

The Sensitizers Nickel Sulfate and 2,4-dinitrofluorobenzene Increase CD40 and IL-12 Receptor Expression in a Fetal Skin Dendritic Cell Line

Ana Luísa Vital,^{1,2} Margarida Gonçalo,^{3,4} Maria Teresa Cruz,^{1,2} Américo Figueiredo,³ Carlos B. Duarte,² and Maria Celeste Lopes^{1,2}

Dendritic cells (DCs) are antigen-presenting cells (APCs) capable of capturing haptens and to process and present them to T lymphocytes. In order to sensitize T cells for contact hypersensitivity (CHS), skin DCs suffer a maturation process with modifications on their surface molecules. The aim of this work was to evaluate changes induced by two contact sensitizers, 2,4-dinitrofluorobenzene (DNFB) and nickel sulfate (NiSO₄), and a non-sensitizer 2,4-dichloronitrobenzene (DCNB), on the protein levels of two activation markers, CD40 and IL-12 receptor (IL-12R), in a mouse skin dendritic cell line (FSDC). The expression of CD40 and IL-12R proteins was evaluated by western blot assay and direct immunofluorescence microscopy. The results showed that CD40 and IL-12R expression increased significantly after cell exposure to NiSO₄ and DNFB, although DNFB exhibited a stronger activity. There was no effect with DCNB. The epidermal cytokine granulocyte-macrophage colony-stimulating factor (GM-CSF), also used in the experiments, slightly increased the expression of both CD40 and IL-12R and when tested together with the sensitizers the effect was partially additive. The results suggest that the sensitizers DNFB and NiSO₄ are directly involved on the changes of the surface markers CD40 and IL-12R in skin DCs, during the sensitization phase of CHS, and this effect may be enhanced by GM-CSF. In contrast, no effect was observed with DCNB.

KEY WORDS: Dendritic cells; CD40; IL-12 receptor; GM-CSF; skin sensitizers; Nickel; DNFB; DCNB.

INTRODUCTION

Contact hypersensitivity (CHS) is a common inflammatory skin disease that occurs as a consequence of increasing exposure to exogenous chemicals. It is a dendritic cell-dependent, T cell-derived and cytokine-mediated skin inflammation (Enk, 1997;

¹Faculdade de Farmácia, Universidade de Coimbra, Rua do Norte, 3000-295, Coimbra, Portugal.

²Centro de Neurociências e Biologia Celular, Universidade de Coimbra, 3004-517, Coimbra, Portugal.

³Faculdade de Medicina (Serviço de Dermatologia), Hospital da Universidade de Coimbra, 3000-075, Coimbra, Portugal.

⁴To whom all correspondence should be addressed. Margarida Gonçalo, Faculdade de Medicina (Serviço de Dermatologia), Hospital da Universidade de Coimbra, 3000-075, Coimbra, Portugal. Fax: + 351-239-480217, E-mail: mmgoncalo@netcabo.pt.

Watanabe *et al.*, 2002). Langerhans cells (LCs) and keratinocytes are epidermal cells that produce, constitutively or upon exogenous stimulation, a wide range of cytokines (Kimber *et al.*, 1995). These cytokines interact with appropriate receptors expressed by LCs becoming crucial in the sensitization phase and in initiation of the CHS (Watanabe *et al.*, 2002).

LCs and other skin dendritic cells (DCs) are antigen presenting cells (APCs) that play an important role in determining the cutaneous response to small chemicals applied at the skin surface (Enk, 1997; Watanabe *et al.*, 2002). These cells capture the small molecular weight exogenous chemical (the hapten) by phagocytosis or by pynocytosis, process it into a small hapten-modified peptide and combine this allergen with a MHC molecule (Lipscomb and Masten, 2002). At this stage skin DCs are very efficient in the uptake and processing of the antigens but poor stimulators of primary T cell responses and, in the skin they would not find naïve T cells for priming. Therefore, they undergo maturation to become potent APCs and migrate to the skin draining lymph nodes searching the appropriate naïve T cells for sensitization (Girolomoni *et al.*, 1995; Pieri *et al.*, 2001; Satthaporn and Eremin, 2001).

