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Hexokinase 2 is a transcriptional target and a positive modulator of AHR signalling

Manon Watzky¹, Solène Huard¹, Ludmila Juricek², Julien Dairou^{3,4}, Caroline Chauvet^{2,3}, Xavier Coumoul^{2,3}, Anne Letessier^{1,*} and Benoit Miotto^{1,*}

¹Université Paris Cité, Institut Cochin, INSERM, U1016, CNRS, UMR8104, F-75014 Paris, France, ²METATOX, T3S, Toxicologie Environnementale, Cibles thérapeutiques, Signalisation cellulaire et Biomarqueurs, INSERM UMR-S1124, F-75006 Paris, France, ³Université Paris Cité, UFR des Sciences Fondamentales et Biomédicales, Paris, France and ⁴Laboratoire de Chimie et Biochimie Pharmacologiques et Toxicologiques, CNRS, UMR 8601, Université Paris Cité, F-75006 Paris, France

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ABSTRACT

The aryl hydrocarbon receptor (AHR) regulates the expression of numerous genes in response to activation by agonists including xenobiotics. Although it is well appreciated that environmental signals and cell intrinsic features may modulate this transcriptional response, how it is mechanistically achieved remains poorly understood. We show that hexokinase 2 (HK2) a metabolic enzyme fuelling cancer cell growth, is a transcriptional target of AHR as well as a modulator of its activity. Expression of *HK2* is positively regulated by AHR upon exposure to agonists both in human cells and in mice lung tissues. Conversely, over-expression of *HK2* regulates the abundance of many proteins involved in the regulation of AHR signalling and these changes are linked with altered *AHR* expression levels and transcriptional activity. *HK2* expression also shows a negative correlation with *AHR* promoter methylation in tumours, and these tumours with high *HK2* expression and low *AHR* methylation are associated with a worse overall survival in patients. In sum, our study provides novel insights into how AHR signalling is regulated which may help our understanding of the context-specific effects of this pathway and may have implications in cancer.

INTRODUCTION

The aryl hydrocarbon receptor (AHR) is a ligand-activated transcription factor belonging to the basic helix-loop-helix/Per-ARNT-SIM (bHLH-PAS) family (1,2). According to classical models, AHR is retained by a chaperone

complex in the cytoplasm and upon binding of its ligand, AHR translocates to the nucleus to form a heterodimer with its most common partner, the AHR nuclear translocator (ARNT, also known as hypoxia-inducible factor 1 β – HIF1 β) (1–3). This new complex then regulates the expression of different set of genes containing consensus AHR-responsive elements in their promoter such as prototypical target genes coding proteins of the cytochrome P-450 family of monooxygenases (e.g. CYP1A1 and CYP1B1) and aldehyde dehydrogenases (e.g. ALDH3A1) (3–7). AHR plays a critical function in the activation of genes involved in the metabolism of xenobiotics and in additional processes including cell proliferation, cell differentiation, immune response and in cancer processes (8–10). AHR is also emerging as a promising therapeutic target in a number of human diseases, including cancer and immune diseases (11–16). It is thus important to better understand how AHR signalling is regulated by environmental signals and intrinsic features to control such diverse functions and transcriptional programs.

A first level of regulation is the activity of *AHR* promoter itself, and hence *AHR* expression. *AHR* is expressed in most tissues, although levels are quite variable across tissues and cell types, with high levels in epithelia (e.g. lung, intestine) and immune cells (17). Several studies have shown that *AHR* promoter is regulated by epigenetic mechanisms including DNA methylation and chromatin marks (18–21). For instance, in human cells *AHR* promoter is susceptible to DNA methylation and heavy DNA methylation of the promoter is correlated with a lower expression of the gene (19).

AHR transcriptional activity is also modulated by a wide variety of exogenous and endogenous ligands. This includes halogenated aromatic hydrocarbon com-

*To whom correspondence should be addressed. Tel: +33 1 44 41 24 35; Fax: +33 1 40 51 64 73; Email: benoit.miotto@inserm.fr
Correspondence may also be addressed to Anne Letessier. Tel: +33 1 44 41 24 35; Fax: +33 1 40 51 64 73; Email: anne.letessier@inserm.fr
Present addresses:

Solène Huard, Institut Curie – PSL Research University, Translational Research Department, Breast Cancer Biology Group, 75005 Paris, France.
Ludmila Juricek, Asfalia Biologics, 18 rue Charcot, 75013 Paris, France.

pounds (HAHs) (e.g. 2,3,7,8-tetrachlorodibenzodioxin, also known as TCDD or dioxin), polycyclic aromatic hydrocarbons (PAHs) (e.g. benzo(a)pyrene (BaP) and 3-methylcholanthrene (3-MC)), and tryptophan derivatives (e.g. formylindolo[3,2-*b*]carbazole (FICZ) and kynurenic acid) (11,12). These ligands exhibit quite different impacts on AHR activity due to their chemical properties and variable affinity for AHR (11,12). For instance, TCDD causes long lasting effects as it persists in tissues following exposure, while FICZ only transiently enhances AHR transcriptional activity due to its rapid metabolism and inactivation in cells (7,12,22).

Finally, AHR activity is regulated by numerous cofactors including proteins that promote its activity such as transcriptional coactivators (e.g. EP300, NCOA3, SRC) (1). On the contrary AHR activity is negatively regulated by repressors such as AHR repressor (AHRR) and TCDD-inducible poly(ADP-ribose) polymerase (TIPARP) (1,23–25). These latter two, in addition to repress AHR signalling, are also induced at the transcriptional level by AHR upon agonists exposure, establishing negative feedback loops that attenuate transcriptional activation of AHR targets (23–25).

AHR is thus regulated at several levels. In this prospect, it is important to characterize additional proteins involved in the regulation of AHR signalling to better understand how it leads to different transcriptional outcomes and target these key regulatory links in disease. In this study, we report a molecular link between hexokinase 2 (HK2) and AHR signalling pathway. HK2 is one of the five hexokinases encoded in the human genome which phosphorylates glucose to produce glucose-6-phosphate, the first rate-limiting step of glycolysis (26). Its over-expression is extensively described to promote metabolic alterations supporting tumour growth and aggressiveness, and its knock-out markedly reduces the growth of genetically-induced tumours in mouse models of lung and breast cancers (26–28). We showed that AHR binds to *HK2* gene in human cancer cells and promotes its expression in response to exposure to a variety of AHR agonists in different cell types. We further demonstrated that over-expression of *HK2* alters the abundance of numerous proteins involved in the AHR network, including regulatory proteins and targets of AHR, and that HK2 also regulates *AHR* gene expression. These regulations are linked with gene-specific changes in AHR transcriptional outcomes. Thus, *HK2* is a target as well as a regulator of AHR signalling, and we showed that the expression of both genes is positively correlated in diverse tumour types suggesting that this regulatory link might play an important function in cancer.

MATERIALS AND METHODS

ChIP-sequencing analyses of AHR binding sites in human cells

AHR ChIP-sequencing data were retrieved from NCBI GEO and fastq files downloaded from EMBL-EBI ENA website (<https://www.ebi.ac.uk/ena>) on the Galaxy web-interface (29). We used Bowtie2 to map sequencing reads

against human reference genome hg19 and produce bam files and ChIP-sequencing peaks (30).

ChIP-sequencing datasets were previously published and prepared as follows. AHR ChIPs in MCF-7 breast cancer cells were performed using an AHR antibody (Santa Cruz, H-211) in triplicate, starting from MCF-7 treated with 10 nM TCDD (2,3,7,8-tetrachlorodibenzodioxin) or DMSO (dimethylsulfoxide) for 45 min and 24 h (GSE90550) (5). AHR ChIPs in GM17212 EBV-immortalised lymphocytes (GSE116638) were performed using an AHR antibody (Cell Signaling Technology (CST); D5S6H) in duplicate, starting from GM17212 cells treated with 1 μ M 3-MC (3-methylcholanthrene) or DMSO for 24 h (6). AHR ChIPs in HepG2 hepatocarcinoma cells were produced using an antibody directed against Flag-Tag (Sigma-Aldrich) from cells engineered to express an AHR flag-tagged protein (GSE127649; ENCSR412ZDC, released on 10 February 2018) (31).

Gene classification and gene-ontology (GO) enrichment were performed using the PANTHER interface (version 16.0 released 2020–12-01) using Fisher's exact *t*-test and false discovery rate (32).

The UCSC Genome Browser and Integrative Genome Viewer (IGV_2.3.92) were used to visualize the ChIP-sequencing data as well as to prepare the different panels in the manuscript (33). The UCSC CpG Islands, GeneHancer and RefSeq gene tracks were used to locate promoters and enhancers as well as genes in the human genome (reference hg19) (33). ENCODE datasets, including transcription factors ChIP-sequencing data and DNaseI sites, are freely usable and were retrieved from the ENCODE download portal (31). Histone ChIP-sequencing data are freely usable and were retrieved from the ChIP-Atlas database (<https://chip-atlas.org/>) (34).

A search for classical recognition motif of the AHR/ARNT complex (5'-GCGTG-3'; 3'-CACGC-5' and its longer version 5'-TNGCGTG-3'; 3'-CACGCNA-5') was performed manually on the sequence of *HK2* gene (coordinates: chromosome 2:75059782–75120481 on human reference genome hg19).

Cell lines, culture and treatments

U2OS human osteosarcoma and HCT116 human colon cancer cells were authenticated at the start of the project by DNA/STR profiling (Eurofins Genomics). 143B osteosarcoma cell line was a gift from Dr Olivia Fromig   (INSERM UMR981, Institut Gustave Roussy, Villejuif, France). All cell lines were routinely tested for mycoplasma contamination by conventional PCR using the mycoplasma detection kit Venor[®] GeM Classic (Minerva Biolabs). Authentication certificates and mycoplasma-test results are available upon request. 143B, U2OS and HCT116 cell lines encode a wild-type AHR protein (35).

U2OS, 143B and HCT116 were cultured at 37°C in a humid chamber with 5% CO₂. U2OS and 143B cells were cultured in DMEM (GlutaMAX, glucose 4.5 g/l and pyruvate) supplemented with 10% foetal bovine serum and 1% penicillin-streptomycin. HCT116 cells were cul-

tured in RPMI-1640 with 10% foetal bovine serum and 1% penicillin-streptomycin. Cells were split every 2 to 3 days.

U2OS-GFP, U2OS-GFP-HK2, 143B-GFP and 143B-GFP-HK2 were maintained in DMEM (GlutaMAX, glucose 4.5 g/l and pyruvate) supplemented with 10% foetal bovine serum, 1% penicillin-streptomycin and experimentally-defined concentrations of neomycin/G418 (Thermo Fisher Scientific; 10131027) as a selection agent (see section 'Plasmids' below).

Cells were seeded at a density of 7800 cells/cm² in six-well plates and after cell attachment to the culture plates, AHR agonists were added to the culture media. TCDD solution in toluene (Supelco; 48599) was added at final concentration of 1 or 10 nM, and toluene (Sigma-Aldrich; 244511) was used as control vehicle treatment. BaP solution (Supelco; 40071) was diluted in DMSO and used at final concentration of 0.5–2 μM. FICZ (Sigma-Aldrich; SML1489) was resuspended at 1 mM in DMSO and used at final concentration of 1 nM to 10 μM. CH-223191 (Sigma-Aldrich; C8124) resuspended in DMSO was used at final concentrations of 0.001–10 μM and added alone or at the same time as TCDD or BaP for 48 h. 5-aza-2'-deoxycytidine (A3656; Sigma-Aldrich) diluted in DMSO was used at final concentration of 10 μM for 48 h and cell culture media was changed every 24 h as previously described (36). DMSO was used as a control for FICZ, BaP, CH-223191 and 5-aza-2'-deoxycytidine exposure experiments. 2-Deoxy-D-glucose (Sigma-Aldrich; D6134) was resuspended in water and used at final concentrations of 2 mM alone or in combination with BaP for 48 h.

