Induction of the Ras activator Son of Sevenless 1 by environmental pollutants mediates their effects on cellular proliferation

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ARTICLE INFO

ABSTRACT

TCDD (2,3,7,8-tetrachlorodibenzodioxin), a highly persistent environmental pollutant and a human carcinogen, is the ligand with the highest affinity for the Aryl Hydrocarbon Receptor (AhR) that induces via the AhR, xenobiotic metabolizing enzyme genes as well as several other genes. This pollutant elicits a variety of systemic toxic effects, which include cancer promotion and diverse cellular alterations that modify cell cycle progression and cell proliferation. Large-scale studies have shown that the expression of Son of Sevenless 1 (SOS1), the main mediator of Ras activation, is one of the targets of dioxin in human cultured cells. In this study, we investigated the regulation of the previously uncharacterized SOS1 gene promoter by the AhR and its ligands in the human hepatocarcinoma cell line, HepG2. We found that several environmental pollutants (AhR ligands) induce SOS1 gene expression by increasing its transcription. Chromatin immunoprecipitation experiments demonstrated that the AhR binds directly and activates the SOS1 gene promoter. We also showed that dioxin treatment leads to an activated Ras-GTP state, to ERK activation and to accelerated cellular proliferation. All these effects were mediated by SOS1 induction as shown by knock down experiments. Our data indicate that dioxin-induced cellular proliferation is mediated, at least partially, by SOS1 induction. Remarkably, our studies also suggest that SOS1 induction leads to functional effects similar to those elicited by the well-characterized oncogenic Ras mutations.

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1. Introduction

The Ras signaling pathway is involved in cell growth and differentiation. Mutations in the Ras proto-oncogene have been described in a large variety of human malignancies [1]. These mutations result in a constitutively active Ras protein, which is one of the most common Ras dysregulations in cancers [1,2]. Three different Ras genes encoding the H-Ras, K-Ras and N-Ras 21 kDa proteins were discovered 30 years ago [1,3]. The activated Ras proteins physically associate with and activate Raf-1, a serine/threonine kinase which triggers a kinase cascade that results in the activation of the extracellular signal-regulated protein kinases (ERKs of the MAPK family). Ras acts as a functional hub by activating multiple downstream pathways that participate in cell growth and differentiation. The activity of Ras is regulated by two sets of proteins: guanine-nucleotide exchange factors (GEFs) and GTPase-activating proteins (GAPs). GEFs stimulate the release of the Ras bound GDP which then allows GTP to bind and activate Ras. Conversely, GAPs stimulate Ras GTPase activity and lead to the basal GDP-bound state.

Several pathways, which lead to the active Ras state, have been described in mammalian cells. The best-characterized pathway employs the activation of tyrosine kinase receptors to elicit binding of various proteins to the phosphorylated receptor, most notably the growth factor receptor bound 2 (Grb2) protein bound to the guanine nucleotide exchange factor SOS1. The complex is targeted to the plasma membrane, allowing the GEF function of SOS1 to activate Ras by GDP/GTP exchange [4–6]. SOS1 can activate other GTP-binding proteins [4] and other effectors also control Ras activity. The most extensively characterized non-physiological conditions under which increased SOS activitiy has been observed result from gain of function mutations. For example, SOS1

mutations have been described in the Noonan syndrome, a developmental disorder that is characterized by short stature, facial dysmorphia, congenital heart defects and skeletal anomalies [7,8]. These gain of function mutations, which lead to the activation of Ras and were the first examples of activating GEF mutations associated with human disease. Surprisingly, the implications of increased expression of SOS proteins have not been addressed and little is known about the regulation of the gene promoter.

Dioxin is a member of the polychlorinated dibenzo dioxins (PCDD) family which is a class of highly persistent environmental contaminants. 2,3,7,8-tetrachlorodibenzodioxin (TCDD), the most extensively characterized dioxin, has been classified in 1997 as a "human carcinogen" by the International Agency for Research on Cancer (IARC) [9]. This pollutant is the ligand with the greatest affinity for the Aryl Hydrocarbon Receptor (AhR). TCDD via the AhR, induces various toxicities including chloracne, wasting syndrome, teratogenicity, immunotoxicity, neurotoxicity, tumor promotion and carcinogenesis [10]. The AhR is a cytosolic, ligand-activated transcription factor which, upon activation, translocates to the nucleus where it forms a heterodimer complex with AhR nuclear translocator (ARNT). This complex binds to specific xenobiotic responsive elements (XRE) which are characterized by a 5′-GCGTG-3′ consensus core and induces specific target genes which include xenobiotic metabolizing enzymes (XME) such as the cytochrome P450 (CYP) 1 family [11]. In addition to detoxification functions, the activation of this receptor has been shown to elicit diverse cellular effects including cell proliferation [12,13]. These cellular alterations might mediate part of the long-term toxicity of the AhR ligands in animals and humans [14]. Indeed, AhR-null mice display several defects of development and proliferation including thymus and liver hypotrophy [15]. Numerous studies using AhR defective cells or AhR invalidating strategies have clearly established a role for this receptor in cell cycle regulation and proliferation [16]. The dysregulation of these cellular pathways may disrupt normal fetal development and favor cancer progression. The AhR gene targets that are involved in the proliferative effects of this receptor are not completely identified. To this end, we have examined large-scale toxicogenomic analyses of dioxin effects and focused on regulatory proteins that control cellular growth, in particular, the main mediator of Ras activation, SOS1.

In the present work, we investigated, for the first time, the regulation of the previously uncharacterized SOS1 gene promoter by the AhR and its ligands in the human hepatocarcinoma cell line, HepG2. We show that several environmental pollutants (AhR ligands) induce SOS1 gene expression via increased transcription. Moreover, considering the critical role of SOS1 in controlling Ras activity, we also demonstrate that induction of SOS1 by AhR ligands leads to a predominant Ras-GTP state, to MAPK activation and to cell growth. The results point towards SOS1 gene induction as being a critical step for the carcinogenic effects mediated by dioxins.

2. Materials and methods

2.1. Cell culture

Human hepatocarcinoma HepG2 cells were cultured in Dulbecco's minimal essential medium (DMEM, Invitrogen, Cergy-Pontoise, France) supplemented with nonessential amino acids and containing 10% fetal bovine serum, 200 U/mL penicillin, 50 μg/mL streptomycin (Invitrogen) and 0.5 U/mL amphotericin B (Bristol-Myers Squibb Co., Stamford, CT) at 37 °C in a humidified atmosphere of 5% CO_2 in air. The day before with the addition of various concentrations of TCDD, cells were cultured in DMEM

without phenol red and supplemented with 3% charcoal-treated (desteroidized) calf serum. Cells were maintained in this medium during all the treatments except for the proliferation studies. TCDD was purchased from LCG Promochem (Molsheim, France). Benzo(a)pyrene, and quercetin were purchased from Sigma–Aldrich (St Louis, United States).

