



HAL
open science

Dynamic regulation of Tgf-B signaling by Tif1 γ : a computational approach.

Geoffroy Andrieux, Laurent Fattet, Michel Le Borgne, Ruth Rimokh, Nathalie Théret

► **To cite this version:**

Geoffroy Andrieux, Laurent Fattet, Michel Le Borgne, Ruth Rimokh, Nathalie Théret. Dynamic regulation of Tgf-B signaling by Tif1 γ : a computational approach.. PLoS ONE, 2012, 7 (3), pp.e33761. 10.1371/journal.pone.0033761 . hal-00851483

HAL Id: hal-00851483

<https://hal.science/hal-00851483v1>

Submitted on 19 Aug 2013

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Dynamic Regulation of Tgf- β Signaling by Tif1 γ : A Computational Approach

Geoffroy Andrieux^{1,2}, Laurent Fattet³, Michel Le Borgne², Ruth Rimokh³, Nathalie Th  ret^{1*}

1 Inserm U1085-IRSET, Universit   de Rennes 1, Rennes, France, **2** Universit   de Rennes 1, IRISA, Rennes, France, **3** Inserm U1052/CNRS 5286, Centre de Recherche en Canc  rologie de Lyon, Lyon, France

Abstract

TIF1 γ (Transcriptional Intermediary Factor 1 γ) has been implicated in Smad-dependent signaling by Transforming Growth Factor beta (TGF- β). Paradoxically, TIF1 γ functions both as a transcriptional repressor or as an alternative transcription factor that promotes TGF- β signaling. Using ordinary differential-equation models, we have investigated the effect of TIF1 γ on the dynamics of TGF- β signaling. An integrative model that includes the formation of transient TIF1 γ -Smad2-Smad4 ternary complexes is the only one that can account for TGF- β signaling compatible with the different observations reported for TIF1 γ . In addition, our model predicts that varying TIF1 γ /Smad4 ratios play a critical role in the modulation of the transcriptional signal induced by TGF- β , especially for short stimulation times that mediate higher threshold responses. Chromatin immunoprecipitation analyses and quantification of the expression of TGF- β target genes as a function TIF1 γ /Smad4 ratios fully validate this hypothesis. Our integrative model, which successfully unifies the seemingly opposite roles of TIF1 γ , also reveals how changing TIF1 γ /Smad4 ratios affect the cellular response to stimulation by TGF- β , accounting for a highly graded determination of cell fate.

Citation: Andrieux G, Fattet L, Le Borgne M, Rimokh R, Th  ret N (2012) Dynamic Regulation of Tgf- β Signaling by Tif1 γ : A Computational Approach. PLoS ONE 7(3): e33761. doi:10.1371/journal.pone.0033761

Editor: Dipankar Chatterji, Indian Institute of Science, India

Received: December 13, 2011; **Accepted:** February 21, 2012; **Published:** March 23, 2012

Copyright:    2012 Andrieux et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Funding: This work was supported by the Institut National de la Sant   et de la Recherche M  dicale (www.inserm.fr) and the Ligue Nationale Contre le Cancer (<http://www.ligue-cancer.net>). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Competing Interests: The authors have declared that no competing interests exist.

* E-mail: nathalie.theret@univ-rennes1.fr

Introduction

Complex signaling by transforming growth factor β (TGF- β) forms a pivotal network that plays an essential role in tissue homeostasis and morphogenesis. At the same time, up-regulation and activity of TGF- β has been linked to various diseases, including fibrosis and cancer, by promoting cell proliferation and invasion and the epithelial-mesenchymal transition [1]. TGF- β signaling occurs through association with a heteromeric complex of two types of transmembrane serine/threonine kinases, the type I (T β RI) and type II (T β RII) receptors. TGF- β binding to T β RII induces recruitment and phosphorylation of T β RI, which in turn transmits the signal through phosphorylation of the receptor-bound R-Smad transcription factors, Smad2 or Smad3. Once phosphorylated, the R-Smads hetero-dimerize with their common partner, Smad4. The resulting complexes then migrate to the nucleus, where they regulate the transcription of TGF- β -target genes in conjunction with other transcription factors [2].

Nuclear Transcriptional Intermediary Factor 1 γ , TIF1 γ (also known as tripartite motif protein TRIM33), is a member of the transcriptional intermediary factor 1 family [3] and was recently identified as a new partner of the Smad-dependent TGF- β signaling pathway. A screen for molecules involved in the specification of the embryonic endoderm first revealed TIF1 γ as a Smad4-binding protein and as a negative regulator of TGF- β signaling [4]. TIF1 γ mono-ubiquitinates Smad4, inducing its nuclear export to the cytoplasm, where the FAM/UPS9x deubiquitinating enzyme was recently shown to allow Smad4 recycling [5]. The role of TIF1 γ as a repressor was also reported in

the control of Smad activity during embryogenesis [6]. In contrast, TIF1 γ was identified as a protein partner for receptor-activated Smad2/3, resulting in an alternative positive regulatory Smad4-independent TGF- β signaling pathway [7].

Whether TIF1 γ down-regulates or promotes alternative TGF- β signaling may be linked to the cellular context. TIF1 γ is a ubiquitous protein and its mRNA has been detected in all tissues [8]. Its loss of expression has been shown to favor Kras^{G12D}-dependent precancerous pancreatic lesions [9], induce cell-autonomous myeloproliferative disorders in mice [10] and potentiate TIF1 α -induced murine hepatocellular carcinoma [11], thereby supporting a protective role of TIF1 γ in cancer. Consistent with this view, a decrease in TIF1 γ expression in human pancreatic cancer and human chronic myelomonocytic leukemia has been reported [9,11] and TIF1 γ silencing in human mammary epithelial cell lines was shown to lead to a strong epithelial-mesenchymal transition mediated by TGF- β 1 [12]. In contrast, a pro-tumorigenesis role for TIF1 γ has been suggested by the observation that its expression prevents Smad4-mediated growth inhibition in response to TGF- β [4]. In line with the uncertain role of TGF- β in cancer, TIF1 γ may differentially affect TGF- β signaling according to the cellular context by acting either as tumor suppressor or promoter.

Several mathematical models have been developed to predict the dynamic behavior of TGF- β signaling. In particular, initial differential models that couple signaling with receptor trafficking have significantly improved our understanding of the plasticity of the TGF- β signaling pathway [13]. Models focusing on Smad phosphorylation [14], Smad nucleocytoplasmic shuttling [15,16]

and Smad oligodimerization [17] have also been developed to understand the dynamics and flexibility of Smad-dependent pathways, while integrative models have coupled receptor trafficking to Smad pathways [18–20]. As the latter models recapitulate the essential components of the canonical Smad-dependent TGF- β signaling pathway, they constitute useful tools to investigate the role of new regulatory components of TGF- β signaling.

We have used an integrative modeling approach to explore the impact of TIF1 γ on the outcome of TGF- β signaling. Taking advantage of mathematical models of receptor trafficking [13] and Smad shuttling [16], we have developed a new TGF- β signaling model that includes TIF1 γ and FAM/UPS9x. Our model, which is based on the transient formation of a ternary complex containing TIF1 γ , Smad4 and Smad2/3, successfully reconciles the different observations reported for TIF1 γ -Smad4 [4] and TIF1 γ -Smad2/3 [7] interactions. We show that TGF- β signaling is highly sensitive to the TIF1 γ /Smad4 ratio, suggesting a critical role for the FAM/UPS9x deubiquitinase. This model also predicts how varying TIF1 γ /Smad4 ratios can modulate the cellular response to transient and sustained TGF- β stimulation, accounting for a highly graded TGF- β response. We discuss how the seemingly opposite roles of TIF1 γ may be resolved by taking into account the dynamic balance of interactions involving Smad4 and Smad2/3.

