

Product Specification



Material 35-1020 & 443666A Material description Agarose, universal

Grade peqGOLD molecular biology grade

ELECTRAN® - DNA pure grade - Electrophoresis grade

CAS Number 9012-36-6 Molecular formula (CIHIII OI)n

Characteristics	Specifications
Performance test	Passes test
Appearance:	
Fine white homogeneous powder	Passes test
DNases	None detected
RNases	None detected
Clarity	≤ 4.00 NTU
Gel strength (1 %; water)	≥ 1000 g/cm ²
Gel strength (1.5 %; water)	≥ 2000 g/cm²
Gelling temperature (1.5 %; water)	34.0 - 37.5 °C
Melting point (1.5 %; water)	86.0 - 90.0 °C
pH (1.5 %; in gel)	6.00 - 9.00
pH (1.5 %; in solution)	6.00 - 9.00
Ignition residue	≤ 1.0 %
Water	≤ 10.0 %
SOI (Sulphate)	≤ 0.15 %

Signature

We certify that this batch conforms to the specifications listed

This document has been produced electronically and is valid without a signature.

Anja Vanhalle, Head of Laboratory - Haasrode VWR International bv; Geldenaaksebaan 464; BE-3001 Leuven; Belgium

For Professional use in Laboratory or Manufacturing. Not for use as an Active Pharmaceutical Ingredient or Food or Animal Feed. Suitability and intended use of the product remains the responsibility of the user



Manual - Universal Agarose

Introduction:

peqGOLD Universal Agarose is ideally suited for use as standard agarose for analytical as well as preparative nucleic acid electrophoresis of fragments from 50 bp to 50 kbp. Even at low concentrations the gel produced is very firm.

Separation range:

DNA: approx. 0.05 kbp – 50 kbp RNA: approx. 0.30 kb – 20 kb

Product numbers:

35-1010 - Universal-Agarose, 100 g 35-1020 + 443666A - Universal-Agarose, 500 g

Quality:

Molecular Biology Grade, DNA pure grade

ELECTRAN® - Electrophoresis grade

Certified free of DNases and RNases

No DNA binding

High lot-to-lot consistency

Characteristics:

High separation properties and sharp band patterns

Easy solubility without foaming

Excellent optical transparency

Analytical specifications:

• Gelling temperature: 34.0 ≤ 37.5

°C

Melting temperature: 86 ≤ 90 °C
Gel strength (1.5 %): ≥ 2000 g /

cm2

Sulphate content: ≤ 0.15 %
Water content: ≤ 10.0 %

Safety instructions:

Always wear eye protection when preparing agarose gel solutions and protect yourself and others against boiling liquids. Refer to the material safety data sheet for further safety and handling instructions.

- Manufactured and quality-controlled in accordance with ISO 9001:2000
- Shipment at ambient temperatures
- Storage at room temperature
- With appropriate storage, stable for minimum 2 years

Note:

For preparing 100 ml of 1% agarose gel solution use 1 g agarose in 100 ml appropriate electrophoresis buffer.

Preparing agarose:

Method 1: Microwave oven

- 1. Pour buffer (approx. 90 % of final volume) into an appropriate flask that can accommodate up to four times the final gel volume and add a magnetic stir bar.
- 2. Put the flask onto a magnetic stirrer and slowly add agarose powder while stirring constantly to avoid clotting.
- 3. Remove magnetic stir bar.
- 4. Add remaining buffer up to the desired final volume.
- 5. Weigh and record the weight of the flask prior to heating. Heat for 1-2 minutes in a microwave oven (600 Watt). Gently swirl the flask to mix the solution. Warning: there may be a delay in the liquid boiling!
- 6. Using the microwave oven, heat in short bursts of 5-10 seconds or until the solution is boiling, with breaks of 10-15 seconds between heating phases to disperse bubbles by gently swirling the flask. Beware of hot glass ware and liquid. Continue until the agarose is completely dissolved.
- 7. Again, weigh the flask and top up lost volume with warm, deionized water. Gently swirl the flask to ensure complete mixing.
- 8. Let the solution cool down at room temperature for 15-20 minutes or until a gel temperature of 50-60 °C is reached.

Method 2: Simmering water bath

- 1. see method 1: steps 1 2 and 4
- 2. Weigh and record the weight of the flask prior to heating. Heat agarose suspension up in a simmering water bath with constant stirring.
- 3. Leave the flask in the water bath for further 15 20 minutes, or until the agarose is completely dissolved.
- 4. Switch off the magnetic stirrer and leave the flask in the bath for further 15 minutes.
- 5. Again, weigh the flask and top up lost volume with warm, deionized water. Gently swirl the flask to ensure complete mixing.
- 6. Let the solution cool down at room temperature for 15-20 minutes or until a gel temperature of 50-60 °C is reached.