FOR MORE INFORMATION, PLEASE CONTACT YOUR AKOYA SALES REPRESENTATIVE.

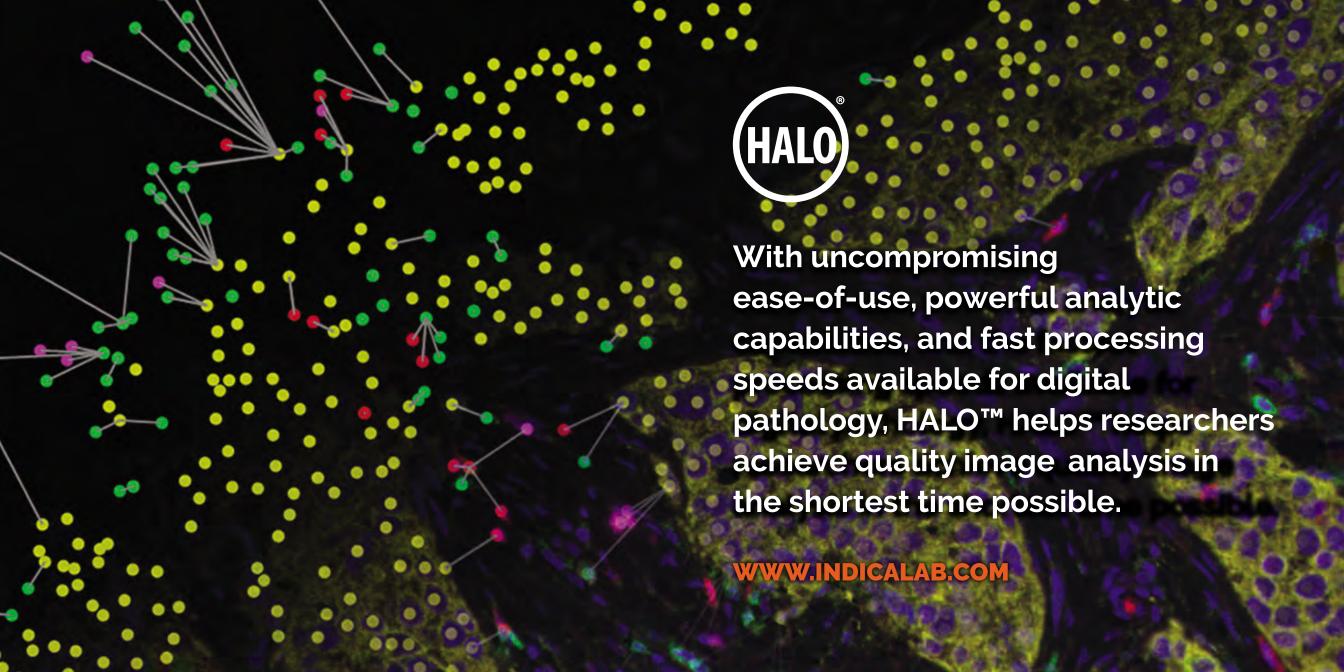




A CUTTING-EDGE IMAGE ANALYSIS
PLATFORM FOR DIGITAL PATHOLOGY



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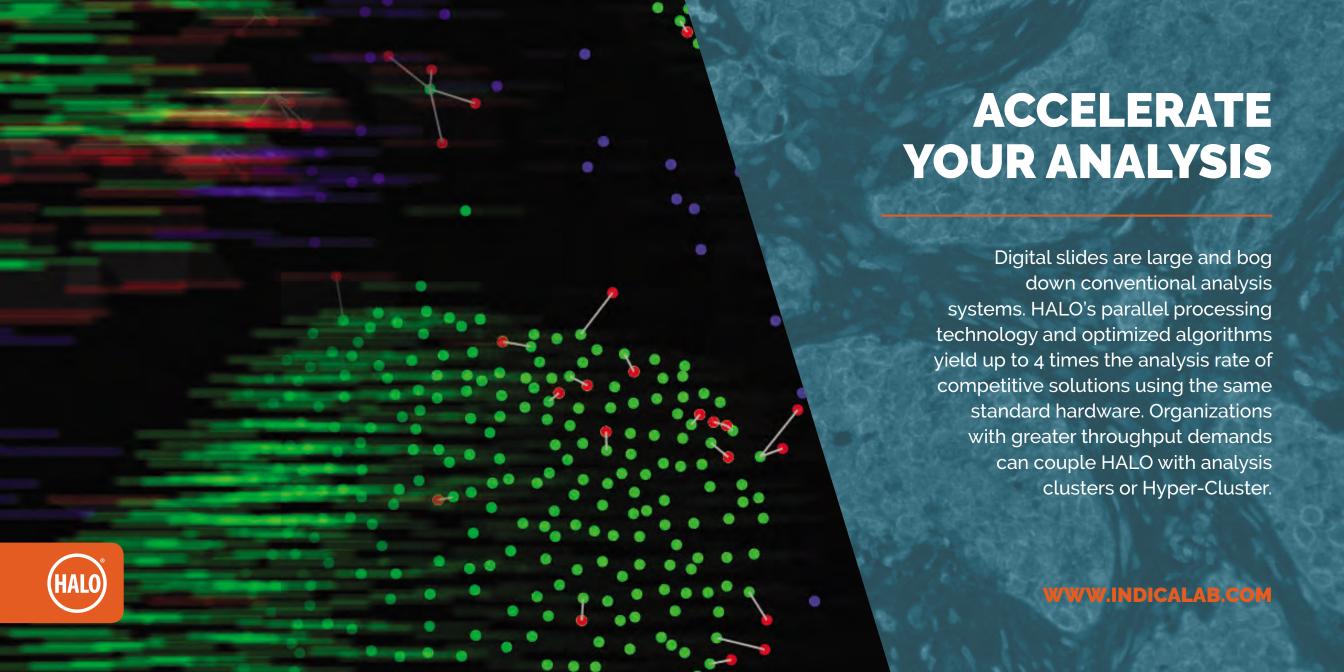


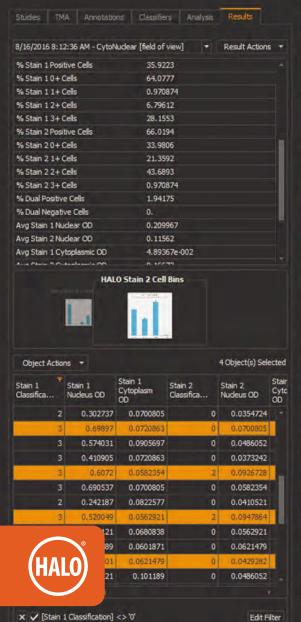


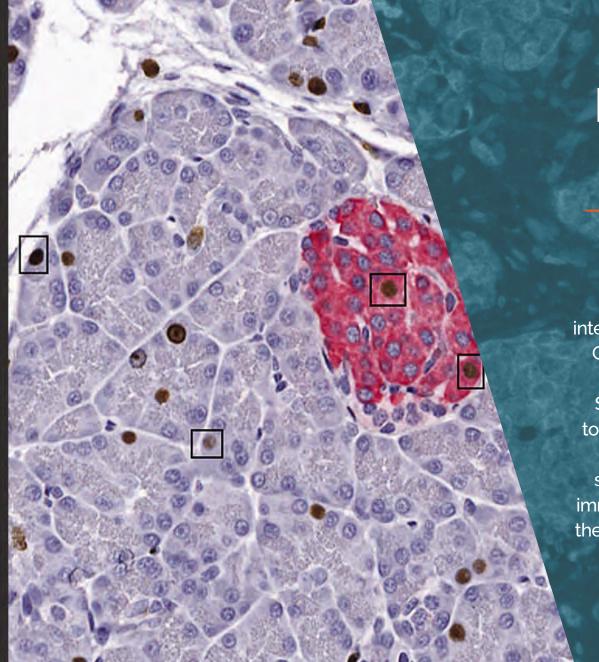


# RAPIDLY ANNOTATE SLIDES

Do you dread annotating digital slides using your current system? Cut annotation time in half with HALO's intuitive, flexible annotation tools. Draw free-hand or use the magnet pen tool to automatically 'snap' annotations to your tissues. Easily redraw, amend, move, rotate or copy-paste existing annotations. Better still, teach the classifier module to automatically select tissues of interest for analysis across all images in a study.







