SIRT1 confers protection against UVB- and H_2O_2-induced cell death via modulation of p53 and JNK in cultured skin keratinocytes

Cong Cao a, b, #, Shan Lu a, #, Rebecca Kivlin a, Brittany Wallin a, Elizabeth Card a, Andrew Bagdasarian a, Tyrone Tamakloe a, Wen-jun Wang c, * Xiuze Song d, Wen-ming Chu b, Nicola Kouttab e, Aie Xu d, Yinsheng Wan a, *

a Department of Biology, Providence College, Providence, RI, USA
b Department of Molecular Microbiology and Immunology, Brown University, Providence, RI, USA
c Department of Chemical and Biochemical Engineering, Zhejiang University, Hangzhou, China
d Department of Dermatology, The 3rd Hospital of Hangzhou, Hangzhou, China
e Department of Pathology, Roger Williams Medical Center, Boston University, Providence, RI, USA

Received: May 23, 2008; Accepted: July 29, 2008

Abstract

SIRT1 is a member of a highly conserved gene family (sirtuins) encoding nicotinamide adenine dinucleotide (NAD)^+ -dependent deacetylases, originally found to deacetylate histones leading to increased DNA stability and prolonged survival in yeast and higher organisms, including mammals. SIRT1 has been found to function as a deacetylase for numerous protein targets involved in various cellular pathways, including stress responses, apoptosis and axonal degeneration. However, the role of SIRT1 in ultraviolet (UV) signalling pathways remains unknown. Using cell culture and Western blot analysis in this study we found that SIRT1 is expressed in cultured human skin keratinocytes. Both UV radiation and H_2O_2, two major inducers of skin cell damage, down-regulate SIRT1 in a time- and dose-dependent manner. We observed that reactive oxygen species-mediated JNK activation is involved in this SIRT1 down-regulation. SIRT1 activator, resveratrol, which has been considered as an important antioxidant, protects against UV- and H_2O_2-induced cell death, whereas SIRT1 inhibitor such as sirtinol and nicotinamide enhance cell death. Activation of SIRT1 negatively regulates UV- and H_2O_2-induced p53 acetylation, because nicotinamide and sirtinol as well as SIRT1 siRNA enhance UV- and H_2O_2-induced p53 acetylation, whereas SIRT1 activator resveratrol inhibits it. We also found that SIRT1 is involved in UV-induced AMP-activated protein kinase (AMPK) and downstream acetyl-CoA carboxylase (ACC), phosphofructose kinase-2 (PFK-2) phosphorylation. Collectively, our data provide new insights into understanding of the molecular mechanisms of UV-induced skin aging, suggesting that SIRT1 activators such as resveratrol could serve as new anti-skin aging agents.

Keywords: SIRT1 • UV • p53 • keratinocytes • apoptosis • skin aging

Introduction

The silent information regulator (SIR) family of genes is a highly conserved group of genes that encode nicotinamide adenine dinucleotide (NAD)-dependent protein deacetylases, also known as class III histone deacetylases. The best characterized of these genes is Saccharomyces cerevisiae SIR2, which is involved in silencing of mating type loci, telomere maintenance, DNA damage response and cell aging [1]. SIRT1, the mammalian orthologue of SIR2, is a protein implicated in regulation of many cellular processes, including apoptosis, cellular senescence, endocrine signalling, glucose homeostasis, aging and longevity [2, 3].

Cellular targets of SIRT1 include acetylated p53 [2, 3], p300 [4], Ku70 [5], forkhead (FOXO) transcription factors [5, 6], PPAR_γ [7] and PPAR_γ coactivator-1α (PGC-1α) protein [7, 8]. Deacetylation of p53 transcription factors and FOXO represses apoptosis and increases cell survival [2, 4–6]. Deacetylation of Ku70 [9] and
namide and sirtinol as well as SIRT1 siRNA enhances UV- and UV/ROS-induced skin cell damage. Support the notion that SIRT1 might be a novel target protecting deacetylates p53 after UV and H2O2 treatment, because nicotinamide as well as SIRT1 RNAi enhance apoptosis. Activation of SIRT1 death, whereas SIRT1 inhibitors such as sirtinol and nicotinamide are extensively studied. However, the possible involvement of signals are involved in this apoptotic process, of which, reactive oxygen species (ROS) production, p53 [10], p38 [11], JNK [11, 12] are extensively studied. However, the possible involvement of SIRT1 in UV-induced skin cell damage is not fully studied.

