Epidermal Growth Factor Immunotherapy: Exploring the Effects of NB-UVB Exposure

E. Costantini1*, C. D’Angelo1, R. Muraro1, M. De Tursi1, C. Natoli1, M. Auriemma2, M. Di Nicola1, P. Amerio2 and Reale Marcella1

1Department of Medical, Oral and Biotechnological Sciences, D’Annunzio University of Chieti–Pescara, Italy
2Department of Medicine and Aging Sciences, D’Annunzio University of Chieti–Pescara, Italy

Received date: June 25, 2018; Accepted date: July 12, 2018; Published date: July 17, 2018

Abstract

Introduction: Inflammatory skin reactions are the most common issue in Cetuximab-treated patients. Many evidences show the correlation with NB-UVB exposure, but the mechanisms are not completely understood. The aim of this study is the evaluation of the effects of NB-UVB exposure on inflammatory cytokines modulation during EGFR immunotherapy.

Materials and methods: Human immortalized keratinocytes, HaCaT cells, were treated with 10 g/ml of Cetuximab for 24 hrs and NB-UVB exposed. Viability, ROS production and pro-inflammatory cytokines gene-expression and release were assessed. An additional, in vivo evaluation of Minimal Erythema Dose in Cetuximab-treated patients was performed.

Results: NB-UVB exposure activates the inflammatory skin response in HaCaT cells, inducing the over-expression of pro-inflammatory mediators. Reduced cells viability, ROS production and expression and release of selected cytokines after Cetuximab-treatment. The co-treatment, with NB-UVB and Cetuximab, show the recovery with baseline conditions. In vivo evaluation underlines that patient’s skin reactions are not changed after NB-UVB exposure.

Conclusion: In conclusion, NB-UVB doesn’t exacerbate papulopustular reactions developed during anti-EGFR treatment, both in vivo and in vitro evaluations. A better understanding of this relationship, between NB-UVB exposure and dermatological side effects, could throw a new light on sun exposure limitation, improving Cetuximab-treated patients QoL and favoring therapeutic adherence.

Keywords: NB-UVB; HaCaT; Cytokines; Cetuximab

Introduction

The Epidermal growth factor receptor (EGFR) is over-expressed in many solid tumors and is associated with poorer prognosis and outcomes [1]. The growing investigation on EGF inhibitors application, in anticancer therapy, has been related to their high specificity for EGFR.

Target therapies for EGFR, with inhibitors, determine a down-regulation of EGFR signal pathway, regulating cell differentiation, proliferation, migration, angiogenesis and apoptosis [2,3] improving the safety profile compared to conventional chemotherapeutic agents, generally interfering with DNA and RNA synthesis.

EGFR-inhibitors can be classified into two categories, the monoclonal antibodies (mAb) and the tyrosine kinase inhibitors (TKI). mAbs acts by a competition with the endogenous ligands, epidermal growth factor (EGF) and transforming growth factor alpha (TGF), leading to receptor dimerization and inhibition of EGFR activity, with a reduction in cancer proliferation, invasion, metastasis, angiogenesis and inhibition of apoptosis [4]. In contrast, TKI mechanism is represented by the competition with the adenosine triphosphate (ATP) to bind the tyrosine kinase region, determining intracellular catalytic activity inhibition of the receptor [5,6].

Cetuximab, an anti-EGFR, is a chimeric human-murine immunoglobulin (IgG1) approved by the American Food and Drug Administration (FDA) for the treatment of EGFR positive metastatic colon rectal cancer, either alone or in combination with chemotherapy [4,7]. Cetuximab, can be used in patients without mutations in RAS gene, since it was demonstrated to inhibit the downstream RAS-signaling cascade and ERK activation [8].

Patients, treated with immunotherapy, show dermatological side effects, with an incidence ranging from 80% to 86% of cases [3,5]. These could be motivated by the highly EGFR expression in the epidermis, at the basal cell layer. Since EGF may control skin immunity [4], the dermatological side effects may be due to an inflammatory response mediated by keratinocyte-derived cytokines, macrophages, mast cells and granulocytes recruitment, followed by long-term cutaneous effects development. Due to its role, in the regulation of barrier function, innate host defense and inflammatory reaction, in epidermal keratinocytes, EGFR inhibition induces the release of inflammatory mediators and consequently infiltration of immune cells.

