A Relay Pathway between Arginine and Tryptophan Metabolism Confers Immunosuppressive Properties on Dendritic Cells

Highlights

- Dendritic cells (DCs) can co-express Arg1 and IDO1 immunosuppressive enzymes
- Arg1 activity is required for IDO1 induction by TGF-β in DCs
- Spermidine, a downstream Arg1 product, but not arginine starvation, induces IDO1 in DCs
- Arg1⁺ myeloid derived suppressor cells (MDSCs) can render DCs immunosuppressive via IDO1

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In Brief

Arginase 1 (Arg1) and indoleamine 2,3-dioxygenase 1 (IDO1) are immunosuppressive enzymes known to operate in distinct immune cells. Mondanelli and colleagues demonstrate that Arg1 and IDO1 cooperate in conferring long-term, immunosuppressive effects to dendritic cells.
A Relay Pathway between Arginine and Tryptophan Metabolism Confers Immunosuppressive Properties on Dendritic Cells

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SUMMARY

Arginase 1 (Arg1) and indoleamine 2,3-dioxygenase 1 (IDO1) are immunoregulatory enzymes catalyzing the degradation of L-arginine and L-tryptophan, respectively, resulting in local amino acid deprivation. In addition, unlike Arg1, IDO1 is also endowed with non-enzymatic signaling activity in dendritic cells (DCs). Despite considerable knowledge of their individual biology, no integrated functions of Arg1 and IDO1 have been reported yet. We found that IDO1 phosphorylation and consequent activation of IDO1 signaling in DCs was strictly dependent on prior expression of Arg1 and Arg1-dependent production of polyamines. Polyamines, either produced by DCs or released by bystander Arg1+ myeloid-derived suppressor cells, conditioned DCs toward an IDO1-dependent, immunosuppressive phenotype via activation of the Src kinase, which has IDO1-phosphorylating activity. Thus our data indicate that Arg1 and IDO1 are linked by an entwined pathway in immunometabolism and that their joint modulation could represent an important target for effective immunotherapy in several disease settings.

INTRODUCTION

Over the course of evolution, the catabolic pathways of L-tryptophan (Trp) and of L-arginine (Arg) have evolved to be primary regulatory nodes in the control of immune responses (Grohmann and Bronte, 2010; Murray, 2016). Trp, an essential amino acid for mammals, is a substrate for indoleamine 2,3-dioxygenase 1 (IDO1), which catalyzes the first, rate-limiting step in the kynurenine pathway, leading to Trp depletion and the production of a series of immunoregulatory molecules collectively known as kynurenines (Grohmann et al., 2003b; Mellor and Munn, 2004; Puccetti and Grohmann, 2007a). Both effects—namely, Trp starvation and kynurenine production—are involved in the conversion of naïve CD4+ T cells into Foxp3+ regulatory T (Treg) cells (Fallarino et al., 2006; Puccetti and Grohmann, 2007a). Moreover, the main IDO1 catalytic product, L-kynurenine (Kyn), has immunoregulatory effects in the absence of Trp starvation, via activation of the Aryl hydrocarbon Receptor (AhR) (Grohmann and Puccetti, 2015; Platten et al., 2012).

High IDO1 expression and catalytic activity occur in dendritic cells (DCs) in response to the cytokine interferon-γ (IFN-γ) (Grohmann et al., 2003b). In DCs stimulated with transforming growth factor β (TGF-β), IDO1 becomes instead phosphorylated in its immune-based inhibitory tyrosine motifs (ITIMs), so to mediate intracellular signaling events in a self-sustaining feedforward loop that leads to durable immunoregulatory effects (Bessede et al., 2014; Pallotta et al., 2014; Pallotta et al., 2011; Volpi et al., 2016). The composite mode of action of IDO1 may well explain its being recognized as an authentic regulator of immunity under several physiopathologic conditions (Orabona and Grohmann, 2011; Puccetti and Grohmann, 2007a).

