Simultaneous Inhibition of the HGF/MET and Erk1/2 Pathways Affect Uveal Melanoma Cell Growth and Migration

Chandrani Chattopadhyay¹, Elizabeth A. Grimm¹, Scott E. Woodman¹,²

¹ Melanoma Medical Oncology, University of Texas, MD Anderson Cancer Center, Houston, Texas, United States of America, ² Systems Biology, University of Texas, MD Anderson Cancer Center, Houston, Texas, United States of America

Abstract

Purpose: Nearly all primary uveal melanoma (UM) that metastasize involve the liver. Hepatocyte growth factor (HGF) is proposed to be an important microenvironmental element in attracting/supporting UM metastasis through activation of MET. The majority (>85%) of UM express mutations in the G-alpha proteins, that drive the MEK-ERK1/2 pathway. Thus, we proposed that the combination of MET and MEK inhibition would inhibit the growth and migration of G-alpha protein mutant versus non-mutant UM cells.

Methods: Western-blots demonstrated the relative protein levels of ERK1/2 and MET in UM cells. Cells were treated with the small molecule inhibitors AZD6244 (MEKi) and/or MK-8033 (METi) and downstream markers evaluated. Further studies determined the effect of combination MEKi and METi treatment on cell growth, apoptosis and migration.

Results: All G-alpha protein mutant UM cell lines express MET mRNA and protein. The level of mRNA expression correlates with protein expression. MEKi, but not METi treatment results in markedly reduced ERK1/2 phosphorylation. Either MEKi or METi treatment alone reduces cell proliferation, but only modest induction of apoptosis. The combination MEKi+ METi results in significant reduction of proliferation in G-alpha protein mutant cells. UM cell migration was blocked by METi, but not MEKi treatment.

Conclusions: MET protein expression showed no correlation with G-alpha protein mutation status. Combining MEKi with METi treatment has added benefit to either treatment alone in reducing G-alpha protein mutant UM cell growth. Combining MEKi with MEKi treatment adds the effect of limiting uveal melanoma cell migration.

Introduction

In Western countries, uveal melanoma is diagnosed in approximately 4–11 cases per million people per year [1,2]. Nearly 30% of patients with uveal melanoma develop metastases by 10 to 15 years after diagnosis, and the metastatic disease is universally fatal. The metastatic disease mortality rate remains unchanged despite advances in treating the primary eye tumor.

More research into the biology of uveal melanoma is needed urgently to understand the critical clinically targetable pathways that will lead to improved patient outcomes.

The recent identification of activating mutations in the G-alpha protein in uveal melanoma has provided a key insight into potential strategies in which to target uveal melanoma cell growth and survival [3,4]. The mutations are somatic and occur in either amino acid sites R183 or Q209 turning GNAQ into a dominant oncogene with constitutive RAS/MEK/ERK1/2 signaling activation [5,6]. However, we and others have shown that although small molecule MEK inhibitors can decrease cell growth in GNAQ mutant cells, MEK inhibition alone often fails to mediate significant apoptotic responses in these cells [4,7]. In addition, similar somatic mutations in R183 or Q209 in the GNA11 gene have been reported, and together with the GNAQ represent approximately 85% of all primary uveal tumors.

It is known that uveal melanoma metastasizes to the liver via the hematogenous route, however the mechanism responsible for liver-predominant metastasis is unknown. Currently, there is no available systemic therapy for preventing or treating uveal melanoma metastases; so our focus on liver biology, and specifically on secreted growth factors like hepatocyte growth factor (HGF) is a high priority. The receptor for HGF, named c-MET (MET), has been found in uveal melanoma tumors and cells previously [8] and our own work on MET in N-Ras mutated cutaneous melanoma has prompted us to explore this system in more detail [9].
Prior studies have indicated that HGF may play an important role in mediating uveal melanoma growth and metastasis. Mallekarjuna et al., showed that primary uveal melanoma tumors that metastasize had higher levels of MET expression than tumors that did not metastasize [10]. In addition, a significant association between MET expression and uveal melanoma specific-mortality was noted. These investigators and others have also shown elevated MET levels in uveal melanoma cells within uveal melanoma tumors [8,10–12].