Materials and Methods

Mathematical modeling

The model consists of a system of nonlinear, ordinary differential equations that merge the ODE models of receptor trafficking [13] and Smad shuttling [16]. Briefly, the receptors described in the Smad shuttling model were replaced by those of the receptor trafficking model using unit conversion in a cell volume of 2.27×10^{-12} L. Model building, parameters, system ordinary equations and description of the model in Systems Biology Markup Language (SBML) are detailed in Tables S1 and S2 and Model S1. Model simulations were implemented with the mathematical Scipy library of Python language programming and the Matplotlib Python 2D plotting library was used to visualize the simulation curves.

Cell culture and siRNA transfection

Human mammary epithelial (HMEC) cells infected with a retrovirus carrying hTERT and the oncogenic H-RasV12 (HMEC-TR) allele were provided by R. A. Weinberg [21] and cultured as previously described [12]. Cells were transfected with 5 nM siRNA and 0.5 μ l/ml lipofectamine RNAiMax (Invitrogen) and further cultured in the presence or absence of 10 ng/ml TGF- β 1 (Peprotech) for the indicated times.

Chromatin immunoprecipitation (ChIP)

Assays were carried out on cells transfected with the PAI-1 p800-Luc construct, as previously described [12], using the kit from Upstate Biotechnology. Briefly, cell lysates were subjected to anti-Smad4 (SantaCruz) or anti-TIF1 γ (Bethyl) immunoprecipitation. Smad4- or TIF1 γ -precipitated genomic DNA was subjected to PCR. The 351-bp PAI-1 promoter region harboring the Smad-binding elements was amplified with primers 5'-AGCCAGA-CAAGGTTGTTG-3' and 5'-GACCACCTCCAGGAAAG-3'. An unrelated genomic DNA sequence (actin) was amplified with primers 5'-AGCCATGTACGTTGCTATCCAG-3' and 5'-CTTCTCCTTAATGTCACGCACG-3'.

Relative quantification of mRNA by real-time PCR

Real-time quantitative PCR was performed using the qPCRTM Core Kit for SybrTM Green I from Eurogentec and the ABI Prism 7700 thermocycler (Perkin-Elmer, Foster city, CA, USA). Primer pairs for target genes were: sense CDH11 (OB-Cadherin), 5'CCC TGA AAT CAT TCA CAA TCC3', antisense 5'AGT CCT GCT TCT GCC GAC T3'; CDH2 (N-Cadherin), sense: 5'GTG CAT GAA GGA CAG CCT CT3', antisense: 5'ATG CCA TCT TCA TCC ACC TT3'; HPRT, sense: 5'TGA CCT TGA TTT ATT TTG CAT ACC3', antisense: 5'CGA GCA AGA CGT TCA GTC CT3'.

Western blot analysis

Cell lysates were subjected to SDS-polyacrylamide gel electrophoresis and transferred onto PVDF membranes. The blots were incubated for 1 hr in Tris-buffered saline containing 0.1% Tween 20 and 5% non-fat dry milk and further incubated for 1 hr with specific primary antibodies (anti-Smad4, SantaCruz biotechnology; anti-TIF1 γ , Euromedex). The bound antibodies were visualized with horseradish peroxidase-conjugated antibodies using the ECL-Plus reagent (Roche).

Results and Discussion

Quantitative models for TIF1 γ -dependent TGF- β signaling

Merging receptor trafficking [13] and Smad cytonucleoplasmic shuttling [16] models through their common receptor-ligand complex in the endosome (LRe), we developed new models that integrate TIF1 γ . Kinetic parameters were estimated according to the experimental data from [5] and [7] and are detailed in Table S1. We first constructed two separate models, each taking into account the different hypotheses regarding Smad/TIF1 γ interactions. The first model is based on the TIF1 γ -dependent negative regulation associated with the ubiquitination of Smad4 ([4,5]; Figure 1A). In this model, TIF1 γ interacts preferentially with Smad4 within phosphorylated Smad2-Smad4 complexes in response to TGF- β , leading to a rapid dissociation of complexes and formation of ubiquitinated Smad4 (Smad4ub) that is exported from the nucleus. Similar to the transient interaction of the phosphatase (PPase) with phosphorylated Smad2 [15,16], the formation of TIF1 γ -Smad complexes was neglected because of fast reaction rates. In the cytoplasm, ubiquitinated Smad4 undergoes deubiquitination by FAM/UPS9x (FAM), thereby recycling Smad4 for TGF- β signaling (Figure 1A). We set the same kinetic parameters for association between TIF1 γ and phosphorylated Smad2-Smad4 complexes in the nucleus (pS2S4n) and association between phosphorylated Smad2 and Smad4. Ubiquitination/deubiquitination and phosphorylation/dephosphorylation kinetics were considered to be similar, as previously described [5]. Export of ubiquitinated Smad4 from the nucleus to the cytoplasm was assumed to be 2-fold higher than entry of Smad4 in the nucleus, based on the observation suggesting that ubiquitinated Smad4 is less efficiently retained in the nucleus [4,5].

Our second model is based on results from He *et al.* [7], who proposed that TGF- β induces a competing interaction between TIF1 γ and phosphorylated Smad2, although an association of TIF1 γ with Smad4 was also detected in the nucleus (Figure 1B). In the absence of conclusive experimental data, we considered the kinetic parameters for association between TIF1 γ and either phosphorylated Smad2 or Smad4 in the nucleus to be similar to those for phosphorylated Smad2 with Smad4. To test this hypothesis, we analyzed the effect of a 2-fold decrease in k_{on}/k_{off}