Arg is a semi-essential amino acid, i.e., it is required by mammals only under special circumstances, such as immune responses. In immune cells, Arg is actively metabolized by arginase 1 (Arg1) or Arg2 to produce urea and L-ornithine (Orn) or it is used for protein biosynthesis (Wu and Morris, 1998). Moreover, nitric oxide synthases use Arg to make nitric oxide—a key anti-microbial gas and signaling molecule—and L-citrulline (Bronte and Zanovello, 2005). Arg consumption by Arg1, rather than Arg2 or nitric oxide synthases, represents a well-known immunoregulatory mechanism exploited by M2 macrophages (Murray, 2016; Sica and Mantovani, 2012), as well as myeloid-derived suppressor cells (MDSCs) in tumor settings (Gabrilovich and Nagaraj, 2009; Marigo et al., 2008). Conversely, the immunoregulatory
role of catalytic products via Arg1 activity (urea and Orn) has been unclear (Grohmann and Bronte, 2010). T helper 2 (Th2) cytokines, such as interleukin-4 (IL-4) and IL-13, represent efficient inducers of Arg1 expression (Gabrilovich and Nagaraj, 2009; Marigo et al., 2008; Sica and Mantovani, 2012), although TGF-β has also been shown to upregulate the enzyme, at least in rat peritoneal macrophages (Boutard et al., 1995). Besides its involvement in mouse dendritic cell (DC) differentiation (Yang et al., 2015), the biology of Arg1 in DCs is still obscure. As a whole, the bulk of the data on Arg1 and IDO1 would suggest that the two amino-acid metabolic enzymes might operate in quite distinct spatial (i.e., cells) and mechanistic modes, namely via either amino acid starvation itself (as is the case for Arg1) or via the combined effects of immunoregulatory Kyn and signaling activity (IDO1). This suggests that the enzymes have separate functions (to possibly cope with distinct environmental needs) and/or have evolved to complement, rather than integrate, each other.

DCs integrate multiple signals and mechanisms in order to promote either adaptive immunity or immune tolerance in response to specific conditions, such as distinct types of cytokinetic milieu (Banchereau and Steinman, 1998; Macagno et al., 2007). By using the main cytokines capable of inducing those metabolic enzymes (i.e., IFN-γ, IL-4, and TGF-β), we here investigated whether Arg1 and IDO1 were co-expressed in DCs and what the functional meaning—if any—of their co-expression would be. We found that (1) TGF-β, but neither IL-4, IFN-γ, nor combination thereof, will induce both Arg1 and IDO1 in DCs, with Arg1 being upregulated before IDO1; (2) Arg1 activity is absolutely required for IDO1-dependent signaling events as initiated by TGF-β; (3) a polyamine—spermidine, resulting from Orn decarboxylation—can replace TGF-β in the activation of the Src kinase that phosphorylates IDO1 and trigger immnosuppressive IDO1 signaling; and (4) DCs can be conditioned by Arg1+ MDSCs to express an IDO1-dependent immunosuppressive phenotype.

RESULTS

TGF-β Induces Co-expression of Arg1 and IDO1 in DCs
We investigated the expression and catalytic activity of Arg1 and IDO1 in splenic CD11c+ DCs stimulated with IL-4 (i.e., the main Arg1 inducer), IFN-γ, or TGF-β, both of which are IDO1 inducers. Combinations of TGF-β and IL-4 or IFN-γ were also tested. In accordance with previous data (Pallotta et al., 2011), transcript analysis at 3 and 18 hr of cytokine incubation revealed that both IFN-γ and TGF-β upregulated Ido1 expression, with IFN-γ inducing the IDO1-encoding gene to a higher extent and more rapidly than TGF-β (Figure 1A). By contrast, IL-4 did not increase Ido1, but it did induce Arg1 expression (>7-fold increase at 3 hr of stimulation). No modulatory effect could be observed for IFN-γ on Arg1 expression. In contrast, TGF-β upregulated Arg1 in a more intense fashion and with faster kinetics than Ido1. Negligible transcript expression for Arg2 and Nos2 (coding for the inducible nitric oxide synthase) could be observed in either unstimulated or TGF-β-stimulated DCs (data not shown). DC incubation with TGF-β plus IL-4 resulted in a synergistic effect on Arg1
Arg1 expression is required for IDO1 induction by TGF-β in DCs

To better appreciate the temporal relationship between Arg1 and IDO1 expression in TGF-β-stimulated DCs, we performed a more comprehensive kinetic analysis—i.e., from 0.5 to 72 hr of cytokine incubation—of transcripts coding for the two amino-acid catabolic enzymes. The results confirmed the earlier induction of Arg1 (3–4 hr; Figures 1A and 2A) as compared to IDO1 transcripts (16–18 hr; Figures 1A and 2A), as well as a higher extent of Arg1 expression that peaked at 24–48 hr (>7 fold increase as compared to unstimulated cells) of TGF-β stimulation. IDO1 expression did not increase more than 3-fold (Figure 2A).

