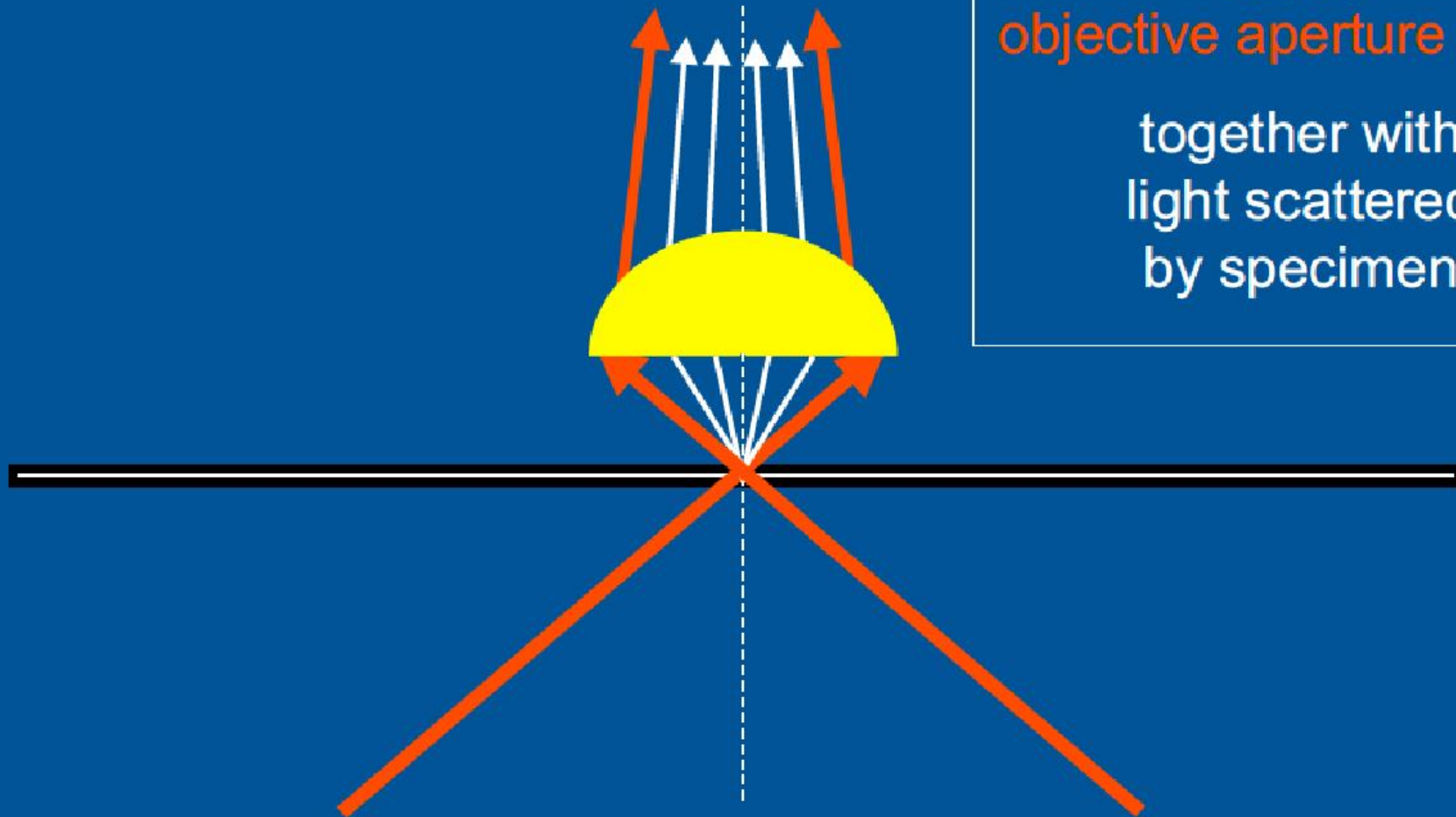


Transmitted-light
Bright field

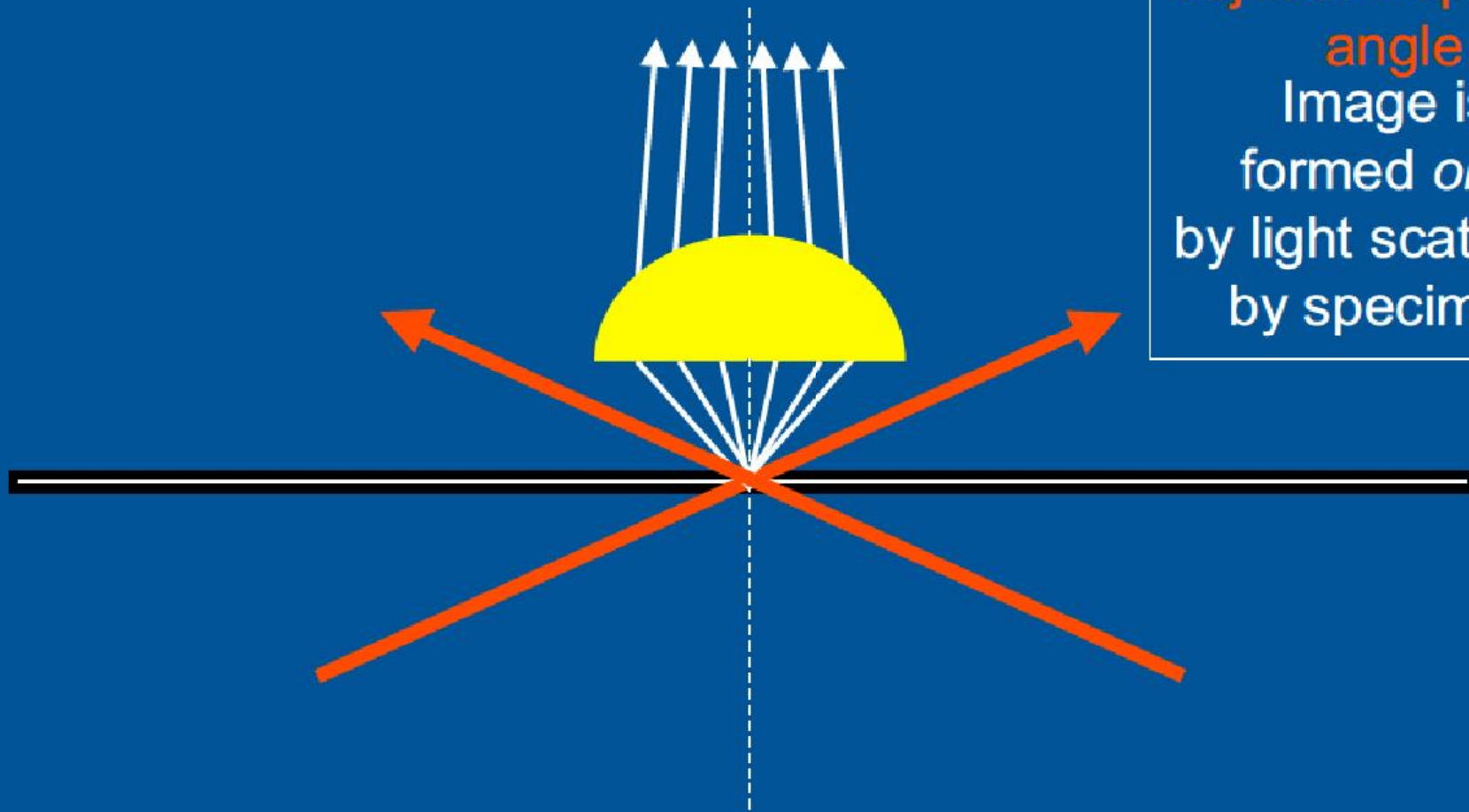
Image formed from
illumination which enters
within
objective aperture angle