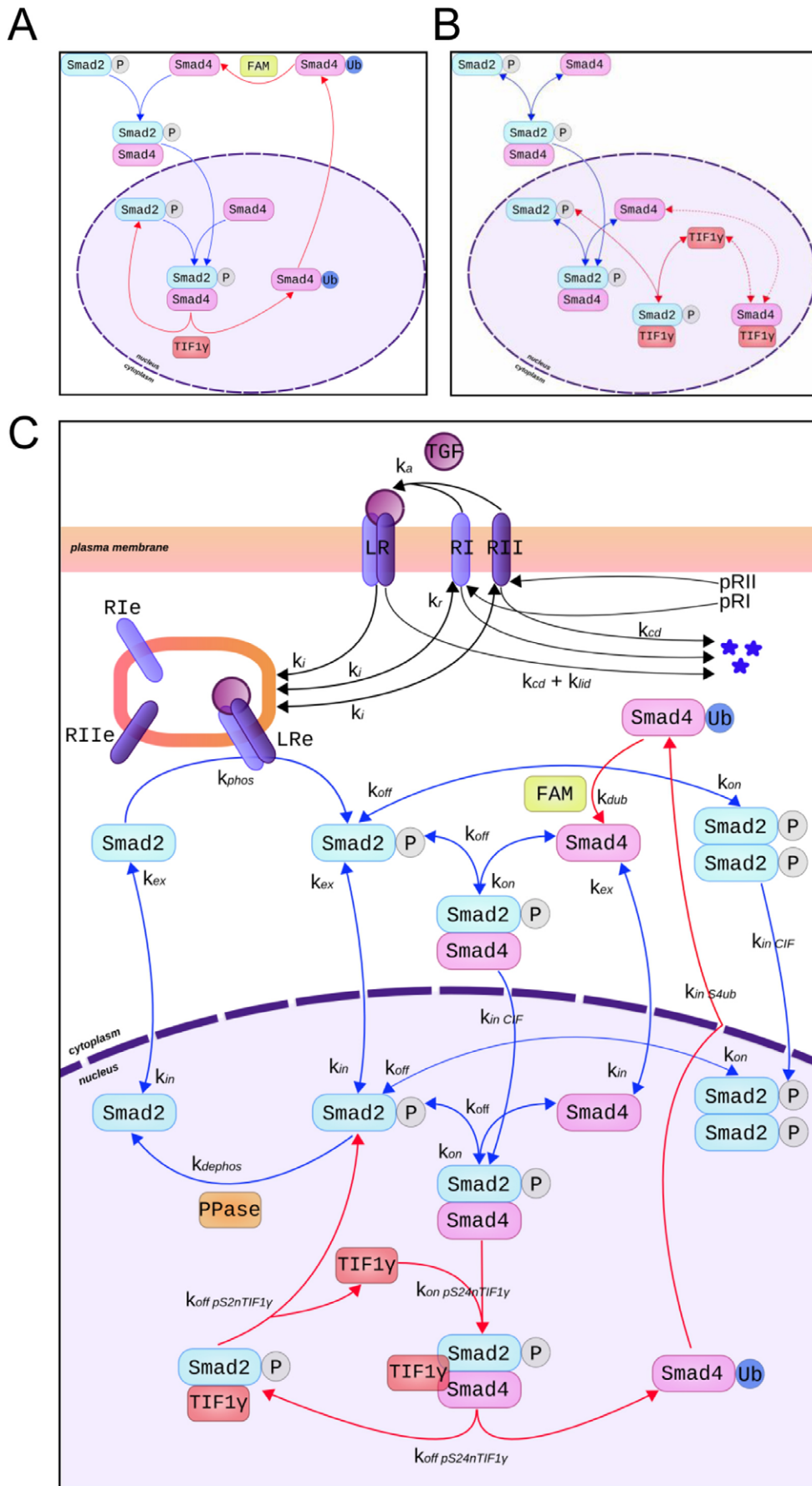


Figure 1. Schematic representation of the models. Detailed information on parameters and entities are given in Tables S1 and S2. A) Model hypothesis from [4]. B) Model hypothesis from [7]. C) Integrated model including TIF1 γ (red rectangle) and FAM (green rectangle). doi:10.1371/journal.pone.0033761.g001

for the association between TIF1 γ and Smad2 or Smad4, which did not modify the TGF- β response in simulation studies.

Finally, we integrated the TIF1 γ and FAM/UPS9x modulators into a unique model that merges all experimental observations (Figure 1C). Unlike the model depicted in Figure 1A, we considered TIF1 γ binding to Smad4 as part of a ternary complex, in which phosphorylated Smad2, Smad4 and TIF1 γ are associated in the nucleus (pS24nTIF1 γ). In this case, note that the interaction of TIF1 γ with Smad2 occurs within phosphorylated Smad2-TIF1 γ (pS2nTIF1 γ) complexes that are generated by dissociation of the ternary complexes in the nucleus. We set the same kinetic parameters for the formation/dissociation of the ternary pS24nTIF1 γ complexes and the formation/dissociation of the phosphorylated Smad2-Smad4 complexes.

Model analysis and simulation

We next performed computational experiments to investigate the dynamics of TGF- β signaling according to each model. TGF- β signaling was expressed as the amount of phosphorylated Smad2-Smad4 complexes in the nucleus (pS24n) because TGF- β target genes are regulated by these heterodimeric complexes. To explore the functional effect of TIF1 γ on the TGF- β transcriptional signal, simulation studies were performed using different concentrations of TIF1 γ varying from 0 to 50 nM, the latter corresponding to the initial concentration of Smad4 (Figure 2, Table S1). These prediction studies showed that each model was either too sensitive, with total inhibition of signaling at low concentrations of TIF1 γ according to the first model (Figure 2A), or too insensitive, with only a slight variation of signaling at higher TIF1 γ concentrations according to the second model (Figure 2B). Each predictive model hence yielded a significant mismatch with the experimental data derived from the other. The strict negative regulatory role of TIF1 γ proposed by Dupont *et al.* [4] is not compatible with the lack of sensitivity of the second model adapted from He *et al.* [7]. Similarly, He *et al.* observed a moderate TIF1 γ effect on TGF- β transcriptional activity that did not agree with the high sensitivity of the first model adapted from Dupont *et al.* In contrast, our integrative model that includes all observations yielded a graded effect of TIF1 γ on pS24n complex formation that is in agreement with the relative abundance of TIF1 γ -Smad complexes reported in both studies, leading to a graded regulation of TGF- β signaling (Figure 2C).

To further explore the robustness of our integrative model, we evaluated the sensitivity of TGF- β signaling to variations in kinetic parameters. As shown in Figure 3, varying the rate of formation (Figure 3A) or dissociation (Figure 3B) of complexes containing TIF1 γ and pS24n had little effect on TGF- β signaling. Similarly, varying the kinetic parameters for the dissociation of phosphorylated Smad2-TIF1 γ complexes (pS2nTIF1 γ) induced only few changes in the concentration of pS24n (Figure 3C). In contrast, TGF- β signaling was highly sensitive to the variation of k_{in} -Smad4ub (Figure 3D), suggesting that the export rate of ubiquitinated Smad4 is a critical component of the regulation of TGF- β transcriptional activity. In addition, the slight alteration in TGF- β signaling induced by changes in the deubiquitination rate of Smad4 (Figure 3E) disappeared with increasing concentrations of the FAM deubiquitinase (Figure 3F), suggesting that changes in FAM expression might be a sensitive marker to predict modulation of TGF- β signaling. Taken together, the results of our simulation studies reveal a new pivotal role of the Smad4 ubiquitination/

