

The Dioxin Receptor Regulates the Constitutive Expression of the *Vav3* Proto-Oncogene and Modulates Cell Shape and Adhesion

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The dioxin receptor (AhR) modulates cell plasticity and migration, although the signaling involved remains unknown. Here, we report a mechanism that integrates AhR into these cytoskeleton-related functions. Immortalized and mouse embryonic fibroblasts lacking AhR (*AhR*^{-/-}) had increased cell area due to spread cytoplasm that reverted to wild-type morphology upon AhR re-expression. The AhR-null phenotype included increased F-actin stress fibers, depolarized focal adhesions, and enhanced spreading and adhesion. The cytoskeleton alterations of *AhR*^{-/-} cells were due to down-regulation of constitutive Vav3 expression, a guanosine diphosphate/guanosine triphosphate exchange factor for Rho/Rac GTPases and a novel transcriptional target of AhR. AhR was recruited to the *vav3* promoter and maintained constitutive mRNA expression in a ligand-independent manner. Consistently, *AhR*^{-/-} fibroblasts had reduced Rac1 activity and increased activation of the RhoA/Rho kinase (Rock) pathway. Pharmacological inhibition of Rac1 shifted *AhR*^{+/+} fibroblasts to the null phenotype, whereas Rock inhibition changed AhR-null cells to the *AhR*^{+/+} morphology. Knock-down of *vav3* transcripts by small interfering RNA induced cytoskeleton defects and changes in adhesion and spreading mimicking those of AhR-null cells. Moreover, *vav3*^{-/-} MEFs, as *AhR*^{-/-} mouse embryonic fibroblasts, had increased cell area and enhanced stress fibers. By modulating Vav3-dependent signaling, AhR could regulate cell shape, adhesion, and migration under physiological conditions and, perhaps, in certain pathological states.

INTRODUCTION

A rapidly increasing number of studies consistently recognize that the aryl hydrocarbon (dioxin) receptor (AhR) serves a physiological function in the cell. AhR has been best characterized by its ability to mediate the toxic and carcinogenic effects of environmental toxicants such as 2,3,7,8-tetrachlorodibenzo-*p*-dioxin (TCDD) (Shimizu *et al.*, 2000; Safe, 2001; Marlowe and Puga, 2005). The mechanism of regulation of AhR during this xenobiotic-dependent process is relatively well known. Under basal conditions, AhR remains sequestered in the cytosol in a heteromolecular complex containing Hsp90, XAP2, p23, and, possibly, additional proteins. On binding to TCDD or other ligands, AhR moves to the nucleus where it interacts with the aryl hydrocarbon

receptor nuclear translocator and with a partially characterized set of coactivators and/or corepressors. These protein complexes regulate the expression of genes having xenobiotic responsive elements (XREs) in their promoters (Whitlock, 1999; Barouki *et al.*, 2007; Furness *et al.*, 2007). As expected, many of the AhR target genes code for xenobiotic-metabolizing enzymes (Hankinson, 1995; Nebert *et al.*, 2004). However, it is now known that AhR regulates other gene subsets with roles not directly linked to xenobiotic responses (Kolluri *et al.*, 2001; Matikainen *et al.*, 2001; Ogi *et al.*, 2001; Santiago-Josefat *et al.*, 2004; Chang *et al.*, 2007; Frericks *et al.*, 2007). The expanding array of AhR target genes, together with increasing evidence of AhR-regulated physiological functions in different animal models (Fernandez-Salguero *et al.*, 1995; Schmidt *et al.*, 1996; Mimura *et al.*, 1997; Barouki *et al.*, 2007; Furness *et al.*, 2007; McMillan and Bradfield, 2007), has fostered research aimed at defining new signaling pathways requiring AhR activity.

Several reports have implicated AhR in the regulation of cell morphology and migration in mammals and in *Caenorhabditis elegans* (Qin and Powell-Coffman, 2004; Barouki *et al.*, 2007), although the mechanisms involved remain rather obscure. Recent data have shown that the effects of AhR on the shape and motility of MCF-7 breast cancer cells are mediated via the c-Jun N-terminal kinase (Diry *et al.*, 2006), a downstream element of the Rac1 pathway (Bustelo

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Abbreviations used: AhR, aryl hydrocarbon (dioxin) receptor; FGM, immortalized mouse fibroblasts; FN, fibronectin; GAP, GTPase-activating protein; GEF, guanosine diphosphate/guanosine triphosphate exchange factor; MEF, mouse embryonic fibroblast; YFP, yellow fluorescent protein.

et al., 2007). We have also recently shown that immortalized mouse fibroblasts lacking AhR expression (T-FGM *AhR*^{-/-}) display a flattened morphology, increased stress fibers, and impaired migration (Mulero-Navarro *et al.*, 2005). Moreover, T-FGM *AhR*^{-/-} cells had lower focal adhesion kinase phosphorylation and decreased Rac1, phosphatidylinositol 3-kinase, and extracellular signal-regulated kinase activities (Mulero-Navarro *et al.*, 2005).

Cell morphology is the result of complex signaling pathways that act in concert to regulate F-actin dynamics and the turnover of cell-cell and cell-substratum interactions (Mitra *et al.*, 2005). The strength and extent of cell-substratum interactions significantly determines cell migration and can be highly relevant in processes requiring major changes in plasticity such as the epithelial-to-mesenchymal transition (EMT) (Li *et al.*, 2005). Many proteins coordinately interact next to the plasma membrane to regulate cell adhesion and migration. Rho/Rac GTPases such as RhoA, Rac1, and Cdc42 are relevant players in this process (Clark *et al.*, 1998; Nobes and Hall, 1999). Thus, it has been shown that RhoA regulates the formation of F-actin stress fibers and focal adhesions (Ridley and Hall, 1992; Chen *et al.*, 2002), that Rac1 promotes the formation of lamellipodia and membrane ruffles (Ridley *et al.*, 1992; Nobes and Hall, 1999), and that Cdc42 induced de formation of filopodia (Fukata *et al.*, 2003; Dianza *et al.*, 2006). To ensure coordinated cytoskeleton changes, there are multiple cross-talks between RhoA, Rac1, and other GTPases. For example, it is known that Rac1 stimulation leads to the down-modulation of RhoA activity and, as a consequence, to the reduction in stress fibers and focal adhesions in certain cell types (Rottner *et al.*, 1999; Sander *et al.*, 1999). As most Ras family members, Rho/Rac proteins have to cycle between an inactive (guanosine diphosphate [GDP]-bound) and an active (guanosine triphosphate [GTP]-bound) states. This cycling is modulated by different regulators. GDP/GTP exchange factors (GEFs) catalyze the exchange of GDP by GTP on the GTPases, thus favoring the rapid transition of these GTPases to their activated state during cell signaling. Instead, GTPase-activating proteins (GAPs) enhance the intrinsic GTPase activity of Rho/Rac proteins, a process that is essential for the hydrolysis of the bound GTP and therefore for the transition from the active to the inactive state at the end of the stimulation cycle. Rho/Rac proteins are also regulated by other processes, such as cytosolic sequestration (via GDP dissociation inhibitor proteins), ubiquitin-mediated degradation, gene expression, and phosphorylation (Bustelo *et al.*, 2007). When in the active state, these proteins bind to effector molecules that modulate many different biological processes such as cytoskeleton changes, gene transcription, vesicle trafficking, and cell proliferation (Bustelo *et al.*, 2007).

To shed further light in the mechanism by which AhR mediates these cytoskeleton-related events, we have studied the effect of *AhR* gene deficiency in cell shape, adhesion, and F-actin structures of both immortalized and primary fibroblasts. Furthermore, the use of microarray- and signaling-based strategies has allowed us to pinpoint the connection of AhR with the *vav3* proto-oncogene product, a phosphorylation-dependent exchange factor for Rho/Rac GTPases.

MATERIALS AND METHODS

Cell Culture

The *AhR*^{-/-} cells used in this study were obtained from AhR-null mice that were generated by gene knockout as reported previously (Fernandez-Salguero *et al.*, 1995). Immortalized T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} mouse fibroblasts were produced and characterized as described previously (Mulero-

Navarro *et al.*, 2005). T-FGM fibroblasts were grown in DMEM/F-12 medium containing 10% fetal bovine serum (FBS), 2 mM L-glutamine, 50 µg/ml gentamicin, and 11 mM D-glucose. Primary *AhR*^{+/+} and *AhR*^{-/-} MEF cells were isolated from 14.5 dpc mouse embryos as indicated (Santiago-Josefat *et al.*, 2001). *Vav3*-null and wild-type mice were produced as reported previously (Sauzeau *et al.*, 2006) and used to isolate primary mouse embryonic fibroblast (MEF) cells. MEFs were cultured in DMEM supplemented with 10% FBS, 2 mM L-glutamine, 100 U/ml penicillin, and 100 µg/ml streptomycin. Culturing conditions were 37°C and 5% CO₂ atmosphere. To induce actin stress fibers and to analyze changes in morphology, cells were maintained for 12 h in the absence of fetal bovine serum before experimentation.

Antibodies and Reagents

Antibody to paxillin was from BD Biosciences Transduction Laboratories (Lexington, KY). Anti-vinculin and anti-β-actin were from Sigma-Aldrich (St. Louis, MO). Anti-RhoA and anti-Rac1 were purchased from Cell Signaling Technology (Beverly, MA). Anti-Net1 was obtained from Santa Cruz Biotechnology (Santa Cruz, CA). Rhodamine-labeled phalloidin was from Invitrogen (Carlsbad, CA). *Taq* DNA polymerase was from Ecogen (Barcelona, Spain). SuperScript II reverse transcriptase was purchased from Invitrogen. SYBR Green I was obtained from Invitrogen and *Q*Taq DNA Polymerase mix from BD Biosciences (San Jose, CA). NSC23766 and Y27632 were obtained from Calbiochem (San Diego, CA). The expression construct for the constitutively active AhR (CA-AhR) was a generous gift from Dr. Fujii-Kuriyama (University of Tsukuba, Japan), and it was produced from the wild-type AhR by deleting the minimal PAS-B motif and by ligating the resulting construct to the green fluorescent protein (EGFP) (Ito *et al.*, 2004).