The migration and maturation process of skin DCs involves profound changes on the molecules that are expressed at their cell surface (Banchereau and Steinman, 1998), namely MHC class II (Herouet *et al.*, 2000) and costimulatory molecules (Coutant KD *et al.*, 1999; Tuschl and Kovac, 2001), and on those secreted to the extracellular milieu, including cytokines (Pichowski *et al.*, 2000; De Smedt *et al.*, 2001) and metalloproteinases (Ratzinger *et al.*, 2002).

In vitro studies, using whole human skin models, cultures of DCs derived from circulating CD34+ cells or immortalized animal skin dendritic cells (Girolomoni *et al.*, 1995; Herouet *et al.*, 2000; Tuschl *et al.* 2000; De Smedt *et al.*, 2001), have shown a direct effect of some sensitizing chemicals, namely nickel, chromium, 2,4-dinitrochlorobenzene (DNCB), 2,4-dinitrofluorobenzene (DNFB) and 2,4,6-trinitrobenzene sulfonic acid (TNBS), on the phenotypic and functional characteristics of DCs. They upregulate DC expression of MHC class II and costimulatory molecules, namely CD54, CD86, CD40, CD83 (Coutant *et al.*, 1999; Herouet *et al.*, 2000; De Smedt *et al.*, 2001; Tuschl and Kovac, 2001). Some reports indicate that sensitizing chemicals may increase IL-1 β and IL-12 secretion by DC cells although this is still a matter of debate, since these responses were not always observed (Coutant *et al.*, 1999; Aiba *et al.*, 2000; Pichowski *et al.*, 2000; De Smedt *et al.*, 2001; Tuschl and Kovac, 2001).

In this work, using a mouse fetal skin dendritic cell line (FSDC), we studied the changes in the expression of membrane-associated receptors induced by chemicals with different skin sensitizing properties, alone or in the presence of GM-CSF. GM-CSF is one of the epidermal cytokines produced by keratinocytes after epidermal application of skin sensitizers, being responsible for DC maturation (Kimber *et al.*, 1995; Pieri *et al.*, 2001). The present work showed that nickel sulfate (NiSO₄), the most prevalent allergen in the human population, and DNFB, a potent experimental sensitizer, upregulated DC expression of the CD40 protein and of the IL-12 receptor (IL-12R). In contrast 2,4-dichloronitrobenzene (DCNB), an analogous of DNFB which is considered a non-sensitizer (Becker *et al.*, 1994), did not exhibit these effects. The epidermal cytokine GM-CSF, which induces the maturation of FSDC (Kimber *et al.*, 1995; Jonuleit *et al.*, 1996), also slightly increased CD40 and IL-12R expression, and this effect was additive with the response to the sensitizers.

MATERIALS AND METHODS

Materials

DNFB was purchased from Sigma–Aldrich Química (Madrid, Spain). DCNB and NiSO₄ were from Sigma Chemical Co (St. Louis, MO, USA). The mouse recombinant GM-CSF was from R & D systems (Minneapolis, MN, USA). The rabbit anti-mouse CD40 immunoglobulin and the rabbit anti-human antibody to IL-12 receptor beta 1 subunit (IL-12Rβ1) were obtained from Santa Cruz Biotechnology (Santa Cruz, CA). The mouse monoclonal antibody against actin and the protease inhibitor cocktail were from Roche (Carnaxide, Portugal). Fetal calf serum was from Biochrom KG (Berlin, Germany) and trypsin from Invitrogen (Paisley, UK). The fluorescein isothiocyanate (FITC)-conjugated swine anti-rabbit immunoglobulin and the swine serum were from DAKO (Copenhagen, Denmark). The mounting medium for fluorescence, Vectashield, was obtained from Vector Laboratories, Inc. (Burlingame, CA, U.S.A.). The ECL western blotting analysis system, the horseradish peroxidase-conjugated donkey anti-rabbit immunoglobulin and the X-ray films were from Amersham Biosciences (Carnaxide, Portugal). The horseradish peroxidase-conjugated goat anti-mouse immunoglobulin was from Pierce (Illinois, USA). Triton® X 100 and paraformaldehyde were from Merck (Lisboa, Portugal). All other reagents were from Sigma Chemical Co (St. Louis, MO, USA).