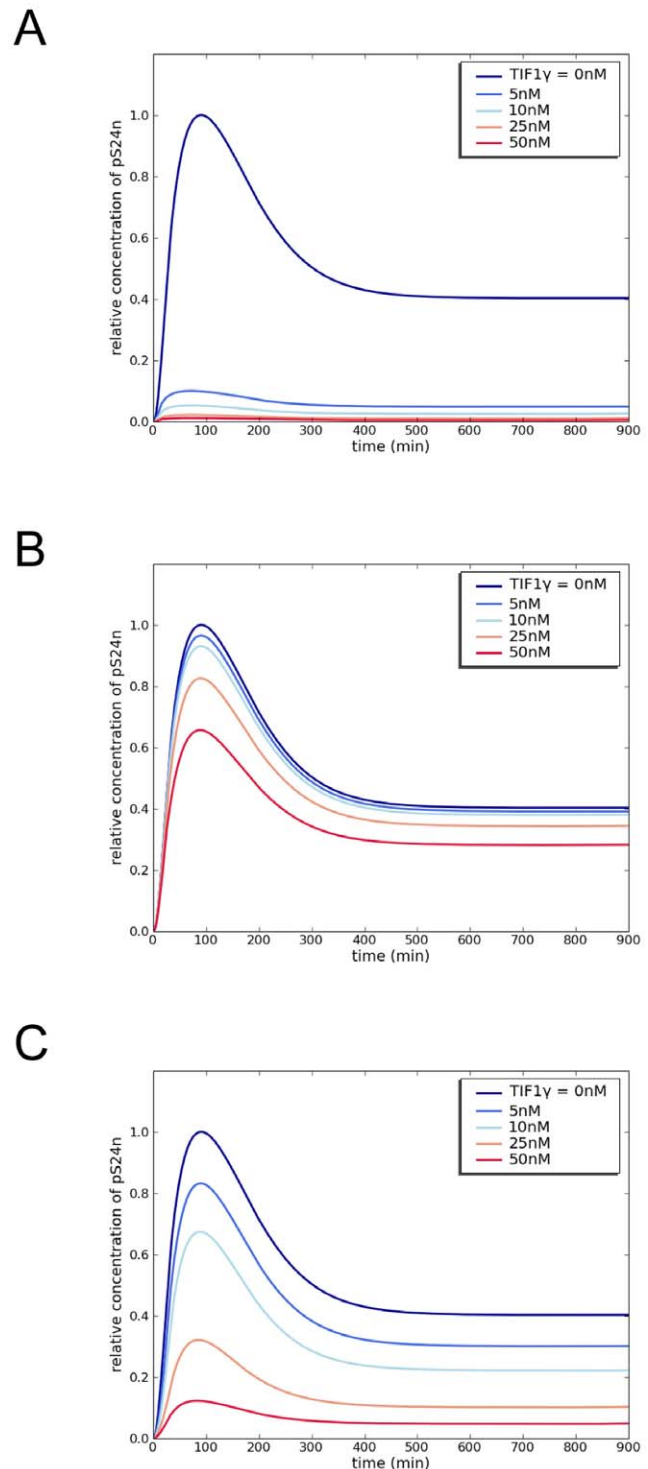


Figure 2. Effect of TIF1 γ on TGF- β signaling. Modeling analysis of the pS24n response to increasing TIF1 γ concentrations at a 10 nM TGF- β input. A) Model according to [4]; B) model according to [7] and C) integrated model. doi:10.1371/journal.pone.0033761.g002

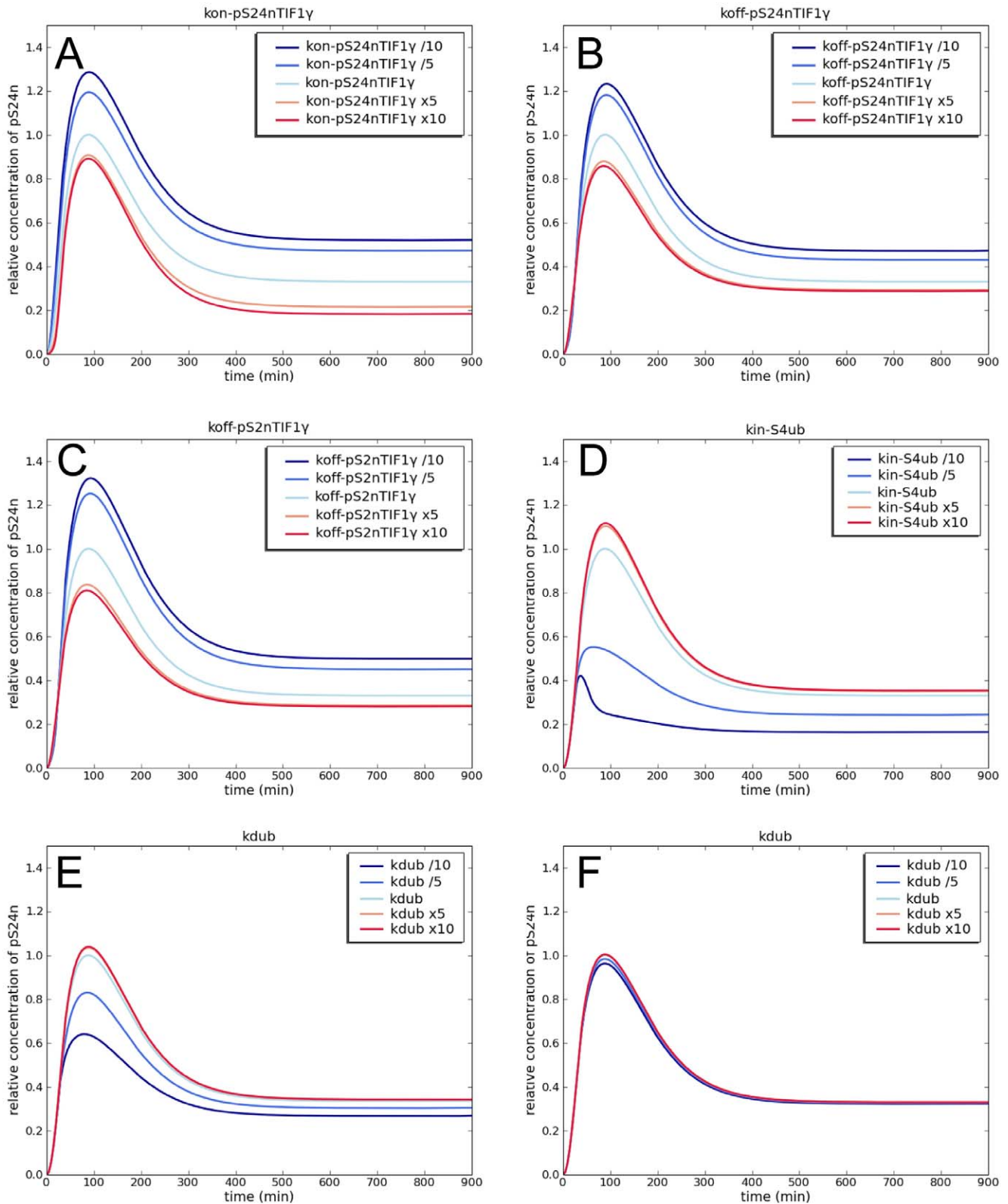


Figure 3. Parameter sensitivity analysis. Modeling analysis of pS24n response to variations of kinetic constants at a 10 nM TGF- β input. A) k_{on} -pS24nTIF1 γ , binding of TIF1 γ to phosphorylated-Smad2/Smad4 complexes; B) k_{off} -pS24nTIF1 γ , dissociation of phosphorylated-Smad2/Smad4/TIF1 γ complexes in the nucleus; C) k_{off} -pS2nTIF1 γ , dissociation of phosphorylated Smad2-TIF1 γ complexes in the nucleus; D) k_{in} -S4ub, nuclear export of ubiquitinated Smad4 in the cytoplasm; E) and F) k_{dub} , deubiquitination of Smad4 according to relative FAM concentrations of 1 nM (E) and 10 nM (F). doi:10.1371/journal.pone.0033761.g003

deubiquitination cycle in the regulation of the dynamics of TGF- β signaling. Of note is the predicted critical regulatory role of FAM in TGF- β signaling through Smad4 recycling.

Experimental validation of the model

A key component of our model is based on the hypothesis that a transient ternary complex is formed, associating Smad4, TIF1 γ and Smad2. To investigate the reality of such an interaction, we performed chromatin immunoprecipitation (ChIP) assays as previously described [12]. As shown in Figure 4, stimulation of cells with TGF- β induced the recruitment of Smad proteins on the promoter sequence of PAI-1, a TGF- β target gene. In the absence of TGF- β stimulation, TIF1 γ showed a significant association with DNA while Smad2/3 was not detected. A faint Smad4 signal could be detected under these conditions. TGF- β stimulation led to the detection of a strong Smad2/3 ChIP signal. Between 30 and 90 min of TGF- β stimulation, the association of all three proteins with DNA appears consistent with the hypothesis that a ternary complex containing Smad4, Smad2/3 and TIF1 γ transiently forms. After 120 min, Smad4 dissociated from DNA whereas Smad2/3 and TIF1 γ remained present on the PAI-1 promoter. This observation is in agreement with our hypothesis that Smad2-TIF1 γ complexes are released from the ternary complexes. Importantly, Dupont *et al.* [4], using a double-immunoprecipitation approach for TIF1 γ and Smad4, previously reported formation of these ternary complexes. More recently TIF1 γ was shown to be present at the promoter region of PAI-1 gene in uninduced cells, whereas an increase in TIF1 γ association with the Smad-binding region of the promoter was also observed upon TGF- β stimulation [22].

