





Introduction

Thank you for purchasing the Euromex Delphi-X Observer

The Delphi-X Observer series has been designed with all kind of Life Sciences applications and great durability in mind. This resulted in a modern, robust and high level microscope for advanced use, equipped with the best optical and mechanical components. An ideal microscope for daily cytology and anatomic pathology use. The 25 mm field of view of the eyepieces and the plan apochromatic objectives enable observations with perfect color rendering at high resolving powers. Specific attention to production methods resulted also in an excellent price/performance ratio. Please read this manual carefully before using this product to ensure correct and save usage

- The contents of this manual are subject to change without notice
- The appearance of the actual product can differ from the models described in this manual
- Not all equipment mentioned in this manual has to be part of the set you have purchased
- All optics are anti-fungus treated and anti-reflection coated for maximum light throughput

Contents

| General safety instructions | 3 |
|---|----|
| Dangers associated with the operation | 3 |
| Photobiological safety LED, important safety instructions | 3 |
| Prevention of biological and infectious hazards | 3 |
| Disinfection and decontamination: | 4 |
| Model with rechargeable batteries | 5 |
| Environment, storage and use | 5 |
| Configuration, construction and controls | 6 |
| Assembling Delphi-X Observer | 8 |
| Operation | 12 |
| Placing the slide | |
| Adjusting tension of X and Y axis control knobs | 12 |
| Switching between light sources | 12 |
| Getting the specimen in focus | 12 |
| Adjusting the coarse focus tension | 13 |
| Setting the focus lock | 13 |
| Switching the fine focus knobs | 13 |
| Adjust interpupillary distance | 13 |
| Adjust diopter of the eyepieces | 14 |
| The correct eye point | 14 |
| Select eyepiece and camera light throughput | 14 |
| Centering the condenser | 15 |
| Using the aperture diaphragm | 15 |
| Using LED WITH frosted filter | 16 |
| Using HALOGEN WITH LBD, ND 6 and ND25 filters | 16 |
| iCare sensor | 16 |
| Replacing the fuse | 16 |
| Cleaning optics | 17 |
| Troubleshooting | 18 |

General safety instructions

Intended use: a non-medical device

This microscope is intended for general observation of cells and tissues, with transmitted/reflected illumination and with the specimen fixed on a slide

Intended use as in vitro medical device class A (Regulation (EU) 2017/746)

The models labelled with suffix /MD may be used as in vitro medical device and are intended for observation and diagnostics of cells and tissues in hospitals or by physicians in private practice in pathology, anatomy and cytology applications. To be used with transmitted/reflected illumination and with the specimen fixed on a slide. Physicians use microscopes to identify the different types of cells and spot abnormal cells. This product helps in identifying and treating diseases

Dangers associated with the operation

- Improper use could result in injury, malfunction or damage to property. It must be ensured that the operator
 informs every user of existing hazards
- Danger of electrocution. Disconnect the power to the entire lighting system before installing, adding or changing any component
- Not to be used in corrosive or explosive environments
- · Avoid direct exposure of eyes to the collimated light beam or direct light from the light guides or fibres
- To avoid a hazard to children, account for all parts and keep all packing materials in a safe place

Photobiological safety LED, important safety instructions

- Avoid direct eye exposure to any LED light source while switched on
- · Before looking through the eyepieces of the microscope, lower the intensity of the LED illumination
- Avoid long and high-intensity exposure to LED light because this may cause acute damage to the retina of the eye

Prevention of biological and infectious hazards

Infectious, bacterial or viral biohazard substances under observation may be a risk to the health of humans and other living organisms. Special precautions should be taken during in vitro medical procedures:

- Biological hazards: keep a logbook of all the biological substances or pathogenic microorganisms that were
 under observation with the microscope and show it to everybody before they use the microscope or before
 they do some maintenance work on the microscope! Agents can be bacterial, spores, enveloped or nonenveloped virus particles, fungi or protozoa
- Contamination hazard:
 - A sample that is properly enclosed with a cover glass never comes in direct contact with the microscope
 parts. In that case prevention of contamination lies in the handling of the slides; as long as the slides
 are decontaminated before use and are undamaged and treated normally, there is virtually zero risk of
 contamination
 - A sample that is mounted on a slide without cover glass, can come in contact with components of the
 microscope and may be a hazard to humans and/or the environment. Therefore, check the microscope and
 accessories on possible contaminations. Clean the microscope surfaces and its components as thoroughly
 as possible. Should you identify a possible contamination, inform the local responsible person in your
 organisation
 - Microscope operators could be contaminated from other activities and cross-contaminate components of the
 microscope. Therefore, check the microscope and accessories on possible contaminations. Clean the microscope
 surfaces and its components as thoroughly as possible. Should you identify a possible contamination, inform
 the local responsible person in your organisation. It is recommended to wear sterile gloves when preparing the
 slides and handling the microscope in order to reduce contamination by the operator
- Infection hazard: direct contact with the focusing knobs, stage adjustments, stage and eyepieces/tubes of
 the microscope can be a potential source of bacterial and/or viral infections. The risk can be limited by using
 personal eyeshades or eyepieces. You can also use personal protections such as operation gloves and/or safety
 goggles, which should be changed frequently to minimize the risk

Disinfectant hazards: before cleaning or disinfecting, check if the room is adequately ventilated. If not, wear
respiratory protective gear. Exposure to chemicals and aerosols can harm human eyes, skin and respiratory
system. Do not inhale vapours. During disinfection, do not eat, drink or smoke. Used disinfectants must be
disposed of according to local or national regulations for health and safety

Disinfection and decontamination:

- Exterior casing and mechanical surfaces must be wiped with a clean cloth, dampened with a disinfectant
- Soft plastic parts and rubber surfaces can be cleaned by gently wiping a clean cloth, dampened with a
 disinfectant. Discoloration can occur if alcohol is used
- The front lens of eyepieces and objectives are sensitive to chemicals. We recommend not to use aggressive
 disinfectants but to use lens paper or a soft fibre-free tissue, damped in cleaning solution. Cotton swabs may
 also be used. We recommend you use personal eyepieces without eyeshades in order to minimize risk
- Never immerse or dip the eyepiece or objective into a disinfectant liquid! This will damage the component
- Never use abrasive compounds or cleaners that may damage and scratch optical coatings
- Properly clean and disinfect all possible contaminated surfaces of the microscope or contaminated accessories before storing for future use. Disinfection procedures must be effective and appropriate
- Leave the disinfectant on the surface for the required exposure time, as specified by the manufacturer. If the
 disinfectant evaporates before the full exposure time, reapply disinfectant on the surface
- For disinfection against bacteria, use a 70% aqueous solution of isopropanol (isopropyl alcohol) and apply for at least 30 seconds. Against viruses, we recommend to refer to specific alcohol or non-alcohol based disinfection products for laboratories

Before returning a microscope for repair or maintenance through a Euromex dealer, an RMA (return authorization form) together with a decontamination statement must be filled in! This document - available from Euromex for any reseller- must be shipped together with the microscope at all times

Reference documents:

World Health Organisation:

https://www.who.int/ihr/publications/biosafety-video-series/en/

Robert Koch Institut:

https://link.springer.com/content/pdf/10.1007/s00103-013-1863-6.pdf

US Centre for Disease Control and prevention

https://www.cdc.gov/infectioncontrol/guidelines/disinfection/index.html

Handle with care

- This product is a high quality optical instrument. Delicate handling is required
- Avoid subjecting it to sudden shocks and impacts
- · Impacts, even small ones, can affect the precision of the instrument

Handling the LED

Note: Always disconnect the power cord from your microscope before handling the LED bulb and power unit and allow the system to cool down approximately 35 minutes to avoid burns

- Never touch the LED with your bare hands
- Dirt or fingerprints will reduce the life span and can result in uneven illumination, lowering the optical
 performance
- Use only original Euromex replacement LEDs
- The use of other products may cause malfunctions and will void warranty
- During use of the microscope the power unit will get hot; never touch it while in operation and allow the system
 to cool down approximately 35 minutes to avoid burns

Dirt on the lenses

 Dirt on or inside the optical components, such as eyepieces, lenses, etc., affects the image quality of your system negatively