The most common form of cutaneous adverse events is the papulopustular acneiform rash [9,10] associated with mAb anti-EGFR,
followed by xerosis, fissures, pruritus, paronychia, and blepharitis [11,12].

The papulopustular acniform rash generally occurs after 1-2 weeks from the onset of therapy (on average 8 to 10 days), peaking at 3 weeks after treatment initiation and decreases in severity after 6-8 weeks. Post-inflammatory skin alterations, such as erythema and hyperpigmentation, are long-standing sequel that can last for months or years. Typically seborrheic areas (face, scalp, trunk) and/or sun-exposed, are preferred, while rarely found in the upper and lower extremities of the abdomen, gluteal region, saving palm plantar sites [3,9]. The histopathological evaluation of papules and pustules shows an asceptic supportive folliculitis: in the earliest stages of treatment with anti-EGFR, the inflammatory infiltrate is mainly composed of T lymphocytes, which is subsequently replaced by an inflammatory infiltration composed mainly of neutrophils distributed in the dermis, particularly around the follicles. Microbiological tests have demonstrated the absence of infection, confirming that this is a sterile process [10]. Papulopustular skin rash may be accompanied by post-inflammatory hyperpigmentation, telangiectasia, and erythema [9].

These features develop as the dose-dependent effect, and the severity of anti-EGFR inhibitors associated rash, was classified using the US Common Terminology Criteria for Adverse Events (CTCAE), in 5 grades [9]. Moreover, a strong correlation between the presence and severity of the cutaneous effects with patient compliance and survival is reported. Patients with dermatological rash appear to be the most responsive to the anti-EGFR treatments [13,14]. Dermatological side effects are persistent during life and have a deep impact on the patient’s quality of life (QoL) and may result in a significant physical and emotional discomfort. Studies have suggested that 30% of patients needs to stop therapy due to cutaneous adverse events [15,16].

Sun exposure could exacerbate anti-EGFR toxicities, as suggested by clinical and experimental data, through the deregulation of cell cycle and cells proliferation induction [17-19].

Based on wavelengths, the solar ultraviolet radiations are classified as ultraviolet A (UVA), ultraviolet B (UVB) and ultraviolet C (UVC). UVB, ranging from 295 to 315 nm is one of the main components of solar UV that reach only the very superficial layers of skin, composed predominantly of keratinocytes [20]. The UVB exposure may be responsible for a dose-related effect, such as a premature skin aging, local and systemic immunosuppression, and promotion of different biological events [17] such as sunburn, inflammation, cell and/or tissue damage, cell death and skin cancer [18,21,22]. Their immunosuppressive properties are due to the interaction with DNA leading to single-stranded DNA breaks and intra-stranded DNA base-cross linking via formation of several photoproducts [21] with immunosuppressive role and production of reactive oxygen species (ROS) responsible for epigenetic alterations [21-25]. In cancer cells, UVB-exposure may be responsible for a conformational change in EGFR, causing an altered EGFR signaling pathway transduction. Moreover, the 280-315 nm wavelengths are commonly used in the clinical application for the treatment of cutaneous conditions due to their immunosuppressive action [21]. Immunomodulatory effect of UVB on the skin is widely used for the treatment of psoriasis and eczema. Phototherapy represents the main and often a favorite therapeutic option for several dermatoses. In vitro studies were conducted to evaluate the beneficial effects of UV-irradiation on skin cells, suggesting that UV-treatment on cell culture activates genes related to cell division and immune responses. UV-exposure might be beneficial for wound healing and restoration of skin homeostasis, besides its anti-inflammatory, antioxidant, and also acting as modulator of keratinocyte-melanocyte cross-talks [26,27]. The balance between negative and positive effect UVB-exposure, remain to be clarified.

It is well known that keratinocytes are able to produce a variety of cytokines and chemokines, according to the inducing stimulus.

In this study, we investigated whether a specific range of light, called “narrow-band ultraviolet” (NB-UVB) that theoretically can reach the optimum response minimizing skin redness if compared with the “broad-band” (BB-UVB) in the range of UVB [28]. Despite NB-UVB has been developed more than 30 years ago, the exact mechanism of its therapeutic action remains poorly understood.