We next devised an experimental approach that could be used to evaluate TGF- β transcriptional activity as a function of variable TIF1 γ /Smad4 ratios. Cells were transiently transfected with siRNAs to silence Smad4 or TIF1 γ expression and were further stimulated or not with TGF- β for the indicated times (Figure 5A).

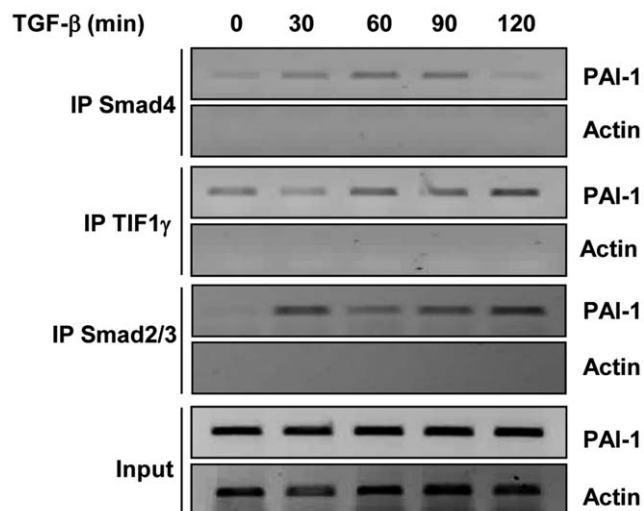


Figure 4. TIF1 γ , Smad2 and Smad4 bind to the PAI-1 promoter. ChIP assays were performed on HMEC cells treated with TGF- β for the indicated times. Cell lysates were subjected to anti-Smad4 (IP Smad4), or anti-TIF1 γ (IP TIF1 γ), or anti-Smad2/3 (IP Smad2/3) chromatin immunoprecipitation. PCR amplification of the endogenous PAI-1 promoter (733/484) was performed to detect protein bound DNA. Primers specific to actin were used as controls. doi:10.1371/journal.pone.0033761.g004

The expression of Smad4 and TIF1 γ was efficiently inhibited since no proteins were detected at day 3 post-transfection compared with cell transfected with non-targeted siRNAs (scr). The efficacy of RNA interference was confirmed at the mRNA level (Figure S1). This effect decreased with time according to siRNA availability and mRNA turnover, leading to the recovery of protein basal levels after several days (Figure 5A, upper panel). Note that silencing Smad4 and TIF1 γ affected the amounts of TIF1 γ and Smad4 proteins, respectively, detected at day 3. The time courses shown in Figure 5 finally allowed us to analyze cells containing variable amounts of endogenous Smad4 and TIF1 γ proteins. For each time point, cell extracts were used for western blot analyses and TIF1 γ /Smad ratios were evaluated by densitometric scanning of blots (Figure 5A, bottom panel). To perform this experimental verification, we quantified the mRNA levels of endogenous TGF- β target genes instead of using the over-expression of reporter genes to estimate transcriptional activities. We selected the CDH2 and CDH11 cadherin genes as they are up-regulated by TGF- β through Smad4- and TIF1 γ -dependent pathways in our cell model (Figure S2). Using the same cell extracts used for western blotting (Figure 5A), the mRNA levels of CDH2 and CDH11 were quantified and TGF- β transcriptional activity was evaluated as the ratio of mRNA levels observed in the presence or absence of TGF- β (Figure 5B). TGF- β -induced expression of CDH2 and CDH11 was correlated with the amount of Smad4 and TIF1 γ proteins. Compared to control cells (scr), low Smad4 expression (Day3) prevented TGF- β -dependent expression of CDH2 and CDH11 while the absence of TIF1 γ led to up-regulation of CDH2 and CDH11.

We then compared these experimental data with results predicted by our integrative model. As shown in Figure 5C, our observations could be fitted to the simulation curves of TGF- β transcriptional signaling, a validation reinforced by the use of physiological parameters. We conclude from these results that TIF1 γ is a new regulator that plays a pivotal role in the control of Smad4-dependent TGF- β transcriptional activity. These data also show that TIF1 γ /Smad4 ratios can determine TGF- β -dependent transcriptional activity. Accordingly, our model supports the hypothesis of fast binding of TIF1 γ to phosphorylated Smad2/Smad4 complexes and the release of both ubiquitinated Smad4 and phosphorylated Smad2-TIF1 γ complexes.

TGF- β dose- and time-dependent responses

The concentration of TGF- β in the cellular microenvironment is highly variable and its increased expression has been reported in numerous pathologies, including inflammation, fibrosis and cancer [23]. However the determination of TGF- β concentrations at the cellular level within tissues remains a difficult task since TGF- β is stocked as a latent form in the extracellular matrix [24]. In addition, its conversion from latent to biologically active forms involves numerous protease- and non protease-dependent mechanisms that differ according to cell type and the physiological context, leading to a complex non-linear delivery [25]. All previous mathematical models are based on biological data obtained from *in vitro* experiments using either TGF- β concentrations (in the nM range) or on/off signal inputs. However, Zi and al. [19] recently developed an integrative model that includes a ligand depletion parameter and demonstrated that cell-fate decision in response to TGF- β stimulation depends not only on its concentration but also on the time course of its delivery. Because we did not integrate ligand depletion in our model, response predictions were insensitive to TGF- β concentration except for concentrations as low as 0.1 nM (Figure 6A) and we routinely used concentrations of 10 nM as the TGF- β input.

