



RAMA  
UNIVERSITY

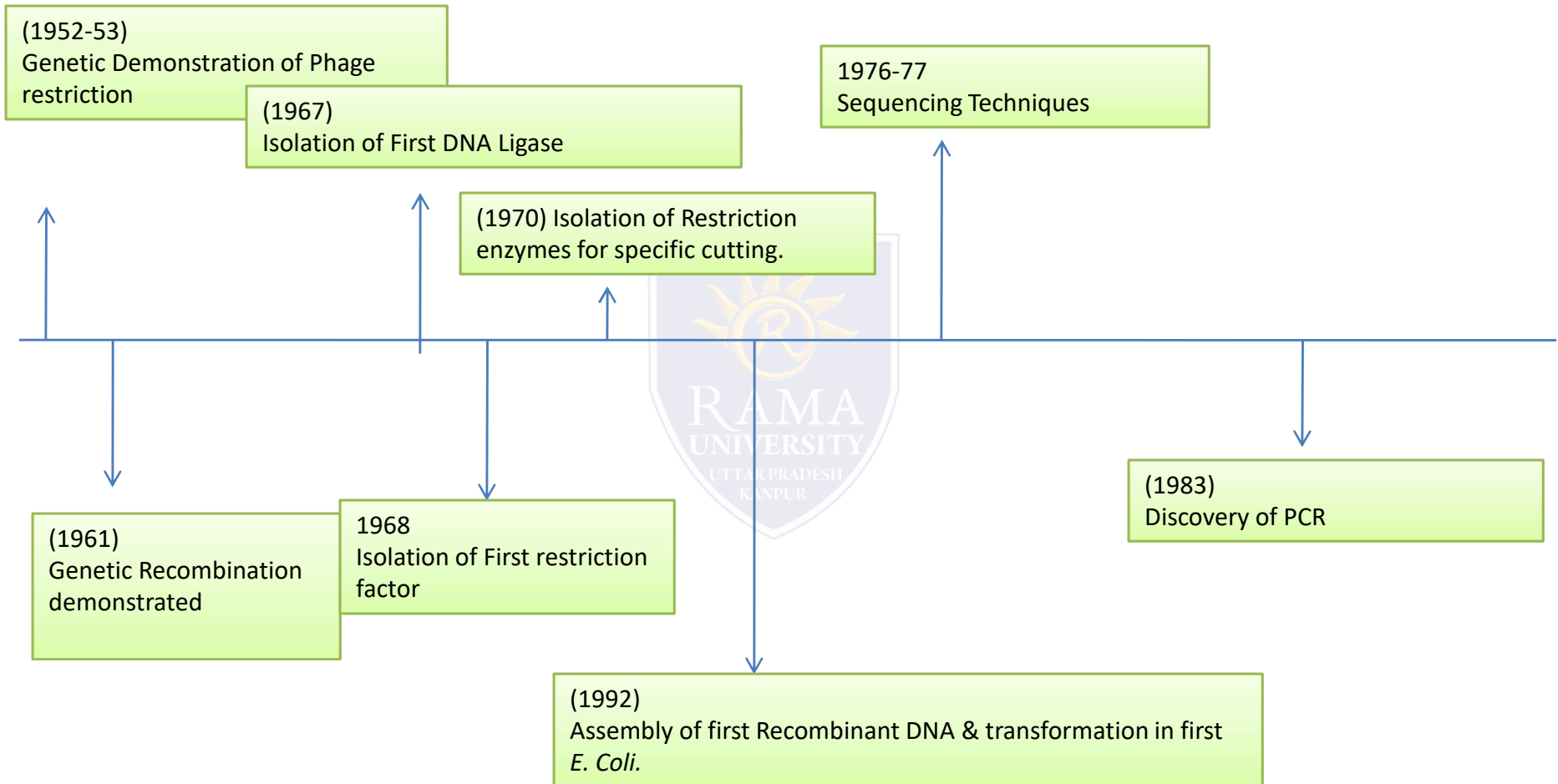
[www.ramauniversity.ac.in](http://www.ramauniversity.ac.in)

