic protein, which prevents normal cell death

Test Interpretation
Results
- Positive – presence of the t(14;18) translocation substantiates a diagnosis of FL
- 20% or more of the cells examined were abnormal
- Negative – absence of t(14;18) translocation

Limitations
- IGH-BCL2 fusion, t(14;18) by FISH has not been validated for tissue fixed in alcohol-based or non-formalin fixatives
- Negative result does not exclude the possibility of translocations involving other partners nor rule out FL