ChIP-qPCR analysis of AHR binding

The chromatin immunoprecipitation (ChIP) was performed using a human AHR specific antibody (Cell Signaling Technology; 13790) following a procedure previously described (37). qPCR analysis was performed with the kit LightCycler[®] 480 SYBR Green I Master (Roche Diagnostics; 04887352001) according to manufacturer recommendations on a Roche LightCycler[®] 480 system (software version 1.5.1.62 SP3). ChIP and input DNAs were amplified and data are presented as percentage of input DNA recovered in each ChIPs.

qPCR primers were as follows: *CYP1A1* promoter 5'-CCT GGG ATC ACA AGG ATC AGG-3' and 5'-CGT ACA AGC CCG CCT ATA AA-3', *HK2* cis-regulatory region A 5'-CCA CTA CCA GGG AAG GCT CA-3' and 5'-TCC TGC CCA GTG ACT AGA GG-3', *HK2* cis-regulatory region B 5'-CAG GGA GCT GGT CAG ATG TG-3' and 5'-AGT GAA GCG GAA TGG GTC AG-3', *HK2* cis-regulatory region C 5'-GAG GTA GTC GGC TCT CAG GA-3' and 5'-TCC AGG TTG CTA CGA ATG CC-3', *HK2* cis-regulatory region D 5'-CAT GCT GGG GTT GGA GAA GG-3' and 5'-TTG GTG CAG GCA TAG GAG TG-3', *HK2* promoter 5'-CAA CAT CGT GTC ACC CAG CT-3' and 5'-GCT AAC TTC GGC CAC AGG AT-3', region 2-kilobases upstream of *HK2* promoter 5'-CCC GGC ATC CCT TGA ATT CT-3' and 5'-TCC AGG CCT GTC TCC AAC TC-3', region 2-kilobases downstream of *HK2* promoter 5'-ATG TAG TGA TGG CGC

GTG AA-3' and 5'-CAG AGC CAC ATC CCA GGA ATT-3'.

RT-qPCR analysis

RNAs were isolated using the TRIzol[™] reagent (Thermo Fisher Scientific; 15596026) followed by DNase treatment and removal (Thermo Fisher Scientific; AM1906) according to a standard protocol previously described (36) or the RNeasy mini-kit according to manufacturer recommendations (QIAGEN; 74106). RNA concentration and purity were assessed using NanoDrop[™] 2000 (Thermo Fisher Scientific). Reverse transcription was performed using the Superscript[™] II reverse transcriptase enzyme with reaction conditions recommended by the manufacturer (Thermo Fisher Scientific; 18064) starting with 500 ng of total RNA preparation and using random hexamers (Thermo Fisher Scientific; N8080127). qPCR analysis was performed using the kit LightCycler[®] 480 SYBR Green I Master (Roche Diagnostics; 04887352001) according to manufacturer recommendations on a Roche LightCycler[®] 480 system (software version 1.5.1.62 SP3).

Primers used for qPCR analyses were as follows for human genes: *HK2* 5'-GAG CCA CCA CTC ACC CTA CT-3' and 5'-CCA GGC ATT CGG CAA TGT G-3', *AHR* 5'-ACA TCA CCT ACG CCA GTC G-3' and 5'-CGC TTG GAA GGA TTT GAC TTG A-3' (used for Figure 2), *AHR* 5'-GCC GGT GCA GAA AAC AGT AAA-3' and 5'-AGC CAA ACG GTC CAA CTC TG-3' (used for Figure 4), *CYP1A1* 5'-TCT TGA GGC CCT GAT TAC CCA-3' and 5'-TTC GGC CAC GGA GTT TCT TC-3', *CYP1B1* 5'-AAC GTA CCG GCC ACT ATC AC-3' and 5'-GCA CTC GAG TCT GCA CAT CA-3', *ALDH3A1* 5'-TGA TCC AGG AGC AGG AGC A-3' and 5'-CCT CTA GGA CGT ACA CCA CC-3'. Gene expression values were normalised using the DeltaCq method ($2^{-\Delta\Delta Cq}$) against reference *MCM3* gene expression in Figures 4A–F, 5A, B, F, G and Supplementary Figure S5 (5'-GCT CCT CTG GAG TGG GTC TG -3' and 5'-TCC TGT TTC CTG GTC TGT GGT-3') and in all other figures against the average value of expression of two reference genes: *B2M* 5'-GGC TAT CCA CGT ACT CCA AA-3' and 5'-CGG CAG GCA TAC TCA TCT TTT T-3' and *MAPK14* 5'-TGC CGA AGA TGA ACT TTG CGA-3' and 5'-TCA TAG GTC AGG CTT TTC CAC T-3'. Cq is the quantification cycle corresponding to the Cp value on Roche LightCycler[®] 480 system.

For *Hk2* murine transcript we used the following primers: 5'-ATG ATC GCC TGC TTA TTC ACG-3' and 5'-CGC CTA GAA ATC TCC AGA AGG G-3'. We normalised *Hk2* expression using the DeltaCq ($2^{-\Delta\Delta Cq}$) method and the average value of expression of two reference genes: *Actb* 5'-CAA TAG TGA TGA CCT GGC CGT-3' and 5'-AGA GGG AAA TCG TGC GTG AC-3' and *Mapk14* 5'-TGA CCC TTA TGA CCA GTC CTT T-3' and 5'-GTC AGG CTC TTC CAC TCA TCT AT-3'.

Western blot analysis

Methods for protein extraction, separation and detection were previously described (36). Proteins of interest were revealed using the following antibodies: AHR (13790),

HK1 (C35C4), HK2 (C64G5), GAPDH (D16H11) and LDHA (C4B5) purchased from Cell Signaling Technology, NQO1 (A180) purchased from Novus Biologicals and CHK2 (ab109413) and Histone H3 (ab1791) purchased from Abcam. α -Tubulin (T6199-100UL), β -Actin (PA1-183) and MCM2 (A300-191A) were purchased from Sigma-Aldrich, Thermo Fisher Scientific and Bethyl Laboratories respectively. Rabbit homemade antiserum against JUN was a gift from Dr Frédérique Verdier (Institut Cochin, INSERM U1016, Paris, France).

Exposition of mouse pups to TCDD

C57BL/6J mice were provided by the CDTA (Center for animal Distribution, Typing and Archiving, CNRS, Orléans, France). Experiments on animals were performed at the animal facility of CDTA and subsequently at the animal core facility of BioMed Tech facilities (Campus Saint Germain des Prés, INSERM US36, CNRS UMS2009, Université Paris Cité, Paris, France). The European Communities Council directive 2010/63/EU on the protection of the animals were followed for the experiments using animals. Animals were treated humanely and with regard for alleviation of suffering. All procedures were given approval by the ethical committees for animal research of CNRS (Orléans, France) and 'Université Paris Cité' (Paris, France) (CEEA34.XC.049.12).

C57BL/6J mice were housed in a temperature-controlled room ($22 \pm 1^\circ\text{C}$) with a relative humidity of $55 \pm 5\%$ and a 12-h light/dark cycle. Water and food (Safe[®] D30) were provided *ad libitum*. Pregnant CD57Bl/6J mice were randomly administered either oral doses of TCDD in vehicle (1 ng TCDD/g body weight; LGC Standards) or *n*-nonane/corn oil vehicle alone (1/24 v/v; Sigma-Aldrich) on embryonic days E7.5, E14.5 and post-natal days P0.5, P7.5, P14.5, P21 in the CDTA animal facility by using a curved gavage probe fitted to the mouse mounted on a 1 ml syringe. The oral route of exposure was used to mimic the major route of exposure in humans. Each administration corresponded to 1 ng/g body weight of TCDD in *n*-nonane (corresponding to 0.2 $\mu\text{l/g}$ body weight) or *n*-nonane alone (0.2 $\mu\text{l/g}$ body weight) diluted in corn oil (10 $\mu\text{l/g}$ body weight). Pups were weaned at 3 weeks of age and allowed to acclimate to the BioMed Tech animal facility from 5 to 9 weeks of age. Five male pups from these two groups of dams were analysed at 9 weeks of age. Their lung tissues were collected and analysed by RT-qPCR and western blot.

The Comparative Toxicogenomics Database (CTD)

The Comparative Toxicogenomics Database (CTD) was interrogated on 1 September 2021 on an updated version of the database including 14 036 unique chemicals (38). Using the search engine 'chemical-gene interaction' tool, 2284 chemical interactions with AHR in human (taxon:9606) were retrieved among which 748 interactions modulate AHR activity (either 'increase', 'decrease' or 'affect with degree unspecified'). Using a similar approach, we identified 180 chemical interactions with HK2 and 158 chemical interactions regulating *HK2* expression in human. We manually curated the lists (to remove duplications) and then com-

pared them to identify the chemicals common to the regulation of *HK2* expression and AHR activity, and we assessed the resulting overlap by hypergeometric test.

Electroporation of small-interfering RNAs (siRNAs)

Transfection of control and *AHR* siRNA duplexes was performed with Neon Transfection System according to the manufacturer's instructions (Thermo Fisher Scientific). Functional *AHR* duplexes were purchased from Thermo Fisher Scientific under references: HSS100337, HSS100336 and HSS100338. Control silencing RNA was also purchased from Thermo Fisher Scientific under reference: 465377. siRNA-electroporated cells were cultured in six-well plates at 37°C under 5% CO_2 in a humid chamber for 24 h prior to treatment.

Plasmids

The following mammalian plasmids pCMV6-AC-GFP and pCMV6-AC-HK2-GFP were purchased from ORIGENE with references PS100010 and RG209482 respectively. pCMV6-AC-HK2-GFP encodes a functional full-length human HK2 protein fused to TurboGFP in its C-terminus. The plasmid backbone further encodes a G418/neomycin selection cassette.

To establish the U2OS-GFP and U2OS-GFP-HK2 cell lines, U2OS cells were transfected with pCMV6-AC-GFP and pCMV6-AC-HK2-GFP respectively, with Neon Transfection System according to the manufacturer's instructions (Thermo Fisher Scientific). Electroporated cells were exposed to a range of G418 from 0.3 to 0.6 mg/ml (Thermo Fisher Scientific, 10131027) and the appropriate concentration empirically determined. 0.6 mg/ml was deemed appropriate to maintain U2OS-GFP-HK2 cells, and 0.4 mg/ml for U2OS-GFP cells.

A similar approach was utilised to produce 143B osteosarcoma cells with stable expression of GFP and GFP-HK2.

pH measurement

The level of media acidification reached by the U2OS-GFP and U2OS-GFP-HK2 cell lines were monitored using a FiveEasy[™] pH meter (Mettler Toledo). One million cells were seeded in regular media and allowed to attach for 6 h to the plate. Then, the cell culture media were replaced with fresh media and the pH of the cells' supernatant measured 4 h later ($n = 6$).

Cell proliferation assay

Fifty thousand cells were seeded in six-well plates. Cells were then numbered 7 days later with a Countess[™] II FL automated cell counter (Thermo Fisher Scientific). Trypan blue (Amresco, K940) staining was used to discriminate living and dead cells in six independent experiments.

Proteomic analysis

Digestion

U2OS-GFP-HK2 and U2OS cell pellets (four samples per group) were lysed during 5 min at 95°C in 100 mM

Tris/HCl pH8.5, 2% SDS. Protein concentration was determined using SDS PAGE of an aliquot and Imagemlab software (Bio-Rad Laboratories). 50 µg of proteins from each lysate were reduced and alkylated with 10 mM tris(2-carboxyethyl)phosphine (TCEP) and 50 mM chloroacetamide for 5 min at 95°C. After cooling to room temperature, extracts were diluted with 300 µl 8 M urea, 50mM Tris/HCl pH 8.5, transferred onto 30 kDa centrifugal filters and prepared for FASP digestion as described previously (39). Proteins were digested overnight at 37°C with 1µg trypsin (V511A; Promega).

Peptide desalting

Peptides were desalted on C18 StageTips, manufactured by stacking six layers of C18 reverse-phase from a disk of 3M Empore Octadecyl C18 High Performance Extraction Disk into a 200 µl micropipet tip.

Peptide fractionation

Peptides were then separated in five fractions using strong cation exchange (SCX) resin (40). Briefly, peptides were loaded into pipette-tip columns made by stacking six layers of a 3M Empore cation extraction disk into a 200 µl micropipet tip. Column conditioning was performed using acetonitrile (ACN). We used 0.1% Trifluoroacetic acid (TFA) for column equilibration. Samples acidified with TFA were loaded on the column and washed with 0.1% TFA. Peptides were finally successively eluted using 20% ACN, 0.05% formic acid, ammonium acetate at 75, 125, 200, 300 mM. The 5th fraction was eluted in 1.4% NH₄OH, 80% ACN.