Nevertheless, in agreement with our previous data (Pallotta et al., 2011), the extent of IDO1 expression remained stable throughout the experiment, i.e., up to 72 hr, after which DCs could no longer be analyzed because of poor viability. Conversely, Arg1 transcripts appeared to decrease from 72 hr onward.

The sequential expressions of Arg1 and IDO1 prompted us to investigate whether Arg1 was required for IDO1 induction by TGF-β in DCs. We examined IDO1 transcript amounts over time using DCs purified from the spleens of Itgax-cre;Arg1fl/fl animals, lacking Arg1 expression in CD11c+ (i.e., DCs) but not in CD11b+ cells (Figure S1). We found that Arg1 deficiency abrogated IDO1 induction by TGF-β but not by IFN-γ in DCs (Figure 2B). Analysis of IDO1 activity (Figure 2C) at 24 hr of TGF-β or IFN-γ stimulation also indicated that lack of Arg1 expression in DCs allowed IFN-γ but not TGF-β to upregulate IDO1. Thus these data identified a cytokine milieu whereby DCs exploit Arg1 expression to increase IDO1.

Orn Upregulates IDO1 in the Absence of Externally Added TGF-β

At variance with IDO1, which exerts immunoregulatory additional effects also via signaling activity (Pallotta et al., 2014; Pallotta et al., 2011; Volpi et al., 2016), Arg1-mediated mechanisms have been shown to occur only via Arg catalytic degradation. We thus investigated whether upregulation of IDO1 transcripts in DCs by TGF-β could be modulated by N-hydroxy-nor-Arg (nor-NOHA), an inhibitor of arginases. Pre-incubation with 50 μM or 100 μM nor-NOHA for 1 hr prior to the addition of
TGF-β significantly impaired the *Ido1*-inducing ability of the cytokine (Figure 3A), indicating a functional role for Arg1 catalytic activity in IDO1 modulation.

We next investigated whether Orn, an Arg1 product, was involved in *Ido1* upregulation. Incubation of DCs with Orn at 50 or 100 μM (Figure 3B) significantly increased *Ido1* expression.

**Figure 3. Orn Induces IDO1 Expression and Activity in DCs**

Kinetic analysis of *Ido1* transcripts in WT DCs stimulated with TGF-β (A and B), Orn (B and C), and/or IFN-γ (C) for different times (indicated) in the presence or absence of nor-NOHA (100 μM for IFN-γ), or medium with standard (i.e., 400 μM) or low (4–40 μM) Arg concentration (F).

Kinetic analysis of *Ido1* transcripts in WT (D, E) or CD11c<sup>−/−</sup> (E) DCs stimulated with Orn in the presence (D) or absence (E) of TGF-β receptor inhibitors. No toxic effects of inhibitors could be observed in DCs at the used concentration of 10 μM (data not shown).

In (A)–(F), data are normalized and presented as in Figure 1A.

(G) Kynurenine concentrations in supernatants of WT DCs stimulated with TGF-β in a standard medium (where not specified) or a medium with low Arg (4 μM), in the presence or absence of nor-NOHA (100 μM) or Orn (100 μM).

*p < 0.05; **p < 0.01; ***p < 0.001 (stimulated samples versus untreated controls). All data are from one experiment representative of three (means ± SD of triplicates).
similar to what observed with the cytokine alone (Figure 3B). In contrast, no modulatory effect in Ido1 expression could be observed in DCs stimulated with IFN-γ in the presence of the Arg1 inhibitor or Orn, both at 100 μM (Figure 3C). To evaluate whether autocrine and/or paracrine TGF-β could be involved in the Ido1-upregulating effects of Orn, WT DCs were incubated with the Arg metabolite for different times after a 1 hr pretreatment with an inhibitor of TGF-β receptor signaling, namely SB-431542, LY2109761, or LY2157299 (also known as galunisertib) (Figure 3D). DCs purified from transgenic CD11cΔArg mice (expressing a truncated form of TGF-β receptor II subunit in CD11c+ cells) (Laouar et al., 2005) were also assayed (Figure 3E). Both pharmacologic and genetic means of TGF-β signaling inhibition indicated that Orn effects do require autocrine and/or paracrine effects of TGF-β in both early inducing and late maintaining phase of Ido1 expression (see also Supplemental Text) and that the cytokine could be produced constitutively by DCs in basal conditions, in accordance to our previous data (Belladonna et al., 2008).

