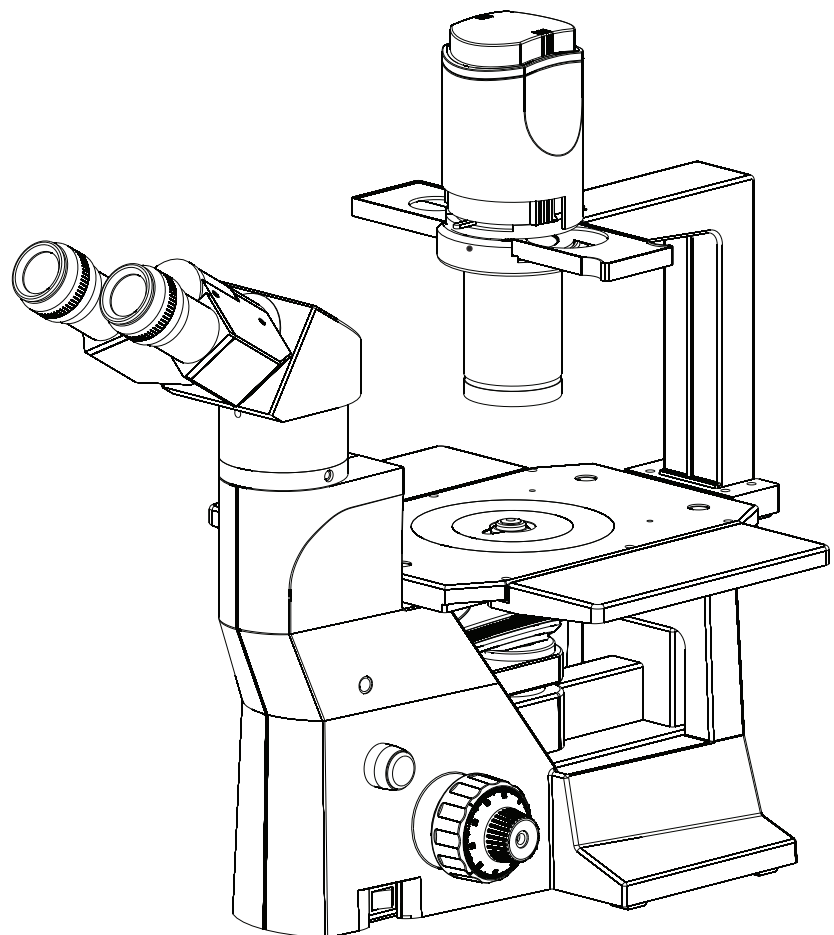


TCM 400 Inverted Tissue Culture Microscopy

User Manual



To ensure proper use of this instrument as well as to avoid injury while operating Instrument, understanding this manual completely before use is highly recommended.

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The TCM 400 inverted microscope sets a new standard for inverted tissue culture microscopes.

It combines high performance and distinctive styling with the high quality craftsmanship for which Labomed is known. The excellent image and comfort of use are the basics of Labomed TCM 400.

The ergonomics of the design, an aesthetically configured TCM400 is the instant choice for Culture studies in the era of bio sciences. Offered with Infinity Corrected True color optical system, choices of observation tubes, soft feel turret movement, stage extenders, micro manipulators giving a paramount performance.

The user should be aware that the protection provided by the equipment may be impaired if used with accessories not provided or recommended by the manufacturer or used in a manner not specified by the manufacturer.

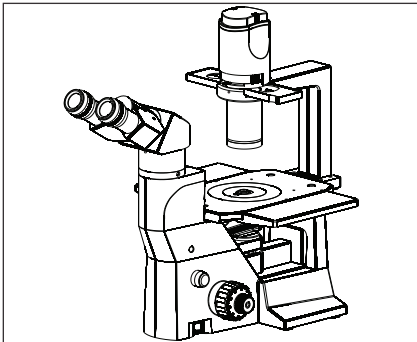


Fig. 1

1. After the microscope has been used for observation of a specimen containing bacteria, clean all parts coming in contact with the specimen to prevent infection.
 - Be sure to remove the specimen before moving this product.
 - In case the specimen is damaged by erroneous operation, it is important to clean all surfaces that may have come in contact with the specimen.
2. To avoid potential electrical hazards when replacing either halogen bulb, turn the microscope main switch to the OFF position and disconnect power cord from wall outlet in advance. Whenever you replace your microscope bulb, allow lamp socket and bulb to cool before touching (Fig.1)

Applicable bulb replacement: 6V30W Halogen bulb P/N EL-455



3. Install microscope on a sturdy, level table or bench and avoid any restriction of air vents in the base of the unit.

Do not place microscope on a flexible surface, as this could result in blocking the air vents and cause overheating.
4. Always use the power cord provided by LABOMED. If the proper power cord is not used, product safety performance cannot be warranted.
5. When installing the microscope, route the power cord away from the microscope frame. Should the power cord come in contact with the microscopes base, the power cord could melt due to overexposure to heat.
6. Always ensure that the grounding terminal of the microscope and that of the wall outlet are properly connected. If the unit is not grounded, LABOMED can not warrant electrical safety.
7. Never allow metallic objects to penetrate the air vents of the microscope frame as this could result in user injury and damage to the microscope.
8. After operation of microscope, be sure to disconnect power cord from connector socket of the microscope or from the wall power outlet.

SAFETY SYMBOLS

The following code categories describe the degree of danger likely to be incurred in the event of user error made in ignorance of these symbols.

Symbol	Explanation
	Hot parts symbol. This symbol is placed on bulb housing and on other parts to indicate that they may be hot. Never touch those parts just after the equipment has been used. Allow sufficient time to cool before touching them. In the event of user error it is possible to injury.
	In the event of user error, death or serious injury is possible.
	In the event of user error, product may be damaged.
	The following describes icon directives This icon denotes "Refer to User Manual" and "Caution".
	This icon represents the productive earth.
	This icon represents alternating current.


	This icon represents the power switch is turned OFF.
	This icon represents the power switch is turned ON.
	Labeling Information
SN	Number following this symbol indicates the serial number of the equipment.
REF	Character following the symbol indicate the catalog number of the equipment.

Caution

If the microscope is used in a manner not specified by this manual, the safety of the user may not be warranted. In addition, the equipment may also suffer damage. Always use the equipment as outlined in this instruction manual.

1 GETTING STARTED

1. A microscope is a precision instrument with delicate glass components. Please handle with care.
2. Do not use the microscope where it is subjected to direct sunlight, high temperature, humidity, dust and Vibrations. (For the operating conditions, see chapter 14. "SPECIFICATIONS")
3. The microscope is ventilated by natural convention. Be sure to leave enough space (10cm or more) around body when installing the unit.

 To prevent damage, do not hold the microscope by the stage or observation tube. Be sure to remove the specimen from the stage clip while transporting unit to avoid damage of the specimen slide.

2 MAINTENANCE AND STORAGE

1. Clean all glass components by wiping gently with cleaning cloth provided. To remove fingerprints or oil smudges, wipe with cleaning cloth slightly moistened with a mixture of petroleum (85%) and isopropanol (15%).

Since solvents such as petroleum and isopropanol are highly flammable, they must be handled carefully. Be sure to keep these chemicals away from open flames or potential Sources of electrical sparks - For Example, electrical equipment that is being switched "ON" or "OFF". Also remember to always use these chemicals only in a well-ventilated room.

