



DIAVERT Inverted Microscope



Instructions



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Design and development of the DIAVERT microscope

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1 Technical description

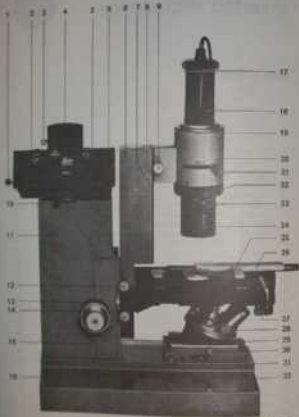


Fig. 1

1. Base plate of the lower column
2. Upper part of the column
3. Packing screw for glass tube
4. Seal screw for the glass tube
5. The tube
6. The tube
7. Seal for the long handle screw (reference No. 2)
8. Seal for the long handle screw (reference No. 2)
9. Packing screw for the glass tube
10. Seal screw for the glass tube
11. Packing screw for the glass tube
12. Packing screw for the glass tube
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32. Packing screw for the glass tube
33. Packing screw for the glass tube
34. Packing screw for the glass tube
35. Packing screw for the glass tube

2 Unpacking the microscope

The following parts are packed in a case (see reference 1):

1. Microscope stand
2. Microscope tube
3. Object stage
4. Microscope handle with upper screw
5. 10X 10X Microscope
6. Lens-cleaning oil
7. Packing material
8. Individual parts such as objectives, eyepieces, condenser, etc.

Assemble the microscope as follows:

1. Assemble the microscope as follows:

2. Assemble the microscope as follows:

3. Assemble the microscope as follows:

4. Assemble the microscope as follows:

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29. Assemble the microscope as follows:

30. Assemble the microscope as follows:

3 Assembling the microscope

3-1 Install the condenser No. 24 for 10x magnification

Align the condenser housing (shown) with the stage clamping guide 12 at the lower end position.

Tighten clamping screw 2.15 (Fig. 2, item 1), insert the sheet-edges into the clamping guide and lower 2 until the condenser sheet matching 2.32 locks into the 2.34. Fix the edge with clamping screw 2.15.

The markings indicate:

- 1 = Long working distance
- 15 = Normal working distance
- 20 = 10.75/10x
- 25 = Vertical displacement with objective + Glass Adjustment length



Fig. 4 Assembling the sheet edge
12 Clamping guide of the condenser housing
15 Condenser housing
16 Cable holder

Tighten clamping screw 2.32, insert the illuminator handle 2.8 in the clamping guide and lower 2 to the stop with the stage bracket. The clamping screw 2.12

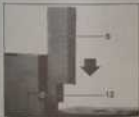


Fig. 7 Assembling the illuminator handle
8 Illuminator handle
12 Clamping guide of the condenser housing

loosen clamping screw 4.6 and insert the later mounting into the clamping guide from above; lower 5 to the figure marking 1 (4.7) and clamp 5 with clamping screw 4.6.

Insert the 63.10x illuminator or lamp housing 30 in the second ring of the lamp mount and lock it by turning 3 clockwise; fix the lamp cable on the cable holder and connect 3 to the stand. Screw the condenser No. 24 (2.36) into the top ring of the lamp mount.



Fig. 6 Assembling the cable holder
2 Cable 1 for the illuminator
3 Clamping guide of the lamp mount



Fig. 5 Assembling the lamp mount



Fig. 8 Assembling the cable housing



Fig. 9 Assembling the illuminator

Push the loading lever 8 to its right, insert the tube vertically and release the loading lever in the position it must be assumed to intake the tube without effort. It is braked by short pressure on the loading lever to the left.

Loosen the clamping screw 6.32 and raise the viewing microscope fitting 6.28. Raise the object stage by means of the coarse adjustment 1.13. Push the viewing microscope 10.26, with objective screwed in position, into the fitting at the top. Fix the objective viewing microscope with set-screw 10.30. Secure the viewing microscope fitting in open and fix it by means of the clamping screw 11.32.