together with
light scattered
by specimen



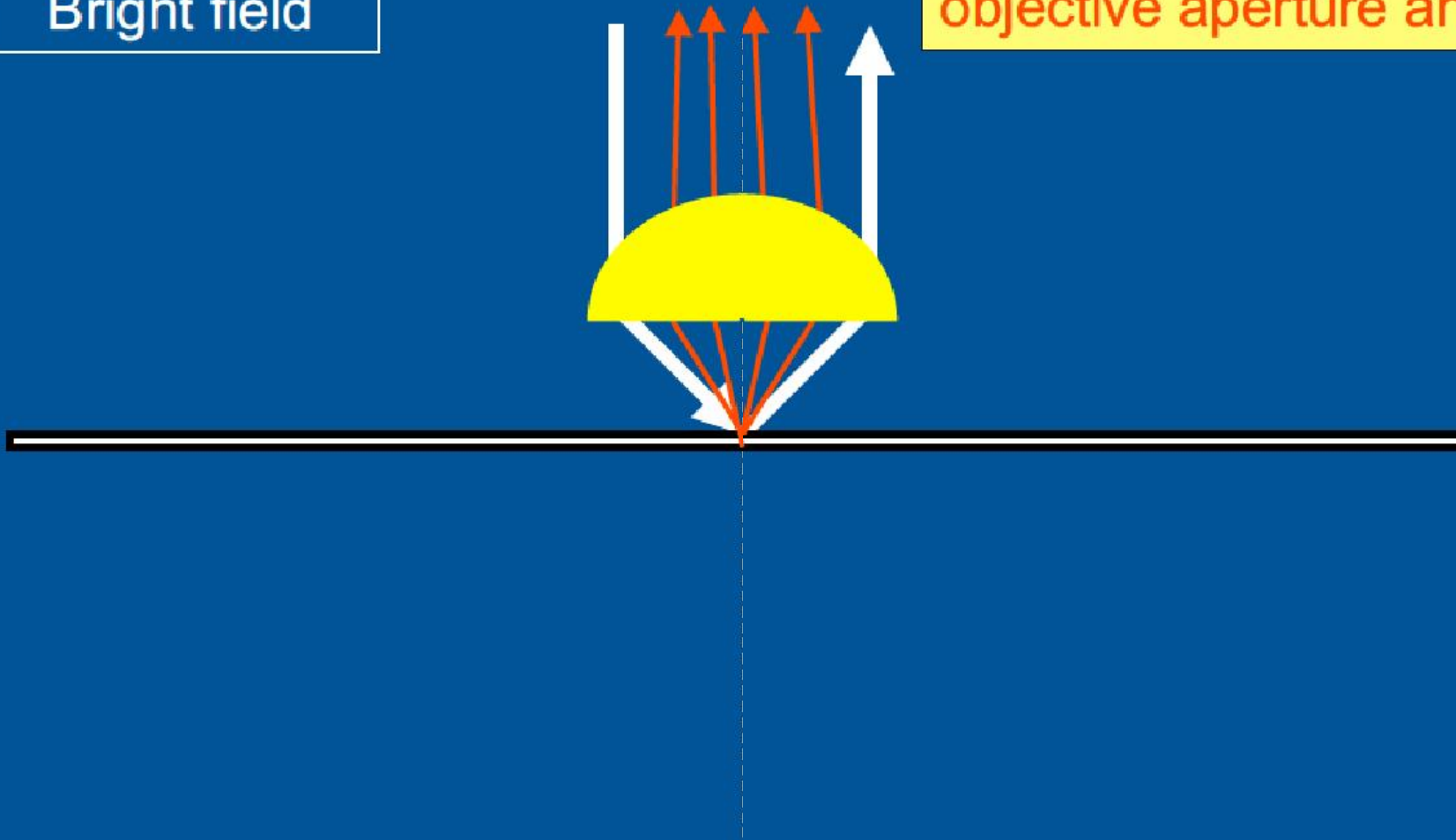
Transmitted-light
Dark field

Illumination falls
on specimen
outside
objective aperture
angle
Image is
formed *only*
by light scattered
by specimen