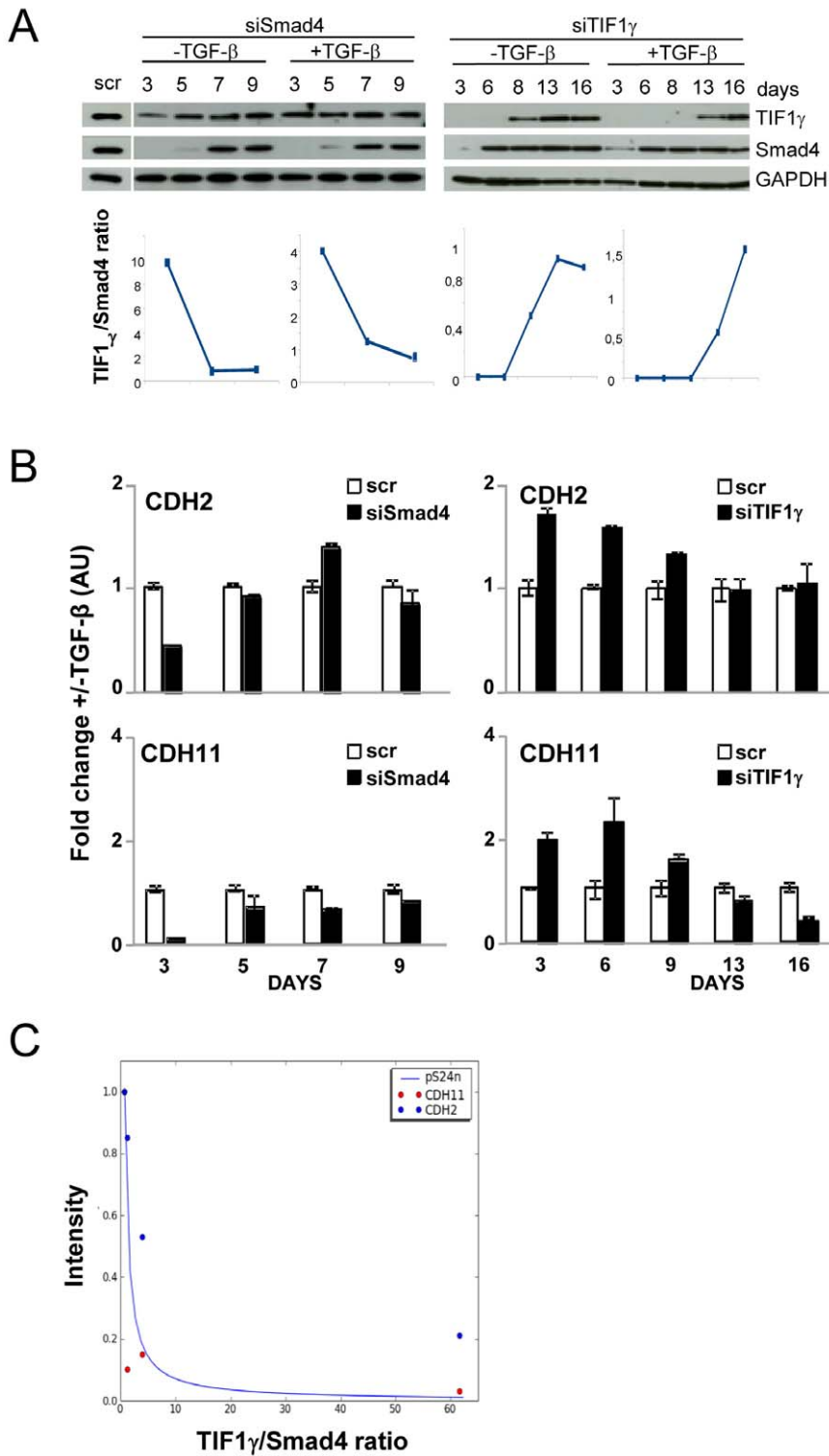


Figure 5. Expression of the CDH2 and CDH11 TGF- β target genes is sensitive to TIF1 γ /Smad4 ratios. HMEC cells were transfected with Smad4 (siSmad4) or TIF1 γ (siTIF1 γ) siRNAs and cultured in the presence (+) or absence (-) of TGF- β for the indicated times (days). Controls were cells transfected with non-targeted siRNA (scr). A) Smad4 and TIF1 γ protein levels were analyzed by immunoblotting (upper panels) and quantified by densitometric scanning (lower panels). B) TGF- β -induced fold changes in CDH2 and CDH11 expression were analyzed by RT-qPCR. All values were normalized to the amount of HPRT mRNA and expressed relative to the value obtained for TGF- β -untreated controls in arbitrary units (AU). Results are expressed as the mean \pm SD of 3 independent experiments. C) mRNA levels of CDH2 (red circles) and CDH11 (blue circles) were plotted against TIF1 γ /Smad4 ratios and were fitted to the predictive equation curve of pS24n relative concentrations. doi:10.1371/journal.pone.0033761.g005

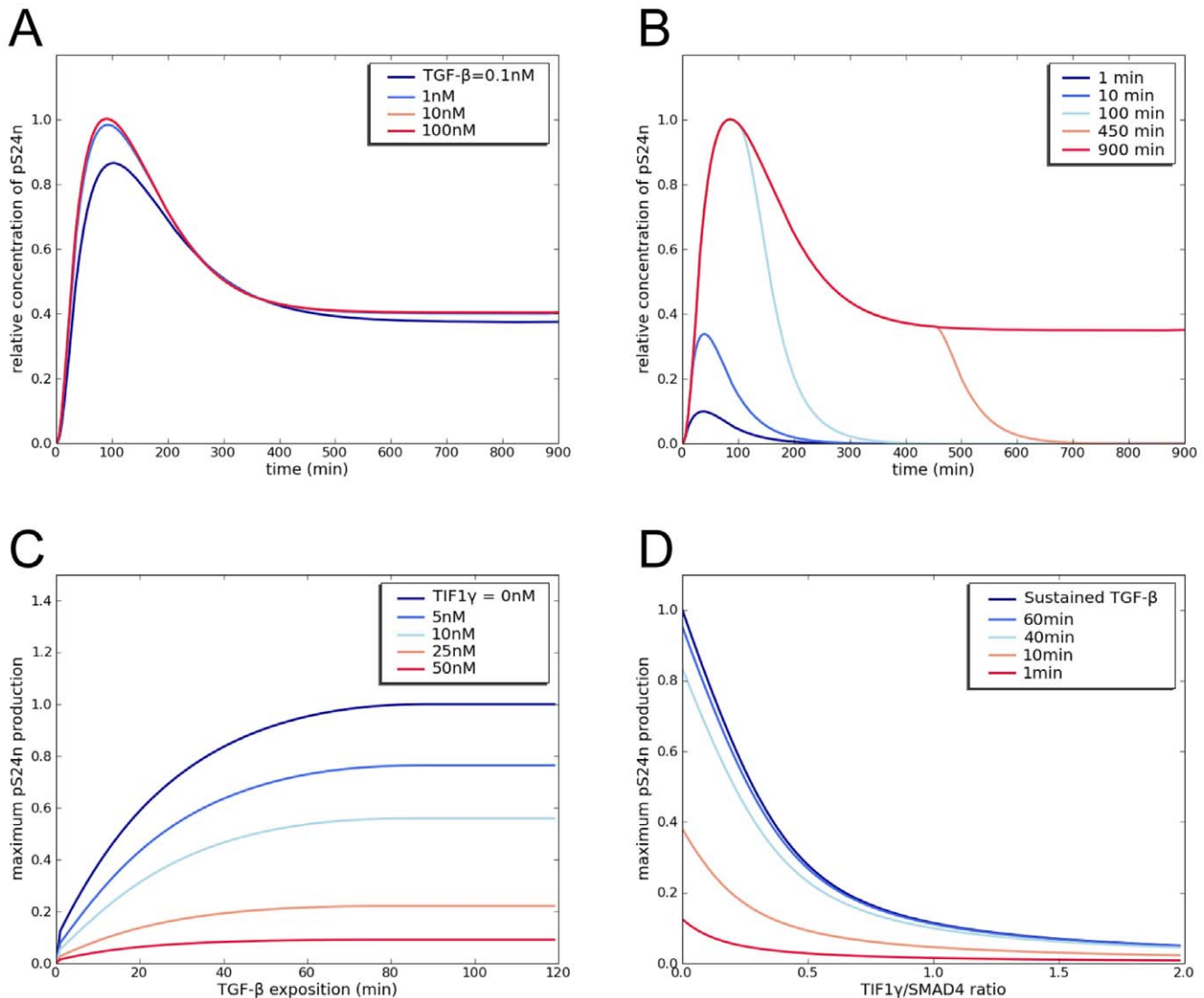


Figure 6. Concentration and time dependence of TGF- β signaling. A) and B) Modeling analysis of the pS24n response to increasing concentrations of TGF- β (A) and duration of stimulation with 10 nM TGF- β (B). C) and D) Modeling analysis of the maximum pS24n response as a function of TGF- β duration of exposure (C) or increasing TIF1 γ /Smad4 ratios (D). doi:10.1371/journal.pone.0033761.g006

When TGF- β depletion was included in our model, both graded short-term and switch-like long-term responses to TGF- β were conserved as reported by Zi *et al.* [19]. However, they were attenuated, suggesting that the presence of TIF1 γ does not affect the signal shape, but only the amplitude of TGF- β signaling (Figure S3).