SDS-Polyacrylamide Gel Electrophoresis (PAGE) and Western Immunoblotting

SDS-PAGE and Western blotting were performed using total cell extracts, as described previously (Santiago-Josefat *et al.*, 2004; Mulero-Navarro *et al.*, 2005).

Measurement of Cell Area, Immunocytochemistry, and Rhodamine-Phalloidin Staining

Cell area and the minor/major axis ratio were measured in *AhR*^{+/+} and *AhR*^{-/-} T-FGM cells and MEFs by blinded analysis in different random fields for each cell culture. Cellular contour and axis were determined using the ImageJ software (National Institutes of Health, Bethesda, MD). Significance of the data was determined as indicated below (see Statistical Analyses). Fluorescence-activated cell cytometry (FACS) was used to determine that T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} had similar cell volume. For immunocytochemistry, cultures were fixed for 20 min at room temperature in 4% paraformaldehyde (Polysciences, Warrington, PA). Vinculin and paxillin immunostaining was used to locate focal adhesions in T-FGM cells and MEFs as reported previously (Santiago-Josefat *et al.*, 2004). To label actin stress fibers, T-FGM fibroblasts and MEF cells were also stained with rhodamine-phalloidin (Invitrogen) as indicated (Mulero-Navarro *et al.*, 2005). Fluorescence microscopy images were taken using an Axioplan epifluorescence microscope (Carl Zeiss, Jena, Germany) equipped with Plan-Neufluor lenses with magnifications of 20× (0.50 aperture) and 40× (0.75 aperture). Pictures were also taken using a Plan Aplanachromat 63× lens (1.40 aperture). Fluorochromes used were Alexa 488, rhodamine, EGFP, and enhanced yellow fluorescent protein (EYFP). Images were captured with a CoolSNAP camera (RS Photometrics, Tucson, AZ) and analyzed using the RS software.

Expression Constructs, Transient Transfections, and RNA Interference

To generate a vector encoding the AhR-EYFP fusion protein, the full-length *AhR* cDNA was cloned into the pEYFP vector as indicated previously (Santiago-Josefat *et al.*, 2001). To generate a mammalian expression vector for CA-AhR-EYFP, a modified *AhR* cDNA encoding a constitutively active version of the receptor was amplified by polymerase chain reaction (PCR) from pEFBOS-mAhR-CA and cloned into pEYFP by using previously described protocols (Santiago-Josefat *et al.*, 2001). Transient transfections were performed in T-FGM by using the Lipofectamine Plus reagent (Invitrogen) and 1 µg of AhR-EYFP, CA-AhR-EYFP, or *Vav3*-EGFP expression vectors (Movilla and Bustelo, 1999). Protein expression was maintained for 24 h, and cultures were analyzed by fluorescence microscopy as indicated below. *vav3* siRNA experiments were done in T-FGM cells by transient transfection of either 20 or 100 nM of commercially available siGenome on-Target plus Smart pool (Dharmacon RNA Technologies, Lafayette, CO). Negative control experiments were also performed in parallel by transient transfection of unspecific scramble siRNAs (Dharmacon RNA Technologies).

Reverse Transcription and Real-Time PCR

Total RNA was isolated from T-FGM fibroblasts and MEF cultures by using the RNeasy kit (QIAGEN, Hilden, Germany). Reverse transcription was performed using random hexamers priming and SuperScript II transcriptase as

indicated previously (Gomez-Duran *et al.*, 2006). Real-time PCR was performed to quantify the expression levels of the *vav3* and *Cyp1a1* mRNAs. Primers for *vav3* were 5'-GGAGTGGAGTCAGCCATCTC-3' (forward) and 5'-ATTGGAACGACCAGCAAATC-3' (reverse). For *Cyp1a1* were 5'-ACAGACAGCCTCATTGAGCA-3' (forward) and 5'-GGCTCCACGAGATAGCAGTT-3' (reverse). We used the expression of the β -actin mRNA as normalization control. This cDNA was amplified using the oligonucleotides 5'-GGTCAGAAG-GACTCTATGTGG-3' (forward) and 5'-TCCTCTCAGCTGTGGTGGT-3' (reverse). Reactions were done using SYBR Green I/QTaq DNA Polymerase Mix (BD Biosciences) on an iCycler equipment (Bio-Rad, Hercules, CA), as described previously (Gomez-Duran *et al.*, 2006).

Cloning of the *vav3* Promoter, Reporter Gene Analyses, and Chromatin Immunoprecipitation

The mouse *vav3* promoter was cloned from the genomic sequence included in the W11-2862P14 phosmid vector (Wellcome Trust Sanger Institute, Cambridge, United Kingdom). Briefly, the W11-2862P14 clone was digested with BglII and a 3574-base pair fragment containing the *vav3* promoter cloned into the pGL2-Basic vector. The resulting clone was then digested with BglII + BstEII and a 2005-base pair fragment isolated and made blunt ended with Klenow DNA polymerase. The 2005-base pair fragment was finally ligated into the SmaI-digested pGL2-Basic vector. The *vav3* promoter sequence was analyzed for potential XRE binding sites by using the consensus element 5'-GCGTG-3'. Correct orientation of the *vav3* promoter was confirmed by digestion with a panel of restriction endonucleases. Luciferase reporter gene assays were done as indicated previously (Roman *et al.*, 2008) by using the dual-glow luciferase kit (Promega, Madison, WI). The pRLTK vector was used as normalization control. Firefly and *Renilla* luciferase activities were determined in a microtiter plate luminometer MLX (Dyex Technologies, Chantilly, VA). Chromatin immunoprecipitation (ChIP) to analyze AhR binding to the *vav3* promoter was performed essentially as described previously (Mulero-Navarro *et al.*, 2006; Roman *et al.*, 2008). Oligonucleotides used for PCR were as follows: XRE1 forward 5'-TCATCCGAGGGTTCACG-3' and reverse 5'-GCACTG-GCTCCGACTG-3'; XRE2 forward 5'-AACCTACTTCTTCCCTCTAC-3' and reverse 5'-GGTCCATCTCAGCCCTTCC-3'; XRE3-4 forward 5'-TCCTAGGCTGATTTATTATTGTT-3' and reverse 5'-GCTGCTGTGTTTCAT-TAGAT-3'.

Site-directed Mutagenesis

The XRE1 site in the proximal *vav3* promoter was inactivated by site-directed mutagenesis changing the consensus sequence for AhR binding 5' CACGC 3' to 5' CAGGA 3' by PCR (Gomez-Duran *et al.*, 2008a) (mutated bases underlined). Briefly, the pGL2-Vav3 full-length construct was partially digested with BglII and the linearized DNA further cut with PstI. This fragment was amplified by PCR using the oligonucleotides forward 5'-CCAGCCAGGGCG-GCGGGCAGGATCCACCCG 3' and reverse 5' CCGGCCGCGCTGCAG-CAGCGAGTGCC 3'. Amplification was carried out for 30 cycles in 50- μ l reaction mixture containing 10 mM Tris-HCl, pH 8.3, 50 mM KCl, 1.5 mM MgCl₂, 0.2 mM each dNTP, 0.5 μ M each primer, and 2.5 U of *Taq* polymerase. Cycling conditions were as follows: denaturation at 94°C for 1 min, annealing at 60°C for 1 min, and extension at 72°C for 1 min. This PCR fragment was then digested with BglII and PstI and cloned into the pGL2-Basic vector cut previously with the same restriction endonucleases. The resulting reporter construct (pGL2-Vav3-mutXRE1) was transiently transfected into Hepa 1c1c7 and c2 hepatoma cell lines as described above.

Cell Adhesion and Cell Detachment Assays

Cell adhesion was performed on plastic or fibronectin-coated culture dishes. Twelve-well plates were coated with 0.04, 0.2, 0.7, 1, 4, or 10 μ g/ml fibronectin. After washing with phosphate-buffered saline (PBS) and culture medium, T-FGM cells and MEFs were seeded at 3×10^5 cells/well. After an incubation of 15 min (T-FGM cells) or 1 h (MEFs), cells were washed in PBS and fixed overnight at 4°C in 3.7% paraformaldehyde. Cultures were then stained for 15 min with crystal violet (20% methanol solution). After washing and drying, attached and expanded cells were counted *de visu* by using light microscopy. Cell adhesion on plastic was done as above using untreated culture plates. Cell detachment experiments were also done on plastic using cultures seeded at the same cell density. After treatment for the indicated periods with a solution containing 4 mM EGTA and 1 mM MgCl₂, cultures were fixed in ice-cold methanol and stained with 4',6-diamidino-2-phenylindole (DAPI). Cells that remained attached were counted using the ImageJ software.

RhoA and Rac1 Activation Assays

Pull-down assays with a glutathione transferase (GST) fusion protein containing the RhoA binding domain of rhotekin (rhotekin-GST) or the Rac1 binding domain of PAK-CRIB (PAK-CRIB-GST) were performed essentially as described previously (Mulero-Navarro *et al.*, 2005). Samples were analyzed for activated and total RhoA and Rac1 by Western immunoblotting using specific antibodies.

Microarray Experiments

Two-color cDNA microarrays containing probes for 15,000 mouse genes were used. Total RNA isolation, labeling, hybridization, normalization, and differential expression analyses were made as described previously (Martinez-Delgado *et al.*, 2004).

Statistical Analyses

Data are shown as mean \pm SEM. Statistical comparison between experimental conditions was done using GraphPad Prism 4.0 software (GraphPad Software, San Diego, CA). One-way analysis of variance followed by Dunn's test was applied. For the analyses of cell area and minor/major axis ratio, the Mann-Whitney nonparametric median statistical analysis was used.

