



## Agilent DNA 1000 Kit

### Agilent DNA 1000 Kit (reorder number 5067-1504)

#### DNA Chips

25 DNA Chips

1 Electrode Cleaner

#### Syringe Kit

1 Syringe

3 Spin Filters

#### DNA 1000 Reagents (reorder number 5067-1505)

● (yellow) DNA 1000 Ladder

● (green) DNA 1000 Markers 15/1500 bp (2 vials)

● (blue) DNA Dye Concentrate\* (1 vial)

● (red) DNA Gel Matrix (3 vials)

\*) "This product is provided under a license by Life Technologies Corporation to Agilent Technologies. The purchase of this product conveys to the buyer the non-transferable right to use the purchased amount of the product and components of the product only as described in accompanying product literature. The sale of this product is expressly conditioned on the buyer not using the product or its components (1) in manufacturing; (2) to provide a service, information, or data to an unaffiliated third party for payment; (3) for therapeutic, diagnostic or prophylactic purposes; (4) to resell, sell or otherwise transfer this product or its components to any third party, or use for any use other than use in the subfields of research and development, quality control, forensics, environmental analysis, biodefense or food safety testing. For information on purchasing a license to this product for purposes other than described above contact Life Technologies Corporation, Cell Analysis Business Unit, Business Development, 29851 Willow Creek Road, Eugene, OR 97402, Tel: (541) 465-8300, Fax: (541) 335-0354."

### Physical Specifications

Type	Specification
Analysis run time	35 minutes
Number of samples	12 samples./chip
Sample volume	1 µl
Kit stability	4 months (Storage temperature see individual box!)

### Analytical Specifications

Type	Agilent DNA 1000 Assay
Sizing range	25–1000 bp
Typical sizing resolution	± 5 bp 25–100 bp ± 5 % 100–500 bp ± 10 % 500–1000 bp
Sizing accuracy	± 10 %* (for ladder as sample)
Sizing reproducibility	5 % CV (for ladder as sample)
Quantitation accuracy	20 %* CV (for ladder as sample)
Quant. reproducibility	25–500 bp: 15 % CV; 500–1000 bp: 5 % CV (for ladder as sample)
Quantitative range	0.1–50 ng/µl
Maximum salt	250 mM for KCl or NaCl, 15 mM for MgCl <sub>2</sub>

\*) Some fragments below 70 bp may deviate from the above specifications.



# Agilent Dna 1000 Kit Quick Start Guide

**Angie Ambers**



## Agilent Dna 1000 Kit Quick Start Guide:

*Hydrogenase: Structure, Function, Maturation, and Application* Stefan Frielingsdorf, Francesca Valetti, Constanze Pinske, Chris Greening, 2023-10-24 Molecular hydrogen H<sub>2</sub> is an important molecule in the metabolism of diverse microorganisms. It is used either as energy source or for the disposal of reducing equivalents depending on environmental conditions. Furthermore, H<sub>2</sub> transmits energy between different species within microbial communities. The enzymes that split or evolve H<sub>2</sub> are called hydrogenases, and these metalloproteins can be divided into three phylogenetically unrelated classes distinguishable by the metal composition of their active sites, namely Fe-FeFe and NiFe hydrogenases. Following a century of hydrogenase research, it is now possible to isolate, handle, and investigate these fragile enzymes. There have been numerous advances in understanding the regulation, function, structures, and maturation of these enzymes, as well as their involvement in important processes such as microbial pathogenesis and biogeochemical cycling. The employment of hydrogenases and hydrogenase-based applications could also potentially facilitate the world's transition to a future sustainable H<sub>2</sub>-based energy economy.

*Sheep and Goat Gene Exploration* Shaobin Li, Rui Su, Xin Wang, 2022-04-25

**Genetics, Genomics and -Omics of Thermophiles, 2nd Edition** Kian Mau Goh, Kok-Gan Chan, Rajesh Kumar Sani, Edgardo Rubén Donati, Anna-Louise Reysenbach, 2019-06-13 This eBook is a collection of articles from a Frontiers Research Topic. Frontiers Research Topics are very popular trademarks of the Frontiers Journals Series; they are collections of at least ten articles all centered on a particular subject. With their unique mix of varied contributions from Original Research to Review Articles, Frontiers Research Topics unify the most influential researchers, the latest key findings, and historical advances in a hot research area. Find out more on how to host your own Frontiers Research Topic or contribute to one as an author by contacting the Frontiers Editorial Office: [frontiersin.org/about/contact](mailto:frontiersin.org/about/contact).

The Urogenital Microbiota in Urinary Tract Diseases A. Lenore Ackerman, 2023-01-10

**Implications of lipids and modified lipoproteins in atherogenesis: From mechanisms towards novel diagnostic and therapeutic targets** Alexander Akhmedov, Chu-Huang Chen, Tatsuya Sawamura, 2023-08-01

*Human Microbiome: Symbiosis to Pathogenesis* Learn-Han Lee, Nurul-Syakima Ab Mutalib, Sunny Hei Wong, Siok-Fong Chin, Vishal Singh, 2021-03-24

*Forensic Genetic Approaches for Identification of Human Skeletal Remains* Angie Ambers, 2022-11-15 Forensic Genetic Approaches for Identification of Human Skeletal Remains: Challenges, Best Practices, and Emerging Technologies provides best practices on processing bone samples for DNA testing. The book outlines forensic genetics tools that are available for the identification of skeletal remains in contemporary casework and historical archaeological investigations. Although the book focuses primarily on the use of DNA for direct identification or kinship analyses, it also highlights complementary disciplines often used in concert with genetic data to make positive identifications, such as forensic anthropology, forensic odontology, and forensic art sculpting. Unidentified human remains are often associated with tragic events such as fires, terrorist attacks, natural disasters, war conflicts, genocide, airline crashes.

homicide and human rights violations under oppressive totalitarian regimes In these situations extensive damage to soft tissues often precludes the use of such biological samples in the identification process In contrast bone material is the most resilient viable sample type for DNA testing DNA recovered from bone often is degraded and in low quantities due to the effects of human decomposition environmental exposure and the passage of time The complexities of bone microstructure and its rigid nature make skeletal remains one of the most challenging sample types for DNA testing Provides best practices on processing bone samples for DNA testing Presents detailed coverage of proper facilities design for skeletal remains processing selection of optimal skeletal elements for DNA recovery specialized equipment needed preparation and cleaning of bone samples for DNA extraction and more Highlights complementary disciplines often used in concert with genetic data to make positive identifications such as forensic anthropology forensic odontology and forensic art sculpting

**Southern California Coastal Water Research Project Annual Report** Southern California Coastal Water Research Project,2013

*Cancer Research* ,2009-07    *Journal of Clinical Microbiology* ,2004    *Tissue Engineering* ,2008    *American Laboratory* ,2000    **International Journal of Oncology** ,2008    *The Journal of Immunology* ,2007-12    *Genetic Engineering News* ,2005    □□□□□□□□ ,2014    **Proceedings of the Entomological Society of Washington** Entomological Society of Washington,2018 List of members in v 1 3 5 14    *Journal of the National Cancer Institute* ,2005  
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