The 90% of NB-UVB irradiation is absorbed by the epidermis, making keratinocytes the most important players in mediating biological activity, moreover NB-UVB also induce systemic effects on different cells. Lymphocytes or Langerhans cells are a primary target of NB-UVB systemic action [21].

Most of NB-UVB effects are attributed to its influence on the immune system, with immunosuppressive effect than BB-UVB on natural killer cell activity, cytokine production and peripheral blood mononuclear cells responses. NB-UVB irradiation it has been shown to determine the reduction in circulating T-helper (Th)1, Th17 and CD8+ cytotoxic T cell levels with an increase in Treg levels, in psoriatic patients compared to healthy control [29].

NB-UVB irradiation is involved in the balance in cytokines network that plays a critical role in the health and in the body’s response to the stimulus. The ability of NB-UVB in the suppression of cytokines production is involved in the shift of the immune response toward T-helper Th2-like responses. Indeed the expression of TNF-α, IL-17A, IL-6, IL-1 β, INF-γ, and IL-10, the most important cytokines involved in processes as wound healing, infections, inflammation, autoimmune and inflammatory skin conditions can be downregulated by NB-UVB therapy [30].

NB-UVB exposure could modulate the expression and production of inflammatory cytokines and chemokines, in human model of keratinocytes, in presence of Cetuximab.

Materials and methods
HaCaT cell culture and treatments
The human immortalized non-tumorigenic keratinocytes cell line (HaCaT) were supplied by Cell Lines Service (CLS, Germany) and growing in High-glucose Dulbecco’s modified Eagle’s medium supplemented with 10% fetal bovine serum, 100 U/ml penicillin and 100 mg/ml of streptomycin (Sigma-Aldrich, USA), in humidified atmosphere containing 5% CO2 at 37°C. For all experiments we used the same passage of HaCaT cells and over-night in serum starving condition, before starting treatments and incubations. Cells were incubated for 24 hrs in presence of EGF (4 nM) or Cetuximab (10 μg/ml) whether or not UVB-irradiation (0.06 J/cm²).

Western Blotting
In HaCaT whole cells lysate, proteins were extracted using ice-cold RIPA lysis buffer (Sigma-Aldrich, USA) added with protease inhibitor cocktail (Sigma-Aldrich, USA). 50 μg of proteins were separated to sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-
PAGE 10%) and then transferred for 1hr to a nitrocellulose membrane (Bio-Rad, USA). EGFR immunoreactive proteins were evaluated using the following antibodies: EGFR 1005 (Santa Cruz Biotechnology, USA), EGFR phosphorytrosine-1068 (Cell Signaling Technology, USA) and horseradish peroxidase-conjugated secondary antibodies, goat anti-rabbit (Santa Cruz Biotechnology, USA). Specific signals were detected and analyzed with ChemidocTM Touch Imaging System (Bio-Rad, USA) using a chemiluminescence system, ClarityTM-ECL Western Blotting Substrate (Bio-Rad, USA).

UVB-exposure

A Sunlamp 70 230 V (CIRCA)-0.8 A-50 Hz; NB-UVB, containing three Philips 20 W ultraviolet-B TL20/01RS bulbs was used for cells irradiation. According to the literature and controlled using a UV light-meter Vario-control (Waldmann, Germany) positioned at the same distance from the UV source as the cells. HaCaT cells were grown in 6-well microplates (Eppendorf, Germany) until confluence. Cells were exposed to UVB-irradiation (0.06 J/cm<sup>2</sup>), replacing the culture medium with 1 ml of Dulbecco’s PBS, w/o CaCl<sub>2</sub> and MgCl<sub>2</sub> (Sigma-Aldrich, USA). Non-irradiated cell controls were also covered with Dulbecco’s PBS for the same time. After irradiation, the culture medium has been replenished and cells were grown for 24 hrs prior testing.