Given that SIRT1 plays important roles in cellular apoptosis and cell survival, we undertook this study to investigate the role of SIRT1 in UV-induced cellular damage. In this study, we found for the first time that SIRT1 is functionally expressed in cultured skin keratinocytes. Both UV and H2O2, two major factors of skin cell damage, down-regulate SIRT1. ROS-mediated JNK activation is involved in UV-induced SIRT1 down-regulation. SIRT activator, resveratrol protects against UV- and H2O2-induced apoptotic cell death, whereas SIRT1 inhibitors such as sirtinol and nicotinamide as well as SIRT1 RNAi enhance apoptosis. Activation of SIRT1 deacetylates p53 after UV and H2O2 treatment, because nicotinamide and sirtinol as well as SIRT1 siRNA enhances UV- and H2O2-induced p53 acetylation. Our study provides evidence to support the notion that SIRT1 might be a novel target protecting UV/ROS-induced skin cell damage.

Materials and methods

UVB light apparatus

As previously reported [13–15], UV-irradiation apparatus used in this study consisted of four F36T12 EREVHO UV tubes. A Kodacel TA401/407 filter (International Light Inc., Newburyport, MA, USA) was mounted 4 cm in front of the tubes to remove wavelengths <290 nm. Irradiation intensity was monitored using an IL443 phototherapy radiometer and a SED240/UV/W photodetector. Before UV irradiation, cells were washed with 1 ml phosphate buffered saline (PBS) and changed to fresh 0.5 ml PBS each well. Cells were irradiated at the desired intensity without plastic dish lid. After UV irradiation, cells were returned to incubation in basal medium with treatments for various time-points prior to harvest.

Cell viability assay (MTT dye assay)

Cell viability was measured by the 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide (MTT) method [15]. Briefly, cells were collected and seeded in 96-well plates at a density of 10^3 cells/ml. Different seeding densities were optimized at the beginning of the experiments (data not shown). After incubation for 24 hrs, cells were exposed to fresh medium containing reagents at 37°C. After incubation for up to 24 hrs, 20 μl of MTT tetrazolium salt (Sigma) dissolved in Hank’s balanced solution at a concentration of 5 mg/ml was added to each well and incubated in CO2 incubator for 4 hrs. Finally, the medium was aspirated from each well and 150 μl of DMSO (Sigma) was added to dissolve formazan crystals and the absorbance of each well was obtained using a Dynatech MR5000 plate (Dynatech Laboratories, Alexandria, VA, USA) reader at a test wavelength of 490 nm with a reference wavelength of 630 nm.

Assessment of the percentage of apoptotic cells

To detect apoptotic cells [15], cells were stained with DNA binding dye Hoechst 33342 (Sigma). After the cells were exposed to UV and the test purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA). Monoclonal mouse anti-p-akt, anti-SIRT1, sirtinol, nicotinamide, resveratrol and Hoechst33342 were obtained from Sigma (St. Louis, MO, USA). Phospho-AKT (Ser473), phospho-ERK (Tyr1068), phospho-JNK (Thr183/185), phospho-AMPK (Thr172), p-acetyl-CoA carboxylase (ACC) (Ser 79) phospho-p38 (Thr180/Tyr182), phosphofructose kinase-2 (PFK-2) (Ser466), SAPK/JNK, p38 antibody and AKT antibody were all from Cell Signaling Technology (Beverly, MA, USA).
was determined by Western blot after 48 hrs. of UV treatment. The final SIRT1 siRNA concentration of 150 nM. SIRT1, protein expression

diluted in 90
dition may be involved in UV- and H2O2-induced skin cell damage. Furthermore, H2O2 also induces SIRT1 down-regulation in a dose (Fig. 1E and F) and a time (Fig. 1G and H) dependent manner. These results demonstrate that both UV radiation and H2O2 down-regulate SIRT1 expression, suggesting that SIRT1 down-regulation may be involved in UV- and H2O2-induced skin cell damage.