In contrast to Orn, media deficient in Arg (4 and 40 μM in the place of standard 400 μM; Figure 3F) did not significantly increased Ido1 expression. Moreover, Orn at 100 μM but not the use of an Arg-deficient medium (i.e., Arg at 4 μM) significantly increased Ido1 activity to an extent comparable to that observed with TGF-β (Figure 3G). Therefore, our data indicated that Orn, a main Arg1 product, but not Arg starvation, can condition DCs to upregulate Ido1 expression and activity if autocrine and/or paracrine TGF-β is present.

**Orn Confers Immunosuppressive Properties on DCs via Ido1 Signaling**

Ido1 immunosuppressive effects include non-enzymic functions, namely intracellular signaling events that, initiated by ITIM phosphorylation in the enzyme, are involved in reprogramming gene expression and in the induction of a stably regulatory phenotype in DCs (Orabona et al., 2012; Pallotta et al., 2011), capable of controlling T cell-mediated autoimmune responses (Pallotta et al., 2014; Volpi et al., 2016). In particular, Ido1’s ITIM phosphorylation is triggered in DCs by TGF-β via a pathway that requires phosphatidylinositol 3-kinase (PI3K) and a tyrosine kinase of the Src family (Bessede et al., 2014; Volpi et al., 2016), which phosphorylates Ido1 ITIMs. In turn, phosphorylated Ido1 ITIMs work as docking sites for tyrosine protein phosphatases, such as SHP-1 and SHP-2, which are concomitantly upregulated by TGF-β and Ido1 (Orabona et al., 2012; Pallotta et al., 2011). These events lead to the activation of an immunoregulatory signaling pathway in DCs that promotes endogenous production of TGF-β and induction of the Ido1 gene, perpetuating Ido1 signaling events and associated immunosuppressive effects over the long term.

Because Orn was capable of upregulating Ido1 in DCs in a sustained fashion comparable to that of TGF-β (Figure 3), we investigated whether Orn could replace the cytokine in activating the Ido1’s signaling and immunosuppressive effects. By means of an antibody specific for the phosphorylated form of the enzyme (Bessede et al., 2014; Pallotta et al., 2011; Volpi et al., 2016), we found that the DC incubation with Orn did induce Ido1 phosphorylation, with a peak at 15 min, i.e., an effect earlier than that triggered by TGF-β (at 30–60 min) (Figure 4A). Moreover, similarly to TGF-β, stimulation with the Arg1 product upregulated transcript expression of Ptpn6 (coding for SHP-1) and Tgfb1 at 18 hr in WT DCs. In contrast, no or less upregulation of Ptpn6 and Tgfb1, respectively, was induced by TGF-β in Itgax-cre;Arg1ΔArg DCs (Figure 4B), further confirming the important role of Arg1 in triggering and maintaining the Ido1 signaling.

To appreciate the immunosuppressive potential of DCs conditioned by Orn, we used the skin test assay, an established protocol for measuring the in vivo induction of antigen-specific
immunoreactivity versus tolerance in DCs (Grohmann et al., 2002; Grohmann et al., 2007; Pallotta et al., 2011; Puccetti et al., 1994). To this purpose, we sensitized wild-type (WT) mice with the HY peptide (containing the H-2D^b epitope of male minor transplantation antigen) presented by WT CD8^-DCs (constituting an immunostimulatory, splenic DC subset) (Grohmann et al., 2003a) administered alone or in combination with a minority fraction of the same cells (5%) purified from either WT or ido1-/- animals after conditioning with TGF-β, Orn, or medium alone for 24 hr. After priming the mice, we assessed immune reactivity at two weeks by intrafootpad challenge with the HY peptide in the absence of DCs, as described (Grohmann et al., 2002; Grohmann et al., 2007; Pallotta et al., 2011; Puccetti et al., 1994). As expected, the default priming ability of immunostimulatory DCs was not affected by the presence of untreated cells. Yet, sensitization together with TGF-β but also Orn-pre-treated WT DCs caused suppression of HY-specific reactivity, an effect not detectable in mice sensitized with ido1-/- DCs, regardless of whichever type of conditioning molecule had been used (Figure 4C).

As a whole, our data indicated that the mere incubation of DCs with an Arg metabolite can activate molecular events traceable to IDO1 signaling as well as IDO1-dependent immunosuppressive outcomes detectable in vivo.