- Always try to prevent your microscope from getting dirty by using the dust cover, prevent leaving fingerprints
 on the lenses and clean the outer surface of the lens regularly
- Cleaning optical components is a delicate matter. Please, read the cleaning instructions further on in this
 manual

Model with rechargeable batteries

- · Always disconnect the power cord from the microscope before you replace the rechargeable batteries
- The rechargeable batteries must not be thrown away as regular trash but should be taken to special waste
 collection sites, according your local or national regulations
- Risk of explosion: when removing the rechargeable batteries, do not throw the batteries into fire or any other heat source
- Do not replace the rechargeable batteries with non-rechargeable batteries
- Avoid extreme environmental conditions and temperatures which could affect the rechargeable batteries and lead to fire, explosion or leakage of hazardous substances
- If the rechargeable batteries have leaked, avoid contact of the chemicals with skin, eyes and mucous membranes
- When in contact with the chemicals, flush the affected areas immediately with plenty of fresh water and seek medical attention

Environment, storage and use

- This product is a precision instrument and it should be used in a proper environment for optimal use
- Install your product indoors on a stable, vibration free and level surface in order to prevent this instrument to
 fall thereby harming the operator
- · Do not place the product in direct sunlight
- The ambient temperature should be between 5 to +40°C and humidity should be within 80% and 50%
- Although the system is anti-mold treated, installing this product in a hot, humid location may still result in the
 formation of mold or condensation on lenses, impairing performance or causing malfunctions
- Never turn the right and left focus knobs in opposite directions at the same time or turn the coarse focus knob
 past its farthest point as this will damage this product
- Never use undue force when turning the knobs
- Make sure that the microscope system can dissipate its heat (fire hazard)
- Keep the microscope away from walls and obstructions for at least approximately 15 cm
- Never turn the microscope on when the dust cover is in place or when items are placed on the microscope
- Keep flammable fluids, fabrick, etc. well out of the way

Disconnect power

Always disconnect your microscope from power before doing any maintenance, cleaning, assembling or replacing LEDs to prevent electric shocks

Prevent contact with water and other fluids

Never allow water or other fluids to come in contact with your microscope, this can cause short circuiting your device, causing malfunction and damage to your system

Moving and assembling

- This microscope is a relatively heavy system, consider this when moving and installing the system
- Always lift the microscope by holding the main body and base of the microscope
- Never lift or move the microscope by its focusing knobs, stage or head
- When needed, move the microscope with two persons instead of one



Configuration, construction and controls

This chapter describes the main parts and functions of the Delphi-X Observer



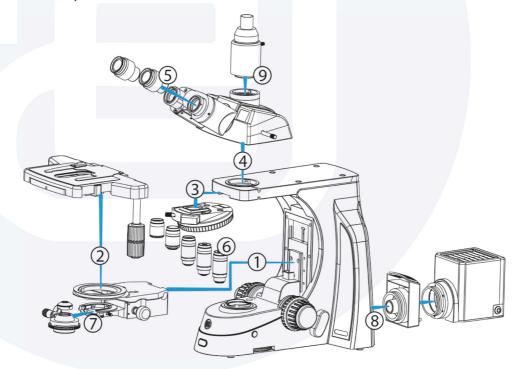
| 1A | Photo port | 1J | Lamp housing unit fastening screw |
|----|--|----|-----------------------------------|
| 1B | Optical light path selector | 1K | Excess cable storage |
| 1C | Nosepiece | 1L | Allen wrench tool |
| 1D | Objectives | 1M | Lamp housing unit |
| 1E | Stage | 1N | On/off switch |
| 1F | Coaxial control X-Y stage movement | 10 | Power socket and fuse holder |
| 1G | Coarse and fine focusing control knobs | 1P | Lamp housing unit plug |
| 1H | Tension control knobs | 1Q | Power connector (not used) |
| 11 | Field diaphragm adjustment wheel | 1R | External grounding rod |



| 2A | Eyepieces | 2L | Label for selecting light path of photo tube |
|----|--|----|--|
| 2B | B Eyepiece tubes | | Head |
| 2C | Screw for fixing extension slot | 2N | Main body |
| 2D | DIC extension slot | 20 | Slide holder |
| 2E | Condenser height control knob | 2P | Condenser |
| 2F | Focusing lock | 2Q | Condenser centering screw |
| 2G | Coarse and fine focusing control knobs | 2R | Coaxial control X-Y stage movement |
| 2H | Filter selector | 25 | Coarse and fine focusing control knobs |
| 21 | Light selector | 2T | iCare on/off button |
| 2J | Light intensity control knob | 2U | iCare sensor |
| 2K | Collector lens | | |

