

Differentiation of normal and cancer cells induced by sulfhydryl reduction: biochemical and molecular mechanisms

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Abstract

We examined the morphological, biochemical and molecular outcome of a nonspecific sulfhydryl reduction in cells, obtained by supplementation of *N*-acetyl-L-cysteine (NAC) in a 0.1–10 mM concentration range. In human normal primary keratinocytes and in colon and ovary carcinoma cells we obtained evidences for: (i) a dose-dependent inhibition of proliferation without toxicity or apoptosis; (ii) a transition from a proliferative mesenchymal morphology to cell-specific differentiated structures; (iii) a noticeable increase in cell-cell and cell–substratum junctions; (iv) a relocation of the oncogenic β -catenin at the cell–cell junctions; (v) inhibition of microtubules aggregation; (vi) upregulation of differentiation-related genes including p53, heat shock protein 27 gene, *N*-myc downstream-regulated gene 1, E-cadherin, and down-regulation of cyclooxygenase-2; (vii) inhibition of c-Src tyrosine kinase. In conclusion, a thiol reduction devoid of toxicity as that operated by NAC apparently leads to terminal differentiation of normal and cancer cells through a pleiaide of converging mechanisms, many of which are targets of the recently developed differentiation therapy.

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Keywords: colon carcinoma; cyclooxygenase-2; junctions; ovary carcinoma; proliferation; c-Src

Abbreviations: HSP27, heat shock protein 27 gene; NAC, *N*-acetyl-L-cysteine; NHEK, normal human epidermal keratinocytes;

NDRG1, *N*-myc downstream-regulated gene 1; NSAIDs, non-steroidal anti-inflammatory drugs

Introduction

Molecular and biochemical evidences attribute several regulatory functions to the redox state of thiol groups. Modulation of protein activity by disulfide bond formation represents a universal and common mechanism to adopt a fast and efficient turn on-off of cellular signaling.¹ In particular, protein tyrosine phosphatases play a pivotal role in the control of cell cycle and differentiation, and were reported to be regulated through the redox state of their cysteine –SH residues.^{2,3} Also protein tyrosine kinases have an essential role in the control of cell proliferation and differentiation. For three of them, Yes, Fyn and c-Src, a redox regulation has been recently reported^{4,5} and particularly for c-Src, it apparently implies a reversible change in the redox state of a group of four cysteine residues close to the catalytic site.^{4–7}

The activity of transcription factors can also be modulated through thiol interactions: Binding of transcription factors to DNA may be sensitive to the redox state of sulfhydryl residues or to the presence of specific soluble thiols.^{1,8,9}

The study of the effect of sulfhydryl reduction is therefore an extremely attractive issue that we pursued by supplementing *N*-acetyl-L-cysteine (NAC) to different cell types, two human carcinoma lines and primary human epidermal keratinocytes. We found for the first time that this reductant treatment is able to induce cell differentiation through the modulation of several mechanisms, with modalities that depend on the cell type. Besides a dose-dependent inhibition of proliferation in the absence of both toxicity and apoptosis, our results show morphological, biochemical and molecular changes related to terminal differentiation. Following the current knowledge on the differentiation process, some of the observed changes can also represent the actual driving mechanism for differentiation.

The results obtained on normal primary cells suggest that the presented redox-driven mechanisms pertain to the general cell physiology and that the grade of cell malignancy could be related to the maintenance of the ability to respond to some of the outlined signals.

Results

Growth inhibition

Different NAC concentrations were tested on two human colon and ovary carcinoma cell lines, Caco-2 and OVCAR3, respectively, and primary normal human keratinocytes (NHEK). The results show that a decrease or a complete block of [³H]thymidine incorporation could be obtained by using effective NAC concentrations that were specific for each

cell type (Figure 1a). The proliferation of Caco-2 cells rapidly decreased after a double supplementation of NAC, at 24 h interval, reaching a block of about 95% at 10 mM, while a single supplementation with 5 mM NAC resulted in a block of about 85%. The OVCAR3 proliferation instead, was reduced to about 40% after two supplementations with 10 mM NAC. The NHEK were the most sensitive to NAC treatment, being almost completely blocked by a single supplementation of 2 mM NAC. When we also tested NAC concentrations lower than 0.5 mM, the cell proliferation was slightly increased, suggesting a trophic effect of NAC at these low concentrations. NAC supplementation did not alter cell viability, as tested by Trypan blue exclusion method, always between 94 and 96%. In addition, the cytometric analysis of Caco-2 and OVCAR3 cells did not show any evidence of apoptosis after

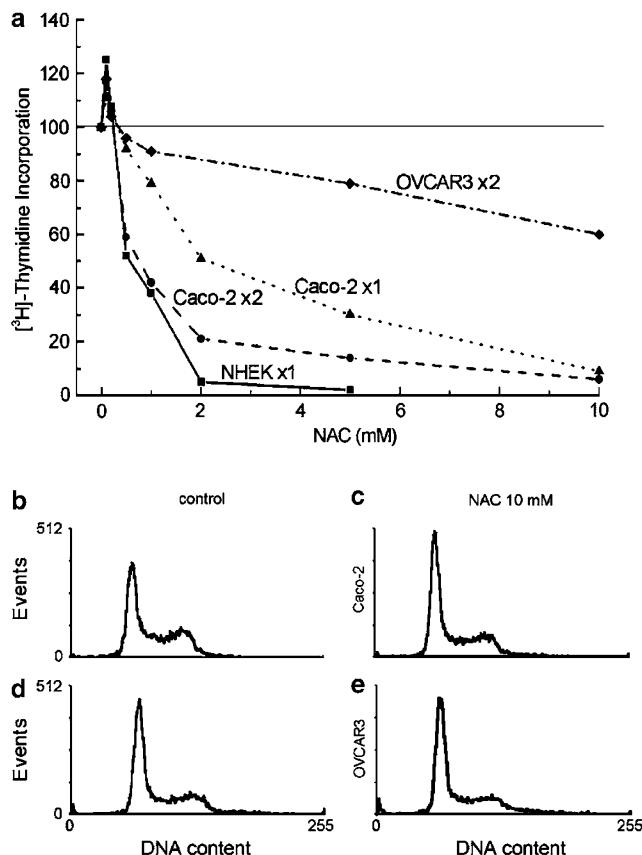


Figure 1 (a) $[^3\text{H}]$ Thymidine incorporation of cells treated with NAC. Data normalized to controls and reported as a function of NAC concentration for a single supplementation to NHEK, for a single (x1) and a double supplementation (x2, at 24 h interval) to Caco-2 and OVCAR3 cells. (b–e): Flow cytometry data of propidium iodide-labeled Caco-2 (b and c) and OVCAR3 (d and e) cells, controls (b and d) and treated with 10 mM NAC (c and e). OVCAR3 cells were treated twice at 24 h interval

supplementation with all tested NAC concentrations, even after two supplementations with 10 mM NAC, at 24 h interval, as in the case of OVCAR3 cells. In the Figure 1b–e we only report the cytometry data relative to controls and to the highest NAC concentration. The data analysis¹⁰ also indicated an increase in the relative concentration of cells in the G1 phase, of about 25 and 10% for Caco-2 and OVCAR3 cells, respectively. This last observation is consistent with all additional evidences of cell differentiation, reported in the following sections.