In this study we investigate the effect of MEKi and/or METi treatment on \(\text{GNAQ}\) mutant and wild-type uveal melanoma cell lines. Our goal is to determine whether any biological basis exists for combined METi and MEKi treatments as a potential targeted therapy option, and whether the effects are more pronounced in mutant G-alpha protein cells or not.

### Materials and Methods

#### Reagents

The small molecule MET inhibitor, MK-8033, was obtained under a material transfer agreement with Merck and Co (SRA # LS2009-00026397JW). AZD6244 was obtained from Selleckchem (Houston, TX, USA). Human HGF was purchased from R&D Systems (Minneapolis, MN). MK-8033 or AZD6244 were dissolved in dimethyl sulfoxide (DMSO) to prepare a stock solution of 10 mM, and then diluted as indicated in fresh medium. In all experiments, the final concentration of DMSO was <0.1%.

#### Cells and Cell Culture

Melanoma cells were grown in RPMI 1640 medium supplemented with 10% fetal bovine serum.

All uveal melanoma cell lines used in this study were a generous gift of Martine Jager (Leiden University, Netherlands), whose laboratory generated the 92.1 cell line [13]. Mel202, Mel270, Mel285, Mel290, OMM2.3, and OMM2.5 originated from Bruce Ksander [14]. OCM1 and OCM3 originated from the laboratory.

![Image](image_url)

**Figure 1.** MET mRNA expression levels in GNAQ mutant and wild-type uveal melanoma cells. (A) MET mRNA expression (top panel) and actin mRNA expression (lower panel) as determined by RT-PCR. Images are from the same gel at the same exposure. (B) MET protein expression levels in GNAQ mutant and wild-type uveal melanoma cells. MET protein expression (top panel) and actin protein expression (lower panel) as determined by western-blot.

doi:10.1371/journal.pone.0083957.g001
of June Kan-Mitchell [15]. Cell line validation was accomplished by short random repeat (STR) DNA fingerprinting techniques and mutational analysis by MDACC Cancer Center Support Grant (CCSG)–supported Characterized Cell Line Core. Cell lines were validated by STR DNA fingerprinting using the AmpFSTR Identifier Kit (Applied Biosystems, Foster City, CA) according to manufacturer’s instructions. The STR profiles were compared to known ATCC fingerprints (ATCC.org), and to the Cell Line Integrated Molecular Authentication database (CLIMA) version 0.1.200808 (http://bioinformatics.istge.it/clima/). The STR profiles matched known DNA fingerprints or were unique [16].

Western Blotting and Antibodies
Cells were lysed in a buffer containing 50 mM Tris (pH 7.9), 150 mM NaCl, 1% NP40, 1 mM EDTA, 10% glycerol, 1 mM sodium vanadate, and protease inhibitor cocktail (Roche Pharmaceuticals, Nutley, NJ). Proteins were separated by SDS-PAGE with 4–20% gradient gels (Bio-Rad Laboratories, Hercules, CA), transferred to a Hybond-ECL nitrocellulose membrane (GE Healthcare Biosciences, Piscataway, NJ), and blocked in 5% dry milk in PBS. The membrane was then incubated with primary and secondary antibodies, and target proteins were detected with ECL detection reagent (GE Healthcare Biosciences).

MET(C-12), FARP and beta actin antibodies were obtained from Santa Cruz Biotechnology, Phospho-Met antibody was purchased from Invitrogen, Phospho Erk1/2 and Erk1/2 antibodies were obtained from Cell Signaling Technology.

Cell Proliferation Assays
Melanoma cells were plated at a density of 1×10⁶ cells/well in triplicate wells in a 24-well plate in RPMI 1640 growth medium, DNA content and cell cycle phase were analyzed using a FACScan at a final concentration of 5 mg/mL PI and 10 mg/ml RNAse.

Cell migration assays were performed in Boyden chambers using uncoated filters (BD Biocoat control inserts, BD Biocoat, San Jose, CA). 2.5×10⁵ cells/well were plated in serum-free medium, with or without a four hour treatment of MK-8033, and the migration assay performed. Stained cells were photographed with a Nikon Eclipse TE2000-U microscope at 20X magnification using NIS Elements advanced research software. To quantify migration, the cells in each filter were counted from five independent fields under the microscope at 40X magnification and the mean cell number/field was calculated. Each assay condition was tested in 2 replicates.