LC-MS/MS

After speed-vacuum drying, fractions were solubilized in 10 µl of 0.1% TFA, 10% ACN. Liquid chromatography and mass spectrometry analyses were performed on an U3000 RSLC nanoflow-HPLC system coupled to a Q-Exactive Orbitrap mass spectrometer (both from Thermo Fisher Scientific). 1 µl of each fraction were concentrated and washed on a C18 reverse-phase precolumn (3 µm particle size, 100 Å pore size, 75 µm inner diameter, 2 cm length, Thermo Fischer Scientific), then separated using a C18 reverse-phase analytical column (2 µm particle size, 100 Å pore size, 75 µm inner diameter, 25 cm length from Thermo Fischer Scientific) with a 3 h gradient starting from 99% of solvent A (0.1% formic acid) to 55% of solvent B (80% ACN and 0.085% formic acid). The mass spectrometer acquired data throughout the elution process and operated in a data-dependent scheme with full MS scans acquired, followed by up to 10 successive MS/MS HCD-fragmentations on the most abundant ions detected. Settings for Q-Exactive were: full MS automated gain control (AGC) target 1.106 with 60 ms maximum ion injection time (MIIT) and resolution of 70 000. The MS scans spanned from 350 to 1500 Th. Precursor selection window was set at 2 Th. HCD normalized collision energy (NCE) was set at 27% and MS/MS scan resolution was set at 17 500 with AGC target 1.105 within 60ms MIIT. Dynamic exclusion time was set to 30 s and spectra were recorded in profile mode.

Identification and quantification

The mass spectrometry data were analyzed using Maxquant version 1.6.2.6 (41). The database used was a concatenation of human sequences from the Uniprot-Swissprot database (Uniprot, release 2018-06) and an incremented list of contaminants. The enzyme specificity was trypsin. The precursor mass tolerance was set to 4.5 ppm and the fragment mass tolerance to 20 ppm. Carbamidomethylation of cysteines was set as constant modification and acetylation of protein N-terminus and oxidation of methionines were set as variable modifications. Second peptide search was allowed and minimal length of peptides was set at seven amino acids. False discovery rate (FDR) was kept below 1% on both peptides and proteins. Label-free protein quantification (LFQ) was done using both unique and razor peptides. At least two ratio counts were required for LFQ. All experiments were analyzed simultaneously with the 'match between runs' option with a match time window of 0.7 min and an alignment time window of 20 min.

Proteome data analysis

Using Perseus software (version 1.6.2.3) (42) false proteins discovery were filtered out, to wit proteins that match with contaminant, to the reverse database and proteins identified only with modified peptide, leading to a matrix of 7278 proteins. LFQ intensity data were transformed into log₂ and a Student's *t*-test was performed using the Benjamini-Hochberg false discovery rate to identify the differentially expressed proteins. Proteins present in at least 3/4 replicates only in U2OS or U2OS-GFP-HK2 cells were manually given the scores: $P\text{-value} = 0$ and $\log_2 \text{ratio} = 10$ (i.e. $\log_2(\text{mean LFQ intensity in U2OS}) - \log_2(\text{mean LFQ intensity in U2OS-GFP-HK2})$). Then Canonical Pathways and Upstream Regulator analyses were generated through the use of Ingenuity Pathway Analysis software IPA (QIAGEN, <https://www.qiagenbioinformatics.com/products/ingenuitypathway-analysis>) version 65367011, considering proteins with a cut-off *P*-value <0.001 and an absolute log ratio > 1.

Cytoscape visualization

The list including the differentially regulated proteins belonging to the AHR signalling pathway, the Upstream Regulators predicted in this pathway and their target proteins was submitted into Cytoscape software (v.3.8.2) for network visualisation (43). Proteins of the 'AHR signalling' were manually highlighted by bold lines. Proteins were colour coded according to their log ratio between U2OS and U2OS-GFP-HK2 cells.

Tryptophan and other metabolites quantification

Metabolites from 2 millions of thawed cells were extracted using a methanol/chloroform extraction method with minor modifications (44). Cold methanol/chloroform (2:1, v/v; 1.5 ml) was added to cell pellets and homogenized on ice. The sample tube was centrifuged at 15 000g for 10 min at 4°C, and the supernatant was transferred to a new sample tube through a 70-mm cell strainer. Ice-cold water (0.6

ml) was added, and the sample tube was vortexed and centrifuged (15 000g, 5 min, 4°C) to obtain phase separation. The upper and lower phases were separately collected in fresh sample tubes with a syringe, taking care not to disturb the interface. The polar (upper) phase (500 ml) was evaporated to dryness in a SpeedVac concentrator (Savant), and then was reconstituted in 100 µl of methanol/water (1:1, v/v). Extracted metabolites were stored at -80°C until analysis. Tryptophan (TRP) and TRP metabolites were measured via HPLC using a coulometric electrode array (ESA Coultronics, ESA Laboratories) and fluorometric detection. Separation of TRP metabolites were done by reversed-phase liquid chromatography using a 20 mM NaH₂PO₄ buffer (not pH adjusted) with 5.0% acetonitrile. The mobile phase was delivered by an HPLC pump (Shimadzu) through a Supelcosil™ LC-18 column (250 mm × 4.6 mm, 5 µm, Supelco) at a rate of 1 ml/min. Quantifications were performed by referencing calibration curves obtained with internal standards.

Cancer Cell Line Encyclopedia (CCLE) data

RNA sequencing data were downloaded from the authors' manuscript through the cBioPortal web interface (<https://www.cbioportal.org>) and retrieved in reads per kilobase per million mapped reads (RPKM) (35,45). Illumina 450K methylation data and reduced-representation bisulfite sequencing (RRBS) data of the *AHR* promoter in osteosarcoma cell lines were retrieved using the CellMiner Cross-Database (CDB) web interface (<https://discover.nci.nih.gov/cellminerfdb/>) (46). Only CpG located in the *AHR* CpG islands with coverage >5 were exported for RRBS data.

Methylated-DNA immunoprecipitation (MeDIP)

Genomic DNA were extracted and purified using a standard phenol/chloroform procedure (36). Methylated DNA immunoprecipitation (MeDIP) was performed with the MagMeDIP kit (Diagenode; C02010021) according to the supplier recommendations starting from 1.2 µg of genomic DNA sonicated to 200–400 bp using a Bioruptor® PLUS sonicator (Diagenode). qPCR analysis was performed using the kit LightCycler® 480 SYBR Green I Master (Roche Diagnostics; 04887352001) according to manufacturer recommendations on a Roche LightCycler® 480 system (software version 1.5.1.62 SP3). Quality of the MeDIP procedure was monitored using control primers for methylated DNA (*TSH2B* gene) and for unmethylated DNA (*GAPDH* promoter) provided by the MagMeDIP kit manufacturer (Diagenode). *AHR* promoter methylation was assessed using the following primers: 5'-GAC CGC CAG CTC AGA ACA G-3' and 5'-CTC CCA GCT TCC GTT CGG-3' (coordinates chr7:17338323–17338373 on hg19) and data are presented as percentage of input DNA recovered after MeDIP (i.e. DNA methylation level).

The Cancer Genome Atlas (TCGA) data

Gene expression, DNA methylation and clinical data in 33 cancer types were obtained from the TCGA and cBioPortal data portal (45,47,48). Correlations were evaluated

with Pearson coefficient values. High and low *HK2* expression groups were defined using the median expression value of *HK2* in each TCGA cancer type. High and low *AHR* expression and promoter methylation level groups were defined using the same criteria. All data used in this study were freely available, freely re-usable and patients' information anonymized. Adrenocortical carcinoma (ACC), bladder urothelial carcinoma (BLCA), breast invasive carcinoma (BRCA), cervical squamous cell carcinoma and endocervical adenocarcinoma (CESC), cholangiocarcinoma (CHOL), colon adenocarcinoma (COAD), lymphoid neoplasm diffuse large B-cell lymphoma (DLBC), esophageal carcinoma (ESCA), glioblastoma multiforme (GBM), head and neck squamous cell carcinoma (HNSC), kidney chromophobe (KICH), kidney renal clear cell carcinoma (KIRC), kidney renal papillary cell carcinoma (KIRP), acute myeloid leukemia (LAML), brain lower grade glioma (LGG), liver hepatocellular carcinoma (LIHC), lung adenocarcinoma (LUAD), lung squamous cell carcinoma (LUSC), mesothelioma (MESO), ovarian serous cystadenocarcinoma (OV), pancreatic adenocarcinoma (PAAD), pheochromocytoma and paraganglioma (PCPG), prostate adenocarcinoma (PRAD), rectum adenocarcinoma (READ), sarcoma (SARC), skin cutaneous melanoma (SKCM), stomach adenocarcinoma (STAD), testicular germ cell tumors (TGCT), thyroid carcinoma (THCA), thymoma (THYM), uterine corpus endometrial carcinoma (UCEC), uterine carcinosarcoma (UCS) and uveal melanoma (UVM).

Osteosarcoma patients' data

Gene expression values were retrieved from the National Cancer Institute (NCI) Therapeutically Applicable Research to Generate Effective Treatments (TARGET) Osteosarcoma (OS) project (accession studies phs000218 and phs000468) and from GSE21257 (49). TARGET Osteosarcoma datasets are available without restrictions on their use in publications since 2019.

Statistical analysis

All the data points as well as the different statistical tests are available in Supplementary Table S5. All error bars represent standard deviation. Student's t-tests and hypergeometric tests were performed. $P < 0.05$ was considered significant. * $P < 0.05$; ** $P < 0.01$ and *** $P < 0.001$. n.s., not significant.

RESULTS

A genomic analysis of AHR binding sites identifies Hexokinase 2, *HK2*, as a potential transcriptional target of AHR

To identify key AHR signalling targets in human cells, we retrieved AHR binding sites identified by chromatin immunoprecipitation followed by deep-sequencing (ChIP-sequencing) in three human cancer cell types. We compared the lists of AHR targets identified from human hepatocellular carcinoma cells (HepG2), from human breast cancer cells (MCF-7) exposed to 10 nM TCDD for 45 min or 24

