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To achieve efficient assembly of PCR fragments into a vector, we suggest using a 15–25 nt overlap with a T_m equal to or greater than 48°C (assuming A-T pair = 2°C and G-

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C pair = 4°C). To prevent errors in primer design it is highly recommended to first perform DNA fragment assembly in silico and create a final sequence file displaying both DNA strands, Fig. 2A, Step I). This virtual ...

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Page 4: Assembly

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