RNA Isolation
Total cellular RNA was extracted according to manufacturer’s instructions using a NucleoSpin RNA II kit (Macherey-Nagel, Bethlehem, PA).

Reverse Transcription PCR (RT-PCR)
To determine the mRNA levels of MET in uveal melanoma cells, we performed first-strand cDNA synthesis with 400 ng of total RNA using a GeneAmp RNA PCR kit (Applied Biosystems, Carlsbad, CA) according to the manufacturer’s protocol. A 2-µL cDNA product was used for each 25-µL PCR reaction. The RT-PCR primers used to detect human MET and beta-actin were adapted from the publication by Fujita and Sugano [17]. The PCR protocol consisted of initial denaturation at 95°C for 5 min; 30 cycles of 95°C for 40 s, 55.5°C for 30 s, and 72°C for 60 s; primer extension at 72°C for 1 min; and a final extension at 72°C for 10 min. We analyzed 20 µL of PCR product on a 1% agarose gel.

Results
MET mRNA and Protein Expression in Uveal Melanoma Cell Lines
Total RNA was extracted from uveal melanoma cell lines and levels of MET mRNA transcript determined by RT-PCR (Figure 1A). All uveal melanoma cell lines, regardless of GNAQ mutation status, exhibited MET mRNA expression, although the 92.1 (GNAQ_Q209L) cell line displayed far less MET mRNA transcript compared to the other cell lines. Each of these cell lines also demonstrated MET protein expression, irrespective of GNAQ mutation background (Figure 1B). Consistent with the lower MET transcript levels in the 92.1 cell line, the MET protein level was also lower in these cells. Of note, the MEL270, OMM2.3 and OMM2.5 cell lines are derived from the same individual [18,19], and show similar MET mRNA and protein expression. Cell lines that have been stated to be of uveal melanoma origin, but for which we and others have documented the presence of the BRAF V600E mutation, [16] do not show significant MET protein expression despite displaying detectable MET transcript levels (Figure S1A&B).

Effect of MEKi and/or METi Treatment on MET and Erk1/2 Phosphorylation in Uveal Melanoma Cells
We have previously demonstrated the capacity of the small molecule MEK inhibitor (AZD6244) to decrease ERK1/2 expression despite displaying detectable MET transcript levels (Figure S1A&B).
phosphorylation in uveal melanoma cell lines at low nanomolar concentrations [20]. To determine the capacity of the small molecule MET inhibitor (MK-8033) to inhibit MET phosphorylation in uveal melanoma cell lines (MEL290), and the lower panel demonstrates the effect of MK-8033 on MET phosphorylation in GNAQ mutant cells (MEL202). Total MET protein expression is shown for each cell line under each condition. (B) Effect of METi (MK-8033) and/or MEKi (AZD6244) treatment on Erk1/2 phosphorylation in GNAQ mutant and wild-type uveal melanoma cells. The upper panel demonstrates the effect of MK-8033 and/or AZD6244 on Erk1/2 phosphorylation in WT uveal melanoma cells (MEL285), and the lower panel demonstrates the effect of MK-8033 and/or AZD6244 on Erk1/2 phosphorylation in GNAQ mutant uveal melanoma cells (MEL202).

doi:10.1371/journal.pone.0083957.g002

Figure 2. Effect of METi (MK-8033) treatment on MET phosphorylation in GNAQ mutant and wild-type uveal melanoma cells. (A) The upper panel demonstrates the effect of MK-8033 at 0, 1 and 2.5 μM on MET phosphorylation in WT uveal melanoma cells (MEL290), and the lower panel demonstrates the effect of MK-8033 on MET phosphorylation in GNAQ mutant cells (MEL202). Total MET protein expression is shown for each cell line under each condition. (B) Effect of METi (MK-8033) and/or MEKi (AZD6244) treatment on Erk1/2 phosphorylation in GNAQ mutant and wild-type uveal melanoma cells. The upper panel demonstrates the effect of MK-8033 and/or AZD6244 on Erk1/2 phosphorylation in WT uveal melanoma cells (MEL285), and the lower panel demonstrates the effect of MK-8033 and/or AZD6244 on Erk1/2 phosphorylation in GNAQ mutant uveal melanoma cells (MEL202).