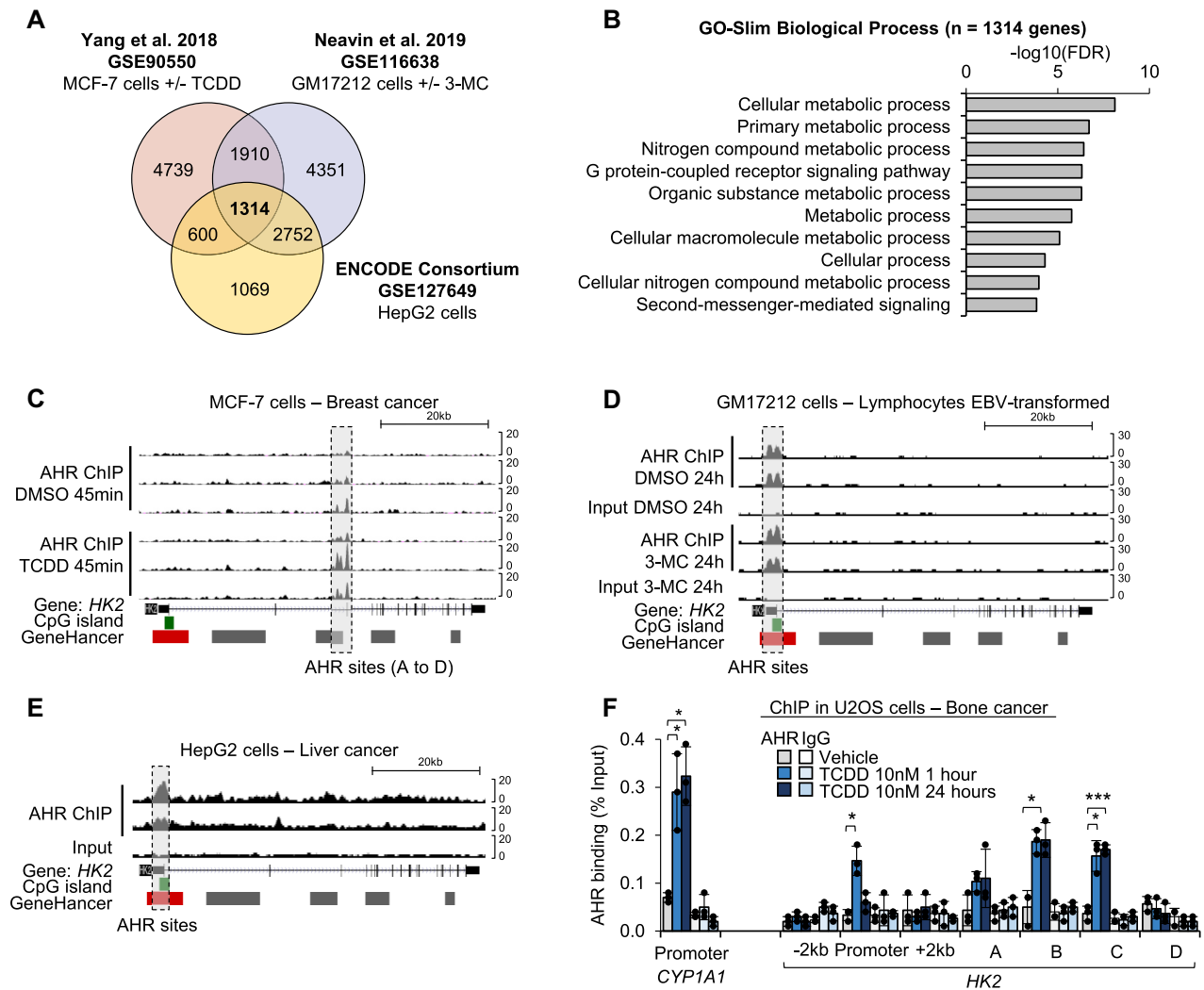


Figure 1. *HK2* is a genomic target of AHR. (A) Venn diagram showing the intersection of AHR-bound genes between AHR ChIP-seencing analyses in MCF-7 breast cancer cells (vehicle and TCDD-treated for 45 min and 24 h), in GM17212 lymphoblastoid cells exposed to vehicle and 3-MC (24 h) and in HepG2 hepatocellular carcinoma cells. Data were retrieved from publicly available datasets deposited at NCBI GEO under accession numbers GSE90550, GSE116638 and GSE127649. (B) Functional classification in biological processes of common AHR targets ($n = 1314$) using the PANTHER GO-Slim Biological Process database. (C–E) Snapshot of ChIP-seencing data showing AHR binding sites at the *HK2* locus (C) in MCF-7 cells treated with TCDD (45 min) compared to vehicle (DMSO) treatment, (D) in GM17212 cells exposed to 3-MC for 24 h or DMSO and (E) in HepG2 cells. The location of CpG island (in green), promoters (in red) as well as enhancers (in grey) from the GeneHancer database are indicated. (F) ChIP-qPCR characterisation of AHR binding sites at the *CYP1A1* promoter and the *HK2* locus in U2OS cells treated with vehicle (i.e. toluene) or 10 nM TCDD for 1 h and 24 h ($n = 3$). P -value as determined by paired Student's t -test (vehicle versus TCDD). * $P < 0.05$; *** $P < 0.001$.

h or vehicle alone, and from human EBV-immortalised lymphocytes (GM17212) treated with 3-methylcholanthrene (3-MC) at 1 μM for 24 h or vehicle alone (5,6,31). As previously described and discussed by the authors, we observed that most AHR binding sites are detectable in non-treated conditions in these cell lines, possibly due to the chronic activation of AHR by endogenous ligands (5,6). We identified 1314 AHR-bound genes common between the different ChIP-seencing datasets and experimental conditions (Figure 1A). We performed a functional annotation of this list of genes using the PANTHER GO-Slim Biological Process database (32) and observed that a large proportion of AHR target genes encode proteins involved in metabolic pathways and processes (Figure 1B and Supplementary Table S1).

We notably observed that *HK2*, a key cancer gene, is an AHR target in all experimental conditions of the three studies which led us to further explore this AHR/*HK2* axis. In MCF-7 cells, AHR binding is detectable in a cluster of *cis*-regulatory elements in the vicinity of *HK2* exon 3 (i.e. introns 2 and 3; that we called A, B, C and D *cis*-regulatory elements hereafter) and the binding is enhanced after 45 min of TCDD exposure (Figure 1C). In GM17212 and HepG2 cells, AHR ChIP-seencing profile at the *HK2* loci is different: AHR binds at the *HK2* promoter and no binding is detectable around exon 3 in all conditions (Figure 1D, E). Consistent with these ChIP-seencing data, a search for AHR DNA binding consensus sequences confirmed the presence of putative AHR-response elements in the promoter of *HK2* and in the *cis*-regulatory regions A

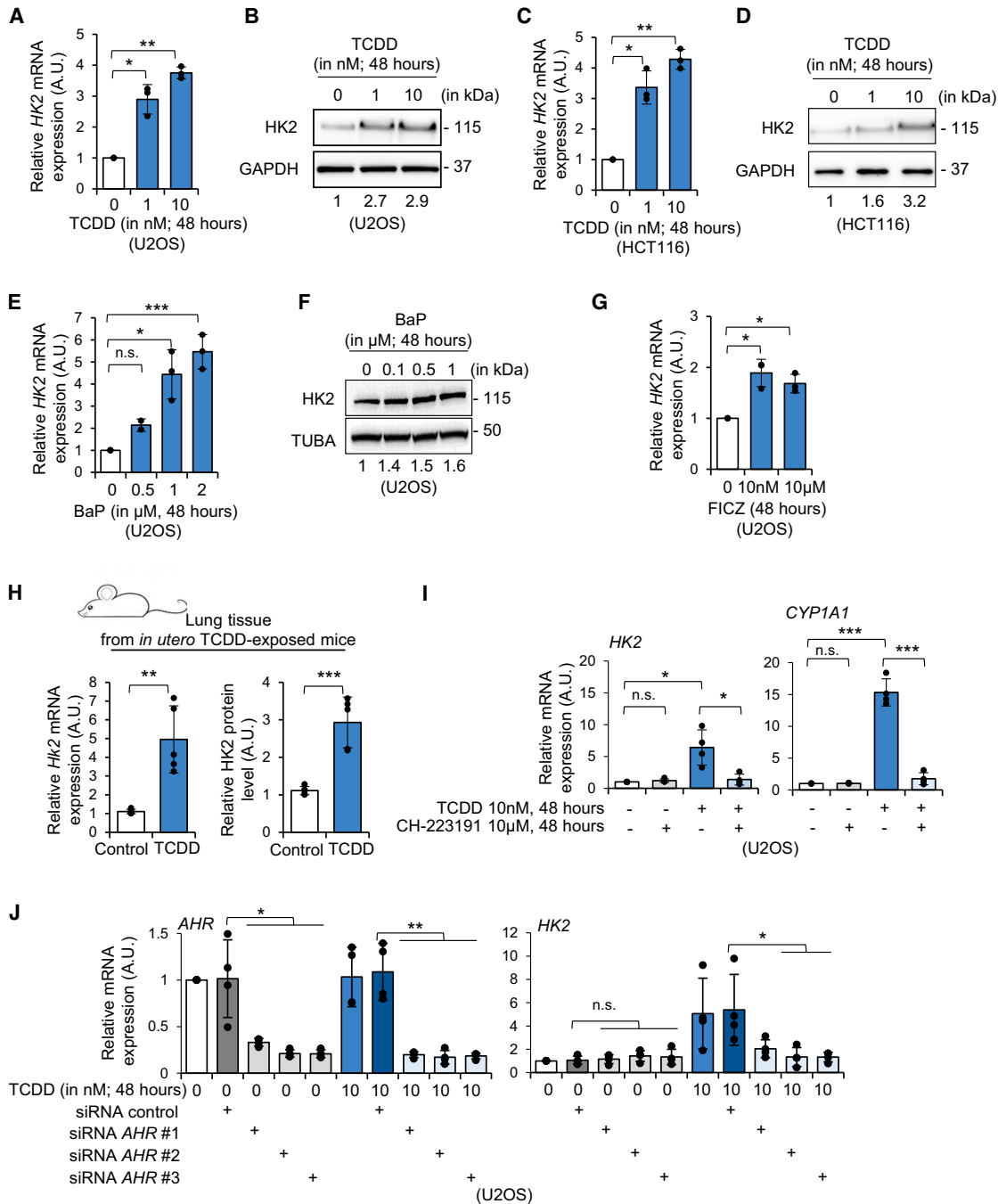


Figure 2. AHR regulates *HK2* mRNA expression. (A) RT-qPCR analysis of *HK2* expression in U2OS cells treated with vehicle (i.e. toluene) or TCDD (1 and 10 nM) for 48 h ($n = 3$). A.U., arbitrary unit. (B) Western blot analysis of *HK2* and *GAPDH* levels in U2OS cells treated with vehicle (i.e. toluene) or TCDD (1 and 10 nM) for 48 h. Relative quantifications are indicated below the blots (versus *GAPDH* control and relative to vehicle condition). (C) RT-qPCR analysis of *HK2* expression in HCT116 human colon cancer cells treated with vehicle (i.e. toluene) or TCDD (1 and 10 nM) for 48 h ($n = 3$). A.U., arbitrary unit. (D) Western blot analysis of *HK2* and *GAPDH* levels in HCT116 cells treated with vehicle (i.e. toluene) or TCDD (1 and 10 nM) for 48 h. Relative quantifications are indicated below the blots (versus *GAPDH* control and relative to vehicle condition). (E) RT-qPCR analysis of *HK2* expression in U2OS cells treated with vehicle (i.e. DMSO) or BaP (0.1 μ M up to 2 μ M) for 48 h ($n = 3$). A.U., arbitrary unit. (F) Western blot analysis of *HK2* and *TUBA* levels in U2OS cells treated with vehicle (i.e. DMSO) or BaP (0.1 μ M up to 1 μ M) for 48 h. Relative quantifications are indicated below the blots (versus *TUBA* control and relative to vehicle condition). (G) RT-qPCR analysis of *HK2* expression in U2OS cells treated with vehicle (i.e. DMSO) or FICZ (10 nM and 10 μ M) for 48 h ($n = 3$). A.U., arbitrary unit. (H) RT-qPCR (left graph) and relative quantification of western blots (right graph) of *Hk2* expression and protein levels in lung tissues from adult male mice exposed to TCDD or vehicle (i.e. n-nonane) *in utero* and during lactation ($n = 5$). A.U., arbitrary unit. (I) RT-qPCR analysis of *HK2* (left graph) and *CYP1A1* (right graph) expression in U2OS cells treated with vehicle (i.e. toluene), TCDD (10 nM), CH-223191, an AHR antagonist (10 μ M) or with TCDD + CH-223191 ($n = 4$). A.U., arbitrary unit. (J) RT-qPCR analysis of *AHR* (left graph) and *HK2* (right graph) expression in U2OS cells transfected with control or *AHR* siRNAs and further exposed to vehicle (i.e. toluene) or TCDD (10 nM) for 48 h ($n = 4$). A.U., arbitrary unit. Paired Student's *t*-test was used to calculate *P*-value in panels A, C, E, G and I, and unpaired Student's *t*-test for panels H and J. n.s., not significant; **P* < 0.05; ***P* < 0.01; ****P* < 0.001.

and D near exon 3 (Supplementary Figure S1A). In addition, profiling of ChIP- and DNaseI-sequencing data in MCF-7 indicates that *cis*-regulatory regions A–D are located in an enhancer region of *HK2* as illustrated by the presence of DNaseI hyper-sensitive sites, enrichment for enhancer-associated histone marks and binding sites for transcription factors including FOXA1 and cMYC, which are transcriptional co-factors of AHR (50,51) (Figure 1C and Supplementary Figure S1B). These observations support the hypothesis that these AHR-bound genomic sites are functional *cis*-regulatory elements of *HK2*.

To confirm that *HK2* is a *bona-fide* target of AHR, we treated U2OS human osteosarcoma cells with 10 nM TCDD or vehicle for 1 h and 24 h before assessing the binding of AHR at genomic sites nearby exon 3 (i.e. regions A, B, C and D), at the *HK2* promoter as well as at negative control regions located ± 2 kb from the promoter. As an additional control we also monitored AHR binding at the *CYP1A1* promoter, commonly assessed as a read-out of AHR transcriptional activity (3). Importantly, using histone marks ChIP-sequencing data produced in U2OS cells, we could confirm the location of *HK2* enhancers, although only AHR-binding sites A, B and C are enriched for enhancer-associated histone marks (Supplementary Figure S1C and D). In U2OS cells treated with vehicle, we did not observe significant AHR binding at the *CYP1A1* and *HK2* promoters, nor at the other regions tested (Figure 1F). By contrast, in TCDD-treated cells we observed a significant binding of AHR at the *CYP1A1* promoter 1 h and 24 h after TCDD treatment as well as a significant enrichment of AHR at the *HK2* promoter and nearby exon 3 in *cis*-regulatory region C 1 h post TCDD-treatment, and in regions B and C 24 h post TCDD-treatment (Figure 1F). No AHR ChIP-qPCR enrichment is detected at negative control sites (i.e. ± 2 kb from *HK2* promoter) monitored in our assays, nor at regions A and D, at either time points (Figure 1F). These results demonstrate that in U2OS cells, AHR binds to the *HK2* promoter and several *cis*-regulatory elements near exon 3 in response to TCDD exposure.