Morphological changes

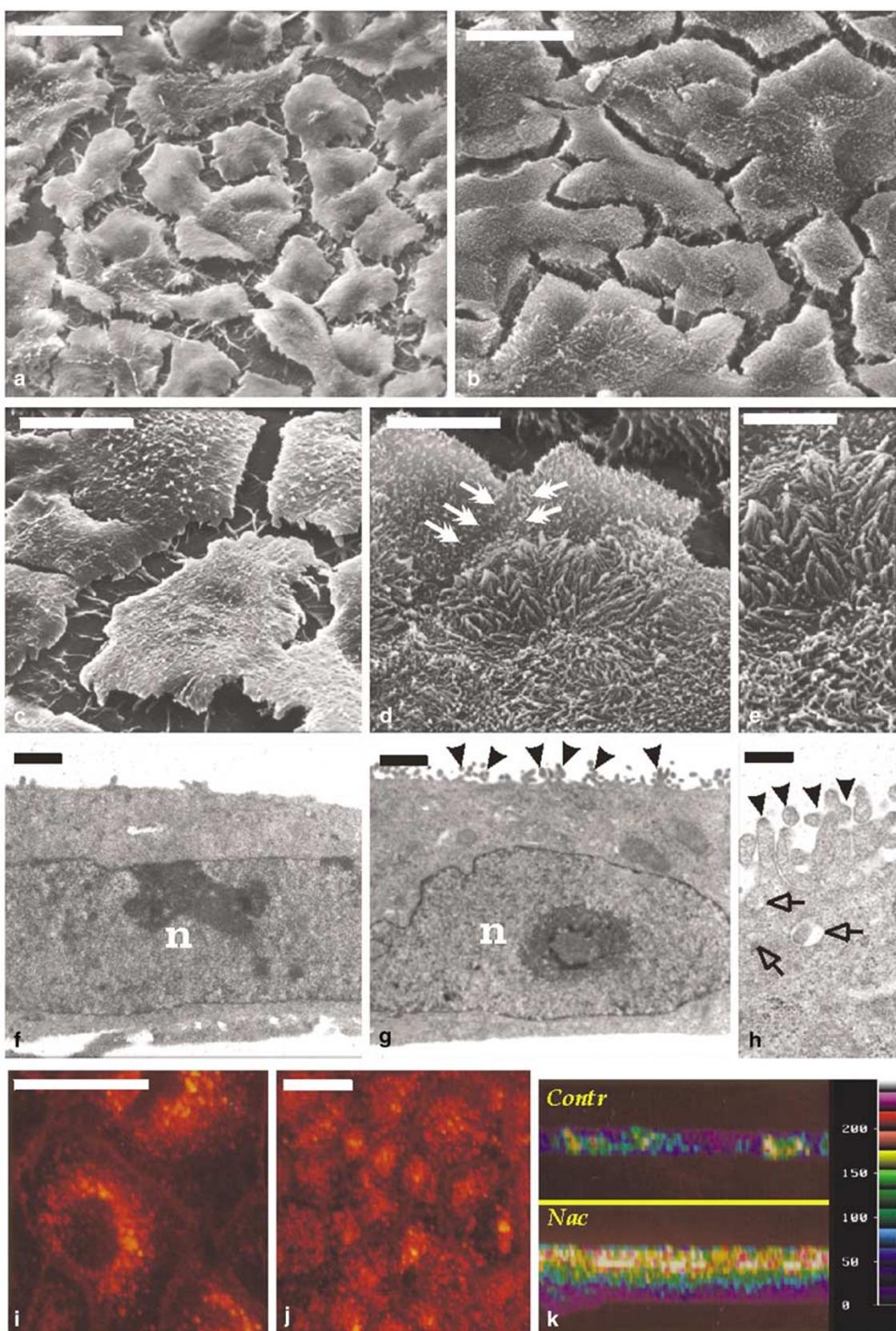
Proliferating Caco-2 cells display an irregular morphology and form several multiple cell layers with scarce microvillous structures and large intercellular space (Figure 2a and c). A single supplement of 2 mM NAC was given to Caco-2 cells, 24 h after seeding. Cells were then grown for three more days and analyzed. NAC-treated Caco-2 cells showed: (i) a regular polygonal morphology typical of end-stage differentiated cells; (ii) an increased thickness, about three times than that of controls (Figure 2a, b and k); (iii) a relevant number of brush border microvilli at the cell surface (Figure 2d, e and h); (iv) a well-developed secretory vesicles apparatus in the vicinity of the microvillous structures (Figure 2h–k); (v) a dramatically reduced intercellular space (arrows in Figure 2d); (vi) the location of the nucleus in the basal portion (Figure 2g), as it is in differentiated cells.

Primary NHEK have a limited life span in culture, of approximately 30 days, with a terminal differentiation achieved at the end of this period, concurrently with the formation of the cornified envelope. During the first passages, proliferating NHEK show a heterogeneous polygonal morphology, with a villous surface, relatively broad intercellular space and multiple cell layers can be observed with increasing cell density (Figure 3a and d). At 3 days after a 2 mM supplementation of NAC, NHEK grew flat in a thin monolayer, with a regular polygonal morphology (Figure 3b) and a smooth surface, losing the fine microvillous structure of proliferating keratinocytes (Figure 3e). We observed a relevant number of cells expelling their nucleus and exfoliating, as in the cornified envelope stage (Figure 3b, c and inset). In addition, NAC-treated cells had dramatically reduced intercellular space (Figure 3e).

Increase in cell–cell and cell–substratum adhesions

NAC-treated Caco-2 showed a remarkable increase in β -catenin and E-cadherin labeling (Figure 4c–f). As analyzed by the NIH ImageJ software, the integrated fluorescence

Figure 2 Effect of NAC treatment on the morphology of Caco-2 cells. (a–e) Scanning electron micrographs of controls (a and c) and of NAC-treated cells (b, d and e). Arrows in (d) point to cell–cell junctions. In (e) is a detail of brush border microvilli. (f–h): Transmission electron micrographs of control (f) and of NAC-treated cells (g and h). Arrowheads in (g) and (h) point to brush border microvilli and open arrows in (h) point to secretory vesicles. (i): Confocal fluorescence micrographs of intracellular secretory vesicles apparatus. Fluorescence intensity of the WGA-TRITC-labeled Golgi apparatus and endoplasmic reticulum in controls (i and top k) and in NAC-treated cells (j and bottom k). The apical distribution of secretory vesicles is well visualized in the longitudinal sectioning along the x/z plane of images (k) with the intensity color scale on the right. Bars: 25 μm in (a, b, i, j); 10 μm in (c and d); 5 μm in (e); 1 μm in (f, g); 300 nm in (h)