doi:10.1371/journal.pone.0083957.g002

phosphorylation in uveal melanoma cell lines at low nanomolar concentrations [20]. To determine the capacity of the small molecule MET inhibitor (MK-8033) to inhibit MET phosphorylation in uveal melanoma cell lines with a GNAQ mutation versus no GNAQ mutation (hereafter “wild-type”), cells were treated with MK-8033 at either 0, 1 or 2.5 μM concentrations (Figure 2A). METi treatment reduced MET phosphorylation in both GNAQ mutant or wild-type cells by a relatively similar proportion, relative to baseline MET phosphorylation levels. To test the combined effect of MEKi and/or METi treatment, GNAQ mutant and wild-type cells were treated with AZD6244 (MEKi) at 25 nM and/or MK-8033 (METi) at 2.5 μM. MEKi treatment resulted in a marked decrease in Erk1/2 phosphorylation in both GNAQ mutant and wild-type cells, although complete inhibition was not observed in wild-type cells. METi treatment up to 2.5 μM failed to reduce Erk1/2 phosphorylation in either GNAQ mutant or wild-type cells. The combination MEKi+METi treatment recapitulated the effect of MEKi treatment alone in GNAQ mutant cells, but served to enhance the reduction of MEKi alone treatment in wild-type cells (Figure 2B). To further analyze the functional significance of MET expression in uveal melanoma, cells were treated with HGF (100 ng/ml), the ligand for MET. HGF treatment caused marked elevation of ERK1/2 phosphorylation which was decreased with co-treatment of low micro-molar concentrations of the METi, showing the efficacy of the METi at a HGF concentration higher than present in serum (Figure 3). Of note, METi treatment also extinguished the HGF induced phosphorylation of AKT.
Effect of MEKi and/or METi Treatment on Uveal Melanoma Cell Proliferation

To determine the effect of MEK and/or MET inhibition on the growth of uveal melanoma, cells were treated with MEKi and/or METi (Figure 4). Relatively lower concentrations of each drug were used in order to optimize the potential of enhanced effects being observed with combination therapy. Treatment with either the METi or MEKi alone resulted in a noted decrease in cell proliferation in GNAQ mutant uveal melanoma cells. MEKi treatment resulted in a modest decrease in proliferation of wild-type cells, but no difference was observed after METi treatment in wild-type cells. The combination MEKi+METi treatment resulted in a further decrease in proliferation of both GNAQ mutant and wild-type cells.

Figure 3. Effect of METi treatment on Erk1/2 and AKT phosphorylation in GNAQ mutant and wild-type uveal melanoma cells following stimulation with HGF (100/1ng/ml). The effect of HGF on uveal melanoma cell Erk1/2 phosphorylation (upper panel) or AKT phosphorylation (lower panel) with or without METi treatment at 0.25 or 2.5 μM. Images are from the same blot at the same exposure. doi:10.1371/journal.pone.0083957.g003

Figure 4. Effect of MEK and/or MET inhibition on the growth of GNAQ mutant versus wild-type uveal melanoma cells. The upper panel shows the effect of 25 nM MEKi and/or 2.5 μM METi treatment on two distinct GNAQ mutant cells, OMM2.3 (left) and MEL202 (right). The lower panel shows the results of the same treatment conditions in GNAQ wild-type cells, MEL285 (left) and MEL290 (right). doi:10.1371/journal.pone.0083957.g004
in an enhanced inhibition of cell growth in GNAQ uveal melanoma cells, but not in wild-type cells.

Targeted siRNA knockdown of MET was performed to better determine the specific effect of MET reduction has on uveal melanoma cellular proliferation. The relative degree of MET knockdown, as determined by MET protein expression, was similar in mutant and wild-type cells (Figure S2A). The siRNA knockdown of MET resulted in a decrease in proliferation in all uveal melanoma cell lines, with a more pronounced effect being observed in mutant cells (Figure S2B).