ROS-mediated JNK activation is involved in UV- and H2O2-induced SIRT1 down-regulation

The above data showed that UV radiation and H2O2 induce SIRT1 down-regulation in cultured human skin keratinocytes, and yet cell signal transduction pathways involved in this process remain unclear. Mitogen-activated protein kinase (MAPK) and PI3K/AKT pathways are known to mediate UV-induced cellular events leading to photoaging [10, 18, 19]. To investigate whether those signalling pathways are also involved in UV-induced SIRT1 down-regulation, various pharmacological inhibitors were utilized in our experiments. Although inhibitors of p38 (SB 203580), MEK/ERK (PD 98059 and U0126) and PI3K/AKT (LY 294002 and Wortmannin) have no effects on UV- and H2O2-induced SIRT1 down-regulation (data not shown), JNK inhibitor (SP 600125, 1 μm, or JNKi) attenuates SIRT1 down-regulation (Fig. 2A–D). This result suggests that JNK activation is involved, at least in part, in UV- and H2O2-induced SIRT1 down-regulation. To further investigate the role of ROS in SIRT1 down-regulations, cells were pre-treated with antioxidants NAC (n-acetyl-l-cysteine). The results showed that NAC protects against UV- and H2O2-induced loss of SIRT1 (Fig. 2E–H). As expected, NAC pre-treatment inhibits UV-induced ROS production (Fig. 2I) and JNK activation (Fig. 2J). Collectively, our data suggest that ROS-mediated JNK activation is involved in UV- and H2O2-induced SIRT1 down-regulation.

SIRT1 modulates UV-induced JNK activation

Because JNK MAPK, principally activated by ROS, mediates UV-induced cell death in keratinocytes, we next examined whether SIRT1 could affect UV-induced JNK phosphorylation. The results showed that treatment of keratinocytes with UV (30 mJ/cm²) leads to a rapid and time-dependent activation of MAPKs (JNK, ERK and p38). Pre-treatment with SIRT1 inhibitors such as sirtinol and nicotinamide, which alone has little effect on MAPK activation (data not shown), enhances UV-induced JNK phosphorylation (Fig. 3A), but has no effects on UV-induced AKT activation, which is known as a pro-survival signal (Fig. 3A and B). To further
confirm the role of SIRT1 in UV-induced JNK activation, SIRT1 siRNA specific knockdown was used. As shown in Fig. 3C, SIRT1 siRNA knockdowns SIRT1 expression in HaCaT cells. Furthermore, SIRT1 siRNA knockdown enhances UV-induced JNK activation, whereas SIRT1 activator resveratrol inhibits it (Fig. 3D and E). These data suggest that SIRT1 inhibits UV-induced JNK activation.

**SIRT1 negatively regulates UV- and H2O2-induced p53 acetylation**

Previous studies have indicated that SIRT1 may function to promote cell survival [3, 20, 21] via direct interactions with several apoptotic proteins, including p53 [3]. We next tested the possible role SIRT1 in UV-induced JNK activation. As shown in Fig. 4A–D, UV-induced p53 acetylation is enhanced by SIRT1 inhibitor sirtinol and nicotinamide (Nico). Furthermore, pre-treatment with SIRT1 activator resveratrol (Rev) almost abolishes sirtinol or Nico’s effects. Similar results are also seen in H2O2-treated cells (Fig. 4E and F). To further confirm the involvement of SIRT1 in UV- and H2O2-induced p53 acetylation, wild-type and p53 knockout MEFs were used in our experiments. As shown in Fig. 4G and H, Nico enhances UV- and H2O2-induced p53 acetylation in wild-type but not in p53 knockout MEFs. Next we tested p53 acetylation in SIRT1 knockdown HaCaT cells. As shown in Fig. 4I, SIRT1 siRNA knockdown enhances UV- and H2O2-induced p53 acetylation in HaCaT cells. Collectively, our results suggest that SIRT1 negatively regulates p53 acetylation in UV- and H2O2-treated skin keratinocytes.