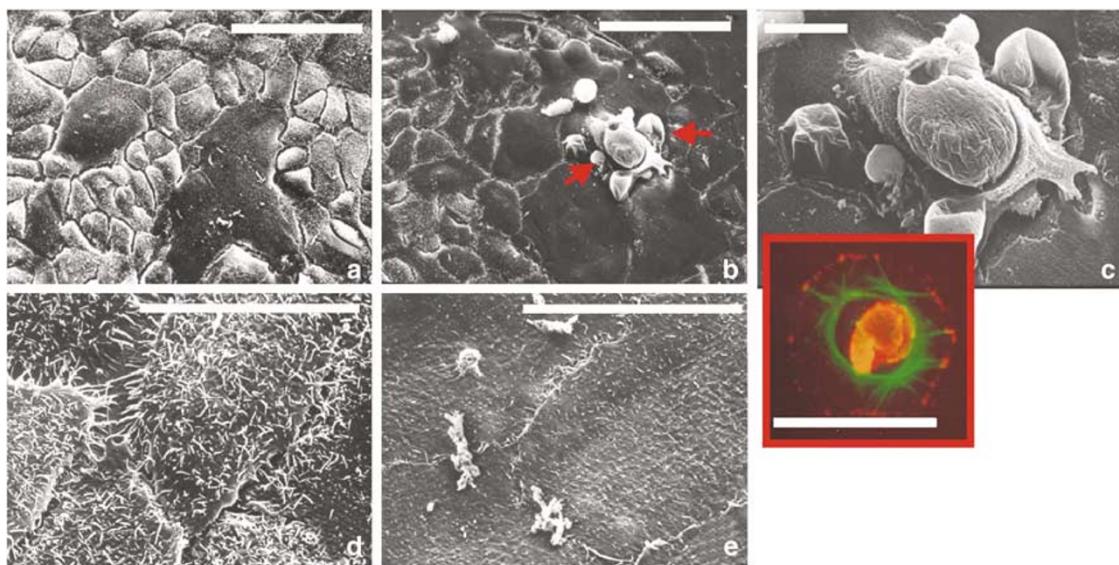


Figure 3 Effect of NAC treatment on NHEK morphology. **(a–e)** Scanning electron micrographs of controls (**a** and **d**) and of NAC-treated cells (**b**, **c** and **e**); arrows in (**b**) point to a cell showing initial stage of the cornified envelope, detailed in **c** at higher magnification. Inset: confocal fluorescence micrograph of an actin–vinculin (green and red, respectively) double-stained cornified cell showing that actin microfilaments participate to the enucleation process. Bars: 100 μ m in (**a** and **b**); 25 μ m in (**c–e**) and inset

intensity of β -catenin and E-cadherin increased of a factor between 7 and 10 in treated *versus* untreated Caco-2 cells. The intracellular localization of β -catenin was also greatly affected by NAC, with redistribution from the nucleus to adherens junctions. Also cell–substratum adhesion plaques increased after NAC treatment, with the formation of several focal adhesions below and around the cell monolayer (Figure 4g and h).

After 3 days from a 5 mM NAC treatment, OVCAR3 cells showed clear morphological changes, from large and flat to smaller and regularly polygonal (Figure 4i–l, upper insets), and with the disappearance of β -catenin labeling from the nucleus while it increased at the level of cell–cell junctions (Figure 4i–l). From the image analysis, in NAC-treated OVCAR3 cells the β -catenin fluorescence intensity at the junctions increased of a factor of 30 with respect to controls (Figure 4i–l).

Also NHEK treated with NAC (2 mM) showed an increased number of intercellular junctions, with β -catenin bound to junctions (Figure 4m and n), and a modified distribution of focal adhesion plaques, as evidenced by vinculin labeling, both below and at the borders of the cell monolayer (Figure 4o and p).

Modification of cell cytoskeleton

NAC treatment induced a reorganization of cell cytoskeleton both in Caco-2 and in NHEK cells, as evidenced by its fragmentation followed by a rearrangement of the relative location of cytokeratin and actin at early (24 h) and later times (72 h), respectively. In Caco-2 cells, 24 h after NAC supplementation, fragmented and curly cytokeratin fibers were observed (Figure 5b). After 3 days, the cytokeratin signal strongly increased (Figure 5c and d) and actin fibers were

located in the upper cell portion, sustaining the brush border microvilli (Figure 5g and h). Untreated control tumor cells showed a random or even opposite location of actin and cytokeratin (Figure 5c). As a consequence and in agreement with increased cell–cell and cell–matrix adhesions shown above, cell motility was dramatically reduced so that the NAC-treated culture was constituted of sparse colonies in monolayers instead of cells covering the whole flask, occasionally with multiple layers, as in controls (Figure 5i–j). Residual proliferating cells were present in our samples, as expected in Caco-2 cells treated with 2 mM NAC (Figure 1a), but we observed that although chromosomes were doubled, for the lack of tubulin polymerization, several cells could not divide and the mitotic spindle could not be formed, so that anomalous prophase features were detected (Figure 5k and l).

In NHEK, at early times after exposure to NAC, actin fibers showed several stress fibers (Figure 6a and b) and cytokeratin was fragmented and curly (Figure 6c and d). This fragmentation was confirmed by electrophoresis of the whole cytoskeleton extract where we observed more bands than those in the cytoskeleton extracted from control cells (Figure 6i). At 3 days after NAC supplementation, images of treated NHEK showed a redistribution of actin and cytokeratin, and an increase in total cytokeratin (Figure 6e–h), while the cytokeratin 14, a marker for the undifferentiated state of keratinocytes, almost completely disappeared in NAC-treated cells (Figure 6j). Concurrently with the formation of the cornified envelope, actin fibers participated in the enucleation process (green labeling in the inset of Figure 3c). Our results extend those of a previous report on the rearrangement of actin cytoskeleton and on the enhancement of adhesion to plastic or vitrogen substrates after NAC supplementation to human epidermoid carcinoma cells A431.¹¹

