MIF Antagonist (CPS1-1306) Protects Against UVB-induced Squamous Cell Carcinoma

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Running Title: MIF inhibitor reduces UVB skin carcinogenesis

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Abstract

Macrophage Migration Inhibitory Factor (MIF) is a homotrimeric pro-inflammatory cytokine implicated in chronic inflammatory diseases and malignancies including cutaneous squamous cell carcinomas (SCC). To determine if MIF inhibition could reduce ultraviolet-B light (UVB)-induced inflammation and squamous carcinogenesis, a small-molecule MIF inhibitor (CPSI-1306) was utilized that disrupts homotrimerization. To examine the effect of CPSI-1306 on acute UVB-induced skin changes, Skh-1 hairless mice were systemically treated with CPSI-1306 for 5 days prior to UVB exposure. In addition to decreasing skin thickness and myeloperoxidase (MPO) activity, CPSI-1306 pre-treatment increased keratinocyte apoptosis and p53 expression, decreased proliferation and phospho-histone variant H2AX (γ-H2AX), and enhanced repair of cyclobutane pyrimidine dimers (CPD). To examine the effect of CPSI-1306 on squamous carcinogenesis, mice were exposed to UVB for 10 weeks, followed by CPSI-1306 treatment for 8 weeks. CPSI-1306 dramatically decreased the density of UVB-associated p53 foci in non-tumor bearing skin while simultaneously decreasing the epidermal Ki-67 proliferation index. In addition to slowing the rate of tumor development, CPSI-1306 decreased the average tumor burden per mouse. While CPSI-1306-treated mice developed only papillomas, nearly a third of papillomas in vehicle-treated mice progressed to micro-invasive SCC. Thus, MIF inhibition is a promising strategy for prevention of the deleterious cutaneous effects of acute and chronic UVB exposure.

Implications: Macrophage Migration Inhibitory Factor (MIF) is a viable target for the prevention of UVB-induced cutaneous squamous cell carcinomas.
Introduction

Chronic inflammation and carcinogenesis are two longstanding processes that are intertwined with and feed into each other (1). While it has been widely recognized that long-standing inflammation (extrinsic to the tumor) can provide a favorable milieu for progression of transformed cells, recent studies have indicated that the tumor cells themselves can induce an inflammatory response (intrinsic) (2). Various components of chronic inflammation foster the growth and progression of tumors by contributing to the generation of reactive oxygen species (ROS) and cytokines and ultimately promoting genomic instability, angiogenesis, cellular migration, invasion, metastasis, and evasion of tumor immunosurveillance.

Cytokines function as the principal mediators between tumor and the various inflammatory cells and may be produced by either or both. Macrophage Migration Inhibitory Factor (MIF) is a pleotropic cytokine with predominantly pro-inflammatory properties that has been proposed to function as the primary link between inflammation and cancer (3, 4). MIF is a homotrimeric molecule that is expressed by a number of epithelial, mesenchymal, and inflammatory cells. MIF has been shown to play seminal roles in several chronic inflammatory diseases including diabetes mellitus (5), multiple sclerosis (6), systemic lupus erythematosus, rheumatoid arthritis and atherosclerosis (7). MIF expression levels are elevated within the tumor tissue and/or in the serum in a number of malignancies not limited to those arising from the lung, stomach, colon, ovary and liver and SCC of esophageal and head and neck origin (8-13). This increased expression of MIF has been correlated with a negative prognosis in most of these malignancies.

MIF has recently been recognized to be an important player in a multitude of skin diseases of infectious, inflammatory and neoplastic etiologies including squamous cell carcinomas (SCC) (14, 15). Ultraviolet-B light (UVB) functions as a complete carcinogen in skin by initiating genetic
mutations through direct DNA damage and production of ROS as well as promoting the survival
and growth of the transformed cells to produce tumors (16). Prolonged exposure to UVB light
also causes chronic inflammation and immunosuppression, contributing to the persistence and
proliferation of transformed cells. In skin, MIF is expressed in and secreted by epidermal
keratinocytes, dermal fibroblasts and infiltrating inflammatory cells (17). MIF is upregulated after
UVB exposure and through its pro-inflammatory and pro-angiogenic properties, MIF can abet
the tumor promoting effects of UVB (17-19). The other mechanisms by which MIF promotes
tumorigenesis have not yet been elucidated.

