

## HISTOLOGY

### Inspector Instructions:

 <b>READ</b>	<ul style="list-style-type: none"> <li>Sampling of histology policies and procedures</li> <li>Sampling of specimen preparation records</li> <li>Sampling of histology QC policies and procedures</li> <li>Sampling of QC records (histochemical)</li> <li>Sampling of records of daily review of histologic slide quality</li> <li>Sampling of immunofluorescence QC records</li> <li>Sampling of IHC policies and procedures</li> <li>Sampling of new antibody validation/verification records</li> <li>Sampling of new reagent/shipment confirmation of acceptability records</li> <li>Sampling of antibody QC records</li> <li>Sampling of buffer pH records</li> <li>Sampling of batch control records</li> </ul>
 <b>OBSERVE</b>	<ul style="list-style-type: none"> <li>Sampling of tissue blocks (identification)</li> <li>Sampling of slides (labeling, quality)</li> </ul>
 <b>ASK</b>	<ul style="list-style-type: none"> <li>How does the histology section ensure specimen identity throughout processing?</li> <li>How does your biorepository validate/verify new antibodies?</li> <li>How does your biorepository confirm the acceptability of new reagent lots?</li> <li>How does your biorepository distinguish non-specific false-positive staining from endogenous biotin?</li> </ul>
 <b>DISCOVER</b>	<ul style="list-style-type: none"> <li>If problems are identified during the review of histology procedures, further evaluate the responses, corrective actions and resolutions</li> <li>Select a representative specimen and follow from receipt in the department through accessioning, grossing, processing, time reported and availability in the LIS</li> </ul>

#### BAP.05330 Specimen Preparation Records

Phase I

**The histology section retains records of the number of blocks, slides, and stains prepared and appropriately denotes the block from which the slide was prepared.**

#### BAP.05332 Cross-Contamination - Histology

Phase II



**The biorepository prevents cross-contamination of specimens in the histology section.**

**NOTE:** The process must address steps to prevent cross-contamination during the various phases of tissue handling including: processing, embedding, microtomy, and slide preparation. Problems with cross-contamination must be addressed in the biorepository quality management system.

*Instruments must be clean and well-maintained (eg, tissue processors, embedding centers, dispensers, floatation baths, staining and coverslipping equipment).*