2. Do not attempt to use organic solvents to clean the microscope components other than the glass components. To clean non-glass components, use a lint-free, soft cloth slightly moistened with a diluted neutral Detergent.
3. Do not disassemble any part of the microscope as this could result in malfunction or mitigated performance.
4. When not using the microscope, ensure that the frame is fully cooled before storing the unit in a dry locker or covering with a dust cover (provided).
5. To clean the condenser, remove the condenser by rotating it anti clockwise. Wipe the front lens of the condenser with optical cleaning solution (mixture suggested above) and lens tissue.
6. Be sure to observe your local rules/regulation for product disposal.

3

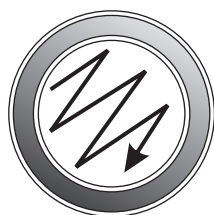
3 CARE & MAINTENANCE

Your microscope had been engineered for a long and safe operational life with the least amount of maintenance required. In general, routine maintenance is limited to keeping the microscope working parts lubricated and optics clean. Always cover the microscope with the provided dust cover when not in use.

Optical Cleaning

1. The objective have been adjusted for a tight fit to prevent any damage during transportation. To remove an objective, rotate it counterclockwise while gripping it with a rubber sheet, etc. to avoid any slip page.
2. To clean the lens surfaces, remove dust using a soft brush or gauze (compressed air dust cans are ideal). For removing finger marks or grease, soft cotton cloth or lens tissue lightly moistened with cleaning solution (85% petroleum ether and 15% isopeopanol) should be used. For cleaning the optics, use Meathanol. Observe sufficient caution in handling Methanol. Place the Objective and/ or eyepieces on a dust free surfaces (e.g. aluminum foil). All other optical components to be cleaned should be as accessible as possible.
3. Blow all loose dust particles away with a dust blower.
4. Remove all water-soluble dirt with distilled water. If this is unsuccessful repeat using a solution of diluted hand soap liquid. Remove any remaining residue with a dry cotton swab.
5. To remove oil, use a solution of diluted hand-soap liquid initially. If this does not produce a satisfactory result, repeat the cleaning using a solvent (Optical Cleaning Solution 85% petroleum ether and 15% isopropanol).
6. Grease must always be removed using a solvent.
7. Cleaning is achieved by using a spiral motion from the center to the rim. Never wipe using zig-zag movements as this will only spread the dirt. With larger optical surfaces (e.g. tube lenses) the spiral motion starts initially at the rim before to the middle and is only then followed by a center to rim cleaning motion. Normally several spiral wipes are recommended.

We recommend pure, volatile petroleum ether or Optical Cleaning Solution as explained in point 3 above.



Zig-zag motion (X)

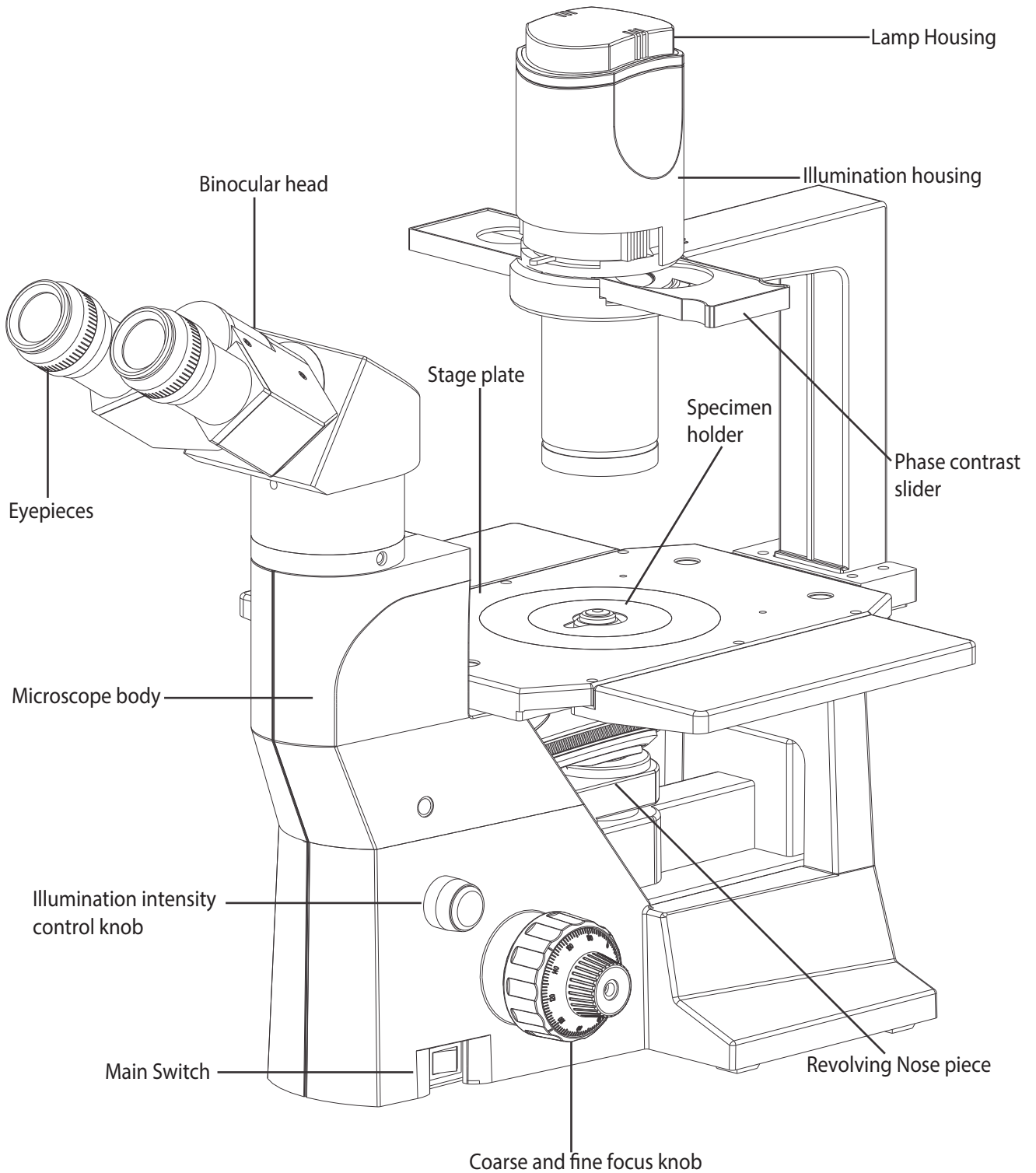


Spiral motion (✓)