RESULTS

Dioxin Receptor Deficiency Induces Changes in Cytoskeleton-related Processes in Both Immortalized and Primary Fibroblasts

We have shown previously that immortalized mouse fibroblasts lacking AhR expression (T-FGM *AhR*^{-/-} cells) had impaired migration rates (Mulero-Navarro *et al.*, 2005). To further characterize the influence of AhR deficiency in this biological process, we decided to study in detail the changes in shape and adhesion of those cells. In addition, and to avoid secondary effects derived from the immortalization process, we also conducted these experiments with primary embryonic fibroblasts (MEFs) derived from *AhR*^{+/+} or *AhR*^{-/-} mice. We observed that, irrespective of the cell type used, AhR deficiency always correlated with the generation of cells with enlarged cell area and reduced polarity (Figure 1A). Quantification and statistical comparisons of the minor/major axis ratio and cell areas corroborated these *de visu* observations (Figure 1B). However, the increase in cell area of *AhR*^{-/-} cells with respect to *AhR*^{+/+} fibroblasts was not the consequence of higher cell volumes as determined by FACS analyses (data not shown). Transfection of AhR-deficient cells with mammalian expression vectors encoding yellow fluorescent proteins (YFPs) fused to either wild-type or constitutively active (lacking the minimal PAS-B domain) versions of AhR restored the typical shape and size of the cells (Figure 2, A and B; and Supplemental Figure S1), confirming that these differences in cell morphology were a direct consequence of the loss of AhR expression. YFP alone did not alter cell shape or size of T-FGM *AhR*^{+/+} or T-FGM *AhR*^{-/-} fibroblasts, further excluding an unspecific effect (Supplemental Figure S2).

To analyze the cytoskeleton in AhR-deficient fibroblasts, we next stained T-FGM cells and MEFs of the indicated genotypes with antibodies to paxillin and rhodamine-labeled phalloidin to monitor the status of focal adhesions and the F-actin cytoskeleton in the presence or absence of AhR expression, respectively. Anti-paxillin immunofluorescence experiments indicated that the AhR deficiency was associated with an increase in focal adhesions (Figure 3, A and B, a and b) and stress fibers (Figure 3, A and B, c and d) both in T-FGM cells and MEFs. Unlike AhR-positive cells, the focal adhesions present in AhR-deficient cells were usually detected around all the cell perimeter rather than been limited and polarized to areas of membrane protrusions (Figure 3, A and B, a and b). The difference in focal adhesions among *AhR*^{+/+} and *AhR*^{-/-} cells could be also influenced by altered expression of focal adhesion (FA)-localized proteins. Although paxillin expression was similar in *AhR*^{+/+} and *AhR*^{-/-} T-FGM cells and MEFs, vinculin levels seemed moderately elevated in T-FGM and MEF *AhR*^{-/-} fibroblasts (Supplemental Figure S3). Considering that vinculin stabilizes FAs and thereby decreases cell migration (Ziegler *et al.*, 2006), the contribution of this protein to both increased

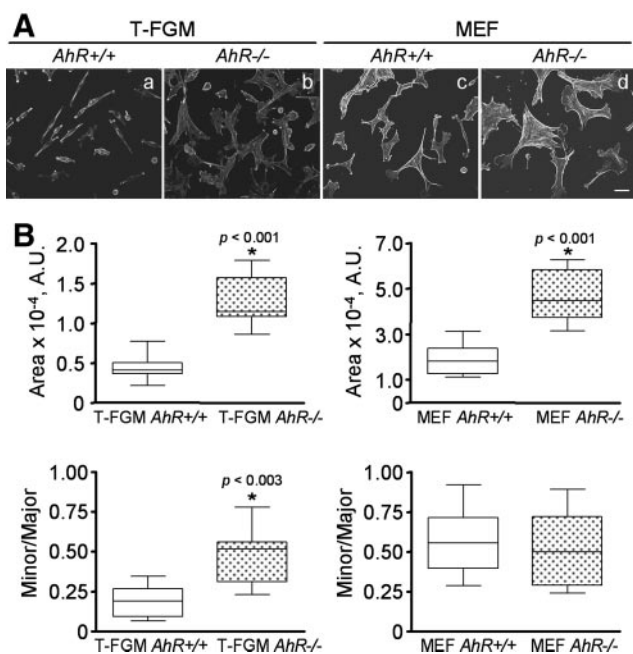


Figure 1. Dioxin receptor expression modulates cell morphology. (A) Immortalized T-FGM cells and MEFs from *AhR*^{+/+} and *AhR*^{-/-} mice were plated at the same cell density for each cell type and starved in the absence of serum for 12 h. After fixation, cultures were stained with rhodamine-phalloidin and the cellular contours were defined using the ImageJ software. (B) Cell area was measured for each genotype and cell type, and data represented as median ± SEM (nonparametric Mann–Whitney *U* test). Minor/major axis ratios were also measured and data analyzed as indicated. At least 15 cells were analyzed for each experimental condition, and the experiments were done in at least two different cultures. The *p* values for statistical comparison between genotypes are indicated. The calibration bar corresponds to 20 μm. A.U., arbitrary units.

FA content and lower migration of T-FGM and MEF *AhR*^{-/-} fibroblast (Mulero-Navarro *et al.*, 2005; Gomez-Duran *et al.*, 2009) deserves further consideration.

The functional relevance of AhR in cytoskeleton-related events was further analyzed by cell spreading and detachment experiments. When plated on fibronectin (FN)-coated dishes, T-FGM *AhR*^{-/-} cells displayed higher spreading rates than their wild-type counterparts (Figure 4, A and B, left). However, these differences became less acute when these experiments were repeated with coverslips coated with higher FN concentrations (i.e., 10 μg/ml; Figure 4, A and B, right). Similar results were observed in *AhR*^{-/-} MEFs, although in this case the difference in spreading with respect to wild-type cells was less pronounced (Figure 4, A and B, right). The lower spreading rates of *AhR*^{+/+} T-FGM cells and MEFs was not due to initial defects in substrate attachment, because total cell counts indicate equal numbers of attached cells to the dishes regardless of the genotype used (Figure 4C). The differences in spreading between *AhR*^{+/+} and *AhR*^{-/-} fibroblasts became even more prominent when the plating step was performed on a plastic substratum. Thus, whereas T-FGM *AhR*^{+/+} cells showed negligible spreading on plastic after 60 min of being plated, 50% of the T-FGM *AhR*^{-/-} cells efficiently spread under identical conditions (Figure 5, A and B, left). Likewise, *AhR*^{-/-} MEFs displayed much higher spreading rates compared with wild-type MEFs (Figure 5, A and B, right). These differences in spreading behavior were detected until at least

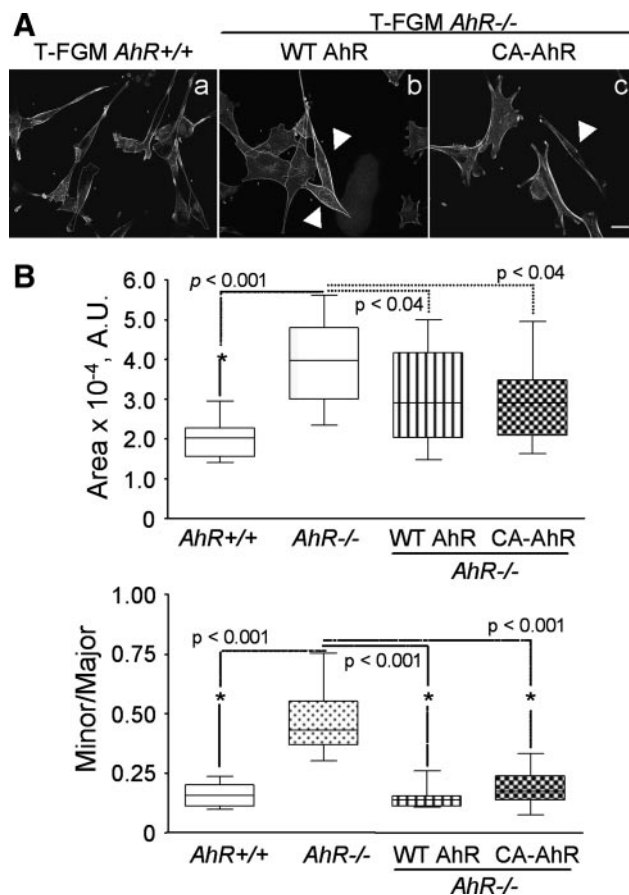


Figure 2. AhR re-expression changes the AhR-null to the wild-type phenotype. (A) T-FGM *AhR*^{-/-} cells were plated and transfected with wild-type AhR-EYFP (WT AhR) or constitutively active AhR-EYFP (CA-AhR) expression constructs. Protein expression was allowed for 24 h under fetal bovine serum deprivation. Cells were then stained with rhodamine-phalloidin and analyzed by fluorescence microscopy. The ImageJ software was used to measure YFP-labeled cells in each experimental condition. Arrowheads indicate cells expressing the ectopic proteins. (B) Cell area and minor/major axis ratios were calculated for each experimental condition and data represented as median ± SEM. At least 15 transfected cells were analyzed for each genotype, and the experiments were done in triplicate. The *p* values for statistical significance are indicated. The calibration bar corresponds to 10 μm. A.U., arbitrary units.

4 h after plating (Figure 5B, right). Additional controls indicated again that these differences were not due to initial defects in cell attachment to the plates (Figure 5C).

To further verify the implication of AhR in cell–substratum interactions, we resorted to measuring their kinetics of detachment upon Ca²⁺ chelation by EGTA treatment (see *Materials and Methods*). As shown in Figure 6A, the kinetic of EGTA-dependent detachment was much slower in T-FGM *AhR*^{-/-} cells than in their wild-type counterparts. Similar results were obtained when *AhR*^{-/-} and *AhR*^{+/+} MEFs were compared, although, similarly to the cell spreading experiments, the differences with respect to wild-type cells were less pronounced than in the case of the immortalized T-FGM cell line (Figure 6B). Together, these data strongly support a role for AhR in the regulation of cell shape, cytoskeleton organization, and adhesion. These observations are also in good agreement with our previous results indicating that T-FGM *AhR*^{-/-} fibroblasts show lower migration activity (Mulero-Navarro *et al.*, 2005).