**Orn Decarboxylation Is Required for Orn Effects Mediated by IDO1 in DCs**

Orn can be further metabolized into putrescine, a polyamine that, in turn, can be transformed into spermidine and then spermine (Figure S2), by Orn decarboxylase (ODC). In tumor cells, where it is often highly expressed, ODC favors proliferative events. More recently, ODC has been shown to be likewise involved in immunoregulatory mechanisms that oppose anti-tumor immunity (Hayes et al., 2014). As a matter of fact, administration of α-difluoromethylornithine (DFMO), an ODC inhibitor, will inhibit tumor growth via impairment of MDSC-mediated suppressive effects (Ye et al., 2016). Whether DFMO-promoted immunity might also depend on impairment of IDO1 activity in DCs is currently unknown.

To evaluate ODC expression in DCs, we conducted a meta-analysis of public microarray data restricted to the expression of the Odc1 gene in several mouse DC subsets purified from both lymphoid and nonlymphoid organs. The results showed that high amounts of Odc1 transcripts were detectable in all DC subsets analyzed so far, including splenic, conventional CD11c^+ DCs, the subject of the current study (Figure 5A). Prompted by these data, we investigated the effects of ODC inhibition on Orn actions in DCs. Because of the obligate requirement for ODC in many cell types (Cervelli et al., 2014; Guo et al., 2005), we addressed this issue by using DFMO. We found that co-incubation of splenic DCs with Orn and DFMO abrogated the capacity of Orn alone to upregulate ido1 expression (Figure 5B). Moreover, by using the skin test assay as in Figure 4C,
we found that no immunosuppressive effects could be conferred by Orn on WT DCs upon co-incubation of cells with DFMO (Figure S5C).

Because stimulation with Orn induced significant upregulation of the IDO1 gene in human DCs as well, and DFMO negated this effect (Figure S5), our data suggest that the ODC catalytic activity basally expressed in DCs exerts important immunoregulatory effects and that metabolites downstream of ODC function might represent the most proximal inducers of IDO1 signaling and IDO1-mediated immunosuppression in DCs.

Spermidine Activates the Src Kinase and Confers IDO1-Dependent, Immunosuppressive Properties in DCs

Polyamines, i.e., the diamine putrescine, the triamine spermidine, and the tetra-amine spermine, are highly bioactive polycations capable of binding nucleic acids and proteins and of modulating several signaling pathways. Polyamine functions have been studied most extensively in tumors, where they are often required for cell transformation and proliferation (Gerner and Meyskens, 2004). Whether polynomials can be produced by, or exert effects on, DCs is currently unknown.

Real-time PCR analyses revealed that DCs expressed high amounts of Smr and Smgs genes, coding for spermidine and spermine synthase, respectively, and that such expression was not modulated by either TGF-β or IL-4 (Figure S4A). To evaluate the effective production of polyamines by DCs, we assessed the profile of Arg metabolites along with that of Trp (Table S1), and that it also promotes phosphorylation of a peptide from IKKβ, a kinase involved in the activation of the pro-inflammatory NF-κB pathway (Table S4). Although less potent when considering the overall effect (Figure S4E), spermidine did promote tyrosine phosphorylation of peptides from protein kinase C (i type) and β-catenin, known substrates of Src and Fyn kinases (Table S1), respectively, but not from IKKβ (Table S4).

In a skin test assay, DC incubation with spermidine (Figure 6D) led to immunosuppressive effects detectable in vivo. Moreover, as observed for Orn (Figure 4C), spermidine effects were lost in /ido1<−/−> DCs (Figure 6D). Importantly, the immunosuppressive effects were also impaired by coincubating WT DCs with spermine and PP2 (a Src kinase inhibitor) but not PP3 (a negative control) (Figure 6D).

Overall, our data suggest that spermidine and its metabolic precursor putrescine are produced by DCs and represent major modulators of their own signaling pathways and function, including the activation of Src kinase. However, only spermidine triggered immunosuppressive IDO1 signaling, an effect possibly lost when using putrescine for DC conditioning because of the concomitant activation of several kinases other than those belonging in the Src family, including proinflammatory IKKβ, and because of the increased expression of proteasome subunits possibly degrading the IDO1 protein.

Arg1<sup>+</sup> MDSCs Condition DCs toward an IDO1-Dependent, Immunosuppressive Phenotype

As evident from their name, the defining feature of MDSCs is their ability to suppress immune cell functions. Main factors
Figure 6. Spermidine Activates IDO1 Immunosuppressive Signaling

(A) Analysis of Arg and Trp metabolite profiles in culture supernatants from DCs incubated with Orn (100 μM) for 4 or 16 hr. Results are expressed in terms of heatmap, in which each square represents the fold change in mass to charge ratio mean value of the relative metabolite as compared to time 0. Annotated targeted metabolites were derived from nontarget raw date and only represent signals of established chemical identities with 5 ppm.