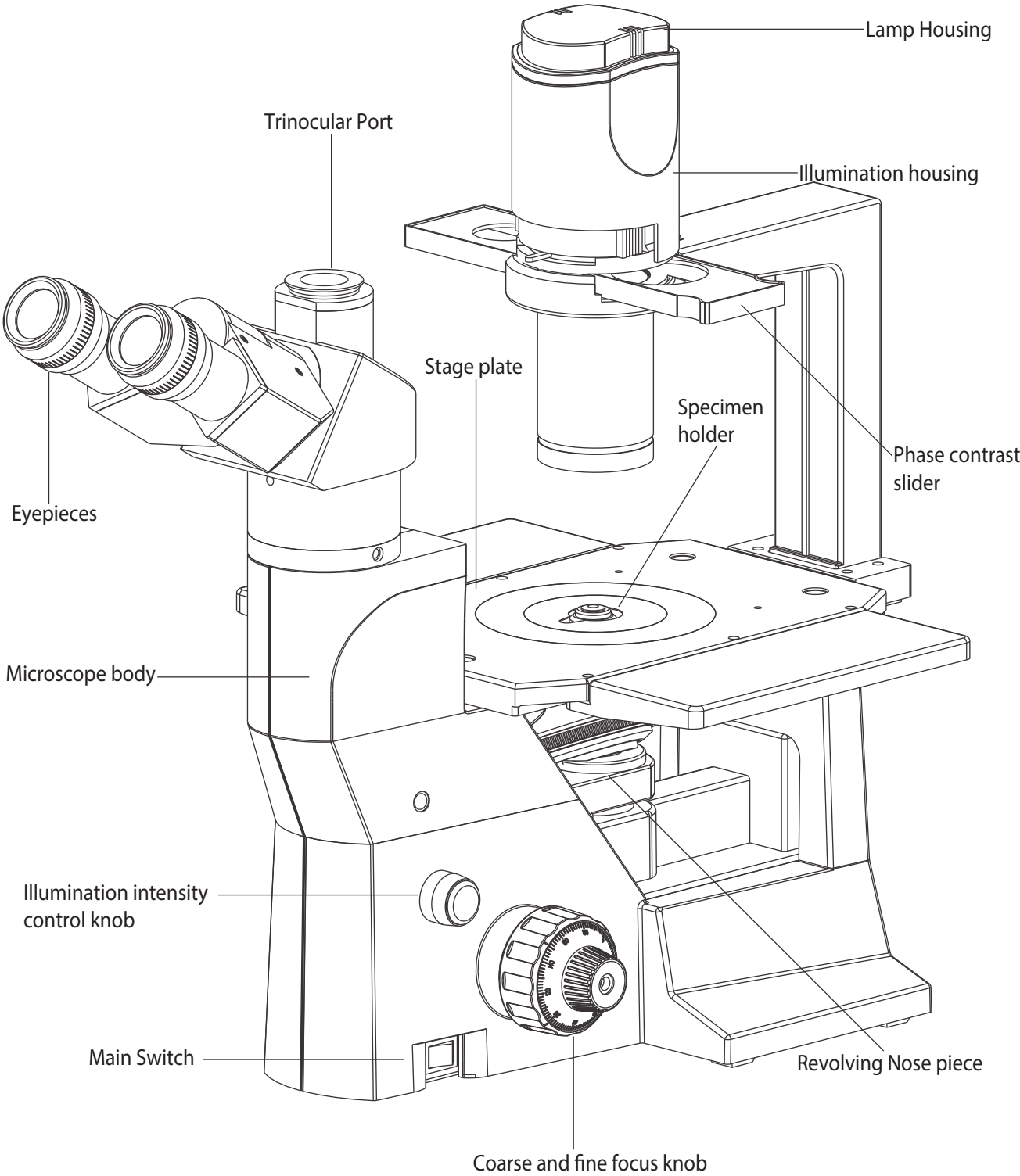
Wipe using a spiral movement. Do not use a zig-zag motion!

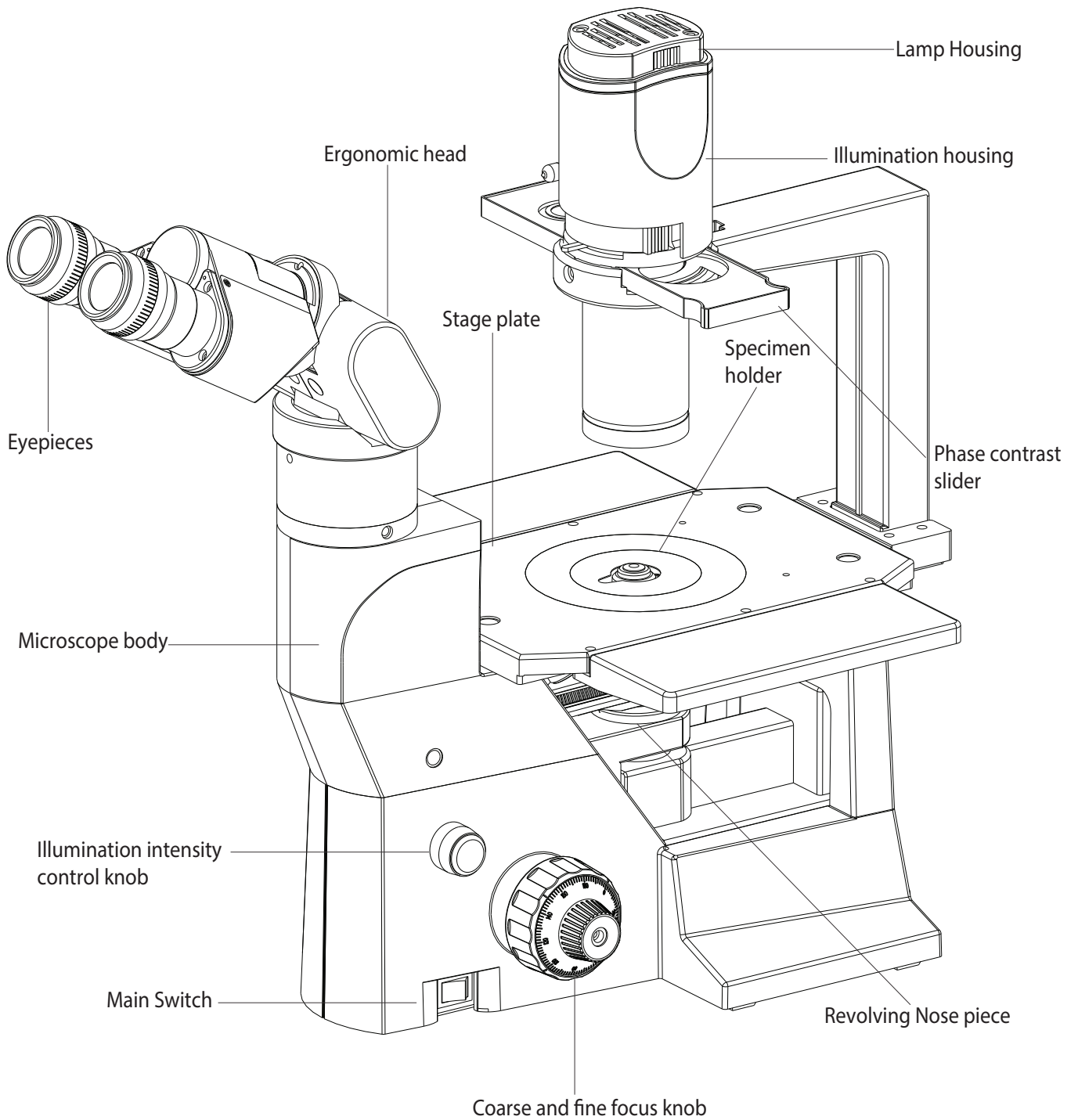
Cleaning of painted surfaces

Avoid the use of any organic solvent (e.g. thinner, xylene, ether, alcohol etc.) for cleaning of painted surfaces of the instrument. Painted surfaces can be cleaned with a very lightly moistened micro fiber cloth. Loose dust and dirt can be removed using a brush of soft hair used exclusively for this purpose.



