



# RipTide™ High Throughput Rapid DNA Library Prep

## User Manual Releases

Current release: Version 1.03 (as of July 18, 2019)

### Updates

Revision:	Distribution Date:	Notes:
1.00	11-03-2017	Initial release
1.01	06-13-2018	<p><b>Section IV :</b> Update to the PCR cycle recommendations</p> <p><b>Section VIII:</b> Updates to the following steps</p> <p><b>Step 1o:</b> Add 1 <math>\mu</math>L of 150 mM EDTA to each product of the A reaction.</p> <p><b>Step 1o (Note):</b> Recommend the use of a collection plate in combination with a plate centrifuge to harvest and combine A reaction products.</p> <p><b>Step 1v:</b> Reduce the total volume of 10 mM Tris-HCl (pH 8.0) added to SPRI Beads I at the end of the SPRI bead purification protocol from 150 <math>\mu</math>L to 100 <math>\mu</math>L.</p> <p><b>Step 3a:</b> Reduce the number of PCR cycles from nine to eight.</p> <p><b>Steps 4a (option 1) and Steps 4a and 4c (options 2,3,4 &amp; 5):</b> Include an extra pipette mixing step during three SPRI Bead II binding steps.</p>
1.02	07-03-2018	<p><b>Section VIII:</b> Updates to the following: Provide clarification regarding the SPRI Bead II volumes listed in Table 1 and in the size selection method summary at the beginning of Section 4 of the protocol.</p>
1.03	07-18-2019	<p><b>Section VIII:</b> Addition of a new step to the protocol: <b>Step 2f:</b> Perform a wash of the Capture Beads with 100 <math>\mu</math>L of 0.1 N sodium hydroxide.</p>