**SIRT1 positively regulates UV-induced AMPK activation**

Some of the metabolic changes caused by resveratrol, a SIRT1 activator, mimic those observed in response to AMPK activation, and our data in this study demonstrated for the first time that UV
Fig. 2 ROS-mediated JNK activation is involved in UV- and H2O2-induced SIRT1 down-regulation. HaCaT cells were pre-treated with JNK inhibitor (SP 600125, 1 μM, or JNKi) for 1 hr, followed by 20 mJ/cm² UV radiation (A and B) or 250 μM of H2O2 (C and D) and incubated for 24 hrs. SIRT1 and β-actin expression were detected by Western blot. HaCaT cells were pre-treated with anti-oxidant NAC (n-acetyl-l-cysteine) (NAC, 400 μM) for 1 hr, followed by 20 mJ/cm² UV radiation (E and F) or 250 μM of H2O2 (G and H) and incubated for 24 hrs, SIRT1 and β-actin expression were detected by Western blot. HaCaT cells were pre-treated with NAC for 1 hr, followed by 20 mJ/cm² of UV radiation or 250 μM of H2O2 and incubated for 1 hr (I), ROS production was detected by FACS as described in methods. HaCaT cells were pre-treated with NAC (400 μM) or JNK inhibitor (SP 600125, 1 μM, or JNKi) for indicated time-points, p-JNK and β-actin were detected by Western blot. The data in figures represent mean ± S.E. of three independent experiments. The symbol ‘*’ means P < 0.05 with untreated group. The symbol ‘#’ means P < 0.05 with UV- or H2O2-treated group.
radiation induces AMPK activation in cultured skin keratinocytes (Fig. 5A and B). We next tested the possible role of SIRT1 in AMPK activation. Our results showed that SIRT1 inhibitors sirtinol and nicotinamide inhibit UV-induced AMPK (Fig. 5A and B) and downstream PFK-2 and ACC phosphorylation (Fig. 5C and D). Furthermore, SIRT1 activator resveratrol alone also induces AMPK activation, and this induction is largely impaired by SIRT1 inhibitor (nicotinamide, or Nico) or AMPK inhibitor (compound C, or AMPKi) (Fig. 5E and F). Collectively, our data suggest that SIRT1 positively regulates AMPK activation in response to UV and resveratrol. At least some of the actions of resveratrol, such as fatty acid oxidation, are mediated by AMPK activation.

Fig. 3 SIRT1 regulates UV-induced JNK activation. HaCaT cells were pre-treated with nicotinamide (Nico, 10 mM) or sirtinol (2 mM) for 1 hr, followed by UV radiation (25 mJ/cm²) and then incubated in DMEM for 0.5, 1.0 and 2.0 hrs, p-JNK, p-ERK, p-p38, p-AKT (Ser 473) and T-p38 were detected by Western blot (A) and JNK phosphorylation was quantified in (B). HaCaT cells with or without SIRT1 siRNA were pre-treated with resveratrol (Rev, 10 μM) for 1 hr, followed by UV radiation for indicated time-points, p-JNK and T-JNK were detected by Western blot. HaCaT cells were treated with 150 nM of SIRT1 siRNA or controls for 48 hrs, SIRT1 and β-actin were detected by Western blot (C). (D) Control or SIRT1 siRNA (150 nM) pre-treated cells were treated with UV radiation (25 mJ/cm²) for indicated time along with or without resveratrol (Rev, 10 μM, 1 hr prior to UV radiation), p-JNK and T-JNK were detected by Western blot. JNK phosphorylation was quantified in (E). The data in figures represent mean ± S.E. of three independent experiments. The symbol ‘#’ means P < 0.05 with UV-treated group.
SIRT1 protects against UV-radiation-induced cell death