4 TCM 400 TRINOCULAR



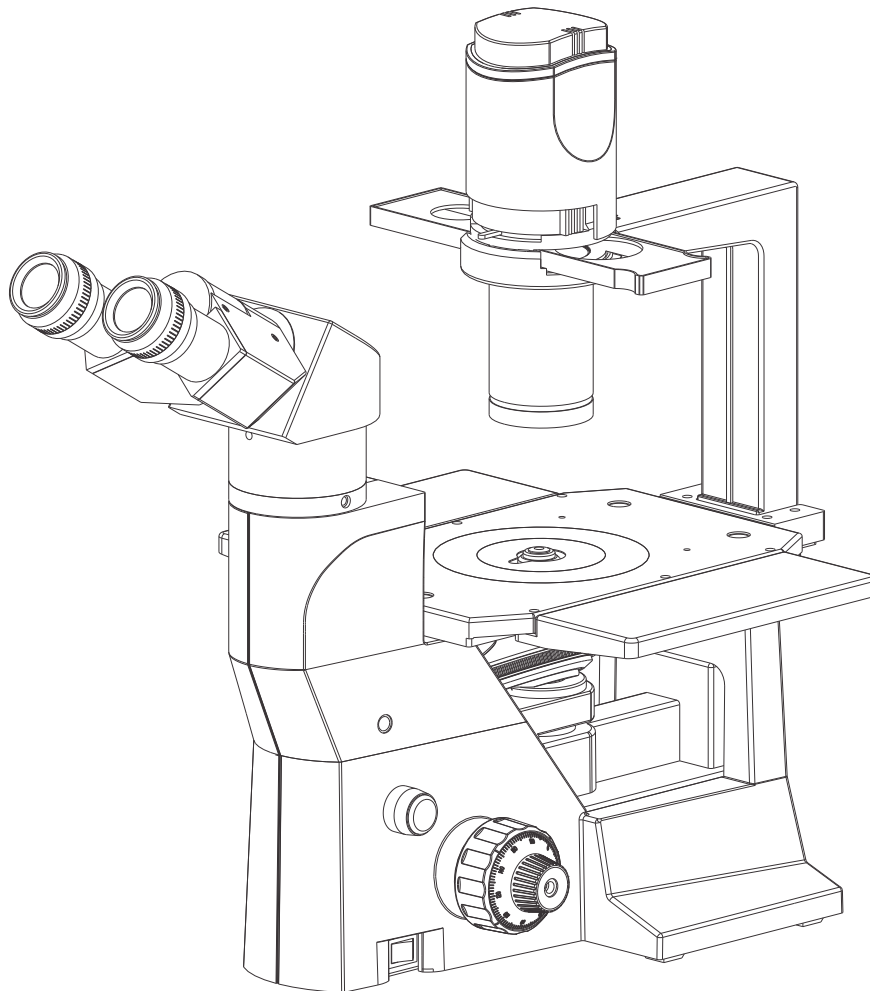


TCM400 comes securely packed in the custom configured foam packing, to prevent it from all shocks and transit damages.

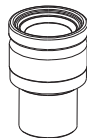
make sure that you receive all the following components before discarding packing material:

- a. Microscope body
- b. Observation tubes, as ordered
- c. Eyepieces
- d. Illumination housing
- e. Stage extension plates
- f. Specimen plate
- g. Phase slider
- h. Phase telescope
- i. Objectives as ordered
- j. Blue filter
- k. Green Filter
- l. Frosted filter
- m. Installation tools
- n. Power cord
- o. Spare fuses, as ordered
- p. Spare bulb, as ordered
- q. Adapter for telescope
- r. Dust cover

After removing your microscope from its packaging, make sure that all of the following contents are present. "Please note that the contents of your microscope may vary as the optional configuration, contrasting method or viewing body opted for may not be of the standard configuration highlighted here".



Daylight (blue) filter



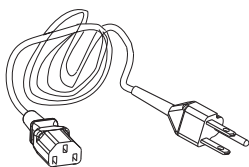
Paired Eyepieces



Allen Wrench 3mm



Frosted filter



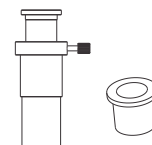
Power Cord



6V 30W Halogen bulb

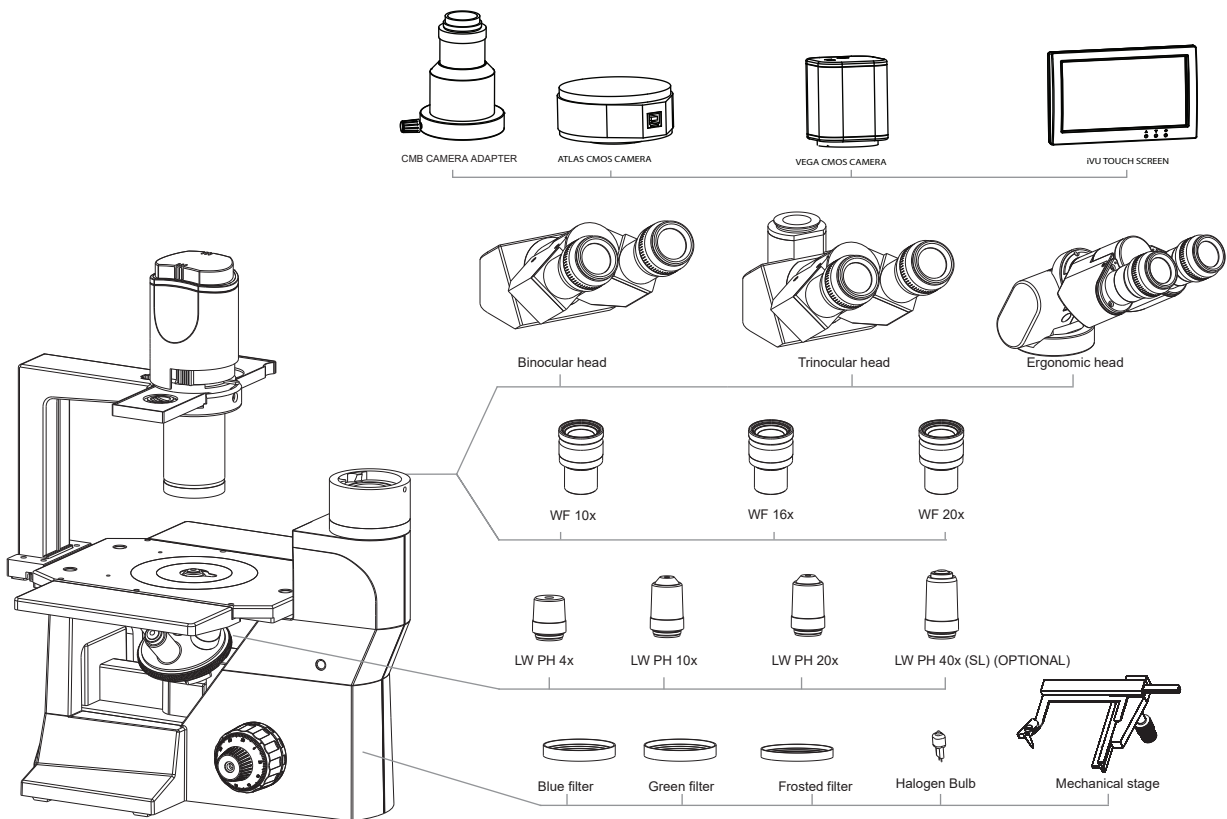


Green filter



Phase Telescope with adapter

SYSTEM DIAGRAM OF OPTIONAL ACCESSORIES



INSTALLATION AND OPERATION OF OPTIONAL ACCESSORIES

1 CAMERA MODULE SYSTEM

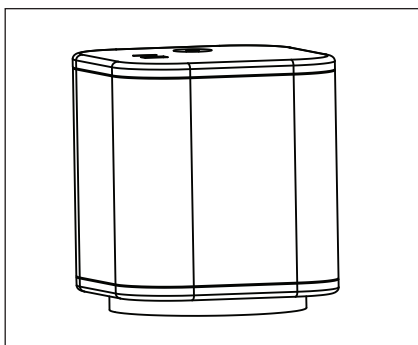


Fig. 2

1. Mount the Video adapter 1/2" on Trinocular observation head.
2. Mount Camera Module System on Video adapter.

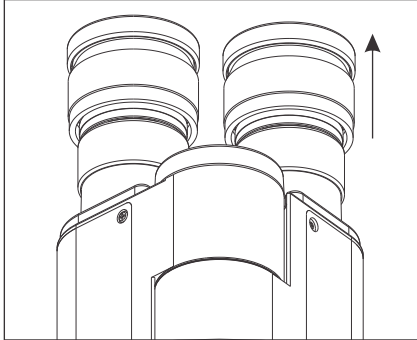
2 OPTIONAL EYEPIECES

Fig. 3.1

10X eyepieces are provided. To replace:

1. Pull out the 10x eyepieces out from the observation heads Ocular tube.
2. Insert desired eyepieces in empty ocular tube.(Fig.3.1)

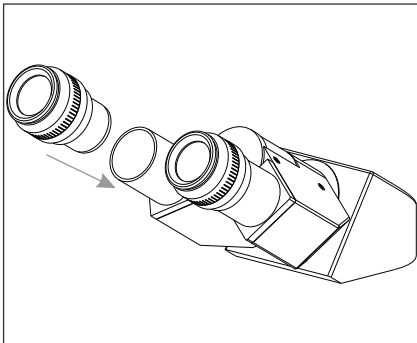


Fig. 3.2

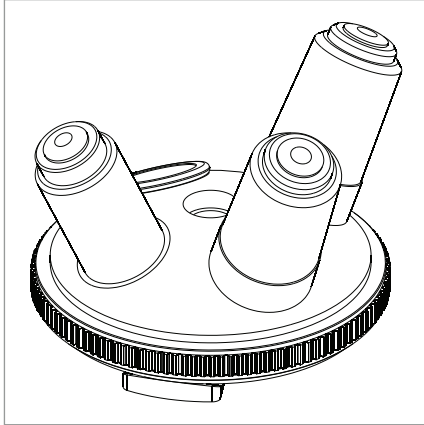
1 MOUNTING THE OBJECTIVES

Fig. 4

Settle down the nosepieces by rotating coarse adjustment knob towards back. Remove the dust prevention caps from the nose pieces.

Screw the objective with the lowest magnification into the revolving nose-piece from left side of the microscope. Turn the nose piece clockwise and mount the remaining objectives in ascending magnification order. It makes easier to change magnification. (Fig.4)

Clean the objective periodically.

- ⚠ Be sure to cover any unused threaded positions with the
- ⚠ dust prevention caps to prevent from dirt and dust getting inside.

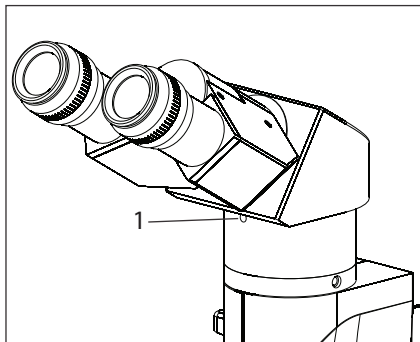
2 OBSERVATION HEAD

Fig. 5

Install the observation head using the following procedure:

1. Using allen wrench 3mm (provided), loosen the Head Locking Screw (1) and remove the dust cover cap provided in dovetail cavity as well as on observation head dovetail.
2. Mount the Observation Head by engaging the dovetail provided at the bottom of the head into the dovetail cavity provided in the microscope arm.
3. Tighten the Head Locking Screw (1) after positioning the Observation Head as desired. See figure5.

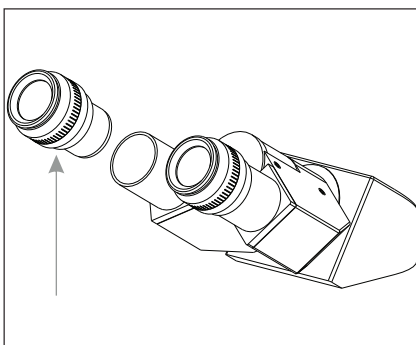
3 MOUNTING OF EYEPIECES

Fig. 6

Insert the eyepieces into the ocular tube of Observation Head using following procedure:

1. Remove the protective caps from the observation tube.
2. Insert 10x eyepieces into the ocular sleeve and tighten the thumb screw.

4.1 BASE UNIT

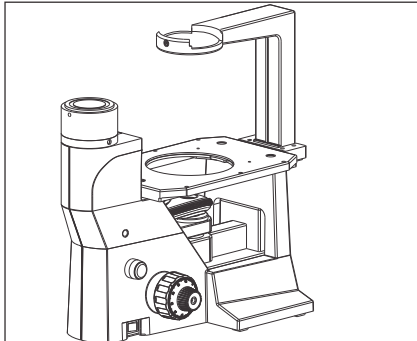


Fig. 7

Base unit is assembled with co-axial focus mechanism, turret, stage plate and the lamp house arm as shown in figure (fig 7) as shown.

4.2 INSTALLING THE TRANSMITTED LIGHT ILLUMINATOR

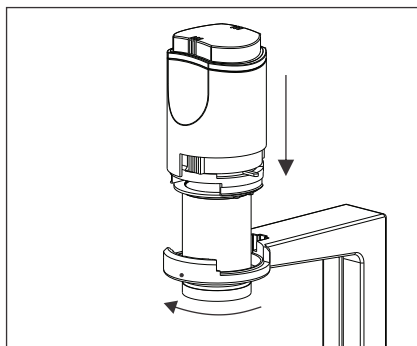


Fig. 8.1

1. Insert the illuminator into the arm opening and rotate the illuminator 90° clockwise so that the letters "NA 0.30" faces directly in front. See fig 8.1

2. Tighten the clamping screw using a allen wrench 3.0mm provided with microscope. See fig 8.2

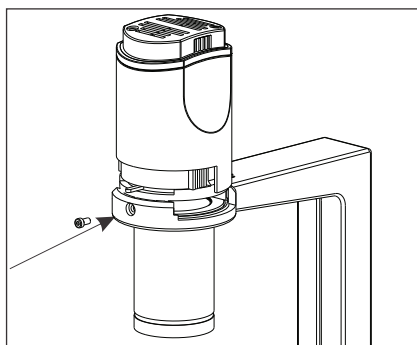
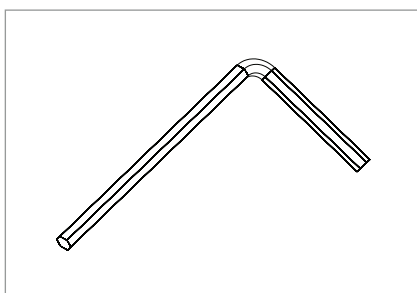


Fig. 8.2



Allen Wrench 3.0mm

4.3 PHASE SLIDER/ FILTER

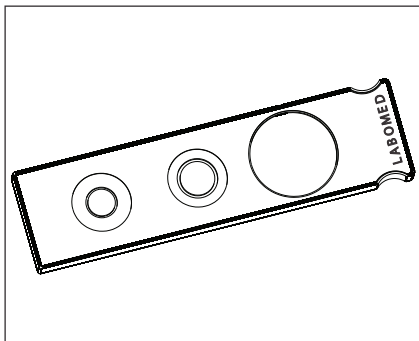


Fig. 9

1. Hold the phase slider face up (LABOMED written facing upward) with the finger on the right and slide it gently into the slot. As shown in Fig. 9.