The stroboscopic object guide can be attached to the object stage from 2 sides. Place the stroboscopic object guide on to the object stage in the desired position and push it into the recesses provided there. Fix it with the bracket screws 10.30.

Insert the object holder into the object guide from the front, ensuring that the recessed side of the object holder fits to the rear and the drilled slots on top. The object holder clamps must engage. Connect the object with the transformer to bring all the connecting cables and the transformer to the water.



Fig. 6. Mounting the microscope tube.
10. Loading lever.



Fig. 7. Mounting the viewing microscope fitting.
20. Clamping fitting for the viewing microscope.
32. Clamping screw for the viewing microscope fitting.



Fig. 8. Mounting the objective viewing microscope.
28. Viewing microscope.
30. Clamping screw for viewing microscope.



Fig. 9. Mounting the object stage.
30. Object stage fit into the object guide.



Fig. 10. Fixing in of the viewing microscope fitting.
30. Clamping screw.



Fig. 11. Mounting the object holder.

2.1 *Start with Series 600 brightfield condenser.*

Series 400 phase contrast condenser and background condenser 0.20-0.35 and 0.40-1.00

The object stage and the illuminator holder are inserted according to Fig. 3 and 4. Loosen the clamping screw 14.27 and insert the condenser fitting 14.20 with lock change in the clamping guide from above and insert it to the stop. Tighten clamping screw 14.37. Insert the condenser in the lock change in the stop. The background lens 14.26 points downwards. Loosen clamping screw 18.3, insert the lamp mount in the clamping guide from above, lower it to the figure marking 2 (0.5) and clamp it. Only with condenser top of long interrupt distance 7 otherwise figure 1. The filter (object camera and object guide) are inserted according to directions pp. 6, 7.



Fig. 3 Inserting the condenser
 36 Condenser top of long interrupt distance



Fig. 4 Inserting the condenser fitting
 40 Condenser fitting
 41 Clamping screw of the condenser holder



Fig. 5 Inserting the lamp mount
 40 Clamping screw of the lamp mount
 41 Hole 2 for long interrupt distance

2.2 *Inserting the XV 10 W lamp*

The lamp is changed as follows:
 1) Loosen the clamping screw 17.20 and pull out the lamp mount.
 2) Take lamp 17.20 out quickly new lamp into the socket 17.51 and turn it anticlockwise for clamping.
 3) Pressed into non-rotated lamp can be used.
 4) Insert a pressed lamp so that the end of the lamp comes to be below the red dot of the lamp socket.
 5) Push the non-rotated lamp into the guide grooves of the socket with the two pins and lock it in position by pushing it in to the right.
 Insert the Lamp Housing 15 in the lamp mount and lock it in position.

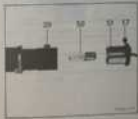


Fig. 6 Inserting the 10 W lamp
 17 Lamp housing
 15 Lamp housing
 17 Lamp
 40 Socket

2.3 *Clamping the 10 W lamp*

Insert the pressing bar in the center of the object stage. Completely turn the aperture stopwheel. Adjust the lamp socket 17.51 with the light axis exactly parallel to the clamping bar. By adjusting the lamp pressing screw 17.17 move the lamp parallel into the center of the clamping bar. After removal of the pressing bar adjust the lamp socket until the red mark points of the objective is exactly illuminated.



Fig. 7 Clamping the 10 W lamp in the microscope

28.1 Inserting the 12.5 MW lamp (lamp lamp)

Remove the shutter cover (19) from the Lamp Housing (2) and remove the lamp from the housing.

Insert the 12.5 MW lamp with protective cap on the shutter and remove adjustment screw.

Insert the bracket on the holder and adjust the angle. Tighten the bracket screw.



Fig. 28.1 Inserting the 12.5 MW lamp into the Lamp Housing (2).

- 19 Shutter cover
- 2 Lamp housing
- 42 12.5 MW lamp
- 43 Adjustment screw



Fig. 28.2 Adjusting the Lamp Housing (2) on the microscope.

28.2 Centering the 12.5 MW lamp (lamp lamp in the Lamp Housing (2))

After each change of lamp, the lamp/lamp housing must be recentered.