2.2. RNA extraction, reverse transcription and quantitative RT-PCR

Total RNAs were extracted using the RNeasy mini kit (Qiagen, Les Ulis France) and reverse transcription was performed with each RNA sample using the cDNA high-capacity archive kit (Applied Biosystems, Courtaboeuf, France) as previously described [17]. Specific oligonucleotides were designed using the OLIGO Explorer software (Molecular Biology Insights, Inc., Cascade, CO). Gene specific primers used for the real time PCR were: SOS1: forward 5′-TGAGAGGCAA-CAGAAAGAGC-3′ and reverse 5′-GAGAAGGGAAATGAAATGGG-3′; AhR: forward 5′-ACATCACCTACGCCAGTCGC-3′ and reverse 5′-TCTATGCCGCTTGGAAGGAT-3′; RPL13A: forward 5′-CCTGGAGGA-GAAGAGGAAAGAGA-3′ and reverse 5′-GAGGACCTCTGTGTATTTGT-CAA-3′. Oligonucleotides were obtained from (Qiagen). Quantitative RT-PCR was carried out in a 10 μL reaction volume containing 40 ng of cDNA, 300 nM of each primer and ABsolute™ QPCR SYBR® Green (Abgene, Villebon sur Yvette, France). Quantitative RT-PCR measurements were performed on an ABI Prism 7900 Sequence Detector system (Applied Biosystems). PCR cycles consisted of the following steps: Taq activation (15 min, 95 °C), denaturation (15 s, 95 °C) and annealing and extension (1 min, 60 °C). The threshold cycle (Ct), which is inversely correlated to the amount of target mRNA, was measured as the number of cycles for which the reporter fluorescent emission first exceeds the background. The relative amounts of mRNA were estimated using the ΔΔCt method with RPL13A as reference.

2.3. Immunoblotting

Whole cell lysates were prepared as previously described [17] from HepG2 cells with M-PER®, Mammalian Protein Extraction Reagent, containing protease and phosphatase inhibitor cocktails (Sigma–Aldrich, St Louis, United States). Equal amounts of total protein were separated by SDS-PAGE and transferred onto nitrocellulose membranes (GE Healthcare, Orsay, France). The membranes were probed with primary antibodies recognizing SOS1 (Upstate, Molsheim, France, 07-337, 1:2000), Actin (Abcam, Paris, France ab37063, 1:10000), AhR (Abcam, ab2770, 1:1000), ERK-Phosphorylated (Cell signaling, Saint Quentin en Yvelines, France 9101, 1:2000), ERK (Cell signaling, 4695, 1:2000) and Ras (Upstate 05-516, 1:10000). Immunoreactive bands were detected with X-ray film using alkaline phosphatase-linked secondary antibody (Applied biosystems, T2191 or T2192, 1:20000) and an enhanced chemiluminescence system CDP-Star® (Applied biosystems). Relative Quantification of the amount of immunreactive material was performed with ImageJ freeware (http://rsb.info.nih.gov/ij/).

2.4. SOS1 mRNA half-life measurement

On day 1, HepG2 cells were seeded at 400,000 cells/well (in 6-well plates) in complete DMEM medium (see above). On day 2, cells were washed with PBS and the medium was replaced by red phenol-free DMEM supplemented with 3% desteroidized fetal bovine serum, 200 U/mL penicillin, 50 μg/mL streptomycin and 0.5 mg/mL amphotericin B. Then TCDD dissolved in medium (1 μM). Fifty microlitre was added to the cells medium to give a final concentration of 25 nM. Twenty-four hours later (on Day 3), cells were treated with 100 μM DRB (5,6-dichloro-beta-D ribofur-

anosyl benzimidazole, Sigma–Aldrich) and lysates were extracted at 0, 2, 4, 6, 10 and 12 h after the addition of DRB using the RNeasy mini kit (Qiagen) Quantitative real-time PCR was then performed.

2.5. In silico analysis and primer extension

In silico analysis was performed with MatInspector® software (Genomatix, Munich, Germany) to identify all the potential consensus responsive elements in the promoter. For the primer extension study, total RNAs were extracted using the RNeasy mini kit (Qiagen) as described. Primer extension experiments were performed with the Primer Extension System-AMV Reverse Transcriptase kit (E3030, Promega, Charbonnières les Bains, France). Positive and negative controls are provided with the Kit from Promega. A specific primer was designed (+100 5′-GGGACCCCGCTTCCCGGCCGCAGC-3′ +123) (+ 1 refers to translation start site) and labeled with [γ^{32}P]ATP (Easytides, BLU502A001MC, PerkinElmer, Courtaboeuf, France). The reverse transcription product was separated on a denaturing acrylamide gel. After migration, radioactive bands were visualized with a phosphorimager (STORM 580).

2.6. siRNA transfection

The day before the transfection, HepG2 cells (4×10^5 cells/well) were seeded in 6-well plates. The medium was replaced the day of the transfection by DMEM, without phenol red, supplemented with 3% charcoal-treated (desteroidized) calf serum. Cells were then transfected with 20 nM SOS1 siRNA, using the Lipofectamine 2000 transfection reagent (Invitrogen) according to the manufacturer's protocol and with 5 nM AhR siRNA using the HiPerfect transfection reagent (Qiagen) according to the manufacturer's protocol. For SOS1 and AhR experiments, the medium was replaced after 24 h of incubation at 37 °C and cells were treated or not with 25 nM TCDD. RNA and protein purification were carried out 48 h after the transfection. siRNA references and/or sequences (Qiagen) were

- siSOS1: Hs_SOS1_5_HP Validated siRNA (ref: SI02655121)
- siAhR: Hs_AHR_5_HP Validated siRNA (ref: SI02780148)
- siControl: GFP Custom siRNA (ref: 1027020)

2.7. Reporter assays

pGLuc basic vector (New England Biolabs, Ipswich, MA) was digested with BglII and HindIII restriction enzymes (New England Biolabs). For the construction of the SOS1 promoter (AF 106953) the −2200, +37 region;(+ 1 refers to the translation start site) was amplified using the following primers: −2200 5′-CTTAGATC-TAAGTGGGGAGTTTTTCAAATGT-3′ −2170 and +37 5′-AAGCTTTCG-CTGAAAAACTCGTAGGG-3′ +17. The PCR product was cut with the same restriction enzymes. Transfection experiments were performed as described in siRNA transfection paragraph, using 1 μg of plasmid. The empty pGLuc basic vector (New England biolabs) was used as transfection control. Twenty-four hours after the transfection, 50 μL supernatant (containing the secreted luciferase) were collected and the medium was changed. The cells were then treated or not with 25 nM TCDD as described above. The supernatant was then harvested at 0, 6, 24 and 48 h after TCDD treatment. A luciferase assay (Gaussia) was performed with a New England Biolabs kit (E3300L) and a luminometer (Lumat LB9507, Berthold, Thoiry, France).