Phase slider has click stops at the three positions i.e. 4x, 10x/20x and bright field stop .

2. 40x objective if ordered is provided with spare slider.

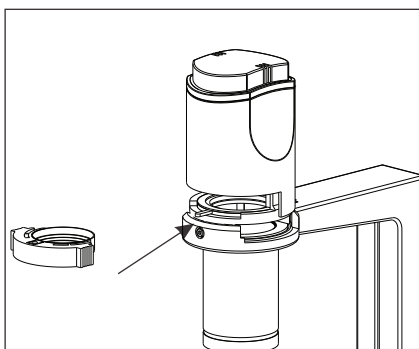


Fig. 10

3. The Blue filter is placed in the filter holder cartridge. The cartridge can be easily fitted in the lamp house. Hold the cartridge with your index finger and thumb and slide it in. See fig 10.

4.4 OBSERVATION HEAD

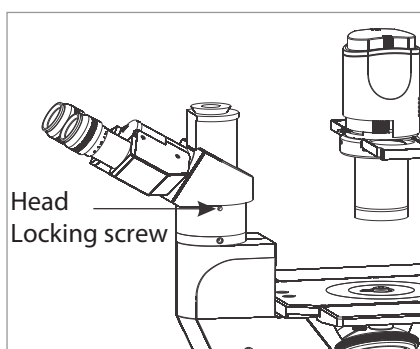


Fig. 11

1. Loosen the Head Locking screw using allen wrench 2.0mm (provided with the microscope) to such a position where it does not come out. See fig. 11.

2. Place the observation head on the stand and engage the dovetail ring provided at the bottom into the engaging recess provided on the stand.

3. Tighten the head locking screw after rotating the Binocular Head to your preferred direction of use.

5.0 ADJUSTING THE INTERPUPILLARY DISTANCE AND DIOPTER

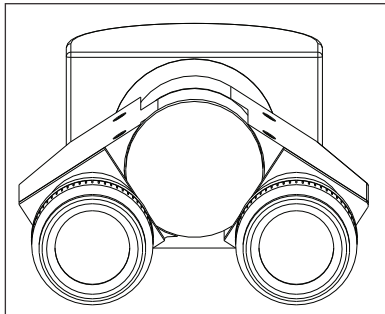


Fig. 12

While looking through the eyepieces, move both eyepieces until the left and right fields of view coincide completely. This matches your IPD.

5.1 ADJUSTING THE DIOPTER

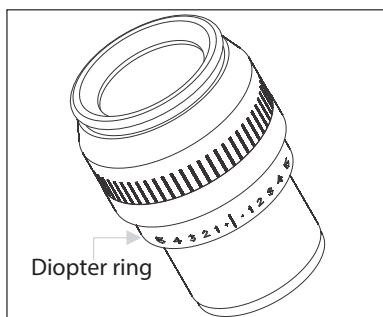


Fig. 13

Adjust the Diopter setting to "0" on the left side eyepiece. While looking through the left eyepiece with your left eye, turn the coarse and fine focus adjustment knobs to bring the specimen into focus.

Adjust Diopter setting to "0" on the right side eyepiece and while looking through the right eyepiece with your right eye, turn only the diopter adjustment ring to focus on the specimen. See figures 12 and 13.

6.0 MOUNTING THE STAGE EXTENSION & MECHANICAL STAGE

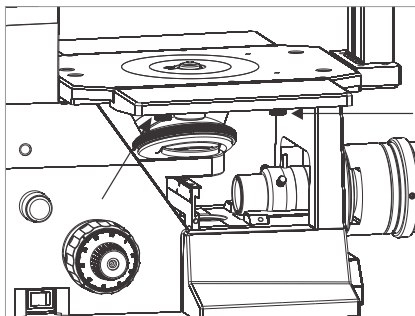


Fig. 14.1

The stage extension plate can be mounted in the left or right side or both sides of the stage to expand the stage surface. However, the stage extension plate and mechanical stage cannot be used simultaneously on the same side.
 Stage Extension Plate: Screw the studded clamping screw into the plain stage from underneath. Tighten them with hand until the plate is securely attached. Refer fig 14.1 and 14.2
 Mechanical Stage: This can be attached on either side of the stage. Attach in the same way as the stage extension plate. Refer to fig 14.3.

