

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

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Panel 1: Mouse C57BL/6 Major Tissue Premade Northern Blot

Catalog #: MN-MT-1-C57

Quantity: 1 blot

Storage Conditions: Store in a sealed bag at room temperature or 4°C. It is good for several months.

Applications: Mouse C57BL/6 Major Tissue Premade Northern Blot

- Analyze gene expression pattern in 19 different mixed (multiple) tissues simultaneously.
- Determine size and relative abundance of specific messages in the mixed tissues.
- Identify alternative spliced forms.

Quality Control: Every step of preparation of the blot, from harvesting tissues and extraction of RNA to the blotting, is carefully monitored to ensure the superior quality and performance.

- Blot is made from total RNA that is treated with RNase-free DNase to remove residual DNA. The purity and integrity of RNA are tested by formaldehyde-denatured agarose gel electrophoresis.
- The efficiency of transfer is checked by staining gel with ethidium bromide.
- The integrity of the blotted RNA is tested by beta-actin specific probe.

Description: Total RNA (20 μg each) was extracted from freshly harvested mouse tissues, fractionated through formaldehyde-denatured agarose gel, transferred to a positively charged nylon membrane, and cross-lined by UV light.

The band sizes of the RNA marker (major bands: 10kb, 6 kb, 4 kb, 3 kb, 2 kb, 1.5 kb, 1 kb, 0.5 kb, and 0.2 kb) are marked on the membrane.

Tissues on the blot: RNAs were loaded in the following left to right order: RNA Marker (1) Brain (whole), (2) Liver (whole), (3) Pancreas, (4) Spleen, (5) Heart, (6) Lung, (7) Esophagus, (8) Stomach, (9) Intestine, (10) Colon, (11) Skeletal Muscle, (12) Kidney, (13) Thymus, (14) Testis, (15) Prostate, (16) Ovary, (17) Placenta, (18) Mammary, (19) Skin

Hybridization: Prior to use, wet blot in water for few minutes. Northern blots can be hybridized using isotopic and non-isotopic DNA or RNA probes. You can follow any general protocol of Northern blot analysis for hybridization, post-washing, and stripping (reusing blot) probe. However, we recommend protocols published in the following books: "Short protocols in molecular biology" by Frederick Ausubel et al. and "RNA Methodologies" by Robert Farrell.

Stripping Probe: Blot is reusable for several times, just strip probe and re-hybridize with a new one. Keep blot wet until the previously hybridized probe has been removed. It is extremely difficult to completely remove hybridized probe from a membrane that has been allowed to dry.

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