2.8. Chromatin immunoprecipitation

HepG2 cells were seeded into 150-mm-diameter dishes (approximately 20×10^6 cells per dish) 48 h before being pro-

cessed for ChIP as previously described [17]. Treatment with TCDD (25 nM) was performed, as described above, for 15- and for 45 min. We used a ChIP-grade antibody from BioMol (anti-AhR antibody, Cat SA-210, BioMol, Plymouth Meeting, Pennsylvania, USA) ChIP DNA was analysed by quantitative PCR using the specific primers covering the region between −984 and −801 bp of the human Sos1 promoter region (forward primer: 5′-GGTTTCGTGACAGAGCACCT-3′ and reverse primer 5′-TAAAGAGGCTCCCCATTGTG 3′) and non-specific primers (forward primer: 5′-TGGGGTAGACTGTGGCAGA-3′ and reverse primer 5′-GGCTAAGGATTCAGCAGCAA-3′).

2.9. Ras activity

On day 1, cells were seeded at approximately 15 million cells per 175-cm² plate. Twenty-four hours later (day 2), the medium was changed and siRNA SOS1 transfection was performed (see above). On Day 3, the medium was changed and TCDD (25 nM) was added as previously described. On day 4, the medium was replaced with a medium without serum containing or not TCCD (25 nM). On day 5, cells were lysed in Magnesium Lysis Buffer (MLB, Upstate) containing protease and phosphatase inhibitor cocktails (Sigma). Equal amounts of total protein extracts, treated or not, were incubated with Raf-RBD GST beads (Cytoskeleton, Le Perray en Yvelines, France RF-02A), to retain only the active form of Ras, (Ras-GTP) following the manufacturer's recommendations (Upsate). The recovered proteins were separated by SDS-PAGE (see above) and western blots (Ras antibody) were performed. Moreover, we used GTP gamma-S as a positive control and GDP as a negative control (Upstate).

2.10. Cell growth

On day 1, cells were seeded at 30,000 cells/well (in 6-well plates) in complete DMEM medium (see above). On day 4, siRNA SOS1 and siRNA AhR transfection was performed (see above). On day 5, cells were treated with TCDD (25 nM) as above. For each condition, the number of cells per well was counted at times corresponding to 0, 24 and 48 h of TCDD treatment. For cell counting, cells in each well were trypsinised with 150 μL of Trypsine-Versene (Invitrogen), which was neutralized, with 150 μL of complete medium.

2.11. Statistical analysis

The data result from at least three independent experiments. The results were expressed as the mean ± standard error. Differences between groups were analyzed by U Mann–Whitney's test (nonparametric comparison of 2 independent series) or H Kruskal–Wallis's test (nonparametric comparison of k independent series) followed by a 1-factor ANOVA test (parametric comparison of k independent series). A p-value < 0.05 was considered as statistically significant.

3. Results

TCDD is known to promote cellular proliferation and cell cycle progression in different cellular models. Since several toxicogenomic large-scale studies, including our own, were performed in the human hepatocarcinoma HepG2 cell line and since this cell line exhibits high levels of AhR expression, we used the HepG2 cells for our gene regulation studies. As will be shown later in this article, TCDD elicited a 20% increase in the number of cells during the exponential phase of growth. Thus, the HepG2 cells constitute an appropriate model to investigate the molecular mechanisms of TCDD-elicited cellular proliferation.
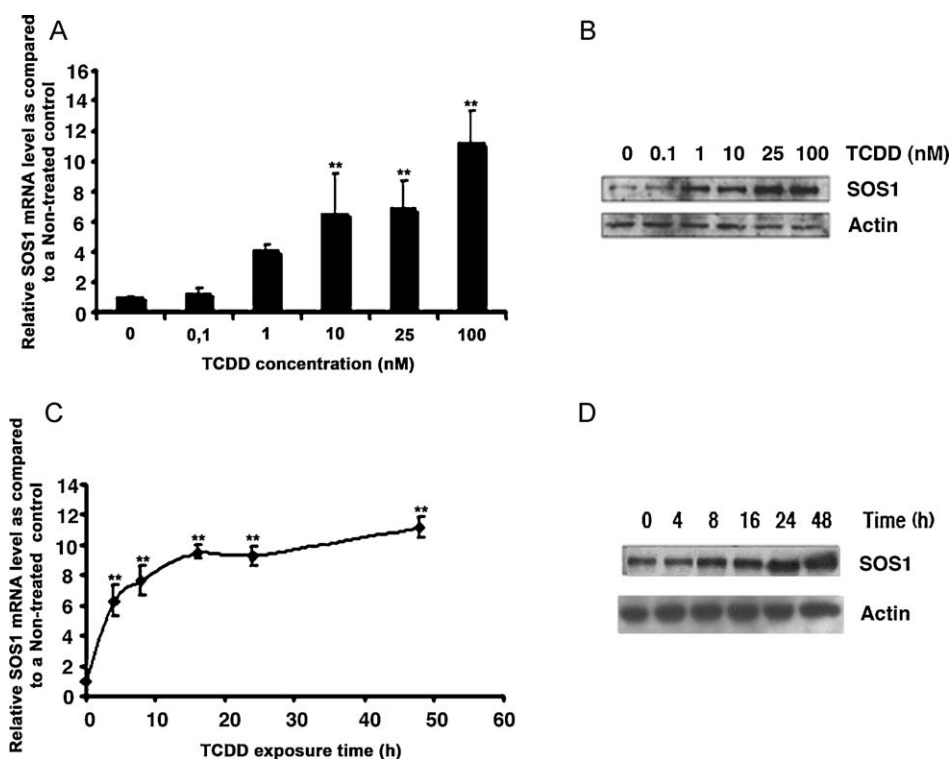
Fig. 1. SOS1 is a target of TCDD in human hepatocarcinoma HepG2 cells. (A) Quantitative real-time PCR (A, C) and western blot analysis (B, D) of SOS1 mRNA and protein levels as a function of TCDD concentration (0–100 nM, 48 h) (A and B) and as a function of time (0–48 h, 25 nM TCDD) (C and D) in human hepatocarcinoma HepG2 cells. Relative mRNA levels were calculated using controls values (0 nM for A or 0 h for C) as references. Actin was used as a loading control for western blots. **, $p < 0.01$ as compared to non-treated controls ($n > 3$).

