User Manual

S A B I A

Software

User Guide

EBIOGEN, Inc.

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I. Introduction

1. Overview of SABIA (Solution for Bio-Image Analysis) Software

SABIA (Solution for Bio-Image Analysis) is an analysis software for cell and tissue image which comes from microscope and other scientific instruments. It calculates automatically cell confluency, cell number, a ratio of different characteristics, area, and length by using color difference. Numerical output makes it easier for researcher to distinguish accurately the experimental results.

Function	Contents	
	Calculation automatically cell confluency	
Cell Confluency	from cell image	
	Real time image capture and to analysis	
	Counting automatically cell number	
Cell Counting	Measurement of each cell length	
	Calculation of aggregation ratio	
	Comparison the ratio of characteristics of	
	tissue	
Image Analysis	Measurement of length and area of cell or	
	tissue	

2. Major Function

3. Computer Specifications

- 1) Microsoft Window XP or above (recommend Window 7 or above)
- 2) Dual-core or higher processor and more than 4GB main memory
- 3) 1600*900 or higher resolution
- 4) Necessity of Internet connection depending on version.
 - (1) License renewal version: check to connect internet every operating program.
 - (2) Free-trial version: check to connect internet every operating program

4. Purchase Software and Technical Support

EBIOGEN, Inc.	Homepage: www-e-biogen.com	Tel. 82-2-3141-0791
MeTooSoft, Inc.	Homepage: www.metoosoft.com	Tel. 82-1661-3687

II. Install

1. Software Download

Visit EBIOGEN's homepage.

www-e-biogen.com > PRODUTS > Software & Data Analysis > SABIA Software Download

2. Installation and Authentication

Open an installation file, then execute 'exe'. Choose between **Online Authentication** and **Free Trial**. Click **Next**.

[Select Authentication Method]



[Online Authentication]

👌 Software Use Aut	hentication
	Eonline Authentication] Authentication Key can be found on authentication e-mail,, The computer is registered with the key once authenticated, Authentication Key has the following format, Authentication Key : XXXX-XXXX-XXXX-XXXX-XXXX ABCDE-FGHIJ-KLMNO-PQRST-UVWXY
	Previous Next (N) Authenticate Cancel

- 1) Enter 25 digits of authentication key that you received when you purchased the software. Click **Authenticate**.
- 2) Please check your Internet connection before Online Authentication.

[Free Trial]

슃 Software Use A	Authentication
	[Free Trial]
	Company/Organization :
	Name : Tel :
	E-mail :
	Request an authentication key by e-mail
	Authentication Key :
	$\ensuremath{\ll}$ Email with Authentication Key will be sent to the email address above.
	X MAC Address : 7427EA497237
	If Free Trial Authentication is not available, copy the MAC Address above and send it to [service@e-biogen.com] with Name, E-mail, and Name of Company/Organization. We will promptly respond to it with Free Trial Authentication Key.
	Previous Next (N) Authenticate Cancel

- 1) 15-day free trial is available before purchasing SABIA software
- Enter the information about Company/Organization, Name, Tel, and E-mail. Click Request an authentication key by e-mail. Authentication key will be sent to e-mail address you entered above.

[ATTENTION] It might be fail to get the authentication key when using company or organization's e-mail because of a security. Try it again with a personal e-mail. If it is still fail despite trying above mentioned way, please contact us with copied MAC address and your information (service@e-biogen.com).

III. Calculate Cell Confluency

1. Overview

SABIA calculates automatically cell confluency on the cell culture dish by using color differences. It produces the numerical output and is helpful to achieve an optimal reproducibility of results. There are two ways for calculating cell confluency. Either saved images or captured images in real time can be used. A tool for capture image is useful when you use microscopy.

2. How It Works

1) Select Folder

At the left side of the screen, select a folder which images that will be analyzed are located. Selected file can be confirmed using **Preview** and **Original Image** tab.



2) Set Calculation Option for Cell Confluency

It does not matter an order of setting between Color Similarity and Excluded Colors.



- (1) Select Excluded Colors
 - Check any excluded color boxes (up to 4) and click checked color box for choosing a color.
 - ② Touch the color from the original image. That color will be recognized as a background color and excluded from the calculation of cell confluency.
 - ③ Unchecked color box will be excluded from the calculation even if you select color.
- (2) Set Color Similarity
 - ① **Color Similarity** means a permissive range of color. The range of color similarity can be set from 0 to 100(%). Both using up & down button or inputting digits are available.
 - 2 Higher color similarity value, realize it more as a background color.
 - ③ Appropriate range depends on image's characteristics or color. In first time, recommended color range is from 20 to 30%. Then, adjust it following the calculated value.

