Gel-electrophoresis

<u>1% Agarose gel</u>

wear Nitril-gloves

- > 0,1 g agarose per 10 ml TAE-buffer (1x) (roughly 80 mL)
- ▶ prepare shot sleeve \rightarrow use matching combs (volume of slot!)
- liquefy mix for 2 min at 495 W in microwave, boil twice (beware of boiling retardation, heat until no unsolved agarose-particles are visible anymore)
- ▶ add 8 µl EtBr/Gelred depending on gel size and concentration of EtBr/Gelred
- \rightarrow caution: EtBr/Gelred evaporates !
- cast whole gel in chamber (immediate cleaning avoids with water)
- > put combs in liquid gel and let the gel cool down
- loading probes (digestion): 25 µL DNA-sample (use big chambers)
- ➢ separate at 70 V for 60-70 Min

Gel-electrophoresis

- experiment date: ______; time: ______
- name of investigator: ______
- name of DNA-fragment /-backbone: ______
- Marker:

____ g Agarose, ____ TAE (___%), ____GELRED (3-6μl),

at

Sample	Lane	Volume in µl	Expected size in kDa
	Sample	Sample Lane	Sample Lane Volume in µl

____Volt, running time:_____minutes