Cell culture

The fetal mouse skin dendritic cell line FSDC, kindly supplied by Dr. G. Girolomoni (Laboratory of Immunology, Istituto Dermopatico dell'Imacolata, IRCCS, Rome, Italy), is a skin dendritic cell precursor with antigen presenting capacity (Girolomoni *et al.*, 1995). The cells were cultured in endotoxin free Iscove's Modified Dulbecco's Medium (IMDM), supplemented with 10% (v/v) fetal calf serum, 1% (w/v) glutamine, 3.02 g/l sodium bicarbonate, 100 µg/ml streptomycin and 100 U/ml penicillin. For western blot analysis, FSDC were plated at 2×10^6 cells/well, in six-well culture plates, whereas for immunofluorescence analysis, FSDC cells were grown on a Lab-Tek chamber slide with cover (0.2×10^6 cells/slide).

The effect of DNFB and NiSO₄ on the expression of CD40 and IL-12R in FSDC was determined by western blot and immunocytochemistry, after incubating the cells with the compounds for 24 hr. The concentrations of DNFB (1 µg/ml) and NiSO₄ (50 µg/ml) used were those that produced maximal effect and no toxicity (data not shown), as evaluated by the reduction of the tetrazolium bromide salt, 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide (MTT) (Mosmann, 1983).

NiSO₄ was dissolved in culture medium. DNFB and DCNB were dissolved in DMSO in a stock solution (10 mg/ml). Before incubation of the cells with these compounds (DNFB and DCNB) the stock solution was diluted in sterile PBS (10 ×) and the final concentrations of DNFB and DCNB were further diluted in the culture medium containing the FSDC cells (1000 ×). Therefore, the concentration of DMSO incubated with cells never exceeded 0.01%. Under these experimental conditions the assay of cellular MTT reduction (a viability assay) in the presence of this concentration of DMSO revealed no toxic effect and this concentration of

DMSO alone was without effect on CD40 and IL-12R expression (data not shown).

Western blot analysis

Total cell lysates were obtained after harvesting the cells in a sonication buffer containing 0.32 M sucrose, 10 mM Tris-HCl (pH 7.5), 1 mM ethylene-diaminetetraacetic acid, 1 mM dithiothreitol, 0.1 mM phenylmethylsulfonylfluoride and the protease inhibitor cocktail. Then, the lysates were incubated on ice for 30 min and sonicated to disrupt the cells. Protein concentration was determined using the bicinchoninic acid method.

In brief, protein samples were denatured and separated on a 10% (v/v) SDS-PAGE, and transferred to a PVDF membrane. The membrane was blocked with 5% (w/v) dry milk in Tris-buffered saline with 0.1% (v/v) Tween 20, for 1 hr. The levels of CD40 and IL-12R β 1 proteins were detected using a rabbit anti-mouse CD40 (1:500) and a rabbit anti-human IL-12R β 1 (1:500) antibody, for 1 hr, followed by incubation with a horseradish peroxidase-conjugated donkey anti-rabbit antibody (1:5000). The immunocomplexes were visualized by the ECL chemiluminescence method. To demonstrate equivalent protein loading the membranes were stripped and reprobed with an anti-actin antibody (1:10000), followed by incubation with a horseradish peroxidase-conjugated goat anti-mouse immunoglobulin (1:25000).

Immunofluorescence microscopy (immunocytochemistry)

