

MANUAL

EXI-600

RESEARCH GRADE INVERTED BIOLOGICAL MICROSCOPE SERIES

EXI-600 Research Grade Inverted Biological Microscope User Manual

This manual is written for EXI-600 research grade inverted biological microscope. To ensure the safety, obtain optimum performance and to familiarize yourself fully with the microscope, it is strongly recommended that you read this manual carefully before operating the microscope and put this manual in a place easy to get for reference.

The warning and notice signal used in this manual

We provide you the most safety and reliable instruments, but improper operate and ignorance of precautions may lead to a personal injury or a loss of property. To insure the right operating method, we hope you read this manual carefully before you use the product. Besides, it's highly recommended you to put this manual in a place easy to get for easy reference.

In this manual the safety notices will be emphasized by the below symbols, please obey the statement of these symbols.

Symbol	Meaning
MARNING	Ignore this symbol may lead to a serious personal injury even death.
NOTICE	Ignore this symbol may lead to a personal injury or a property loss.

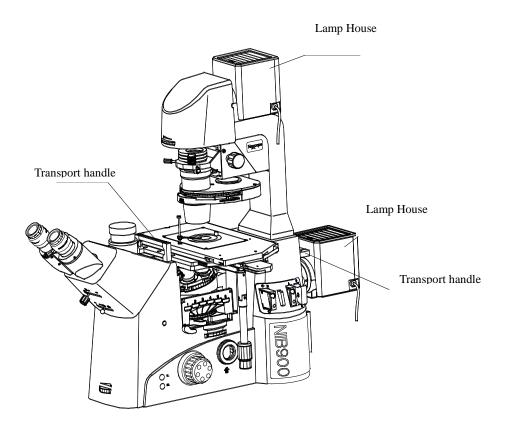
The meaning of symbols on the product

Symbol	Meaning	
Ð	Conductor protect terminal	
<u><u><u>s</u></u></u>	Caution of hot! Remind you of this situation: The lamp house and spaces nearby will get hot when using. Do not touch the lamp house in 30 minutes when it's closed and when it's on. The lamp house may still extremely hot after it's closed, please ensure enough cooling time before replacing the bulb	
	This symbol appears in the electrical name board to remind you to make sure the voltage of your area is the same to the input voltage	
_	Turn on the power, rotate the brightness knob to adjust the brightness of the field of view	
\bigcirc	Turn off the power	
	UV radiation	
	Turn off the power before open the device	

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Safety Precautions



- 1. Avoid placing the microscope in a place exposed to direct sunlight, with high temperature, with high humidity or easily vibrated, make sure the platform is flat, horizontal and stable enough. (weight of the mainframe is 29.5kg)
- 2. When moving the microscope, hold the transport handle tightly and keep a distance between the microscope and the platform before actually moving it.(shown in picture above)
- 3. If bacterial solution or water is dripped over the stage, objective or observe tube, turn off the power supply immediately and wipe out the solution, otherwise the instrument maybe damaged.
- 4.



When operating, the lamp house can be considerably hot, make sure there is enough space around the lamp house for heat dissipation to avoid the heat accumulates and damage the instrument.



Before turn on the power of lamp house, make sure the power supply is connected properly. Before replacing the bulb or fuse, turn off the power and wait until the lamp house cooled completely.

★the indicate bulb: 12V 100WHAL high brightness halogen bulb (OSRAM)

★ the fuse should load the correct melting current, do not use a temporary fuse in case of the broken circuit.

6. Connect the power wires correctly, make sure the instrument is grounded in case of lightning stroke.

7. Use the dedicated electric wires.



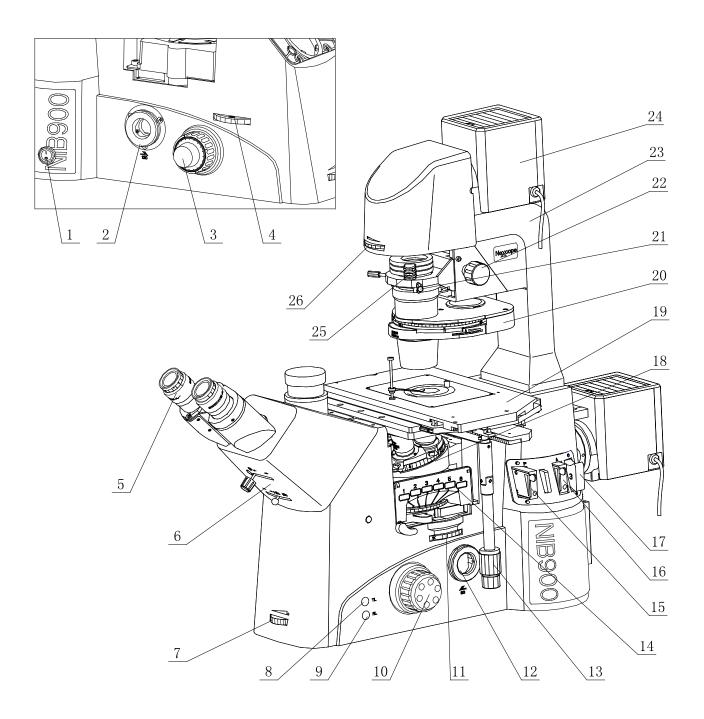
The halogen lamp can radiate UV light thus may cause a burn of eyes and skins. Do not see through the light directly and take protective measures when using the microscope as much as possible.