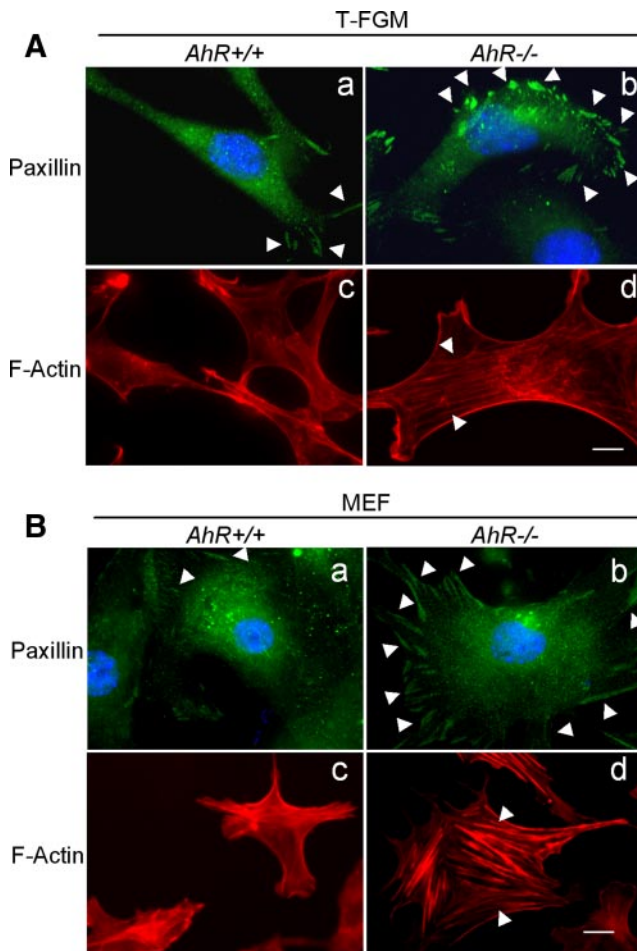


Figure 3. AhR-null cells have unpolarized focal adhesions and increased actin stress fibers. (A) T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells were plated, starved for 12 h by fetal bovine serum deprivation and analyzed for focal adhesions content and distribution using anti-paxillin immunostaining and an Alexa 488-labeled secondary antibody (a and b) or for F-actin by using rhodamine-phalloidin staining (c and d). (B) The same experiments were performed in MEF cultures for focal adhesions (a and b) or actin stress fibers (c and d). The experiment was repeated in at least four different cultures of each cell type and genotype with same results. Bar, 5 μ m. Arrowheads mark focal adhesions in a and b or actin stress fibers in c and d.

The GTPases Rac1 and RhoA Are Involved in the AhR-dependent Cytoskeleton Changes

Because cytoskeleton-dependent events such as cell shape and adhesion are regulated by members of the Rho/Rac GTPase subfamily (Nobes and Hall, 1999; Fukata *et al.*, 2003; Bustelo *et al.*, 2007), we next explored whether variations in the expression and/or activity of these GTPases could be the cause of the phenotype detected in our fibroblast cells. Initial experiments indicated that the expression of Rac1, RhoA, and Cdc42 did not change at the mRNA or protein level (Figure 7A; data not shown), suggesting that AhR does not affect the expression of these important cytoskeleton regulators. However, pull-down assays using a PAK-CRIB-GST fusion protein revealed that T-FGM *AhR*^{-/-} cells had reduced levels of activated GTP-Rac1 compared with wild-type T-FGM fibroblasts (Figure 7A, left). Because it has been reported that Rac1 inhibits RhoA activity in a number of cell systems (Rottner *et al.*, 1999; Sander *et al.*, 1999), we also

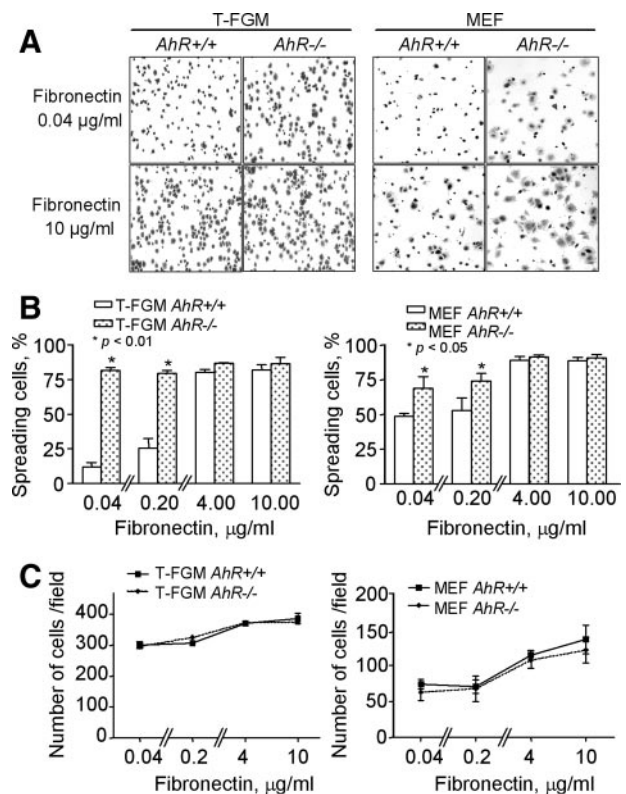


Figure 4. The absence of AhR increases cell spreading on fibronectin. (A) T-FGM cells and MEF of the indicated genotypes were cultured on dishes previously coated with increasing concentrations of FN (0.04, 0.2, 4.0, or 10 μ g/ml). Representative results obtained at 0.04 and 10 μ g/ml are shown for each genotype and cell type. (B) Cell spreading was quantified at 15 min (T-FGM cells) or 60 min (MEFs) by cell counting after crystal violet staining. Cytoplasmic extensions indicative of spreading were determined by blinded analysis. Percentages of spread cells were referred to the total number of cells within the same microscopy field. Between 250 and 300 T-FGM cells or 100 MEFs were counted in six independent fields for each experimental condition. The p values for statistical comparison between genotypes are indicated. (C) The total number of cells attached to the plates was counted for each FN concentration. Data are shown as mean \pm SEM from cells counted in at least two different cultures of each genotype and cell type.

evaluated whether the increase in stress fiber formation and the reduction in Rac1 activity detected in T-FGM *AhR*^{-/-} cells could be due to increased activity of the RhoA pathway. To this end, we analyzed the activation status of RhoA by using pull-down assays with a Rhotekin-GST fusion protein. These experiments showed that T-FGM *AhR*^{-/-} did have higher levels of GTP-bound RhoA than the wild-type cells (Figure 7A, right), indicating that the former cell line has higher levels of RhoA activity than the latter one.

To verify whether the activity of the Rac1 and RhoA signaling pathways was altered, we treated T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells with NSC23766, a drug that blocks the interaction of Rac1 with upstream GDP/GTP exchange factors and, thereby, inhibits its activation in vivo (Gao *et al.*, 2004). NSC23766 efficiently inhibited Rac1 activation in T-FGM *AhR*^{+/+} cells (see Figure 10C, bottom) and, interestingly, we observed that the addition of this compound to T-FGM *AhR*^{+/+} cells induced a morphologically phenotype similar to that found in T-FGM *AhR*^{-/-} cells, whereas it did not significantly affect the phenotype of AhR-

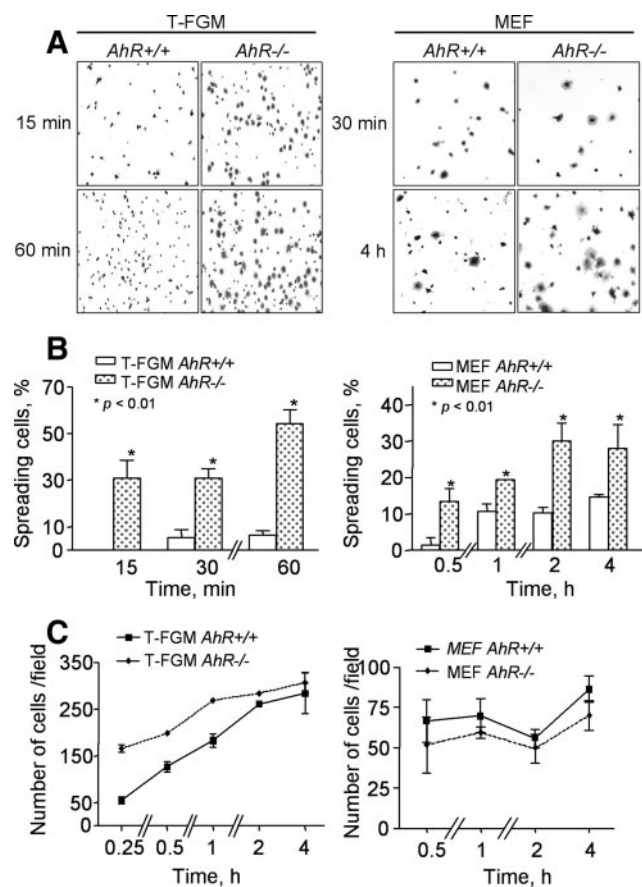


Figure 5. The absence of AhR increases cell spreading on plastic. (A) T-FGM cells and MEFs from *AhR*^{+/+} and *AhR*^{-/-} mice were cultured on untreated plastic dishes for the indicated times and cells stained with crystal violet. (B) Cell spreading was quantified for each experimental condition as indicated in the legend for Figure 4. The p values for statistical comparison between genotypes are indicated. (C) The variation with time of the total number of attached cells was also determined for each cell type and genotype. Data are shown as mean ± SEM from cells counted in at least two different cultures of each genotype and cell type.

null cells (Figure 7B, compare a and b). Those changes seen in T-FGM *AhR*^{+/+} cells included the prototypical increase in both the cell area and stress fiber formation, a result that

suggest that the phenotype induced by AhR deficiency may be mediated by decreased signals from the Rac1 pathway. Next, we treated *AhR*^{+/+} and *AhR*^{-/-} T-FGM cells with Y27632, a specific inhibitor of the serine/threonine kinase Rock (Uehata *et al.*, 1997). This kinase is a downstream element of RhoA that mediates the formation of stress fibers and the actomyosin ring (Riento and Ridley, 2003; Shimokawa and Rashid, 2007). Y27632 induced the rapid disassembly of the stress fibers and, in addition, the acquisition of a T-FGM *AhR*^{+/+}-like morphology in the *AhR*-null cells (Figure 7C, compare a and c). These results indicate that the AhR-dependent changes in cell morphology in these fibroblast cell lines rely, at least in part, in the differential engagement of the GTPases Rac1 and RhoA.