3.1. Effect of dioxin treatment on SOS1 mRNA and protein levels in HepG2 cells

In a large-scale gene expression study, SOS1 mRNA was found to be induced by TCDD [18]. First, we validated those observations in HepG2 cells. We performed kinetic and dose–response experiments and we extracted both total RNA and total protein. SOS1 expression was assessed using both quantitative real-time PCR (Fig. 1A and C) and immunoblotting (western blots) (Fig. 1B and D). Dose–response experiments showed that TCDD up-regulated SOS1 mRNA expression (Fig. 1A) and protein level (Fig. 1B and Supplementary Fig. 1A). The response was significant for concentrations higher than 10 nM TCDD. Consequently, 25 nM TCDD was used in the following experiments. This concentration of TCDD is within the usual concentration range used to treat human cells (5–100 nM). Fig. 1C and D and Supplementary Fig. 1B show that SOS1 was significantly up-regulated by 25 nM TCDD in a time-dependent manner. In addition to HepG2 cells, we have also observed a significant up-regulation of SOS1 mRNA in several mouse tissues (C57/Bl6) following TCDD treatment (Supplementary Fig. 2). These results indicate that SOS1 is up-regulated upon TCDD treatment and suggest that the SOS1 gene might be an AhR transcriptional target.

3.2. SOS1 mRNA and protein induction is AhR-dependent

We examined the contribution of the Ah Receptor using different AhR ligands including TCDD, Benzo(a)pyrene and quercetin (Fig. 2A) and a RNA interference knock down strategy (Fig. 2B–D). Fig. 2A shows that the AhR ligands, 25 nM TCDD and 5 μM BaP, significantly increased SOS1 mRNA; although an increase in SOS1 mRNA was also observed after treatment of cells with 25 μM Quercetin, this increase was not significant. We then knocked down AhR expression using specific siRNA (Fig. 2B),

which decreased AhR mRNA by approximately 60–70% (Fig. 2B). AhR levels remain knock-downed throughout the time course of the experiment (Supplementary Fig. 3). Fig. 2C and D shows that AhR siRNA treatment elicited a statistically significant 50% decrease in both SOS1 mRNA and protein induction by TCDD. In this experiment, we did not expect to completely knock-down AhR levels but we aimed to correlate the relationship between both AhR and SOS1 levels; we observed such a correlation (AhR by the time of treatment, 0 h, 50% decreased levels and SOS1 at the end of the experiment, 50% decrease) and also a 60% decrease in CYP1A1 induction which is classically measured to prove AhR activation (data not shown). These experiments demonstrate that the activated AhR regulates SOS1 mRNA and protein levels.

3.3. The SOS1 gene promoter is a direct target of the AhR

To further characterize the mechanism of SOS1 induction by TCDD, we blocked RNA polymerase II with DRB and measured SOS1 mRNA levels upon TCDD treatment, as well as under untreated conditions. Fig. 3 shows that the rate of SOS1 mRNA decline was not significantly different in treated and untreated conditions suggesting that a decreased rate of SOS1 mRNA degradation is not involved in the TCDD up-regulation of SOS1 mRNA (Fig. 3A). Very recently, the exon-intron structure of the SOS1 gene was displayed in the Pubmed Ace View Database. Exon 1 is shown to start 280 bases upstream of the translation start site for the longest variant. Other mRNA variants differing in the length of the 5′ ends have been found in different tissues but little information was given on their abundance. To further define the length of the 5′ UTR in our cell line model, we performed a primer extension study, which confirmed the presence of a major start site approximately 150 bp upstream from the translation start site, which is in line with the displayed gene structure (data not shown). A minor transcription start site was also observed downstream from the major one,
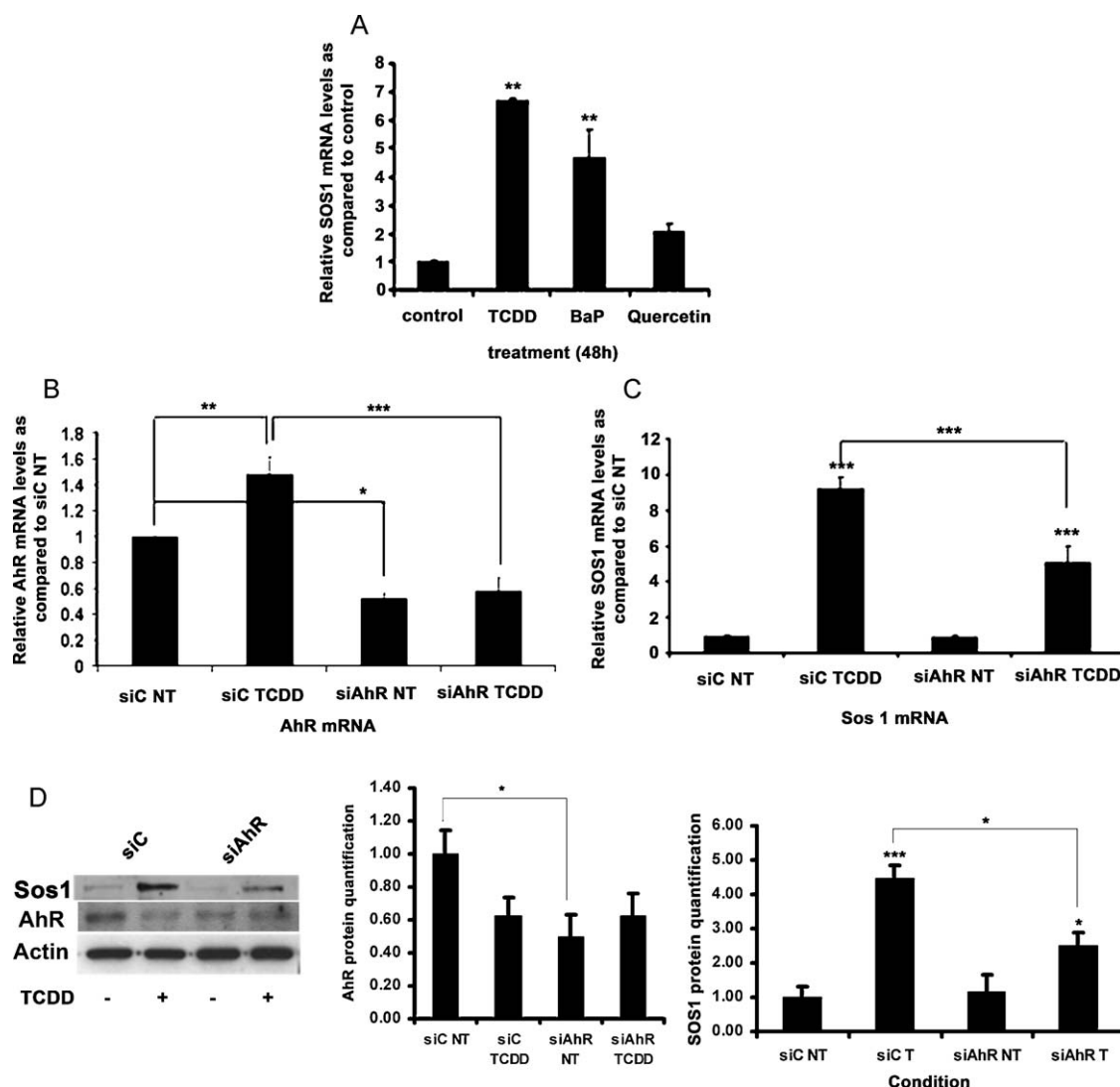
Fig. 2. Involvement of the AhR in the regulation of SOS1 mRNA and protein expression in HepG2 cells. (A) SOS1 mRNA levels measured by quantitative real-time PCR after 48 h-treatment of HepG2 cells with various AhR ligands (TCDD, 25 nM; Benzo(a)pyrene or BaP, 5 μM; Quercetin, 25 μM). Relative mRNA levels were calculated using control value as reference (B) AhR mRNA levels measured by quantitative real-time PCR in AhR (siAhR) or control (siC) siRNA-transfected HepG2 cells (NT: non-treated; TCDD: TCDD-treated). (C) SOS1 mRNA levels measured by quantitative real-time PCR in AhR (siAhR) or control (siC) siRNA-transfected HepG2 cells (NT: non-treated; TCDD: TCDD-treated). (A, B, C: ***, $p < 0.001$; **, $p < 0.01$ and *, $p < 0.05$ compared to controls ($n > 3$)). (D) Western blot analysis and relative quantification of SOS1 and AhR protein levels in AhR (siAhR) or Control (siC) siRNA-transfected HepG2 cells upon TCDD (+) treatment. Actin was used as a loading control for western blots ($n = 3$).