In a BALB/c mouse model of UVB induced SCC, lack of MIF decreased the acute inflammatory
response and dermal edema. Compared to their wild-type (WT) counterparts, these mice also
had diminished tumor number and burden, slower tumor progression and lower VEGF levels
(20). On the other hand, transgenic mice that overexpressed MIF were highly susceptible to
UVB induced carcinogenesis. Not only did they develop tumors earlier than their WT littermates,
but they also developed more and larger tumors (21). In both of these mouse models, there was
an inverse correlation between MIF expression and p53 levels. While MIF is expressed at
moderate levels in human skin (within keratinocytes and fibroblasts), actinic keratoses and
SCCs express high levels of MIF. In addition, exposure to UVB radiation increased MIF
expression in keratinocyte and SCC cell lines (19). These studies highlight the significance of
MIF in the development and progression of cutaneous SCC. The following experiments were
designed to examine whether inhibition of MIF via the administration of a CPSI-1306 can
alleviate the inflammatory and carcinogenic effects of acute and chronic UVB exposure.

**Materials & Methods**

Mice
The experiments were performed using 6–8 week old female Skh-1 hairless mice obtained from Charles River Laboratories (Wilmington, MA). The mice were housed in cages in groups of 5 in the vivarium at The Ohio State University according to the requirements established by the American Association for Accreditation of Laboratory Animal Care. All procedures were approved by the Ohio State University Institutional Animal Care and Use Committee before the initiation of the studies.

Drug administration

An orally bioavailable small molecule inhibitor of MIF, CPSI-1306 (Cytokine PharmaSciences, King of Prussia, PA), was used at a dose of 20mg/kg/day per mouse. The mice were orally fed 100µL per day of vehicle (15% DMSO, 0.5% methyl cellulose in water) or CPSI-1306 (in vehicle). The CPSI-1306 solution was prepared weekly based on the average mouse weight per cage.

Acute UVB exposure study

Mice (n=5 treated with vehicle or drug were used for each time point, 50 mice total) were treated with vehicle or CPSI-1306 for five consecutive days. 24 hours after the last gavage, they were either euthanized (no UVB group) or dorsally exposed to 2240J/m² (1 minimal erythemic dose, MED) of UVB light after which they were euthanized at 30 minutes, 6, 24 or 48 hours. UVB light was emitted by Philips FS40 UV lamps (American Ultraviolet Company, Lebanon, IN) and a UVX meter (UVP Inc., Upland, CA) was used to determine the UVB dose. The entire dorsal skin of the mouse was processed as follows: approximately 0.5cm² portion from the upper back was fixed in 10% neutral buffered formalin for histologic examination and immuno-stains (2 hours at room temperature (RT)). The fixed tissues were processed and embedded in paraffin blocks. A 10 mm diameter punch was flash frozen for the myeloperoxidase assay and all remaining skin was snap frozen.
Chronic UVB exposure study

A total of 36 mice (n=5 non-UVB exposed vehicle-treated, n=3 non-UVB exposed CPSI-1306-treated, n=13 UVB exposed vehicle-treated and n=15 UVB exposed CPSI-1306-treated) were used in this study. Mice belonging to the UVB groups were exposed to 1MED of UVB three times weekly on non-consecutive days for 10 weeks. At the end of UVB exposure, vehicle or CPSI-1306 solution was fed to the respective groups for 5 consecutive days per week for 4 weeks. During the next 4 weeks, the frequency of feeding was reduced to three times weekly on non-consecutive days. The mice were observed for the next 7 weeks and euthanized at the end of the 25-week study (Figure 1). The individual tumors measuring at least 1mm on each mouse was measured weekly to quantify tumor number, size, and total tumor area. The tumors were obtained and fixed separately in 10% neutral buffered formalin (4 hours at RT) and the rest of the tissues were processed as above.

Tumor grading

Hematoxylin and Eosin (H&E) stained sections of the tumors were examined in a blinded manner by a board certified veterinary pathologist. As previously described, tumor grades were assigned as follows: benign papilloma (grades P1-3), malignant micro-invasive squamous cell carcinoma (grades MI1-3), or fully invasive squamous cell carcinoma (SCC). Papillomas were exophytic tumors that showed no evidence of stromal invasion; while squamous cell carcinomas had a more endophytic appearance, with stromal invasion evidenced by loss of basement membrane continuity and development of an inflammatory stromal response. A grade 1 papilloma (P1) was composed primarily of epithelium without a pronounced papillary pattern; a grade 2 papilloma (P2) was a well-differentiated papillary mass; a grade 3 papilloma (P3) was similar to a grade 2 papilloma, except that a few finger-like projections of atypical cells at the
base of the mass were present. Micro-invasive squamous cell carcinomas (MI1-MI3) were subcategorized by depth of penetration into the dermis. Only tumors that invaded the panniculus carnosus were classified as fully invasive SCC. All grades of papilloma were considered benign (premalignant lesions), while all grades of micro-invasive squamous cell carcinomas and squamous cell carcinomas were considered malignant (malignant tumors) (22).

