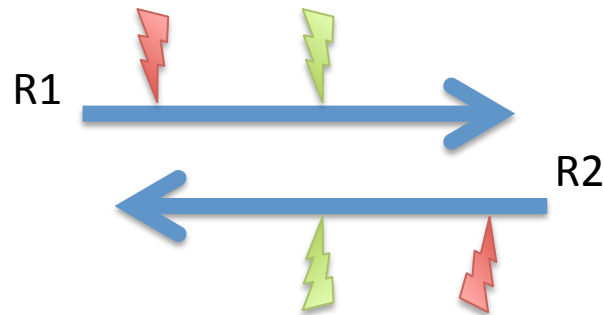


# Alternative Library Prep Methods

And Considerations for libraires with  
less errors

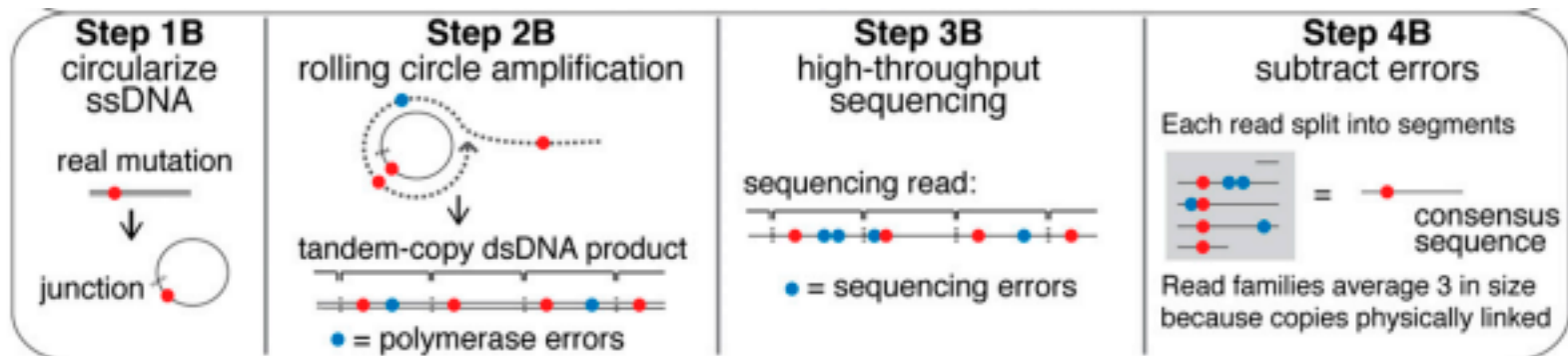
# “Double Read” strategy

- Error rates are per base, or sequence specific meaning you can leverage the relationship between read1 and read2 to do error correction.
- Error rate:  $1 \times 10^{-4}$  to  $1 \times 10^{-6}$  (probably underestimated)



# Circle Sequencing

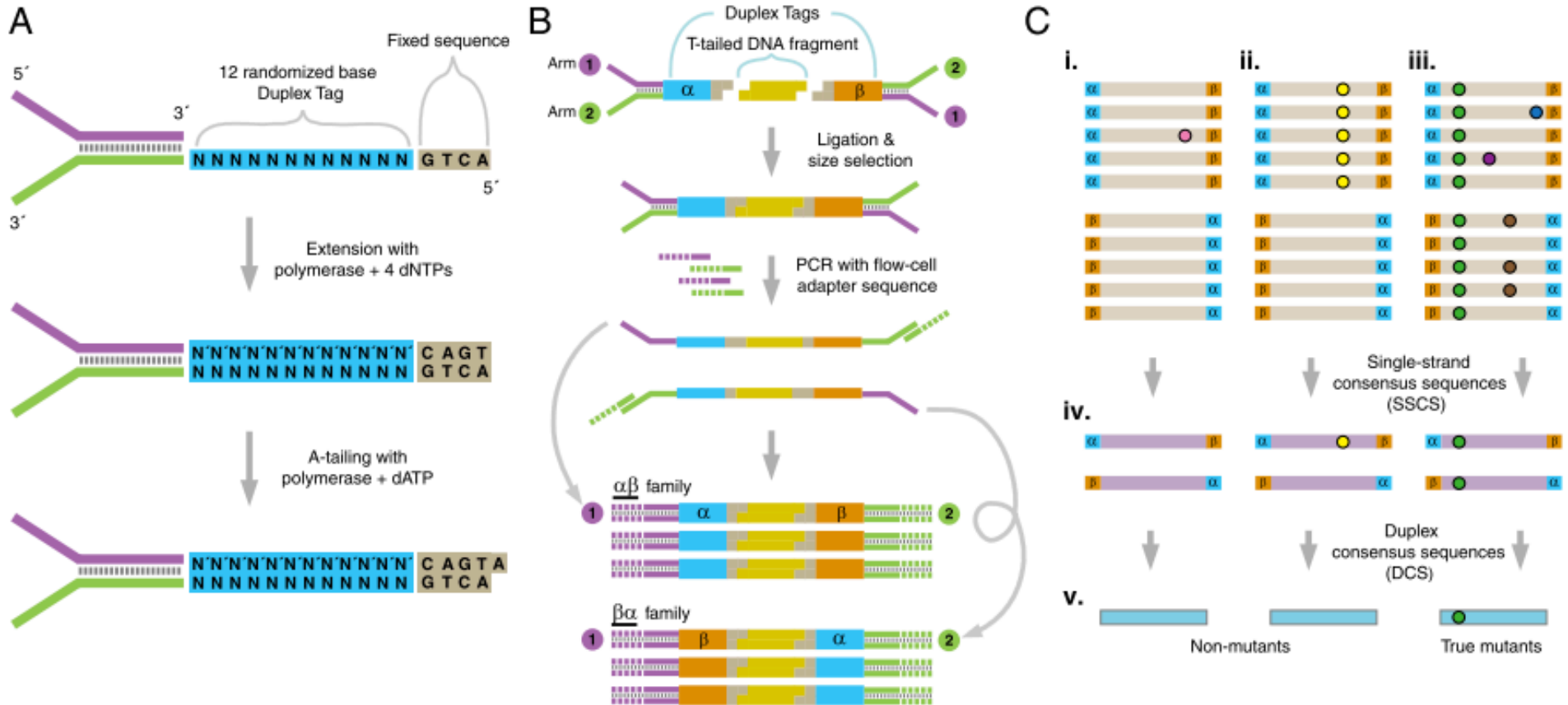
- Circularize smaller fragments of DNA.
- Read each fragment multiple times.
- Discard stochastic errors.
- Error Rate:  $7.6 \times 10^{-6}$
- Reference: [Lou et al 2013](#)



# Duplex Sequencing

- Uses a modified adapter which contains a  $N_{12}$  region which gets read before the DNA fragment.
- Reads containing the same  $N_{24}$  region are stacked together, and discrepancies removed.
- Error Rate: SSCS:  $3.4 \times 10^{-5}$  Duplex Seq:  $3 \times 10^{-6}$

# Duplex Seq Method



# Common Features

- All work on the principle that reading the same template multiple times in different ways.
- Each still subject to different types of error.