[ATTENTION] If value of color similarity is too big, all color would be the excluded color resulting in the color of cell would be recognized as the background.

3) Calculate Cell Confluency: Click Calculate Cell Confluency burton on the menu bar.



- (1) Calcultated cell confluency will be showed beside its file list. Excluded color that you selected will be changed to purple color. This means that it was recognized as a background and is not contain the result of cell confluency. **Converted Image** helps to divide clearly between cells and background.
- (2) If the result of cell confluency is not satisfied, follow next two steps.
 - ① Change the percentage of **Color Similarity**. Then, recalculate it.
 - Check Excluded Colors boxes and add more color following aforementioned protocol.
 It is recommended that choosing another color excepting purple color in converted image. Then recalculate it.
- (3) Setting of **Excluded Colors** and **Color Similarity** will be saved when existing the program and can use it next analysis of other image. (More than an image which came from the same experimental condition can be calculated by **Calculate Total Cell Confluency** with the calculation option.)
- (4) For saving the result of calculation, click **Copy List**, then paste it to Excel or Notepad.
- (5) For saving the converted image, press Save Converted Image. It will be saved in a sub-

folder where the original image is located.

(6) Click **Calculate Total Cell Confluency** to calculate simultaneously all images in the selected folder.

3. Example of Cell Confluency Calculation

Better result can be produced by only changing **Color Similarity**. 61.3% of cell confluency is calculated when select 'an **Excluded Colors'** and '20% of **Color Similarity**.

1) For the deletion of background in the yellow part, the percentage of **Color Similarity** will be increased.



- 2) Only changing **Color Similarity** from 20% to 40%, 30% of cell confluency was obtained.
- 3) Higher **Color Similarity** makes the clearer delineation of cell in both areas, selected yellow area and whole area.



4. Set Capture Area

General video, microscopic video, or any kind of image can be used directly by using capture



1) Image Capture

- (1) Click Set Capture Area, then follow a pop-up instruct on monitor.
- (2) Click and drag a certain area to be analyzed.

- (3) Press E to exist Set Capture Area. For switching monitor when using dual monitor Press M.
- (4) Press **Esc** to cancel of image capture.
- (5) Captured image would be appeared on **Preview** and it will be automatically used after saving image. To save image, enter an image name, then click **Save**.
- (6) Captured image will be saved on a subfolder of the current folder.

IV. Cell Counting

1. Overview

A main function of cell count is an auto calculation of the number of cell (nuclear). Not only counting the number of cells in any kinds of cell images, it enables to measure the length of each cell. It provides reliable and reproducible data and disencumbers from both laborious and time-consuming tasks.

The basic concept of handling program is the same with the calculation of cell confluency. For cell counting, one of added options is Include Selected Color of which function works in opposition to Exclude Selected Color. Chosen color from Include Selected Color will be counted as cells, not be considered as background. Second option is to specify the area where cell counting takes place. Last option is to measure the cell length and aggregation.

2. How It Works

1) **Select Folder**: Open the folder which cell images are located by clicking **Select** in the upperleft side of screen. Select an image to be calculated.



1) Calculation Option for Cell Counting

(1) Set Color

There are two options, **Excluded Colors** and **Included Colors**. Excluded Colors is recommended when image's color is single. Included Colors is recommended when image's color is more than two.

- ① **Excluded Colors**: Choose a color that will be excluded from cell counting (up to 6 colors).
- ② Included Colors: Choose a color that will be included in cell counting (up to 6 colors)

(2) Set Cell Counting Options

- Color Similarity: Color Similarity means the permissive range of color for cell counting

 The range of color similarity can be set from 0 to 100(%). Both using up & down
 button or inputting digits are available.
 - When using Exclude Selected Color, try it first in the range of 10-20%
 - When using Include Selected Color, try it first in the range of 40-60%
- ② Minimum Cell Size: Set min Cell Size which is considered as a cell. The range of cell size can be set from 0 to 100 by using up & down button.
- ③ **Marking Color**: Click **Marking Color** and choose a color that will used to mark counted cells.
- ④ Set Analysis Area: It is useful if you want to count cell on a certain part, not whole image. Details are described in p.14
- (3) **Length Measurement Option**: Set **Length Measurement Option** to measure the length of counted cells. Details are described in p.15
- (4) **Aggregation Count Options**: Set Aggregation Count Options to calculate an estimation of aggregated cells. Details are described in p.17
- (5) **Enlarged View**: It is a screening magnifier. The image could be magnified up to 4 times by using it. Details are described in p.18
- 3) Execute Cell Counting

Click Count. The number of cells will be appeared beside the file name



(1) Example of **Excluded Colors** Option.