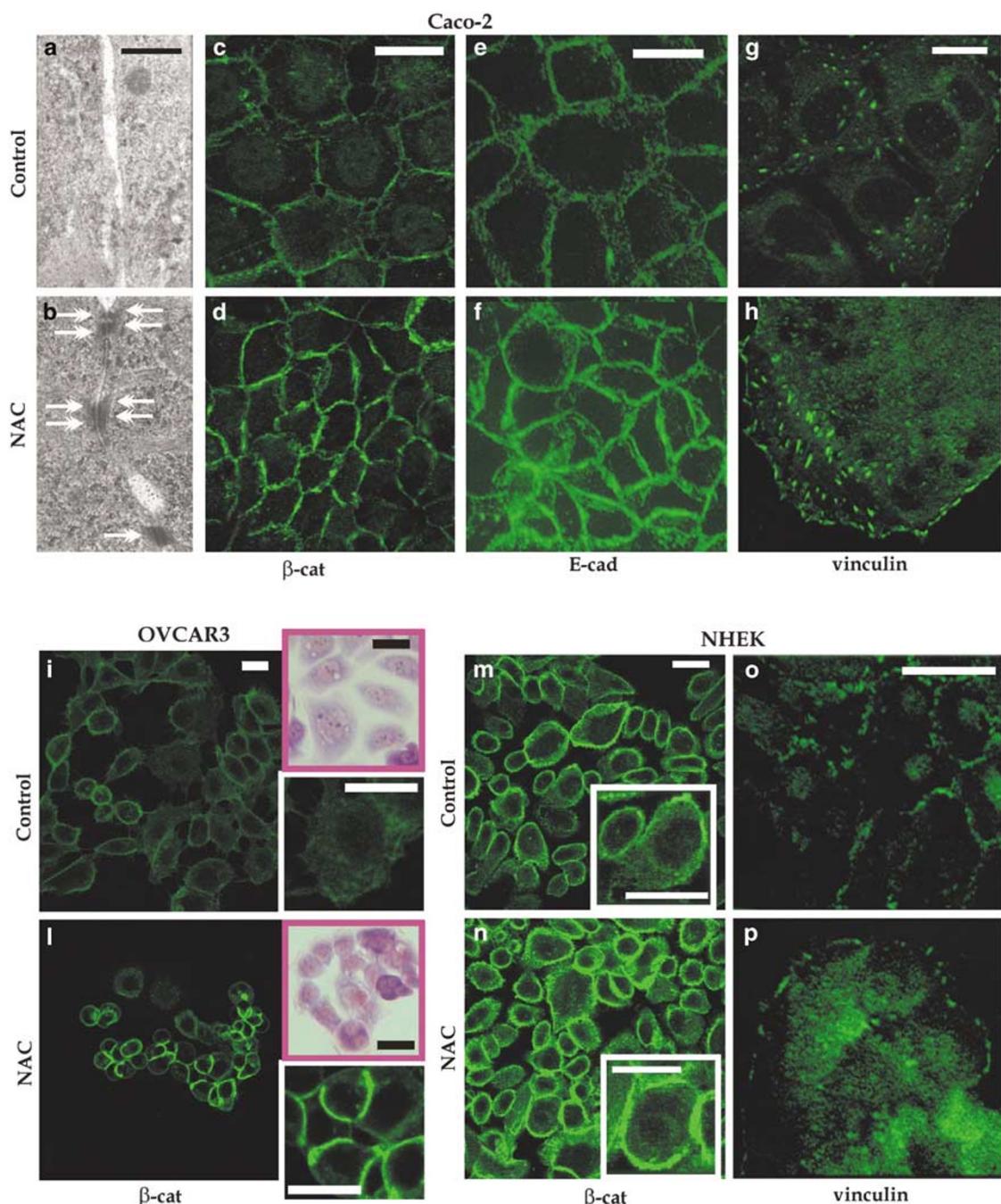


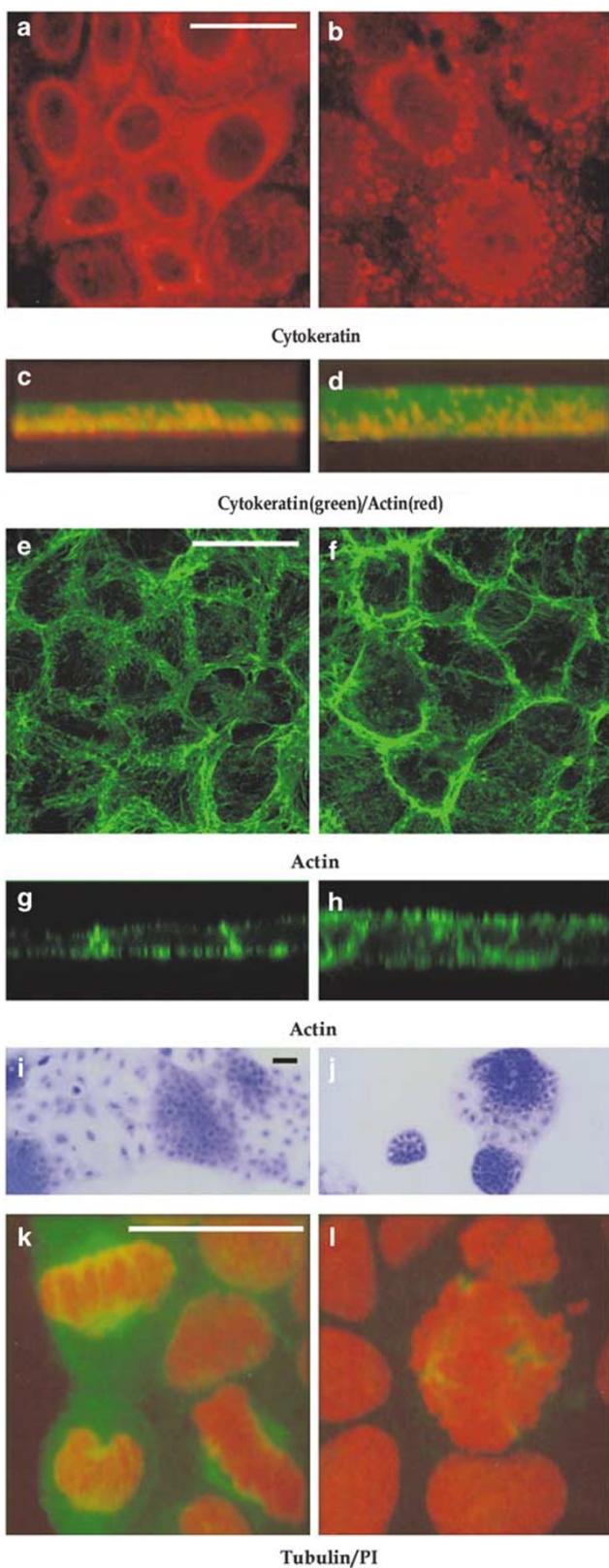
Figure 4 Effect of NAC treatment on cell–cell and cell–substratum adhesions. (a–h): Caco-2 cells; (a and b): transmission electron micrographs of cell–cell contacts in control (a) and in NAC-treated (b) cells. Arrows point to aligned junctions. (c–h): Confocal fluorescence micrographs of β -catenin (c, d), E-cadherin (e, f) and vinculin (g, h) in controls (c, e, g) and in NAC-treated (d, f, h) cells. (i and l): OVCAR3 cells; confocal fluorescence micrographs of β -catenin distribution in controls (i) and in NAC-treated cells (l); in the lower insets, details at higher magnification of β -catenin distribution; in the upper insets, optical microscopic images of cells stained with Wright Giemsa. (m–p): NHEK cells; confocal fluorescence micrographs of β -catenin (m, n) and vinculin (o, p) distribution in controls (m, o) and in NAC-treated (n, p) cells; in the insets, details at higher magnification. Bars: 500 nm in (a and b); 25 μ m in (c–h)

Real-time RT-PCR analysis – NHEK and Caco-2

A global gene expression profile of Caco-2 cells and of NHEK after NAC treatment has been performed and will be published elsewhere (Gustafsson AC, Kupershmidt I, Edlundh-Rose E, Parasassi T, Lundeberg T, Bracci-Laudiero

L, Greco G, Krasnowska EK, Romano MC, Serafino A, Lundeberg J. Global gene expression analysis in time series following N-Acetyl L-Cysteine induced epithelial differentiation of human normal and cancer cells *in vitro*, submitted on line to *BMC Cancer*; Edlundh-Rose E, Kupershmidt I, Gustafsson AC, Parasassi T, Bracci-Laudiero L, Greco G, Krasnowska

EK, Romano MC, Serafino A, Lundeberg T, Nilsson P, Lundeberg J. Gene expression analysis of human epidermal keratinocytes over time following the addition of the anti-



oxidant, *N*-acetyl L-cysteine using spotted cDNA microarrays, submitted to *Pathobiology*). For the present study, some selected genes were also analyzed using real-time RT-PCR and differential expressions are reported relatively to the untreated control samples (Table 1). In Caco-2 cells, the time sequence of differential gene regulation induced by NAC showed an early (12 h) upregulation of the p53, (HSP27) and N-myc downstream-regulated gene 1 (NDRG1) genes, followed by the upregulation of E-cadherin gene at 24 h. The upregulation of p53 following NAC treatment is in agreement with a previous report¹² where microscopic evidence of apoptosis was also reported. Our extensive cytometry investigation excludes the occurrence of apoptosis, reasonably because of the upregulation of the HSP27 gene whose product is reported to prevent apoptosis associated to growth arrest.^{13–15} The upregulation of HSP27 and NDRG1 genes was maintained at 24 h and up to 48 h, respectively, while the p53 gene expression returned to control levels after 24 h. Interestingly, the COX-2 was downregulated at a late time, 48 h. NHEK cells displayed quite a different behavior than Caco-2 cells, with a delayed upregulation of the p53 and E-cadherin genes, at 24 and 48 h, and a constant upregulation of the HSP27 gene at all times. The NDRG1 and COX-2 genes were not affected by NAC treatment.