To further test the role of SIRT1 in UV-induced cell death, we pre-treated cells with various SIRT1 inhibitors and activators and then exposed the cells with UV radiation. As shown in Fig. 6A and B, as expected, UV radiation induces HaCaT cell death in a dose-dependent manner, whereas pre-treatment with resveratrol protects against UV-induced cell death and apoptosis, nicotinamide and sirtinol enhance this process. Furthermore, nicotinamide and sirtinol pre-treatment aggravate UV-induced Bcl-xl degradation,
Fig. 5 SIRT1 positively regulates AMPK activation in cultured skin keratinocytes. HaCaT cells were pre-treated with sirtinol (2 mM) or nicotinamide (Nico, 10 mM) for 1 hr, followed by 20 mJ/cm² of UV for indicated time, p-AMPK (Thr 172) and total-AMPK activation were detected by Western blot (A and B). HaCaT cells were pre-treated with sirtinol (2 mM) or nicotinamide (Nico, 10 mM) for 1 hr, followed by 20 mJ/cm² of UV and incubated for 0.5 and 2.0 hrs, p-ACC (Ser 79), p-PFK-2 (Ser 466) and β-actin were detected by Western blot (C), p-ACC was quantified in (D). HaCaT cells were treated with nicotinamide (Nico, 10 mM) or AMPK inhibitor Compound C (AMPKi, 10 μM) for 0.5, 2.0 hrs, followed by resveratrol (Rev, 10 μM) treatment for 0.5, 1.0 and 2.0 hrs, p-AMPK (Thr 172), p-ACC (Ser 79) and T-AMPK were detected by Western blot (E and F). The data in figures represent mean ± S.E. of three independent experiments. The symbol # means P < 0.05 with UV- or H₂O₂-treated group.
whereas resveratrol delays the process (Fig. 6C). Next we tested
the possible role of p53 in SIRT1-induced protective effects using
p53 knockout cells. As demonstrated in Fig. 6D, p53 knockout
MEFs are resistant to UV-induced cell death and SIRT1-induced
protective effects are almost abolished in p53 knockout MEFs. To
furthermore confirm these protective effects, SIRT1 siRNA was
used in our experiments. As shown in Fig. 6E, SIRT1 knockout
HaCaT cells were more sensitive to UV-induced cell death. Taken
all together, our data demonstrated that SIRT1 protects against
UV-induced cell death, at least in part, via modulation of p53.

Discussion

In response to UV radiation, p53 tumour suppressor is activated
and exerts anti-proliferative effects, including growth arrest, apo-
tosis, and cell senescence [22]. Following DNA damage, p53 pro-
tein is protected from rapid degradation and acquires transcrip-
tion-activating functions, largely as a result of post-translational
modifications [23]. Activation of p53 protein as a transcriptional factor allows it, in turn, to up-regulate the expression of genes
whose products promotes cell cycle exit, such as p21WAF1 gene
[24], or of genes that favour apoptosis [25]. The p53 protein is
phosphorylated in response to DNA damage by ATM at residue
Ser15 [26] and at residue Ser20 by Chk1/2 kinases [26].

However, recent studies suggest that Ser15 phosphorylation
does not lead directly to the functional activation of p53 protein.
Instead, it increases the affinity of the p300 acetylase for p53 [27].
This association leads to the acetylation of p53. Indeed, p53 is
acetylated in vitro by p300 at Lys 370–373, 381, and 382 [28].
Moreover, at least two of these sites, namely residues 320 and
382, are found to be acetylated in vivo in response to DNA dam-
age [29]. Among other factors that can affect acetylation of p53
are MDM2 protein and SIRT1, which are involved in the negative
regulation of p53 [30] and are able to block acetylation of p53 pro-
tein by p300 [31].