Assembling Delphi-X Observer

This chapters describes the steps that need to be taken to assemble the Delphi-X Observer microscope. Euromex Microscopes will always try to keep the number of assembly steps for their customers as low as possible but there are some steps that need to be taken. The steps mentioned on the following pages are not always necessary but described for your convenience nonetheless:



The diagram shows the order of each component's installation

| Step 1 | Attaching the focus cassette | Step 5 | Placing the eyepieces |
|--------|--|--------|--------------------------------|
| Step 2 | Attaching the mechanical X/Y stage | Step 6 | Mounting the objectives |
| Step 3 | Attaching Nosepiece | Step 7 | Placing the condenser |
| Step 4 | Step 4 Placing the microscope head, C-mounts and photo ports | | Attaching the LED lamp chamber |
| | | Step 9 | Attaching the phototube |

Step 1 | Attaching the focus cassette

- Attach the focus cassette according to the path shown in figure 1
- The dovetail slot needs to be aligned with the slot of focusing cassette
- Slide it down until it reaches the locking pin
- Then use the hex wrench tool to tighten the screw shown as I (in figure 2)





Step 2 | Attaching the mechanical X/Y stage

- Turn the coarse focus knob until the elevating section is brought to the lowest position
- · Attach the mechanical object stage according figure 3 by aligning the stage above the ring of the focus cassette
- Fix the mechanical stage into place with screw (figure 4)





Step 3 | Attaching Nosepiece (figure 5)

- Slide the nosepiece into the slot
- Fix into place with screw (II)



Step 4 | Placing the microscope head (figure 6)

- Place the head by loosening the screw (III)
- Mount the head in its position inside the microscope arm
- Secure it by tightening the screw again



Step 5 | Placing the C-mount or photo port, onto the microscope head (figure 7)

- loosen the screw (IV)
- Place either the C-mount or photo port and tighten the screw



Step 6 | Placing and mounting the eyepieces (figure 8)

- First remove the dustcover of eyepiece tubes
- Insert the eyepieces into the eyepiece tubes



Step 7 | Placing the condenser (figure 9)

- Use the condenser height control knob (V) to lower the condenser holder to the lowest position
- Insert the condenser into the holder as shown in figure 9
- Then secure the condenser by fixing the screw indicated
- Centering the condenser is described later in this manual



Step 8 | Attaching the LED Lamp HOUSING UNIT (figure 10A)

- Slide the lampunit (Halogen or LED) into position at back of microscope base
- Use the wrench screw tool to secure bolt (VI)



Step 9 | Connecting the power cord

The Delphi-X Observer microscopes support a wide range of operating voltages: 100 to 240 V. *Please use a grounded power connection*

- Make sure the power switch is off before connecting.
- Insert the connector of the power cord into the Delphi-X Observer power socket (figure 10B), and make sure it connects well
- Insert the other connector into the mains socket, and make sure it connects well
- Put the power switch to ON



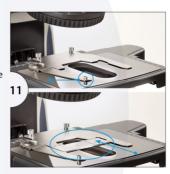
Do not bend or twist the power cord, it will get damaged. Use the special cord supplied by Euromex. If it is lost or damaged, choose one with the same specifications

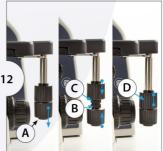
Operation

Placing the slide

(figure 11)

- Lower the condenser slightly from the uppermost position by turning the condenser focus knob
- Open both aperture and field diaphragm entirely
- Bring the 4x objective (or lowest objective in your configuration) into the optical path by rotating the nosepiece untill the right objective clicks into position
- Pull back the spring clamp of the specimen holder and gently place the slide into position
- Gently release pressure from the spring clamp so it softly moves back in
 position securing the slide
- Use the X and Y axis control knobs of the mechanical stage to move to area of interest of the slide into the light path