Inactivation of c-Src

Proteins in cell–cell and in cell–matrix adhesions, as well as proteins in the cytoskeleton, are substrates of the nonreceptor proto-oncogene tyrosine kinase c-Src.^{16–18} In particular, tyrosine phosphorylation of E-cadherin and β -catenin induces their disassembly from adhesion junctions. We therefore investigated whether the observed increase in cell–cell and cell–substratum adhesions was dependent on an inhibition of c-Src kinase after NAC treatment. The modulation of c-Src activity is, in turn, mainly achieved by a selective phosphorylation of two tyrosine residues, with phospho-Y419 corresponding to active c-Src and phospho-Y530 corresponding to inactive c-Src. We examined c-Src phosphorylation at the Y419 and Y530 sites, in Caco-2 and OVCAR3 cells lysates, and the results are shown in Figure 7. In both cell types, after 15 min from NAC treatment we observed a noticeable decrease in the Y419 phosphorylation, while the Y530 phosphorylation was unaffected. Our results well fit with a mechanism of c-Src inactivation induced by sulfhydryl reduction that decreases Y419 phosphorylation without affecting the phosphorylation of Y530.^{4–7}

Figure 5 Effect of NAC treatment on cytoskeleton components of Caco-2 cells. (a–h, k and l): Confocal fluorescence micrographs of controls (a, c, e, g and k) and of NAC-treated cells (b, d, f, h and l). Pan-cytokeratin staining in (a and b); longitudinal sectioning showing the relative redistribution of actin and cytokeratin along the x/z plane: actin (red)/cytokeratin (green) double staining in (c and d). Actin staining in (e–h) with a longitudinal sectioning in (g and h) showing the distribution of actin microfilaments that, in NAC-treated cells, are mainly located in the upper portion of the cell sustaining the brush border microvilli. (k and l): Tubulin staining (green) and counterstaining of nuclei with propidium iodide (red). (i and j): toluidine blue staining of control (i) and of NAC-treated (j) monolayers. Bars: 25 μ m in (a–h, m, n); 100 μ m in (i and j)

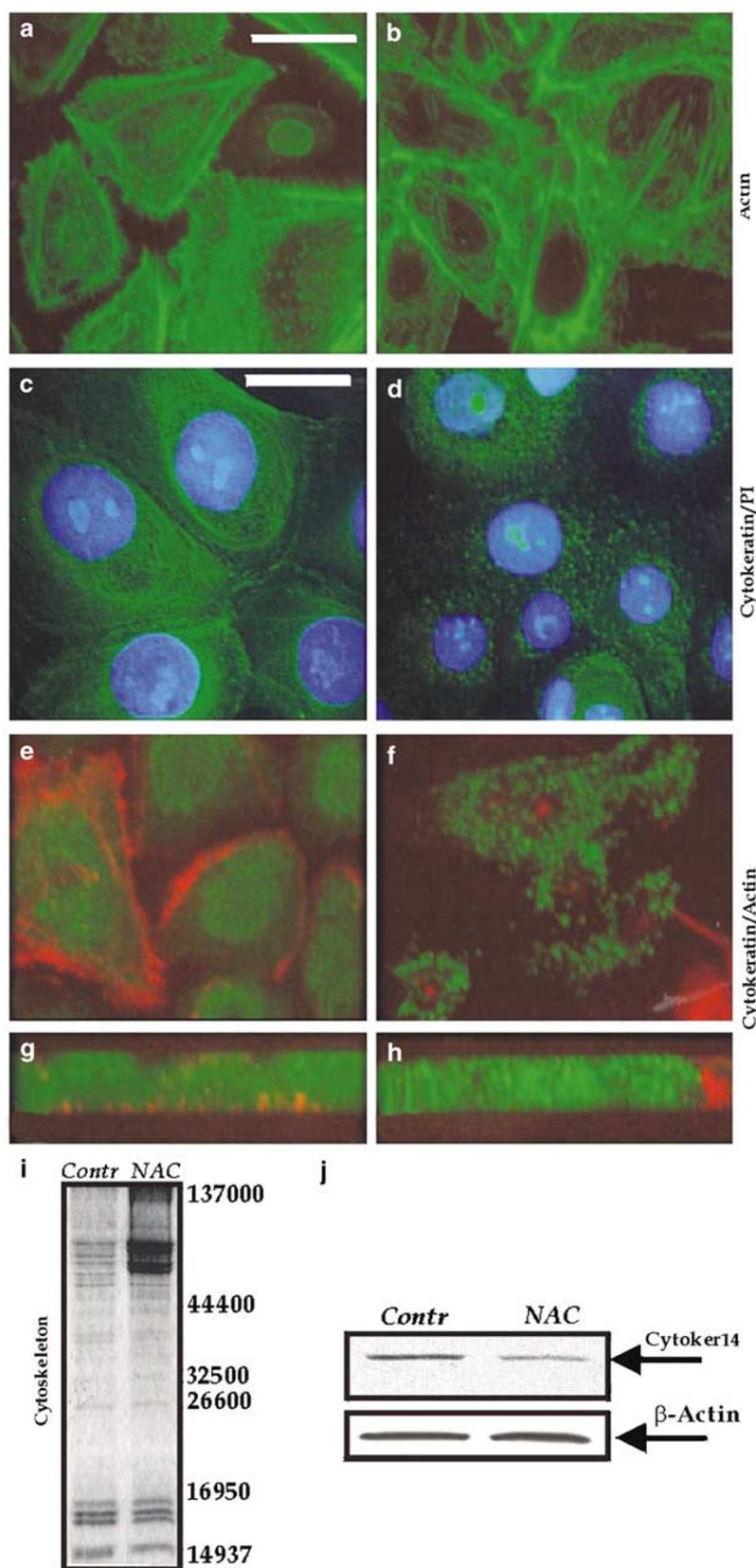


Figure 6

Table 1 Relative gene expression levels measured by real-time RT-PCR in NAC treated as compared to untreated Caco-2 and NHEK cells

Gene	Caco-2			NHEK		
	12 h	24 h	48 h	12 h	24 h	48 h
p53	Up (4.90)	Nc (1.51)	Nc (1.47)	Nc (1.27)	Up (2.57)	Up (2.32)
Hsp27	Up (2.84)	Up (2.19)	Nc (0.77)	Up (2.60)	Up (3.79)	Up (3.58)
NDRG1	Up (5.40)	Up (6.83)	Up (4.56)	Nc (1.35)	Nc (1.25)	Nc (0.96)
E-cadherin	Nc (1.33)	Up (2.67)	Nc (1.08)	Nc (1.57)	Up (5.22)	Up (2.63)
COX-2	Nc (0.91)	Nc (0.59)	Down (0.41)	Nc (0.87)	Nc (0.69)	Nc (0.61)

