

plexWell™ Technology

Library Preparation Simplified

Discover the Benefits and Features of “True Multiplexing”

plexWell’s unique sequential barcoding allows the creation of balanced library pools without the need for sample or library normalization. The simplified 3-hour workflow allows for efficient paralleled multiplexing of 10s to 1000s of samples in a single sequencing run without sacrificing library quality.



Precision Multiplexing



Scalable Workflow

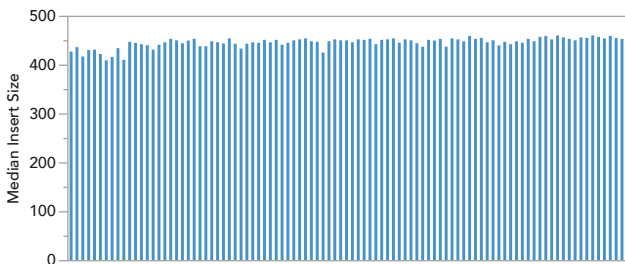


Integrated Normalization

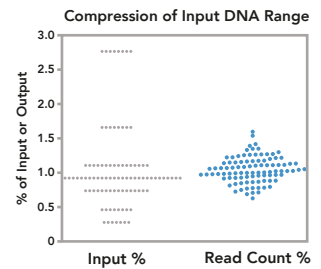
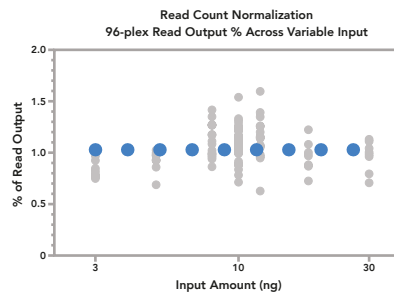
Precision Multiplexing with plexWell

Uniform Depth of Coverage and Insert Size on More Samples

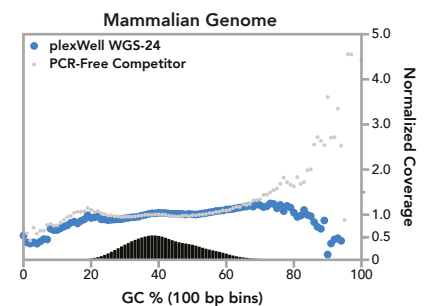
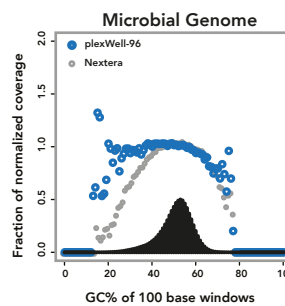
plexWell technology yields balanced multiplexed libraries containing highly uniform insert size distributions and sample read counts.



>90% of input samples at 0.5% or higher output representation; >80% of input samples within 4-fold output range; 1.5 log normalization effect within support input range (100-fold input range reduced to 2.5 output range). E. coli library preparation with plexwell gives consistently longer median inserts, maximizing the per read usable data. Insert sizes are consistent over a range of inputs.



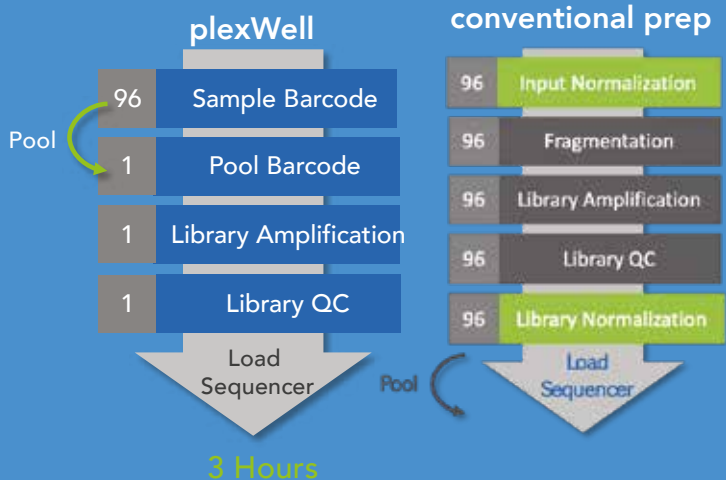
Minimal Bias for Diverse Applications



plexWell Applications:

- Small & Large Whole Genomes
- Microbial Genomes
- Amplicons & Plasmids
- Low-Pass Plant/Animal/Human
- BAC/Fosmid/Cosmids
- RNAseq
- 16s or ITS

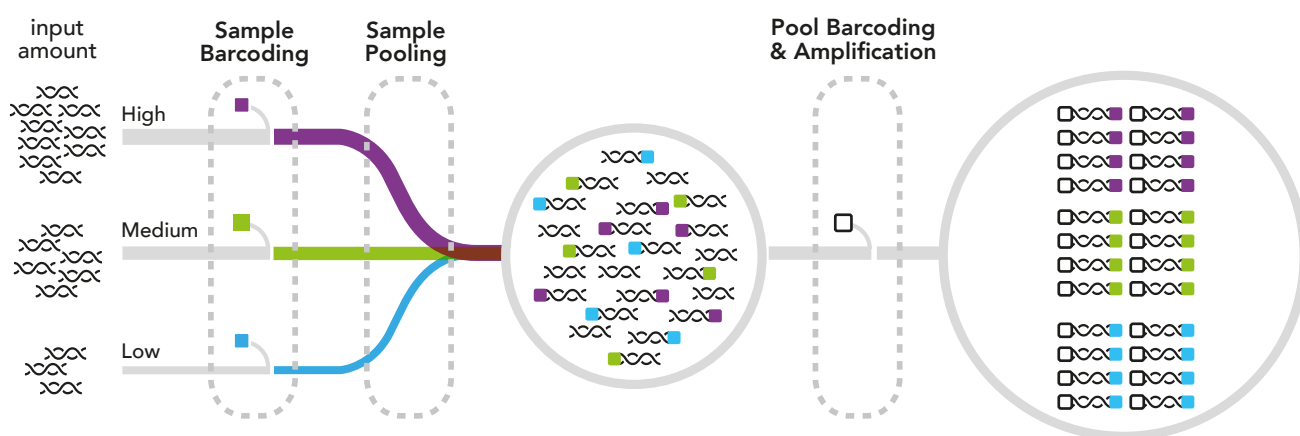




Simple and Scalable Workflows

- 3 hours total time from samples to sequencer loading
- No quantification of sample DNA
- Eliminate library normalization
- Multiplex thousands of samples in a day
- No upfront or additional equipment cost

Normalizing Iterative Barcoding Technology



Sample Barcoding

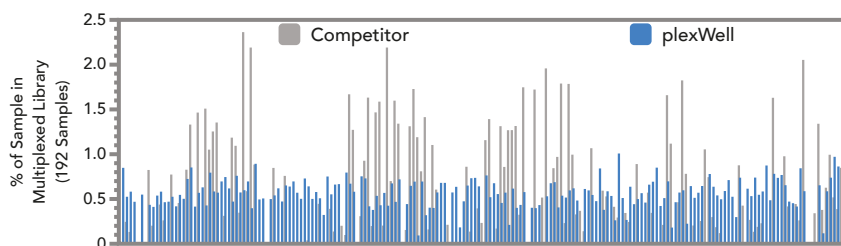
Multiple input samples are tagged with limiting quantities of sample-specific, barcodes containing P7-adapters.

Pool Barcoding

Tagged samples are pooled into a single tube and undergo plate barcoding using excess P5-adapters.

Normalized Library

Following final library amplification and QC, the auto-normalized library is ready for sequencing.



plexWell achieves a significantly better level of multiplexing uniformity for highly multiplexed sequencing applications. At left, sequencing results obtained for sequencing 192 samples of amplified single-cell cDNA with plexWell (blue) and Competitor reagents (gray). Input DNA for Competitor was pre-normalized, whereas plexWell library was made from un-normalized amplified cDNA. Read count variation for plexWell showed 27% C.V. versus 71% for Competitor.

Time and Cost Savings for Multiplexed Sequencing Applications



35 Minutes
Hands-on Time



3000 Samples
Per 8 Hour Day



as low as €7
Per Sample



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Ready to integrate plexWell™ technology into your NGS workflow?
Contact your local distributor Sopachem to learn more



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