Adjusting tension of X and Y axis control knobs

(figure 12)

- The degree of tension on the X and Y axis control knobs can be adjusted
- To do that, draw down hand wheel (A) and find two adjusting rings (B,C)
- By rotating these rings the movement of the knobs can be set lighter and 12 heavier
- Ring B is used for adjusting the X direction
- Ring C is used for adjusting the Y direction

Switching between light sources

(figure 13)

Next to the intensity controller there is a button for switching between transmitted and reflected illumination. The standard brightfield configuration used for this manual does not have this option

- When the button is pushed in, the light is set to reflected mode
- When the button is pushed out, the light is set to transmitted mode (standard)



Getting the specimen in focus

(figure 14)

- Use the coarse control knobs to adjust the focus quickly and roughly
- Get the specimen into sight through the eyepieces
- Then use the fine focus control knob to adjust the focus in detail







Adjusting the coarse focus tension

(figure 15)

Next to the right side coarse focus know there is a ring for adjusting the coarse focus tensions This can be used to make the coarse control move lighter or heavier, according to user preference



Setting the focus lock

(figure 16)

Next to the left side coarse focus know there is a ring setting the focus lock. The focus lock can be used to limit the maximum position of the stage at a certain height. This is ideal for preventing objectives to get damaged, slides from breaking or to set the stage at a reference height

- Move the stage to the desired height then fix the ring tight to lock the mechanical stage's maximum height
- The stage still can be lowered but the highest position is now limited to the set position
- Release the ring to undo the focus lock



Switching the fine focus knobs

(figure 17 and 18)

The fine focus knobs can be switched from left to right to meet user preference

- Pull the knobs with moderate force to release the magnet which is holding the knobs onto the stand
- Attach the magnets onto the holder and let it grab the knobs again to mount them onto the holder







Adjust interpupillary distance

The Delphi-X Observer has an interpupillary distance range of 47 to 78 mm. The correct interpupillary distance is reached when one round image is seen in the field of view



Field of view before adjustment

Field of view after adjustment

This distance can be set by either pulling the tubes towards each other or pulling them from each other. This distance is different for each observer and this should be set individually. When more users are working with the microscope it is recommended to remember your interpupillary distance for a quick setup during new microscopy sessions

Adjust diopter of the eyepieces

(figure 19)

In order to compensate for human eye differences, distortion, thickness differences in cover glasses and tune for the best parfocality between objectives, one can use the diopter to do so. Take a good prepared slide for your reference:

- Set (both) the diopter adjustments of the eyepieces to "0"
- Select the 10x objective, look for a interesting area on the specimen and focus on this area
- Select the 40x objective and focus on the specimen

Warning: don't change the coarse and fine adjustment anymore

- With your dominant eye open (close your other eye), rotate the diopter adjustment from "+" to "-" untill the selected area get as sharp as possible
- If during this operation the image becomes unsharp, take your eyes from the eyepieces and turn the diopter adjustment, without looking into the eyepieces, a few divisions back from "-" to "+".
- Look into the eyepiece again and turn the diopter adjustment from '+' to '-' untill the selected area on your specimen gets the optimal sharpness
- Repeat for your non-dominant eye, and with the second diopter

Verification:

- Take your eyes from the eyepieces and look for 2 seconds to a far point in the room in order to "reset" your eyes
- Look again into the eyepieces. If the adjustment is not good, repeat the operation till you reach the same sharpness for the 10x and 40x objective without touching the coarse and micrometric adjustments







The correct eye point

(figure 20)

The eye point is the distance from the eyepiece to the user's pupil. To obtain the correct eye point, move the eyes towards the eyepieces until a sharp image is reached at a full field of view

Select eyepiece and camera light throughput

(figure 21)

The Delphi-X Observer gives users the option to select out of three output types, giving large flexibility when using cameras. The push/pull rod on the side of the microscope head can be set to 3 positions:

POSITION 1 | The optical light path is sent to the eyepieces only. Ideal for when no camera is used