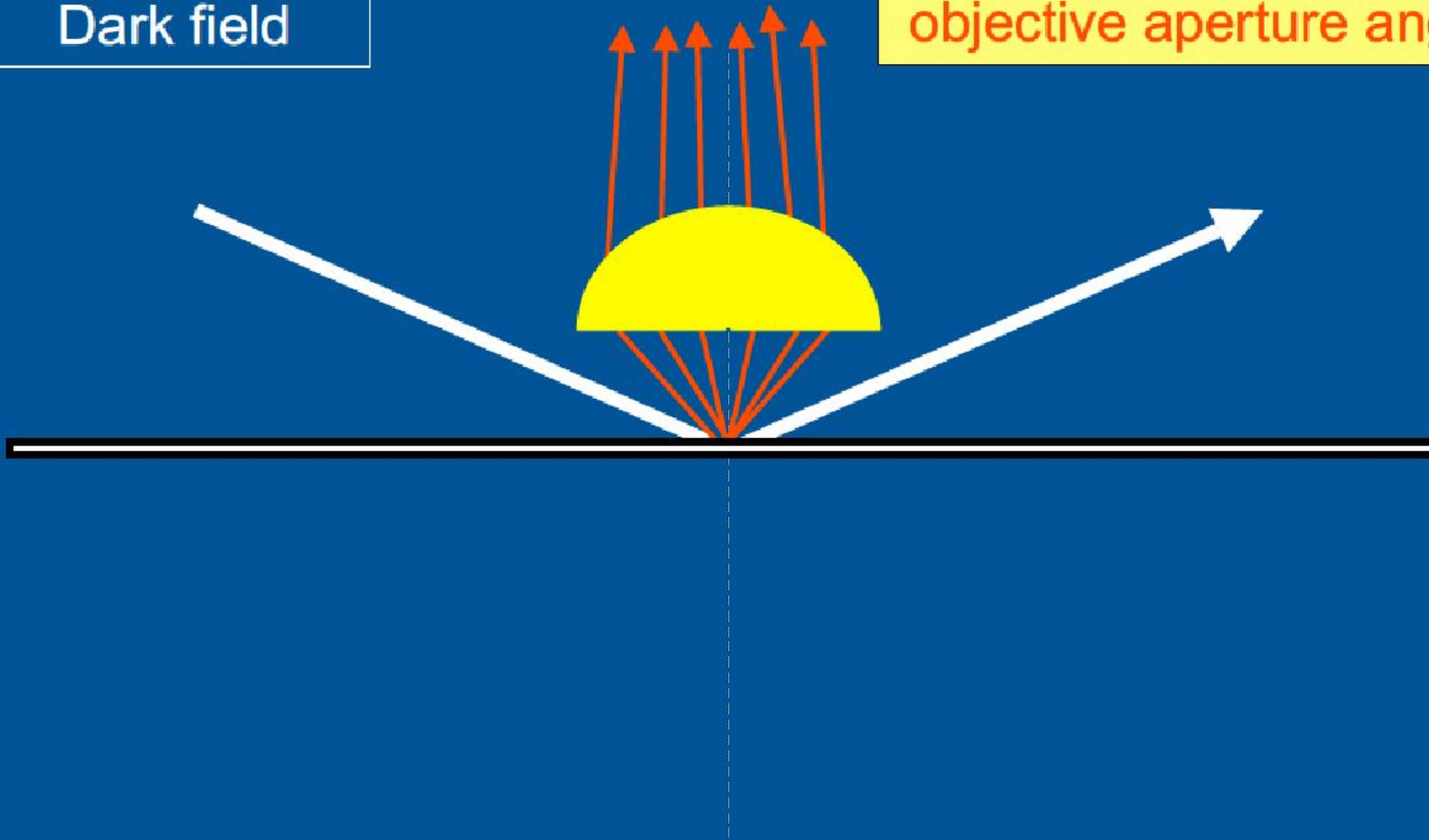
Reflected-light
Bright field

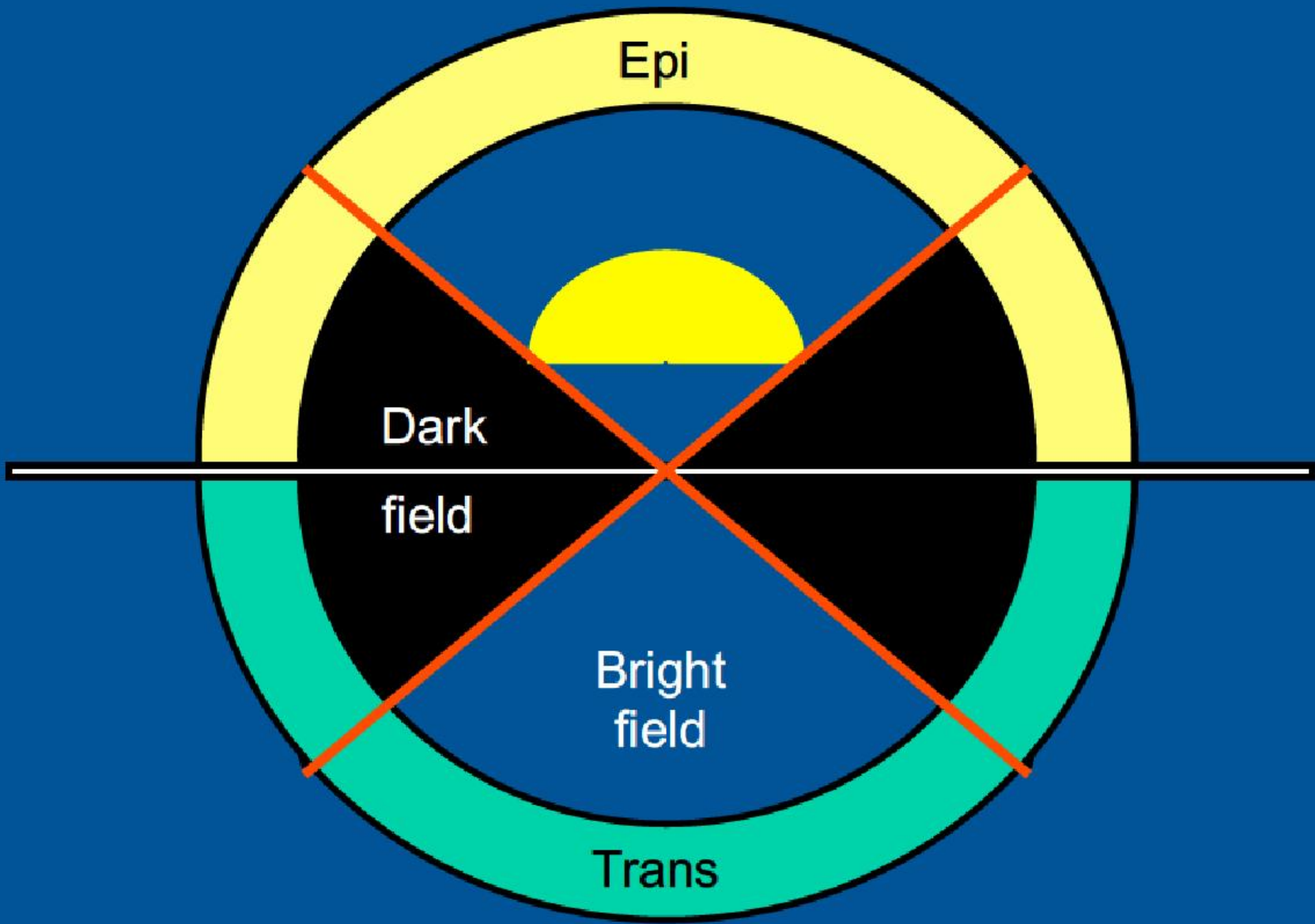
Bright field:
Illumination from *within*
objective aperture angle



Reflected-light
Dark field

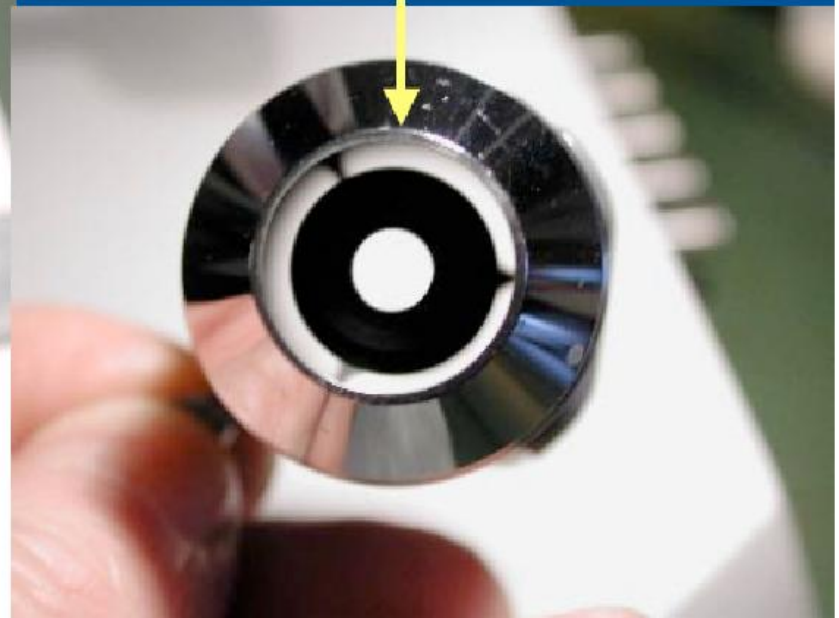
Dark field:
Illumination from *outside*
objective aperture angle

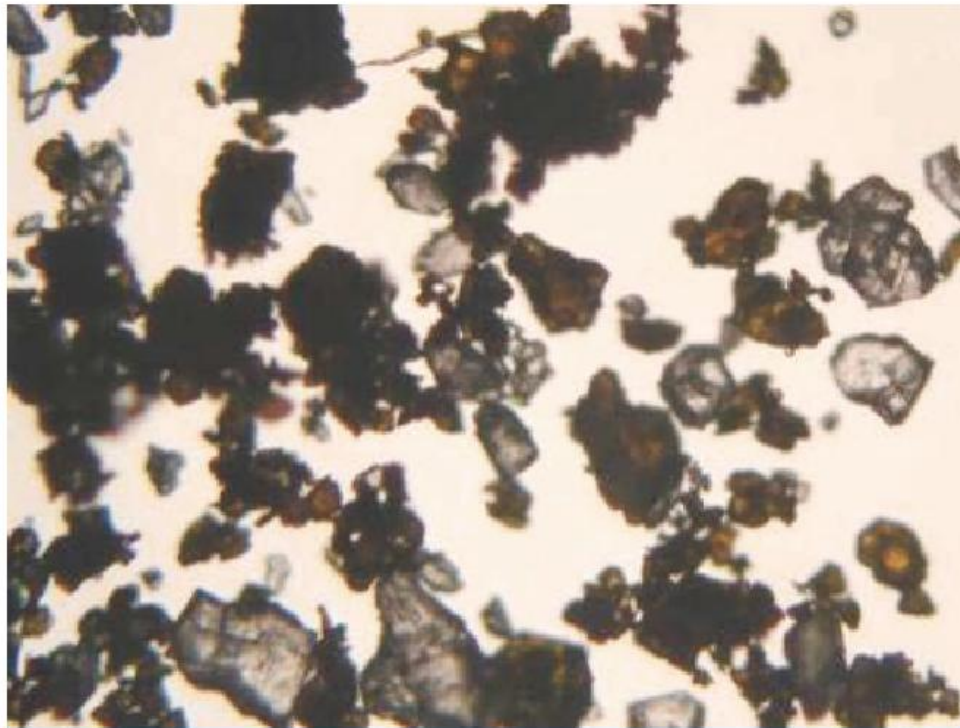




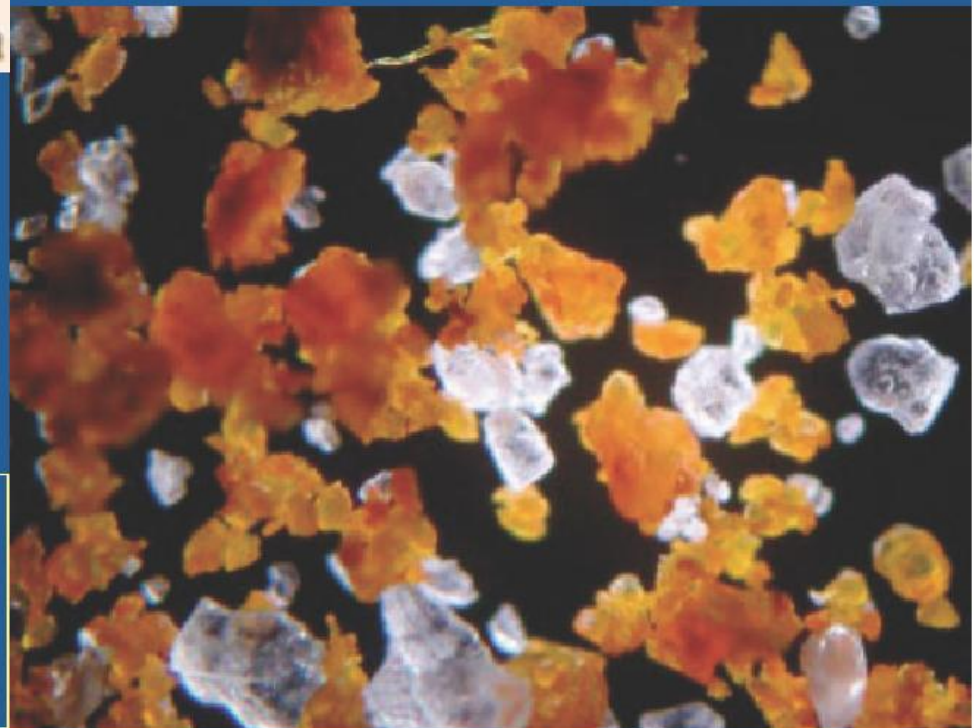
Normally-functioning objective lens

Surround with reflecting end delivering light at oblique angle on to surface of specimen