Significant fold change: $0.5 < 2^{-\Delta\Delta C_T} > 2$ for down- and upregulated genes, respectively. nc, no change. Numbers in parentheses indicate the measured fold change

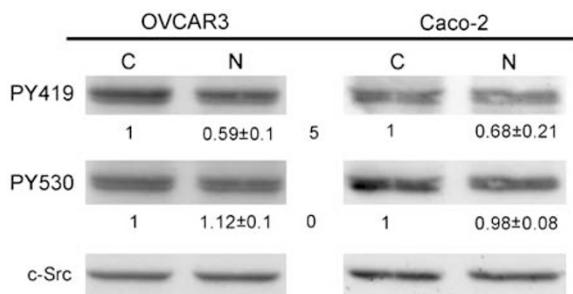


Figure 7 Inactivation of c-Src induced by NAC treatment. Proteins from lysate of control cells (C) or of cells treated with NAC (N), 15 min after the treatment, were probed with antiphosphoY419 or Y530 Abs. For the same representative experiment, we also report the bands of the c-Src protein. After antiphosphoY419 probing, membranes were stripped, reprobed with anti-c-Src, stripped again and reprobed with antiphosphoY530. The numbers below bands represent the average intensities of five independent experiments, with S.D. For each experiment, the antiphospho-tyrosine intensity has been normalized to the intensity of its c-Src protein, then each value of a NAC-treated sample has been normalized to its control. Band intensity was read using the Gel-Doc image analyzer system (BioRad, Hercules CA, USA). OVCAR3 cells were treated once with 20 mM NAC, and Caco-2 cells were treated once with 10 mM NAC

Discussion

The thiol-bearing compound NAC has several experimental and pharmacological uses related to its antioxidant and detoxifying actions.^{19,20} The *in vitro* differentiating action of NAC reported for the first time in the present work is not related to its radical scavenging properties. Indeed, neither antiproliferative effect nor differentiation were induced by the paradigmatic scavenger Trolox C, supplemented to the cell growth medium at concentrations ranging from 0.1 to 0.5 mM. Therefore, NAC action has to be attributed to a modification of protein thiol/disulfide status and to thiol-mediated changes in gene activation.¹ We observed that also L-cysteine can induce an increase in cell–cell junctions, however using a concentration one order of magnitude higher than that of NAC and therefore with a clearly detectable toxic effect on cells (not shown).

The first result on cell proliferation obtained by sulphydryl reduction was a dose-dependent and biphasic effect. Both in normal and cancer cells, a low concentration range, below 0.5 mM NAC, stimulated cell proliferation, suggesting a trophic effect. Instead, the mid to high NAC dose range, from 0.5 to 10 mM, had a dose-dependent antiproliferative effect in the absence of any variation of cell viability. The inhibition of cell proliferation was also cell type dependent, with a relatively low dose, of 2 mM, able to induce a complete block of normal keratinocytes proliferation, while higher doses and, eventually, multiple supplementations were necessary to arrest the colon and the ovary carcinoma cells proliferation. This dose- and cell type-dependent antiproliferative – and differentiating – effect of NAC can explain the unsatisfactory results obtained after a 2-year clinical trial on patients with head, neck, or lung cancer, performed with low dosages, that is, 600 mg/day.²¹ The decreased proliferation of NAC-treated cells is not dependent on apoptosis. In agreement with the generally observed inhibition of apoptosis due to NAC,²⁰ our cytometric analysis of NAC-treated Caco-2 and OVCAR3 cells did not show any evidence of apoptosis. In addition, in our confocal microscopy observations, after propidium iodide staining (Figures 5k, l and 6c, d) we never observed fragmented nuclei. Consistently, our gene expression analysis in colon carcinoma cells and in normal keratinocytes showed an early upregulation of both p53, a proliferation suppressor gene, and of HSP27. The HSP27 gene product is reported to prevent apoptosis associated to growth arrest, possibly by binding to cytochrome c and to procaspase-3.^{13–15} The cytometric analysis, instead, suggested that the antiproliferative effect of NAC has to be related to an increase of the cell population in the G1 phase, finally to a terminal differentiation, as demonstrated by all our other results.

The decrease in proliferation after sulphydryl reduction resulted in the induction of differentiation in colon and ovary carcinoma cells, and in a 10-fold acceleration of normal keratinocytes differentiation. At a morphological level, in NAC-treated cells we showed: (i) a regular polygonal morphology, as opposed to the mesenchymal morphology of proliferating cells; (ii) a *de novo* formation of the brush border microvillar

Figure 6 Effect of NAC treatment on NHEK cytoskeleton components. (a–h) Confocal fluorescence micrographs of controls (a, c, e and g) and of NAC-treated cells (b, d, f and h); (a and b): actin staining; (c and d): Pan-cytokeratin staining (green) and nuclei (blue) counterstained with propidium iodide; (e–h): actin (red)/cytokeratin (green) double staining; (e and f): transversal sectioning at the level of cell–substratum adhesion site; (g and h): longitudinal sectioning showing the relative redistribution of actin and cytokeratin along the x/z plane and the increase in the fluorescence intensity of labeled cytokeratin. Bar: 25 µm. (i) Electrophoresis of the whole cytoskeleton extracts. (j) Western blotting for the determination of cytokeratin 14 content, and for monitoring the amount of loaded protein by β-actin, in control and NAC-treated NHEK

structure in colon carcinoma cells and of the cornified envelope in keratinocytes; (iii) the basal relocation of the nucleus in Caco-2 cells together with the formation of a robust apical vesicular apparatus; (iv) an overall rearrangement of cytoskeleton to fulfill the requirements of newly formed structures; (v) the appearance of several cell-cell and cell-substratum junctions; and, finally, (vi) the disappearance of the oncogenic β -catenin from the nucleus and its only presence at the cell-cell junctions. Taken together, the relocation of β -catenin at the level of junctions and the increased concentration of E-cadherin represent both markers and inducers of differentiation^{22–26} suggesting the establishment of a positive feedback in the differentiation process. The increase in cell-cell and cell-substratum adhesions identifies a more adhesive and less motile phenotype. Also, an increased cell-substratum adhesion has been indicated as a differentiation marker.²⁷

At a molecular level, the upregulation of HSP27 is related to differentiation.^{13–15} Induction of differentiation in Caco-2 cells after NAC treatment was also shown by the early upregulation of the NDRG1 gene, originally identified as a marker of differentiation^{28–31} and usually not found in colon cancer cells.³² In tumors, the NDRG1 gene is downregulated while restoration of its expression in neoplastic cells results in growth inhibition and in morphological differentiation of colon cancer cells, accompanied with the increase of differentiation markers such as alkaline phosphatase and E-cadherin.^{33,34} As expected, the NDRG1 gene expression did not change in NHEK cells after NAC treatment. In agreement with the dramatic increase of cell-cell junctions evidenced by immunofluorescence labeling, an upregulation of E-cadherin gene after NAC treatment was evidenced both in Caco-2 and in NHEK. It is worth noticing that in Caco-2 cells the downregulation of COX-2 gene after sulphydryl reduction well fits the recent efforts in the treatment of colon carcinoma by using nonsteroidal anti-inflammatory drugs (NSAIDs).^{35,36}

The genes we analyzed in this work represent a selection from a more extensive gene expression analysis after NAC treatment in colon carcinoma cells and in normal keratinocytes, performed in a time series by the Affymetrix Gene-Chip™ Human Genome U95Av2 chip, which contains approximately 12 000 previously characterized sequences, and by a spotted cDNA array, comprising approximately 30 000 features. On a whole, the obtained results, to be published elsewhere, showed the downregulation of genes involved in proliferation and the upregulation of genes involved in differentiation-related pathways.