After incubation in the absence or in the presence of the appropriate stimuli, as indicated in Figs. 2c and 3c, the cells were washed with PBS and fixed with PBS containing 4% (w/v) sucrose and 4% (w/v) paraformaldehyde, for 15 min. FSDC were then permeabilized in PBS containing 1% (v/v) Triton® X 100, for 10 min. Non-specific binding was blocked by incubation of the cells with PBS supplemented with normal swine serum (1:20) and 0.5% (w/v) BSA-PBS, for 45 min, at room temperature. Cells were then incubated with a rabbit anti-mouse CD40 immunoglobulin (1:50) or with a rabbit anti-human IL-12R β 1 antibody (1:50), for 90 min at room temperature. After rinsing with PBS, the cells were incubated with FITC-conjugated swine anti-rabbit immunoglobulin (1:40) in 0.5% BSA-PBS, for 45 min. The cells were rinsed again and mounted with the mounting medium for fluorescence, Vectashield. Cells labeled with FITC-anti CD40 or with FITC-anti IL-12R β 1 antibodies were photographed on a Zeiss Axiovert 200 microscope. Control experiments consisted of processing the same preparations as described, except for the omission of the primary antibody, and resulted in no specific staining.

Data analysis

Results are presented as mean \pm SEM of the indicated number of experiments. Mean values were compared using one-way ANOVA and the Bonferroni's multiple comparison test. The significance level was * p < 0.05, ** p < 0.01 and *** p < 0.001.

RESULTS

Effect of NiSO₄, DNFB and DCNB on CD40 expression in FSDC

Stimulation of the cells with NiSO₄ (50 µg/ml) or DNFB (1 µg/ml), for 24 hr, increased significantly the expression of CD40, by 32 and 50%, respectively (Fig. 1a, lanes 3 and 4) as demonstrated by western blot. In order to determine how FSDC maturation affects the expression of CD40 induced by sensitizers, the cells were pre-incubated with GM-CSF (100ng/ml), for 24 hr, before stimulation with NiSO₄ or DNFB. GM-CSF was previously shown to induce the differentiation and maturation of DCs progenitors (Kimber *et al.*, 1995; Jonuleit *et al.*, 1996; Banchereau and Steinman, 1998) and, in particular, of this cell line (Girolomoni *et al.*, 1995; Pieri