FACULTY OF ENGINEERING & TECHNOLOGY  
DEPARTMENT OF BIOTECHNOLOGY

## BACTERIAL HOST USED IN GENETIC ENGINEERING

Sr. No.	Heterologous host	Advantages	Disadvantages
1	<i>E. coli</i> (Gram-negative)	<ul style="list-style-type: none"> <li>•Fast growth rate</li> <li>•Extensive genetic tools</li> <li>•Clean chemical background for downstream natural product detection and separation</li> <li>•Comprehensive knowledge of native metabolic networks</li> </ul>	<ul style="list-style-type: none"> <li>•Lacks necessary biosynthetic machinery and precursors</li> <li>•Extensive genetic manipulation may be required for production of actinomycete natural products</li> </ul>
2	<i>P. putida</i> (Gram-negative)	<ul style="list-style-type: none"> <li>•Fast growth rate</li> <li>•Well-developed genetic tools</li> <li>•Good adaptability to different physicochemical and nutritional conditions</li> <li>•Good xenobiotics tolerance</li> <li>•High NADPH regeneration rate</li> <li>•Versatile intrinsic metabolism with diverse enzymatic capacities</li> </ul>	<ul style="list-style-type: none"> <li>•Low productivity yield of PKs/NRPs</li> <li>•Lack of advanced expression strategies for large BGCS</li> <li>•Limited knowledge of native metabolic networks</li> </ul>
3	<i>B. subtilis</i> (Gram-positive)	<ul style="list-style-type: none"> <li>•Fast growth rate</li> <li>•Thorough genetic characterization</li> <li>•Well-developed recombinant methods</li> <li>•Suitable host for a wide assortment of biologically active small molecules from <i>Bacillus spp.</i></li> </ul>	<ul style="list-style-type: none"> <li>•Lack of autonomous plasmids to facilitate cloning, transfer and heterologous expression of large BGCs</li> </ul>
4	<i>Streptomyces spp.</i> - (Gram-positive)	<ul style="list-style-type: none"> <li>•Rich in metabolic precursors and enzymatic mechanisms supporting most biosynthetic pathways</li> <li>•Versatile intrinsic metabolism supporting unique posttranslational modifications required for PKS and NRPS function</li> <li>•Suitable for expression of most proteins from actinomycetes</li> </ul>	<ul style="list-style-type: none"> <li>•Slow growth rate</li> <li>•Lack of genetic parts and advanced genetic manipulation tools</li> <li>•Endogenous competing BGCs</li> </ul>

# HISTORY OF MOLECULAR CLONING



# TRADITIONAL CLONING

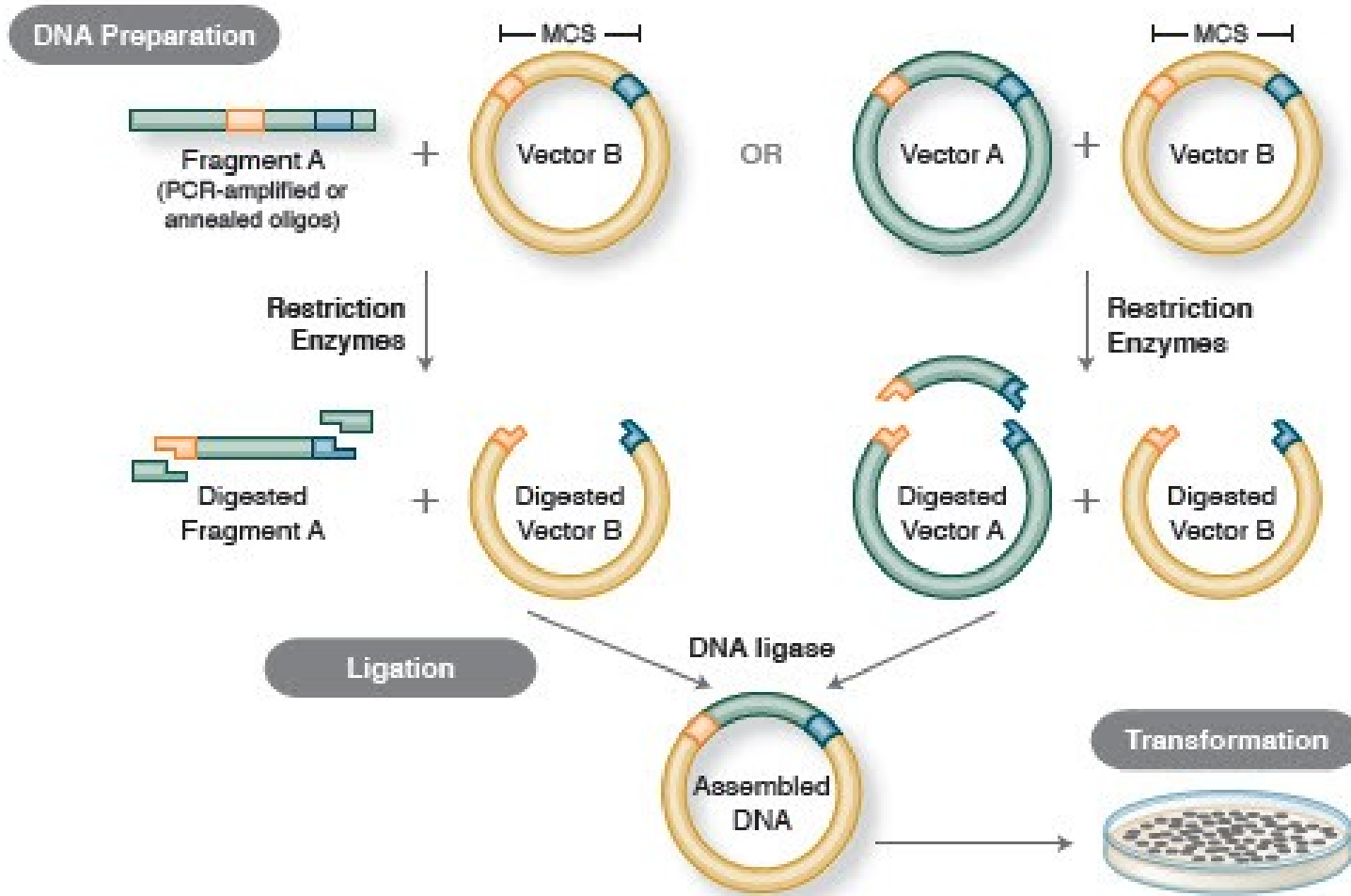


Image taken from the website of New England Biolabs