AhR Regulates the Constitutive Transcription of *Vav3*, which Is a Major Intermediate Modulating the AhR-dependent Phenotype

Because the expression levels of Rac1 and RhoA did not change depending on the AhR status of the cell (see above), we speculated that the variations in the activities of their signal transduction pathways could be mediated by alterations in upstream molecules (GEFs), negative regulators (i.e., GAPs), and/or downstream elements. Given the large number of putative AhR targets, we decided to conduct microarray experiments to obtain a global, genome-wide view of the transcriptomal differences present in T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells. The functional annotation of the transcriptomal changes indicated that the AhR deficiency induced the down-modulation of a subset of RhoA (Net1) (Alberts and Treisman, 1998), Rac1 (Sos1, Vav3, and Abr) (Chuang *et al.*, 1995; Nimnual *et al.*, 1998; Bustelo, 2000), and Ras (Sos1) (Boguski and McCormick, 1993) GEFs as well as of two Rab family members (Rab1 and Rab9) (Figure 8A). In addition, it up-regulated the expression of two Rho/Rac GAPs (ArhGAP18 and ArhGAP20) and the Rab8 GTPase (Figure 8A). Two main regulators of RhoA and Rac1 GTPases that seemed markedly repressed in T-FGM *AhR*^{-/-} cells were Vav3 and Net1 (Burrige and Wennerberg, 2004). Given previous results indicating the importance of Vav3 in cytoskeleton organization (Movilla and Bustelo, 1999; Hornstein *et al.*, 2004; Couceiro *et al.*, 2005) and its marked down-regulation in T-FGM *AhR*^{-/-} cells, we decided to verify whether the modulation of this exchange factor by AhR could be a component of the cytoskeleton-related effects of this transcriptional factor. To this end, we first investigated whether the *vav3* proto-oncogene was indeed a potential

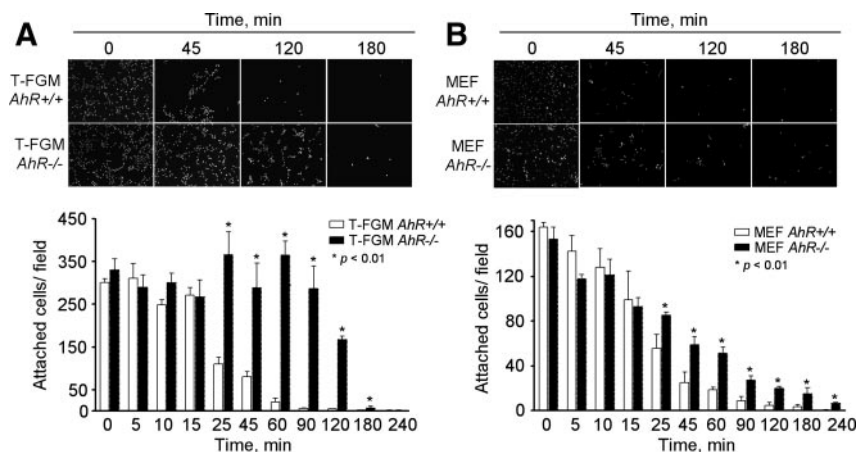


Figure 6. AhR expression decreases cell adhesion to the substratum. (A) T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells were seeded at the same cell density on plastic dishes and then induced to detach by using treatments with EGTA for the indicated times. Nuclei were stained with DAPI, and attached cells counted using the ImageJ software. (B) MEFs of the indicated genotypes were cultured at the same initial cell density and processed and analyzed as described in A. Data are shown as mean ± SEM for measurements made in duplicate in at least two independent cultures for each genotype and cell type. The p values for statistical comparison between genotypes are indicated.

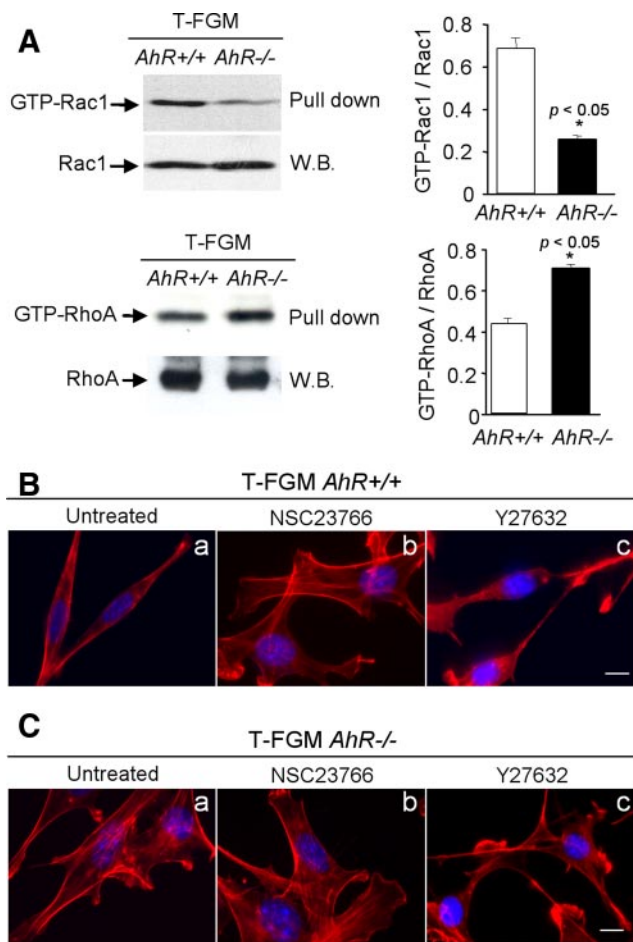


Figure 7. Rac1 and RhoA contribute to the AhR-dependent control of cell morphology. (A) Levels of Rac1 (GTP-Rac1) and RhoA (GTP-RhoA) activation were measured in T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells by pull-down assays by using the PAK-CRIB-GST or Rhotekin-GST target proteins, respectively. Total cellular levels of Rac1 and RhoA were also determined by Western blot (W.B.) using anti-Rac1 and anti-RhoA antibodies. The levels of GTP-RhoA and GTP-Rac1 were quantified and normalized by total RhoA and Rac1 expression, respectively. Data are shown as mean \pm SEM from at least two independent cultures. The p values for statistical comparison between genotypes are indicated. (B) T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells were plated and treated for 6 h with the Rac1 inhibitor NSC23766 (5 μ M) or (C) with the Rock inhibitor Y27632 (10 μ M). Cells were then fixed, stained with rhodamine-phalloidin, and photographed. The experiment was done in three different cultures with the same results. Bar, 5 μ m.

AhR transcriptional target. Consistent with the microarray results, we observed that the *vav3* transcript was remarkably decreased in *AhR*^{-/-} T-FGM and MEFs cells, as determined by quantitative, real-time PCR analysis (Figure 8B, left and middle). Importantly, unlike other classical AhR targets such as *Cyp1a1* (Figure 8B, right) (Hankinson, 1995; Whitlock, 1999; Nebert *et al.*, 2004), the expression of the *vav3* mRNA could not be enhanced by the addition of the AhR ligand TCDD to T-FGM *AhR*^{+/+} cells (Figure 8B, left and middle). These results indicate that AhR functions as a transcription factor committed to maintain the constitutive expression of *vav3*, and suggest a relevant role for the physiological levels of Vav3 in the regulation of the AhR-dependent phenotype. Regarding Net1, we have analyzed protein expression by Western immunoblotting and found that, although re-

pressed in our cDNA expression arrays, Net1 had similar protein levels in T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} (Figure 8C). Even though the regulation of Net1 could deserve further analysis, based on its similar protein levels in T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} fibroblasts, we focused our study on Vav3.

We decided to isolate the mouse *vav3* gene promoter and to further characterize its regulation by AhR. A 2005-base pair upstream genomic fragment containing the *vav3* promoter was cloned that contained four XRE consensus elements for AhR binding. These were located at -369 (XRE1), -823 (XRE2), -1484 (XRE3), and -1499 (XRE4) with respect to the translation start codon ATG (Figure 9A, top). We then analyzed the functionality of this promoter sequence by luciferase reporter assays using wild-type mouse hepatoma Hepa 1c1c7 cells and its variant c2 clone (expressing <10% wild-type levels of AhR). In agreement with an AhR-dependent process, *vav3* promoter activity was higher in wild-type 1c1c7 than in AhR-deficient c2 hepatoma cells (Figure 9A, right). Furthermore, the reexpression of AhR by the pcDNA-AhR expression vector in c2 cells restored *vav3* promoter activity to values close to those observed in wild-type 1c1c7 cells (Figure 9A, right). To further demonstrate that AhR contributes to the transcriptional regulation of *vav3* expression, ChIP assays were conducted. AhR binding to the XRE1, XRE2, and XRE3-4 elements was analyzed by PCR in AhR immunoprecipitates. The results showed that AhR was recruited to the proximal XRE1 element on the *vav3* promoter (Figure 9B), thus supporting the fact that *vav3* is an AhR-regulated gene in the mouse. Moreover, the XRE1 element is indeed necessary for *vav3* promoter regulation since its inactivation by site-directed mutagenesis abrogated the ability of the *vav3* promoter to drive luciferase reporter activity in AhR-expressing wild-type 1c1c7 cells (Figure 9C).