- 9. The product should store in a place with shelter and no acid gas, alkali, organic solvent or any harmful substances around.
- ★ a 3-pin plug is reserved for the machine for safety concern, the machine is grounded through the 3-pin plug. Do not use any other adapter plug or e the safety performance may decrease.

 \star Do not put the instrument in a place where is hard to cut off the melting current.

 \bigstar The protective offered by the device may be damaged if the device is not used in the way indicated by our company.

I Structure and Names



Pic. 1

Component names:

- 1 Power switch
- 2 Left camera port
- 3 Left coarse/fine focusing handwheel
- 4 Side ports converter(light path convert—left and right side ports/observe)
- 5 Eyepiece
- 6 Trinocular tube
- 7 Brightness adjustment knob
- 8 Transmitted illumination shutter
- 9 Reflected illumination shutter
- 10 Right coarse/fine focusing handwheel
- 11 Middle magnification converter
- 12 Right camera port
- 13 X/Y axis handwheel of sample stage
- 14 Multifunction module turntable
- 15 Field diaphragm of reflected light
- 16 Aperture diaphragm of reflected light
- 17 3-hole filter spile
- 18 Obejective converter
- 19 Sample stage
- 20 Turntable condenser
- 21 Condenser centring screw
- 22 Lifting handwheel of condenser
- 23 Transmitted illumination support
- 24 12V 100W illumination lamp house
- 25 Filter support
- 26 Field diaphragm of transmitted light

II Applications

The EXI-600 inverted biological microscope is mainly used to study the cells, cultured organization and sediment in culture bottles and culture dishes using transmitted and reflected light.

This device can be used in bright field, dark field, phase contrast, differential interference contrast, polarized light and fluorescence observation.

Main application of this device: study of human blood and tissue samples, observation of the connection, activity and growth between live cells, drug reaction, minimally invasive, external fertilization, toxicity test, digital record, automatic continuous time-point observation, single molecule detection, etc.

III Assembly

Preparation before assembly and operation of the microscope

Tear down the packaging of the mainframe and the accessories.

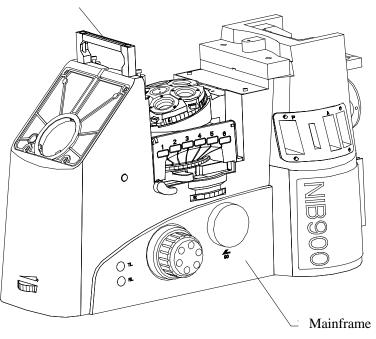
The packaging include of mainframe, eyepieces, objectives, condenser support, lamp house and other accessories such as filters, DIC blocks, dust-proof shield, tools and user manual. The optional accessories will be packaged individually.

- 1. Tear down all the packaging, check the goods and confirm the accordance with the products you purchased.
- 2. Dismantle the transport handles shown in pic.2.

Put the mainframe in a vibrate-proofed platform and then dismantle the handle with 4mm hexagon screwdriver.

 \star Please take proper preservation of the handles.

Transport handle



Pic.2

Assembly

1. Trinocular tube

①Use a 2mm hexagon screwdriver to loosen

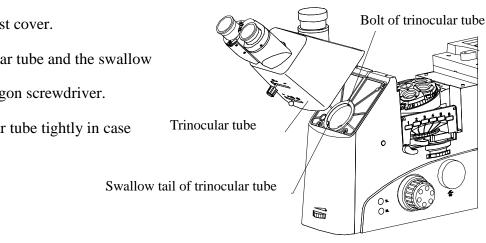
the bolt, take out the dust cover.

⁽²⁾Calibrate the trinocular tube and the swallow

tail, fix it by 2mm hexagon screwdriver.

 \star Do hold the trinocular tube tightly in case

of fallen down.



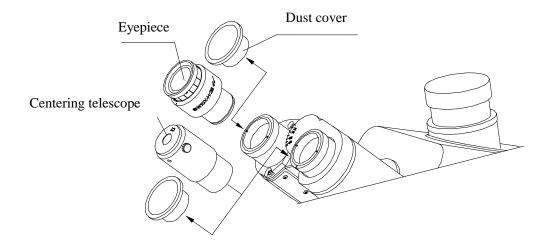
Pic.3

2. Eyepiece and centering telescope

①Take off the dust cover, insert the 2 eyepieces into the tube(the eyepiece should fully contact with the tube).