Effects on signaling molecules were also observed after sulphydryl reduction, including the relocation of the oncogenic β -catenin from the nucleus to the cell-cell junctions.^{24,25,37} Among other functions related to the proliferation/differentiation transition, the nonreceptor tyrosine kinase c-Src has a pivotal role in β -catenin relocation.^{38,39} Robust *in vitro* evidences bind the modulation of c-Src kinase activity to the redox state of a group of four cysteine residues close to its catalytic site, a key target for a mutually counteracting reversible control operated by oxidizing and reducing agents.^{4–7} The direct involvement of —SH residues has been demonstrated by the use of a nitric oxide generator, known to bind to sulphydryl residues on proteins, and by reversing this

activation, only *in vitro*, by 2-mercaptoethanol or by dithiothreitol.^{4,5} c-Src was also activated when sulphydryl cysteine residues were specifically oxidized by mercuric ions.⁶ Mercuric ions have the unique property to react with two adjacent sulphydryl groups and its action was specifically inhibited by NAC, as a sulphydryl-donating agent. Which cysteine residues are involved in the redox regulation of c-Src was proven by the use of a series of Src mutants in which cysteines were replaced to alanines.⁷ This modulation of c-Src activity operating through the redox state of —SH groups implies, for inactivation, a dephosphorylation at the auto-catalytic Y419 site, with no effect on the phosphorylation of at the Y530 site. This characteristic phosphorylation pattern is alternative to the well-known regulation operating through phosphatases and kinases acting on the Y530. We hereby demonstrated that the sulphydryl reductant action of NAC induces the inhibition of c-Src in living cells, yielding the same signature pattern than that obtained in immunoprecipitated c-Src treated with mercaptoethanol or dithiothreitol,⁴ that is, a dephosphorylation of Y419 without affecting Y530. It is worth noticing that in comparison with these other sulphydryl reductants, NAC is not toxic. Although also protein tyrosine phosphatases should be activated by sulphydryl reduction,³ the lack of Y530 dephosphorylation excludes a role of phosphatases in this NAC-mediated c-Src inactivation. More experiments are in progress to better elucidate this mechanism for c-Src inactivation and to quantitatively determine the oxidized/reduced —SH groups. Indeed, the NAC-mediated inhibition of c-Src deserves a particular attention when considering that: (i) the inherited or acquired deregulation of this kinase has been implicated in the pathogenesis of cancer; (ii) the inhibition of c-Src has been proposed as a promising strategy in anticancer treatment;³⁹ and (iii) the recent approval of Imatinib mesylate prompted the investigation of new molecules to inhibit others nonreceptor protein kinases through the design of new drugs able to interfere with the binding of ATP or of the substrate to c-Src.^{40,41}

Together with c-Src inactivation, NAC also downregulates the COX-2 gene in Caco-2 cells. The inhibition of cyclooxygenase-2 represents a target for most recent drug design in cancer therapy. The COX-2 gene is upregulated in several human tumor cells, and antitumor properties of NSAIDs have been demonstrated in epidemiological studies and in experimental animal models.^{35,42–44} Among the evolving family of selective COX-2 inhibitors, some are already in clinical trials for the prevention of colorectal carcinoma and promising results have been reported.⁴⁴

In summary, by treating proliferating normal and cancer cells with NAC we obtained a nonspecific sulphydryl reduction that shifted the overall cell activity to a differentiated state, through a series of converging mechanisms. The 10-fold acceleration of differentiation observed in primary normal keratinocytes highlights the intervention of sulphydryl reduction in a physiological switch that can be (partly) conserved after transformation. A further elucidation of involved processes can well be of both physiological and therapeutic interest. Apparently, sulphydryl reduction can achieve some of the goals of the recent differentiation therapy for cancer treatment.^{45–47} An appropriate drug design would be desirable, to select compounds and delivery methods able to

minimize adverse alterations of the thiol status in areas external to the targeted regions.

Materials and Methods

Cell culture and NAC treatment

The Caco-2 human colon carcinoma cells were seeded at a density of 9×10^3 cells/cm², grown in Dulbecco's-modified Eagle minimum essential medium (DMEM, GIBCO Labs, Grand Island, NY, USA), supplemented with 10% (v/v) heat-inactivated fetal calf serum (GIBCO Labs), L-glutamine (2 mM), penicillin (50 IU/ml) and streptomycin (50 µg/ml). OVCAR3 cells at a density of 2×10^4 cells/cm², were grown in RPMI 1640 medium with 10% fetal bovine serum, 10 mM Hepes, 1 mM sodium pyruvate, 4.5 g/l glucose, 0.01 mg/ml insulin, penicillin (50 IU/ml) and streptomycin (50 µg/ml). Normal human epidermal keratinocytes (NHEK, Cambrex Corp., San Diego, CA, USA), plated at a density of 8×10^3 /cm², were grown in KGM medium plus KGM Single Quots (Cambrex Corp.). *N*-acetyl-L-cysteine (NAC, Sigma Chem. Co., St. Louis, MO, USA) stock solution (20 or 100 mM in growth medium), stored at 4°C and in the dark, was used within 1 week from preparation and sterilized by filtration before addition to the cell culture. A proper aliquot of NAC stock solution was added to the cell culture 24 h after seeding. For the proliferation assay (below) the total volume of NAC stock solution never exceeded 20% of the growth medium.

Cell proliferation and viability assay

Overnight incorporation of 0.5 µCi/well [³H]-thymidine (Amersham Biosciences, Freiburg, Germany) by cells plated on 96-wells dishes was determined by liquid scintillation counting in Rach Beta scintillation counter (LKB Wallac 1214 Rack Beta, Bromma, Sweden). Parallel counting of dead cells was performed by Trypan blue (0.2%) staining. Cells were seeded at a density of 8×10^3 /cm² for Caco-2 and NHEK cells and 2×10^4 /cm² for OVCAR3.

Flow cytometry measurements

Caco-2 and OVCAR3 cells were treated once or twice, at 24 h interval, with different NAC concentrations, from 2 to 10 mM, as described above. At 3 days after NAC supplementation, cells were fixed in 70% ethanol and stained with propidium iodide (50 µg/ml; Sigma Chem. Co.) in PBS containing RNAase A (75 kU/ml; Sigma Chem. Co.), and measured by an EPICS 541 (Coulter Hialeah, FL, USA) flow cytometer. DNA histograms were analyzed by a suitable mathematical model¹⁰ to estimate the percentage of cells in the various phases of the duplication cycle.