Measurement of cell viability and ROS production

HaCaT cells viability was determined by using the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay (Sigma-Aldrich, USA). HaCaT were plated in 96-well microplates in quadruplicate and exposed to the three different treatments, EGF (4 nM), Cetuximab (10 μg/ml) and UVB (0.06 J/cm<sup>2</sup>), for 24 hrs. After treatments, the MTT-based assay was developed according to the procedures indicated by the manufacturer. Color absorbance was quantified at 570 nm using the spectrophotometer module of the GloMax<sup>®</sup>-Multi Microplates Detection Reader (Promega Corporation, USA). Cell viability was also estimated using a Bürker counting chamber with trypan blue, vital dye. To assess, reactive oxygen species (ROS), 10 μM of carboxy-2,7′-dichloro-dihydro-fluorescein diacetate (DCFH-DA) fluorescent probe has been added to the cells treated with Cetuximab. The plate was irradiated with UVB (0.06 J/cm<sup>2</sup>) and intracellular ROS production was evaluated at 485 nm (excitation) and 527 nm (emission) wavelengths with the GloMax<sup>®</sup>-Multi Microplates Detection Reader (Promega Corporation, USA).

Enzyme-linked immunosorbent assay

Cytokine and chemokine concentration, in the supernatant obtained from HaCaT cells, after NB-UVB and Cetuximab treatments, were determined using ELISA assay. The Quantikine ELISA kit for Human CXCL8/IL8 and CCL5/RANTES (R&D Systems, USA) were used following manufacturer’s instructions. Relative absorbance was measured at 450 nm and the cytokines and chemokines concentration calculated using a reference standard curve.

Extraction of nucleic acid, cDNA synthesis, and real-time PCR

Total RNA was extracted from HaCaT according to the classical protocol phenol-chloroform, using QIAzol Lysis Reagent (Qiagen, Germany). After extraction, the total RNA concentration and purity were assessed using 1 μl of the sample and performed with NanoDrop 2000 (Thermo Scientific, USA). 1 μg/μl of cDNA, in a total volume of 20 μl, was obtained using the QuantiTect Reverse Transcription Kit (Qiagen, Germany) following the manufacturer’s instruction. Template for the real-time PCR was 1:2 diluted and evaluated for gene expression of IL-1β, IL-6, IL-8, CCL-2, and CCL-5, using RPS18 as housekeeping gene (Table 1). All primers sequences were designed using the AlleleID Primer Biosoft International (Premier Biosoft, USA) and IDT SciTools, Inc. (Tema Ricerca, Italy). Real-time PCR was carried out using the Mastercycler<sup>®</sup> ep realplex (Eppendorf, Germany) according to the manufacturer’s protocol. The cycle conditions for real-time PCR were 95°C for 10 minutes, followed by 40 cycles of 95°C for 15 seconds, and 60°C for 1 minute. Each reaction was performed in triplicate.

### Table 1: Gene sequences

<table>
<thead>
<tr>
<th>Forward Primer Sequence (5’-3’)</th>
<th>Reverse Primer Sequence (5’-3’)</th>
<th>Amplicone size</th>
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</tr>
<tr>
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</tr>
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</tr>
<tr>
<td>CCL-2</td>
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### Statistical analysis

All data were summarized as the mean and standard deviation (SD) from three independent experiments. For each real-time experiment to give an indication of the precision of the fold change, calculated with 2^-ΔΔCt method, the 95% confidence interval (95% CI) was determined. Statistical comparison between values from exposed to non-exposed cells was analyzed by Student t-test for unpaired data. P-values were corrected for multiple comparisons when appropriated.

Analysis of variance (ANOVA) test was applied to evaluate statistical significance differences in CCL-5 and IL-8 release between different experimental conditions.
In all statistical tests, the threshold of statistical significance will be assumed equal to $p=0.05$. Data were analyzed using R-2.9.2 (R Core Development System: http://www.r-project.org).

**Patients**

Ten patients with colorectal or pulmonary carcinoma, candidates for Cetuximab-treatment were recruited at the Department of Oncology of the Clinical “SS. Annunziata” Hospital of Chieti. Eligibility criteria included patients who were scheduled to start first-line chemotherapy (oxaliplatin-/irinotecan-based chemotherapy) combined with Cetuximab, administered on a weekly schedule, with a 400 mg/m$^2$ initial loading dose, followed by 250 mg/m$^2$ weekly infusions until disease progression, and presenting skin Fitzpatrick’s Phontotype less than IV, as assessed by the Dermatological Clinic of the SS Annunziata Hospital University of Chieti. This study was conducted in accordance with the guidelines from the International Conference on Harmonization-Good Clinical Practice and the Declaration of Helsinki. The study protocol was approved by the Ethical Committee of the University “G d’Annunzio”, Chieti-Pescara, Italy. All patients provided written informed consent before participating in the study.