Myeloperoxidase activity assay

Myeloperoxidase (MPO) is an enzyme produced by neutrophils and the levels reflect the degree of acute inflammation in a tissue. The activity was determined using a chemical assay as described previously (23).

Immunohistochemistry

The following primary antibodies and dilutions were used for immunohistochemical staining of the skin: p53- 1:500 dilution, 60 minutes at RT [clone CM5p; Novocastra (Leica Microsystems), Buffalo Grove, IL]; Ki67- 1:200 dilution, 4°C overnight [clone TEC-3; Dako, Carpinteria, CA]; cleaved caspase-3- 1:750 dilution, 4°C overnight [clone Asp175; Cell Signaling Technology, Danvers, MA] and phospho-histone H2A.X- 1:250 dilution, 4°C overnight [clone Ser139; Cell Signaling Technology]. Five µm thick skin sections were deparaffinized and subjected sequentially to endogenous peroxidase blocking with 3% hydrogen peroxide, and antigen retrieval in Antigen Unmasking Solution (Vector Laboratories, Burlingame, CA) for 15 minutes in a microwave. The slides were cooled for 20 minutes and blocked using the Avidin- Biotin blocking kit (Vector Laboratories – for cleaved caspase-3 and phospho-histone H2A.X only). The slides were then blocked with 1x casein (Vector Laboratories) for 30 minutes at RT, followed by incubation with the specific primary antibodies as mentioned above. After incubation with their respective biotinylated secondary antibodies, the stains were developed using the Vectastain ABC (Vector Laboratories) and Diaminobenzidine (Vector Laboratories) kits, followed
by Hematoxylin counterstain. The slides were rinsed in 1x phosphate buffered saline containing 0.05% Tween-20 between the incubations.

**Immunohistochemistry Image Analysis**

The percentage of positive staining cells was calculated by manually counting positive and negative cells (ten 60x magnification images for the acute study- p53, Ki67, cleaved caspase-3; five 60x magnification images for chronic study- Ki67). For phospho-histone H2A.X, the percentage of positive epidermal area was determined using ImageJ from ten 60x magnification images. The results are expressed as the average per group for the above. The number of p53 foci in the chronic study was determined by manually counting through the entire length of the dorsal skin section and expressed as average number per 20x field.

**CPD Dot Blot**

Epidermal DNA was extracted using the PureLink Genomic Extraction Kit (Invitrogen) according to manufacturer directions. DNA (100ng) was chemically denatured with NaOH, heat denatured at 95C for 10min, then neutralized with ammonium acetate. DNA was mixed with SSC and blotted onto nitrocellulose membranes via gentle vacuum pressure in triplicate. Membranes were dried for 1hr at 80C then washed 3 times in TBST at room temperature for 5 minutes each. Membranes were blocked in TBST containing 5% BSA for 1 hr at room temperature and incubated overnight at 4C with a 1:500 dilution of anti-thymine dimer antibody (Kamiya Biomedical). Membranes were washed as above then incubated with secondary antibody at a 1:10,000 dilution for 45min at room temperature. Finally, membranes were washed, incubated with Lumiglo (Invitrogen), photographed, washed, incubated with SYBR Gold (Invitrogen) and photographed. Image J analysis was used to generate pixel intensities for CPD and SYBR gold and these values were expressed in a ratio to determine relative intensities.
Statistical analysis

For the acute exposure experiments, linear mixed effects models were applied to the data to allow for correlations among observations from the same mouse. For all models, the interaction between UV exposure time and outcome (cleaved caspase-3, Ki67, phospho-histone H2A.X, p53, CPD formation, skin thickness, and MPO) was assessed. In the presence of significant interaction, differences in outcome between CPSI-1306–treated mice and vehicle mice were estimated at each UV exposure time, with 95% confidence intervals (CI); otherwise, an overall group difference with 95% CI was estimated. For outcomes with non-zero values at baseline, interaction contrasts were also used to assess the relative effect of CPSI-1306 in the presence of UVB exposure vs. no UVB exposure. Note that cleaved caspase-3, CPD formation, skin thickness, and MPO were first natural-log transformed to stabilize variances; for cleaved caspase-3 a small number (0.01) was added before log transformation due to zero values.