- Yellow background color was selected as Excluded Colors and Color Similarity was set at 30%.
- ② As a result, the number of cells are 209 which were marked with red color on the Converted Image.
- 3 Enlarged View on the below left helps to see details of the result.
- (2) Example of **Included Colors** options.

SABIA/Solution for Automatic Blo-Image Analysis	
File(F) Cell Confluency Cell Count Image Analysis SABIA Abo	utta)
Coll Count x	
Count Copy List Analysis Options	Polar False
Select Folder Location	Select Coor
Polities construction and select	Subtact Selected Color C
File name Count Average Hemocytumeter w0103031Wc2 Jpp	Cold Call Defons Cold Call Defons Cold Call Call Call Call Call Call Call Ca
m m r m m r m m r m m r m r m r m r m r m r m r m r m r m r m r m m r m m	
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- Included Colors is useful when cells' colors are more than two. For example, live cells and dead cell in an image can be separately counted.
- ② Check **Included Colors** box, choose color from the image. Then, adjust **Color Similarity** and click **Count**.
- ③ Above example image has both red and blue colors cells. It shows that Included Colors is marked by red, resulting in only red color cell are counted.



3. Adjusting after Completing Cell Count (if necessary)

SABIA/Solution for Automatic Blo-Image Analysis		- 6 - X-
File(F) Cell Confluency Cell Count Image Analysis SABIA Ab	about(A)	
👌 Cell Count x		•
💼 Count 🛛 🖓 Copy List 👘 🍘 Analysis Options		
Select Folder Location	Select Color Save Image	
Folder C: WProgram Files (x86) WMeTooSoft Select	Subtract Selected Color	
File name Count Average Hemocytometer W2(193)1 Weig	Cell Card Options Color Similarly(s) 200 (9) Min Cell Size : (9) Cell Color : • • • • • • • • • • • • • • • • • •	
WLangtocourt-Kf WLangtocourt-Sf WLangtocourt-Sf		
z · · · · · · · · · · · · · · · · · · ·		
Add Add Decrease		2010.04.01

1) Adjusting-needed circumstance.

- (1) When actual cells were not counted.
- (2) When not actual cells were recognized as cells
- (3) When counted cells needs modification of its size
- (4) When the mark of counted cells were not matched with actual location.

2) Adjusting Ways

- (1) Click with the right burton of mouse or double click cell which you want to adjust. Popup menu will be appeared.
- (2) Add: It makes a new cell which was not counted even if it is real cell.
- (3) Increase: it increases the number of the already counted cell.
- (4) **Decrease**: it decreases the number of duplicated cell which is actually a cell. If the number of cell is one, it will be showed Delete, not Decrease.
- (5) **Delete**: it deletes a cell which is not actually a cell.
- (6) **Number**: It means the size of cell. Default value is the same with Minimum Cell Size on the Cell Counting Option. Cell size can be adjusted by put an exact number.
- (7) **Move**: The spot of cell can be moved by clicking, dragging, and dropping.

4. Set Analysis Area

1) Check **Set Analysis Area**, then click and drag a certain area to be analyzed. Choosing multiple areas is available but overlapping areas is not allowed.



2) Set Select Color and Cell Count Options, then execute Count.



- (1) The number of cells in each area will be appeared in the black circle. An average of cells of all areas will be appeared beside the file list.
- (2) After completing cell count, the results like cells' number or size can be modified as above mentioned in **Adjusting Ways**.
- (3) **Dilution Factor**: This value is for the calculation of **Hemocytometer**.

5. Measurement of Length of Cell

Check Cell Length Option, then set Standard Pixel, Standard Length, and Standard Color. Cell length data will be automatically showed on the left side when counting cell.

[ATTENTION] Scale bar must be needed for this option.



- 1) Details of Length Measurement Options
 - (1) **Standard Pixel**: Pixel about the scale bar by using standard bar.
 - (2) Standard Length: The value about the scale bar on the image.
 - (3) Unit : Unit of scale bar (example image's unit is micrometer)
 - (4) The example image indicates 100 micrometer is equal to 82 pixel. Using this pixel, the length of cell will be calculated and its unit will be micrometer.
- 2) Setting Standard Value for Length Measurement
 - (3) Click **Display Scale Setting** to create a standard bar on the image.
 - (4) Drag the standard bar and match exactly the end of standard bar and the end of scale bar on the image. Using **Enlarged View** on the bottom left side helps to adjust precisely.
 - (5) Another way, click the right mouse button on the standard bar, then adjust the location of stand bar by using X and Y coordinates.
 - (6) For adjusting the stand bar size, keep clicking an end of stand bar then drag it to extend or enter an exact digit into **Standard Pixel**.
- 3) Result of Cell Length Measurement

A result of measurement of cell length will be showed on the left side. It can be copied using **Copy List** or dragging and copy & paste directly the result on the left side to excel or notepad (see below).