(B) Real-time PCR analysis of Ido1 transcripts in DCs incubated with Orn, putrescine (Put), spermidine (Spd), or spermine (Spm; all at 20 μM) or medium alone for 24 hr. Data are normalized and presented as in Figure 1A. *p < 0.05 and **p < 0.01 (unpaired Student’s t test; Orn- or polyamine-treated versus untreated samples).

(C) Kinetics of Src phosphorylation in WT DCs incubated with Put, Spd, or Spm at 20 μM for different times. Cell lysates were analyzed by sequential immunoblotting with antibody to phosphorylated Src (pSrc), anti-Src, and β-tubulin.

(D) In vivo suppression assay with a minority fraction (5%) of WT or Ido1−/− CD8+ DCs with no conditioning (Unt) or conditioned in vitro with 20 μM spermidine in the presence or absence of PP2 or PP3 at 5 μM; analysis of skin reactivity is as in Figure 4C. **p < 0.01 and ***p < 0.001 (paired Student’s t test as in Figure 4C). Data are from one experiment representative of two (A) and three (B–D; means ± SD of triplicates in B and six samples in D). Please see also Figure S4.
implicated in MDSC-mediated immune suppression include high expression of Arg1, among others (Marvel and Gabrilovich, 2015). In addition to their inherent immunosuppressive activity, MDSCs might amplify regulatory properties of other immune cells, particularly in tumor microenvironments. Although some mechanisms underlying MDSC-macrophage interaction have been established (Ugel et al., 2015), the cross-talk between MDSCs and DCs is still unclear (Ostrand-Rosenberg et al., 2012).

Prompted by the finding that an extracellular polyamine can condition DCs toward an IDO1-dependent, immunosuppressive phenotype, we investigated whether MDSCs could also exert similar effects on DCs via the Arg1 pathway. Arg1+ MDSCs were obtained from the bone marrow of WT mice as described (Youn et al., 2008), pre-treated with nor-NOHA, DFMO (100 μM and 1 mM, respectively), or medium alone for 1 hr, extensively washed, and cocultured with DCs in separate chambers of transwells (Figure 7A). After 24 hr, DCs were recovered and analyzed for Ido1 transcript expression by real-time PCR as well as immunosuppressive potential in vivo by skin test assay as in Figure 4C and Figure 6D. Controls included DCs cultured in transwells without MDSCs (medium; Figure 7A). We found that the separate coculture with untreated MDSCs significantly upregulated Ido1 expression (Figure 7B) and conferred in vivo immunosuppressive properties on DCs (Figure 7C). However, both effects were lost when either Arg1 or ODC catalytic activity had been inhibited in MDSCs before their co-culturing with DCs. Moreover, no in vivo immunosuppressive phenotype could be acquired by
immunoregulatory functions by “simply” extending the ancient
mechanism of starvation to the control of activation and prolifer-
ation of T lymphocytes. This is even more true if one considers
that, for instance, mouse macrophages stimulated with IL-4
will upregulate Arg1 expression by 100- to 1,000-fold (Pauleau
et al., 2004). In contrast, in our study, the increase in Arg1 tran-
scripts in DCs incubated with IL-4 was no higher than 8-fold,
suggesting that the Arg1 function in DCs might be considerably
different as compared to the profound Arg starvation determined
by Arg1+ macrophages. Nevertheless, the co-presence of IL-4
and TGF-β further upregulated Arg1 transcripts more than
20-fold, suggesting that maximal expression of Arg catabolism
would require multiple signals in DCs. In contrast, IDO1, the
Trp catabolizing enzyme mainly operating in immune cells such
as DCs, has been shown to exert immunoregulatory effects via
Trp starvation but also via its catalytic products, i.e., kynure-
nines, and, perhaps most importantly, via a non-enzymic
signaling activity (Chen, 2011; Grohmann et al., 2003b; Puccetti
and Grohmann, 2007a). In fact, by means of its ITIM domains
(phosphorylatable by Src tyrosine kinases activated in the pres-
ence of TGF-β), IDO1 establishes an intracellular signaling
network in DCs that leads to the long-term expression of Ido1 it-
self and subsequent, sustained control of adaptive immunity.

At variance with Arg1, mouse Ido1, and human IDO1 as well, might
be the result of duplication of the more ancient Ido2 (IDO2 in hu-
mans) gene, an Ido1 (IDO1) paralog characterized by low effi-
ciency in Trp degradation and inability to act as a signaling mole-
cule (Orabona et al., 2012; Yuasa et al., 2007). This represents
the successful repurposing of a very ancient enzyme. Trypto-
phan 2,3-dioxygenase (TDO), another ancient enzyme mainly
expressed in the liver and responsible for the degradation of
Trp entered by diet, does not contain ITIMs.