Completely open the aperture diaphragm, insert the centering disk at the stage aperture.

a) Focus the mirror image of the lamp filament by adjusting the reflector (25.4). Open an image and a mirror image of the lamp filament on the tanking disk by closing ring (26.4).

b) Move the image into central position of the filament into the center by adjusting screw (27.4).

c) Turn screw (28.4) for the vertical adjustment of the tube into the stage and mirror image on the filament over the stage the edge.

Remove the centering disk and remove it by the aperture.

Rotate the light microscope horizontally using observing the two focal planes of the objective, parallel of the filament from the top view with the two focal planes clearly distinguished.



Fig. 28.3 Lamp Housing (2).

- 25 Light reflector adjustment
- 26 Locking screw for the filament
- 27 Reflector
- 28 Vertical screw for the filament adjustment of the lamp
- 42 Stage aperture
- 43 Filament screw for the vertical adjustment of the lamp



Fig. 28.4 Centering the lamp/lamp (2) on the microscope.

4 Technical hints

4.1 Binocular tube 2

The tube can be set to the individual interpupillary distance of the observer. This requires a corresponding correction of the tube length, which is carried out on the eyepiece tubes as follows: set the interpupillary distance by pushing or pulling with both hands so that the 2 barrels coincide in the microscope. But only a single circular image is seen. Bend the microscope tube on the side engraved in the front plate of the tube and transfer it to the left eyepiece tube.



Fig. 10. Binocular tube 2

4.2 Binocular tube FSA

The binocular tube FSA has a ringed beam-splitting prism, which splits up the light intensity at a ratio of 50:50 (40% for photography, 10% for visual observation) or directs the entire light flux into the eyepiece tubes for visual observation. The interpupillary distance is set on the tube with both hands by simply pulling or pushing.

If the interpupillary distance is not known, the tube is adjusted during binocular observation until only a single, circular and evenly conveyed field of view appears. Correction of visual defects must be carried out with the aid of the focusing systems of one of the two PERPLAN objectives.

4.3 Objective matching microscope

The matching microscope has 2 numbered threads for the objectives. An objective/eyepiece pair always carries every such 1/2 indicator, which can be used to transfer the threads to which the individual objectives are matched.

4.4 2 knob operation

The fine adjustment is operative for about 2 turns, and then activates the mechanism of the coarse adjustment when the rotating direction is reversed. Fine adjustment will automatically be re-engaged. One short rotation of the fine adjustment corresponds to about 2 μ m.

Microscope, 100x - normal phase contrast, 10x and 40x - phase contrast. Objectives without object of observation are withdrawn. Objective for phase contrast is engaged. "Phase" immersion objectives from the word "oil" and a brass ring engraved on the mount.

The letter 1 indicates ring setting distance.

The following approximate data can be used with the microscope for 30 objective (30):

Setting	Obj. No. 30	Obj. No. 40	Obj. No. 60	Obj. No. 100
Ax.	1.650	1.650	1.650	1.650
F ₁	17.5	17.5	17.5	17.5
F ₂	17.5	17.5	17.5	17.5
Working distance	17.5	17.5	17.5	17.5
Resolution	0.25	0.25	0.25	0.25
Depth of field	1.7	1.7	1.7	1.7
Field of view	17.5	17.5	17.5	17.5
Image diameter	17.5	17.5	17.5	17.5
Image height	17.5	17.5	17.5	17.5
Image area	17.5	17.5	17.5	17.5

4.5 Objectives

The data engraved on the microscope objectives apply to the LEITZ objective indicator:

175

distance in mm from the flange of the objective to the top of the tube (mechanical tube length).

837

is the coverglass thickness to be used in mm. Instead of the figure 0.17 a steel foil spacer, which means 0.17, will free objectives specimens can be observed with or without coverglass.

Because the tube length and coverglass thickness indicated in the microscope data (dimensional data of microscope image) and the numerical aperture of the objective 100/30 are engraved in abbreviated form.