 \star ②When observing with phase contrast, replace one eyepiece with the centering telescope, make the phase contrast ring focusing accurately.

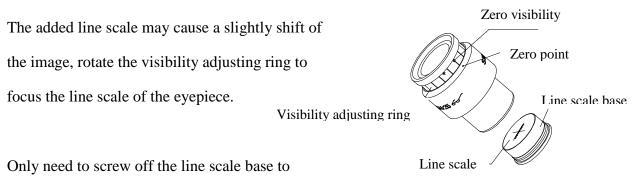
\bigstar when using centering telescope



Pic.4

3. Graduated ocular

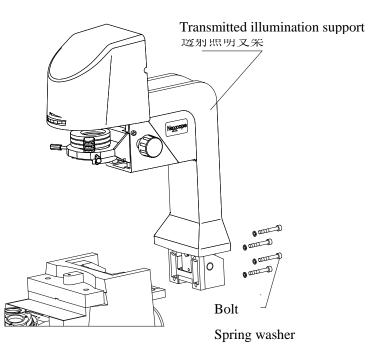
Change the line scale.





4. Transmitted illumination supportPut the support at the back ofmainframe and tighten it withM5X35 bolt and spring washer.

The transmitted support does not need any adjustment.



5. Objective

Dust cover

Screw off the dust cover on the converter, screw the

objective into the trapped hole

The objectives should match the lettering on the

converter by their magnification.

Screw the dust cover into other vacancies.

6. Sample platform

Assemble the platform 135X85R/L

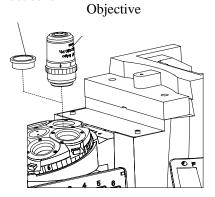
with the mainframe by three bolt M4X10

 \bigstar The platform can be assembled

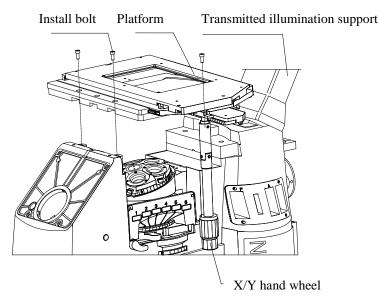
at either right or left. (meanwhile

the X/Y hand wheel can be at either right or left)

 \bigstar The transmitted support can be tilt a bit to the back to assemble the platform.

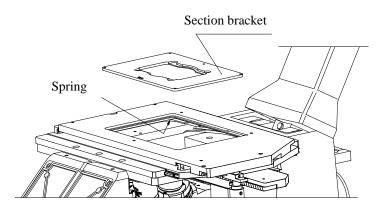


Pic. 7



Pic. 8

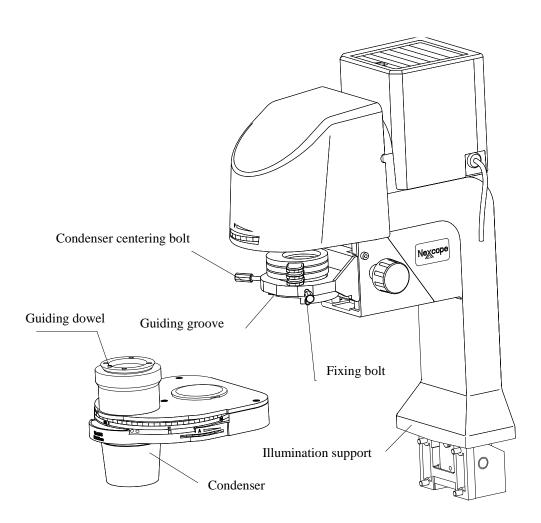
Put the section bracket close to the spring on the edge of the platform and press it to the platform levelly.



Pic. 9

7. Turntable condenser

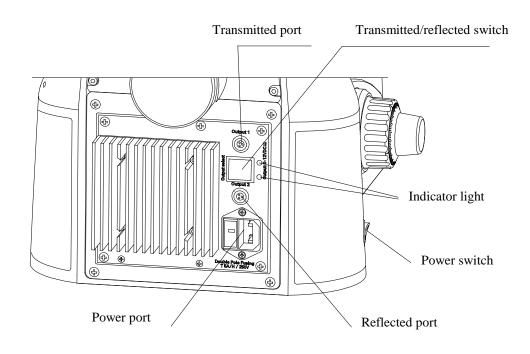
Align the guiding dowel of the condenser to the guiding groove of the support and fix the condenser with the bolts after inserting it into the transmitted illumination support.