POSITION 2 | The optical light path is sent to the eyepieces for 20% only. Ideal standard setting for when a camera is used

POSITION 3 | The optical light path is sent to the camera only. Ideal for when camera is used at low light imaging

These positions are indicated on the head as well for user convenience

| Icon | Action | Eyepiece / camera |
|-----------------------|-----------------------------|-------------------|
| H 4 | Push rod in completely | 100 / 0 |
| ⊢ ■ (© | Pull rod towards the middle | 20 / 80 |
| | Pull rod out completely | 0 / 100 |

Centering the condenser

(figure 22)

- Move the condenser to the top position (1)
- Focus on a specimen using the smallest objective (f.e. 4x or 10x objective)
- Close the field diaphragm (2)*
- Use screws (figure 23) to move the field diaphragm into to view center
- Open the field diaphragm carefully to the outside of the field of view to ensure the field diaphragm is in the center and so the condenser has been centered properly

Using the aperture diaphragm







The aperture diaphragm (figure 24/3) should be used to adjust the numerical aperture, not to adjust image brightness. When the aperture diaphragm is opened to the $70 \sim 80\%$ of objective aperture the ideal position is reached

The simplest way to do this is to use the markings on the condenser

Example: when a 40x objective with N.A. 0.65 is used, one can set the condenser to 70 – 80% of 0.65 which is 0.45 to 0.58

^{*} The iris can only be closed in brightfield position, in the phase or DIC position the iris is blocked from closing for best image quality

Using LED WITH frosted filter

(figure 25)

For LED models there is only 1 push button

• Push the button in for placing the frosted filter into the light path



LED version with frosted filter

Using HALOGEN WITH LBD, ND 6 and ND25 filters

(figure 26)

The halogen version has three filter options:

- 1. LBD is a filter for increasing color temperature
- 2. ND25 is a filter with 25% light transmittance
- 3. ND6 is a filter with 6% light transmittance



Halogen version with LBD and two ND filters

iCare sensor

(figure 27)

The unique iCare Sensor is developed to avoid unnecessary loss of energy. The illumination of the microscope automatically switches off shortly after the user steps away from his or her position

- Pushing the iCare button will re-active the light
- The iCare function is turned on by default
- To turn off the iCare function push the iCare button for 4 seconds
- The function will be deactivated and the bright LED will dim to indicate the function has be turned off
- Repeating this step will turn the function back on



Replacing the fuse

(figure 28)

The fuse is placed in a drawer

- To open it push the drawer aside with a screwdriver
- Take out the drawer and replace the fuse gently



Cleaning optics

How to keep the optics clean?

Dust and dirt particles have a negative affect on the image quality. Keeping the optical system of your microscope clean is essential for the best image quality and overall lifetime of your microscope. Dust and dirt on optical elements such as lenses, prisms and filters that are left unattended can become difficult - or even impossible - to remove and may cause mold

FIGURE A

- Place your objective or eyepiece on a secure location
- Objectives can be screwed into the cover of an objective case
- Eyepieces can be placed in the microscope box
- Condensers and collector lenses can remain in place in the microscope

FIGURE B |

 To prevent scratches on coatings and optical glass try to remove dirt and dust that sticks to the optical surface first with an air-blower or with pressurized dry air (oil-free and under moderate pressure version only)

FIGURE C |

- Use an absorbent lens paper or cotton swap.
- Dampen a swap or towel with a small amount of lens cleaning fluid or cleaning mixture (either pure iso-propanol or a mixture of 7 parts ether and 3 parts alcohol)

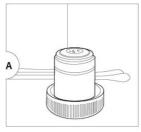
FIGURE D

- Clean the lens by using the tip of the cotton swap or the lens paper. Use enough lens paper so that solvents do not dissolve oils from your hands which can make their way through the paper on to the coated surface
- When cleaning a large lens surface, wipe with little pressure from the centre towards the periphery in a circular motion. Do not use zig-zag motion
- Discard each lens paper or cotton swap after a single use

FIGURE E

- Wait until the cleaning fluid is evaporated, or speed up this process by using pressurized dry air
- Check if the surface is clean by using a magnifying glass
- Place the cleaned item back on the microscope