In addition the show of correction is also given. It = Selenia system. Age =

For work with biological and other weak preparations and normally with a bottom of 2.75 mm thickness, an LEITZ specialized light objective light up to an immersion can be used.

Now the use of a Series 800 objective condenser is recommended for phase contrast investigations. A Series 800 condenser is used. Ensure that the top plate of the lens is identical to the 800 or 802 Series condenser.

4.1 Eyepiece

Dimensions of eyepieces under 15mm can be found in the DataSheet table. The diameter of the field of view is dictated by the observed magnification.

Eyepiece, 14mm eye-ring (dia. 23.2mm)	
Magnification	Field of view
2.2x	16
2.2 x M	18
5x	19
10 x*	14
20 x	9
High-power eyepieces	
4.2	16
11 x	11
15 x M	11
15 x MP 1/2	16
Aperture	
Diameter = 100 microns	

* Eyepieces with 14mm eye-ring will require a 100 microns objective.
 ** Eyepieces with 14mm eye-ring will require a 100 microns objective.

the diameter of the object area to be surveyed is obtained.

Field of view (mm)
 Objective magnification x tube factor

Standard eyepieces P225P, A4, M, MP (dia. 23.2mm)	
Magnification	Field of view
MP 15 x	18
MP 15 x M	18
OP 15 x	18
OP 15 x M	18
OP 15.5 x	18
OP 15.5 x M	19
OP 12.5 x MP 1/2	18
OP 16 x	15
OP 25 x	10
OP 25 x M	11
Aperture	
Diameter = 100 microns	

* with parallel illumination as in eyepieces series with 100 microns field of view (checklist series)

Standard eyepieces (dia. 23.2mm)

Magnification	Description
4.2 x	Primer eyepiece
5.5 x	Double primer eyepiece
MP 15 x	Eyepiece with field pointer (engraved in yellow)
MP 15 x*	Eyepiece with field pointer (engraved in yellow)
MP 15 x*	Eyepiece with field pointer (engraved in yellow)
MP 15 x	Comparison eyepiece
OP 15 x	Eyepiece for photomicrography (red dot engraved)
15.5 x	Series microscope eyepiece
15.5 x	Series microscope eyepiece for monochromatic light

4.7 Condensers on the DIASERT

When the condenser No. 51 of long working distance is used in simple method of illumination is required, then the condenser is arranged so that with optimum lens combination even the resolution of the microscope image is ensured. The iris diaphragm of the holder serves as aperture diaphragm. Depth of field (area receiving power) and contrast of the microscope image can be changed with it. In most cases best results are obtained if the field aperture and objective aperture have a ratio of 2.5, i.e. if about 2/3 of the diameter of the rear focal plane of the objective is illuminated. This can easily be checked when the eyepiece is removed from the eyepiece tube. More than one stopping diaphragm may result in noticeable diffraction effects and adversely affect the quality of the image.

With the system condenser (Series 800 and 800 K) the method of illumination is used, here the iris diaphragm built into the lamp mount serves as the field diaphragm. When the condenser is correctly set (see p. 26) a sharp image of the diaphragm is formed in the object plane. Related to the size of the microscope image field, the field diaphragm limits the ray cone in the object. Only the cone of rays required for the formation of the image will reach the objective. This permits a degradation of the image owing to flare, loss of contrast and disturbing reflections. At the same time unnecessary heating of the specimen is prevented.

The optometric system condensers of Series 900 consist of a standard Series No. 50 with condenser lens which is the condenser for low powers. The

aperture diaphragm for the complete condenser and microscope system can be set at various positions, when high definition and contrast. The condenser lens can be swung out to the beam. The condensers are interchangeable in a horizontal standard guide and can be vertically adjusted by means of a lock and pinion.

