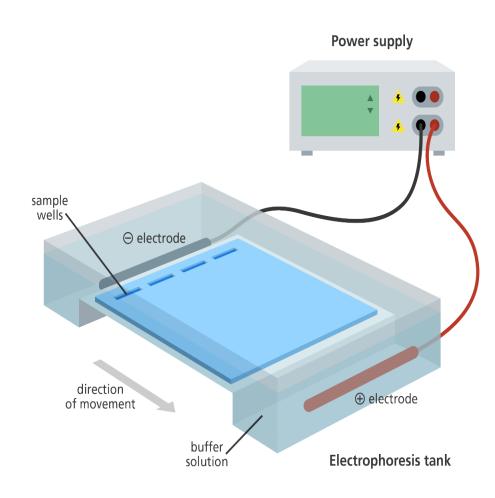


Introduction:

- Migration of a charged particle under the influence of electric field
- Electro-charged particle
- Phoresis-movement
- This electrokinetic phenomenon was observed for the first time in 1807 by Russian professors Peter Ivanovich Strakhov and Ferdinand Frederic Reuss (Moscow State University)
- Migration of clay particles dispersed in water on application of a constant electric field

Principle:

- Laboratory technique to separate macromolecules (DNA, RNA and Proteins based on size.
- Charged species such as protein, DNA and RNA migrate towards the oppositely charged electrode.
- Gel acts as a size filter/ Seive, with smaller fragments migrating faster than larger fragments.
- Gels also serves to maintain the finished separation, so that a post electrophoresis stain can be applied.



https://infograph.venngage.com/p/220088/gelelectrophoresis-technique

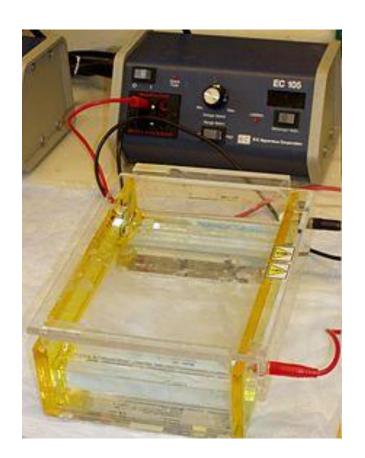
Instrumentation:

1. Electrophoretic Apparatus:

- Power Supply: voltage of 5 volts per cm (distance between electrodes) of gel.
- Staining Box: ethidium bromide
- **Trans-illuminator:** UV light box to visualize stained DNA in gels.

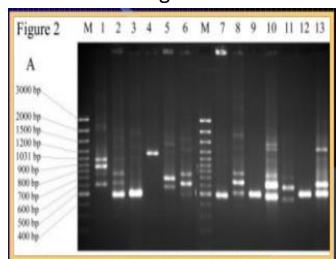
2. Chemical Components:

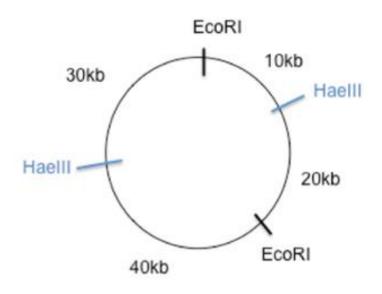
- Supporting Media: Cuts down convection current. Polyacrylamide gel & agarose
- Buffer: reduces pH changes due to the electric field. Tris/Acetate/EDTA (TAE) & Tris/Borate/EDTA (TBE)



Applications:

- DNA Gel electrophoresis is usually performed for analytical purposes, often after amplification of DNA via polymerase chain reaction (PCR)
- DNA foot-printing: how proteins bind to DNA.
- To separate proteins by size, density and purity.
- It can also be used for plasmid analysis, which develops our understanding of bacteria becoming resistant to antibiotics.





RESULTS OF GEL ELECTROPHORESIS

EcoRI	HacIII	EcoRI + HaeIII	Molecular Weight Standards	Kilobase Pairs
				100
				90
				80
				70
				60
				50
				40
				30
				20
				10

Thank you