Transmitted-light
Bright field



Reflected-light
Dark field

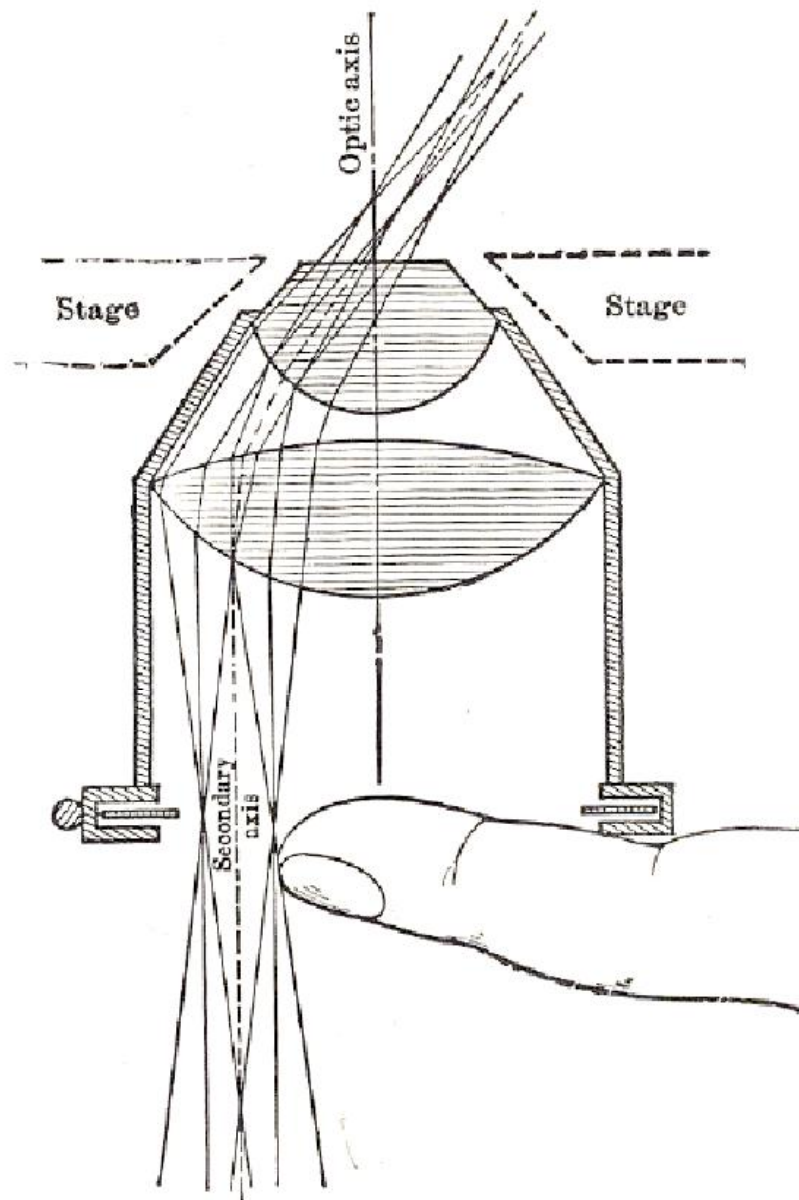
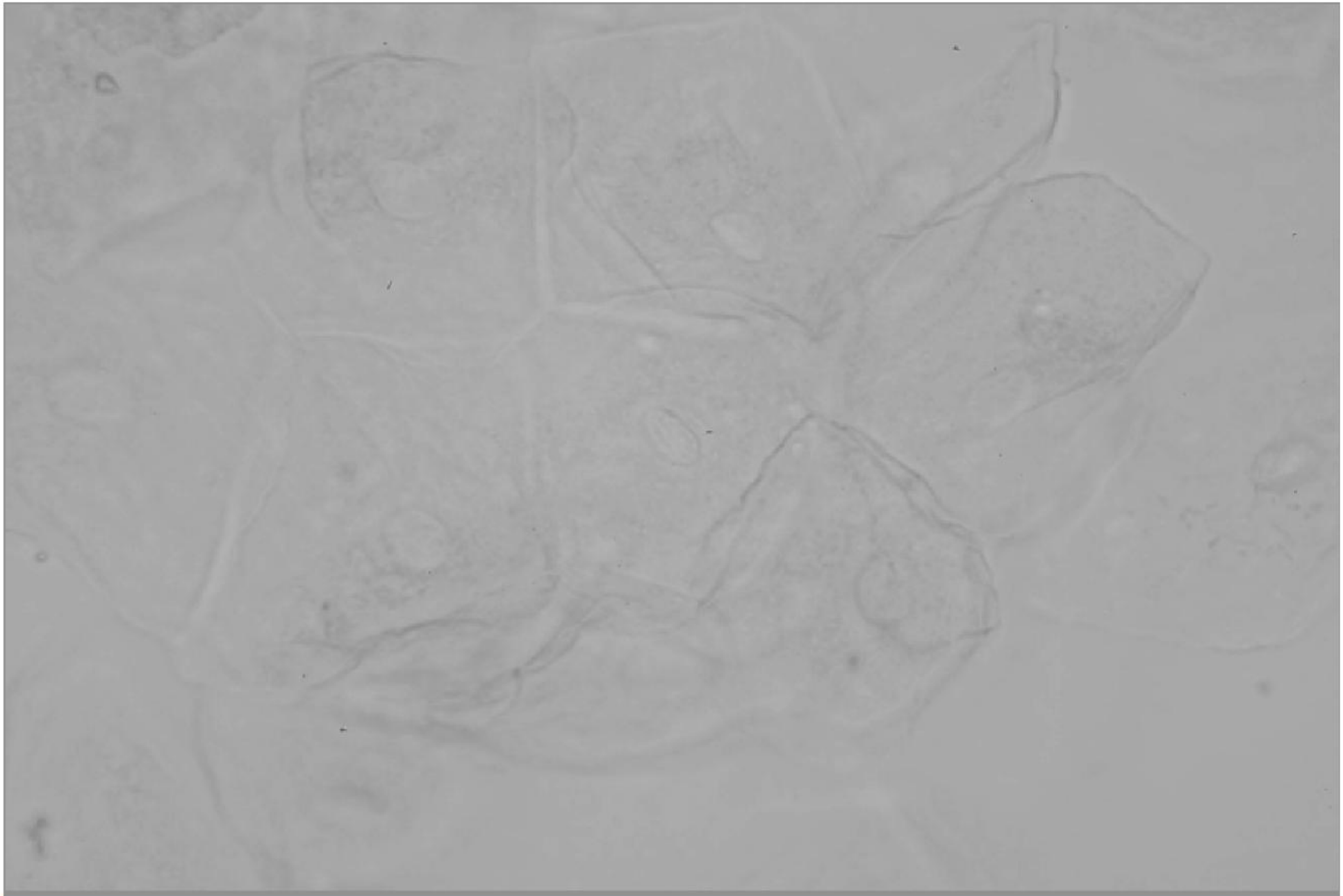


FIG. 62. OBLIQUE LIGHT WITH A CONDENSER.
(From Chamot).