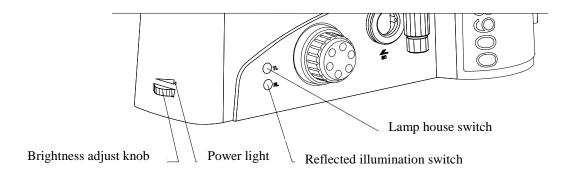


Pic. 10

Please off the power before installing any parts and accessories.

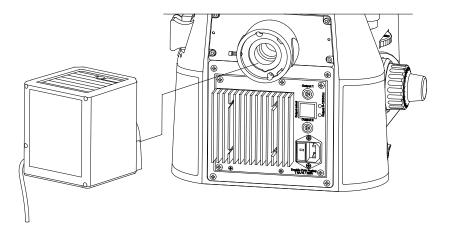
- \star The TL lamp house switch can control the transmitted lamp house without turning off the power.
- \star The RL lamp house switch can control the reflected lamp house without turning off the power.





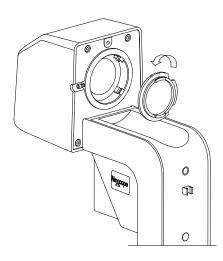
Insert the swallow tail of the lamp house to the socket, make the lamp house parallel to the stable table and then fixed it with bolts Connect power wires by the port position shown in pic.11.

Check the connection and open the power supply, rotate the potentiometer knob and see if the brightness changes.



Assemble the diffuser

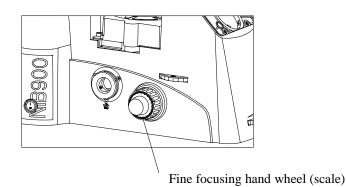
Locate the two round grooves of the diffuser to the two bulged on the swallow tail socket and screw the diffuser into the socket groove. Screw it reversely when taking the diffuser out.



Pic.13

IV Operating and adjusting

 Coarse and fine focusing hand wheel move range: 10mm coarse focusing hand wheel: 2mm/cycle fine focusing hand wheel: 0.2mm/cycle





★Focusing hand wheel is set at both side Of the mainframe

The left hand wheel has scales

The right hand wheel has no scales

2. Side port converter/light path converter—observe from left, right port and ocular

The side port converter has three instruction of different splitting ratio.

100%vis: 0% camera 20%vis: 80% camera left

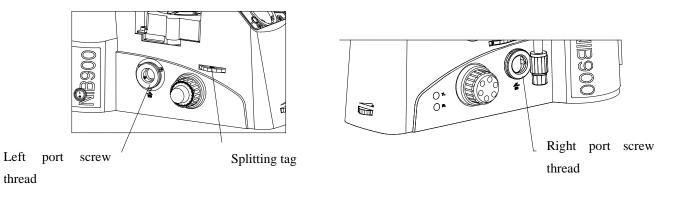
0% vis: 100% camera right



100

 \star : vis---observe by ocular

camera---photonic





3. Turntable condenser

(transmitted light)

NA:0.55

WD: 26mm

Condenser turntable/6 positions

Bright field: H

Phase contrast: PH1, PH2, PH3

Pic. 16

Adjust the aperture diaphragm: rotate the diaphragm plate

6 position adjust: rotate the condenser tunable, if you turned the bright field/H into the light path, make the mark "H" face you.

Centering bolt

Phase contrast ring centering bolt: use 1.5mm hexagon screwdriver to adjust the phase contrast ring to match the objective.

Replacing DIC prism

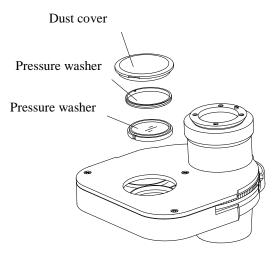
Take out the condenser and put it on the plate form before replacing the DIC prism

Take out the dust cover, screw out the pressure washer with matched tools, and turn over the condenser to let the DIC prism slip off.

Put the new DIC prism to the groove of condenser, screw the pressure washer in and cover the dust cover, shown in pic.17

★ attention the direction of the DIC prism and clip it into condenser carefully to protect the prism surfaces.

 \star make sure the scale ring in the tunable is right.



Aperture diaphragm plate

Condenser turntable



4. Transmitted light field diaphragm

When using transmitted illumination, screw the tunable to adjust the field diaphragm.

Screw the condenser lift hand wheel to lift the condenser and set the observe height.

The condenser is fixed by fixing bolts.

Screw the 2 condenser centering bolts to move the condenser to the center of the light path.

The filter bracket can hold different filters.