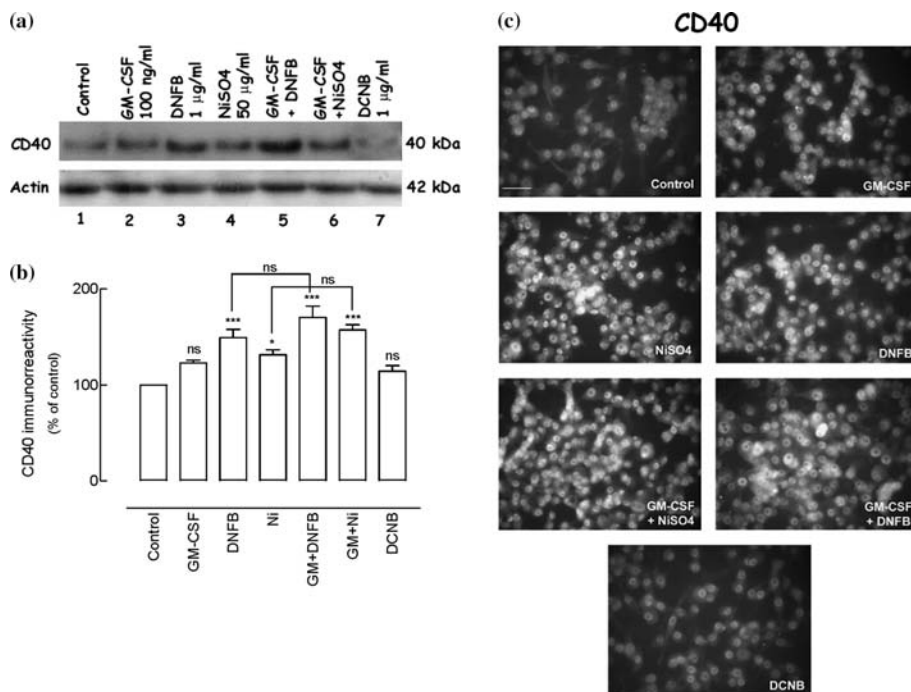


Fig. 1. Effect of GM-CSF on NiSO₄- and DNFB-induced CD40 protein expression. (a) Western blot analysis of CD40 protein. FSDC cells (2×10^6 cells) were incubated in culture medium in the absence (control, lane 1) or in the presence of GM-CSF (100 ng/ml, lane 2), DNFB (1 µg/ml, lane 3) or NiSO₄ (50 µg/ml, lane 4). Where indicated, the cells were stimulated with GM-CSF in the presence of DNFB (1 µg/ml, lane 5) or NiSO₄ (50 µg/ml, lane 6). Total cell extracts were electrophoresed through SDS-PAGE and subjected to western blot analysis using an anti-CD40 antibody, as described in “Materials and Methods”. The blot shown is representative of three blots yielding similar results. The blot was digitally generated using a HP ScanJet 5p and processed in the Adobe Photoshop 7.0 program. (b) The bands were quantified with an image analyser. The values are means \pm SEM from three independent experiments, where **p* < 0.05, ***p* < 0.01 and ****p* < 0.001, as determined by one-way ANOVA with Bonferroni’s multiple comparison test. (c) Immunofluorescence analysis of CD40 protein. FSDC cells (0.2×10^6 cells) were incubated on a Lab Tek chamber with cover, under the same conditions as described in A. Immunostaining was performed as described in “Materials and Methods”. Immunoreactivity means CD40 protein expression (% of control). Scale bars = 50 µm.

et al., 2001). As shown in Fig. 1a and b, incubation of FSDC cells with GM-CSF, for 48 hr, increased the expression of CD40 by 23%, but this effect did not reach statistical significance (Fig. 1a, lane 2). However, when FSDC cells were incubated with NiSO₄ (50 µg/ml) or DNFB (1 µg/ml), for 24 hr, together with GM-CSF, after a 24 hr pre-incubation with the cytokine, the increase in the expression of CD40 was slightly higher but not statistically different from that observed in the presence of the sensitizers alone (58 and 70%, respectively) (Fig. 1a, lanes 5 and 6). Interestingly, the small effect of GM-CSF on the expression of CD40 appears to be additive with the response to the sensitizers. When the cells were incubated with the non-sensitizer DCNB the observed effect was not different from the control (Fig. 1a, lane 7).

In Fig. 1c, immunofluorescence microscopy experiments performed under the same experimental conditions confirmed all these results, demonstrating an upregulation of CD40 expression induced by the sensitizers NiSO₄ and DNFB, and a slightly higher increase in the expression of the protein in cells pre-incubated with GM-CSF. Immunofluorescence also confirmed that the non-sensitizer DCNB had no significant effect on this model.

Effect of NiSO₄, DNFB and DCNB on IL-12R expression in FSDC

The effect of NiSO₄, DNFB and DCNB on the expression of IL-12R in FSDC cells is summarized in Fig. 2a and b. In FSDC cells incubated with NiSO₄ (50 µg/ml) or with DNFB (1 µg/ml), for 24 hr, the expression of IL-12R was increased by 45 and 72%, respectively, when compared with control cells, i.e., cells incubated with culture medium (Fig. 2a, lanes 1, 4 and 5). DCNB had no effect on this model (Fig. 2a, lane 7). In order to determine how FSDC maturation affects the expression of the IL-12R induced by sensitizers, the cells were pre-incubated with GM-CSF (100 ng/ml), for 24 hr, before stimulation for an additional 24 hr with NiSO₄ or DNFB, together with the cytokine. When the cells were incubated only with GM-CSF, for 48 hr, we observed a non-statistically significant increase of IL-12R expression of 23% (Fig 2a, lane 2). When FSDC cells were incubated with the sensitizers NiSO₄ (50 µg/ml) or DNFB (1 µg/ml), for 24 hr, together with GM-CSF, after a 24 hr pre-incubation with the cytokine, the increase in IL-12R expression (66 and 90%, respectively) was higher but not statistically different from that observed in cells stimulated only by the sensitizers (Fig. 2a, lanes 3 and 6). As observed for the expression of CD40, the small effect of GM-CSF on IL-12R expression was additive with the response to the sensitizers. When the cells were incubated with the non-sensitizer DCNB the observed effect was not different from the control (Fig. 2a, lane 7).

In Fig. 2c, immunofluorescence microscopy confirmed all of these results, demonstrating an upregulation of IL-12R expression induced by the sensitizers NiSO₄ and DNFB in cells pre-incubated or not with GM-CSF. The immunofluorescence also confirmed that the non-sensitizer DCNB did not affect the expression of the IL-12R.