**Evaluation of photosensitivity**

To determine the photosensitivity of the patients, a solar simulator (model 16S SolarLight Co) was used. The device is capable of delivering an intensity of 2 mW/cm$^2$ to simulate a solar exposure at doses gradually increasing of UVB on the surface of the forearms. Patients undergoing stimulation with the solar simulator had phenotype according to Fitzpatrick <IV . The timing of NB-UVB exposure was reported in Table 2. Evaluation of photosensitivity was carried out by handheld skin color probe (CORTEX TECHNOLOGY) for melanin detection (pigment present volume index), erythema (reddening index based on the amount of hemoglobin) and CIE*L*a*B*, three relative parameters for complexity (CIE*L) and skin coloring (CIE* and CIE*B). Color Space Color Lab (CIELAB) software is used to read colorimetric data.

<table>
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<tr>
<th>J/cm$^2$</th>
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<th>0.16</th>
<th>0.2</th>
<th>0.25</th>
<th>0.31</th>
<th>0.39</th>
<th>0.49</th>
</tr>
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<tbody>
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<td>1'40''</td>
<td>2'</td>
<td>2'35''</td>
<td>3'15''</td>
<td>4'</td>
</tr>
</tbody>
</table>

**Table 2:** Timing of irradiation

**Results**

**EGFR phosphorylation in HaCaT cells**

Presence and functionality of EGFR in HaCaT cells were assessed after stimulation with EGF (4 nM), UVB (0.06 J/cm$^2$) exposure and Cetuximab-treatment. Figure 1a shows EGFR expression that appears to be down-regulated after EGF-treatment. Instead, EGFR phosphorylation (EGFR phosphotyrosine-1068), evaluated by Western Blotting analysis, was increased. UVB exposure enhanced levels of phosphorylation of EGFR were observed. No effects after Cetuximab-treatment were noticed. Interestingly, a combination of NB-UVB exposure and Cetuximab resulted in increased EGFR activation, higher than Cetuximab or NB-UVB alone (Figure1b). Our results indicate that EGFR is constitutively expressed in HaCaT cells and they represent a good model to study, in vitro, the effects of Cetuximab-treatment.

![Figure 1: Effect of EGF-stimulation on EGFR expression and activity. (a) EGFR gene-expression in EGF-stimulated HaCaT cells, data are shown as expression levels, relative to control cells. (b) Representative image of Western blotting for EGFR, p-EGFR, and β-actin, using protein extracted after EGF (E), Cetuximab (C), UVB (U) and their association.](image)

**Cells viability and ROS production**

Cells viability was evaluated by 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay, and by cell counts by Bürker chamber in HaCaT cells, as described above. EGF-treatment, NB-UVB and their combination induce an increased cell survival and proliferation, concurrently inducing the increase of EGFR phosphorylation. Cetuximab-treatment causes a reduction of cells growth, and the association between NB-UVB and Cetuximab show no variation in cell viability (Figure 2). Taken together, these results suggest that NB-UVB can counteract the Cetuximab-dependent reduction of cells viability.

Oxidative stress and production of ROS, was also investigated. NB-UVB exposure stimulates ROS generation, while Cetuximab-treatment causes about a 5% of the reduction in NB-UVB induced ROS generation (data not shown).
Figure 2: Effect of treatments on HaCaT cell viability. MTT assay was performed on HaCaT cell after 24 hrs of stimulation with EGF, UVB, Cetuximab. Control experiments were performed setting the 100% survival. All experiments were run in triplicate. Data are expressed as mean ± SD.