Pic. 18

Field diaphragm tunable

Condenser

centering bolt

Filter bracket

Fixing bolt

Nexcope

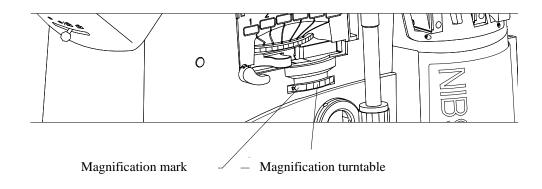
Condenser lifting

hand wheel

5. Middle magnification converter

The middle magnification mark: 1X/1.5X

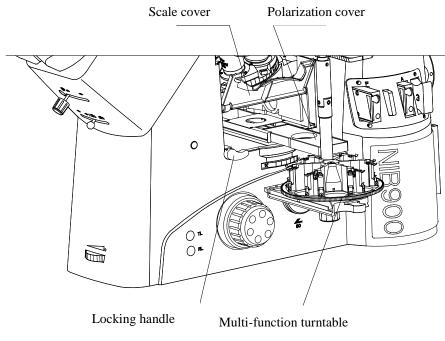
You can directly convert the middle magnification the change the observe magnification.



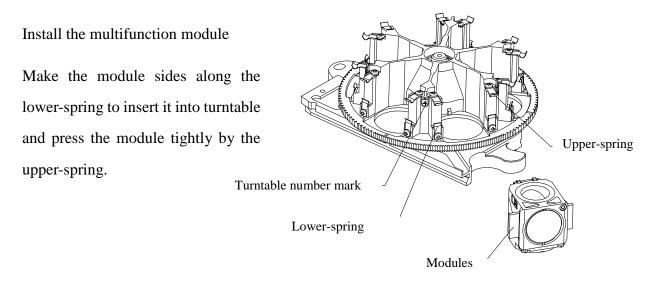
Pic. 19

6. Multi-function modules

6 modules optional .Turn over the scale cover plate, turn down the locking handle, you can insert and extract the turntable.

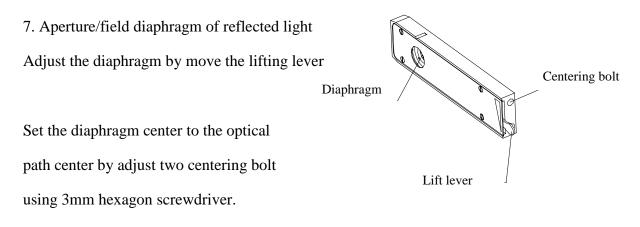


Pic. 20



★ Notice that the module mark should match the number mark on the turntable.

Pic. 21



Pic.22

8. Trinocular tube

Trinocular tube:

45°/25 trinocular tube equipped with vis/doc slip prism, Bertrand lens and manual vis switch.

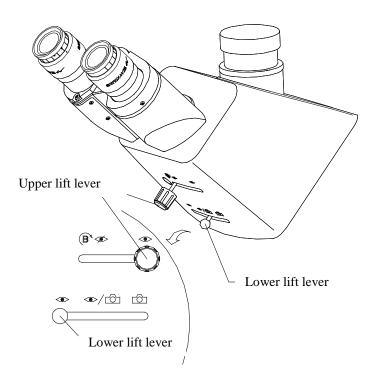
0% vis: 100% camera 50% vis: 50% camera 100% vis: 0% camera

Upper lift level-a lift block to focus the Bertrand lens

100% vis 0 Bertrand lens



∕ ┌∕⊜`



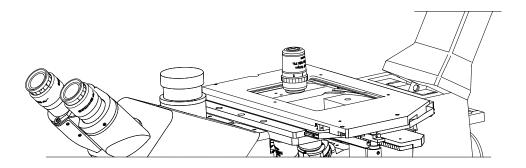
Pic. 23

Cautions in observation

1. Objective

- ① When rotating the converter to change objectives, a sound of click can be heard if the objective is right at the center of the light path.
- ② when operating, firstly search and focus the sample by a low magnification objective(4X or 10X) and then change to a high magnification objective to observe according to your needs.
- ③ The objectives can also be changed by install through the aperture in the platform.

Notice that the magnification should match the scale on the converter, in normal cases when you rotate the converter clockwise, the magnification will get higher.



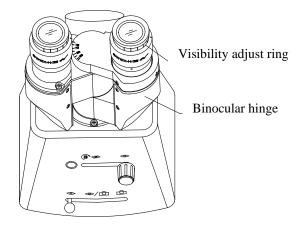


2. Trinocular tube

①Visibility adjustment

Observe one eyepiece by one eye and rotate the coarse focusing hand wheel to focus on sample, then observe another eyepiece by another eye. If the image is not clear, use the visibility adjust ring to it clear by both eyes.

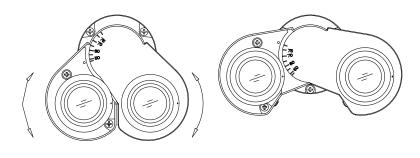
★ There are ± 5 diopters on the visibility adjust ring of the eyepiece, the value which align at the dots on eyepiece holder is the visibility of eyes.