which is not surprising in this GC-rich promoter. Then we performed a MatInspector® analysis of the 2200 base pairs located upstream of the transcription start sites to identify consensus AhR sequences (Xenobiotic Responsive Elements or XRE); Fig. 3B shows the positions and the sequences of 2 putative XREs (AhR binding).

Several experiments were carried out in order to determine whether the SOS1 gene promoter is a direct target of the AhR, including luciferase reporter gene assays (Fig. 4A and B) and Chromatin immunoprecipitation (ChIP) assays using a specific AhR antibody (Fig. 4C). We subcloned the (−2200/+37) fragment of the SOS1 gene promoter (which includes the two putative XREs) upstream of the Gaussia luciferase reporter gene. The activity of this gene product can be readily monitored in the medium. Following transfection with this recombinant reporter gene plasmid, HepG2 cells were treated with TCDD and the medium was recovered at different time points for luciferase assay (Fig. 4A). A significant increase in luciferase activity can be observed 24 and 48 h following TCDD treatment. These experiments confirm that the (−2200/+37) SOS1 gene fragment exhibits promoter activity as compared to the empty vector (Fig. 4B), and

suggest that this gene promoter fragment mediates part of the inducing effect of TCDD. Similar experiments were performed in control siRNA and AhR-siRNA treated cells (Fig. 4B). We observed a similar 2- to 3-fold increase in the activity of the SOS1 gene promoter in siControl-transfected cells upon TCDD treatment. However, basal and induced SOS1 promoter activities were significantly reduced in AhR-depleted cells suggesting a contribution of this receptor to SOS1 gene promoter activation.

Finally, we performed a Chromatin immunoprecipitation experiment using a SOS1 promoter fragment (−984/−801) to confirm the direct binding of AhR to the endogenous SOS1 gene promoter (Fig. 4C). AhR antibody and real-time PCR were used to provide a quantitative measurement of receptor binding. Cells were treated for 15 and 45 min with TCDD. Several controls were used for the experiments (inputs, mock antibody). We observed a time-dependent, specific and potent increase of AhR binding to the SOS1 gene promoter. Altogether, these experiments clearly show that the TCDD-mediated transcriptional regulation of the SOS1 gene is related to the rapid direct binding of the AhR to the promoter.
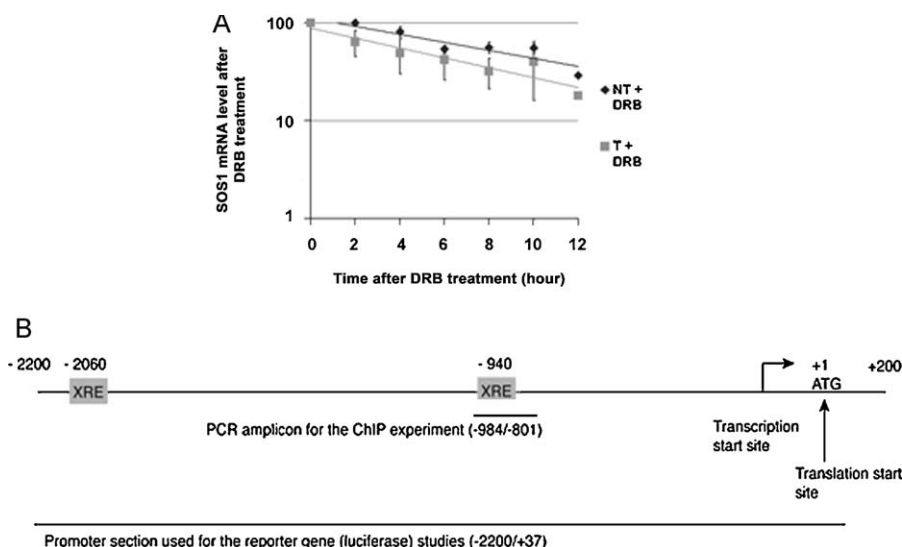
Fig. 3. The AhR dependent-SOS1 regulation is transcriptional. (A) SOS1 mRNA levels measured by relative quantitative real-time PCR in the presence of DRB, an inhibitor of RNA polymerase II in non-treated (NT + DRB) and TCDD-treated (T + DRB) HepG2 cells. Cells were treated or not with 25 nM TCDD for 16 h then with DRB and subsequently harvested (between 0 and 12 h). We use a semi-logarithmic representation. (B) Diagram of the SOS1 promoter ($-2200/+37$) using the translation start site as reference (+1). Localizations of both XRE consensus sites (-940 and -2060 bp) and the promoter section used for the reported gene (luciferase) experiments (see below).