Cytokines and chemokines gene expression in NB-UVB exposed and cetuximab-treated cells

The expression levels of pro-inflammatory cytokines and chemokines, in HaCaT cells treated with Cetuximab, alone or in association to NB-UVB, were evaluated by Real-time PCR assay, after 24 hrs. Cetuximab-treatment reduces the expression of IL-1β (p=0.0001), IL-8, CCL2 and CCL-5, a very slow increase of IL-6 expression, while NB-UVB exposure is responsible for the opposite expression, increasing significantly IL-1β (p=0.003), IL-8 (p=0.002) and CCL-5 (p=0.001), in relation to unexposed cells. Association between NB-UVB and Cetuximab-treatments on keratinocytes cell culture, increased significantly CCL-2 expression, compared to control cells. IL-1β, IL-6, IL-8 and CCL-5 expression levels were not influenced (Figure 3).

Figure 3: Gene expression levels of pro-inflammatory cytokines/chemokines. Cells were incubated with Cetuximab (10 µg/ml), UVB (0.06 J/cm²) or their combination, for 24 hrs. Expression levels of each cytokine in HaCaT are reported in relation to untreated cells. Data are representative of pooled experiments and expressed as means of expression levels and relative 95% confidence interval (95% CI). *p<0.05 Student's t-test vs. control.
Chemokines quantification

CCL-5 and IL-8, the major chemokines involved in the chemotaxis and inflammatory process, supporting acneiform reactions, were also quantized in cell cultures supernatant. The mean concentration of IL-8 decrease in presence of Cetuximab, compared to untreated cells, on the other hand, increased production of CCL-5 was observed. After NB-UVB exposure, levels of IL-8 and CCL-5 concentrations, any significant variation was detected. Co-stimulation is responsible for an increased amount of both chemokines (Figure 4).

Figure 4: ELISA analysis of cell-free supernatants. IL-8, CCL-5 levels are expressed as the mean ± SD.

Photosensitivity of patients

In order to assess the sensitivity to solar light (Solar simulator model 16S Solar Light Company, USA), a pivotal experiment on oncologic patients, was performed. Patients enrolled were evaluated for photosensitivity before, during and after Cetuximab-treatment. We did not find differences in the intensity of erythema after NB-UVB exposure in all patients (n=10). The minimal erythema dose (MED) did not change in any patients, from baseline thought out the treatment time (Table 3).

<table>
<thead>
<tr>
<th>Exposure time</th>
<th>T2-T1</th>
<th>p-value</th>
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</thead>
<tbody>
<tr>
<td>1'</td>
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</tr>
<tr>
<td>1'20''</td>
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<td>0.92</td>
</tr>
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<td>1'40''</td>
<td>0.5</td>
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</tr>
<tr>
<td>2'</td>
<td>1.5</td>
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<td>2'35''</td>
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<tr>
<td>3'15''</td>
<td>0.2</td>
<td>0.89</td>
</tr>
<tr>
<td>4'</td>
<td>1.4</td>
<td>0.63</td>
</tr>
</tbody>
</table>

Table 3: MED evaluation

Discussion

The anti-EGFR drugs represent a recognized treatment for a variety of tumour types and have established efficacy as a single agent and in combination with chemo/radio/hormone therapy. The efficacy of anti-EGFR therapies was related to EGFR and the activated signalling cascades involved in cell proliferation, differentiation, and apoptosis. These drugs are able to increase the progression-free survival, overall patient survival, to improve clinical outcome and quality of life in cancer patients. Unfortunately, besides its needed effects, also some unwanted effects were observed during anti-EGFR therapy. The adverse events may interfere with the quality of life of patients, reducing the compliance of the therapeutic regimen [16]. Sometimes, due to the severity of side effects, temporary suspension of the biological therapy could be recommended and, in 8%-17% of the cases, the complete suspension of the therapy is required. Anti-EGFR dose reduction or discontinuation, affect the treatment outcomes.

Since EGFR was expressed in the keratinocytes, the follicular epithelium, the most common side effects of anti-EGFR drugs are skin problems such as an acne-like rash on the face and chest during treatment. To ensure patient compliance, it could be helpful to identify the anti-EGFR role in cutaneous adverse events development. Treatments for dermatological side effects may include topical [17] and systemic antibiotic, steroids or topical immune modulator agents [31,32].

