



# Agilent DNA 1000 Kit

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**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)

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**Physical Specifications**

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

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**Analytical Specifications**

Type	Agilent DNA 1000 Assay
Sizing range	25–1000 bp
Typical sizing resolution	$\pm$ 5 bp 25–100 bp $\pm$ 5 % 100–500 bp $\pm$ 10 % 500–1000 bp
Sizing accuracy	$\pm$ 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/ $\mu$ l
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 Frielingdorf,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

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*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.

## **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  
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