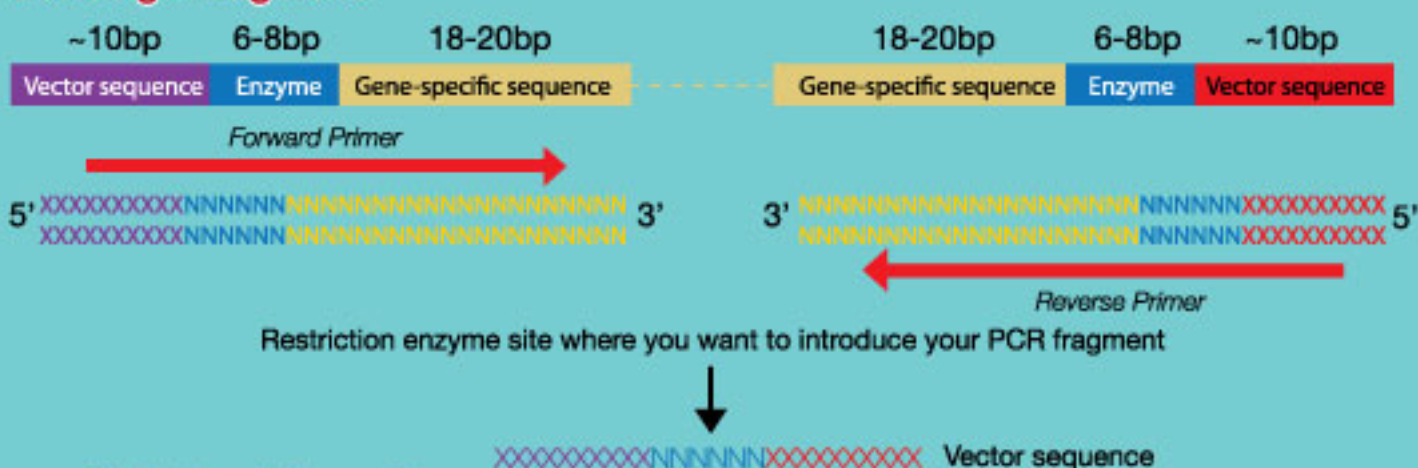


**Clone** Any PCR Fragment,  
into Any Vector, at Any Site, in One Day

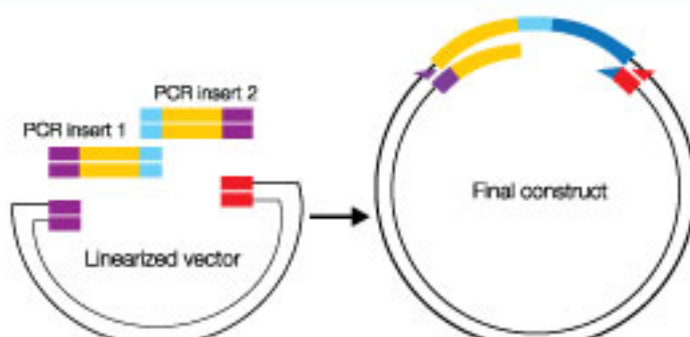
The application protocol is simple. The PCR fragments can be generated by PCR Polymerases (Taq DNA Polymerase) with primers that are designed to have at least 10 bases of homology at their linear ends. No additional treatment of the PCR fragment is required (such as restriction digestion, ligation, phosphorylation, or blunt-end polishing). The linearized vector can be generated by PCR or restriction enzymes (single or double cut). *Flex-C Enzyme* joins PCR fragments and linearized vectors accurately and efficiently by recognizing the 10bp overlap at their ends.

- ▶ Clone any insert, at any site within any vector
- ▶ Restriction enzyme, phosphatase and ligase free system
- ▶ Joining multiple fragments at once
- ▶ Broad PCR size up to 10kb
- ▶ Good for 5' overhangs, 3' overhangs, blunt end
- ▶ Precise insertion at a desired orientation
- ▶ High Efficiency with > 95% positive clones
- ▶ Multiple applications:
  - adding adaptor, linker and tag before or after the insert
  - mutation generation
  - gene synthesis
- ▶ High throughput application

## Primer Design



Catalog No	Description	Pack Size
MELP01	Flexi-Cleave Cloning Kit	20 applications



8 of 9 clones randomly picked from the plate showing a correct insert

8 of 8 clones randomly picked from the plate showing two correct inserts