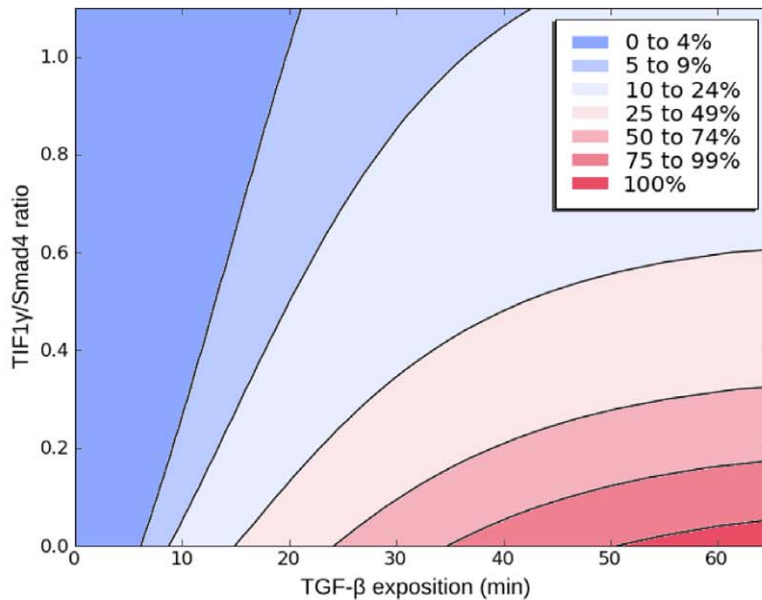
In contrast, we observed that, in our model, the length of stimulation modified the cell response. This was particularly true for short times (Figure 6B), maximum pS24n complex formation being highly dependent on TIF1 γ concentration (Figure 6C and 6D). This indicates that the magnitude of the cellular response to TGF- β depends on both TIF1 γ /Smad4 ratios and time-dependent stimulation, predicting a broad range of responses according to TGF- β cellular content and availability in the microenvironment (Figure 7A). Note that the alternative pS2TIF1 γ transcription complexes proposed by He *et al.* [7] displayed an opposite profile that required high TIF1 γ /Smad4 ratios and longer stimulation times to be fully active (Figure 7B).

In agreement with Zi *et al.* [19], our model showed that periodic short pulses of ligand stimulation yielded an outcome similar to that produced by sustained ligand stimulation, whereas an increase in the duration between pulses prevented a continuous response. These observations support the memory concept of ligand-receptor complex (LCR) activity (Figure S4A). When TIF1 γ was added, the shape of the response was similar, albeit attenuated, suggesting that, in our model, TIF1 γ does not affect LCR recycling (Figure S4B).

Conclusions

Taking into account the seemingly contradictory observations of Smad4-TIF1 γ and Smad2/3-TIF1 γ interactions, we propose an integrative model based on the formation of Smad2-Smad4-TIF1 γ ternary complexes. Validation of our hypotheses by *a posteriori* biological experiments provides strong support for our model, which shows that the TIF1 γ /Smad4 ratio serves as a regulator of TGF- β signaling that may affect determination of cell

A



B

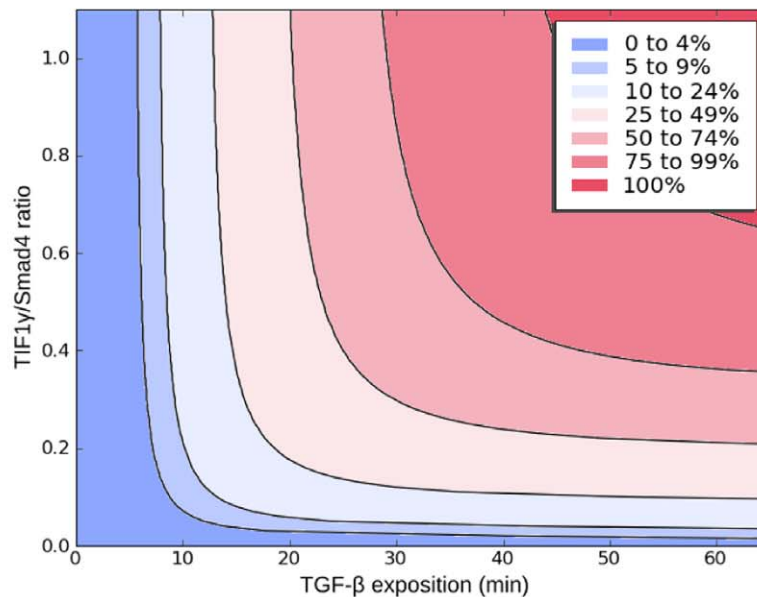


Figure 7. TGF- β time-dependent pS24n and pS2nTIF1 γ response profiles as a function of TIF1 γ /Smad4 ratios. Results are expressed as percentage of the maximum production of pS24n (A) or pS2nTIF1 γ (B).
doi:10.1371/journal.pone.0033761.g007

fate. We demonstrate that the response to TGF- β signaling is highly sensitive to TIF1 γ /Smad4 ratios, especially for short stimulation times that mediate higher threshold responses. A critical role for the TIF1 γ /Smad4 ratio in the regulation of TGF- β signaling is supported by the antagonistic role of TIF1 γ and Smad4 in the epithelio-mesenchymal cell transition [12], embryonic patterning and trophoblast stem-cell differentiation

[6], suggesting that TIF1 γ acts as a negative regulator of higher TGF- β threshold responses.

Our results emphasize the significance of TIF1 γ in orchestrating the pleiotropic effects of TGF- β signaling according to the cellular context. Its sensitivity to Smad4 levels and stimulation times suggests that TIF1 γ helps define a broad landscape of TGF- β responses. We note that Agricola *et al.* recently proposed a new

model for TIF1 γ ubiquitin ligase activity that requires binding to histones [22], thus implicating chromatin dynamics in the control of Smad localization at the promoter of TGF- β target genes. According to these results, epigenetic events contribute to the transcriptional regulation of TGF- β target genes *via* acetylation and methylation processes [26–28]. In order to understand the complexity of TGF- β -dependent gene regulation and to predict cellular responses, we believe that future models will need to integrate not only the Smad canonical pathway but also Smad-independent pathways and epigenetic events. Because of the lack of quantitative data, such an ambitious goal will require the development of different modeling-based approaches that utilize discrete models [29,30].

Supporting Information

Figure S1 Effects of Smad4 and TIF1 γ knockdown on gene expression. HMEC cells were transfected with Smad4 (siSmad4) or TIF1 γ (siTIF1 γ) siRNAs and cultured in the presence (+) or absence (–) of TGF- β for the indicated times (days). Controls were cells transfected with non-targeted siRNA (Scr). Smad4 and TIF1 γ gene expression was quantified by RT-qPCR. All values were normalized to the amount of HPRT mRNA and expressed in arbitrary units (AU). Results are expressed as the mean+SD of 3 independent experiments. (PDF)

Figure S2 Expression of the CDH2 and CDH11 is induced by TGF- β through TIF1 γ - and Smad4-dependent pathways. HMEC cells were transfected with Smad4 (siSmad4) or TIF1 γ siRNAs (si TIF1 γ) and cultured in the presence (+) or absence (–) of TGF- β for 2 days. Control cells transfected with non-targeted siRNA (Scr). CH2 and CDH11 gene expression was quantified by RT-qPCR. Results are normalized to the amount of mRNA in untreated cells and expressed as the mean+SD of 3 independent experiments. (PDF)

Figure S3 TIF1 γ does not affect short-term and switch-like long-term responses to TGF- β . TGF- β depletion was

added to the integrated model and modeling analysis of the pS24n response was performed using either increasing concentrations of TGF- β (A) or increasing concentrations of TIF1 γ in the presence of 1 nM (B) 5 nM (C) and 10 nM TGF- β (D). (PDF)

Figure S4 TIF1 γ does not modify the pS24n response to a pulsed exposure to TGF- β . Model prediction of the pS24n response in the absence (A) or presence (B) of 10 nM TIF1 γ to sustained TGF- β (10 nM) stimulation (blue curve), continuous short pulses at 30-minute intervals (green curve) or 3-hour intervals (red curve), as previously described experimentally (Zi et al 2011). We use 10 nM TIF1 γ as an average dose of tested concentrations. Concentrations up to 50 nM TIF1 γ did not modify the behavior of the signal but only reduced the signal range. (PDF)

Table S1 System parameters. (PDF)

Table S2 System of ordinary differential equations. Equations in black are from Vilar *et al.*, 2006 and Schmierer *et al.*, 2008; equations in red are estimated from biological experiments from Dupont *et al.*, 2005, 09 and He *et al.*, 2006. (PDF)

Model S1 Description of the model in Systems Biology Markup Language (SBML). (PDF)

Acknowledgments

The authors would like to thank Dr. Emmanuel Käs (LBME, CNRS/ Université Paul Sabatier) for help in writing this manuscript.