Pic. 25

⁽²⁾Interpupillary distance adjustment

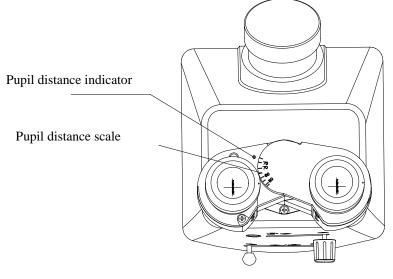
Rotate the binocular hinge to adjust the pupil distance and unity the field of two eyes, also adjust the height of exit pupil.



Pic. 26

Range of pupil distance: 55~75mm

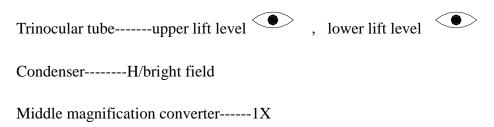
The scale on the pupil distance board corresponded with the dot " \cdot " is the value of pupil distance.



Pic.27

Transmitted illumination

1. Bright field observation



Multi-function modules ----transmitted module (number 3)

2. Phase contrast observation

Trinocular tube----- upper lift level , lower lift level

Condenser-----PH1/PH2/PH3

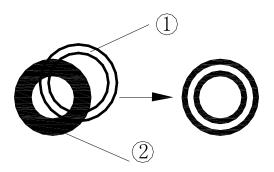
Middle magnification converter-----1X

Multi-function modules ----transmitted module (number 3)

- ① The magnification of phase contrast objective should match the condenser phase contrast mark. PH1/10X-20、PH2/40X、PH3/60X
- ② Centering phase contrast ring shown in pic.28.

If the phase contrast ring is not centered, use 1.5mm hexagon screwdriver to adjust by following steps:

- a. Put a sample on the platform and focus it.
- b. Take out the eyepiece and insert the centering telescope into the eyepiece tube.
- c. Make sure the matched phase contrast ring 2(in the objective) and phase contrast ring 1(in the condenser) have shifted in the light path.
- d. Observe through centering telescope, adjusting the phase contrast ring by 1.5mm hexagon screwdriver until the 2 centers overlaps.
- f. The above methods apply to objectives of different magnification.
- \star If the phase contrast ring is not centered properly, the best effect can't be get.
- ★ The phase contrast ring may shifted after remove the sample or replace it with a thicker sample, if this occurs, repeat the above steps until the 2 centers overlap.
- ★ If the glass slide or vessel is not flat, for a bigger contrast, repeat the above steps to adjust the 2 centers overlap.



Pic.28

3. Differential Interference Contrast (DIC) observation

Trinocular tube----- upper lift level

level

Condenser-----DIC、DIC II

Middle magnification converter-----1X

Multi-function modules ----transmitted module (number 3)

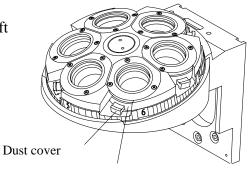
Side port converter----- (**O**)

The condenser DIC mark should match the DIC prism

in the socket of objective converter.

The polarizer should be put into the light path. First install the polarizer into the top most filter holder above the condenser - the pin on the bottom of the polarizer should fit into the large slot of the filter holder.

The analyzer slider should be put into the light path. Insert the slider



lower lift

DIC prism socket

Pic. 29







Reflected illumination

1. Bright field observation

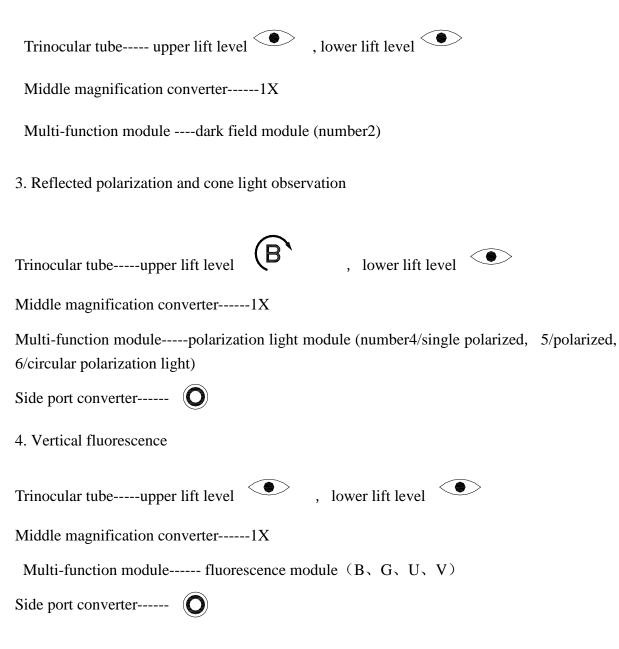
Trinocular tube----- upper lift level , lower lift level