Although NB-UVB phototherapy represents one of the main treatments for an inflammatory skin condition, such as psoriasis and vitiligo, with minimal systemic toxicities [33]. According to different guidelines [33] and patient-oriented skin, adverse effects care treatments, exposure to UVB is not recommended in patients during immunotherapy. Several studies reported that following Cetuximab or others anti-EGFR treatment, patients may be sensitized to UVB exposure [3,34,35], and solar ultraviolet exposure has been linked to the increased skin inflammatory observed in these patients. Recently, the specific pathways activated by UVB-phototherapy, have been better understood, highlighting the link with the alteration of the cytokines profile at systemic level [36]. It is possible to explain that the increased sensitization could be due to the ability of the UVB to determine the inflammation onset, with associated vasodilatation and immune cells
recruitment at the skin level. UV-radiation also activates VEGF signalling involving EGF/PI3K pathway [37,38] and MAPK pathway [3].

The HaCaT cells, due to the common characteristic with normal keratinocytes, may allow a widely used in vitro model for skin pattern, producing a well-organized and differentiated pseudo-stratified epithelium [16,34].

To evaluate if NB-UVB exposure may modulate inflammatory mediators in anti-EGFR treated cells, we have analyzed the expression of pro-inflammatory cytokines in HaCaT cells, exposed to NB-UVB or Cetuximab alone and in association.

Our study indicates that the NB-UVB and Cetuximab treatments play a different role in defining the activation and functionality of keratinocytes. Following NB-UVB exposure there is increase cells viability and an important reduction of Cetuximab-induced cell death. First, we underline the higher EGFR phosphorylation UVB-induced, and then the induction of intracellular ROS formation. In accord with Yoshizumi et al. [39], the increase of IL-1, IL-8, CCL-2 and CCL-5 production and expression in HaCaT cells, was observed after 24 hrs of NB-UVB exposure. Otherwise the inhibition of EGFR by incubation with Cetuximab, in our in vitro skin model showed a reduction in cytokines and chemokines expression and release [40]. Inhibition of EGFR reduced also ROS production, likely with the downstream of Ras/Nox1 cascade [41,42]. Thus, in our model, Cetuximab reduced inflammatory mediators, in accord with previous studies conducted by El-Abaseri et al. and other researchers we observed that NB-UVB exposure may revert the Cetuximab-dependent EGFR protein reduction, followed by restoration of IL-1, IL-6 and IL-8 expression to basal levels [3,19,23].

CCL-5 and CCL-2, involved in the early inflammatory response [43-45] and in neutrophils, basophils, and monocytes recruitment in the inflammatory site, was not significantly reduced by the combination of NB-UVB exposure and Cetuximab-treatment.

Collectively, these data shows the tighter regulation of inflammatory response following NB-UVB exposure in Cetuximab-treated cells. The effect of NB-UVB, as immune-modulator, during Cetuximab-treatment, may open new perspectives for patients treated with EGF immunotherapy.

Our in vitro results are supported by the preliminary in vivo evaluation of the MED. Evaluation of patient’s phenotype and photosensitivity, before and after Cetuximab administration and exposure to the solar simulator device, show any additional harmful effects.

In conclusion, our in vivo and in vitro result, seem to deny the additive effect of NB-UVB exposure on dermatological side effects, during anti-EGFR therapy. Nevertheless, to better understanding the pathogenesis of skin reactions during anti-EGFR treatment, and effects of NB-UVB exposure, further in vivo studies must be performed. The knowledge of the pathogenic mechanisms of skin reactions could promote new strategies to prevent and manage adverse reaction in cancer patients, improving their QoL and favoring therapeutic adherence.

Acknowledgment

It is gratifying to honor the memory of Dr. Giovina Vianale with this work, which was substantially inspired by long dialogues with her. This study was supported by grants from the Università degli Studi G. D’Annunzio di Chieti-Pescara (Italy). The following statements should be used: E.C. and P.A. conceived and designed the experiments; M.D.T. and C.N. collected the samples; C.D.A and E.C. performed experiments; M.A. performed the in vivo exposure; M.R. and P.A. analyzed the data; M.R. and P.A. contributed reagents/materials/analysis tools; E.C., P.A., M.R. wrote the paper; R.M. contributed to discussion.

References