Author Contributions

Conceived and designed the experiments: GA MLB RR NT. Performed the experiments: GA LF. Analyzed the data: GA MLB RR NT. Contributed reagents/materials/analysis tools: MLB RR NT. Wrote the paper: NT.

References

1. Massague J (2008) TGFbeta in Cancer. *Cell* 134: 215–230.
2. Schmierer B, Hill CS (2007) TGFbeta-Smad signal transduction: molecular specificity and functional flexibility. *Nat Rev Mol Cell Biol* 8: 970–982.
3. Venturini L, You J, Stadler M, Galien R, Lallemand V, et al. (1999) TIF1gamma, a novel member of the transcriptional intermediary factor 1 family. *Oncogene* 18: 1209–1217.
4. Dupont S, Zucchini L, Cordenonsi M, Soligo S, Adorno M, et al. (2005) Germ-layer specification and control of cell growth by Ectoderm, a Smad4 ubiquitin ligase. *Cell* 121: 87–99.
5. Dupont S, Mamidi A, Cordenonsi M, Montagner M, Zucchini L, et al. (2009) FAM/USP9x, a deubiquitinating enzyme essential for TGFbeta signaling, controls Smad4 monoubiquitination. *Cell* 136: 123–135.
6. Morsut L, Yan KP, Enzo E, Aragona M, Soligo SM, et al. (2010) Negative control of Smad activity by ectoderm/TIF1gamma patterns the mammalian embryo. *Development* 137: 2571–2578.
7. He W, Dorn DC, Erdjument-Bromage H, Tempst P, Moore MA, et al. (2006) Hematopoiesis controlled by distinct TIF1gamma and Smad4 branches of the TGFbeta pathway. *Cell* 125: 929–941.
8. Yan KP, Dolle P, Mark M, Lerouge T, Wendling O, et al. (2004) Molecular cloning, genomic structure, and expression analysis of the mouse transcriptional intermediary factor 1 gamma gene. *Gene* 334: 3–13.
9. Vincent DF, Yan KP, Treilleux I, Gay F, Arfi V, et al. (2009) Inactivation of TIF1gamma cooperates with Kras to induce cystic tumors of the pancreas. *PLoS Genet* 5: e1000575.
10. Aucagne R, Droin N, Paggetti J, Lagrange B, Largeot A, et al. (2011) Transcription intermediary factor 1gamma is a tumor suppressor in mouse and human chronic myelomonocytic leukemia. *J Clin Invest* 121: 2361–2370.
11. Herquel B, Ouararhni K, Khetchoumian K, Ignat M, Teletin M, et al. (2011) Transcription cofactors TRIM24, TRIM28, and TRIM33 associate to form regulatory complexes that suppress murine hepatocellular carcinoma. *Proc Natl Acad Sci U S A* 108: 8212–8217.
12. Hesling C, Fattet L, Teyre G, Jury D, Gonzalo P, et al. (2011) Antagonistic regulation of EMT by TIF1gamma and Smad4 in mammary epithelial cells. *EMBO Rep* 12: 665–672.
13. Vilar JM, Jansen R, Sander C (2006) Signal processing in the TGF-beta superfamily ligand-receptor network. *PLoS Comput Biol* 2: e3.
14. Clarke DC, Betterton MD, Liu X (2006) Systems theory of Smad signalling. *Syst Biol (Stevenage)* 153: 412–424.
15. Melke P, Jonsson H, Pardali E, ten Dijke P, Peterson C (2006) A rate equation approach to elucidate the kinetics and robustness of the TGF-beta pathway. *Biophys J* 91: 4368–4380.
16. Schmierer B, Tournier AL, Bates PA, Hill CS (2008) Mathematical modeling identifies Smad nucleocytoplasmic shuttling as a dynamic signal-interpreting system. *Proc Natl Acad Sci U S A* 105: 6608–6613.
17. Nakabayashi J, Sasaki A (2009) A mathematical model of the stoichiometric control of Smad complex formation in TGF-beta signal transduction pathway. *J Theor Biol* 259: 389–403.
18. Chung SW, Miles FL, Sikes RA, Cooper CR, Farach-Carson MC, et al. (2009) Quantitative modeling and analysis of the transforming growth factor beta signaling pathway. *Biophys J* 96: 1733–1750.
19. Zi Z, Feng Z, Chapnick DA, Dahl M, Deng D, et al. (2011) Quantitative analysis of transient and sustained transforming growth factor-beta signaling dynamics. *Mol Syst Biol* 7: 492.
20. Zi Z, Klipp E (2007) Constraint-based modeling and kinetic analysis of the Smad dependent TGF-beta signaling pathway. *PLoS One* 2: e936.
21. Elenbaas B, Spirio L, Koerner F, Fleming MD, Zimonjic DB, et al. (2001) Human breast cancer cells generated by oncogenic transformation of primary mammary epithelial cells. *Genes Dev* 15: 50–65.

22. Agricola E, Randall RA, Gaarenstroom T, Dupont S, Hill CS (2011) Recruitment of TIF1 γ to chromatin via its PHD finger-bromodomain activates its ubiquitin ligase and transcriptional repressor activities. *Mol Cell* 43: 85–96.
23. Bierie B, Moses HL (2006) Tumour microenvironment: TGF β : the molecular Jekyll and Hyde of cancer. *Nat Rev Cancer* 6: 506–520.
24. Hyytiäinen M, Penttinen C, Keski-Oja J (2004) Latent TGF- β binding proteins: extracellular matrix association and roles in TGF- β activation. *Crit Rev Clin Lab Sci* 41: 233–264.
25. Annes JP, Munger JS, Rifkin DB (2003) Making sense of latent TGF β activation. *J Cell Sci* 116: 217–224.
26. Bruna A, Darken RS, Rojo F, Ocana A, Penuelas S, et al. (2007) High TGF β -Smad activity confers poor prognosis in glioma patients and promotes cell proliferation depending on the methylation of the PDGF-B gene. *Cancer Cell* 11: 147–160.
27. Barter MJ, Pybus L, Litherland GJ, Rowan AD, Clark IM, et al. (2010) HDAC-mediated control of ERK- and PI3K-dependent TGF- β -induced extracellular matrix-regulating genes. *Matrix Biol* 29: 602–612.
28. Hannigan A, Smith P, Kalna G, Lo Nigro C, Orange C, et al. (2010) Epigenetic downregulation of human disabled homolog 2 switches TGF- β from a tumor suppressor to a tumor promoter. *J Clin Invest* 120: 2842–2857.
29. Assmann SM, Albert R (2009) Discrete dynamic modeling with asynchronous update, or how to model complex systems in the absence of quantitative information. *Methods Mol Biol* 553: 207–225.
30. Sreenath SN, Cho KH, Wellstead P (2008) Modelling the dynamics of signalling pathways. *Essays Biochem* 45: 1–28.