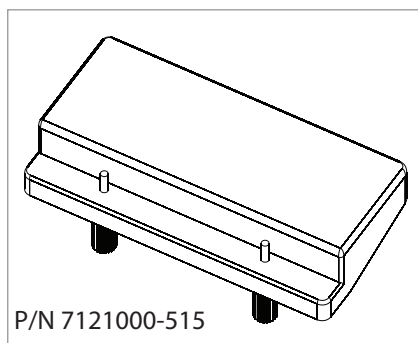


Fig. 14.2

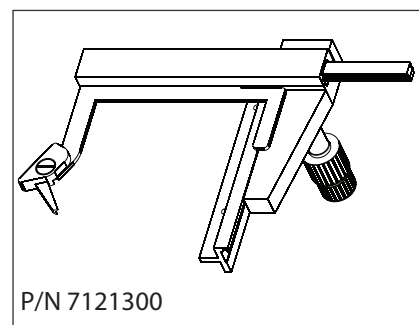


Fig. 14.3

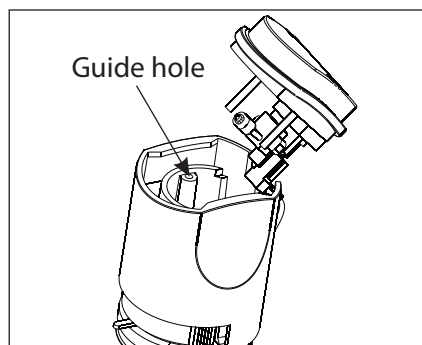
7.0 BULB REPLACEMENT

Fig. 15

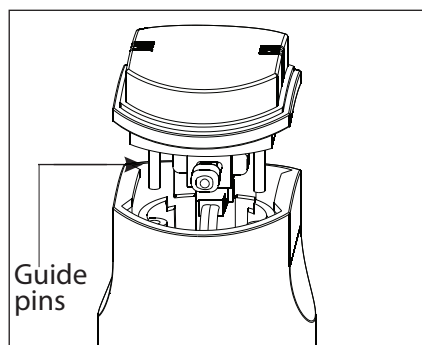


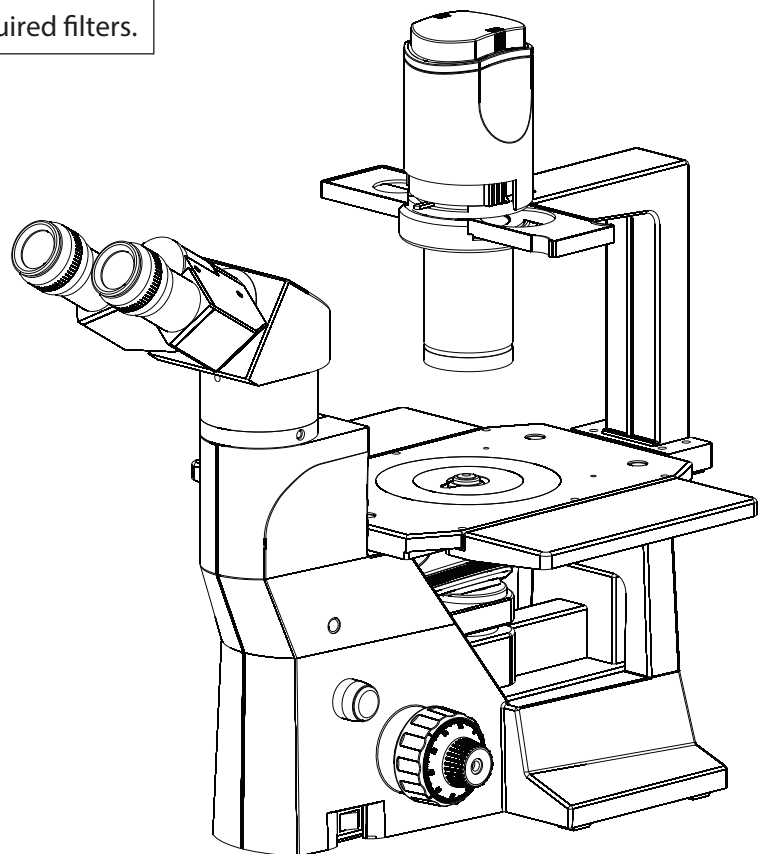
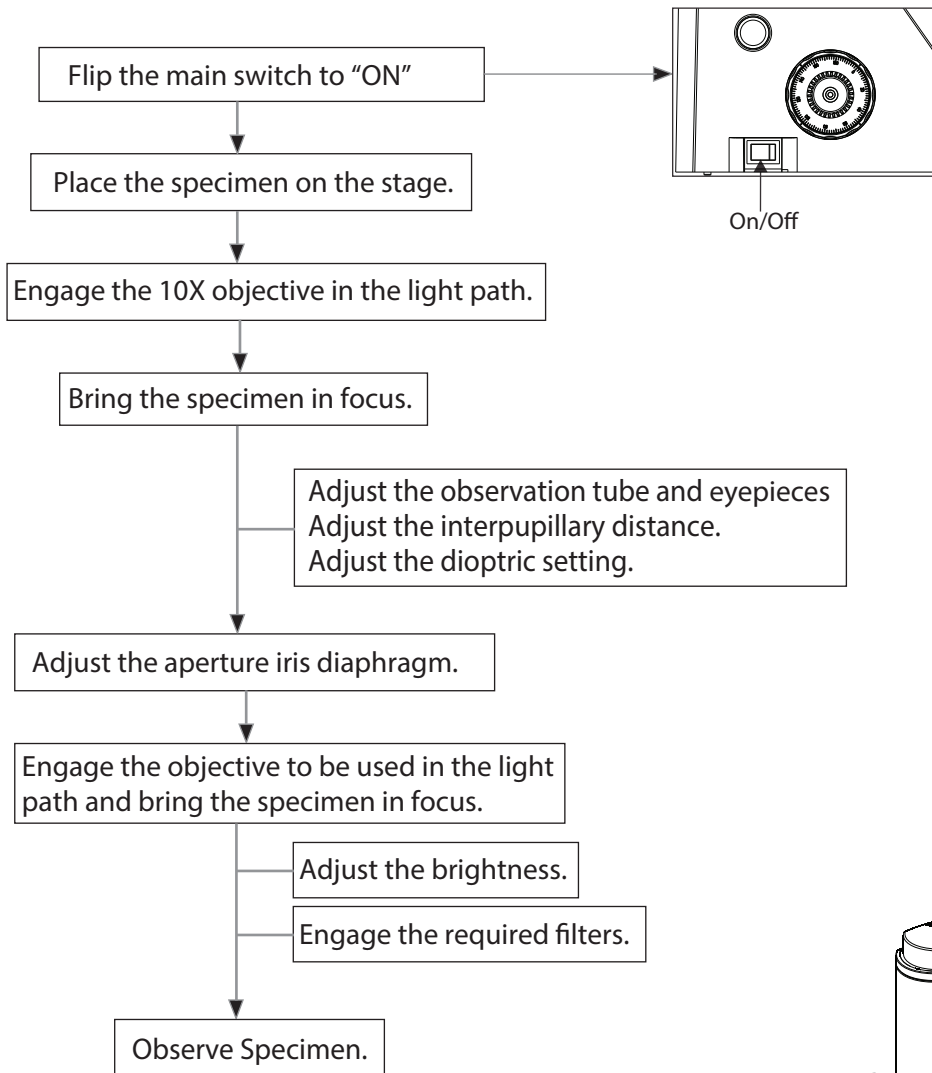
Fig. 16

Use only recommended 6V-30W high intensity Halogen bulb.

Pull open the Lamp socket and pull out the blown bulb using a soft cloth. Gently insert the new bulb. Do not squeeze or press the bulb too much which can cause damage to the bulb and hand.

Fit the lamp socket by aligning the guide pins with condenser guide holes. Push the lamp socket gently into the transmitted light illuminator. (Fig. 15, 16)

SUMMARY OF BRIGHT FIELD OBSERVATION PROCEDURE



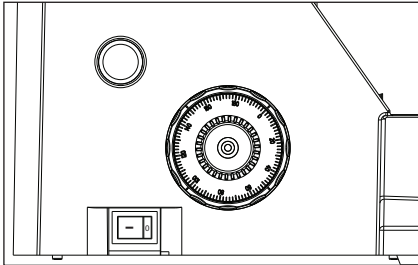
1 TURNING THE LAMP ON

Fig. 17

1. Flip the main switch to "I" (ON) as shown in Figure 17.
2. Rotating the light intensity adjustment knob in the direction of the arrow increases brightness and rotating knob in the opposite direction decreases brightness. The intensity bar on the knob indicates the direction of intensity level.

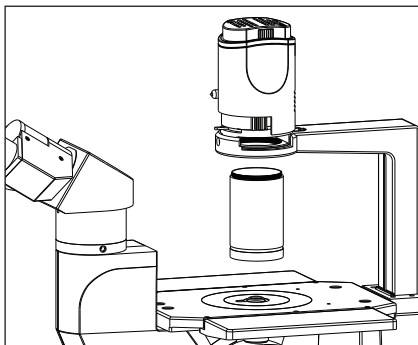
2 PLACING SPECIMEN ON THE STAGE

Fig. 18

Place the specimen on the stage plate. In case the specimen is in Petri dish, keep it on the petri dish holder.

In case you are using high Flask, remove the condenser front lens by rotating it anti clock wise to fit in the flask. As shown fig. 18

In case your microscope has mechanical stage mounted and you are using either glass slide or 90 well plate, use appropriate holders.

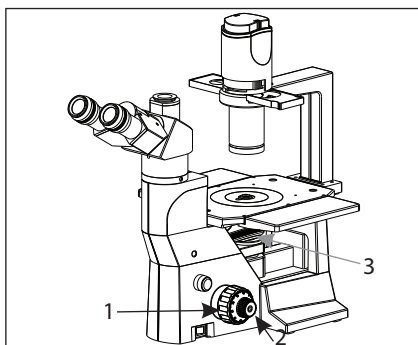
3 ADJUSTING THE FOCUS

Fig. 19

Focusing Procedure (Figure 1 9)

1. Rotate the coarse adjustment knob (1) clockwise so that the objective (3) is as close as possible to the specimen (We recommend starting with 10X).
2. While observing the specimen through the eyepieces, slowly rotate the coarse adjustment knob (1) counterclockwise to lower the objectives.
3. When coarse focusing of the specimen is obtained (an image is observed), rotate the fine adjustment knob (2) for fine detail focusing.

Working Distance (WD)

The WD refers to the distance between each objective and the Specimen, when acute focus of the specimen is obtained.