Effect of MEKi and/or METi Treatment on PARP Cleavage in Uveal Melanoma Cells

To determine the effect of MEK and/or MET inhibition on apoptosis in uveal melanoma, cells were treated with the MEKi and/or METi and PARP cleavage evaluated (Figure 5). Each cell line had little to no detectable cleaved PARP at baseline. Similarly increased levels of cleaved PARP were observed in GNAQ mutant uveal melanoma cells following either MEKi or METi treatment alone. The level of cleaved PARP was more enhanced by MEKi+METi combination treatment in GNAQ mutant cells. The MEL285 wild-type cell line (WT1) showed a modest increase in cleaved PARP to each agent alone, without a clear enhancement to combination treatment. The MEL290 wild-type cells (WT2) failed to show significantly increased levels of cleaved PARP with MEKi and/or METi treatment. Although low concentrations of each drug were employed in order to identify additive effects, the sub-G0 cell fraction in flow cytometry-based cell cycle analysis was observed to be slightly increased following combination AZD6244+MK-8033 treatment in the GNAQ mutant cells, consistent with the cleaved PARP results (data not shown).

Effect of MEKi and/or METi Treatment on Uveal Melanoma Cell Migration

To determine the effect of MEK and/or MET inhibition on cell migration, uveal melanoma cells with a GNAQ mutation were treated with MEKi and/or METi. Figure 6A shows images of stained migrated GNAQ mutant or wild-type cells following 0, 1 and 2.5 μM METi treatment. A marked reduction in cell migration was observed following METi treatment at micromolar concentration levels in all uveal melanoma cells, regardless of mutation or wild-type status (Figure 6B). Of note, METi treatment had no significant effect on cell migration in the BRAF mutant cell lines tested (Figure S3A), consistent with the lack of MET protein expression in these cells as shown in Figure S1B. MEKi treatment had no significant effect on cell migration in any of the cell lines tested (Figure S3B).

Discussion

Approximately 85% of uveal melanoma cells harbor activating mutations in G-alpha proteins. When activated, these G-alpha proteins clearly signal through the MEK-ERK1/2 pathway. We have shown that small molecule inhibition of MEK is insufficient to drive cellular death in the majority of cells with GNAQ/11 mutations, suggesting that other important pathways contributing to cell survival and are active in cells harboring these mutations [20].

In this study we demonstrate that uveal melanoma cells with a GNAQ mutation express the MET receptor at high levels. Employing small molecule inhibitors to both the MEK and the MET receptor, we show that MET inhibition enhanced the growth inhibitory effect of the MEKi. The METi had the added effect of inhibiting the migration of uveal melanoma cells regardless of GNAQ mutation or wild-type status in all cell lines expressing the MET receptor, suggesting that the migration inhibitory effects of METi are present across all uveal melanoma...

Figure 5. Effect of MEKi and/or METi treatment on PARP cleavage in uveal melanoma cells. The upper panel shows the effect of MEKi at 25 nM and/or METi at 2.5 μM on two distinct GNAQ mutant cells, OMM2.5 (left) and MEL202 (right). The lower panel shows the results in GNAQ wild-type cells, MEL285 (left) and MEL290 (right).
doi:10.1371/journal.pone.0083957.g005
cell lines with MET expression. An interesting observation in the current study was that the METi alone treatment had growth inhibitory effects on \(GNAQ\) mutant cells, but not wild-type cells, at the concentration of METi used in this study. This observation is consistent with the findings of Abdel Rahman et al. who showed the MET inhibitor SU11274 to more markedly inhibit the proliferation of the uveal melanoma cell lines 92.1 and MEL202, compared to other cell lines tested [8]. It has since been determined that the 92.1 and MEL202 cell lines harbor \(GNAQ\) mutations, and the other cell lines treated in that study have neither \(GNAQ\) nor \(GNA11\) mutations [16].

At the doses used in the current study, METi treatment resulted in marked loss of MET phosphorylation in both \(GNAQ\) mutant and wild-type cells, although complete extinguishment of MET phosphorylation was not observed in wild-type cells. MET inhibition did not result in marked loss of ERK1/2 phosphorylation in either \(GNAQ\) or wild-type cells, thus the growth inhibiting effect of METi treatment in \(GNAQ\) mutant uveal melanoma cells may be mediated through a distinctly different signaling pathway. The MET receptor has been shown to signal through a myriad of other pathways (e.g., PI3K/AKT, STAT, Src, PLC) and is proposed to have cross-signaling effects on cell proliferation [21].