Categorized tumor burden and the proportion of mice with at least one tumor at week 18 were compared between CPSI-1306 and vehicle groups using Fisher’s exact test. Similar to the acute experiments, linear mixed effects models were used to assess differences in Ki67 and p53 between the chronic exposure CPSI-1306 and vehicle groups. For Ki67, an interaction contrast was used to assess the relative effect of CPSI-1306 in the presence of chronic exposure vs. no UV exposure. For p53, group comparisons were made at various cells per field (1-2 cells, 3-5 cells, 6-10 cells, 11-20 cells, >20 cells).

All analyses were performed using SAS/STAT software, version 9.2 (SAS Institute Inc., Cary, NC).
Results and Discussion

Acute UVB exposure study

To determine if inhibition of MIF could protect against the deleterious epidermal effects of UVB exposure, we utilized a small molecule MIF inhibitor, CPSI-1306 (6). This molecule prevents the association of MIF monomers and prevents the generation of the biologically functional MIF homotrimer. We gavaged Skh-1 hairless mice with CPSI-1306 for 5 consecutive days and then exposed them to 1 MED (minimal erythemic dose) of UVB. Exposure to UVB causes significant epidermal keratinocyte apoptosis in the form of sunburn cells, which is accompanied by an increase in p53 expression (24). In severe cases, the keratinocyte apoptosis may be extensive, resulting in full thickness epidermal necrosis, such as in blistering sunburns. Towards the end of apoptosis or soon after, the lost keratinocytes are replaced by enhanced epidermal proliferation. Disruption of these processes such as by decreased apoptotic clearance and/or increased epidermal proliferation can lead to accumulation of DNA damage. UVB exposure also leads to the release of pro-inflammatory cytokines, leading to a late inflammatory response, which may in turn induce additional cell damage through ROS production. We euthanized the mice at specific time-points from 30 min to 48 hours post-UVB exposure and analyzed the levels of key epidermal markers of UVB-induced damage.

Inhibition of MIF decreased acute UVB-induced epidermal proliferation

To examine the net effect of acute UVB exposure on epidermal proliferation, we compared the levels of cleaved caspase-3, an apoptotic protein and Ki67, a marker of proliferation between vehicle-treated and CPSI-1306-treated mice. No significant differences were observed in the levels of either marker between the non-UVB exposed vehicle- and non-UVB exposed CPSI-1306-treated mice (Figure 2A and 2B). UVB-associated keratinocyte
apoptosis peaked at 24 hours post UVB exposure, as shown by the increased level of cleaved caspase-3. CPSI-1306-induced keratinocyte apoptosis could be appreciated as early as 30 minutes after a single UVB exposure (Figure 2A). At 6, 24 and 48 hours following UVB exposure, the CPSI-1306-treated mice showed significantly increased expression of cleaved caspase-3 compared to the vehicle-treated mice (p<0.0001). Epidermal proliferation as determined by Ki67 staining was significantly decreased by CPSI-1306 at 24 hours (p=0.0206) and 48 hours (p=0.0010) compared to vehicle treated mice at each time point (Figure 2B). Though not statistically significant, suppression of Ki67 by CPSI-1306 compared to vehicle-treated mice within each time point could also be observed at 30 minutes and 6 hours after UVB exposure (Figure 2B). Thus, inhibition of MIF led to enhanced keratinocyte apoptosis and decreased epidermal proliferation following UVB exposure.

