



Agarose PFGE

Agarose for gel with very high gel strength and higher exclusion limit

Product Description

Name :	Agarose PFGE			
Catalog Number	: A2X2G0, 100g A2X2G1, 500g	3		
Specifications :	Moisture	$\leq 7\%$	Gel Strength 1.5% (g/cm 2) \geq 3200	
	Ash	$\leq 0.25\%$	Gelling Temperature 1.5% (°C) 36±1.5	
	EEO (electroendosmosis)	≤ 0.12	Melting Temperature 1.5% (°C) 88±1.5	
	Sulfate	$\leq 0.12\%$	DNAse/RNAse activity	None detected
	Clarity 1.5% (NTU)	≤ 4	DNA resolution ≥ 1000 bp	Finely resolved
	Gel Strength 1% (g/cm 2)	≥ 1800	Gel background	Very low
Applications :	 Conventional Electrophoresis: can be used in • Blotting a wide range of concentrations. • Pulsed Field Gel Electrophoresis: because of its higher exclusion limit, larger molecules can be separated. • Conventional Electrophoresis: can be used in • Blotting • Agarose Beads preparation. • Cell and enzyme immobilization. 			
Storage: F	Room temperature (Z			

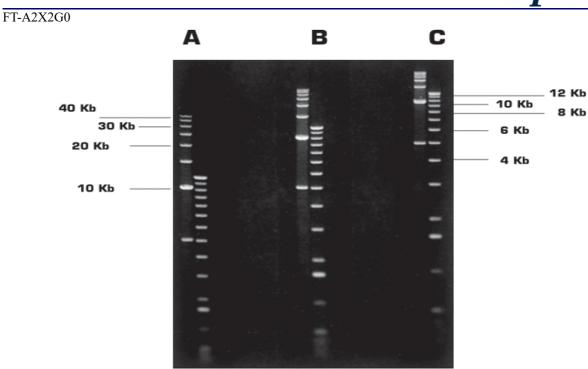
Introduction

Agarose PFGE is a linear polymer with a very high molecular weight, giving gel structures unlike those of traditional agaroses. This characteristic, added to the very low sulfate content, produces an strong intercatenary interaction, yielding a gel with very high gel strength and higher exclusion limit.

As we can see in the following photographs, Agarose PFGE is suitable for a **wide variety of ranges**, just by modifying its concentration.







2 Agarose PFGE gels in 1 X TAE. A-0.3%, B-05%, C-08%. Markers: lane 1-5 kb, lane 2-1 kb ladder. Electrophoresis conditions: submarine gel, 16 hours, 1 V/cm. in IXTAE buffer.

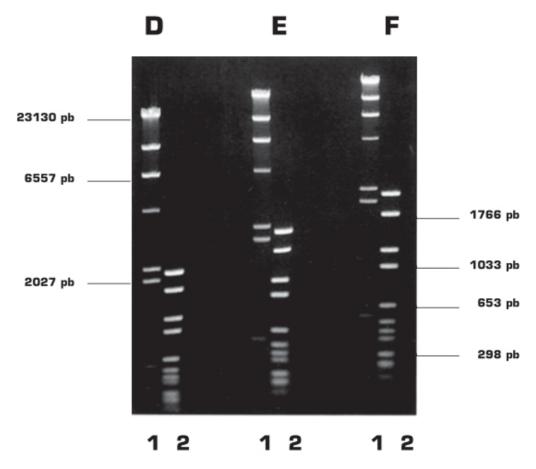
1

1

2

2

1



Agarose PFGE gels in 1 X TAE. D-0.5%, E-1%, F-1.5%. Markers: lane 1-Lambda DNA. HindIII, lane-2-pBR328DNA. Bgll + pBR328DNA. Hindfl. Electrophoresis conditions: submarine gel, 2 hours, 4.5 V/cm. in 1XTAE buffer.

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Technical and Scientific Information

- Extremely high gel strength allowing for lower gel concentrations (0.3%), enabling it to be used not only with high molecular weight nucleic acids, including chromosomes, but also with large sized particles like viruses and ribosomes.
- High electrophoretic mobility. DNA mobility is greater when compared with Agarose, regular use. Electrophoresis times are reduced depending upon buffer and agarose concentration used.
- Easy preparation of the gel by simple dissolution in aqueous buffers either by standard boiling or microwaving.
- Greater thermal stability due to high hysteresis (difference between gelling and melting temperatures).
- Exceptionally low absorption of staining agents.
- Absence of toxicity (the alternative is polyacrylamide which can be toxic).

Ordering information

Catalog size quantities and prices may be found at <u>http://www.interchim.com</u>. Please inquire for higher quantities (availability, shipment conditions).

For any information, please ask : Uptima / Interchim; Hotline : +33(0)4 70 03 73 06

Order on-line or Contact your local distributor

Disclaimer: Materials from Uptima are sold **for research use only**, and are not intended for food, drug, household, or cosmetic uses. Uptima is not liable for any damage resulting from handling or contact with this product.

