# Two New Solutions for Nucleic Acid Fragmentation

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#### the pre-analytical advantage™

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### **Covaris – the established standard for DNA shearing**

Overview	S-series	<b>E-series</b>	L-series	• 500 kHz Transducer	<ul> <li>Circular transducer</li> </ul>	<ul> <li>Linear Transducer</li> </ul>
<ul> <li>The Covaris Adaptive Focused Acoustics (AFA<sup>™</sup>) technology enables unprecedented iso-thermal control over DNA shearing forces in the sample.</li> </ul>		Covaris	Covaris	technology	• Serial processing	Parallel processing
<ul> <li>The development of clinical applications requires high recovery, high reproducibility, robust systems and methodology, and</li> </ul>			LE220	Sample Vessel Focal Zone		Focal Zone
automation capabilities.	• Manual	Automated      Batch	<ul> <li>Linear transducer – Parallel</li> <li>processing</li> </ul>	Water Bath		

• With the increasing resolution of genomic analytical techniques, high recovery, bias free DNA shearing becomes critical.

• Single

• Circular transducer

• Single sample processing

• Circular transducer

• 24 to 96 sample processing

Robot integration for full

automation

• Highly uniform field along a row treats a full 96-w plate 8x faster



## g-TUBE – DNA fragmentation from 6kb to 20kb



**Versatile:** Ideal for direct sequencing, mate-pair libraries, and other applications that require longer DNA fragments. Selectable Fragment Size: g-TUBE shears DNA in user-selectable fragment sizes ranging from 6kb to 20kb. Highly Reproducible: DNA shearing results with g-TUBE are reproducible assay-to-assay, lab-to-lab, day-to-day. Fast and Scalable: Shear 6kb - 20kb fragments in 2 minutes or less. Runs multiple samples simultaneously. **Efficient:** High sample recovery (90%+) with a closed vessel process.

**Economical:** Use an Eppendorf<sup>®</sup> "MiniSpin<sup>®</sup> plus" microcentrifuge - no other equipment needed.

Ladder kbp		1	2	3	4	23 Ladder kbp	
10	-			H	-	9.4	4 611 / 44500
8	See	11	п	н	4.6	6.6	1 – 6 kbp / 14500 rpm
6	tink.	23		10			2 $0 $ $k $ $h $ $n $ $1 $ $0 $ $1 $ $0 $ $n $ $m $ $m$

#### From DNA to sequencing with Covaris g-TUBEs and PacBio RS

E coli DNA was sheared to 10 kbp with both Covaris g-TUBE and Hydroshear. Experiment was repeated for 2 different inputs of DNA.

Libraries built following PacBio RS protocols and sequenced on PacBio RS instrument.



Easy to follow protocol: Less than 3 minutes to process up to 24 g-TUBEs

Eppendorf* MiniSpin plus – Speed (RPM)							
Targeted size	6 kbp	8kbp	10 kbp	20 kbp			
Mass of DNA							
4 μg	14,500	9,400	8,000	5,500			
8 µg	14,500	9,400	8,000	6,500			
15 μg	-*	11,200	9,400	7,600			
30 µg	-*	13,000	11,200	9,600			
Processing time (seconds)	30 sec	60 sec	60 sec	60 sec			

Easy to follow protocol: Required centrifuge speed is determined by the targeted fragment size and mass of input DNA.



Ideal companion for bench-top NGS instruments

• Comparable to industry-standard AFA instruments

• Provides isothermal and bias-free DNA shearing

Sample ID		Starting ng		% Yield
g-TUBE 10K SB 7.5ug - 10uL		5000		14%
g-TUBE 10K SB 15ug - 10uL		5000		13%
Hudrochoor 10K SP 7 Eug P1 10ul		5000		1 / 0/
Hydroshear IOK 3B 7.50g KI - IOUL		5000		1470
Hydroshear 10K SB 15ug - 10uL		5000		11%
	gTL	JBE	Hydroshear SC13	
	10K 7.5ug	10K 15ug	10K 7.5ug	10K 15ug
lob Metric	Value	Value	Value	Value
% Adapter Dimer (0-10bp)	0.16	0.67	0.56	1.09
6 Short Insert (11-100bp)	0.02	0.08	0.06	0.12
t of Movies	3	3	3	3
Pre-Filter#of Bases	323492608	400785167	393724278	377533951
Post-Filter # of Bases	172001266	192748644	125869867	117074824
Pre-Filter # of Reads	225459	225459	225459	225459
Post-Filter#of Reads	48707	47989	30202	28672
Pre-Filter Mean Readlength	637	686	441	409
Post-Filter Mean Readlength	2899	3154	3226	3148
Pre-Filter Mean Read Quality	0.177	0.173	0.108	0.102
Post-Filter Mean Read Quality	0.819	0.811	0.801	0.802
f Post-Filter Reads	48707	47989	30202	28672
Aean Mapped Subread Readlength	2072	2173	2355	2247
of Mapped Reads	47374	46045	28825	26550
Mean Mapped Readlength	2769	3025	3099	3001
For Mapped Subreads	62407	62948	3/3//	34838
tof Menned Read	/155	120202402	/ 858	70666646
or wapped Bases	131175408	139293492	12097	79666616
Acon Monnod Subroad Acouracy	04.06	12/0/	13087	12408
lean Depth of Coverage	04.20	03.59	02.53	02.00
real Depth of Coverage	20.21	21.19	0	15.75
	0		0	0



Post sequencing metrics for both Hydroshear and g-TUBEs libraries. Mean depth of coverage (top) and mean of maximum subread length (bottom)

### M220 Personal Fragmenter – Compact instrument for 150bp to 3kb

15ml Water bath volume

 Ready to use in <10 minutes after power-up</li> Integrated Peltier thermo-electric cooling



— 300 bp —— 500 bp — 1000 bp - 3000 b

• Pre-loaded shearing conditions: click and go

• Small footprint, no external chiller

**Versatile:** AFA provides fine control of the energy delivered to the sample and enables tight fragment size distributions with a mean selected between 150bp and 3 kb. Left side, electropherogram from Agilent Bioanalyser 2100 plotted on a linear scale. On the right, same data plotted on a logarithmic scale.

 Compatible with Covaris microTUBES and miniTUBES

 Robust construction and simple user interface

Biosciences, Menlo Park, CA



High reproducibility: 10 replicates of DNA samples sheared with M220 following a 300bp protocol