SIRT1 is a member of a highly conserved gene family (sirtuins)
encoding NAD+-dependent deacetylases. SIRT1 remains one of
the most important cell signalling molecules that are associated
with cell survival and longevity [2, 3]. In this study, we have evi-
dence showing that SIRT1 plays protective role in UV-induced skin
cell damage. We found that UV radiation and H2O2 induce p53
acetylation in cultured skin keratinocytes and MEFs cells (Fig. 4),
SIRT1, as a deacetylase, negatively regulate UV-induced p53
acetylation (Fig. 7A), because SIRT1 inhibitors sirtinol and nicoti-
namide as well as SIRT1 siRNA enhance UV-induced p53 acetyla-
tion, whereas SIRT1 activator resveratrol enhances it (Fig. 4).
Considering recent study showing that acetylation is indispensa-
able for p53 activation, we suggest that SIRT1 protects from UV-
induced cell death, at least in part, by negatively regulating p53
acetylation (Fig. 6D).

Previous studies have demonstrated that UV radiation induces
down-regulation of a number of cellular proteins such as collagen
and water channel protein aquaporin-3 or AQP-3 in both skin ker-
atinocytes and fibroblasts [19, 32]. Interestingly, in this study, we
observed that UV radiation also induces down-regulation of SIRT1
in cultured human skin keratinocytes (Fig. 1). Given the important
functions of SIRT1 discussed above, one can easily envision the
consequence of SIRT1 down-regulation in response to UV radia-
tion. Systematic studies revealed that ROS-mediated JNK activa-
tion is involved in SIRT1 down-regulation in response to UV radia-
tion. JNK inhibitor and antioxidant NAC could recover SIRT1 lost
due to UV radiation (Figs. 2 and 7B). Interestingly, SIRT1 inhibits
UV-induced JNK activation (Figs. 3 and 7C). Collectively, these
data suggest that SIRT1 serves as a negative regulator against UV-
induced JNK activation, probably by de-acetylation and inhibition
of one or more of JNK upstream signals, which may serve as
another mechanism to protect against UV-induced cell death.
However, the detailed mechanisms through which SIRT1 nega-
tively regulates JNK activation needs further investigation. Our
data have provided more insights into understanding of the
molecular mechanism through which resveratrol, the SIRT1 acti-
ator, acts as an important anti-aging agent. Our data may also
help to develop better cosmetics products against UV-induced
human skin photaging and even skin cancer.

Our data also suggest that SIRT1 positively regulates AMPK
activation in response to UV and resveratrol (Fig. 5). At least
some of the actions of resveratrol, such as fatty acid oxidation,
are mediated by AMPK activation [33]. These results are consis-
tent with previous studies which demonstrated that resveratrol

Fig. 6 SIRT1 protects against UV-induced skin cell damage. HaCaT cells were pre-treated with resveratrol (Rev, 10 μM), sirtinol (2 mM), nicotinamide (Nico, 10 mM), Rev + Nico or Rev + sirtinol for 1 hr, followed by 10, 20 or 30 mJ/cm² of UV radiation and incubated in DMEM for 24 hrs, cell viability was detected by MTT assay (A). HaCaT cells were pre-treated with nicotinamide, resveratrol, or Rev + Nico for 1 hr, followed by 20 mJ/cm² of UV radiation and incubation in DMEM for 24 hrs, cell apoptosis was detected by Hoechst assay (B). HaCaT cells were treated with indicated treatments for 24 hrs, Bcl-xl and β-actin were detected by Western blot (C). Wild-type and p53 knockout MEFs were treated with UV, UV + Nico or UV + Rev and incubated in DMEM for 24 hrs, cell viability was detected by MTT assay (D). Control or SIRT1 siRNA pre-treated cells were treated with UV (20 mJ/cm²) or UV + Nico (Nico, 10 mM) for 24 hrs, cell viability were detected by MTT assay (E). The data in figures represent mean ± S.E. of three independent experiments. The symbol * means P < 0.05 with UV- or H2O2-treated group, the symbol ** means P < 0.05 with UV plus Nico or sirtinol treated group, the symbol *** means P < 0.05 with p53 knockout or SIRT1 siRNA group.
acts as an activator of AMPK in both neuron and whole brain [33], and the most recent study which indicated that AMPK activation might be involved in resveratrol’s calorie resistant effect [34]. However, the detailed mechanism through which resveratrol activates AMPK and the biological function of these effects warrant further investigation.