DISCUSSION

FSDC is a fetal mouse skin derived dendritic cell line, which represents a model of a skin DC precursor. These cells are sensitive to and require cytokine activation to

become functionally efficient APCs (Girolomoni *et al.*, 1995). We have previously shown that FSDC are sensitive to the epidermal cytokine GM-CSF (Cruz *et al.*, 2001; Vital *et al.*, 2003), which is involved on DC differentiation and maturation (Kimber *et al.*, 1995; Jonuleit *et al.*, 1996; Banchereau and Steinman, 1998).

In this work, we studied the effect of two strong contact sensitizers, DNFB and NiSO₄, on the expression of two membrane-associated receptors, CD40 and IL-12R, in FSDC. The results show that both skin sensitizers upregulated the expression of the two cell surface molecules, CD40 and IL-12R, whereas the non-sensitizer DCNB had no significant effect (Figs. 1 and 2). In another set of experiments we tested the effect of pre-incubating FSDC with GM-CSF, which induces FSDC maturation (Kimber *et al.*, 1995; Jonuleit *et al.*, 1996; Pieri *et al.*, 2001), on the response to the

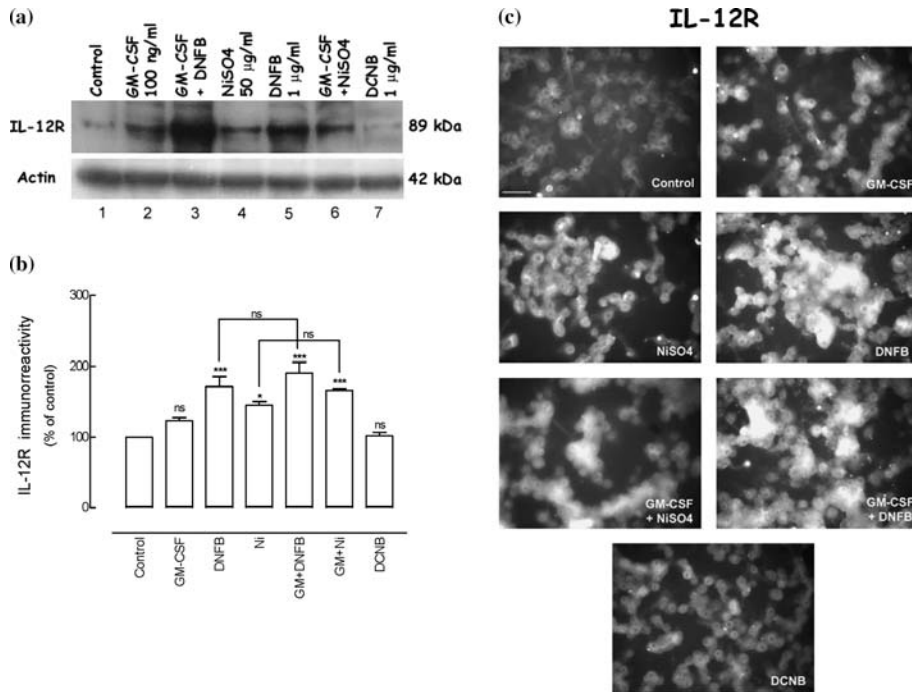


Fig. 2. Effect of GM-CSF on NiSO₄ and DNFB-induced IL-12R expression. (a) Western blot analysis of IL-12Rβ1 protein. FSDC cells (2×10^6 cells) were incubated in the absence (control, lane 1) or in the presence of GM-CSF (100 ng/ml, lane 2), NiSO₄ (50 μg/ml, lane 4) or DNFB (1 μg/ml, lane 5). Where indicated, the cells were stimulated with GM-CSF in the presence of DNFB (1 μg/ml, lane 3) or NiSO₄ (50 μg/ml, lane 6). Total cell extracts were electrophoresed through SDS-PAGE and subjected to western blot analysis using an anti-IL-12Rβ1 antibody, as described in materials and methods. The blot shown is representative of three blots yielding similar results. The blot was digitally generated using a HP ScanJet 5p and processed in the Adobe Photoshop 7.0 program. (b) The bands were quantified with an image analyser. The values are means \pm SEM from three independent experiments, where * $p < 0.05$, ** $p < 0.01$ and *** $p < 0.001$, as determined by one-way ANOVA with Bonferroni's multiple comparison test. (c) Immunofluorescence analysis of IL-12R protein. FSDC cells (0.2×10^6 cells) were incubated on a Lab Tek chamber with cover, under the same conditions as described in A. Immunostaining assay was performed as described in materials and methods. Immunoreactivity means IL-12R protein expression (% of control). Scale bars = 50 μm.