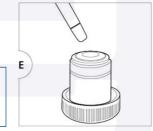
Please note that cleaning of the optical surfaces indicated in this instruction only applies to external surfaces of objectives, eyepieces, filters and condensers. Internal surfaces must always be done by your Euromex microscopes distributor











Troubleshooting

Proper use and maintenance ensure best performance of your Delphi-X Observer. If problems occur this chapter explains how to resolve most issues. Please make sure this chapter is read and checked before contacting your Euromex distributor for service. If a problem is not described in this list or the suggested solution does not bring the result needed, please contact your Euromex distributor

| | No power The bulb is not inserted The bulb is defective e transmitted-reflected illumination switch is in the wrong position The fuse has blown here is no power from the mains connection Poor bulb quality | Check if the power cable is connected well, try other power cable Remove the bulb and place it back Remove the bulb and place it back Change position of the switch Replace the fuse | |
|---------------------------------------|--|---|--|
| The lamp burns out | The bulb is defective e transmitted-reflected illumination switch is in the wrong position The fuse has blown here is no power from the mains connection | Remove the bulb and place it back Remove the bulb and place it back Change position of the switch | |
| The lamp burns out | The bulb is defective e transmitted-reflected illumination switch is in the wrong position The fuse has blown here is no power from the mains connection | Remove the bulb and place it back Change position of the switch | |
| The lamp burns out | e transmitted-reflected illumination switch is in the wrong position The fuse has blown here is no power from the mains connection | Change position of the switch | |
| The lamp burns out | in the wrong position The fuse has blown here is no power from the mains connection | - ' | |
| The lamp burns out | here is no power from the mains connection | Replace the fuse | |
| The lamp burns out | | | |
| | Poor bulb quality | Replace it | |
| Juducilly | 1 ooi balb quality | Use the specified lamp to replace it. If the problem is not solved, contact your reseller | |
| The bulb flickers or | The bulb is at the end of it's life span | Replace the bulb | |
| the brightness is The vertiginous | e bulb is not inserted entirely into the holder | Remove the bulb and replace it | |
| not turn off the | here are other things in front of microscope (within 1 meter) | Clear all object in a 1 meter radius | |
| illumination | The iCare function is turned off | Press and hold the iCare button for 3 seconds to activate the function | |
| OPTICAL SYSTEM | | | |
| Problem | Cause | Action | |
| The edge of the Tield of view is dark | The nosepiece is not in the located position (objective and light path are not coaxial) | Locate the nosepiece properly where it clicks | |
| or the brightness is | The image of the lamp is not centered | Center the lamp | |
| not uniform | The lens (objective, condenser, eyepiece or collector) is dirty | Clean it thoroughly | |
| | irst rotate the eyepieces, if the dust moves: | Clean the eyepieces | |
| in the field of view | Next move the stage with slide, if the dust moves: | Clean the slide or replace the slide | |
| Ne | ext move the condenser up and down, if the dust moves (using 4 or 10x objective): | Clean the top of the condenser | |
| N | ext change objective, if the dirt is no longer visible: | Clean the bottom lens of the objective | |
| | If problem remains: | Clean the collector lens | |
| Image quality is not | There is no cover slip on the specimen | Add cover slip | |
| optimal (resolution | The cover slip is too thick or too thin | Use the standard coverslip (0.17mm) | |
| or contrast) | The specimen is placed upside down | Turn slide around | |
| The | ere is oil on a non-oil lens, this often happens to the 40x objective | Clean the objective | |
| cc | There are stains on the lens (including ondenser, objective, eyepiece and collector) | Clean the optical elements | |
| | No oil is used for the 100x oil objective | Use Euromex immersion oil (PB.5255) | |
| | There are bubbles in the oil | Try to remove the bubbles or create a new slide | |
| | Wrong oil is used | Use Euromex immersion oil (PB.5255) | |
| Th | ne size of the aperture diaphragm is too large | Close the diaphragm | |
| Th | e size of the aperture diaphragm is too small | Open the diaphragm | |