Biophysical condenser system 800

Model	Aperture	Numerical Aperture	Field of View (mm)	Use
800	Variable	0.10	100	For use with the microscope for low magnification
801	Variable	0.18	50	For use with the microscope for low magnification
802	Variable	0.30	30	For use with the microscope for low magnification
803	Variable	0.45	20	For use with the microscope for low magnification
804	Variable	0.60	15	For use with the microscope for low magnification
805	Variable	0.75	10	For use with the microscope for low magnification
806	Variable	0.90	7	For use with the microscope for low magnification
807	Variable	1.00	5	For use with the microscope for low magnification
808	Variable	1.10	4	For use with the microscope for low magnification
809	Variable	1.20	3	For use with the microscope for low magnification
810	Variable	1.30	2	For use with the microscope for low magnification
811	Variable	1.40	1.5	For use with the microscope for low magnification
812	Variable	1.50	1	For use with the microscope for low magnification
813	Variable	1.60	0.8	For use with the microscope for low magnification
814	Variable	1.70	0.6	For use with the microscope for low magnification
815	Variable	1.80	0.5	For use with the microscope for low magnification
816	Variable	1.90	0.4	For use with the microscope for low magnification
817	Variable	2.00	0.3	For use with the microscope for low magnification

Phase contrast condenser system 800 according to Zeiss

Model	Aperture	Numerical Aperture	Field of View (mm)	Use
820	Variable	0.10	100	For use with the microscope for low magnification
821	Variable	0.18	50	For use with the microscope for low magnification
822	Variable	0.30	30	For use with the microscope for low magnification
823	Variable	0.45	20	For use with the microscope for low magnification
824	Variable	0.60	15	For use with the microscope for low magnification
825	Variable	0.75	10	For use with the microscope for low magnification
826	Variable	0.90	7	For use with the microscope for low magnification
827	Variable	1.00	5	For use with the microscope for low magnification
828	Variable	1.10	4	For use with the microscope for low magnification
829	Variable	1.20	3	For use with the microscope for low magnification
830	Variable	1.30	2	For use with the microscope for low magnification
831	Variable	1.40	1.5	For use with the microscope for low magnification
832	Variable	1.50	1	For use with the microscope for low magnification
833	Variable	1.60	0.8	For use with the microscope for low magnification
834	Variable	1.70	0.6	For use with the microscope for low magnification
835	Variable	1.80	0.5	For use with the microscope for low magnification
836	Variable	1.90	0.4	For use with the microscope for low magnification
837	Variable	2.00	0.3	For use with the microscope for low magnification

5. Operation of the microscope

Since the tube of the DIAVENT can be easily rotated on the stand, the microscope can be operated very precisely and conveniently in the lateral position (see Fig. 24).

The left hand grips the rear viewing tube to the stand in the adjustment. The right hand rotates the stand 90°, in the specimen slowly, which sets the cover-glass.



Fig. 24. Lateral operation of the microscope.

5.1 Observation with the condenser No. 21

Place object to be investigated on 2-Petl slide directly on the glass stage or in the object guide. Choose the objective of lowest power for first observation. The base of the aperture diaphragm is on the right-hand side (left on the illumination). Ensure that the lamp holder is in position 1. Focus the specimen by means of the coarse and fine adjustment.

Corrected and carried out as follows. Look through the eyepiece with both eyes open with one eye (e.g. right-hand eyepiece/light eye). Focus the condenser with the fine adjustment. Now look through the eyepiece with following eyepiece with the left eye.

Adjust the system until the same object is in the object and eyepiece sharp. Do not interfere with the fine adjustment.

Take care of the correct illumination for the specimen.

First check correction of the lamp cone (p. 5, diagram 2.2).

Remove the eyepiece from the eyepiece tube, release the clamping screw, and move the lamp mount with Lamp Housing 20 (the lamp condenser) until the rear focal plane (nucleus aperture of the objective) is evenly illuminated.

5.2 Observation with the Series 400 wing-and-condenser

Place the object to be investigated (e.g. Petl slide) either directly on the glass stage or into the object guide. Focus the specimen by means of the coarse and fine adjustment. Carry out correction for visual defects (see para. 5.1).

Ensure that the lamp mount is in position 1. For the condenser of long wavelength (distance set the lamp housing in position 2.