(1) **Pixel Length**: the number of pixels which are occupied on the screen.

- (2) Actual Length: actual length of cell that was measured by the scale bar.
- (3) **Search Selected Cell's Length**: place mouse cursor on counted cell. Its length will be marked with blue color on the left list.
- (4) Search Cell's Location: double click the length data which you want to find the location of cell. Its location will be marked with white color on the converted image.

6. Calculation of Aggregation

Aggregation can be estimated by using the average size and by comparing to each size.



- 1) Details of Aggregation Count Options
 - (1) Mean Size: the average size of all marked cell (automatically calculate)
 - (2) Mode Size: most frequent size among all among marked cell (automatically calculate)
 - (3) Median Size: middle value from all marked cell (automatically calculate)
 - (4) **User Input:** the value that you want to input.

2) Interpretation of Aggregation Result (list on the right)

- (1) **Pixel Length**: the maximum length of marked cell
- (2) Area: total number of pixel marked cell. It indicates its area
- (3) Estimated Number: the estimated cell number from the counted cell. Estimated number
 = area(total pixel) / area which is measured by Calculation Standard value. It is able to sort list in ascending or descending order.

Pixel Length	Area (Total ~ Pixels)	Estimate Count	•
22	282	3.8	
20	277	3.7	_
26	277	3.7	-
22	259	3.5	
22	252	3.4	
20	206	2.8	_
18	199	2.7	
22	192	2.6	
16	168	2.3	
18	162	2.2	
16	160	2.2	

7. Function of Analysis Options

Cell counting options can be saved by **Analysis Options**. It is convenient and fast. Saved cell counting options will be retrieved and applied next cell counting tasks. Multiple cell counting options can be also saved.

Manage Analysis Options
Option Name Cell-1, Cell-2 condition Lengtgcount-6 condition
Option Name : Lengtgcount-6 condition
Apply Remove Add Edit

- 1) Option Name: easily rememberable name is better
- 2) Apply: apply the selected Analysis Option
- 3) **Delete**: delete the selected Analysis Option
- 4) Add: add a new Analysis Option
- 5) Edit: modify the name of selected Analysis Option

8. Enlarged View (Screen Magnifier)

- 1) An image can be magnified up to 4 times.
- 2) For more big size, dragging Enlarged View window. For original location, drag Enlarged View to its original location or use mouse right button and click dock.





9. Examples of Cell Count from Various Conditions.



[Single Color Background]



[Grid Lines]



[Single Color Background 2]



[Set Analysis Area]

V. Image Analysis - Tissue Ratio

1. Overview

A purpose of Tissue Ratio analysis is to calculate the ratio of tissue characteristics (different colors) on an image. Not only the tissue's different characteristics can be verified, and also the thing how its characteristics are clear can be observed by using Color Intensity. A certain area can be specified as well.



- 1) **Select Folder**: Open the folder which images are located by clicking **Select** in the upper-left side of screen. Select an image to be calculated.
- 2) Set Tissue Ratio Calculation Option
 - Select Color: Select colors into A and B boxes to calculate the ratio of tissue's characteristics. Recommend Color Similarity starts at 50% then adjust it depending on the result.
 - (2) Fill Color: select colors into A and B boxes to mark calculated result (if necessary)
 - (3) **View Intensity:** check **View Intensity** and choose one A or B box. Its value will be showed on the bottom of image and the average intensity will be recorded on the left side(if necessary).
- 3) **Tissue Ratio Calculation:** Click **Calculate Tissue**. The ratio of two different colors will be calculated based on the **Tissue Ratio Calculation Option**.

3. Result of Tissue Ratio Calculation



- 1) Based on **Select Color** and **Color Similarity**, the results of calculation will be showed on the **Converted Image** with Fill Color.
- 2) If color is not included in two colors, it will be excluded from the Tissue Ratio Calculation
- 3) The ratio of tissue characteristics will be appeared on the left side.



4. Calculation for A Specific Area

- 1) Check **Select Area**, then specify a certain area to be calculated. The ratio of tissue characteristics will be calculated within the selected area.
- 2) If check **View Intensity**, the intensity of A or B box will be showed on the bottom of image instead of Fill Color. Results of intensity value and pixel size will be appeared beside of file name.