**CPSI-1306 reduced acute UVB-induced keratinocyte DNA damage**

UVB exposure induces DNA damage directly through generation of cyclobutane pyrimidine dimers (CPD) and pyrimidine (6-4) pyrimidone photoadducts (25) as well as indirectly via production of ROS resulting in 8-oxo-2'-deoxyguanosine adduct formation (16). To determine the effect of MIF inhibition on UVB-induced keratinocyte DNA damage, we examined the repair kinetics of CPD by dot blot analysis. UVB-induced CPD DNA adducts were more abundant at every time point examined in vehicle versus CPSI-1306-treated epidermis (Figure 3A, p=0.0001). Phospho-histone H2A.X has been shown to be a surrogate marker for double-stranded DNA breaks (DSB) induced by UVB (26). Following UVB exposure, the levels of phospho-histone H2A.X peaked at 6 hours in both vehicle- and CPSI-1306-treated epidermis, following which the levels returned to nearly baseline levels. At every time point, CPSI-1306 treatment reduced the expression of phospho-histone H2A.X in epidermal keratinocytes (Figure 3B, p<0.0001). Even the baseline phospho-histone H2A.X levels were decreased in the non-UVB exposed epidermis of CPSI-1306 compared to vehicle treated mice.
MIF has been shown to antagonize the expression as well as function of wild-type p53 (27, 28). An increased level of p53 after UVB exposure ensures appropriate repair of DNA damage and elimination of irreversibly damaged cells (29). This enhanced production of p53 occurs principally through stabilization of the p53 protein, which prevents its proteasome-mediated degradation. To examine the effect of CPSI-1306 on p53, we examined its expression by immunohistochemistry. As expected, induction of p53 is an early event following UVB exposure, detectable within 30 minutes (Figure 3C), and reaching a maximum between 6 to 24 hours. Inhibition of MIF activity enhanced p53 levels in epidermal keratinocytes at 30 minutes (p=0.0009) and 6 hours (p=0.0001) after UVB exposure (Figure 3C). After this, there was no significant difference in the expression of p53 between the vehicle- and CPSI-1306-treated mice (Figure 3C). Thus, our findings are consistent with the idea that inhibiting MIF resulted in a decrease in the accumulation of DNA damage via a p53 mechanism.

**CPSI-1306 reduced UVB-induced acute inflammation**

The acute inflammatory response induced by UVB exposure contributes to secondary epidermal changes and accumulation of DNA damage. This inflammation is characterized by production of pro-inflammatory and chemotactic factors including MIF from damaged keratinocytes and fibroblasts, leading to vasodilation resulting in dermal edema and influx of neutrophils, followed by other inflammatory cells such as macrophages. To determine whether inhibition of MIF could reduce this inflammation in murine skin, we compared the average myeloperoxidase (MPO) levels, a surrogate for the number of neutrophils that infiltrate into the skin following UVB exposure (Table 1). The MPO levels increased progressively up to 48 hours after UVB irradiation at which time point the CPSI-1306-treated mice showed significantly lower MPO levels (p=0.0293). We then compared the average bi-fold thickness of back skin, a physical manifestation of UVB-induced cutaneous dermal edema (Table 1). The skin thickness
increased progressively, reaching a maximum at 48 hours in both groups. Pretreatment with CPSI-1306 reduced the average skin thickness at 48 hours, at an almost statistically significant level (p=0.0656). Thus, there is some evidence that inhibition of MIF may reduce the UVB-associated acute inflammatory response. Interestingly, the skins of the vehicle-treated mice were significantly thinner than the CPSI-1306-treated mice at 30 minutes following UVB exposure (p=0.0003).

**Chronic UVB exposure study**

To determine whether inhibition of MIF can protect against the deleterious effects of long-term UVB exposure, we exposed Skh-1 mice to 1 MED of UVB 3 times a week for 10 weeks. The mice were then treated with CPSI-1306 for five consecutive days per week for 4 weeks. Due to a limited supply of CPSI-1306, the frequency of CPSI-1306 feeding was reduced to three non-consecutive days per week during the subsequent 4 weeks. After 7 additional weeks of observation without CPSI-1306 treatment, the mice were euthanized at the end of 25 weeks (Figure 1). Upon prolonged exposure to UVB, there was increased tumor burden in vehicle versus CPSI-1306 treated mice (Figure 4C) corresponding to increased accumulation of DNA mutations as indicated by mutant p53 foci (Figure 5B) and enhanced epidermal proliferation (Figure 5A).