1. Close the front diaphragm completely to the lamp mount.

2. Lower the condenser by means of the vertical adjustment until a sharp image of the back diaphragm is formed in the plane of the specimen.

3. Centre the image of the back diaphragm with the fine centring screws.

4. Tilt the front diaphragm so that it just touches the edge of the field of view.

Check lamp correction and illumination as under para. 2.2 a and 2.2 (nucleus aperture).



Fig. 20. Observing the back diaphragm

5.1 Investigation of phase contrast according to Zeiss

Two methods are available for phase contrast investigation:

1. Phase contrast with wide condenser No. 21 for ring working distances (magnification range 10 x to 200 x)
2. Phase contrast with special Series 400 condensers for normal working distances (magnification range 62.5 to 1000 x)

5.2 Phase contrast with condenser No. 21

For this device the lighting system WOT is setting focus is already built into the ring mount. In addition is the phase contrast objective.

Phase C 100/25

Phase C 200/50

Phase C 300/80

A light ring diaphragm (Phase I) and a focusing telescope are required.



Fig. 20 Light ring diaphragm

Setting the phase contrast with

Some of the Phase objectives into the working microscope. Insert the illumination the changing guide and clamp it in the working position. Focus the specimen (objective 100/25), insert the light ring diaphragm in the well marked "Phase" in the stop. Completely open the eyepiece diaphragm.

Remove any eyepiece from the eyepiece tube and insert the focusing telescope. Remove the ocular screw on the focusing telescope and adjust the upper part until both the light and the phase ring are exactly in focus.

Check whether both rings are concentric and superimposed. If not, introduce this position by means of the two turning screws.

The color of the image formed of the light ring is changed in phase contrast observation of objects in liquids when the level of the liquid varies. The necessary compensation can be carried out by adjustment of the illuminator tube (uniform superimposition of light and phase ring).

Replace the focusing telescope with the eyepiece. If necessary repeat correction after magnification change.

A further stack of the image formation of the ring is no longer necessary, since the objective, once set, is permanent.

For a rapid change-over between phase contrast and brightfield illumination, merely remove or insert the light ring diaphragm in its holder.

5.3 Phase contrast with condenser with Series 400 condenser

In addition to the phase contrast objective, a Series 400 phase contrast condenser, a focusing telescope, and two setting keys are required.

Setting the phase contrast image

Insert the Phase objectives into the working microscope, insert the microscope into the changing guide and clamp it in the working position. Bring the Phase 100/25 objective into the mount and set phase ring 1 in the Phase condenser.

Focus the specimen.

Close the field diaphragm.

Lower the condenser to focus of its vertical adjustment, until a sharp image of the field diaphragm is formed in the field of view. Center the image of the diaphragm with the two condenser centering screws.

Open the field diaphragm so that its outer circumference is beyond the edge of the field of view.

Insert the focusing telescope in one of the eyepiece tubes.

Adjust the focused scale on the focusing microscope and adjust its top until a sharp image of both the light and the phase rings is seen.

When the rings are off center, insert the setting keys into the two rear apertures of the condenser and rotate the light and phase rings clockwise by turning the setting keys.

Take out the setting keys and check the type of extinction for all other objectives.

No further check of the image formation of the ring is now necessary, when the object, once focused, set is permanent. For further details see instructions Phase Contrast Illumination according to Zeiss, No. 973 84.



Fig. 21 Series 400 phase contrast illumination

54 ULTRAFLEX halogen-light illuminator

Mount the lamp housing (2) in the hole of the stand and clamp it with screw (3).

Push the ULTRAFLEX with camera plate into the sliding guide and lock it in the working position, insert the illuminator.

Open the Lamp Housing (2) in the forward position and remove it by turning it. Connect the Lamp Housing with the main via the 120 volt transformer. Further directions are contained in the instructions, ULTRAFLEX No. 135-90.

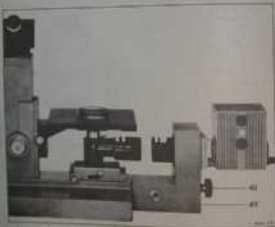


Fig. 54. ULTRAFLEX with ULTRAFLEX halogen light illuminator.