| TI W CH I I I | A 12 |
|---|--|
| | Adjust the position |
| | Swing out the top lens of the condenser |
| , | |
| | Open the diaphragms |
| , | Take out lamp unit and re-install |
| Incorrect position of the light path switching lever | Set to right position |
| The nosepiece is not in the right position | Turn the nosepiece until it "clicks" into position |
| The condenser is not centered correctly | Center the condenser |
| The condenser is placed inclined in it's holder | Install the condenser again and center it |
| The nosepiece is not in the right position | Turn the condenser until it "clicks" into position |
| Diaphragm is not centered | Center diaphragm |
| The condenser is placed inclined in it's holder | Install the condenser again and center it |
| The stage is tilted | Re-install the stage making sure it is levelled |
| The specimen slide is not placed flat on the stage | Replace the slide on the stage |
| | Turn the nosepiece until it "clicks " into position |
| | Try a specimen of known quality and confirm |
| | Release the focus lock, focus and lock again |
| | helease the locas lock, locas and lock again |
| | Re-install the stage making sure it is levelled |
| The stage is not instance correctly | ne mstan the stage making sare to sievenea |
| Interpupillary distance has not been set correctly | Perform interpupillary adjustment |
| Dioptre adjustment has not been done correctly | Perform diopter adjustment |
| | |
| | |
| Internunillary distance has not been set correctly | Perform interpupillary adjustment |
| | Perform diopter adjustment |
| | Adjust brightness using intensity control knob or |
| brightness is not correct | filters |
| Too low intensity set on intensity controller | Increase the intensity by rotating the controller |
| The size of the aperture diaphragm is too small | Adjust again |
| The position of the condenser is too low | Adjust the position |
| Poor bulb quality | Use the specified lamp |
| Diaphragm(s) closed too far | Open the diaphragms |
| Light path selector set in wrong position | Select the 100:0 or 20:80 position |
| The bulb is at the end of it's life span | Replace the bulb |
| The Köhler incident light is not in the centre | Adjust the bolt of Köhler incident light |
| Too high intensity set on intensity controller | Decrease the intensity by rotating the controller |
| The size of the aperture diaphragm is too large | Adjust again |
| The position of the condenser is too high | Adjust the position |
| Too low or too high intensity set on intensity | Increase or decrease the intensity by rotating the |
| controller (Halogen illumination only) | controller, and use the ND filters |
| The bulb is at the end of it's life span | Replace the bulb |
| The cover slip is too thick | Use the standard coverslip (0.17 mm) |
| | |
| The specimen is placed upside down | Turn slide around |
| | The nosepiece is not in the right position The condenser is not centered correctly The condenser is placed inclined in it's holder The nosepiece is not in the right position Diaphragm is not centered The condenser is placed inclined in it's holder The stage is tilted The specimen slide is not placed flat on the stage The nosepiece is not in the right position The specimen slide is not prepared well Focus lock system is secured at the wrong position The stage is not installed correctly Interpupillary distance has not been set correctly Dioptre adjustment has not been done correctly Brightness is not correct Too low intensity set on intensity controller The size of the aperture diaphragm is too small The position of the condenser is too low Poor bulb quality Diaphragm(s) closed too far Light path selector set in wrong position The bulb is at the end of it's life span The Köhler incident light is not in the centre Too high intensity set on intensity controller The size of the aperture diaphragm is too large The position of the condenser is too high Too low or too high intensity set on intensity controller The bulb is at the end of it's life span |

| The objective | The cover slip is too thick | Use the standard coverslip (0.17 mm) | |
|---|--|---|--|
| touches the specimen when the magnification is being changed | Focus lock system is secured at the wrong position | Release the focus lock, focus and lock again | |
| Large focus deviation | An objective is placed incorrectly, not screwed in all the way | Make sure the right objective is used and screw it all the way into the revolving nosepiece | |
| while changing objectives | The tension of the X/Y controls of the stage are set too tight | Adjust tension to proper setting | |
| | Diopter adjustment has not been done correctly | Perform diopter adjustment | |
| The slide does not move, or moves too | The specimen is not placed between the specimen holder | Place specimen between the holder | |
| heavily | The tension of the X/Y controls of the stage are set too tight | Adjust tension to proper setting | |