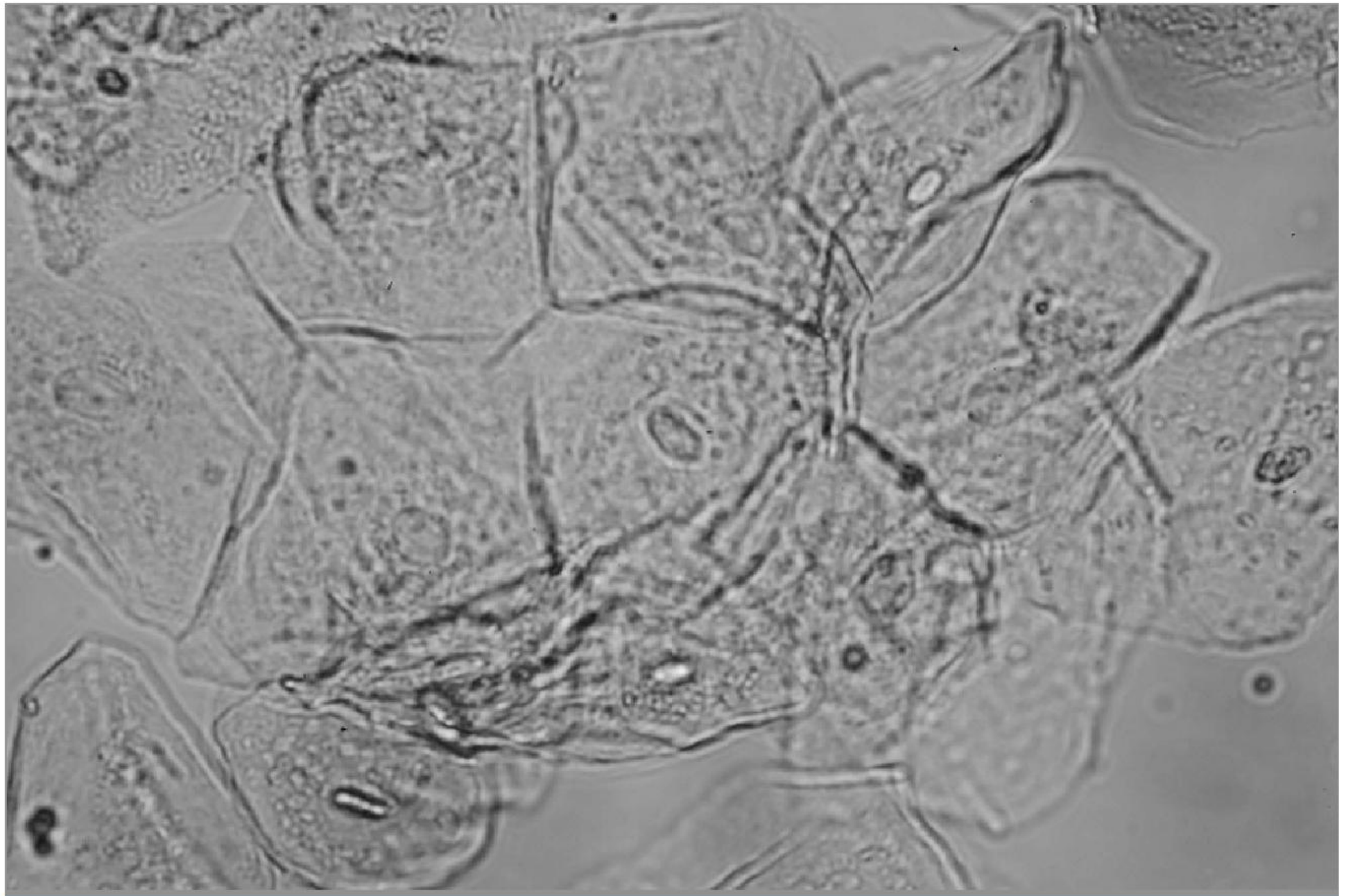
‘Digital’
oblique illumination

- ie using your finger!

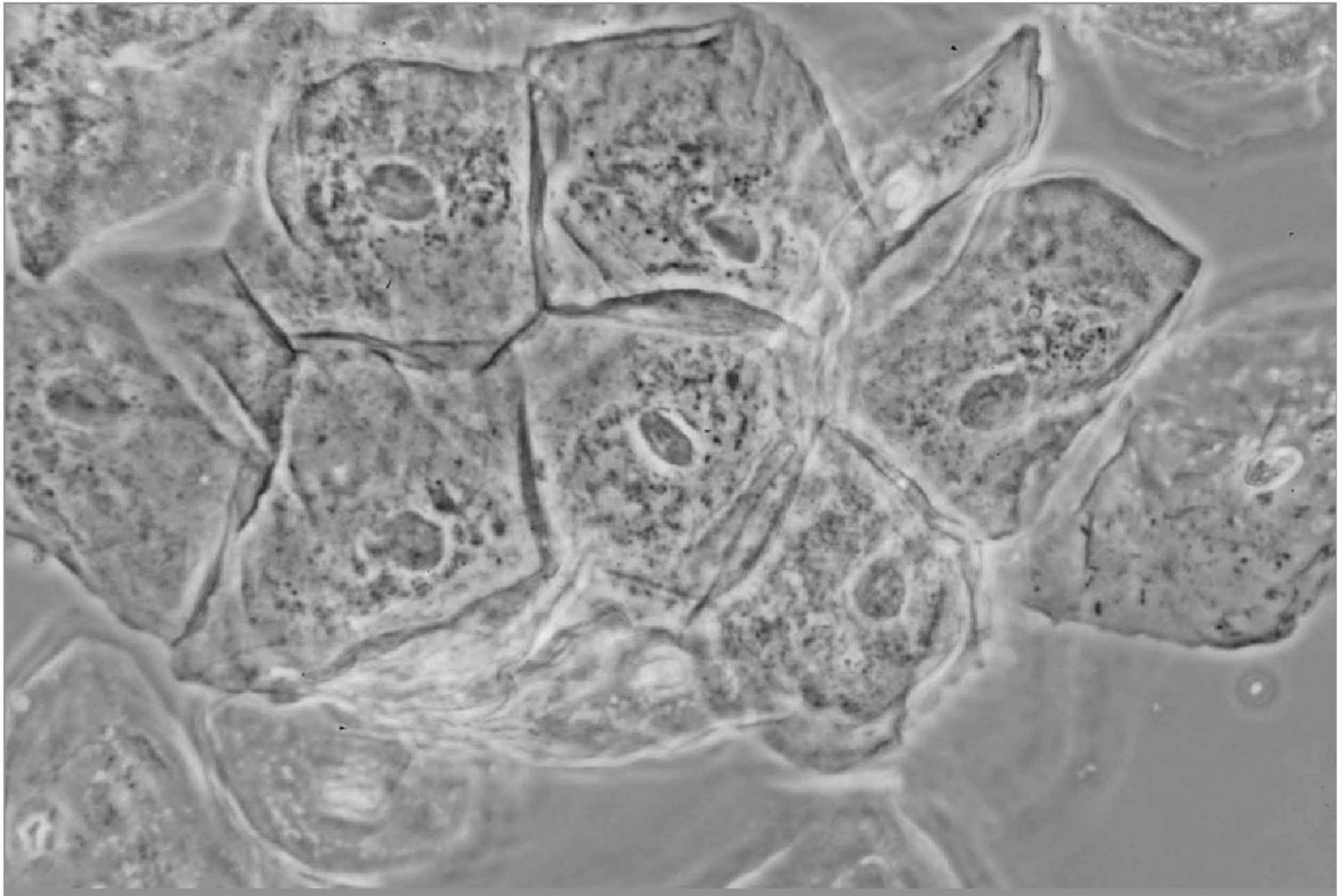
A ‘no-cost’ option
for most microscopes



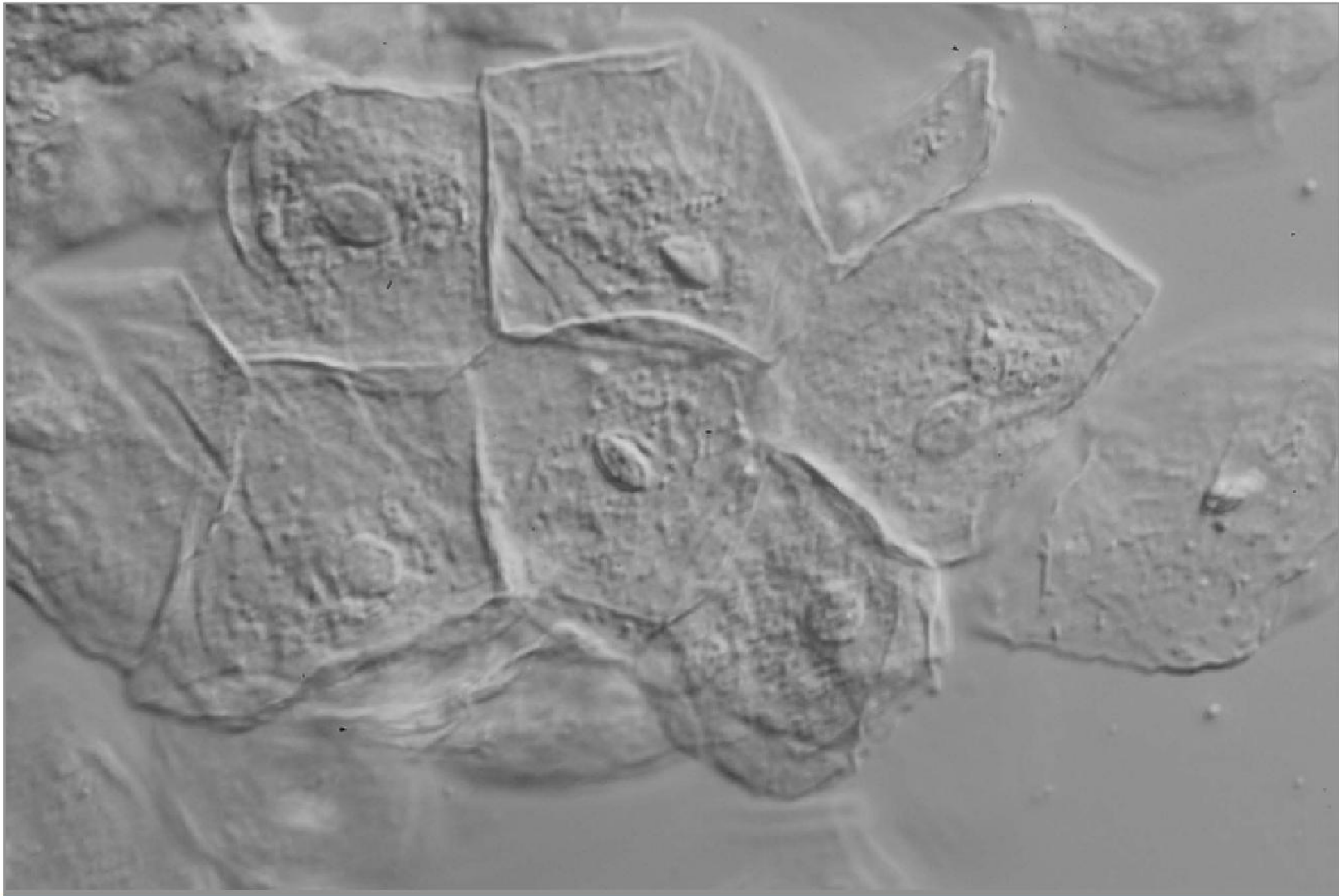
Bright field, full illuminating aperture