We here showed that, in the presence of TGF-β but not IL-4,
IFN-γ, or combinations thereof, Arg1 and IDO1 expressions co-
exist in DCs. If anything, the co-presence of other cytokines
rather impaired the TGF-β upregulating action on both amino-
acid catabolizing enzymes, an effect possibly due the antago-
nistic effects of IL-4 (Musso et al., 1994) and IFN-γ (Bronte and
Zanovello, 2005) (G. Natoli, personal communication) on Ido1 and
Arg1 gene expressions, respectively. The apparently tran-
sient, yet intense, induction of Arg1 enzymatic activity by TGF-β
was mandatory for the subsequent IDO1 upregulation in terms
of both catalytic and signaling mechanisms. These Arg1 effects
were not mediated by Arg deprivation (as one might have ex-
pected, due to the poor or absent proliferative capacity of DCs)
but rather by its downstream enzymatic catabolates, namely,
the polyamine spermidine, generated downstream of decarbox-
ylation of Orn, one of the direct Arg1 catalytic products. More-
over, our data indicated that this polyamine can promote IDO1
phosphorylation and signaling events in DCs (Bessedette et al.,
2014; Pallotta et al., 2011; Volpi et al., 2016), possibly via direct
activation of the Src kinase. Spermidine might therefore repre-
sent a two-sided node responsible for an intersection between
the immunometabolic pathways of Arg1 and IDO1. Perhaps
most importantly, this relay pathway would allow the immune sys-
tem to translate an initial short-term (Arg1-mediated; typical
of early-acting MDSCs and regulatory macrophages; Goldszmid
et al., 2014) into a sustained regulatory response (via IDO1; typical
of long-term acting tolerogenic DCs; Morelli and Thomson, 2007).

TGF-β is a cytokine that appeared in metazoans and, by virtue
of its marked pleiotropy and multiple effects, plays a prominent
role in the logic of communicative networks in multicellular organ-
isms (Massagué and Comis, 2006). Although the output of a
TGF-β response is highly contextual, the presence of this cytokine
in microenvironments often favors local immunosuppression, in-
hibiting anti-tumor immunity (Gorelik and Flavell, 2002; Pickup
et al., 2013; Tu et al., 2014). The TGF-β potential for inducing both
enzymes, i.e., Arg1 in macrophages (Boutard et al., 1995) and
IDO1 in DCs (Belladonna et al., 2009; Pallotta et al., 2011)
was already known. However, we here demonstrated that the
immunoregulatory effects of TGF-β in DCs would go beyond
the mutually exclusive upregulation of the Arg1 and IDO1 enzymes
(observable in the presence of IL-4 and IFN-γ, respectively) by al-
lowing the establishment of a network involving both Arg1 and
IDO1 in DCs, further underlining the functional plasticity of these
cells. Moreover, because MDSCs themselves are an abundant
source of TGF-β production (Bierie and Moses, 2010), the
network established by the triad constituted by TGF-β, Arg1, and
IDO1 might be highly relevant to establishing potent immunono-
suppressive environments if also DCs are present as well.

In conclusion, our data, besides further underlining the impor-
tance of critical pathways linking metabolism and immunity in
multicellular animals, suggest that the appearance of a highly
evolved form of biologic response such as immune regulation
would rely on a network based on the co-option of two ancient
pathways, i.e., Arg and Trp catabolisim (Anderson et al., 2016)
as reinforced by TGF-β. A consequence of this could be that tu-
mors, considered to be the result of an evolutionary process (Billi-
aud and Santoro, 2011), have become particularly apt to co-opt
metabolic and immunosuppressive networks to propel their gen-
eration and progression. Thus our data might predict that the
simultaneous inhibition of two immune checkpoints such as
Arg1 (https://clinicaltrials.gov/show/NCT02903914) and IDO1
(Buqué et al., 2016) could represent a successful strategy in tu-
mor immunotherapy.