sensitizers. The results show that GM-CSF alone slightly increased CD40 and IL-12R expression, and this effect was additive with the result of cell stimulation only with the allergens (Figs. 1 and 2).

The upregulation in the expression of CD40 observed here (Fig. 1) is in agreement with the results obtained in other DCs (Coutant *et al.*, 1999). CD40 is a membrane-associated receptor belonging to the tumor necrosis factor-receptor (TNF-R) family that is expressed broadly, including in DCs (van Kooten and Banchereau, 2000). CD40 interacts with the CD40 ligand (CD40L) on naïve T cells in a process important for initiating primary immune responses (Banchereau *et al.*, 2000). CD40 activation by CD40L is critical for maintaining DC viability, to promote their maturation through the upregulation of costimulatory molecules and to control their migration to the draining lymph nodes (Moodycliffe *et al.*, 2000; Mann *et al.*, 2002). CD40 ligation also activates cytokine gene expression in DCs and promotes IL-6 and IL-12 secretion, which is important in driving T cell development and in the initiation of primary immune responses (Mann *et al.*, 2002; Wesa and Galy, 2002). Therefore, CD40 could be regarded as a surface marker of DC activation. In the present work we also found that DNFB induced a higher increase in CD40 expression than NiSO₄ (Fig. 1), which is in agreement with the *in vitro* and *in vivo* stronger activity of this skin sensitizer (Yokozeki *et al.*, 1995).

FSDC were cultured with GM-CSF in the absence or in the presence of the contact sensitizers, in order to mimic the *in vivo* events. During CHS, epidermal keratinocytes provide a wide range of cytokines that influence the differentiation of cells of the dendritic lineage, in particular Langerhans cells. GM-CSF is one of these cytokines, relevant for the maintenance of the differentiated state of LCs within epidermis (Pieri *et al.*, 2001). Also, a variety of environmental stimuli, including epicutaneous contact with the skin sensitizers used in the present work, induce epidermal keratinocytes to release inflammatory and chemotactic cytokines and growth promoting cytokines, namely GM-CSF (Corsini and Galli, 1998). In fact, our results show that GM-CSF alone slightly increased CD40 expression, a marker of DC maturation, which is in agreement with a previous report showing that GM-CSF slightly upregulated CD40 expression in purified LCs (Salgado *et al.*, 1999). The slightly higher CD40 upregulation in the presence of GM-CSF together with sensitizers indicate that the effect of these two stimuli was additive. The gene that codes for CD40 contains NF- κ B binding sites in its enhancer promoter regions (Yoshimura *et al.*, 2001; Nguyen and Benveniste, 2002), and this may constitute the mechanism whereby CD40 protein levels are upregulated by allergens and by this epidermal cytokine. Indeed, GM-CSF, NiSO₄ and DNFB activate the NF- κ B transcription factor in FSDC (Cruz *et al.*, 2001; Cruz *et al.*, 2002; Vital *et al.*, 2003). NiSO₄ induces the translocation of p65, RelB and cRel from the cytosol to the nucleus, whereas GM-CSF induces the translocation of p50, p52 and RelB proteins (Cruz *et al.*, 2001; Cruz *et al.*, 2002). The differential activation of NF- κ B components by GM-CSF and NiSO₄ may explain the additive effect of GM-CSF and NiSO₄ on CD40 protein levels. In contrast, GM-CSF and DNFB activate the same NF- κ B elements, suggesting that the regulation of CD40 by the cytokine and the allergen may involve additional signaling events.