3.4. TCDD stimulates Ras activity and ERK1/2 phosphorylation through induction of SOS1

Since the Guanine nucleotide Exchange Factor SOS1 regulates the Ras-MAPK pathway, we hypothesized that increased amounts of SOS1 could lead to the activation of this pathway. Thus, we monitored Ras activity (the active GTP-bound form) and ERK phosphorylation following TCDD treatment of HepG2 cells. To monitor Ras activity, we performed affinity pull-down experiments. Beads grafted with the domain of Raf, which specifically interacts, with the GTP-bound form of Ras were mixed with different cellular extracts. The bound fraction was then eluted and analyzed with a Ras antibody. Fig. 5A shows that the active form of Ras was increased following TCDD treatment. In parallel, the phosphorylated fraction of ERK was also increased by TCDD treatment (Fig. 5B). To assess the implication of SOS1 in these processes, we down-regulated SOS1 expression using a specific siRNA and observed a clear decrease of TCDD-induced Ras-GTP levels as shown in Fig. 5C. SOS1 protein levels remain knocked down throughout the time course of the experiment. These results show that SOS1 is a critical mediator of TCDD-dependent RAS-MAPK pathway activation. Importantly, for the first time, these observations show that increased amounts of SOS1 are sufficient to activate the downstream pathway indicating that, in HepG2 cells, the amount of SOS1 is limiting. It is noteworthy, however, that the Ras pathway activation by TCDD is delayed as compared to the time course of the response which can be traditionally achieved by Tyrosine Kinase receptor activation [19].

3.5. SOS1 and AhR are critical for dioxin-dependent changes in growth

Since the Ras-ERK pathway in implicated in cell growth, we tested the effect of TCDD treatment on HepG2 cells proliferation. As shown in Fig. 6A and B, TCDD elicited a statistically significant 20% increase in cellular growth in the absence or in the presence of a control siRNA. Following the knock down of SOS1 gene expression (Fig. 6A) using two specific siRNAs, the effect of TCDD was completely abolished. A partial decrease in basal cellular proliferation was also noted, which is not surprising given the implication of the SOS1-Ras-ERK pathway in cellular proliferation.

In addition, AhR-specific siRNAs also prevented the TCDD effects (Fig. 6B). We conclude that SOS1 induction plays a critical role in the TCDD-elicited increase in cellular proliferation.

4. Discussion

AhR activation by polyaromatic and polyhalogenated hydrocarbons has traditionally been associated with induced expression of phase I and phase II xenobiotic metabolizing enzymes, which promote the clearance of pollutants and drugs from metazoan organisms [20]. However, recent toxicogenomic studies, as well as knock-out studies in mice, have unveiled new AhR target genes [13,21–23] and suggested novel functions for this receptor including the regulation of cell proliferation and migration. Several large-scale studies in HepG2 cells [21,22,24] indicated that the SOS1 gene could be a target of TCDD. SOS1, a Guanine Exchange Factor, is essential for the activation of the Ras and Rac signaling pathways, which lead to the regulation of some of the most important cellular functions including cell division, differentiation and migration [4]. Despite these critical roles, neither the regulation of SOS1 expression nor the characterization of its gene promoter has been carried out previously. We recently observed that AhR ligands increase HepG2 cell proliferation and hypothesized that increased SOS1 expression might be involved in this biological effect. We first validated our transcriptomic experiments and showed that the AhR and its ligands positively regulate SOS1 gene expression. Then, we characterized and subcloned, for the first time to our knowledge, the human SOS1 gene promoter. We showed that TCDD and other AhR ligands regulate, via transcriptional mechanisms their activities upon AhR activation followed by direct promoter binding of AhR to XRE sites. Interestingly, the level of induction of SOS1 elicited by each ligand is different. This is in line with one of our former studies showing that the transcriptional response of the AhR depends on the nature of its ligand [25] and, more generally, with the SAhRM concept [26]. Finally, we focused on the implications of the regulation of SOS1 expression by environmental pollutants. Using a siRNA strategy, we demonstrated that increased SOS1 expression is directly linked to Ras and ERK activation, which promotes cell division. This study demonstrates that the activation of the Ras-
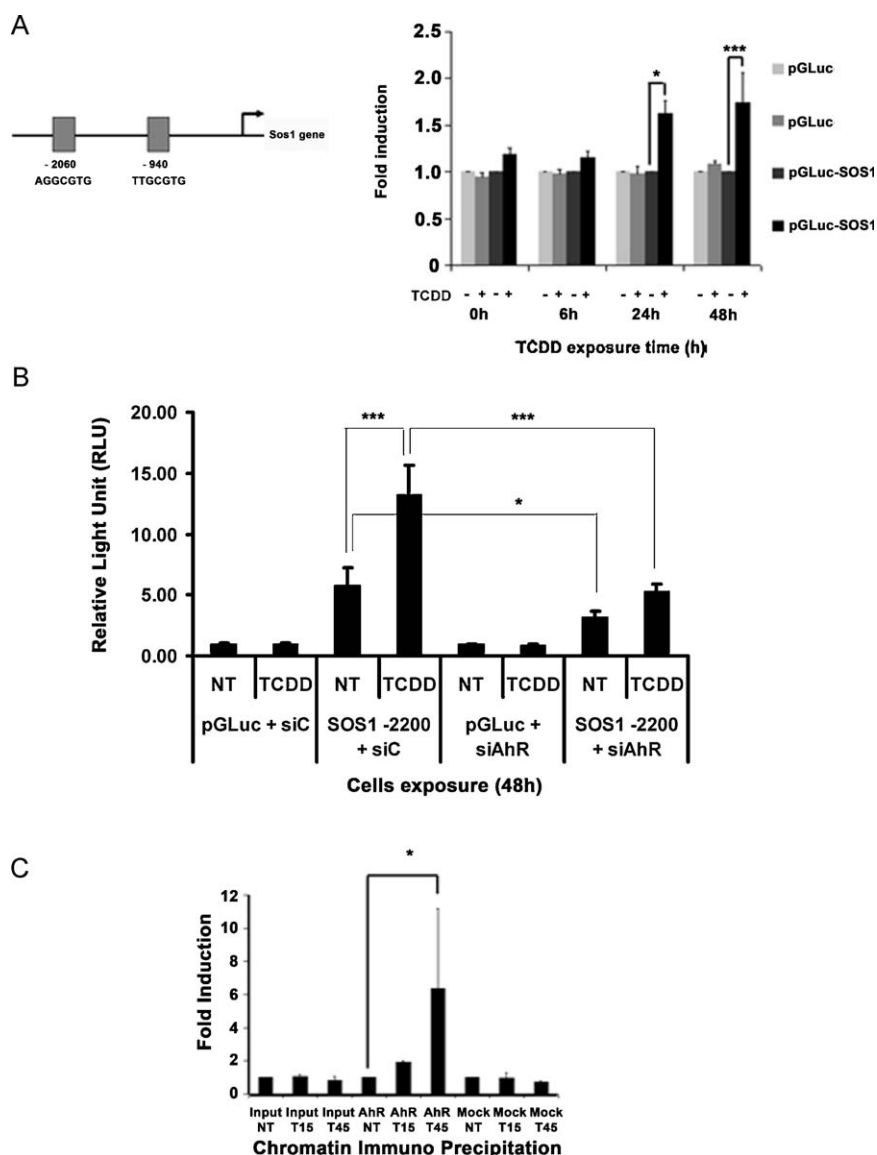
Fig. 4. SOS1 regulation depends on the binding of the AhR to SOS1 promoter. (A) Diagram of the SOS1 promoter subcloned in the Gaussia luciferase basic vector (with both XRE consensus sites at −940 and −2060 bp) and activity of the SOS1 promoter measured after transfection of HepG2 cells with a vector (pGLuc-SOS1) containing the −2200/+37 theoretical promoter region of SOS1 upstream from the Gaussia luciferase reporter gene (+1 refers to the translation start site). The cells were treated (+) or not (−) with TCDD. The medium was sampled after 0, 6, 24 and 48 h. An empty vector (pGLuc) was used as a control. Relative luciferase activities are reported on the y-axis and are calculated using non-treated conditions (−) as control reference for both vectors (B) Activity of the SOS1 promoter measured after transfection of HepG2 cells with a control (siC) or AhR targeting (siAhR) siRNA and the vector containing the −2200/+37 theoretical promoter region of SOS1 upstream from the Gaussia luciferase reporter gene (pGLuc-SOS1). The cells were treated (+) or not (−) with TCDD for 48 h. The medium was sampled after 48 h. An empty vector (pGLuc) was used as a control. Relative luciferase units (RLU) are reported on the y-axis (C) Chromatin immunoprecipitation performed with a mock or a specific AhR antibody (15 or 45 min of TCDD treatment of HepG2 cells, respectively T15 or T45). Specific primers surrounding the proximal XRE (−940, see Figs. 3 and 4A) of the Sos1 gene promoter were used for the study. Input refers to non-precipitated samples and is used to ensure proper "loading" controls before precipitation. The mock antibody is used to measure unspecific precipitations. Fold induction refers to relative levels calculated using non-treated (NT) conditions as reference. ***, $p < 0.001$ and *, $p < 0.05$ as compared to controls ($n = 3$).