3) Chart for Color Intensity

Color Intensity uses gray scale (difference of brightness) to calculate the degree of dyeing. Differences between same tissue, but the difference of the amount of dyeing or changes in tissues characteristics by time interval can be analyzed by using color intensity.



- (1) In chart of **View Intensity**, X-axis indicates the value of intensity and Y-axis indicates the area of calculated tissue(pixel).
- (2) Higher intensity value, closer to black. Smaller value, on the contrary, closer to white.
- (3) Results of Color Intensity can be copied to excel.

V. Image Analysis - Area

1. Overview

Image Analysis for area is to measure area. A measurement of area is calculated by color differences. A certain part is specified to measure area and Color Intensity is also available in this part. The basic concept is the same with other parts. An important thing is that scale bar is needed to measure area likewise Cell Counting.



2. How It Works

1) **Select Folder**: Open the folder which images are located by clicking **Select** in the upper-left side of screen. Select an image to be calculated.

2) Set Color Options

- (1) Select Color: select a color from the image for measuring Area and Color Intensity.
- (2) Fill Color: select color to represent area after measuring (if necessary).
- (3) **View Intensity**: check **View Intensity**, color intensity and pixel will be showed instead of Fill Color (if necessary).
- (4) **Color Similarity**: Recommendation is starting at 50%. Then adjust the range of Color Similarity base on the results.

3. Color Distribution and Color Intensity



After you execute **Calculate Area**, distribution of selected color will be changed with Fill Color and the sum of pixel will be appeared beside of file name. This is without the calculation of area.

4. Area Calculation



Check **Area Calculation Option** and **Display Scale Setting**. Set the standard bar options as described in chapter IV-5, Set Standard Bar.

Click Calculate Area resulting in measured area will be appeared beside of file name.

5. Calculate Area in A Specific Area.

Specifying a certain area is available to calculate area and color intensity. It has the same

procedure to total area calculation.



- 1) Check Select Area, draw a part where you want to calculate area by using mouse.
- If Select Color is unchecked, total area of selected part will be calculated. If Select Color is checked, selected color of selected part will be calculated. Either Fill Color or View Intensity is needed to check.
- 3) Click Calculate Area
- 4) If Select Color is unchecked (total area 883 with checking View Intensity)



5) If Select Color is checked (total area 131 with checking Select Color).



6. Chart for Color Intensity: It is the same as described in chapter V. Calculation Tissue Ratio 4-3

7. Example of Color Intensity

Area calculation can compare between the same sample with different time point or the same sample with different dyeing ways. The numerical value acquires objectivity and university in comparison to manual effort.

1) Image A: relatively light condition (Average Color Intensity 125)





2) Image B: relatively dark condition (Average Color Intensity 167)





V. Image Analysis - Length

1. Overview

An actual length can be measured when an image has scale bar and the measurement ways of length are various. Using straight line or adjustable curved line enable to measure the length. The basic concept is similar to other functions.



2. How It Works

- 1) **Select Folder**: Open the folder which images are located by clicking **Select** in the upper-left side of screen. Select an image to be calculated.
- Set Standard Bar: Click Display Standard Bar and set Length Measurement Option. Handling options is the same to the calculation of standard bar (See IV-5 Cell Length Calculation).
- 3) **Other Options:** If necessary, **Line Color**, **Label Fill Color**, and **Font Color** can be changed by clicking the mouse right button.
- 4) Unit of Length

Unit is equal to Scale bar's unit.

3. Methods of Length Measurement

The left burton of mouse means the start of line setting and the right burton of mouse means the end of line setting.



- 1) Connect Two Points
 - (1) Click two points which you want to measure length, then click the right mouse burton to end setting.
 - (2) Straight line will be appeared. Horizontal, vertical, and diagonal line can be drawn.

2) Dragging Mouse

- (1) Keep clicking the left button of mouse and drawing the line. Then click the right button of mouse to finish setting.
- (2) It is useful when you want to draw the adjustable curved line.
- 3) Using Ctrl key
 - Keep pressing Ctrl key and clicking the left mouse button, then move to where you want. Release the left mouse button and click the right mouse button to finish setting.
 - (2) It is useful when you want to draw the straight line.
- 4) Finish and Cancel Setting
 - (1) To finish setting, click the right button of mouse. A label of line will be appeared at the point of line and the result will be added in the left side of screen.
 - (2) To cancel line drawing, Press Ctrl + Z keys. Drawn lines and results will be disappeared.

5) Click **Copy Length List** to copy results of length measurement to clipboard.

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