The sensitizers DNFB and NiSO₄ also increased the expression of IL-12R, as shown in Fig. 2. In addition, GM-CSF upregulated this effect when was tested

together with the sensitizers. The IL-12R complex is composed of two beta-type cytokine receptor subunits, IL-12R β 1 and IL-12R β 2 (Presky *et al.*, 1996; Nagayama *et al.*, 2000) and has been described in dendritic cells, T cells, NK cells and other PBMC (Desai *et al.*, 1992; Chua *et al.*, 1994; Chua *et al.*, 1995; Presky *et al.*, 1996). IL-12R β 1 mediates the signaling events induced by IL-12 on DCs, which promote their functional activation (Grohmann *et al.*, 1998; Nagayama *et al.*, 2000). However, the effect of skin sensitizers on IL-12R expression had never been addressed before.

Previous reports have shown that contact sensitizers like DNCB, which has a similar sensitizing potential as DNFB, can induce IL-12p40 subunit production by some APCs (Warbrick *et al.*, 1999). Therefore, DC upregulation of IL-12R induced by the sensitizers could be fundamental for the engagement of IL-12 produced by the DC itself or by other cells in the vicinity, thus promoting IL-12 signaling and functional DC activation. Accordingly, studies concerning the regulation of IL-12R have shown that an increase in IL-12 production and release is sufficient to induce expression of the two subunits of IL-12R, beta 1 and beta 2, by an autocrine mechanism in T cells (Rogge *et al.*, 1997; Kim *et al.*, 2001). One of the signaling pathways activated by IL-12 after binding to its receptor involves members of NF- κ B in DCs (Grohmann *et al.*, 1998; Yoshimura *et al.*, 2001) and, therefore, it is possible that the expression of IL-12R is dependent on NF- κ B. Since GM-CSF, NiSO₄ and DNFB also activates NF- κ B in FSDC (Cruz *et al.*, 2001; Cruz *et al.*, 2002; Vital *et al.*, 2003), this may be the mechanism whereby the cytokine and the sensitizers upregulate IL-12R protein levels in the cells.

IL-12 is a pivotal cytokine during antigen presentation, inducing functional activation of DCs, by the upregulation of the expression of MHC-II molecules (Grohmann *et al.*, 1998; Kelleher and Knight, 1998; Nagayama *et al.*, 2000) and the secretion of several cytokines, namely GM-CSF, IL-1 β , IL-6, TNF- α , IFN- γ and IL-12p40 subunit itself (Grohmann *et al.*, 1998; Kelleher and Knight, 1998; Nagayama *et al.*, 2000). Since haptens induce IL-12 production by antigen presenting cells, namely DCs (Warbrick *et al.*, 1999), the upregulation of IL-12R in FSDC reported here suggests that the sensitizers cause a general upregulation of IL-12 function. This also suggests that IL-12 may play a key role in the response of DCs to the presence of the sensitizers.

In summary, we observed that the two skin contact sensitizers, DNFB and NiSO₄, increased FSDC expression of the two membrane-associated receptors, CD40 and IL-12R, whereas the non-sensitizer DNCB had no significant effect as expected. GM-CSF also increased in a small extent the expression of both proteins, and this effect was additive with the effect of the allergens. These results suggest that these skin contact sensitizers induced early specific phenotype changes that may represent an early-activation state of the DCs, which could be responsible for the initiation of the sensitization phase of CHS *in vivo*. In addition, these results also suggest that the epidermal cytokine GM-CSF is involved on these changes. Moreover, we interestingly observed that the higher expression of CD40 and IL-12R was obtained with the stronger sensitizer DNFB, whereas NiSO₄ induced a lower expression of those proteins. Our results suggest that upregulation of CD40 and IL-12R expression could also be a promising readout in prescreening for strong and moderate contact sensitizers and could help to develop an *in vitro* assay to reduce the

number of experimental animals used for ascertaining the sensitizing potential of environmental chemicals.

ACKNOWLEDGMENTS

This work was supported by the Portuguese Foundation for Science and Technology, project number 36373/99.

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