Bright field, small illuminating aperture

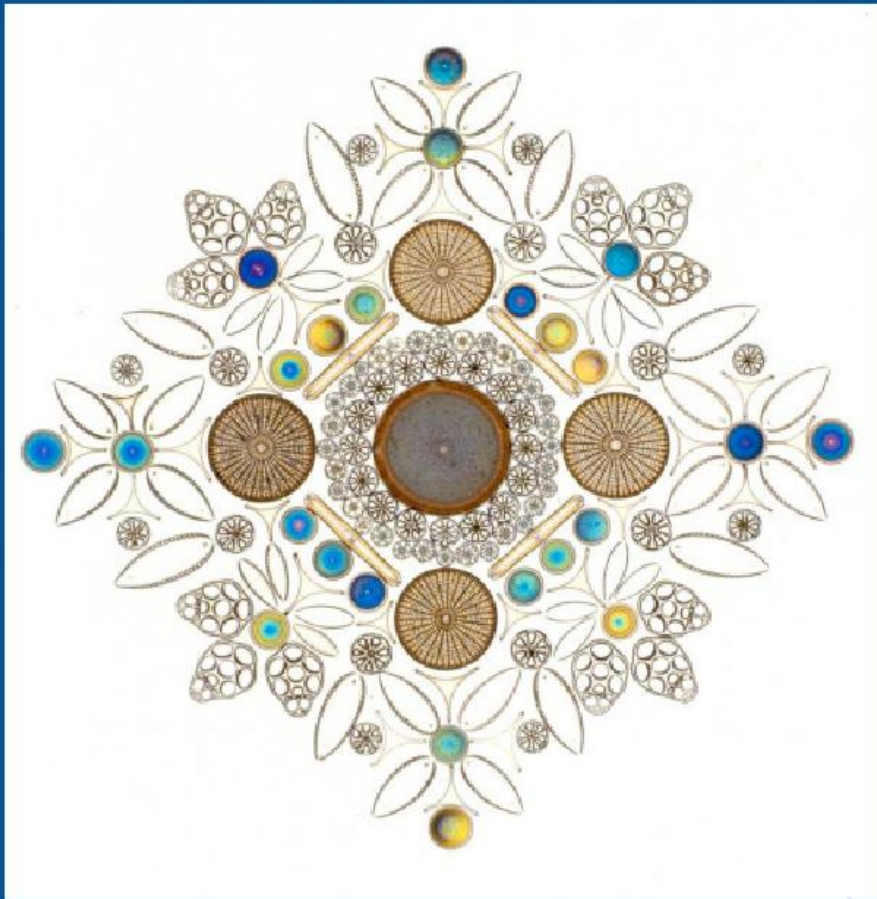


Phase contrast

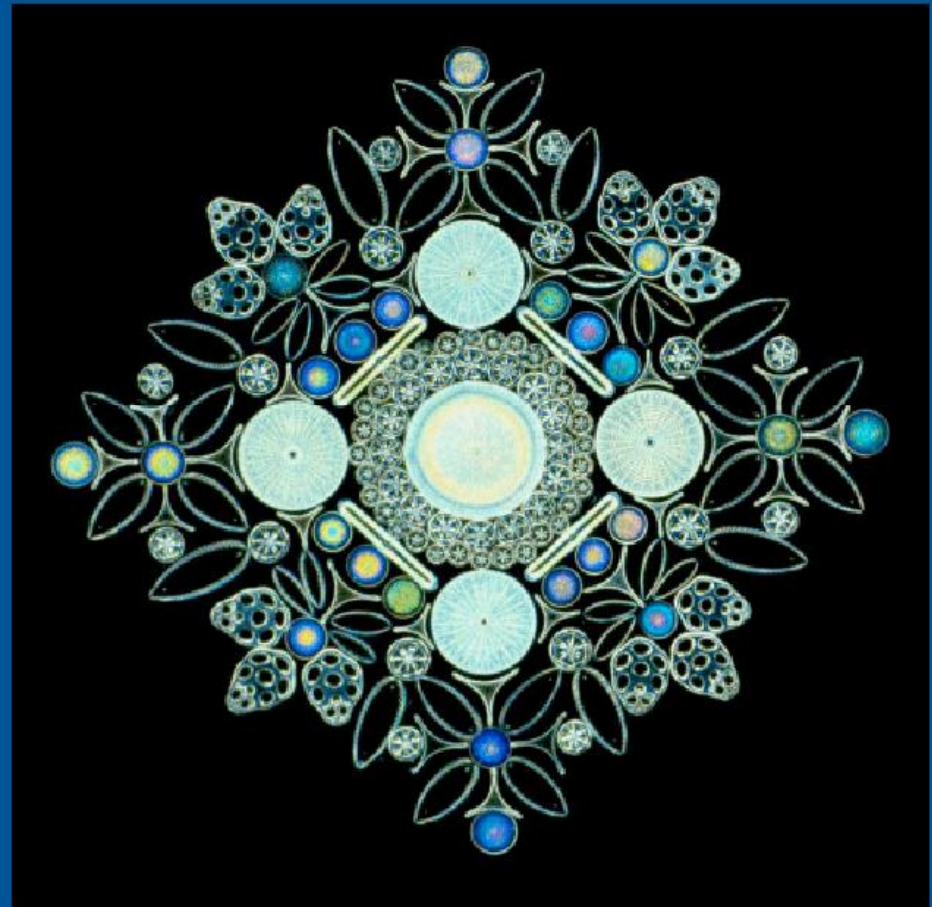


Differential interference contrast

Diatom arrangement: transmitted-light