**Inhibition of MIF after chronic UVB exposure protected against SCC development**

Following chronic UVB radiation, the mice were monitored continuously for skin tumor development. Though the difference in the number of tumors between the groups was less pronounced at the end of 25 weeks, at all time points examined, CPSI-1306 treated mice had fewer tumors compared to vehicle controls (Figure 4A). At the completion of CPSI-1306 treatment (18 weeks), 62% of vehicle-treated mice were tumor free compared to 93% of CPSI-1306-treated mice (Figure 4B, p=0.0691). At the end of 25 weeks, 62% of vehicle-treated mice
were tumor free compared to 60% of CPSI-1306-treated mice (Figure 4B, p=0.9337). While the average tumor burden in the CPSI-1306-treated mice was visibly lower throughout the entire study compared to vehicle treated mice (Figure 4C), due to the lack of tumor development in some mice and the inherent variability of this tumor model, this differential tumor burden response did not achieve statistical significance. However, when classified based on size (≤10 mm², 11-20 mm², ≥20 mm²) the categorized tumor burden in CPSI-1306-treated mice at the end of CPSI-1306 treatment (18 weeks) was significantly lower than that of the vehicle-treated mice (Supplementary table 2, p=0.0131). After 7 weeks of no CPSI-1306 (25 weeks), categorized tumor burden in CPSI-1306 compared to vehicle treated mice was no longer significantly different (Supplementary table 2, p=0.5573). These data suggest that the efficacy of CPSI-1306 may be dependent upon continuous treatment. Of the tumors that developed in the vehicle-treated mice, 28% were micro-invasive SCCs (14% MI1 and 14% MI2) while none of the tumors of the CPSI-1306-treated mice developed into micro-invasive (MI) or frank SCCs (Figure 4D). Thus, inhibition of MIF prevented UVB-induced cutaneous squamous carcinogenesis. In our model, the beneficial effects of CPSI-1306 were more apparent at 18 weeks than at 25 weeks. This data suggests that continued inhibition of MIF through administration of CPSI-1306 could have further prevented UVB-induced squamous carcinogenesis.

**CPSI-1306 treatment after chronic UVB exposure diminished epidermal proliferation**

To examine whether CPSI-1306 could affect chronic UVB-induced epidermal proliferation, the average Ki67 proliferation index of the non-tumor bearing skin was determined. CPSI-1306 decreased the percentage of Ki67 positive epidermal keratinocytes in both non-UVB exposed and UVB exposed skin. This decrease was statistically significant in the skin of UVB-exposed mice relative to no UV exposure (Figure 5A, p=0.0168). Thus, inhibition of MIF decreased epidermal proliferation associated with chronic UVB-exposure.
CPSI-1306 inhibition following chronic UVB exposure reduced the number of p53 foci

Chronic UVB exposure causes mutations of the p53 gene and the resultant abnormal p53 protein is resistant to proteasome-mediated degradation and can be easily detected (30, 31). Cells with the non-functional p53 protein are frequently impervious to normal DNA-damage checkpoints and can persist for long periods of time. They may expand clonally to form UVB induced ‘p53 foci’ that can be detected by immunohistochemistry. These clones may accumulate additional DNA mutations and progress to develop squamous papillomas and SCCs. In fact, the risk for SCC development correlates with the density of p53 foci (32, 33).

To examine if CPSI-1306 could affect the epidermal accumulation of mutant p53, we examined the expression of p53 protein in non-tumor bearing skin by immunohistochemistry. The average number of epidermal p53 foci was significantly reduced in the CPSI-1306-treated mice compared to the vehicle-treated mice (Figure 5B, p=0.0014). The p53 foci were further classified based on their size, i.e. number of cells per foci as follows: 3-5, 6-10, 11-20 and > 20 cells. The majority of the foci contained less than ten cells. While there was a statistically significant decrease only in the number of the smaller p53 foci, i.e. those with 3-5 and 6-10 cells (p<0.0001, p=0.0578, respectively, data not shown), there were fewer foci of all sizes in the skin of CPSI-1306 treated mice compared to vehicle treated mice (Figure 5B, p=00014). Thus, inhibition of MIF led to a lower density of p53 foci.

Our data indicate that inhibition of MIF prior to acute UVB exposure reduced epidermal proliferation while enhancing apoptotic removal of cells. This was accompanied by increased and early expression of p53 protein and an overall decrease in DNA damage and DNA double strand breaks. In addition to these beneficial effects on epidermal keratinocytes, CPSI-1306 also lessened the dermal edema and acute inflammation associated with UVB exposure. Thus,
MIF inhibition protected against both the direct and indirect effects of acute UVB exposure induced skin changes. Treatment of chronically UVB exposed skin with CPSI-1306 decreased the density of p53 foci and significantly reduced epidermal proliferation. Inhibition of MIF also reduced tumor burden and malignancy rate. These effects were present even after cessation of CPSI-1306. Taken together, our data indicate that inhibition of MIF was protective against both the acute and chronic effects of UVB exposure.