Real-time RT-PCR

Gene-specific real-time PCR primers for six human transcripts (Transferrin receptor: NM_003234, COX-2: NM_000963, n-myc downstream regulated: NM_006096, p53:NM_000546, E-cadherin:NM_004360, hsp27:NM_001540) were designed using the Primer Express software (Applied Biosystems, Foster City, CA, USA). The analysis was performed in triplicate using ds cDNA (diluted 1:50) synthesized from mRNA obtained from Caco-2 cells or NHEK cells grown for 12, 24 and 48 h, with or without supplementation of NAC, as template. The total RNA was extracted from cell cultures using Trizol (GIBCO Labs) according to the manufacturers instructions. Thereafter, mRNA was extracted by oligo dT Dynabeads (Dynal, Oslo, Norway) and the quality of mRNA was validated using the Bioanalyzer 2100 (Agilent technologies, Waldbrunn, Germany).

The Transferrin receptor gene was used as internal standard. A PCR mastermix was prepared using the SYBR® Green PCR Core Reagents (Applied Biosystems) and aliquoted into microplate wells together with 1 µl template and 5 pmol of each primer for a final volume of 25 µl per reaction. The iCycler iQ™ Multicolor Real-Time PCR Detection System (Bio-Rad Laboratories, Hercules, CA, USA) was used for PCR and for the detection of fluorescent signal. Standard curves (C_T versus log concentration) were generated for each primer pair using duplicate cDNA samples in a series of consecutive five-fold dilutions. Efficiency calculations ($E = 10^{(-1/\text{amplificationslope})} - 1$) were performed to validate compatibility of investigated genes with the internal control. The compatibility of all pair-wise compared amplification efficiencies were confirmed with a maximal deviation of 10% (data not shown). The specificity of all individual amplification reactions was confirmed by meltcurve analysis (data not shown). For data evaluation, the comparative threshold cycle (C_T) method was used, and the level of significance was set to a two-fold relative difference between samples, that is, significant fold change $0.5 < 2^{-\Delta C_T} > 2$ for down- and upregulated genes, respectively.

Scanning electron microscopy

Cells grown on coverslips were fixed with 2.5% glutaraldehyde in 0.1 M Millonig's phosphate buffer (MPB) at 4°C for 1 h. After washing in MPB, cells were postfixed with 1% OsO₄ in the same buffer for 1 h at 4°C and dehydrated in increasing acetone concentrations. The specimens were critical-point dried using liquid CO₂ and sputter-coated with gold before examination on a Stereoscan 240 scanning electron microscope (Cambridge Instr., Cambridge, UK).

Transmission electron microscopy

Cells in flask were fixed for 1 h at 4°C with 2.5% glutaraldehyde in 0.1 M Millonig's phosphate buffer (MPB) containing 2% sucrose, and postfixed for 1 h at 4°C with 1% OsO₄ in the same buffer. Samples were then scrapped out of flasks, dehydrated in increasing ethanol concentrations and embedded in Spurr epoxy resin (Agar Scientific LTD, Stansted, Essex, UK). Ultrathin sections were stained with uranyl acetate and lead citrate and observed by a Philips CM12 transmission electron microscope operating at 80 kV.

Optical microscopy

Treated and untreated OVCAR3 or Caco-2 cells, grown on coverslips, were stained with Wright Giemsa or toluidine blue, respectively, and observed by optical microscopy.

Immunofluorescence labeling and confocal microscopy observations

Cells grown on coverslips were fixed with 4% paraformaldehyde for 10 min and permeabilized with 0.2% Triton-X 100 in PBS for 5 min. Indirect immunofluorescence staining was performed using the following primary antibodies: (i) mAb to human vinculin (Sigma Chem. Co.); (ii) mAb to E-cadherin (Sigma Chem. Co.); (iii) mAb to β-catenin (Transduction Labs, Lexington, KY). Primary antibody detection was obtained by reaction with fluorescein (FITC)-conjugated IgG (Sigma Chemical Co.). Actin microfilaments were stained with FITC-conjugated phalloidin (Sigma Chem. Co.), cytokeratin was stained by mAbs to FITC-labeled Pan-cytokeratin and

microtubules were stained by rabbit polyclonal antibody raised against chicken tubulin plus FITC IgG. Cell nuclei were stained with 2 µg/ml propidium iodide (Sigma Chem. Co.) in the presence of 0.1 mg/ml RNAase (Sigma Chem. Co.). For Golgi apparatus and endoplasmic reticulum staining, cells were incubated with TRITC-conjugated wheat germ agglutinin (WGA-TRITC; Sigma Chem. Co.) for 1 h at room temperature.

Fluorescently labeled samples were imaged by a confocal LEICA TCS 4D microscope (Leica, Heidelberg, Germany) equipped with an argon/krypton laser. Confocal sections were taken at 0.5–1 µm intervals. The excitation and emission wavelengths were 488 and 510 nm, respectively, for FITC-labeling, and 568 and 590 nm, respectively, for propidium iodide and TRITC-labeling. The NIH ImageJ software was used to calculate the integrated fluorescence intensity of interest, after subtracting a constant background.

Whole cytoskeleton extraction

NHEK were washed twice with PBS containing 1 mM Ca²⁺ and Mg²⁺, and incubated for 20 min at 4°C with the extraction buffer (50 mM Tris/HCl, pH 7.5, 40 mM KCl, 5 mM EGTA, 5 mM MgCl₂, 1% Triton X-100, 1 mM PMSF, 40 µg/ml aprotinin, 2 mM benzamidine). Cells were then scraped, centrifuged at 4000 × g at 4°C for 20 min. The pellet was washed twice with the extraction buffer and resuspended in 8 M urea/1% SDS. Samples of 6 µg protein were used for PAGE on 15% slab gel under nonreducing conditions. For immunoblotting, proteins from SDS-PAGE were transferred into nitrocellulose filter (Amersham Biosciences), using 4% milk protein in Tris-buffered saline (TBS: 25 mM Tris, pH 7.5, 0.5 M NaCl), then exposed to the mouse anticytokeratin peptide 14 (Sigma Chem. Co.). After incubation for 1 h with sheep anti-mouse biotinylated IgG (Amersham Biosciences), the blot was transferred to a solution of streptavidin conjugated with horseradish peroxidase (Amersham Biosciences), washed three times with TBS and developed by the Bio-Rad system. The blot was simultaneously incubated with anti-β-actin antibody to monitor the amount of protein.

Immunoblotting

Cells were washed twice with ice-cold PBS containing 1 mM Na₃VO₄ and lysed with RIPA buffer (20 mM Tris-HCl pH 7.4, 1% Triton X-100, 150 mM NaCl, 5 mM MgCl₂, 0.5% sodium deoxycolate, 0.1% SDS, 50 mM NaF and 1% protease-phosphatase inhibitor cocktail (Sigma Chem. Co.)) for 20 min at 4°C. The lysate was centrifuged 10 min at 1000 × g at 4°C and protein concentration was determined (Bradford assay, Bio-Rad Labs.). Proteins were resolved on SDS 10% polyacrylamide and transferred to a nitrocellulose membrane. The membrane was incubated with anti-Src mouse monoclonal antibody (Upstate Biotechnology, Lake Placid, NY, USA) and with anti-Src phosphospecific Y419 and Y530 rabbit polyclonal antibodies (Calbiochem, Cambridge, MA, USA; where human Y419 and Y530 correspond to the Y416 and Y527 chicken c-Src numbering, respectively) for 1 h at room temperature. Proteins were then probed with the appropriate horseradish peroxidase-conjugated antibodies for 1 h at room temperature and visualized by ECL-Western blotting detection reagents (Amersham Biosciences).

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