Collectively, our data demonstrate that the transcription factor AHR binds to *HK2* gene in different human cell types.

Agonists of AHR promote *HK2* expression in human cells and mice lung tissue

We next investigated whether *HK2* expression was regulated in response to AHR agonists exposure. To do so, we treated human U2OS cancer cells with TCDD and monitored by RT-qPCR and western blot the expression level of *HK2*. We observed that *HK2* mRNA and protein levels are significantly increased at 48-h post-treatment with 1 and 10 nM TCDD compared to control cells exposed to vehicle alone (Figure 2A, B). In addition, we observed that *HK2* expression was up-regulated as soon as 24 h post-TCDD (10 nM) treatment (Supplementary Figure S2A). We also assessed *HK2* expression level upon TCDD treatment in the colon cancer cells HCT116 and we observed that TCDD exposure significantly increases *HK2* expression at mRNA and protein levels (Figure 2C, D). We next assessed whether other AHR agonists could promote *HK2* expression. We

treated U2OS cells with benzo[*a*]pyrene (BaP) or with 6-formylindolo[3,2-*b*]carbazole (FICZ) at different concentrations and for different durations. We observed that *HK2* expression was up-regulated in a time- and concentration-dependent manner upon BaP treatment at mRNA and protein levels (Figure 2E, F, Supplementary Figure S2B). Upon FICZ treatment, *HK2* mRNA expression was slightly up-regulated but no difference was detectable at protein level compared to vehicle-treated cells (Figure 2G and Supplementary Figure S2C, D). Altogether, these data indicate that different agonists of AHR induce *HK2* expression in human cells in a time- and concentration-dependent manner, although the kinetic and amplitude of induction are quite variable.

We also investigated whether *HK2* expression was regulated upon TCDD exposure *in vivo*, in tissues expressing high levels of AHR, such as the lung (18). To do that, we studied *Hk2* expression by RT-qPCR and western blot in lung tissues from adult male mice exposed to TCDD *in utero* and during lactation. We observed significant higher *Hk2* mRNA and protein levels compared to vehicle-treated mice tissues (Figure 2H). *Hk2* is thus also regulated by TCDD exposure *in vivo*.

Finally, to gain a broader insight into the association between modulators of AHR activity and the expression of *HK2*, we interrogated the Comparative Toxicogenomics Database (CTD) (38). Among the 266 chemicals (or mixtures) interactions reported to modulate AHR activity in human cells, we identified 18 compounds that also regulate *HK2* expression (fold enrichment: 8.05; $P(X = 18) = 8.06 \times 10^{-12}$) (Supplementary Figure S2E and Table S2). Among these associations, we retrieved complex chemical mixtures such as cigarette smoke as well as several HAH and PAH compounds. This data mining analysis further supports that *HK2* is transcriptionally regulated by several modulators of AHR pathway in human cells, although it appears context (agonist- and cell type-) dependent.

AHR regulates *HK2* mRNA level upon activation by TCDD

We next investigated whether AHR activity was involved in the regulation of *HK2* expression upon AHR agonists exposure. We first used a competitive inhibitor of AHR called CH-223191 (52). We tested the effect of different concentrations of CH-223191 in U2OS cells on *HK2* expression. We did not observe any change in *HK2* expression at the mRNA and protein levels with concentrations up to 10 μ M for 48 h (Supplementary Figure S2F–G). We then treated U2OS cells with CH-223191, at the time of TCDD treatment, and monitored *HK2* and *CYP1A1* expression. We observed that both genes were induced by TCDD as expected, and that co-treatment with CH-223191 abolishes their induction (Figure 2I).

To strengthen these observations, we next transfected U2OS cells with *AHR* small interfering RNAs (siRNAs) and 24 h later treated the cells with 10 nM TCDD for 48 h. RT-qPCR analyses indicated low levels of *AHR* expression in U2OS cells and confirmed the knock-down of *AHR* expression (Figure 2J). We also observed that depletion of *AHR* prevents *HK2* mRNA up-regulation in response to TCDD treatment, while it has no detectable effect on *HK2*

mRNA expression in control treated cells (Figure 2J). All these results show that AHR regulates *HK2* expression in response to its activation by TCDD.

Over-expression of *HK2* promotes an imbalance in AHR signalling pathway proteins abundance

We then investigated the consequences of *HK2* over-expression in human cells by establishing U2OS cells stably expressing either GFP-*HK2* (U2OS-GFP-*HK2* cells) or GFP alone as control (U2OS-GFP cells). By western blot, we observed that 48 h post-transfection, GFP-*HK2* is expressed at its expected size in U2OS-GFP-*HK2* cells (Supplementary Figure S3A). Numerous studies have previously reported, notably in osteosarcoma cells (53,54), that the stable over-expression of *HK2* increases culture media acidification (due to increased release of lactate, an end-product of glycolysis) and cell proliferation. We thus monitored these two parameters. We confirmed that the pH of the culture medium of the U2OS-GFP-*HK2* cells is lower than that of U2OS-GFP cells (Supplementary Figure S3B) and that expression of GFP-*HK2* promotes cell proliferation compared to GFP alone (Supplementary Figure S3C). These results validate that GFP-*HK2* is a functional protein in our experimental conditions.

We next wanted to get deeper insight into the signalling and molecular pathways regulated by *HK2* over-expression. We used a quantitative proteomic approach to characterise and compare the proteome of U2OS and U2OS-GFP-*HK2* cells (Figure 3A). We detected 7278 proteins by mass spectrometry and as anticipated, we observed that the amount of *HK2* is significantly higher in U2OS-GFP-*HK2* compared to U2OS cells ($P < 0.001$) (Supplementary Figure S3D). We also confirmed previous reports (27) showing that *HK2* over-expression promotes an imbalance in glycolytic enzymes abundance with for instance lower levels of *HK1* ($P < 0.001$) and *GAPDH* ($P < 0.001$), and heightened level of *LDHA* ($P < 0.001$) (Supplementary Figure S3E). Following downstream statistical analyses, we identified the differentially expressed proteins (log ratio > 1 and P -value < 0.001) between the two cell lines. A functional annotation of this list using the QIAGEN Ingenuity Pathway Analysis (IPA) software revealed several pathways deregulated upon *HK2* over-expression, among which ‘AHR signalling’ ($P = 8.32 \times 10^{-8}$) with 29 associated proteins. This list includes proteins involved in phase-I and phase-II detoxification mechanisms (*ALDH1B1*, *ALDH3A1*, *ALDH3A2*, *ALDH6A1*, *GSTK1*, *GSTM3*, *GSTP1*, *GSTZ1*, *NQO1* and *NQO2*); cell cycle regulators (*CCND2*, *CCND3*, *CCNE1*, *CDK4*, *CDK6*, *CDKN2A*, *CHK2* and *RB*); chromatin factors (*EP300*, *JUN*, *NCOA3* and *NFKB2*) and signalling proteins (*HSPB1*, *MAPK1*, *MAPK8*, *RXRA*, *SRC*, *TGFB2* and *TGM2*) (Figure 3B). Of note, additional proteins of the AHR network were detected in our proteomic analysis although not robustly enough to be considered by IPA, including most of the CYP enzymes and AHR itself (Supplementary Table S3). As an orthogonal validation of the proteomic data, we performed western blot analyses. We validated that *HK2* over-expression leads to the down-regulation of *HK1* and *JUN* and the up-regulation of *LDHA*, *CHK2* and *NQO1* at pro-

tein levels in U2OS-GFP-*HK2* compared to U2OS-GFP cells (Figure 3C).

Our analysis, using the IPA software, also indicated that 12 of these 29 proteins are predicted as ‘upstream regulators’, suggesting that they potentially regulate additional proteins identified as mis-regulated by *HK2* over-expression (Supplementary Table S4). For instance, *CDKN2A* is predicted to regulate 32 proteins regulated by *HK2* over-expression in our analysis ($P = 1.52 \times 10^{-8}$). This complementary analysis led us to identify 166 downstream proteins including 16 which are annotated as belonging to ‘AHR signalling’ (Figure 3D, E). We also observed that 24 of the 29 genes coding the proteins of the *HK2*/AHR signalling were previously identified as AHR ChIP targets in other cell lines such as MCF-7, HepG2 or GM17212 cells (Figure 3F). These correlative associations suggest that *HK2* might actually regulate AHR signalling at different levels and that several proteins of the *HK2*/AHR network might cross-regulate each other functions.

Over-expression of *HK2* enhances a subset of AHR transcriptional target genes expression

Based on our proteomics findings, we tested whether AHR transcriptional activity was altered in U2OS-GFP-*HK2* cells. To do so, we compared by RT-qPCR the expression level of prototypical AHR target genes *CYP1A1*, *CYP1B1* and *ALDH3A1* between U2OS-GFP and U2OS-GFP-*HK2* cells. We observed higher expression levels of *CYP1B1* and *ALDH3A1* in U2OS-GFP-*HK2* cells compared to U2OS-GFP, while the level of expression of *CYP1A1* was similar between the two cell lines (Figure 4A). We repeated such analyses in another osteosarcoma cell line, 143B, stably expressing GFP or GFP-*HK2*. We could confirm the expression of the GFP-*HK2* fusion protein in these cells (Supplementary Figure S3F) and observed again gene-specific effects, with increased expression of *CYP1B1* in 143B-GFP-*HK2* cells compared to 143B-GFP control cells, while *ALDH3A1* expression was not detected in both cell lines and *CYP1A1* expression was similar between the two cell lines (Figure 4B). These analyses revealed that over-expression of *HK2* and subsequent perturbation of the AHR network alters the expression levels of a subset of AHR target genes in both cell lines.

We wondered whether these changes in gene expression could be a consequence of a change in the intracellular abundance of known endogenous ligands of AHR such as tryptophan (TRP) derivatives (11,12). We observed no major difference in the levels of 3-OH-TRP, TRP, kynurenic acid and 3-OH-kynurenine between U2OS-GFP and U2OS-GFP-*HK2* cells, arguing against such a scenario (Supplementary Figure S4).

Over-expression of *HK2* promotes a better response to AHR agonists in specific cell lines

We then investigated whether *HK2* over-expression might also alter the response to AHR agonists. To do so, we treated cells over-expressing GFP and GFP-*HK2* with BaP or FICZ and monitored by RT-qPCR the expression level of *CYP1A1*, *CYP1B1* and *ALDH3A1*. We observed that

