ADVANTAGES OF LIGHT AND Light microscopes	Eectron microscopes
Material can be prepared easily for	Preparation of material for examination always
examination. Often, a sample can simply be	involves a long series of procedures. These
placed on a slide with a few drops of water and	take several days to complete and often involve
• • • • • • • • • • • • • • • • • • • •	the use of toxic chemicals.
a cover slip. An image can be obtained within	
seconds.	
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Living material can be examined, so do not	Living material cannot survive in the specimens
always have to be. killed. There is less danger	vacuum inside electron microscopes. Tissues
of artificial structures appearing and causing	therefore have to be killed as the first stage in
confusion if the specimen is still alive.	the preparation of them for examination
Movement can be observed if living material is	No movement can be observed as the materia
examined, including the flow of blood,	is always dead. Movement can only be deduce
streaming of cytoplasm inside cells and the	indirectly by complex experiments, often
locomotion of microscopic organisms	involving radioactive tracers.
locomotion of mileroscopic organization	
Colours can be seen - both natural colours and	Only monochrome images are produced, with
artificial colours caused by staining	black, white and shades of grey
artificial colours caused by stamma	
	Only a small field of view can be examined at
e field of view (the area which can be	once - in a TEM the maximum uninterrur d
observed at once) is relatively large	view is about 100 µm across
<ul> <li>2 mm across at low power with typical</li> </ul>	VIOLE SECTION
microscopes.	
g d'	
The resolution of light microscopes is relatively	The short wavelength of electrons gives very
poor - about 0.25 µm so the maximum useful	good resolution - about 0.25 nm. This allows
magnification is only x 600. Many structures	magnification of about up to x 500 000. Very
within cells cannot be seen clearly.	small objects therefore become visible
	including many of the details of cell structure