**Functional antagonism between MIF and p53**

MIF was identified as one of the negative regulators of p53 activity and its inhibitory effect on p53 is manifold; including suppression of p53 transcriptional activity, reduction in p53-dependent apoptosis and evasion of cell-cycle arrest (27). The exact mechanism(s) by which this repression is mediated is still being elucidated and could be context dependent. MIF contributes to the progression of endotoxic shock by repressing p53 function and up-regulating Cox-2 activity (34). On the other hand, NM23-H1, a suppressor of metastasis, physically interacts with MIF and relieves suppression of p53 (35). Jab1, a modulator of intracellular signaling, interacts with several cell cycle regulators including AP1, MIF, and p53 and as such may function as a mediator of functional interactions between MIF and p53 (36, 37). In addition to these indirect associations, MIF has been shown to interact directly with p53 through cysteine residues independent of Jab1 (38) and this interaction is determined by the redox state of the amino acid residues involved. The interaction of MIF and p53 promotes the association of p53 with Mdm2 and thus leading to the ubiquitination of p53 and its proteasome-mediated degradation. Thus, MIF and p53 play contradictory roles in cell cycle regulation, inflammation and tumorigenesis.

**Proposed mechanism of action of CPSI-1306 in our model**
After acute UVB exposure, there is an induction of MIF expression and trimerization in the skin (19) which promotes p53 degradation thus, antagonizing the function of p53 (39), resulting in increased inflammation and accumulation of DNA damage through increased epidermal proliferation and decreased keratinocyte apoptosis. CPSI-1306 prevents the formation of the MIF homotrimer and renders MIF biologically non-functional and incapable of blocking p53 protein expression and activity (Figure 6A). In chronically UVB-exposed skin, there is accumulation of DNA damage indicated by increased formation of p53 foci. Inhibition of MIF activity alleviates its inhibitory effect on p53. Increased p53 activity slows cell cycle progression, allowing time for the repair of DNA damage and thus, decreasing p53 foci density. These processes together diminish the carcinogenic transformation and progression of transformed cells (Figure 6B). Thus, our studies demonstrate that the inhibition of MIF protects against the deleterious effects of acute and chronic UVB exposure principally by relieving its inhibitor effects on p53.
Acknowledgements

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References


Table 1. CPSI-1306 decreases UVB induced acute inflammation.

<table>
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<tr>
<th>UV exposure</th>
<th>Group</th>
<th>Myeloperoxidase Activity: Mean (Standard Deviation)</th>
<th>Bi-fold skin thickness (mm): Mean (Standard Deviation)</th>
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<td>No UV</td>
<td>Vehicle</td>
<td>0.0041 (0.0015)</td>
<td>0.64 (0.05)</td>
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*p=0.0293, *p=0.0656, ^p=0.0003
Figure Legends

Figure 1. Schematic of Chronic UVB exposure and CPSI-1306 treatment schedule. Female Skh-1 mice were exposed to UVB 3 times per week on non-consecutive days for 10 weeks. UVB treatments were then halted and mice were gavaged with 20mg/kg/day of CPSI-1306 or vehicle 5 days per week for 4 weeks, then 3 days per week for an additional 4 weeks then left untreated for the final 7 weeks, after which they were sacrificed.

Figure 2. Inhibition of MIF increases keratinocyte apoptosis, while decreasing proliferation. Female Skh-1 mice were per-treated with vehicle or CPSI-1306 for 5 days, exposed to 1 MED of UVB and sacrificed at 30min or 6, 24, or 48h after UVB exposure. Dorsal skin was examined for expression of the pro-apoptotic protein cleaved caspase-3 or the proliferation marker Ki67 by immunohistochemistry. The percentage of positively staining epidermal cells was determined (CPSI-1306 vs. vehicle). A, Left: CPSI-1306 significantly enhanced epidermal cleaved caspase-3 mediated apoptosis at 6, 24, and 48 hours compared to vehicle (*p<0.0001). Right: representative cleaved caspase-3 immunohistochemistry images (60X) at 24 hours post UVB-exposure when the maximum difference was observed between CPSI-1306 and vehicle treated mice. Arrows indicate cleaved caspase-3 positive epidermal cells. B, Left: CPSI-1306 significantly decreased epidermal proliferation as examined by Ki67 immunohistochemistry at both 24h (*p=0.0206) and 48h (*p=0.0010), relative to vehicle. Right: representative Ki67 immunohistochemistry images (60X magnification) at 48 hours post UVB-exposure when the maximum difference was observed between CPSI-1306 and vehicle treated mice. Bars represent standard deviation, * p<0.05.