# EASILY EXPLORE CELLULAR DATA

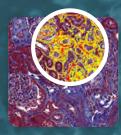
expression data on a cell-by-cell basis across entire tissue sections and maintains an interactive link between cell data and cell image. Click on any cell in the image and immediately see analysis outputs for that specific cell. Sorting and filtering capabilities allow the user to mine millions of cells while visually assessing corresponding cell populations. For example, sort cells according to biomarker intensity and immediately locate cells with highest intensity in the image. Just imagine the endless possibilities.



# GROW WITH HALO

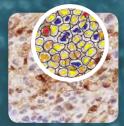
HALO offers a modular platform that can expand with your needs. Start with a few modules, and add more as your needs change. Use HALO on a single workstation or ramp up to a server-based license for your entire group. HALO is flexible enough for any budget.

## **GENERAL BRIGHTFIELD APPLICATIONS**



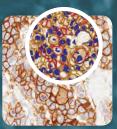
#### **AREA QUANTIFICATION**

Measure positive stain area and optical density for up to three stains, categorize pixels according to stain density (negative, weak, moderate, or strong positive), and measure percentage colocalization of stains. Included free with any MALO bundle.



#### **SINGLE & DUAL STAIN IHC**

Measure cellular positivity for one or two IHC stains, measure stain localization within subcellular compartments, and categorize each cell according to stain optical density (negative, weak, moderate, or strong positive) and co-expression of stains on a per cell basis.



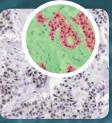
#### **MEMBRANE IHC**

Quantify intensity of membrane stains on a per cell basis and categorize cells according to stain optical density (negative, weak, moderate, or strong positive).



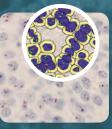
#### **OBJECT QUANTIFICATION**

Count microvessels and other macro-cellular objects. Measure one or two differently stained objects in a single tissue, quantify object area, optical density, and colocalization of object(s).



#### TISSUE CLASSIFICATION

Separate tissues classes, such as tumor and stroma, across whole slides using a learn-by-example approach. Can be used in conjunction with all other modules to select specific tissue classes for further analysis. Available for both brightfield and fluorescent applications.

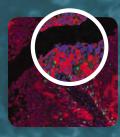


#### ISH QUANTIFICATION

Analyze CISH, SISH, RNA ISH, or virtually any type of 'spot' within the nucleus and/or cytoplasm. Quantify one or two probes and report number of probe signals per unit area or per cell.

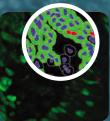


### **GENERAL FLUORESCENCE APPLICATIONS**



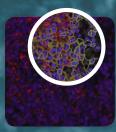
#### **AREA QUANTIFICATION FL**

Measure positive dye area and intensity for up to three dyes, categorize pixels according to stain intensity (negative, weak, moderate, or strong positive), and measure percentage colocalization of dyes. Included free with any HALO bundle.



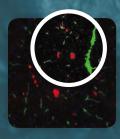
#### **SINGLE & MULTIPLEXED IF**

Measure cellular positivity for up to three fluorescently-labeled cytonuclear markers. Measure localization within sub-cellular compartments, categorize cells according to stain intensity (negative, weak, moderate, or strong positive), and measure co-expression of iomarkers on a per cell basis.



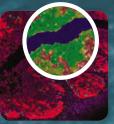
#### **MEMBRANE IF**

Measure intensity of fluorescently-labeled membrane markers on a per cell basis and categorize cells according to intensity (negative, weak, moderate, or strong positive).



#### **OBJECT QUANTIFICATION IF**

Count microvessels and other macro-cellular objects. Measure one or two objects stained with different fluorophores, quantify object area, intensity, and colocalization.



#### **TISSUE CLASSIFICATION**

Separate tissues classes, such as tumor and stroma, across whole slides using a learn-by-example approach. Can be used in conjunction with all other modules to select specific tissue classes for further analysis. Available for both brightfield and fluorescent applications.