These results prompted us to further analyze the contribution of Vav3 to the AhR-dependent phenotype. We hypothesized that if a reduction in the physiological levels of Vav3 expression in AhR-deficient cells were involved in determining their characteristic morphological phenotype, the knockdown of the *vav3* mRNA in wild-type fibroblasts should induce an *AhR*-null-like phenotype. To address this possibility, we transfected *vav3*-specific small interfering RNAs (siRNAs) in T-FGM *AhR*^{+/+} cells, and, 72 h later, we monitored the morphology, polarity, and cytoskeleton changes in mock- and *vav3* siRNA-transfected cells by using microscopy techniques. As an additional control, we used in these analyses untransfected T-FGM *AhR*^{-/-} cells. We initially determined by Western immunoblotting that T-FGM *AhR*^{-/-} cells expressed lower Vav3 protein levels and that *vav3*-specific siRNAs, but not unspecific scramble siRNAs, consistently reduced Vav3 protein expression to ~30% its basal levels in T-FGM *AhR*^{+/+} cells ($p < 0.05$) (Figure 10A). Notably, *vav3* knockdown induced a shift in the morphology of T-FGM cells from the wild-type to the *AhR*-null-like phenotype (Figure 10B). This change included the increase in cell area (Figure 10B, compare a with c and d and e with g and h), the promotion of stress fiber formation (Figure 10B, compare a with c and d), increased numbers of focal adhesions that localize around the cell edges (Figure 10B, compare e with g and h), and cell depolarization (Figure 10B, compare e with g and h). Transfection of unspecific scramble siRNAs, on the contrary, did not change these phenotypes in T-FGM *AhR*^{+/+} cells (Figure 10B, b and f). We then analyzed whether knockdown of Vav3 expression in fact affected its downstream target proteins RhoA and Rac1. Consistent with the role proposed for Vav3 in the regulation of the Rac1 and RhoA signaling pathways, down-modulation

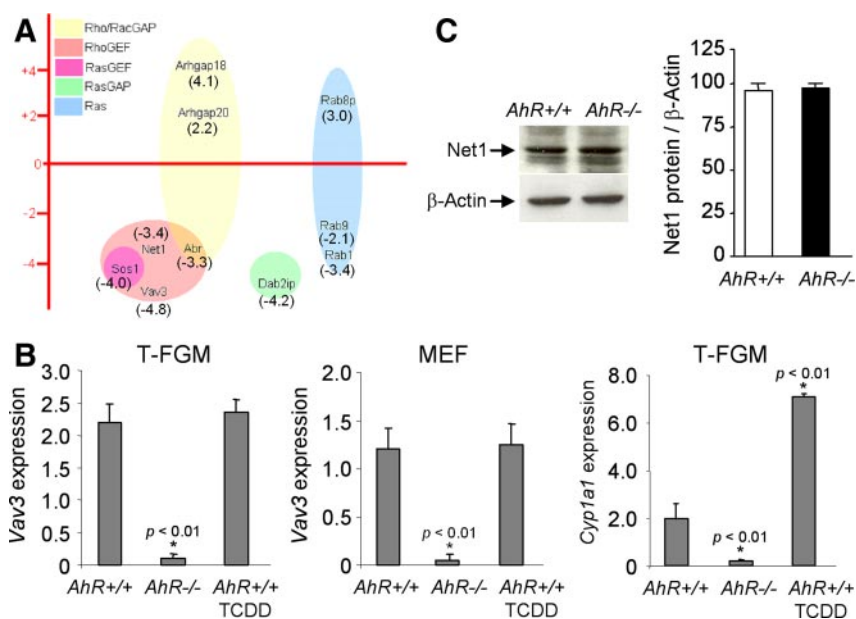


Figure 8. AhR regulates constitutive *vav3* mRNA expression. (A) Total RNA purified from T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells was analyzed by microarrays as indicated in *Materials and Methods*. Genes above and below the horizontal dotted line are over-expressed or repressed, respectively. (B) *vav3* mRNA expression was determined by real-time PCR in T-FGM cells and MEFs of the indicated genotypes under basal conditions or after a 24-h-long treatment of TCDD (10 nM). As positive control for TCDD-dependent AhR activation, mRNA levels of the target gene *Cyp1a1* were determined in parallel. Gene expression was normalized among the different experimental conditions using as comparative control the expression of the β -actin gene. (C) *Net1* protein expression was analyzed in T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} by Western immunoblotting using 15 μ g total protein and a specific antibody. The expression of β -actin was used as loading control. Data are shown as mean \pm SEM from triplicate measurements. The experiments were done in two different cultures of each genotype and cell line. The p values for statistical comparison between genotypes and between

untreated and TCDD-treated *AhR*^{+/+} T-FGM cells are indicated.

of *Vav3* protein levels by *vav3* siRNAs had an effect on these GTPases, and thus RhoA activation was increased, whereas Rac1 activation was decreased in transfected T-FGM *AhR*^{+/+} cells (Figure 10C). As indicated above, transfection of unspecific scramble siRNAs did not affect RhoA or Rac1 activation in T-FGM fibroblasts of either genotype (Figure 10C). We also confirmed that Rac1 has a prominent role in the phenotypic changes induced by NSC23766 in T-FGM *AhR*^{+/+} cells (Figure 7B) because this molecule decreased Rac1 activity in our pull-down assays (Figure 10C, bottom right). These results support that AhR, through the control of constitutive *Vav3* expression and by modulating the activation of its downstream target proteins RhoA and Rac1, has a relevant role in determining fibroblast cell phenotype.

In addition, because *Vav3* can be regulated by phosphorylation (Bustelo, 2002; Llorca *et al.*, 2005), we also analyzed whether T-FGM *AhR*^{-/-} cells had a lower phosphorylation potential that could contribute to a reduction in *Vav3* signaling. T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells were transfected with a myc-tagged *Vav3* expression construct and the levels of p*Vav3* determined by Western immunoblotting in myc immunoprecipitates. Both cell lines had a similar potential to phosphorylate *Vav3* (Supplemental Figure S4), further supporting that decreased *Vav3* protein expression is a major determinant in the AhR-dependent phenotype. We also performed overexpression experiments in which T-FGM *AhR*^{-/-} and T-FGM *AhR*^{+/+} fibroblasts were transfected with the wild type or with a constitutively

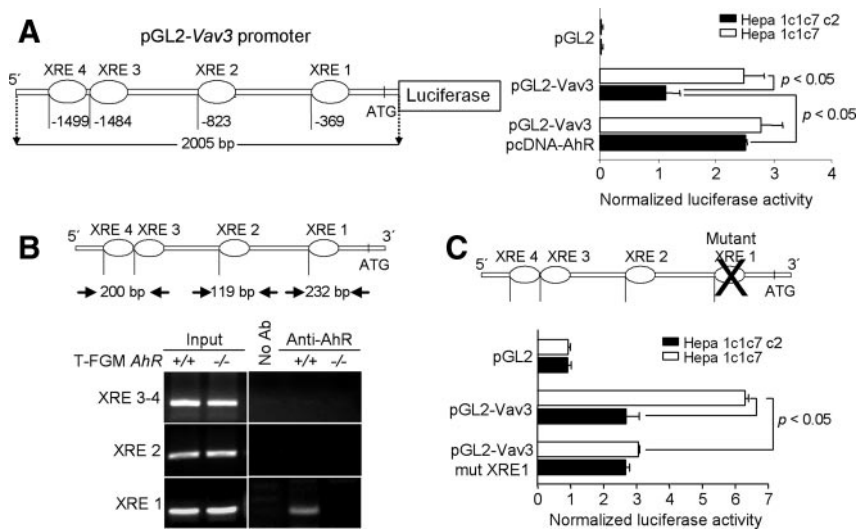


Figure 9. *vav3* is a new transcriptional target of the AhR. (A) A 2005-base pair fragment of upstream genomic sequence containing the *vav3* mouse promoter was cloned from the W11-2862P14 phosmid into the pGL2-Basic vector. The resulting construct was transfected in wild-type Hepa 1c1c7 cells or in the AhR-deficient c2 line (expressing <10% wild-type levels of AhR) and firefly luciferase activity measured and normalized by *Renilla* luciferase activity. (B) Four XRE consensus elements for AhR binding were located in the *vav3* promoter. AhR binding to these XREs was determined by ChIP in T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells. The location of the oligonucleotides used to amplify by PCR the regions of the *vav3* promoter containing XRE sequences is indicated. Positive controls were performed using the input fractions whereas negative controls were made in the absence of antibody. (C) Site-directed mutagenesis was used to inactivate the XRE1 site in the *vav3* promoter. The mutated form of the promoter was cloned into the pGL2-Basic vector. Mutant *vav3* promoter activity was analyzed by luciferase assays as indicated above.

and the resulting construct transfected in wild-type Hepa 1c1c7 or in AhR-deficient c2 cells. Mutant *vav3* promoter activity was analyzed by luciferase assays as indicated above.