ERK pathway could be achieved by increasing the amount of the SOS1 protein. These results suggest that, under our cell culture conditions, the amount of SOS1 protein is limiting.

The activation of the Ras ERK pathway has been studied extensively in the past and several mechanisms have been reported and characterized [19]. Receptor activation followed by SOS1 recruitment is a classical mechanism under physiological and certain pathological conditions and is one paradigm for cellular signaling [6]. In cancer cells, considerable research has been devoted to the upregulation of this pathway through activating Ras mutations. Similarly, the function of SOS1 has been considerably investigated [27–29]. Recent studies also have correlated SOS1 activating mutations to several diseases of development including

Noonan syndrome, a developmental disorder characterized by short stature, facial dysmorphia, congenital heart defects and skeletal abnormalities [7,8,30] as well as hereditary gingival fibromatosis type 1, a benign overgrowth condition of the gingiva [31]. Despite considerable attention to these gains of function mutations, SOS1 gene regulation has remained poorly characterized [32,33]. We speculate that the findings reported here on the upregulation of SOS1 gene expression by environmental pollutants might constitute one mechanism, which accounts for TCDD and other AhR ligand-mediated carcinogenesis. We expect this effect to be particularly relevant in cancerous cells, which do not exhibit Ras mutations. Indeed, continuous upregulation of SOS1 in these cells may mimic Ras mutational activation and lead to the permanent
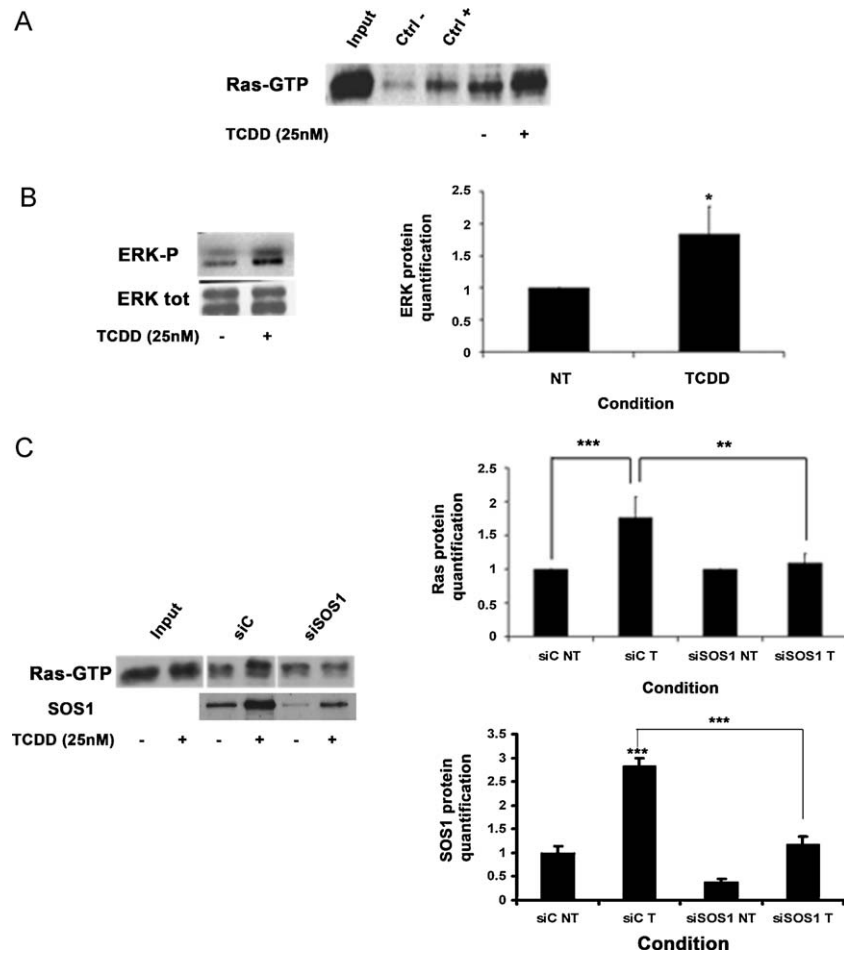
Fig. 5. Increased expression of SOS1 after TCDD treatment stimulates Ras activity and ERK1/2 phosphorylation (A) Western blot analysis of Ras-GTP protein levels after a Ras-GTP specific pull down assay in untreated (−) or 25 nM TCDD-treated (+) HepG2 cells upon 48 h treatment. Input Ras levels show the levels of Ras before precipitation. Precipitation is undertaken using a protein, which specifically interacts with Ras-GTP (Raf), linked to GST-beads. Negative (Ctrl −) and positive (Ctrl +) control respectively refers to extracts over-treated with GDP or GTP-ϒS before precipitation (see Material and methods for more details) (B) Western blot analysis of phosphorylated ERK levels (ERK-P) in untreated (NT) or TCDD-treated (T) HepG2 cells. Total ERK was used as a loading control. The quantification of three independent experiments is presented in the right panel. Fold induction refers to relative levels calculated using the NT condition as reference. *, $p < 0.05$ compared to controls ($n = 3$). (C) Western blot analysis of Ras-GTP and SOS1 protein levels after Ras-GTP specific pull down assays of untreated (−, NT) or 25 nM TCDD-treated (+, T) HepG2 cells extracts after SOS1 (siSOS1) or control siRNA (siC) transfection. The quantification of three independent experiments is presented in the right panel for both SOS1 and Ras-GTP proteins. Fold induction refers to relative levels calculated using NT conditions as reference. ***, $p < 0.001$ and **, $p < 0.01$ compared to controls ($n = 3$).