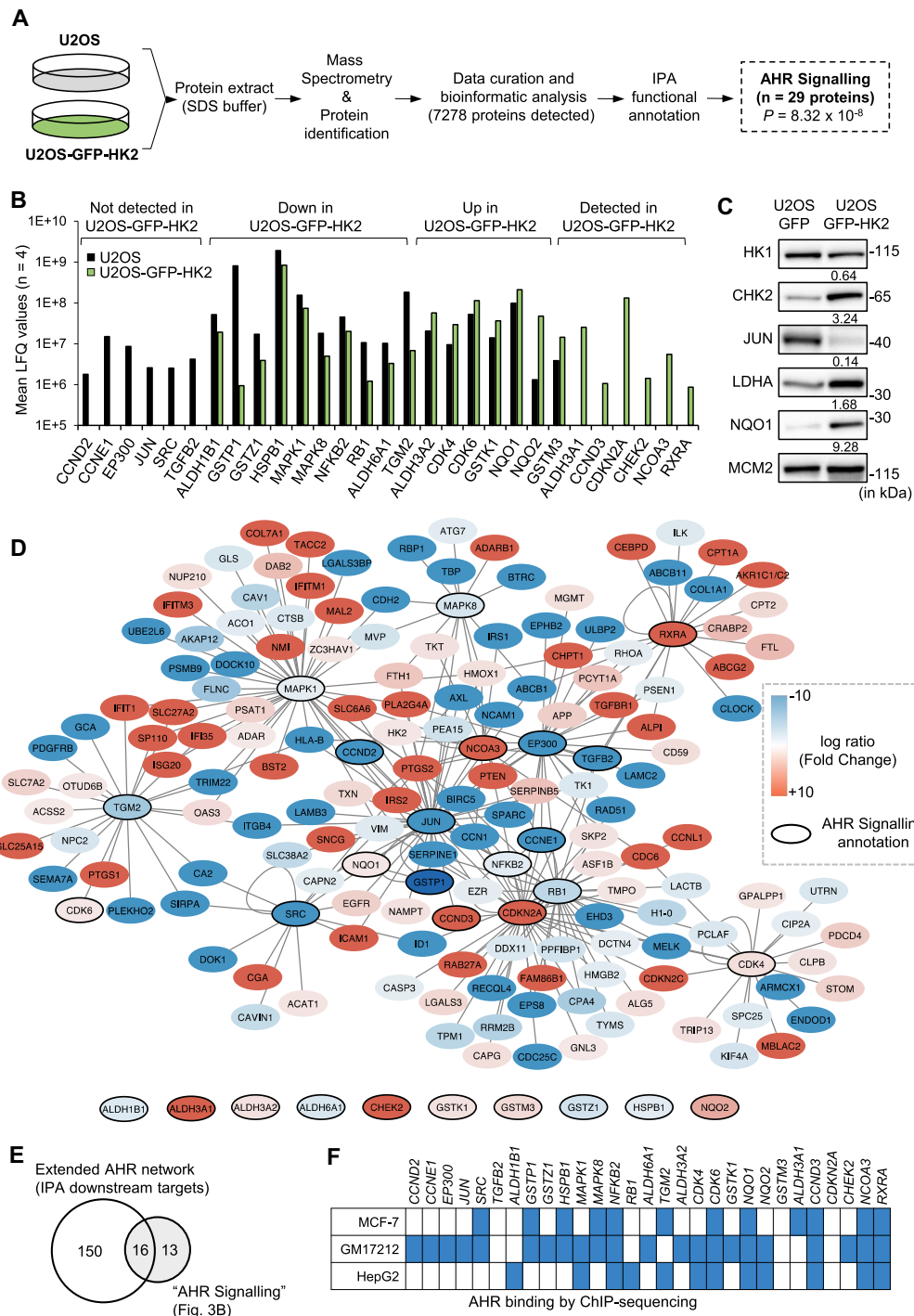


Figure 3. Over-expression of *HK2* alters the abundance of a subset of proteins involved in the AHR network. (A) Schema of the protocol used to characterise U2OS and U2OS-GFP-HK2 proteomes. Four samples of each cell lines were analysed by mass spectrometry. (B) Mean of label free quantification (LFC) values of proteins annotated by Ingenuity Pathway Analysis (IPA) software as belonging to the ‘AHR signalling pathway’ and identified in the proteomic approach with significant changes in abundance between U2OS and U2OS-GFP-HK2 cells (n = 4). Targets are grouped in four classes according to their abundance levels in U2OS and U2OS-GFP-HK2 cells. (C) Western blot analysis of a subset of proteins identified in the proteomic analysis and belonging to the ‘AHR signalling pathway’. Relative quantifications between cell lines are indicated below the blots (relative to MCM2 levels which is not differentially expressed between U2OS and U2OS-GFP-HK2 cells according to the proteomic data). (D) Cytoscape visualization of proteins deregulated in U2OS-GFP-HK2 compared to U2OS cells and annotated as ‘AHR signalling pathway’ or ‘downstream targets’ (i.e targets of predicted Upstream Regulators) using the IPA software. Proteins are colour-coded with up-regulated proteins in U2OS-GFP-HK2 depicted in red and proteins down-regulated depicted in blue. Proteins annotated as ‘AHR signalling pathway’ are highlighted by bold lines. (E) Venn diagram showing the overlap between ‘AHR signalling pathway’ proteins and ‘downstream targets’. (F) Genes encoding proteins of the HK2/AHR network identified as AHR genomic targets in MCF-7, GM17212 and HepG2 cells (either in presence or absence of agonist). Blue indicates the presence of an AHR binding sites in the vicinity of the promoter and/or in the coding sequence of the gene.

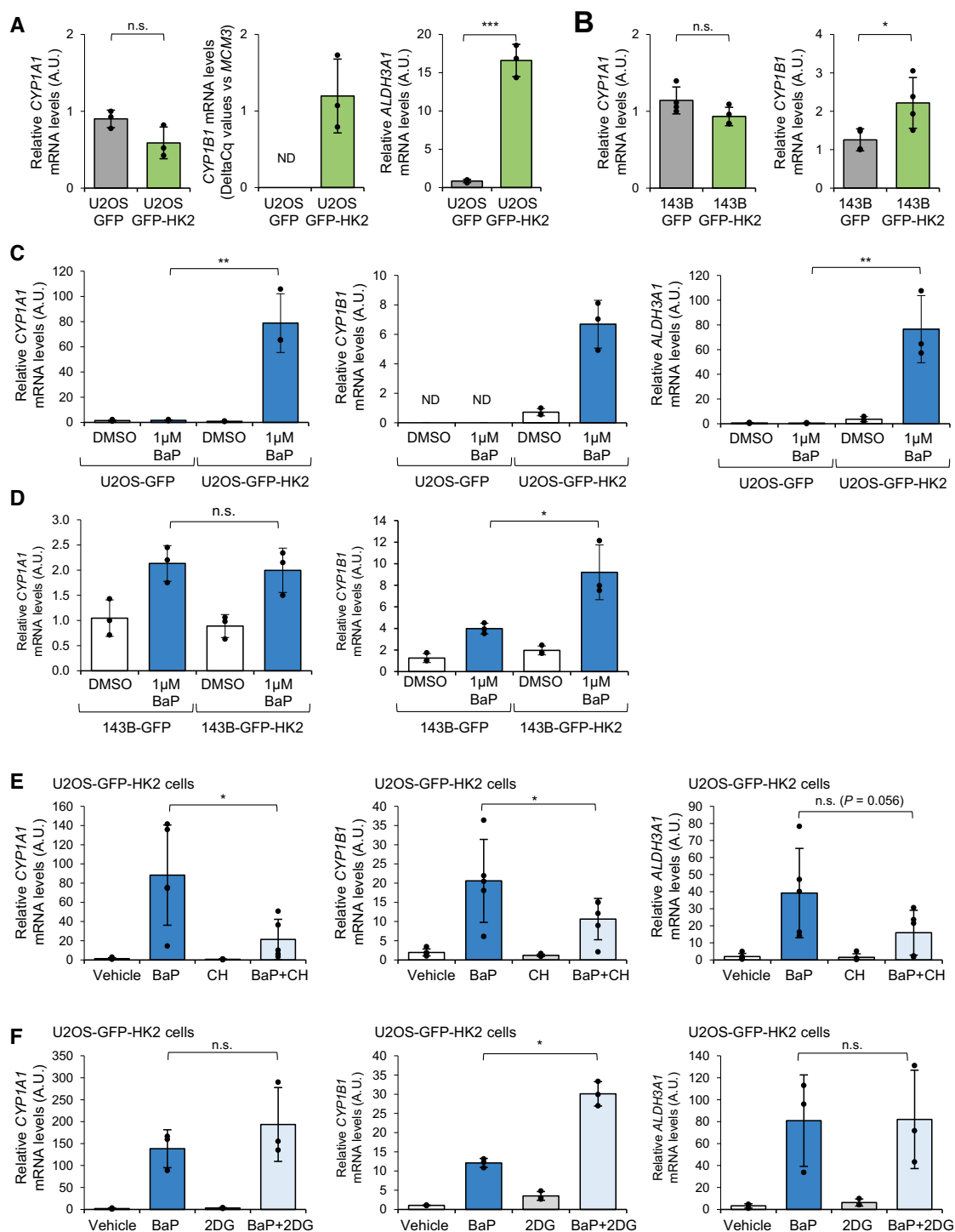


Figure 4. Over-expression of *HK2* enhances a subset of AHR transcriptional target genes expression and potentiates the AHR transcriptional response to BaP exposure. (A) RT-qPCR analysis of *CYP1A1* (left graph), *CYP1B1* (middle graph) and *ALDH3A1* (right graph) expression in U2OS cells expressing GFP or GFP-HK2 ($n = 3$). A.U., arbitrary unit. ND, not detected by RT-qPCR. (B) RT-qPCR analysis of *CYP1A1* (left graph) and *CYP1B1* (right graph) expression in 143B-GFP and 143B-GFP-HK2 cells ($n = 4$). A.U., arbitrary unit. Of note, *ALDH3A1* is not detected in these cell lines. (C) RT-qPCR analysis of *CYP1A1* (left graph), *CYP1B1* (middle graph) and *ALDH3A1* (right graph) expression in U2OS cells expressing GFP and GFP-HK2 and treated with BaP 1 μ M or vehicle (i.e. DMSO) for 48 h ($n = 3$). ND, not detected by RT-qPCR. (D) RT-qPCR analysis of *CYP1A1* (left graph), *CYP1B1* (right graph) expression in 143B cells expressing GFP and GFP-HK2 and treated with BaP 1 μ M or vehicle (i.e. DMSO) for 48 h ($n = 3$). (E) RT-qPCR analysis of *CYP1A1* (left graph), *CYP1B1* (middle graph) and *ALDH3A1* (right graph) expression in U2OS-GFP-HK2 cells treated with vehicle (i.e. DMSO), BaP (1 μ M), AHR antagonist CH-223191 (10 μ M) or BaP + CH-223191 for 48 h ($n = 5$). A.U., arbitrary unit. (F) RT-qPCR analysis of *CYP1A1* (left graph), *CYP1B1* (middle graph) and *ALDH3A1* (right graph) mRNA levels in U2OS-GFP-HK2 cells treated with vehicle (i.e. DMSO), BaP (1 μ M), hexokinase inhibitor 2-DG (2mM) or BaP + 2-DG for 48 h ($n = 3$). A.U., arbitrary unit. P -value was calculated using unpaired Student's t -test for panels A, B, C, D, and paired Student's t -test for panels E and F. n.s., not significant; * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$.

