# RNA and Protein from one Lysate

undivided samples parallel isolation

# NucleoSpin® RNA/Protein

### Rapid Purification of total RNA and Protein from Cells and Tissue

Studies of gene expression on transcriptional and translational level are often complicated by small sample sizes and incompatible techniques for RNA and protein isolation.

The NucleoSpin® RNA/Protein Kit enables the parallel isolation of RNA and protein from one lysate and a broad variety of starting materials.

### Procedure Homogenization of sample Cell lysis Filtration of lysate NucleoSpin® Filters Bind RNA/DNA RNA/DNA bound to silica membrane Protein in column flow-through **Protein Purification RNA Purification** Precipitate Protein Membrane desalting (Special Protein Precipitator) **DNase I incubation** directly on the membrane at RT Wash Protein Pellet Wash RNA Redissolve pellet in PLB Elute highly pure RNA (Protein Loading Buffer) SDS-PAGE/ All typical downstream applications Western blot

## **Features**

✓ RNA and Protein from one lysate simple and fast procedure: no phenol, no chloroform, no acetone



High quality RNA

RNA is suitable for all common downstream applications e.g. RT-PCR, TaqMan®, blotting, or microarray

High Protein yield

high protein concentration, suitable for SDS-PAGE and Western blot analysis

✓ Parallel DNA purification possible

parallel DNA purification is possible in combination with the optional NucleoSpin® RNA/DNA buffer set



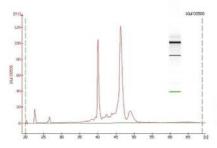
# **RNA and Protein from one Lysate**

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# **Applications**

✓ gene expression profiling on transcription and translation levels

## **Application data**



# High quality of RNA

#### Sample material:

106 HeLa cells

Elution was done using 100 µl RNase free water, 10 µl were analyzed on Agilent Bioanalyzer according to the standard protocol.

High quality of RNA proven by Bioanalyzer analysis!

## Quantitative **Protein Isolation**

Sample material: A: 106 HeLa cells

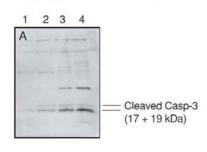
B: 30 mg liver

C: 100 mg garden cress seedlings

#### Just 1.4 % of the total isolated protein loaded per lane!

This corresponds to protein from 14 000 HeLa cells (A), 0.43 mg liver (B), and 1.43 mg garden cress seedling (C), respectively per lane.

NucleoSpin® RNA/Protein procedure results in sufficient protein for SDS PAGE analysis



Erk2 (41 kDa)

Expression analysis of cleaved caspase-3 and Erk2 in carcinoma cell lines upon treatment with an apoptosis inducing DNA damaging agent

Sample material: carcinoma cell line Sample amount: approx. one million cells

Precipitated volume of column flow-through for protein isolation: 200 µl

Protein resolubilization volume: 200 µl PLB Sample volume loaded per lane: 16 µl

A: Western-blot probed with anti-cleaved caspase-3

B: Western-blot probed with anti-Erk2

- 1: untreated
- 2: 24 h upon treatment
- 3: 48 h upon treatment
- 4: 120 h upon treatment

Data was kindly provided by Steffen Naumann and Prof. B. Kaina, Department of Toxicology,

University of Mainz, Germany

#### Visualization of changes in protein level possible!

#### Ordering Information:

Product	Cat. No.	Distributed by:	0.2005
NucleoSpin® RNA/Protein (10/50/250 preps)	740 933.10 / .50 / .250		NuSpinRNA/Protein 1 Printed in Garmany
NucleoSpin® RNA/DNA buffer set (100 preps)	740 944		

For more information regarding the use of MN products, please contact your local representative or visit MN directly: www.mn-net.com.

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