

# **GELDOC UVITEC-1D**

License-free software for unlimited users. Multi-user platform for advanced image acquisition, editing and analysis.











**APPLICATIONS** 

- Software comes with default protocols for easier, faster use:
  - > Default protocols are already configured, optimized and ready-to-use
  - > Default protocols cannot be modified nor deleted, to avoid loss or confusion
  - > Default protocols will automatically appear when modules are incorporated into the system
  - > Choose from numerous default protocols: UV Fluorescence, Gel Marker, Epi-Blue (with FireReader V10 only), Epi-UV (with FireReader V10 only) depending on system configuration
- Create single or multi-channel/sequence protocol
- Define/modify/customise/delete as many user-defined protocols as desired:
  - > All parameters easily customisable to users' sample requirements and preferences
  - > Change/adjust lighting, emission filter, lens aperture, camera sensibility, focus, display enhancements, image palette...
  - > Use the camera's full resolution or binning modes to increase sensitivity
  - ➤ Define a specific location for every single protocol to automatically save pictures
- Extended resolution: effective pixels technology allows extension of camera's native resolution by software interpolation

#### **ACQUISITION**

- Exposure of 0.1 second to 160 minutes or more
- 3 different acquisition modes to choose from, and easily switch from one to another
  - Auto acquisition mode:
    - ➤ Automatically selects the appropriate filters, light sources and camera settings for the required application
    - > 1-click image acquisition process (fully-automated) for extremely easy and fast image acquisition
    - > Fully-automated capture: optimal exposure time is automatically calculated through advanced algorithm
  - Manual acquisition mode:
    - > Exposure time is entered as desired and can be easily adjusted at all times
    - > Live & live-3D modes during acquisition with all fluorescence applications (live-picture displayed during acquisition)
  - Preview acquisition mode:
    - Advanced live mode to preview sample and image direct visualisation
    - > Easily position sample on the tray/table, and adjust the lens aperture and focus while positioning the sample

### **VISUALISATION**

- Autofit: untick the autofit button to view the whole image, regardless of the window size
- Zoom in or out on the image
- Visualise any picture in 3D:
  - ➤ Instantly check levels of signal and saturation
  - Apply a grid on 3D view to find out about the intensity of signals (shades of grey)
  - Quickly adjust 3D image colours and levels with the palette button
- Compare images together:
  - Compare as many images as desired with one simple click
  - Compare images in 3D
  - Normalise all images to the designated reference one

- Click the saturation button to highlight the image's over-saturated areas in red
- Check the image dynamic at a simple glance
- Change the image display by modifying the image greyscale using 5 image pre-sets (100%, 80%, 60%, 40%, 20%)
- Invert display option, to inverse the grey level of the image
- GLP (Good Laboratory Practice): retrieve all technical details about the image (date, exposure time, system, aperture...), print, save, export

## **EDITING / POST PROCESSING**

- Save images in tiff, tagged tiff, jpeg or bitmap format, 16-bit or 8-bit images
- Rotate the image with different tools: 90° left, 90° right, horizontal mirror, vertical mirror, rotate by angle...
- Add text: annotate images with text and arrows, modify font colour, select the text orientation
- Insert object: use an arrow, symbol or a rectangle to emphasize on a particular area
- Add the current time, or the GLP-file time to the image
- Crop the image: cut out a portion of an image as desired
- Merge or paste a marker: choose and adjust the intensity and proportion of marker and signal
- Background subtraction: separate foreground objects from the background clutter
- Artefact correction: easily remove dust or small defect from an image
- Flat field: correct uneven illumination intensity for both background and signal

#### **ANALYSIS**

- Free, unlimited analysis tools for Quantification, Molecular Weight, Colony Counting and Distance Calculations
- Analyse DNA/RNA gels, protein gels, colony dishes, western blots, dot blots, fluorescent gels and blots...
- Quantification with volume %, volume, intensity and area:
  - Quickly position boxes on top the bands of interest
  - Auto-save boxes selection, order and position for next quantification
  - > Background subtraction option with user-friendly 1D graph, either rolling ball or linear
  - Add as many band separations as desired for one single box or apply to all bands at once to save time and use the same area across the image
  - > Normalise by modifying the quantity of reference and choosing the band(s) of reference to compare others with
- Molecular weight:
  - ➤ For either horizontal or vertical gels
  - Easily select the desired number of lanes and optional gap between lanes
  - Automatic band detection, manually adjustable with user-friendly 1D graphs
  - > Quickly create a new marker, save or load an existing marker
  - > Easily choose the marker lane, then automatic or manual marker assignment
- Colony Counting:
  - > Choose the area of analysis as desired: circular, rectangular or customised, allowing you to easily draw the area of interest
  - > Select the desired calculation method: automatic or manual
  - > Automatic software counting based on the requested detection intensity settings
  - > Calculates green and red colonies
  - Change/adjust detection settings (shades of grey) to include or exclude a certain intensity, thus specific colonies
- Distance Calculation:
  - Easily select the desired number of lanes and optional gap between lanes
  - Automatic band detection, manually adjustable with user-friendly 1D graphs
  - > Set Reference (origin) and Front (final) values to calculate distances
- All results and data exportable to Excel at a simple click
- Print, copy, save data