Figure 6. Effect of METi treatment on \(GNAQ\) mutant or wild-type uveal melanoma cell migration. (A) Representative images of distinct \(GNAQ\) wild-type (WT1 = Mel285, WT2 = Mel290) or mutant (Mut1 = Mel270, Mut2 = Mel 202) uveal melanoma cell line migration, stained following 0, 1 and 2.5 \(\mu M\) METi treatment. (B) Percent of \(GNAQ\) wild-type or mutant uveal melanoma cells that migrated following 0, 1 and 2.5 \(\mu M\) METi treatment.

doi:10.1371/journal.pone.0083957.g006
The elimination of the HGF induced PI3K phosphorylation following METi treatment (Figure 3) suggests a significant role for MET – PI3K/AKT signaling in growth and survival in uveal melanoma. Recent data supports a role for MET in uveal melanoma. The small molecule tyrosine kinase inhibitors XL147 or Crizotinib, which target MET, have been reported to show efficacy in mouse xenograft models of metastatic uveal melanoma [22,23].

GNAQ mutant cells clearly demonstrated the induction of cleaved PARP with either of the MEKi or METi treatment doses employed. However, combination MEKi+METi treatment resulted in greater cleaved PARP levels than either treatment alone in these cells. Among the wild-type cells, the MEL 285 demonstrated low levels of cleaved PARP with all treatments, with no enhancement observed with combination treatment, and GNAQ wild-type MEL 290 failed to demonstrate evidence of cleaved PARP with MEKi and/or METi treatment. Thus, GNAQ mutant uveal melanoma cell lines appear more sensitive to the combination of MEK and MET. These findings are consistent with the studies of Wu et al., who failed to demonstrate cleaved PARP induction with two distinct MET inhibitors (PHA-665752 and PF-02341066) or with shRNA knockdown of MET in two cell lines that have neither GNAQ nor GNA11 mutations (C918 and OCM1) [24]. Neither GNAQ nor GNA11 mutant uveal melanoma cell lines were tested in that study. The observation in the current study that PARP cleavage occurred primarily in GNAQ mutant cells with MEKi and/or METi treatment suggests that the pathways driven by these molecules are important for cell survival, in a manner distinct from wild-type cells. However, only modest increases in the sub-G0 fraction of cells was noted, indicating the presence of counteracting anti-apoptotic mechanisms [25].

Two prior studies that investigated the use of single agent MET inhibitors, and used cell lines that did not harbor GNAQ mutations (e.g., OCM1, OCM3, OCM8, C918); but rather have BRAF mutations [published and unpublished data] [8,24]. Using standard sequencing techniques BRAF mutations are not observed in primary uveal melanoma tumors, although two groups have identified BRAF mutations in a small number of primary uveal melanoma samples using more sensitive techniques [26,27]. The clinical relevance of these findings awaits further investigation.

Prior to this study it was not known whether combined inhibition of MEK and MET caused further growth inhibition in G-alpha protein mutant uveal melanoma cells or whether the combination treatment further diminished the migration in these cells. Our data show that combined treatment of uveal melanoma cells with small molecule MEK and MET inhibitors simultaneously enhances the growth inhibitory effect of each agent in a GNAQ mutation-dependent manner, and yet the small molecule METi also served as a powerful blocker of migration in both GNAQ mutant and wild-type cells. This study greatly enhances our knowledge of the role played by the MET pathway in uveal melanoma cells, with new information proposing the combination treatment of small molecule MEK and MET inhibitors in cells harboring G-alpha protein gene mutations. Thus a combination of MEK and MET inhibitors can control uveal melanoma cell growth and migration better than individual treatment with each of these inhibitors alone.

Finally, the data presented in this paper may also be placed in the greater context of primary drug resistance given the observation by Straussman et al, that HGF derived from stromal cells results in reactivation of the mitogen-activated protein kinase (a.k.a., Erk1/2) and phosphatidylinositol-3-OH kinase (PI(3)K)-AKT signaling pathways, in BRAF inhibitor treated cutaneous BRAF mutant melanoma cells [28]. Likewise, the HGF-MET axis may play an important role in mediating resistance to MEK inhibition, and thus combination MEKi and METi treatment would be a rationale strategy under these conditions.