Bright field

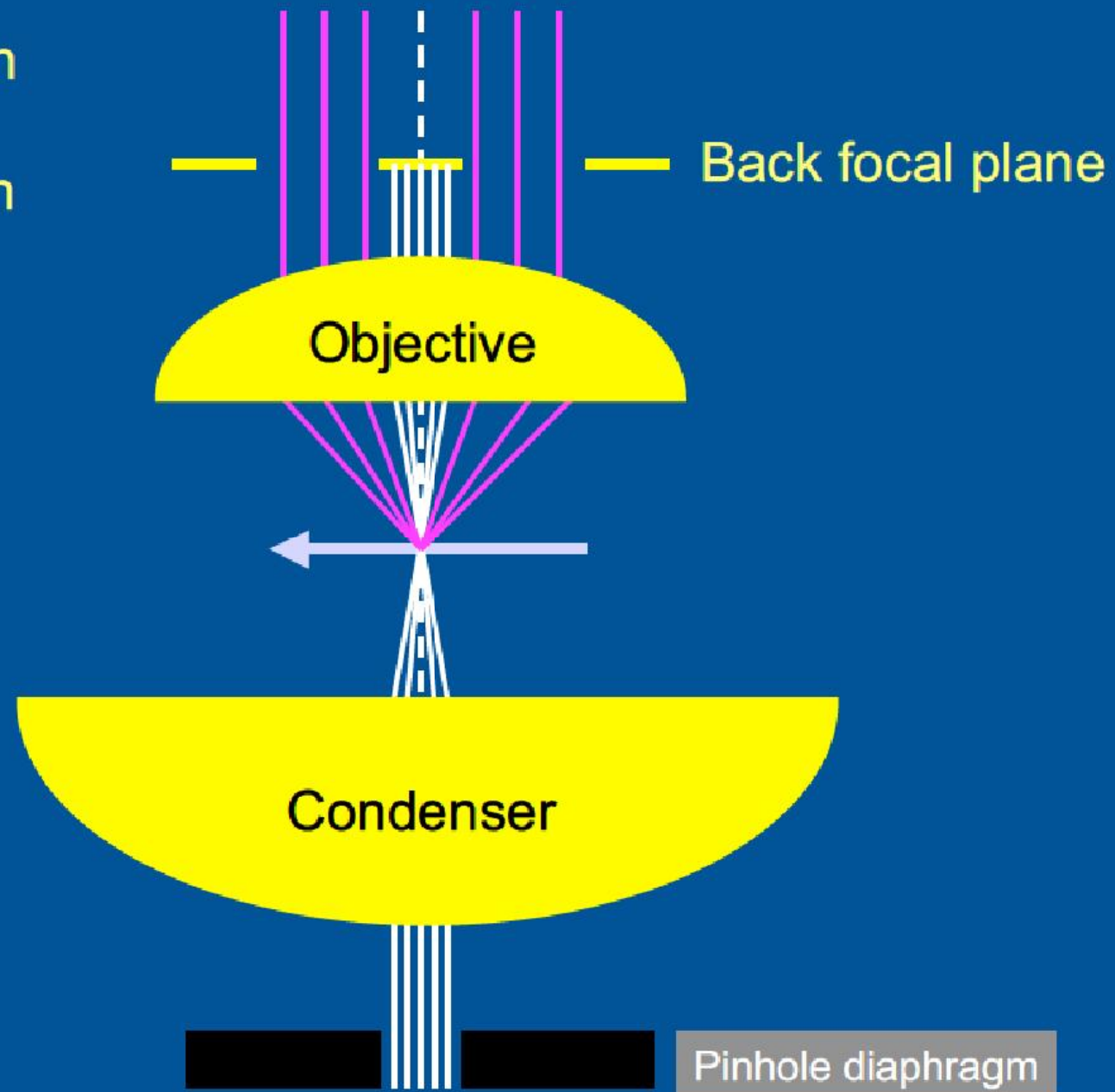


Dark field

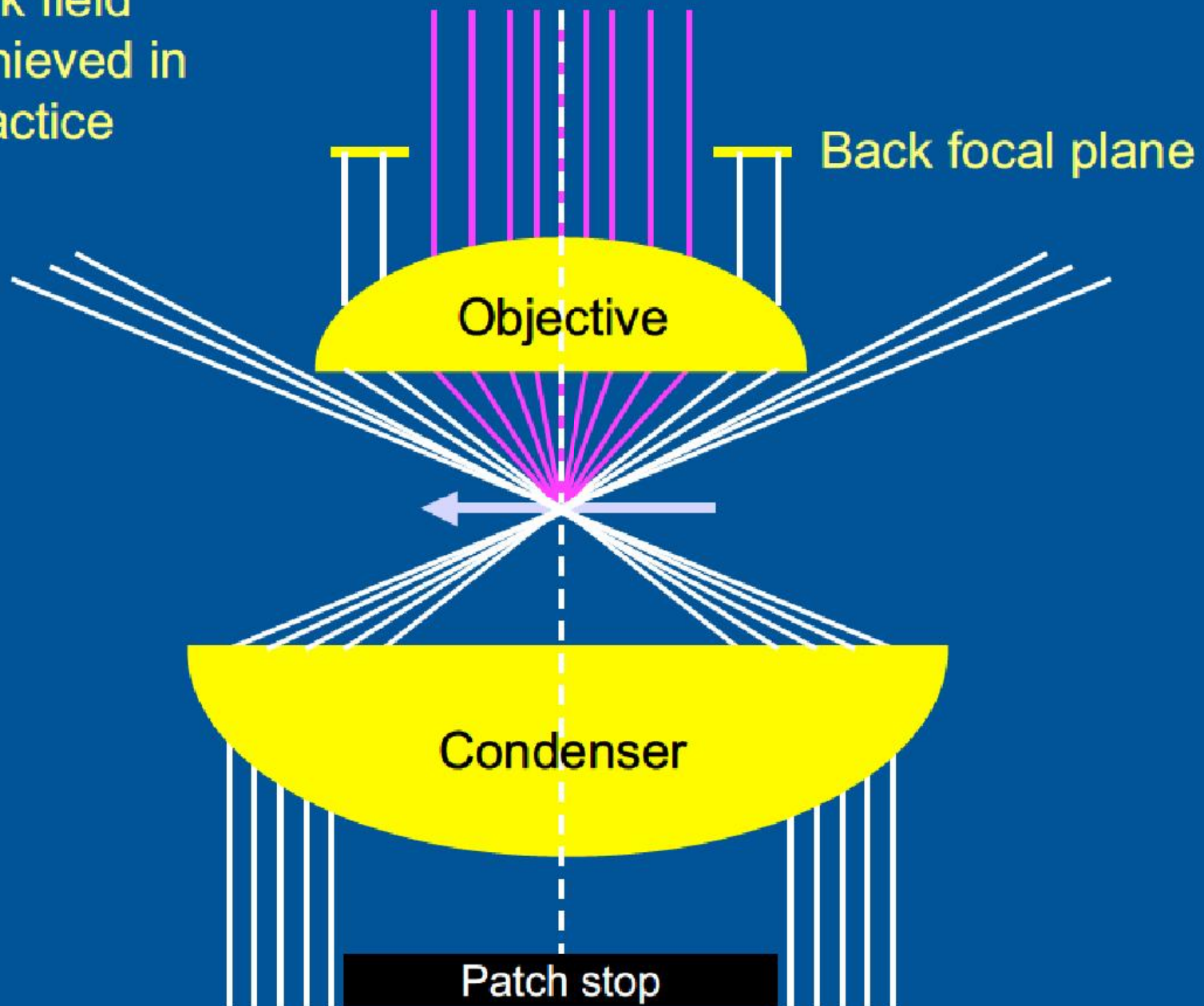
Bryozoa: dark field



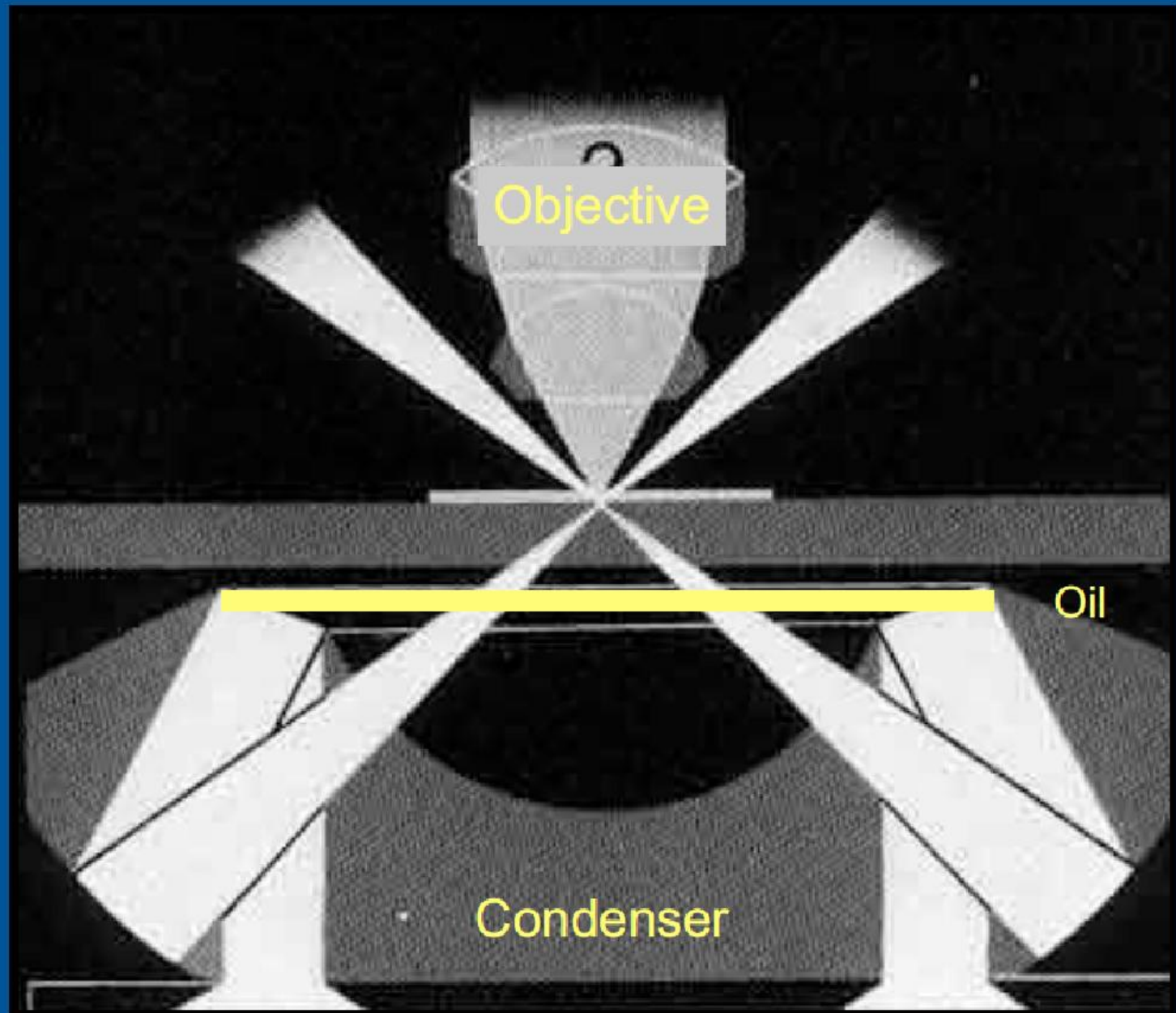
Dark field
as achieved in
diffraction
demonstration