Middle magnification converter-----1X

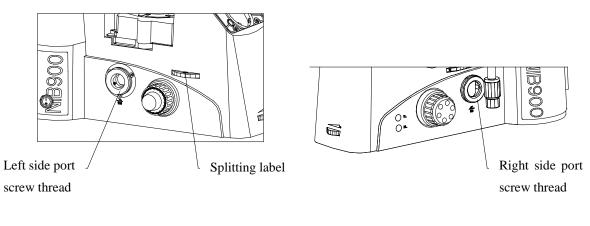
Multi-function module ----bright field module (number1)

2. Dark field observation



VI Microscopy imaging

1. Side port imaging

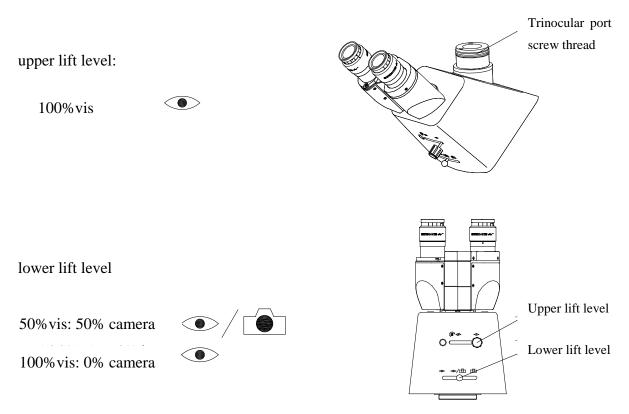




20% vis: 80% camera left 0% vis: 100% camera right



2. Trinocular tube imaging



VII Technical specification

Optical system	Infinite system		
Observe tube	Hinge binocular tube, 45° tilt		
Eyepiece	10X wide field eyepiece with line field of Φ 25mm		
Objective converter	Six-hole converter		
Objective	Infinite plan phase contrast: 10X, 20X, 40X, 60X		
Focusing	Coarse and fine coaxial focus		
	Coarse focusing 2mm/cycle		
	Fine focusing 0.2mm/cycle		
	Stroke (from the plat focal point):up 7mm, down 2mm		
Platform	Range of move: 135(length)X 80(wide)mm		
Illumination	12V 100W halogen tungsten lamp with center preset and brightness adjustable continently		
Condenser	Working distance LD26mm, NA0.55		
Operating environment	 Used indoor Max altitude: 2000m Environment temperature: 5°C~40°C(41°F~109°F) Max relative humidity: 80% at temperature of 31°C(88°F) Decreased linear with temperature 70% at temperature of 34°C(93°F) 60% at temperature of 37°C(99°F) 50% at temperature of 40°C(104°F) Pollution level: level 2 Power supply: 100-240V AC ±10%, 50/60HZ Consumed power: 100W Fuse: T5A/250V Φ5X20mm Air pressure: 80kPa~106kPa 		

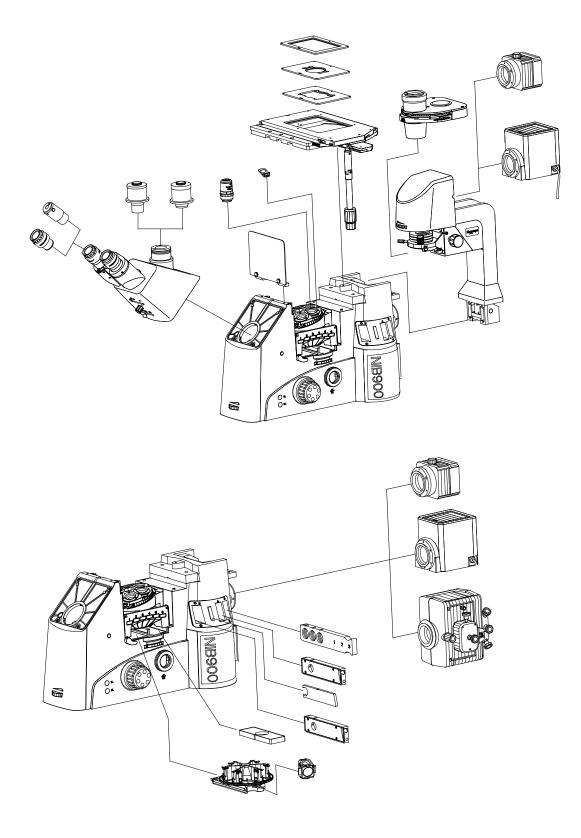
1. Main technical specification

2. Objective parameters

Class	Magnification	Number	Working	Conjugate	Parfocal	Glass
		aperture	distance	distance	distance	Slide
		(N.A)	(mm)	(mm)	(mm)	thickness
						(mm)
Infinity plan	10X	0.30	7.4			1.2
apochromatic	20X	0.45	7.5-8.8	xo	60	0-2
objective	40X	0.60	3-4.4			0-2
	60X	0.70	1.8-2.6			0.1-1.3

VIII Configuration diagram of EXI-600 inverted biological

microscope



Trouble shooting

In particular cases, the performance of this device may be affected reversible by some nondeficit factors. If trouble occurs, take the proper solution listed by the following chart. If the trouble is still not solved, please contact the sales department of our company.