Supporting Information

**Figure S1 MET mRNA expression levels in BRAF mutant uveal melanoma cells.** (A) The top panel shows MET mRNA expression and the lower panel shows actin mRNA expression as determined by RT-PCR. Images are from the same gel at the same exposure. (B) MET protein expression levels in BRAF mutant uveal melanoma cells. The top panel shows MET protein expression and the lower panel shows actin protein expression as determined by western-blot. (TIF)

**Figure S2 Effect of siRNA knockdown of MET protein and cell viability.** (A) Effect of non-targeting versus MET targeting siRNA on MET protein level as determined by western-blot. (B) Effect of MET targeting siRNA on cell viability relative to non-targeting siRNA control. (TIF)

**Figure S3 Effect of METi treatment on BRAF mutant uveal melanoma cell migration.** (A) Representative images of migrated BRAF mutant (OCM1 or OCM8) uveal melanoma cells stained following 0 or 1 μM METi treatment. (B) Effect of MEKi treatment on GNAQ wild-type or mutant uveal melanoma cell migration. Images of migrated wild-type (WT1 = Mel295, WT2 = Mel290) or mutant (Mut1 = Mel270, Mut2 = Mel 202) GNAQ uveal melanoma cells stained following 0 or 50 nM MEKi treatment. (TIF)

Author Contributions

Conceived and designed the experiments: CC EAG SEW. Performed the experiments: CC EAG SEW. Performed the experiments: CC. Analyzed the data: CC EAG SEW. Contributed reagents/materials/analysis tools: CC EAG SEW. Wrote the paper: CC EAG SEW.

References

1. van den Bosch T, Kilic E, Paridaens D, de Klein A (2010) Genetics of uveal melanoma and cutaneous melanoma: two of a kind? Dermatol Res Pract 2010: 360136.

2. Singh AD, Turell ME, Topham AK (2011) Uveal melanoma: trends in incidence, treatment, and survival. Ophthalmology 118: 1881–1885.

3. Oaken MD, Wooley LA, Long MD, Duan S, Council ML, et al. (2008) Oncogenic Mutations in GNAQ Occur Early in Uveal Melanoma. Investigative Ophthalmology & Visual Science 49: 5230–5234.

4. Van Raamsdonk CD, Bezrookove V, Green G, Bauer J, Gaugler L, et al. (2009) Oncogenic Mutations in GNAQ Occur Early in Uveal Melanoma. Investigative Ophthalmology & Visual Science 49: 5230–5234.

5. Van Raamsdonk CD, Liu JY, Holby SE, Weil MH, Eaves DL, et al. (2009) Mutations in GNA11 in Uveal Melanoma. New England Journal of Medicine 360: 2191–2199.

6. Woodman SE (2012) Metastatic Uveal Melanoma: Biology and Emerging Treatments. The Cancer Journal 18: 148–152 110.1097/PPO.1090- b1015e31824bed3126.

7. Khalili JS, Yu X, Wang J, Hayes BC, Davies MA, et al. (2012) Combination Small Molecule MEK and PI3K Inhibition Enhances Uveal Melanoma Cell Death in a Mutant GNAQ and GNA11 Dependent Manner. Clinical Cancer Research.

8. Moch A, Hofmann F, Harbott J, Staib H, Schrader J, et al. (2009) Frequent somatic mutations of GNAQ in uveal melanoma and blue nevi. Nature 457: 599–602.

9. Van Raamsdonk CD, Liu JY, Holby SE, Weil MH, Eaves DL, et al. (2009) Mutations in GNA11 in Uveal Melanoma. New England Journal of Medicine 360: 2191–2199.

10. Frequent somatic mutations of GNAQ in uveal melanoma and blue nevi. Nature 457: 599–602.

11. Woodman SE (2012) Metastatic Uveal Melanoma: Biology and Emerging Treatments. The Cancer Journal 18: 148–152 110.1097/PPO.1090-b1015e31824bed3126.

12. Moch A, Hofmann F, Harbott J, Staib H, Schrader J, et al. (2009) Frequent somatic mutations of GNAQ in uveal melanoma and blue nevi. Nature 457: 599–602.
10. Mallikarjuna K, Pushparaj V, Biswas J, Krishnakumar S (2007) Expression of Epidermal Growth Factor Receptor, Ezrin, Hepatocyte Growth Factor, and c-Met in Uveal Melanoma: An Immunohistochemical Study. Current Eye Research 32: 281–290.