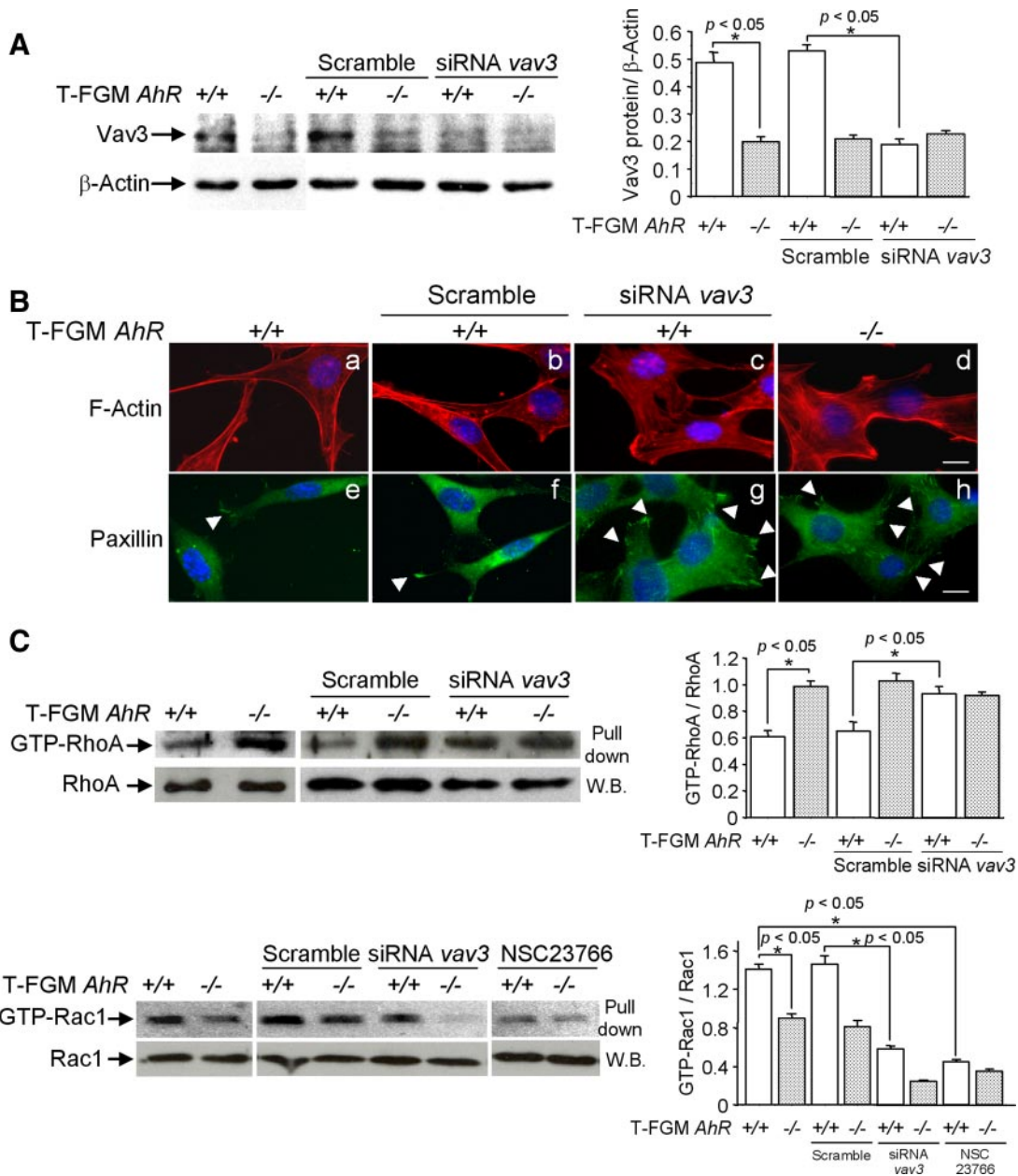


Figure 10. The knockdown of the *vav3* mRNA phenocopies the cytoskeleton-related changes observed in T-FGM *AhR*^{-/-} cells. (A) Vav3 protein expression was determined in T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells by Western blot with an anti-Vav3 antibody. In parallel experiments, *vav3* siRNAs or unspecific scramble RNAs were transfected in T-FGM cells of both genotypes and Vav3 protein expression determined as described above. The expression of β -actin was used as control. Vav3 protein levels were normalized by β -Actin for each experimental condition and the results obtained represented on the right panel. (B) T-FGM *AhR*^{+/+} cells were transfected with *vav3* siRNAs or with unspecific scramble RNAs and, 72 h later, stained with rhodamine-phalloidin (Rho-Pha) to visualize the F-actin cytoskeleton (a-c) or used in anti-paxillin immunofluorescence experiments to analyze their pattern of focal adhesions (e-g). T-FGM *AhR*^{-/-} cells were also stained under the same conditions for comparison (d and h). Actin stress fibers and focal adhesions are indicated by arrowheads in c and d and g and h, respectively. Bar, 5 μ m. (C) T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells were grown and cell extracts prepared and used in pull-down assays for RhoA and Rac1 by using the rhotekin-GST and the PAK-CRIB-GST fusion proteins as targets, respectively. T-FGM cells of both genotypes were also transfected with *vav* siRNAs or with unspecific scramble siRNAs as described above and used in pull-down assays as described in B. In additional experiments, T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells were treated with NSC23766 as indicated in the legend for Figure 7, and its effect on the levels of Rac1 activation was determined by pull-down assays as described above. Protein levels for RhoA and Rac1 were also determined by Western blotting (W.B.) in the same cell extracts. Levels of GTP-RhoA and GTP-Rac1 were normalized by total RhoA and Rac1 protein, and the values are plotted (right). Data are shown as mean \pm SEM from experiments performed in two different cultures of each genotype. The p values for statistical comparison between genotypes under the different experimental conditions are indicated.

active version of Vav3 (Δ 1-184 deletion mutant). We found that Vav3 overexpression produced the same morphological

alteration in T-FGM fibroblasts regardless on their AhR status (Supplemental Figure S5), suggesting that conditions

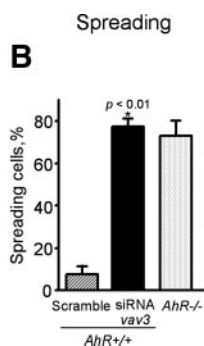
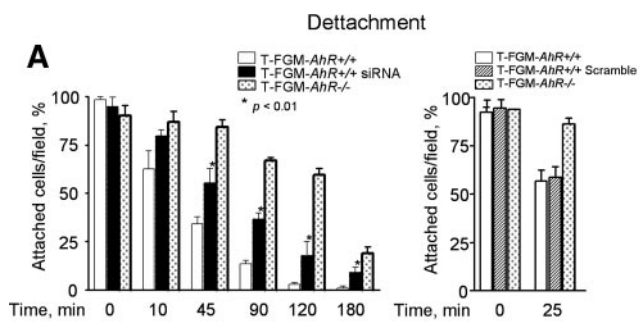


Figure 11. Vav3 down-regulation increases the spreading and attachment efficiency of T-FGM *AhR*^{+/+} cells. (A) *vav3* siRNA- or un-specific scramble siRNAs-transfected T-FGM *AhR*^{+/+} fibroblasts were used in EGTA-induced detachment experiments. At the indicated times, nuclei were stained and counted as indicated in Figure 6. The p values for statistical comparison between *vav3* siRNA-transfected T-FGM *AhR*^{+/+} and untransfected wild-type cells are indicated. (B) T-FGM *AhR*^{+/+} cells were transfected with *vav3* siRNAs and their spreading efficiency determined after crystal violet staining. Between 250 and 300 T-FGM cells were counted in six independent fields for each

experimental condition. The p values for statistical comparison in the increase of cell spreading with time in T-FGM *AhR*^{+/+} cells are indicated. Data are shown as mean ± SEM from duplicate experiments in two different cultures.

in which Vav3 activity largely exceeds its constitutive cellular levels, could abolish the physiological control of cell morphology making the process AhR independent. We used an additional experimental approach to further support that overactivation of Vav3 signaling alters cell morphology independently of AhR. Because Vav3 overactivation should increase the activity of its downstream client protein Rac1, transfection of a constitutively active form of Rac1 should also induce a phenotypic change regardless of AhR expression. T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells were transfected with a wild-type or a constitutively active form of Rac1 and their effects on cell morphology analyzed (Supplemental Figure S6). Transfection of wild-type Rac1 did not significantly change the basal morphology of T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells (Supplemental Figure S6A). However, the constitutively active version of Rac1 (V12Rac1; del Pozo *et al.*, 1999) induced a change in cell area and polarity that was of similar magnitude in both T-FGM cell lines (Supplemental Figure S6B). Thus, overactivation of Vav3-dependent signaling overrides the influence of AhR in the control of fibroblast cell phenotype. Together, these data support a mechanism in which highly regulated control of constitutive Vav3 expression by the transcriptional activity of AhR has an important role in modulating fibroblast shape and cytoskeleton under physiological conditions.

To investigate whether the reduced Vav3 expression also participated in the enhanced adhesion properties of T-FGM *AhR*^{-/-} cells, we measured the effect of the *vav3* knockdown in the detachment and spreading properties of wild-type T-FGM cells. As shown in Figure 11A, the *vav3* knockdown increased the attachment efficiency of T-FGM *AhR*^{+/+} in the presence of EGTA to values significantly higher than those observed in untransfected wild-type cells. Transfection of un-specific scramble siRNA, on the contrary, did not affect the attachment efficiency of T-FGM *AhR*^{+/+} cells (Figure 11A, right). Moreover, *vav3* siRNA-transfected T-FGM *AhR*^{+/+} cells did gain ability to spread on plastic, reaching values close to those found in T-FGM *AhR*^{-/-} cells 60 min after plating (Figure 11B). Together, these data indicate that Vav3 is a bonafide AhR target implicated in the cytoskeleton-dependent effects induced by this transcriptional factor.