In summary, we found for the first time that SIRT1 is functionally expressed in cultured skin keratinocytes. Both UV and H$_2$O$_2$, two major factors of skin cell damage, down-regulate SIRT1 in a time and dose-dependent manner. Systematic studies revealed that ROS-mediated JNK activation is involved in UV-induced SIRT1 down-regulation (Fig. 7). SIRT activator, resveratrol which has also been considered an important antioxidant, protects against UV- and H$_2$O$_2$-induced apoptotic cell death, whereas SIRT1 inhibitors such as sirtinol and nicotinamide as well as SIRT1 RNAi enhance apoptosis. Our study provides evidence to support the notion that SIRT1 might be a novel target protecting UV/ROS-induced skin cell damage leading to skin photoaging and skin cancer.

**Acknowledgements**

This research was supported in part by a grant from NIH (P20 RR016457 from INBRE Program of the National Center for Research Resources), a grant for biomedical research from Rhode Island Foundation and a grant from Slater Center for Environmental Biotechnology.
References

1. Kaeberlein M, McVey M, Guarente L. The Sir2/3/4 complex and Sir2 alone promote longevity in Saccharomyces cerevisiae by two different mechanisms. Genes Dev. 1999; 13: 2570–80.

2. Vaziri H, Dessain SK, Ng Eaton E, et al. hiSir2(SIRT1) functions as an NAD-dependent p39 deacetylase. Cell. 2001; 107: 149–59.

3. Luo J, Nikolaev AY, Imai S, et al. Negative control of p53 by Sir2alpha promotes cell survival under stress. Cell. 2001; 107: 137–48.

4. Bouras T, Fu M, Sauve AA, et al. SIRT1 deacetylation and repression of p300 involves lysine residues 1020/1024 within the cell cycle regulatory domain 1. J Biol Chem. 2005; 280: 10264–76.

5. Brunet A, Bonni A, Zigmond MJ, et al. Akt promotes cell survival by phosphorylating and inhibiting a Forkhead transcription factor. Cell. 1999; 96: 857–68.

6. Motta MC, Divecha N, Lemieux M, et al. Mammalian SIRT1 represses forkhead transcription factors. Cell. 2004; 116: 551–63.

7. Picard F, Kurtev M, Chung N, et al. Sirt1 promotes fat mobilization in white adipocytes by repressing PPAR-gamma. Nature. 2004; 429: 113–8.

8. Rodgers JT, Lerin C, Haas W, et al. Nutrient control of glucose homeostasis through a complex of PGC-1alpha and SIRT1. Nature. 2005; 434: 113–8.

9. Jeong J, Juhn K, Lee H, et al. SIRT1 promotes DNA repair activity and deacetylation of Ku70. Exp Mol Med. 2007; 39: 8–13.

10. Cui R, Widlund HR, Feige E, et al. Central role of p53 in the sultan response and pathologic hyperpigmentation. Cell. 2007; 128: 853–64.

11. Bode AM, Dong Z. Mitogen-activated protein kinase activation in UV-induced signal transduction. Sci STKE. 2003; 2003: RE2.

12. Tournier C, Hess P, Yang DD, et al. Requirement of JNK for stress-induced activation of the cytochrome c-mediated death pathway. Science. 2000; 288: 870–4.

13. Wan YS, Wang ZQ, Shao Y, et al. Ultraviolet irradiation activates PI 3-kinase/AKT survival pathway via EGFR receptors in human skin in vivo. Int J Oncol. 2001; 18: 461–6.

14. Fisher GJ, Kang S, Varani J, et al. Mechanisms of photoaging and chronological skin aging. Arch Dermatol. 2002; 138: 1462–70.

15. Cao C, Healey S, Amaral A, et al. ATP-sensitive potassium channel: a novel target for protection against UV-induced human skin cell damage. J Cell Physiol. 2007; 212: 252–63.