11. Economou MA, All-Ericason C, Bykov V, Girnita L, Bartolazzi A, et al. (2005) Receptors for the Liver Synthesized Growth Factors IGF-1 and HGF/SF in Uveal Melanoma: Intercorrelation and Prognostic Implications. Investigative Ophthalmology & Visual Science 46: 4572–4575.

12. Topçu-Yılmaz P, Karタイトル H, Sağlam A, Stylenreazouli F, Hacıcak G (2010) Correlation of clinicopathological parameters with HGF, c-Met, EGFR, and IGF-IR expression in uveal melanoma. Melanoma Research 20: 126–132 110.1097/CMR.1090b1013e328335a328916.

13. De Waard-Siebinga I, Blom DJ, Griffioen M, Schrier PI, Hoogendoorn E, et al. (1995) Establishment and characterization of an uveal-melanoma cell line. Int J Cancer 62: 155–161.

14. Chen PW, Murray TG, Salgaller ML, Ksander BR (1997) Expression of MAGE genes in ocular melanoma cell lines. J Immunother 20: 265–275.

15. Kan-Mitchell J, Mitchell MS, Rao N, Liggett PE (1989) Characterization of uveal melanoma cell lines that grow as xenografts in rabbit eyes. Invest Ophthalmol Vis Sci 30: 629–634.

16. Griewank KG, Yu X, Khalili J, Sozen MM, Stemke-Hale K, et al. (2012) Genetic and molecular characterization of uveal melanoma cell lines. Pigment Cell & Melanoma Research 25: 182–187.

17. Fujiwara S, Sugano K (1997) Expression of c-met proto-oncogene in primary colorectal cancer and liver metastases. Jpn J Clin Oncol 27: 327–332.

18. Folberg R, Kadkol SS, Frenkel S, Valyi-Nagy K, Jager MJ, et al. (2008) Authenticating cell lines in ophthalmic research laboratories. Invest Ophthalmol Vis Sci 49: 4697–4701.

19. Griewank KG, Yu X, Khalili J, Sozen MM, Stemke-Hale K, et al. (2012) Genetic and molecular characterization of uveal melanoma cell lines. Pigment Cell Melanoma Res 25: 182–187.

20. Woodman SE, Stemke-Hale K, Williams M, Jager MJ, Ksander B (2010) Targeting Genetic Aberrations in Uveal Melanoma. ARVO Annual Meeting; Ft Lauderdale, Florida, 2010 869/D1101.

21. Trusolino L, Bertotti A, Comoglio PM (2010) MET signalling: principles and functions in development, organ regeneration and cancer. Nat Rev Mol Cell Biol 11: 834–848.

22. Surriga O VR, Ambrosini G, Schwartz G (2013) Abstract 2706: Crizotinib, a selective c-Met inhibitor, prevents metastasis to the liver and lungs in a primary uveal melanoma mouse model. Cancer Research 73: Supplement 1.

23. Yeh I GK, Ding VW, Bastian BC (2011) Abstract 5387: XL184: c-Met inhibition is effective in a mouse xenograft model of metastatic uveal melanoma. Cancer Research 71: Supplement 1.

24. Wu X, Zhou J, Rogers AM, Jaine PA, Benedettini E, et al. (2012) c-Met, epidermal growth factor receptor, and insulin-like growth factor-1 receptor are important for growth in uveal melanoma and independently contribute to migration and metastatic potential. Melanoma Research 22: 123–132 110.1097/CMR.1090b1013e328335a328916.

25. Tang HL, Tang HM, Mak KH, Wu S, Wang SS, et al. (2012) Cell survival, DNA damage, and oncogenic transformation after a transient and reversible apoptotic response. Molecular Biology of the Cell 23: 2240–2252.

26. Maat W, Kliche E, Luyten GPM, de Klein A, Jager MJ, et al. (2008) Pyrophosphorolysis Detects B-RAF Mutations in Primary Uveal Melanoma. Investigative Ophthalmology & Visual Science 49: 23–27.

27. Janssen CS, Sibbett R, Henriquez FL, McKay IC, Kemp EG, et al. (2008) The T1799A point mutation is present in posterior uveal melanoma. Br J Cancer 99: 1673–1677.

28. Straussman R, Morikawa T, Shee K, Barzily-Rokni M, Qian ZR, et al. (2012) Tumour micro-environment elicits innate resistance to RAF inhibitors through HGF secretion. Nature 487: 500–504.