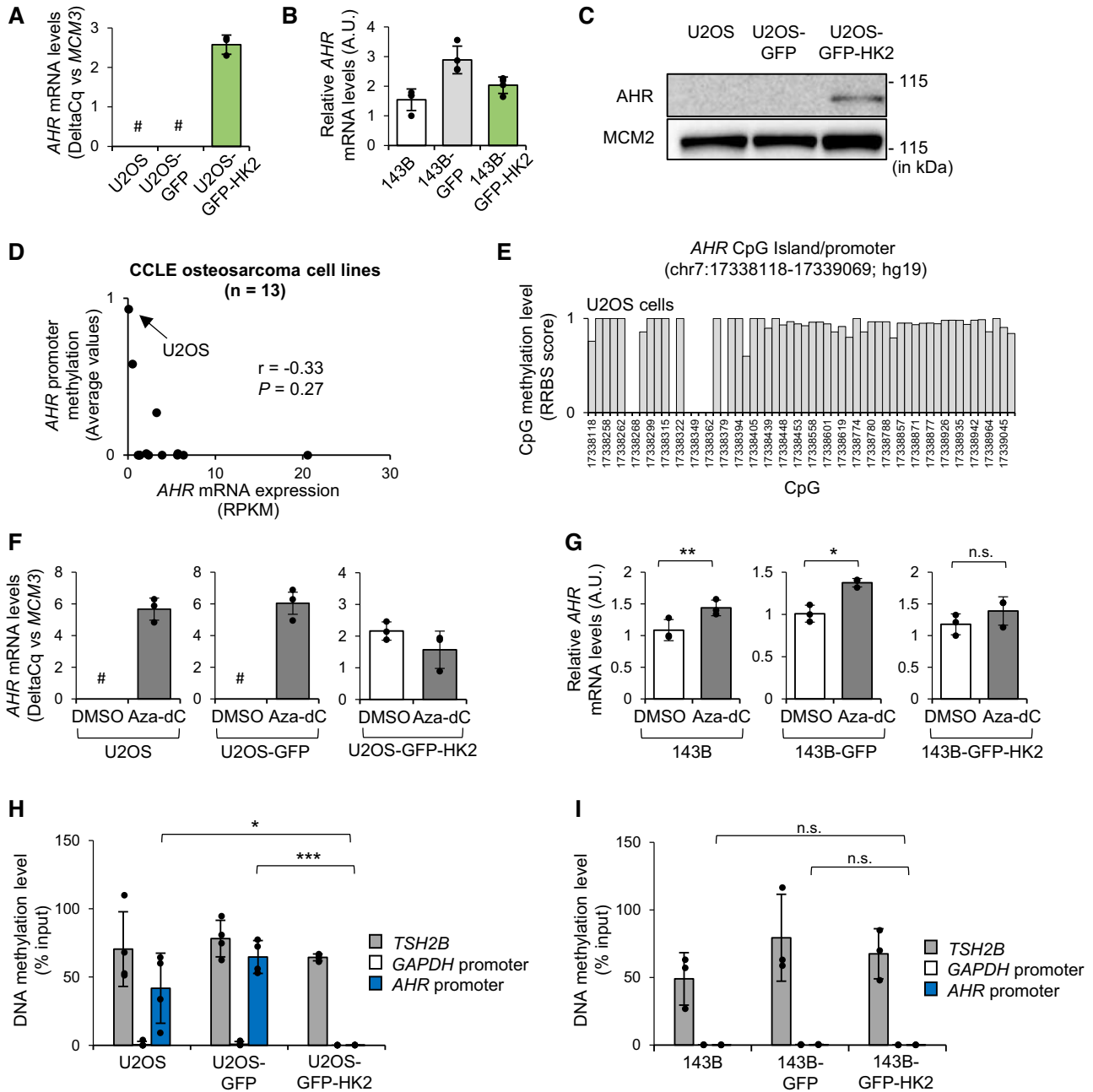


Figure 5. Over-expression of *HK2* enhances *AHR* expression and it correlates with the DNA demethylation of *AHR* promoter in U2OS cells. (A) RT-qPCR analysis of *AHR* expression in U2OS, U2OS-GFP, U2OS-GFP-HK2. Data are presented as DeltaCq values versus *MCM3* for U2OS cell lines ($n = 3$; #, high Cq values for *AHR* (>38)). (B) RT-qPCR analysis of *AHR* expression in 143B, 143B-GFP and 143B-GFP-HK2 cells ($n = 4$). A.U., arbitrary unit. (C) Western blot analysis of AHR and MCM2 levels in U2OS, U2OS-GFP and U2OS-GFP-HK2 cells. MCM2 is used as loading control as it is not differentially expressed between U2OS and U2OS-GFP-HK2 cells according to the proteomic data. (D) Graph showing the correlation between *AHR* promoter methylation status and *AHR* expression from the CCLC osteosarcoma cell lines repository. DNA methylation are presented as average beta-value for *AHR* promoter (0: no methylation; 1: complete methylation). (E) Individual CpG methylation levels at the *AHR* CpG island promoter in U2OS cells defined by RRBS score (0: no methylation; 1: complete methylation). (F) RT-qPCR analysis of *AHR* expression in U2OS, U2OS-GFP and U2OS-GFP-HK2 cells after treatment with vehicle (i.e. DMSO) or with the DNA demethylating agent 5-aza-2'-deoxycytidine (10 μ M; Aza-dC) for 48 h. Data are presented as 'DeltaCq' values versus *MCM3* ($n = 3$). A.U., arbitrary unit. #, high Cq values for *AHR* (>38). (G) RT-qPCR analysis of *AHR* expression in 143B, 143B-GFP and 143B-GFP-HK2 cells after treatment with vehicle (i.e. DMSO) or with the DNA demethylating agent 5-aza-2'-deoxycytidine (10 μ M; Aza-dC) for 48 h ($n = 3$). A.U., arbitrary unit. (H, I) MeDIP-qPCR analysis of the methylation level of *AHR* promoter in (H) U2OS, U2OS-GFP and U2OS-GFP-HK2 cells ($n = 4$) and (I) 143B, 143B-GFP and 143B-GFP-HK2 cells ($n = 3$). *TSH2B* region is used as reference of heavily methylated sequence, while *GAPDH* region is used as a reference of unmethylated DNA. Paired and unpaired Student's *t*-test were used to calculate *P*-value respectively in panels G and H-I. n.s., not significant; * $P < 0.05$; *** $P < 0.001$.

the induction of each gene was markedly higher in U2OS-GFP-HK2 cells compared to U2OS-GFP cells upon BaP or FICZ treatment (Figure 4C and Supplementary Figure S5A). *HK2* over-expression is thus potentiating AHR transcriptional activity in U2OS cells, in response to agonists. Intriguingly, we did not observe a similar effect in the 143B cell line. We observed that the induction of *CYP1A1* mRNA expression in response to BaP and FICZ treatments is similar in 143B-GFP and in 143B-GFP-HK2 (Figure 4D and Supplementary Figure S5B). In the case of *CYP1B1*, the gene was induced by BaP and FICZ and levels of expression were much higher in 143B-GFP-HK2 cells upon treatments compared to 143B-GFP cells. Nonetheless, since the *CYP1B1* gene was already expressed at higher levels in untreated 143B-GFP-HK2 cells, the amplitude of induction was actually similar in the two cell lines (Figure 4D and Supplementary Figure S5B).

To confirm that the up-regulation of these AHR prototypical target genes is mediated by AHR, we co-treated U2OS-GFP-HK2 cells with BaP and CH-223191. We observed that CH-223191 limits the induction of *CYP1A1*, *CYP1B1* and *ALDH3A1* upon BaP treatment, consistent with AHR involvement (Figure 4E). We also tested whether inhibition of hexokinase activity would affect the expression of these AHR targets upon BaP treatment in U2OS-GFP-HK2 cells. We co-treated these cells with BaP and with 2-deoxy-D-glucose (2-DG), an allosteric inhibitor of hexokinase activity (55). In BaP + 2-DG co-treated cells, we did not observe lower expression levels of *CYP1A1*, *CYP1B1* and *ALDH3A1* compared to BaP treated cells (Figure 4F). These data indicate that 2-DG treatment does not block the induction of AHR target genes upon BaP exposure.

All these data show that over-expression of *HK2* amplifies the transcriptional levels of AHR target genes in response to AHR agonists, and that this response is AHR dependent and not prevented by 2-DG treatment.

Over-expression of *HK2* is associated with the DNA demethylation of *AHR* promoter and higher *AHR* expression in U2OS cells

We hypothesize that the cellular context, including *AHR* expression level, might explain the differential regulation of AHR activity in the two osteosarcoma cell lines. We thus monitored *AHR* expression by RT-qPCR in the different U2OS and 143B cell lines we established (Figure 5A, B). We detected higher *AHR* mRNA levels in U2OS-GFP-HK2 compared to respective control cells while *AHR* levels were quite similar between 143B, 143B-GFP and 143B-GFP-HK2 cells (Figure 5A, B). This observation suggests that a dramatic change in *AHR* transcriptional status between U2OS/U2OS-GFP and U2OS-GFP-HK2 cells is linked to the dramatic change in AHR transcriptional outcomes in response to agonists exposure. Importantly, by western blot, we confirmed higher levels of AHR protein in U2OS-GFP-HK2 cells compared to control cells (Figure 5C).

As *AHR* promoter contains CpG sequences susceptible to DNA methylation (19), we investigated whether DNA methylation was involved in the regulation of *AHR* expression. We first assessed the methylation status and the expression level of *AHR* in a panel of osteosarcoma cells us-

ing the Cancer Cell Line Encyclopedia (CCLE) datasets (35,46). We observed that the methylation of *AHR* promoter is negatively correlated with *AHR* mRNA expression in this set of cell lines and that in U2OS cells *AHR* promoter is highly methylated and *AHR* mRNA expression is low (Figure 5D) (data are not available for the 143B cell line in this database). The methylation of *AHR* promoter in U2OS cells was further confirmed by interrogating single CpG methylation analyses from the CCLE (Figure 5E). We thus treated U2OS, U2OS-GFP and U2OS-GFP-HK2 cells with DNA demethylating agent 5-aza-2'-deoxycytidine for 48 h and monitored *AHR* expression by RT-qPCR. We detected *AHR* expression in 5-aza-2'-deoxycytidine-treated U2OS and U2OS-GFP cells at 48 h while it was hardly detectable in vehicle-treated cells. In contrast, expression of *AHR* was detectable in U2OS-GFP-HK2 cells in treated and untreated conditions, at roughly similar levels (Figure 5F). These results indicate that *AHR* expression is regulated by DNA methylation in U2OS and U2OS-GFP cells. Consistent with this conclusion, RT-qPCR analyses showed that in 143B, 143B-GFP and 143B-GFP-HK2 cells, which already express *AHR* in untreated conditions, 5-aza-2'-deoxycytidine treatment for 48 h moderately increases *AHR* expression (Figure 5G).

To further confirm the role of DNA methylation in the regulation of *AHR* expression we assessed the methylation status of the *AHR* promoter by methylated-DNA immunoprecipitation (MeDIP) in U2OS, 143B and their derivative cell lines (Figure 5H, I). As controls, we used *GAPDH* promoter as an unmethylated region and *TSH2B* promoter as a heavily methylated region as indicated by the manufacturer (Diagenode). The methylation status of control regions was in line with what was expected (Figure 5H, I). For the *AHR* promoter region we observed high levels of CpG methylation in U2OS and U2OS-GFP cells and low methylation levels in U2OS-GFP-HK2 cells as well as in 143B, 143B-GFP and 143B-GFP-HK2 cells (Figure 5H, I). These data indicate that over-expression of *HK2* is associated with the demethylation of the *AHR* promoter in U2OS cells which may explain the enhanced expression of *AHR*.

HK2 expression negatively correlates with the methylation of *AHR* promoter in several cancers

We next explored the relevance of the *HK2*/*AHR* axis in cancer by investigating the correlation between the expression level of *HK2* and the expression level of *AHR* using the data of 33 types of cancer available through The Cancer Genome Atlas (TCGA) (47,48,56). We found that *HK2* and *AHR* mRNA expression levels are positively correlated in cancer, notably in nine cancer types (Figure 6A). In addition to TCGA datasets, we also analysed two different osteosarcoma (OS) cohorts (NCI/TARGET/OS and GSE21257) publicly available and commonly used by the community (49). We found no significant correlation between *AHR* and *HK2* mRNA expression in OS tumours in the two cohorts (Supplementary Figure S6).

Focusing on the nine cancer types exhibiting a clear correlation between *HK2* and *AHR* expression, we then investigated a possible correlation between *HK2* expression and the methylation of *AHR* promoter. We observed a signifi-

