

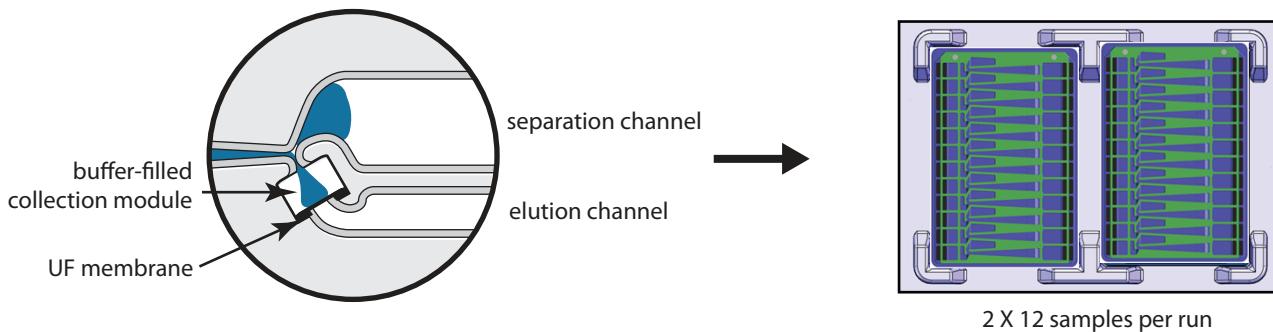
Collect up to 24 Size-Selected DNA Fractions in 30 Minutes

Benefits:

- Generates higher-quality libraries for improved sequence informatics
- Fragment sizes and ranges are reproducibly collected, providing consistent results, run-to-run
- Flexible programming allows collection of multiple size ranges, or up to 24 collections of the same fragment range



Automated Preparative Electrophoresis



Sage Science's proprietary technology, featuring electro-elution from agarose, has been configured to run 12 samples on an SBS-footprint gel cassette— **for higher-throughput workflows and with a lower cost per sample.**

Specifications:

Sample Capacity	Maximum Sample Load	Run Times	Target Range	Min. Size Distribution as Expressed by (CV)	Accuracy*	Reproducibility**
12 samples/cassette	1.5 µg	25-50 minutes	90 - 2000 bp	<8%	≥ 90%	≥ 90%
2 cassettes (24 samples)/run		40 min for 500 bp				

* 100% minus the deviation of actual target value (Agilent Bioanalyzer) from software input value divided by the actual value.

** 100% minus 2X standard deviation of replicate samples.

Take Good Care of Your Library.



Automated size selection: An indispensable tool for NGS

Selected References Citing Pippin DNA Size Selection

Paired-End Sequencing

Sheaffer, K.L. et al., DNA methylation is required for the control of stem cell differentiation in the small intestine. *Genes Dev.* 28: 652-664 (2014)

Mate-Pair Sequencing

Srivastava, A. et al., Discovery of transgene insertion sites by high throughput sequencing of mate pair libraries. *BMC Genomics.* 15:367 (2014)

Long-Read Sequencing

Benson, M.A. et al., Evolution of hypervirulence by a MRSA clone through acquisition of a transposable element. *Molecular Microbiology* 93(4), 664-681 (2014)

miRNA Isolation

Singhal, R. et al. FOXO1 regulates expression of a microRNA cluster on X chromosome. *AGING*, 5:5. (2013)

ChIP-seq

Wharton, K., et al., Methylation and Next-Generation Sequencing of free circulating DNA from human plasma. *BMC Genomics*, 15:476 (2014)

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Москва
ул. Магаданская, д. 7, к. 3 ■ тел./факс: (495) 745-0508 ■ sales@dia-m.ru

www.dia-m.ru

С.-Петербург
+7 (812) 372-6040
spb@dia-m.ru

Новосибирск
+7(383) 328-0048
nsk@dia-m.ru

Воронеж
+7 (473) 232-4412
vrn@dia-m.ru

Йошкар-Ола
+7 (927) 880-3676
nba@dia-m.ru

Красноярск
+7(923) 303-0152
krsk@dia-m.ru

Казань
+7 (843) 210-2080
kazan@dia-m.ru

Ростов-на-Дону
+7 (863) 303-5500
rnd@dia-m.ru

Екатеринбург
+7 (912) 658-7606
ekb@dia-m.ru

Кемерово
+7 (923) 158-6753
kemerovo@dia-m.ru

Армения
+7 (094) 01-0173
armenia@dia-m.ru