stimulation of the downstream signaling pathway, which controls cellular proliferation. Thus, our data imply that increased SOS1 expression may constitute an alternative to the activating mutations of Ras and that it should be relevant to explore and characterize alterations of the expression of this gene in cancer cells [34–36]. Furthermore, it is unclear at this stage whether these two mechanisms can be additive or not. An important question, in this respect, is whether gene induction-mediated chronic activation of the SOS1/Ras pathway can be achieved readily in humans. Because of the high contamination levels and widespread distribution of AhR ligands in our environment, chronic exposure to such pollutants is very common, especially in industrialized countries. In addition, long-term effects could also be triggered by acute high-level exposure to these toxins. Indeed, pollutants such as dioxins tend to resist metabolism and accumulate in human and animal adipose tissue, leading to a permanent internal exposure [37].

Genetic and developmental studies in flies and mice have suggested different and unexpected connections between the AhR and the small G proteins network. A recent study by Carvajal-Gonzalez et al. using AhR KO mice models, has unveiled a link between the receptor and the Vav3, a guanosine diphosphate/guanosine triphosphate exchange factor for Rho/Rac GTPases [38].

Moreover, Son of Sevenless (SOS) has been shown to be a major player in the development of Drosophila eye ommatidia. SOS mutations change ommatidia organization and the color perception of the fly [39,40]. Spineless, the Drosophila AhR ortholog, is involved in the ommatidal mosaic formation as well as SOS1 [41]. Thus, these two genes converge in the control of eye development. It is tempting to speculate that this genetic interaction may be related to a possible conservation of the control of SOS gene expression by spineless in this invertebrate species. Interestingly, invertebrate AhR orthologs (Spineless and AHR-1 from *Caenorhabditis elegans*) display structural and functional similarities with vertebrate AhR such as transcription factor activity and XRE binding. However, they do not bind dioxins or other hydrocarbon ligands and, at this stage, their possible receptor functions and their activation mechanisms have not been characterized. It is intriguing to consider that, while Spineless and AHR-1 are implicated in fundamental developmental processes, recent *in vivo* and *in vitro* evidences have established direct connections between the mammalian AhR and developmentally relevant pathways such as cellular proliferation, cellular migration and epithelial mesenchymal transition [42,43]. Thus, it is likely that several of those AhR functions including SOS1 regulation, have
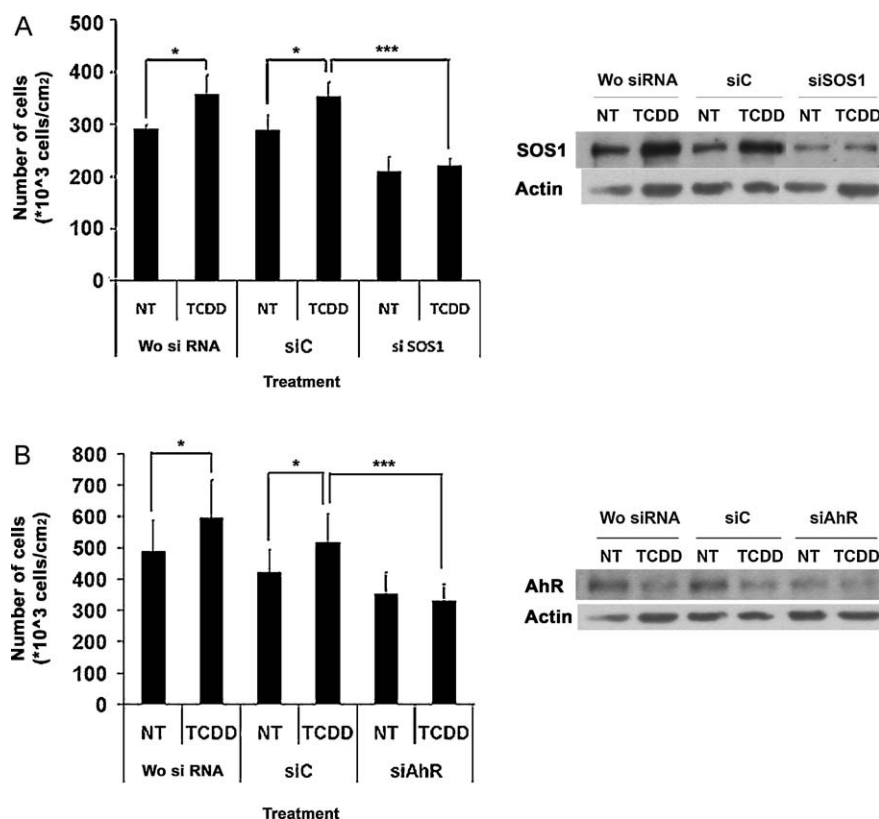
Fig. 6. TCDD-stimulated HepG2 cell growth depends on both AhR expression and increased expression of SOS1. (A) HepG2 cells were either not transfected (wo si RNA) or transfected with control siRNA (siC) or SOS1 siRNA (siSOS1). One day after, they were treated (T) or not (NT) with 25 nM TCDD for 24 h. Cell numbers, expressed in $*10^3$ cells/cm^2, were quantified for each condition. In the right panel of the figure, a western blot-analysis of SOS1 protein levels is presented. (B) HepG2 cells were either not transfected (wo si RNA) or transfected with siControl (siC) or siAhR (siAhR). They were treated (T) or not (NT) with 25 nM TCDD for 48 h. Cell numbers, expressed in $*10^3$ cells/cm^2, were quantified for each condition. In the right panel of the figure, a western blot-analysis of AhR protein levels is presented $*$, $p < 0.05$ compared to controls ($n = 3$).

been conserved through evolution and the present study as well as several others suggest that one mechanism of pollutants toxicity would be to disrupt such basic developmental functions of this receptor.

Conflict of interest statement

All authors disclose any actual or potential conflict of interest including any financial, personal or other relationships with other people or organizations within three (3) years of beginning the work submitted.

Acknowledgements

Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at doi:10.1016/j.bcp.2010.10.003.

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