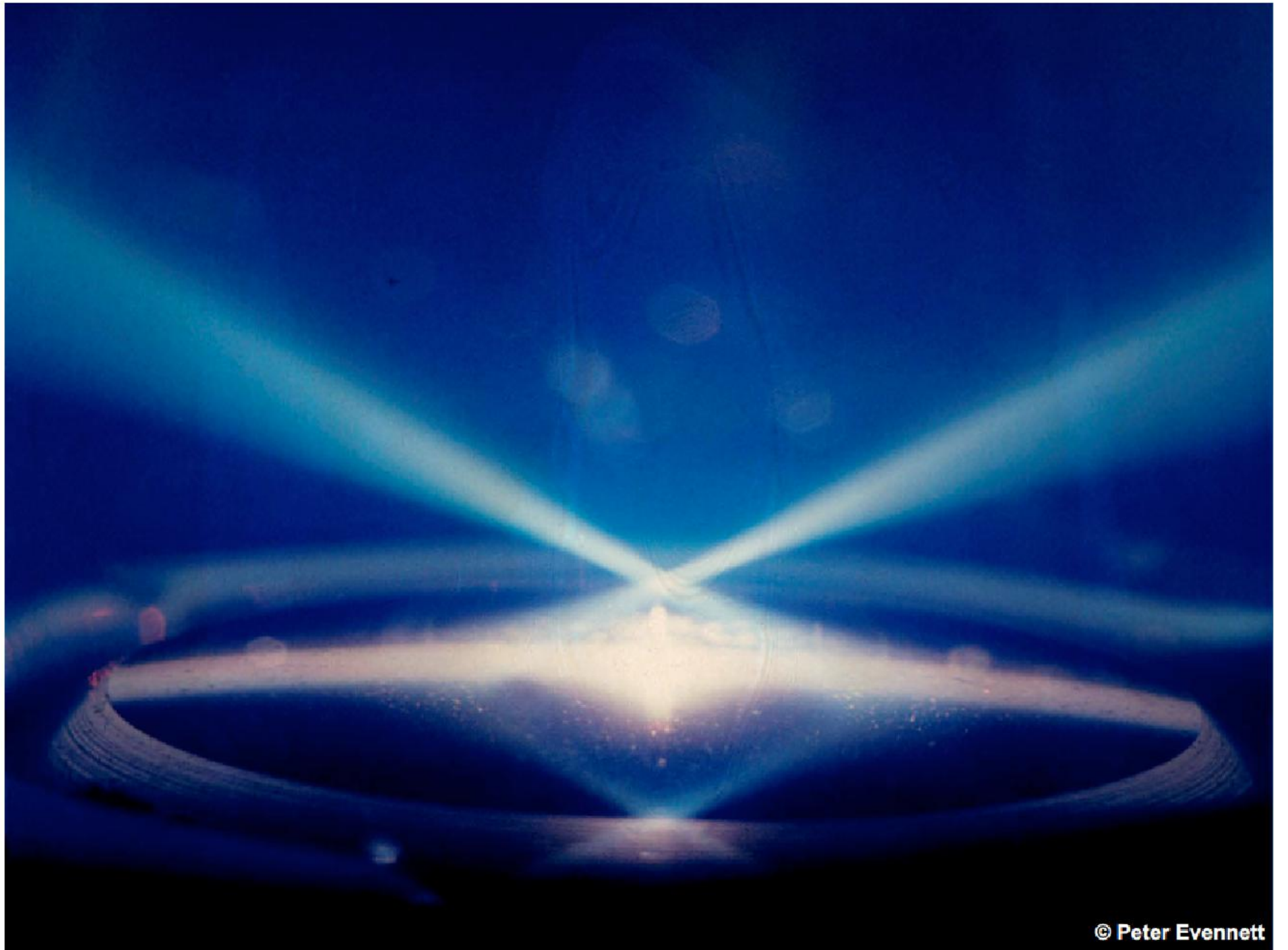


Dark field
as achieved in
practice

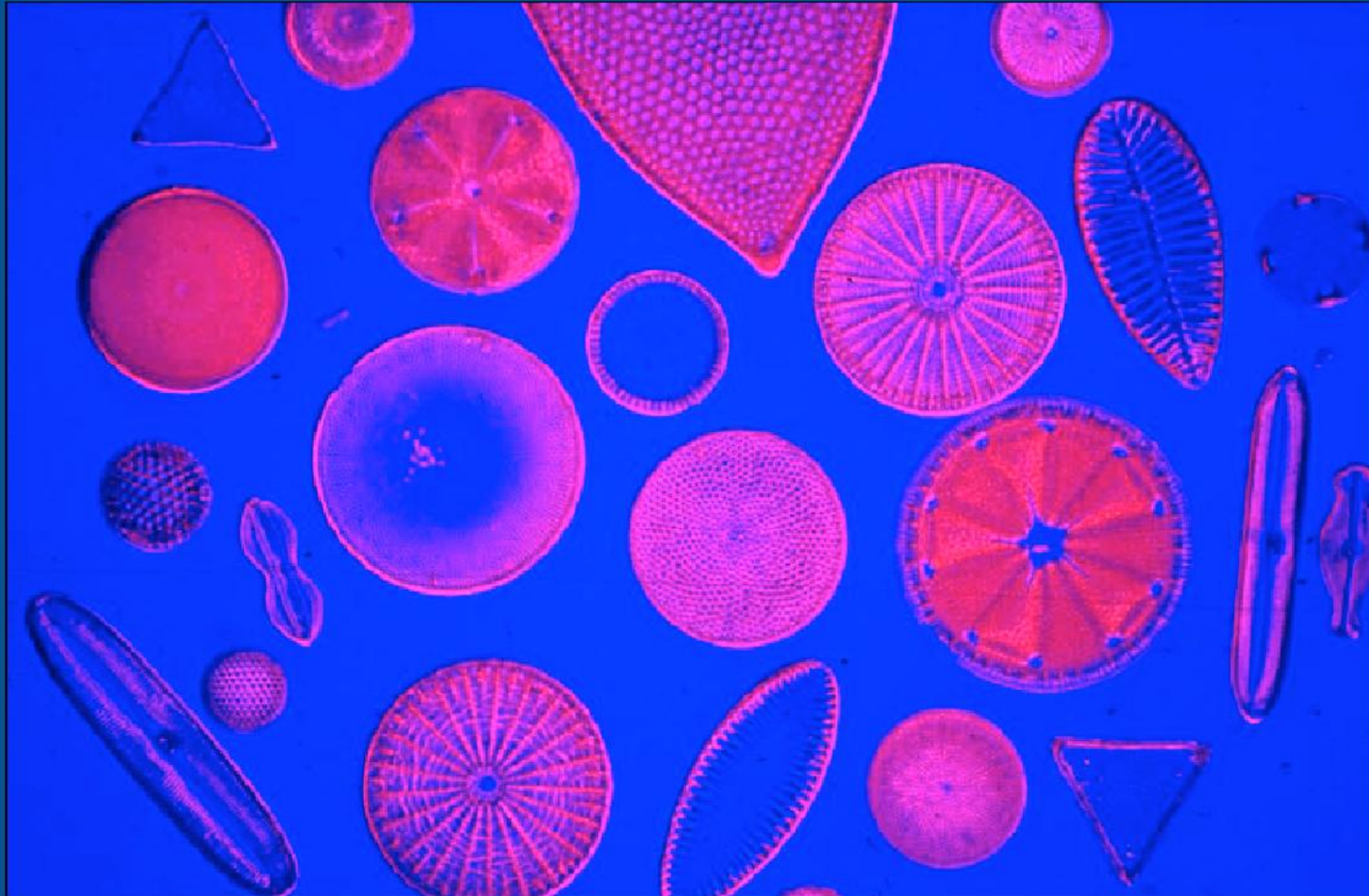


Dark field
condenser
operating
by reflection





Diatoms: Rheinberg illumination



Rheinberg illumination