Objective Magnification	4X	10X	20X	40X
WD (mm)	16	7.2	5.35	2.0

4 ADJUSTING THE INTERPUPILLARY DISTANCE (IPD)

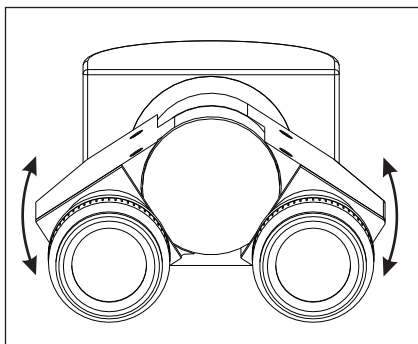


Fig. 20

The inter-pupillary distance adjustment consists of regulating the two eyepieces to align with both eyes' pupils so that you can observe a single microscopic image through two eyepieces in stereo vision. This greatly helps to reduce fatigue and discomfort during observation.

While looking through the eyepieces, move both eyepieces laterally until the left and right fields of view coincide completely. The position of index dot (•) indicates the inter-pupillary distance value.

Note your interpupillary distance so that it can be quickly referred to in the future. This is happen when multiple users work with the microscope.

5 ADJUSTING THE DIOPTER

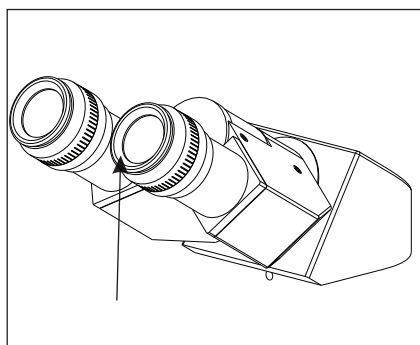


Fig. 21

Adjust the Diopter setting to "0" on the left side eyepiece. While looking through the left eyepiece with your left eye, turn the coarse and fine focus adjustment knobs to bring the specimen into focus.

Adjust the Diopter setting to "0" on the right side eyepiece and while looking through the right eyepiece with your right eye, turn only the diopter adjustment ring to focus on the specimen.

Using the Eye Guards

When Wearing Eyeglasses

Use with the eye guards in the normal, folded-down position. This will prevent the eyeglasses from being scratched.

When Not Wearing Eyeglasses

Extend the folded eye guards outwards (direction of the arrow) to prevent ambient light from entering into your line of vision.

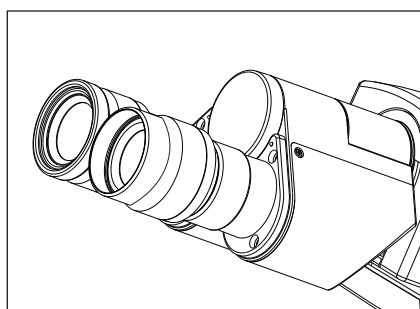






Fig.22

Under certain conditions, performance of the unit may be adversely affected by factors other than defects. If problems occur, please review the following list and take corrective action as needed. If problem Persists, please contact LABOMED or your local LABOMED dealer.

OBSERVATION	CAUSE	REMEDY
1. Although the illumination is on, the field of view is dark.	The socket pin is not connected to the illumination column at the back of the instrument. The bulb is burned out. The light intensity control is set to too low. The objective is not in position. Fuse is blown.	Connect it securely. Replace the bulb. Set the illumination to desired position. Make sure the objective is clicked properly. Replace fuses (250V / 5 Amps)
2. Dirt or dust is visible in the field of view.	Dirt/dust on the specimen. Dirt/dust on the eyepieces.	Clean the specimen. Clean them thoroughly.
3. Visibility is poor. - Image is not sharp. - Contrast is not sharp. - Details are indistinct. - Phase contrast effect cannot be obtained.	The objective is not correctly engaged. The aperture iris diaphragm is opened or stopped down too far in brightfield observation. The light annulus of the condenser does not match the phase annulus and phase	Replace it with a clean specimen. Clean eyepieces gently. Adjust the aperture. Overlap / adjust Phase annuli position.
4. One side of image is blurred.	The revolving nosepiece is not correctly engaged The specimen is not correctly mounted on the stage. The optical performance of the culture vessel bottom plate is poor.	Make sure that the revolving nosepiece clicks properly into place. Place it correctly on the stage. Use a vessel with a good profile Irregularity characteristic.
5. The bulb flickers and the Brightness is unstable.	The line voltage fluctuates The bulb is nearly burned out. The power cord is not connecting securely.	Use a voltage stabilizer. Replace the bulb. Connect it securely.
6. The coarse adjustment knob is too Difficult to rotate.	The tension adjustment ring is tightened too much. The tension adjustment ring is loosened too much.	Loosen it appropriately. Tighten it appropriately.
7. The field of view of one eye does not match that of the other.	Incorrect interpupillary distance adjustment. Incorrect diopter adjustment.	Adjust the interpupillary distance. Adjust the diopter.

1. Illumination	Built-in illumination system Halogen			
2. Focusing mechanism	Stage height adjustment mechanism Fine adjustment scale: 0.002 per graduation Fine adjustment stroke: 0.2mm per turn Total stroke: 28mm Co-axial coarse and fine focusing on ball drive Coarse adjustment travel per rotation			
3. Revolving nose piece	Quadruple positions fixed			
4. Observation tube		Binocular	Trinocular	Ergonomic
	Field number	22	22	22
	Tube tilting angle	30°	30°	0° - 25°
	Interpupillary distance adjustment range	52-75	52-75	52-75
5. Stage	Size	240 X 160mm (with mechanical stage)		
	Movement range	115 x 75mm		
	Specimen holder	Slide and Petri dish		
6. Condenser	Type	Incident condenser (daylight filter detachable)		
	N.A.	0.30		
	Aperture iris diaphragm	Built-in		
7. Dimensions & Weight	495.0mm (L) x 300.0mm (W) x 470.mm (H); 9.5 kg net			
8. Electrical	Lamp	Halogen 6V-30W		
	Lamp life	Upto 500 hours		
	Input	100V-240V AC, 50/60 Hz		
9. Operating environment	<p>Indoor use Altitude: Max. 2000 meters Ambient temperature: 5° to 40°C (41° to 104° F) Maximum relative humidity: 80% for temperature up to 31°C (88°F), Decreasing linearly through 70% at 34°C (93°F), to 50% relative humidity at 40°C (104°F) Supply voltage: 100V AC to 240V AC Supply voltage fluctuations: Not to exceed ±10% of the normal voltage. Power consumption: 50W; Fuse: F5A/250V Power supply to the halogen lamp: 6V AT 30W Pollution degree: 2 (in accordance with IEC60664) Installation/Over voltage category: II (in accordance with IEC60664)</p> <p> CAUTION: HIGH VOLTAGE FUZE REPLACEMENT</p> <p> WARNING: HIGH VOLTAGE POWER INLET</p> <p> EARTH</p> <p> WARNING: HIGH TEMPERATURE</p>			

Revision History

Rev. No.	Date of Release	DCR #	Change	App. By
1.3	April 14, 2022	DCR/06/22	Reviewed & Updated for typo error	S.Bal



Labo America Inc.
920 Auburn Court
Fremont, CA
94538
U.S.A.

Phone: 510-445-1257

Fax: 510-991-9862

Email: sales@laboamerica.com

www.laboamerica.com

EU REP.

Labomed Europe

Essebaan 50

NL-2908 LK Capelle a/d IJssel

The Netherlands

Tel: +31 (0)10 4584222

Fax: +31 (0)10 4508251

E-mail: info@labomedeuropa.com



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