Vav3 Null MEFs Had Increased Cell Area and Enhanced Actin Stress Fibers Content

To provide even further support to our hypothesis that Vav3 has an important contribution to the AhR-dependent phenotype in fibroblast cells, *vav3*^{-/-} MEFs were isolated and their morphology and ability to generate actin stress fibers

was analyzed. Cell area and minor/major axis ratio were determined as indicated in Figure 1. *vav3*^{-/-} MEFs had a significant increase in cell area without a change in minor/major axis ratio (Figure 12A), a phenotype closely mimicking that found in *AhR*^{-/-} MEFs (see Figure 1). F-Actin staining with rhodamine-labeled phalloidin was used to analyze actin stress fibers content. Loss of Vav3 expression produced an increase in stress fibers with respect to *vav3*^{+/+} MEFs (Figure 1, A and B, b) that resembled the effects

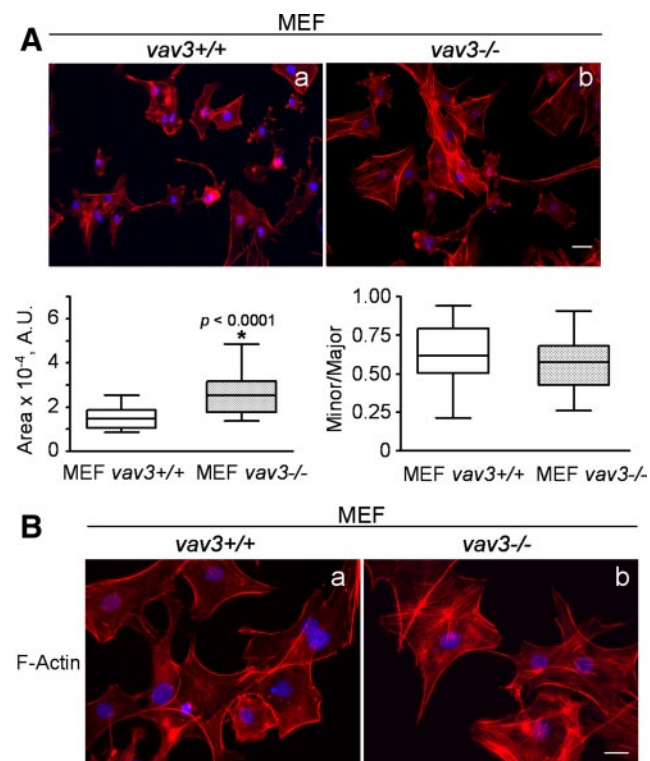


Figure 12. *vav3*^{-/-} MEFs have increased cell area and enhanced actin stress fibers content. (A) MEF cells were isolated from *vav3*^{+/+} and *vav3*^{-/-} mice and analyzed for cell size and polarity (minor/major axis ratio) as indicated in the legend for Figure 1. (B) The F-actin stress fibers were revealed by rhodamine-labeled phalloidin as indicated in the legend for Figure 3. At least 20 cells were analyzed for each experimental condition, and the experiments were done in two different MEF cultures. The p values for statistical comparison between genotypes are indicated. Bar, 10 μm (A) and 5 μm (B). A.U., arbitrary units.

described in AhR-null MEF cells (Figures 3, 7, and 10). The fact that genetically modified *vav3*^{-/-} MEFs share common phenotypes regarding cell morphology and actin stress fibers with *AhR*^{-/-} MEFs, strongly support our premise that Vav3 is a downstream target of AhR and that both proteins have a role in determining the phenotype of fibroblastic cells.

DISCUSSION

AhR is a critical intermediate in xenobiotic-induced toxicity and carcinogenesis (Fernandez-Salguero *et al.*, 1996; Peters *et al.*, 1999; Shimizu *et al.*, 2000) and many studies have analyzed its role in cell cycle, differentiation, apoptosis, and genome integrity (Lahvis *et al.*, 2005; Barouki *et al.*, 2007; Furness *et al.*, 2007; McMillan and Bradfield, 2007). Despite these known functions, emerging evidence indicates that AhR may be also involved in cytoskeleton-related processes. Thus, it has been shown that cell detachment leads to the transcriptional activation of AhR in several adherent cells such as keratinocytes (Sadek and Allen-Hoffmann, 1994a), hepatoma Hepa I cells (Sadek and Allen-Hoffmann, 1994b), and C3H10T1/2 fibroblasts (Cho *et al.*, 2004). Moreover, it has been observed that AhR has a predominantly nuclear distribution in cells located at the migrating edge in wound healing experiments (Ikuta and Kawajiri, 2006). It was also reported that both the ligand-dependent and the constitutive activation of AhR in the immune system leads to enhanced migration of CD4⁻CD8⁻ double-negative lymphocytes from the thymus (Esser *et al.*, 2004; Temchura *et al.*, 2005). Finally, a recent work has shown that either dioxin or a constitutively active form of the AhR induce morphological changes that affect the F-actin cytoskeleton, cell-cell interactions, and the motility of breast tumor cells (Diry *et al.*, 2006). Despite this knowledge, the contribution of AhR to cell morphology, adhesion, and migration under physiological, xenobiotic-free conditions, remains largely unknown.

Consistent with a link between AhR and cytoskeleton-related processes, we report here that AhR status affects cell shape, F-actin cytoskeleton, adhesion, and spreading of murine fibroblasts. A first indication of the mechanism through which AhR can modulate cell morphology and adhesion comes from our results with the constitutively active AhR protein. The expression of constitutively active AhR in *AhR*^{-/-} fibroblasts rescues the morphological phenotype of these cells, suggesting that the process requires the transcriptional activity of the receptor. The spread morphology of *AhR*^{-/-} cells was associated to prominent actin stress fibers and to increased numbers of focal adhesions with a depolarized distribution around the cell area. Focal adhesions are major structures for cell-substratum interaction (Sieg *et al.*, 1999; Schaller, 2001) and, indeed, spreading and attachment were significantly enhanced in AhR-null with respect to wild-type cells. These are relevant data that could help interpreting the lower migration rates observed in T-FGM *AhR*^{-/-} fibroblasts (Mulero-Navarro *et al.*, 2005) and MEF *AhR*^{-/-} cells (Carvajal-Gonzalez, unpublished data).

Rac1 and RhoA play crucial roles in cell migration, polarity, adhesion, and cytoskeleton reorganization (Nobes and Hall, 1999; Bishop and Hall, 2000). The importance of these GTPases in determining AhR-dependent morphology was confirmed by two facts: 1) more spread and highly attached T-FGM *AhR*^{-/-} cells had higher RhoA and lower Rac1 activation than wild-type fibroblasts and 2) the Rac1 inhibitor NSC23766 changed the morphology of *AhR*^{+/+} fibroblasts to that of *AhR*^{-/-} cells, whereas the RhoA kinase

inhibitor Y27632 did the opposite and transformed AhR-null cells into the wild-type phenotype. Moreover, these results are in close agreement with previous data indicating that Rac1 and RhoA have antagonistic effects in different cell types (Leeuwen *et al.*, 1997; Sander *et al.*, 1999). Interestingly, however, protein expression for Rac1 and RhoA was similar between T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells, suggesting that AhR regulated these GTPases at the level of their activation and/or downstream elements. To find possible targets, we carried out expression microarrays in T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells and searched for GTPases and/or GTPase regulators. This analysis indicated that the phosphorylation-dependent exchange factors Vav3 and Net 1 (Movilla and Bustelo, 1999; Hornstein *et al.*, 2004) were significantly down-regulated in *AhR*^{-/-} cells. We focused on Vav3 because Net1 protein levels did not significantly differ between T-FGM *AhR*^{+/+} and T-FGM *AhR*^{-/-} cells. Cloning of the murine *vav3* promoter revealed that AhR expression increases *vav3* promoter activity and that AhR is recruited to a proximal XRE element in the *vav3* promoter. Interestingly, the high-affinity AhR ligand TCDD did not increase the AhR-dependent *vav3* expression, suggesting not only that AhR is committed to regulate constitutive gene transcription of *vav3* but also that the physiological levels of Vav3 could be relevant to mediate the effects of AhR on cell phenotype.

The constitutive transcription of Vav3 has been related to the reorganization of actin stress fibers and to the formation of lamellipodia and membrane ruffles (Movilla and Bustelo, 1999; Bustelo, 2000) and, as such, represents a candidate target gene whose regulation by AhR could modulate cell morphology, adhesion, and spreading. We hypothesized that the down-regulation of *vav3* mRNA and protein in fibroblasts could account for the cytoskeleton-dependent changes observed in AhR-deficient cells. Indeed, siRNA-based experiments, which down-regulate the physiological levels of target genes, demonstrated that this exchange factor represents an important intermediary in the AhR-dependent signaling pathways determining cell morphology. Thus, we observed that down-regulation of Vav3 in T-FGM *AhR*^{+/+} cells induced a T-FGM *AhR*^{-/-}-like phenotype regarding cell shape, cytoskeleton organization, polarity, adhesion, and spreading that was consistent with the changes observed in the activation of the Vav3 downstream GTPases Rac1 and RhoA. The fact that overexpression of the wild-type or a constitutively active Vav3 or a constitutively active form of Rac1 changed cell morphology independently on AhR expression, further support the hypothesis that the physiological level of Vav3 is a highly regulated parameter in the AhR-dependent modulation of cellular morphology. Indeed, *vav3*^{-/-} MEFs mimicked the *AhR*^{-/-} phenotype with respect to increased cell area and enhanced actin stress fibers content, strongly indicating not only that Vav3 is a downstream target of AhR but also that constitutive Vav3 levels are required to maintain fibroblast morphology and cytoskeleton. Therefore, the AhR-dependent phenotype for cell morphology, adhesion, and spreading is controlled, at least in part, by the precise regulation of the physiological levels of the exchange factor Vav3. This is a relevant example of how AhR can modulate physiologically defined cellular properties through the transcriptional regulation of an endogenous target gene that controls migration-related signaling pathways. Although these data attribute a relevant role for the transcriptional activity of AhR in determining cell phenotype, additional, transcription-independent mechanisms, cannot yet be excluded. For example, as a cullin 4B ubiquitin ligase (Ohtake *et al.*, 2007), the liganded AhR could

regulate the levels of migration-related proteins through the proteasome by a mechanism similar to that demonstrated previously for sex steroid receptors (Ohtake *et al.*, 2007).

It is worth noting that our data do not exclude the participation of other signaling molecules in the AhR-mediated cytoskeleton effects in fibroblasts. For example, scaffolding proteins such as vinculin or other GEFs and GAPs are also dependent on the presence of AhR, at least in the case of fibroblasts. It is therefore possible that these additional proteins could cooperate with, or participate in parallel pathways, to those already engaged by Vav3. Further work in this area will address this possibility and unveil the level of cross-talk and cooperativity between these pathways. Given the important role of Rho/Rac pathways in pathologies such as cancer, immunodeficiencies or cardiovascular disease, it will be also interesting to see whether AhR may contribute to the regulation of those pathways in the context of these pathological states.

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