55 Background illumination

For the background illumination of objects requiring a long working distance, ring diagrams of the Photo-Studio in conjunction with the condenser No. 91 are used for low-power work.

For this purpose the sample stage is pushed into the slot marked Photo of the lamp holder.

For large objects such as culture vessels a special holder with a 20° 25W Photo-Studio is available.



Fig. 55. ULTRAFLEX with special ring holder for large objects.

General Note

It is essential to observe the following points for a good photomicrograph:

1. The precise setting of the illuminator (para 3.3)
2. Critical focusing of the image.
3. Accurate determination of the exposure time.
4. Sufficient cleanliness of all the optical faces, e.g. deflecting mirror, shut glass, condenser, objective, tube lens, eyepiece etc., accessible to the user.



Fig. 10. Microscope for the electron.

Special attention must also be paid to the question of useful magnification, the choice of a right filter for the correct spectral rendering, ion beam- and white light of the specimen, the setting of the correct colour temperature of the irradiation lamp (with colour photography) and the choice of suitable exposure materials for photomicrography.

The following attachments are available for photomicrography with the ZEISSORION OPTACONARF 10.5 µm electron microscope camera.
 Micro-attachment for the ZEISSORION with electron beam
 LEITZ System Camera
 LEITZ COMBINATORSM attachment
 system camera
 Please consult the manufacturer about these attachments from their catalogue or through our office.



Fig. 11. ZEISSORION with System Camera and LEITZ MIC.



Fig. 12. ZEISSORION with COMBINATORSM attachment and system camera.

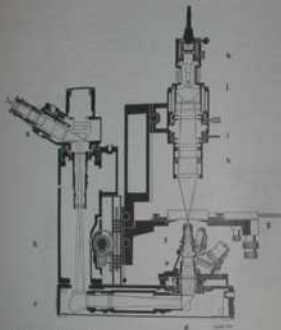


Fig. 10. Main parts of the microscope and
numbering for it.

- A. Revolving nose-piece
- B. Objective lens
- C. Eyepiece lens
- D. Eyepiece tube
- E. Eyepiece
- F. Eyepiece ring
- G. Eyepiece
- H. Eyepiece
- I. Eyepiece
- J. Eyepiece
- K. Eyepiece
- L. Eyepiece
- M. Eyepiece
- N. Eyepiece
- O. Eyepiece
- P. Eyepiece
- Q. Eyepiece

6 Care and maintenance

For protection against dust the microscope should always be covered with its flexible hood after use. Prior to use the slide should be cleaned with a piece of linen or chemical paper. Methylated spirit must be used to clean the slide for that purpose since it attacks the varnish. Petrol, on the other hand, is generally suitable for the cleaning of varnished parts.

Light patches on the slide stage caused by petrol can be removed by treatment with liquid paraffin or acetone spirit.

Special caution is necessary during re-assembly, avoiding the use of sand, lumps of ivory and similar porous materials. Direct contact between optical components and metal and brass elements must be avoided at all costs and all parts should be cleaned thoroughly and immediately after use.

The optical parts of the microscope must be kept continuously clean. Dust on glass surfaces is removed with a fine, dry white brush, lightly blown across the surface as you apply the brush.

Bad contamination requires cleaning with a soft lens or camera tissue and with water or alcohol as solvent.

For cleaning, attention must not be directed to any internal part of the microscope that is exposed to the light.

Special care is necessary during the cleaning of anti-reflection coated surfaces. The external surfaces of the lens tubes and the front lens of the slide stage are covered with coats of dried glass particles. They must be cleaned with as much care as finished glass surfaces. The parts covered with coats of aluminium and silver, however, may only be cleaned with care. They must be very gently blown off it is necessary to be careful to clean internal surfaces of eyepieces.

Correct treatment prevents the perforation of a slide microscope by dust. If a damaged instrument becomes necessary, the dust spirit or oil for lens will be able to help.