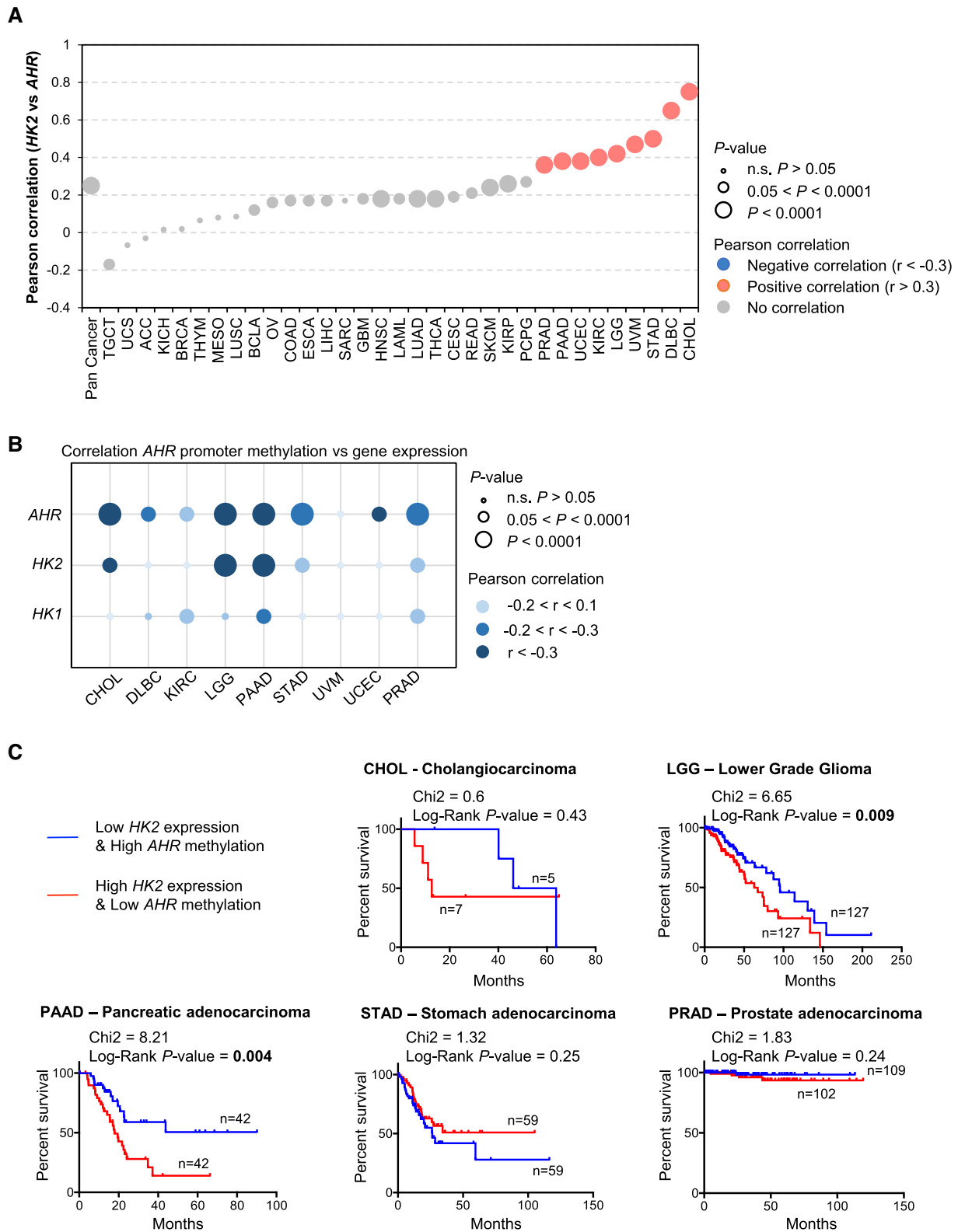


Figure 6. Relationship between the expression of *HK2* and the methylation of *AHR* promoter in cancer. (A) Relationship between the expression levels of *HK2* and *AHR* retrieved in 33 different types of cancer from the TCGA. Significant correlation, Pearson *r* coefficient >0.3 or <-0.3 are indicated in colours (blue, negative correlation; salmon, positive correlation) (B) Relationship between the expression levels of *AHR*, *HK2* and *HK1* and the methylation level of *AHR* promoter in cancers. (C) Prognostic value of the *AHR*/*HK2* axis in five different types of cancer exhibiting a negative correlation between *HK2* expression and *AHR* promoter methylation.

cant negative correlation between *HK2* levels and *AHR* promoter methylation in cholangio-carcinoma (CHOL), lower grade glioma (LGG), pancreatic cancer (PAAD), stomach cancer (STAD) and prostate cancer (PRAD) (Figure 6B). Furthermore, we did not find such correlation between the expression level of *HK1* and the methylation of *AHR*, indicating some level of specificity (Figure 6B). Finally, in lower grade glioma (LGG) and pancreatic cancer (PAAD), tumours exhibiting high levels of *HK2* and low levels of *AHR* promoter methylation are correlated with shorter overall survival (Figure 6C).

These data indicate that *HK2* expression negatively correlates with the methylation of *AHR* promoter in specific cancer types, which may have clinical implication.

DISCUSSION

It is well recognized that AHR signalling activation has variable effects on the transcription of its targets. Nonetheless, the details of the proteins and pathways involved in the regulation of AHR signalling activity are still far from complete and it might depend on features proper to each cellular context such as possible genetic mutations, exposure to exogenous cues and stage of cell differentiation (7). In here, we provide evidences that *HK2* is a transcriptional target of AHR, as suggested by previous large-scale gene expression studies (22,57–59), and studies indicating that AHR might be an important regulator of glycolytic genes expression and glycolysis end-points (3,7,51,60–62). In MCF-7 cells, we identified a cluster of AHR binding sites in the vicinity of exon 3 of *HK2* gene; and binding of AHR at the promoter of *HK2* in two other cell lines GM17212 and HepG2. Using a ChIP-qPCR analysis we demonstrated similar AHR binding sites in U2OS cells treated with TCDD including binding sites nearby exon 3 and at the promoter of *HK2*. The binding at the promoter is detected 1 h after TCDD addition, but not at a later time point (24 h), indicating a transient binding of AHR at *HK2* promoter. In contrast, the binding nearby exon 3, and at the *CYP1A1* promoter, is observed up to 24 h and might reflect *cis*-regulatory elements mediating long-term and persistent effects of TCDD. It would thus be interesting to further dissect the role of these different *cis*-regulatory elements in the *HK2* locus to better define their functions in the regulation of *HK2* expression upon AHR agonists exposure. We indeed showed that in response to AHR agonists exposure, *HK2* expression is enhanced, although the fold-increase is quite variable depending on the duration of the exposure, the concentration and the nature of the ligand. All these findings highlight that the regulation of *HK2* expression by AHR is context-specific and might rely on the binding of additional cofactors of AHR, either at the promoter or nearby exon 3 (1,50,51). Using ENCODE ChIP-sequencing datasets we identified binding sites for cMYC and FOXO factors which may cooperate with AHR in the regulation of *HK2* expression (50,51). The context-specific regulation of *HK2* by AHR may alternatively be explained by our findings that *HK2* regulates AHR transcriptional activity and thus that AHR or some of its targets might further modulate the kinetic and induction of *HK2* expression upon AHR activation. For instance, transglutaminase 2 (TGM2), which is

less abundant in U2OS-GFP-*HK2* cells compared to controls, is also a known regulator of *HK2* expression (63,64). Similarly, retinoblastoma protein 1 (RB1), also affected by over-expression of *HK2*, is a known partner of AHR in the regulation of gene expression (65). Thus, different layers of regulation, both intrinsic to cell type (i.e. mutations) or exogenous (i.e. environmental cues) may have different outcomes on the *HK2*/AHR axis and *HK2* expression.

By investigating the molecular consequences of *HK2* over-expression we observed that numerous proteins contributing to AHR pathway activity, or regulated by this pathway, are mis-regulated upon *HK2* over-expression including cell cycle regulators, metabolic enzymes, signalling proteins and AHR targets. Whether additional AHR targets are also regulated by *HK2* remains possible as the proteomic approach is not exhaustive. For instance, it has been shown that TCDD regulates expression of RANKL, CXCR4, CXCL2, COX2 and PGE2 in osteosarcoma MG63 cancer cells (66). None of these proteins as well as most of the CYP enzymes and AHR itself, were efficiently detected, quantified and thus analysed in our study. This is most likely due to technical limitations as peptides for some of these proteins were actually detected in a limited number of samples. It is thus unclear whether such classes of AHR targets are also mis-regulated in U2OS-GFP-*HK2* compared to control cells. Following up on this proteomic data we unveiled that U2OS-GFP-*HK2* cells present heightened expression levels of AHR protein and of its target genes, as well as enhanced response to BaP and FICZ, compared to U2OS-GFP control cells. Intriguingly, in 143B-GFP-*HK2* cells we also observed heightened levels of AHR target genes expression under basal condition but did not observe a better response to AHR agonists. There might thus be a direct link between *HK2* gene expression and AHR transcriptional activity. In that scenario, expression levels of *HK2* might mitigate, or in contrary amplify, the consequences of AHR agonists on cell fate. Intriguingly, it has been shown that glucose can modulate AHR activity in aortic endothelial cells and that pyruvate kinase muscle isoform M2 (PKM2) regulates AHR transcriptional activity (67,68). It is thus likely that *HK2* might alter AHR activity at multiple levels, as suggested by our data, to regulate cell biology and fate. Importantly, we observed that 2-DG treatment, which inhibits hexokinase activity and glycolysis, did not block the transcriptional response to AHR agonists, suggesting that *HK2* role in glycolysis may have minor effect on the regulation of AHR signalling.

In addition, we documented a function of *HK2* on the regulation of the activity of the *AHR* promoter. It has previously been shown that *AHR* promoter is regulated by DNA methylation in specific cell types (19). We obtained compelling evidences that over-expression of *HK2* leads to *AHR* promoter demethylation in U2OS cells, and that it correlates with *AHR* expression level. This observation probably explains the strong response of U2OS-GFP-*HK2* cells to AHR agonists tested, such as BaP and FICZ, compared to control cells. While *HK2* over-expression triggers DNA demethylation, indicating a link between *HK2* and *AHR* expression we did not clearly pinpoint the mechanism, although we can speculate based on previous studies that DNA demethylation might facilitate the binding of

specificity protein 1 (SP1) that activates *AHR* expression (19,69). Intriguingly, numerous glycolytic enzymes can actually shuttle into the nucleus including glyceraldehyde 3-phosphate dehydrogenase (GAPDH), PKM2 and phosphoglycerate kinase 1 (PGK1) and regulate nuclear processes as diverse as gene expression, DNA replication and chromatin organisation (68,70–72). It is tempting to speculate that HK2 might also shuttle into the nucleus, and as PKM2, regulate the transcriptional activity of AHR or the activity of its promoter. Consistent with such scenario, upon pro-inflammatory stress and glucose deprivation, HK2 translocates in the nucleus of glioblastoma cells and regulates the expression of xanthine oxidoreductase gene in combination with transcription factor NRF2 (73). This remains the sole example of a potential role of human HK2 in the nucleus, needless to say in a quite specific context. Intriguingly, in yeast, low glucose environment also induces the nuclear localisation of the human HK2 homolog, Hxk2, where it regulates gene expression (74). Thus, further studies will be needed to clarify whether HK2 might regulate gene expression in high glucose environment as in our cellular models and how it may control DNA methylation levels at the *AHR* promoter.

This new axis HK2/AHR might also be important in the context of cancer. Indeed, using TCGA datasets, in 9 of the 33 cancer types that we analysed, we found that *HK2* expression shows a positive correlation with *AHR* expression and in a subset of these 9 cancer types a negative correlation with *AHR* promoter methylation levels. Furthermore, tumours with high *HK2* expression and low *AHR* methylation exhibit a poor prognosis in lower grade glioma (LGG) and pancreatic cancer (PAAD). Importantly, we did not see similar correlation with *HK1* which shares high degree of similarity at the protein level and at the biochemical level with HK2 (26). We also did not document a recurrent binding of AHR at the promoter of *HK1* indicating again a preferential relationship between HK2 and AHR. *HK1* is expressed in most tissues while *HK2* expression is restricted to muscle and adipocytes and it is over-expressed in most cancers. Accordingly, a large body of literature has demonstrated that *HK2* is regulated by numerous micro-RNAs and transcription factors including oncogenes (26). Here, we revealed and characterized an additional regulator of *HK2* expression, AHR, which can either act as an oncogene or tumour suppressor in different cancer types (13–15). Intriguingly, in osteosarcoma (OS) tumours, we did not observe a correlation between *HK2* and *AHR* expression in two independent cohorts of patients. A possible explanation to this observation might be that OS tumours are quite heterogeneous at the genomic and cellular level and that additional criteria need to be taken into consideration to unveil a function of the AHR/HK2 axis in these tumours.

Our study is thus adding *HK2* to the list of AHR targets that also regulate its function. It is well known that CYP enzymes are induced upon AHR agonists exposure and that they contribute to the detoxification or activation of most of these AHR agonists, which directly regulates AHR activation over time (1,10,14). Similarly, AHR repressor (AHRP) and TCDD inducible poly(ADP-ribose) polymerase (TIPARP) are induced by AHR upon activation and contribute to reduce AHR transcriptional activity

by protein/protein interaction and/or protein degradation (1,23–25). Strikingly, as opposed to these later examples, *HK2* is a direct target of AHR that positively contributes to AHR activity and/or expression depending on cell context under basal condition and in response to AHR agonists. Our study further highlights the need to identify and characterize additional AHR signalling factors to fully comprehend the molecular basis of AHR context-dependent cellular effects.

DATA AVAILABILITY

AHR ChIP-sequencing data were previously published and are publicly available on NCBI GEO servers with accession numbers GSE90550, GSE127649 and GSE116638. All software used for data analysis and visualization are freely usable and their references are indicated in the materials and methods. The databases used to collect and generate gene-chemical relationships and gene expression, chromatin and DNA methylation information in human cell lines and cancer patients (e.g. ENCODE, ChIP-Atlas, CTD, CellMinerDB, TCGA, TARGET Osteosarcoma) are publicly available and the publication embargoes on the different datasets included in our analyses have expired. The proteomics data produced in this study have been deposited to the ProteomeXchange Consortium via the PRIDE partner repository with accession number PXD028992. All values and statistical tests reported in the figures and tables of the manuscript are accessible in Supplementary Table S5.

SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online.

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