

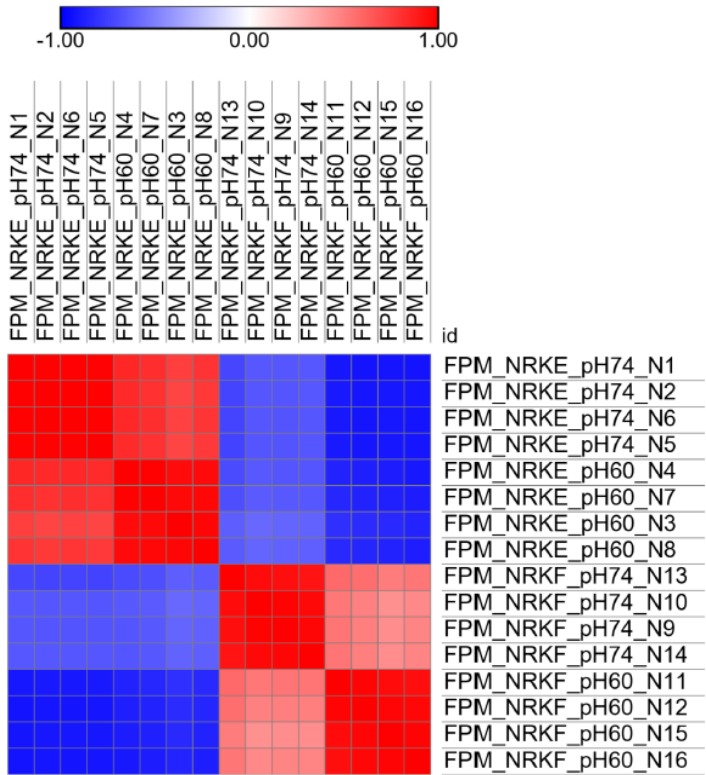
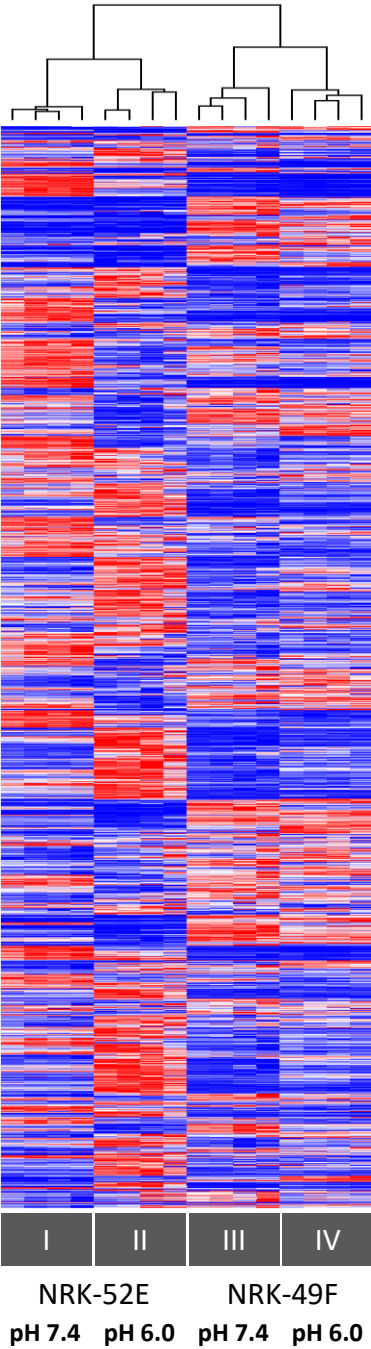
Supplementary figures

Supplementary figure S1

<https://software.broadinstitute.org/morpheus/>:

Clustering:
Euclidian distance
Linkage method = average

Similarity matrix:
Pearson correlation



all, FPM>10, 11122 genes

Supplementary Table S1. Buffer composition

EDTA buffer	<ul style="list-style-type: none">• 136.8 mM NaCl
	<ul style="list-style-type: none">• 2.68 mM KCl
	<ul style="list-style-type: none">• 8.1 mM Na₂HPO₄ x 2 H₂O
	<ul style="list-style-type: none">• 2 mM KH₂PO₄
	<ul style="list-style-type: none">• 0,7 mM EDTA
	<ul style="list-style-type: none">• pH 7,2
MOPS-Triton buffer	<ul style="list-style-type: none">• 10 mM TRIS base
	<ul style="list-style-type: none">• 20 mM MOPS
	<ul style="list-style-type: none">• 100 mM NaCl
	<ul style="list-style-type: none">• 1 mM EDTA
	<ul style="list-style-type: none">• 0.01 % Triton X-100
	<ul style="list-style-type: none">• pH adjusted to 7.5
6xRedmix/Lämmli buffer	<ul style="list-style-type: none">• 124.8 mM TRIS HCl (AppliChem GmbH, Darmstadt, GER)
	<ul style="list-style-type: none">• 6 % SDS (Carl Roth GmbH&CoKG, Karlsruhe, GER)
	<ul style="list-style-type: none">• 1,42 mM β-Mercaptoethanol (Sigma-Aldrich Co, St. Louis, USA)
	<ul style="list-style-type: none">• 4,7 mM Glycerol (Sigma-Aldrich Co, St. Louis, USA)
	<ul style="list-style-type: none">• 155.2 mM Bromphenol Blue (SERVA Electrophoresis GmbH, Heidelberg, GER)
	<ul style="list-style-type: none">• pH 6.8
running buffer	<ul style="list-style-type: none">• 25 mM TRIS

	<ul style="list-style-type: none"> • 3,5mM SDS • 192mM glycine
	<ul style="list-style-type: none"> • 20 % MeOH
transfer buffer	<ul style="list-style-type: none"> • 25mM Tris • 192mM glycine
	<ul style="list-style-type: none"> • 154mM NaCl • 2.7 mM KCl • 8.2mM Na₂HPO₄ • 1.5mM KH₂PO₄
trypsin solution	<ul style="list-style-type: none"> • 0,7mM EDTA • 0,2mM streptomycin • 0,18mM penicillin • 0.02mM trypsin • pH7,1-7,3
	<ul style="list-style-type: none"> • 3mM TRIS base • 140mM NaCl
1xTBS TWEEN	<ul style="list-style-type: none"> • 0.17mM TRIS-HCl • 1 % TWEEN 20 • pH7.4
equilibration buffer	<ul style="list-style-type: none"> • 10 mM TRIS (pH 7.3)

	<ul style="list-style-type: none">• 10 mM KCl• 1.5 mM MgCl• 0.5 mM β-Mercaptoethanol• 1 : 100 protease inhibitor
lysis buffer	<ul style="list-style-type: none">• equilibration buffer• 0.4 % nonidet P40
	<ul style="list-style-type: none">• 20 mM HEPES• 0.4 M NaCl
extraction buffer	<ul style="list-style-type: none">• 1 mM EDTA• 1 mM DTT• 1 : 100 protease inhibtor

Supplementary Table S2. Primer sequences and annealing temperatures

Gene name	Accession number	Forward 5'-3'	Backward 5'-3'	Annealing °C
<i>Hprt1</i>	NM_012583.2	ACCAGTCAACGGGG- GACATA	TTGGGGCTG- TACTGCTTGAC	60 °C
<i>Nrf2</i> (<i>Nef2l2</i>)	NM_031789.2	ACTACAGTCCCAG- CAGGACA	GTTTGG- GAATGTGGG- CAACC	60 °C
<i>Tkt</i>	NM_022592.2	TCCATACCATGCGC- TACAAG	ATAAGATGG- GAGCTGCATGG	60 °C

Abbreviations: *Hprt* - Hypoxanthine Phosphoribosyltransferase 1, *Nrf2* (*Nef2l2*) - Nuclear factor erythroid 2-related factor 2, *Tkt* - transketolase