Figure 3. CPSI-1306 treatment decreases UVB-induced DNA damage. Epidermal DNA isolated from the dorsal skin of mice treated with vehicle or CPSI-1306 for 5 days prior to a single UVB exposure was examined by dot blot analysis to determine levels of cyclobutane
pyrimidine dimers (CPD).  A. Left: At all time points examined, CPSI-1306 significantly decreased UVB-induced CPD levels compared to vehicle treated skin (p=0.0001).  Right: dot blot for CPD; each dot represents DNA from one mouse and the various treatment groups are labeled and outlined for comparison between CPSI-1306 and vehicle treated mice.  Dorsal skin was also examined immunohistochemically to determine changes in levels of phospho-histone H2A.X and p53.  B.  Left: CPSI-1306 treatment decreased epidermal levels of phospho-histone H2A.X compared to vehicle treated mice at baseline and at all time points following UVB exposure (p<0.0001).  Right: representative phospho-histone H2A.X immunohistochemistry images (60X magnification) at 6 hours post UVB-exposure when the maximum difference was observed between CPSI-1306 and vehicle treated mice.  C, CPSI-1306 significantly increased epidermal p53 levels at both 30 min (p=0.0009) and 6hr (p=0.0001) following UVB exposure relative to vehicle.  Right: representative p53 immunohistochemistry images (60X magnification) at 6-hour post UVB-exposure when the maximum difference is observed between CPSI-1306 and vehicle treated mice.  Bars represent standard deviation, * p<0.05.

Figure 4. Inhibition of MIF decreases UVB-induced squamous carcinogenesis. Skh-1 mice were exposed to UVB 3 times per week on non-consecutive days for 10 weeks. UVB treatments were halted, then mice were gavaged 5 days per week with 20mg/kg/day CPSI-1306 for 4 weeks then 3 days per week for an additional 4 weeks, and finally monitored for an additional 7 weeks without further treatment.  All tumors were counted and measured weekly after cessation of UVB exposure, excised at the end of the experiment, and histologically graded by a board certified veterinary pathologist to determine the grade and extent of invasion.  A, Inhibition of MIF decreased tumor multiplicity compared to vehicle; B, CPSI-1306 treatment delayed tumor onset relative to vehicle; C, Treatment with CPSI-1306 decreased tumor burden compared to vehicle; D, CPSI-1306 treatment inhibited tumor progression compared to vehicle; P1-P3:

Figure 5. CPSI-1306 treatment reduces proliferation and p53 foci formation in non-tumor epidermis. Non-tumor bearing dorsal skin harvested from chronically UVB-irradiated mice treated with either vehicle or CPSI-1306 was examined immunohistochemically to determine the percentage of Ki67 positive epidermal keratinocytes as well as the number of p53 foci. A, Left: Treatment with CPSI-1306 significantly inhibited epidermal proliferation in non-tumor bearing skin compared to vehicle, relative to no UV exposure (p=0.0168); Right: representative Ki67 immunohistochemistry images (60X magnification) of chronic UVB-exposed CPSI-1306 and vehicle treated mice. B, Left: MIF inhibition after chronic UVB exposure significantly decreased the density of epidermal p53 foci (p=0.0014); Right: representative p53 immunohistochemistry images (20X magnification) of chronic UVB-exposed CPSI-1306 and vehicle treated mice. Bars represent standard deviation. * p<0.05.

Figure 6. A proposed model for the role of MIF in UVB-mediated inflammation and skin carcinogenesis. A, Acute UVB exposure induces MIF expression and trimerization in the skin which inhibits p53 function by enhanced degradation of p53, increased inflammation and epidermal proliferation, and decreased apoptosis. CPSI-1306 treatment reverses the above-mentioned effects of MIF, decreasing the accumulation of DNA damage. B, Chronic UVB exposure results in increased dermal MIF expression and trimerization leading to increased epidermal proliferation and decreased apoptosis. As a consequence of genomic DNA damage and loss of p53 function, there is increased tumor formation and progression in the vehicle-
treated mice. Treatment with CPSI-1306 prevents the trimerization of MIF, thus reducing the accumulation of DNA damage and eventually the squamous carcinogenesis.
UVB exposure                          Gavage                  Observation
   2240 J/m² 3x per week
CPSI-1306 or Vehicle
   5× per week 3× per week
0 weeks                              10              14             18                           25

Euthanasia
**Figure 2**
Figure 3
Figure 4
Figure 5
A. Effect of MIFi on acute UVB induced skin changes

![Diagram showing the effect of MIFi on acute UVB induced skin changes]

B. Effect of MIFi on chronic UVB induced skin changes and tumorigenesis

![Diagram showing the effect of MIFi on chronic UVB induced skin changes and tumorigenesis]

Figure 6
MIF Antagonist (CPSI-1306) Protects Against UVB-induced Squamous Cell Carcinoma

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