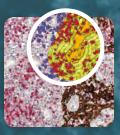


#### **FISH QUANTIFICATION**

A collection of modules for quantifying RNA and DNA fluorescence in situ hybridization. Quantify two probes (amplification-deletion FISH and break-apartfusion FISH) or four probes (multiplexed DNA and RNA FISH) and report number of probe signals per cell.

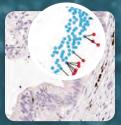


### **ONCOLOGY APPLICATIONS**



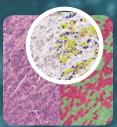
#### **IMMUNE CELL QUANTIFICATION**

Quantify cells labeled with common immune cell markers (CD3, CD4, CD8, PD1, PDL1). When used in combination with a second stain, the proximity of immune cells to the second stain can also be measured. For example, proximity of CD8+ immune cells to cytokeratin + tumor area.



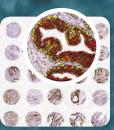
#### **SPATIAL ANALYSIS**

Plot cells on the same image or on registered serial sections together to measure nearest neighbor distances, count target cells within a user-defined proximity of reference cells, or count number of cells within a certain distance of a manually annotated interface line.



#### **SERIAL SECTION ANALYSIS**

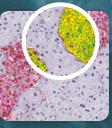
Using registration, serial section analysis allows you to transfer tissue classification masks to serial sections. For example, use H&E to classify tumor from stroma and use resulting classifier mask to quickly select tumor for analysis on registered IHC stained sections.



#### TISSUE MICROARRAY SEGMENTATION

Easily segment TMA slides into individual spots for further analysis. Create TMA maps manually or import from spreadsheet, remove spots with insufficient tissue or artifacts, and run analysis across one or all TMA spots in batch.

### **METABOLISM APPLICATIONS**



#### **ISLET QUANTIFICATION IHC**

Count islets stained with up to two stains in addition to nuclear counterstain, quantify number of stained cells in each islet, and determine area of islets stained with each islet-specific stains. For example, count insulin – and glucagon-positive cells per islet.



#### **ISLET QUANTIFICATION FL**

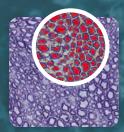
Count islets stained with up to three dyes in addition to nuclear dye, determine number of cells positive for each dye within each islet, and quantify islet area and total tissue area that is positive for each dye. For example, count number of insulin, glucagon, and somatostatin-positive cells per islet.



#### **VACUOLE QUANTIFICATION**

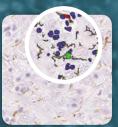
Measure vacuole area, diameter, perimeter and number of vacuoles per cell. Quantify lipid droplets in any tissue, including liver (steatosis) and adipose tissue.

## NEURO | MUSCULAR | VASCULAR APPLICATIONS



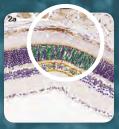
#### **AXON QUANTIFICATION**

Quantify axons in cross sections. Calculate axon area, myelin area, myelin thickness, inner and outer axon diameter, and G-ratio.



#### MICROGLIAL QUANTIFICATION

Quantify microglial activation based on length and thickness of microglial processes. Outputs total number of active and inactive microglia, average number of processes per cell and area, length, and thickness of processes per cell.



#### **LAYER THICKNESS**

Measure thickness of specific tissues or tissue areas, such as the retinal nuclear layers or the epidermal strata. **Included free with any HALO bundle**.



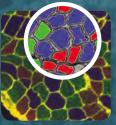
#### **BRANCH STRUCTURE**

Quantify labeled branched objects, for example retinal vascularization or nerve branching. Measure number of branch points and end points, length, thickness and area of branches. Available for both brightfield and fluorescent applications.



#### **MUSCLE FIBER QUANTIFICATION**

Quantify area, perimeter and mean/median/maximum diameter of muscle fibers stained with laminin or other stain to highlight fiber membrane. Count muscle fibers with central nucleus (indicative of regeneration).



#### **MUSCLE FIBER QUANTIFICATION FL**

Count, measure and classify muscle fibers based on expression (or co-expression) of up to three markers.



# SCIENCE + Cur = Better RESULTS

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