Problem	Cause	Solution		
1.Optical part				
a. Although the illumination is on, the field of view is still dark.	Lamp holder pin has not connected to the illumination device	Connect them correctly		
	The bulb burns	Exchange a new bulb		
	Brightness adjusted too low	Adjust the brightness properly		
	Too many filters used	Minimize the filters according to actual needs		
	The bulb is not indicate	Use 12V 100W halogen bulb		
b. The edge of the field of view has shadow or	The objective converter is not in the correct position	Make sure the converter is in the correct position		
not evenly illuminated	Filter stops in middle way	Push it completely		
	Phase contrast plate is not in the correct position	Push the phase contrast plate to the located position		
c. See dusts or pollution in the field of view	There are dust or pollution on the sample	Exchange a clean sample		
	There are dust or pollution on the eyepiece	Wipe the eyepiece		
d. There is ghost image	Aperture diaphragm is too small	Increase the aperture diaphragm		
e. Resolution troublesThe image is not	The objective converter is not in the correct position	Make sure the converter is in the correct position		
obvious;The contrast is not	Aperture diaphragm is too small or too big in bright field observe	Adjust the aperture properly		
good; ● The detail is not	Lenses (condenser, objective, eyepiece or culture dish) is soiled	Make them fully cleaned		
clear;No phase contrast	The thickness of culture dish bottom is over 1.2mm in phase contrast observation	Use a culture dish with the thickness of bottom less than 1.2mm		

effect gained	The bright field objective is used	Use the phase contrast objective
	The condenser ring is not match the objective ring	Adjust the condenser ring to match the objective ring
	The light ring is not centered with phase contrast	Adjust the centering bolts
	The objective is not compatible with the phase contrast observe	Use a objective compatible with the phase contrast observe
	When observing the edge of culture dish, the phase contrast ring deviate the light ring	Move the culture dish to get the phase contrast effect
f. The image blurs at one side	The objective converter is not in the correct position	Make sure the converter is in the correct position
	The sample is not correctly positioned on the platform	Positioned the sample correctly
	The optical performance (such as the profile) of the culture dish bottom is poor	Use a well profiled culture dish

2. Electrical part

a. Bulb didn't shine	No power supply	Check and connect the power wire
	Bulb isn't installed correctly	Install the bulb correctly
	Bulb burns out	Exchange the bulb
b. Bulb burns out frequently	Non-indicate bulbs used	Use indicate bulbs
c. Brightness is not	Non-indicate bulbs used	Use indicate bulbs
enough	The brightness adjust knob is not used correctly	Use it correctly
d. The light twinkles	Bulb is nearly burned out	Exchange the bulb
	Power wire is badly connected	Connect the power wires correctly

3. Observe tube

The field of one eye is not	Pupil distance is wrong	Adjust the pupil distance
coincide with the field of another eye	Diopter is not correctly adjusted	Adjust the diopter
	Have not adapt the microscope	When observing from the eyepiece, observe the
	observation yet	whole field of view before concentrate on the
		sample, it is benefit to look up or faraway before
		observe.

4. Microscopy imaging

a. Image defocused	Focusing incorrectly	Adjust the focal length to make the double cross line and sample visible clearly.
b. Edge of image blurred	The using achromatic objective can't focus the edge	Blur is inevitable
c. The image of window or light appeared	The outside beam entered the eyepiece or viewfinder is reflected	Cover the objective and viewfinder of the illumination system

X Maintenance

1. Use gauze to swipe glass components slightly. If you want to remove the fingerprints and oil stains, use thimbleful alcohol and diethyl ether mixed liquor (3:7) or dimethylbenzene to wipe.

★ Diethyl ether and alcohol are both extremely flammable, DO NOT get them near to open flames or any potential electro sparks such as the switch of electrical device. Use these chemical products in well ventilated rooms as much as possible.

2. Do not use organic solvent to wipe any non-optical components. If you want to clean these components, use a hairless and soft cloth to dip a little of neutral cleaner to wipe.

3. If the microscope is wetted by liquid when operating, cut off the power supply immediately and wipe the liquid.

4. Do not separate any parts of the microscope otherwise the function and performance of the microscope may be affected.

5. The hole on the objective converter must be covered with dust cover if it is not occupied with objective in case of dusts or culture solution enter the inside of the device.

6. The microscope should cover with dustproof shield when it is unused. Wait until the lamp house fully cooled before covering the dustproof shield.

7. The check and replacement of components of this product should be executed by our company or the indicated agency which also provide the accessories.