16. Li Y, Bi Z, Yan B, Wan Y. UVB radiation induces expression of HIF-1alpha and VEGF through the EGFR/PI3K/DEC1 pathway. Int J Mol Med. 2006; 18: 713–9.

17. Wan Y, Wang Z, Shao Y, et al. UV-induced expression of GADD45 is mediated by an oxidant sensitive pathway in melanoma cells. Biochem J. 2006; 400: 225–34.

18. Brunn A, Sweeney LB, Sturgill JF, et al. Retinoic acid inhibits induction of c-Jun protein by ultraviolet radiation that occurs subsequent to activation of mitogen-activated protein kinase pathways in human skin in vivo. J Cell Mol Med. 2000; 6: 683–8.

19. Fisher GJ, Taiwar HS, Lin J, et al. Retinoic acid inhibits induction of c-Jun protein by ultraviolet radiation that occurs subsequent to activation of mitogen-activated protein kinase pathways in human skin in vivo. J Clin Invest. 1998; 101: 1432–40.

20. Brunet A, Sweeney LB, Sturgill JF, et al. Stress-dependent regulation of FOXO transcription factors by the SIRT1 deacetylase. Science. 2004; 303: 2011–5.

21. Cohen HY, Miller C, Bitterman KJ, et al. Sirt1 promotes cell survival by inducing the SIRT1 dependent p53 deacetylase. Cell. 2004; 116: 551–63.

22. Li Y, Bi Z, Yan B, Wan Y. Ultraviolet irradiation activates PI 3-kinase/AKT survival pathway via EGFR receptors in human skin in vivo. Int J Oncol. 2001; 18: 461–6.

23. Wan YS, Wang ZQ, Shao Y, et al. Ultraviolet irradiation activates PI 3-kinase/AKT survival pathway via EGFR receptors in human skin in vivo. Int J Oncol. 2001; 18: 461–6.

24. el-Deiry WS, Tokino T, Velculescu VE, et al. WAF1, a potential mediator of p53 tumor suppression. Cell. 1993; 75: 817–25.

25. Lin Y, Ma W, Benchimol S, Pidd, a new death-domain-containing protein, is induced by p53 and promotes apoptosis. Nat Genet. 2000; 26: 122–7.

26. Siliciano JD, Canman CE, Taya Y, et al. DNA damage induces phosphorylation of the amino terminus of p53. Genes Dev. 1997; 11: 3471–81.

27. Lambert PF, Kashchanchi F, Radonovich MF, et al. Phosphorylation of p53 serine 15 increases interaction with CBP. J Biol Chem. 1998; 273: 33048–53.

28. Gu W, Roeder RG. Activation of p53 sequence-specific DNA binding by acetylation of the p53 C-terminal domain. Cell. 1997; 90: 595–606.

29. Abraham J, Kelly J, Thibault P, et al. Post-translational modification of p53 protein in response to ionizing radiation analyzed by mass spectrometry. J Mol Biol. 2000; 295: 853–64.

30. Oren M. Regulation of the p53 tumor suppressor protein. J Biol Chem. 1999; 274: 3601–4.

31. Kobet E, Zeng X, Zhu Y, et al. MDM2 inhibits p300-mediated p53 acetylation and activation by forming a ternary complex with the two proteins. Proc Natl Acad Sci USA. 2000; 97: 12547–52.

32. Cao C, Wan S, Jiang Q, et al. All-trans retinoic acid attenuates ultraviolet radiation-induced down-regulation of aquaporin-3 and water permeability in human keratinocytes. J Cell Physiol. 2008; 215: 506–16.

33. Dasgupta B, Milbrandt J. Resveratrol stimulates AMP kinase activity in neurons. Proc Natl Acad Sci USA. 2007; 104: 7217–22.

34. Baur JA, Pearson KJ, Price NL, et al. Resveratrol improves health and survival of mice on a high-calorie diet. Nature. 2006; 444: 337–42.