**EXPERIMENTAL PROCEDURES**

**Mice**

Eight- to ten-week-old female C57BL/6 mice were obtained from Charles River
Breading Laboratories. Ido1Δ+/Δ mice were purchased from The Jackson Labora-
tory and bred at Charles River. To obtain Itgax-cre;Argfl/fl mice (i.e., lacking Arg1
expression in CD11c+ cells), C57BL/6-Arg1fl/fl mice were bred to the strain
C57BL/6J-Tg (Itgax-cre,-EGFP)4097Ach/U (The Jackson Laboratory), with
inducible Cre recombinase expression in the CD11c+ cells (Figure S1). CD11ccre+ mice
on C57BL6 background (Lauar et al., 2005) were bred and maintained at
the animal facility of the University of Michigan School of Medicine.

**Isolation and Treatments of DCs and MDSCs**

Spleenic DCs were purified using CD11c MicroBeads (Miltenyi Biotec), as
described (Grohmann et al., 2002). Purity of DCs is detailed in Supplemental
Experimental Procedures. MDSCs were obtained from bone marrow cells,
as previously described (Youn et al., 2009). Details of MDSC purification and
*treatments can be found in Supplemental Experimental Procedures. For all
in vitro studies, CD11c+ or CD8+ DCs were cultured at 1 × 10⁶ cells per well
in 24-well plates in Iscove’s Modified Dulbecco’s medium (IMDM, Thermo
Fisher Scientific) or, in selected experiments, in Dulbecco’s Modified Eagle
Medium (DMEM, Thermo Fisher Scientific), containing low or standard Arg
levels (4, 40, or standard 400 μM) and completed by adding l-asparagine
(120 μM) and l-lysine (790 μM). Recombinant human TGF-β (R&D System),
Orn, and polyamines were used at the final concentration of 20 ng/ml.
50–100 μM, and 20 μM, respectively. In specific experiments, DCs were conditioned by co-culture for 24 hr with MDCSCs (either such or pretreated for 1 hr with inhibitors) using transwell cell culture inserts (Nunc).

Real-Time RT-PCR, Western Blotting, and Determination of Arg1 and IDO1 Catalytic Activity

All these procedures are detailed in Supplemental Experimental Procedures. Arg1 protein expression was investigated in DCs by immunoblot with a goat polyclonal anti-mouse Arg1 antibody (Abcam). IDO1 and pIDO1 expressions were investigated with a rabbit monoclonal anti-mouse IDO1 antibody (cv152) (Kornami et al., 2008) or a rabbit polyclonal antibody to the phosphorylated ITIM1 motif of IDO1 (Pallotta et al., 2011), respectively.

Immunization and Skin Test Assay

A skin test assay was used for measurements of major histocompatibility complex class I-restricted delayed-type hypersensitivity (DTH) responses to the HY peptide (WMHNMDLI) in C57BL/6 female recipient mice, as described (Pallotta et al., 2011). For in vivo immunization, 3 × 10⁷ peptide-loaded CD8⁺ DCs, combined with a minority fraction (5%) of peptide-loaded C57BL/6 or Ido1⁻/⁻ CD8⁺ DCs, were injected subcutaneously into recipient mice. Two weeks later, a DTH response was measured to intrafootpad challenge with the eliciting peptide, and results were expressed as the increase in footpad weight of peptide-injected footpads over that of vehicle-injected (internal control) counterparts. The minority cell fraction, constituted by WT or Ido1⁻/⁻ CD8⁺ DCs, was left untreated or treated overnight with specific reagents as above.

Meta-analysis of DC Gene Expression Data, Metabolomic Analyses, and Kinome Profiling Analyses

All these procedures are detailed in Supplemental Experimental Procedures.

Statistical Analyses

Unpaired Student’s t test was used for in vitro analyses, using at least three values from 2–3 experiments per group, whereas for the skin test assay paired Student’s t test was used (using at least six mice per group). Differences were considered significant with p < 0.05.

SUPPLEMENTAL INFORMATION

Supplemental Information includes five figures, four tables, and Supplemental Experimental Procedures and can be found with this article online at http://dx.doi.org/10.1016/j.immuni.2017.01.005.

AUTHOR CONTRIBUTIONS

G.M., C. Volpi, and U.G. designed the study. P.P., C. Volpi, and U.G. supervised the study as a whole. G.M. performed the majority of in vitro experiments and prepared most of the figures. R.B. and C.O. performed in vivo experiments. M.T.P. and A.M. supervised IDO1 signaling experiments. E.A. performed and supervised IDO1 signaling experiments. E.A. performed and supervised, respectively, metabolomics analyses. A.V. and M.P. performed and supervised, respectively, kinomic analyses. E.M.C.M. and S.B. performed and supervised, respectively, meta-analyses of Odc1 expression. U.G. wrote the manuscript.

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Supplemental Information

A Relay Pathway between Arginine and Tryptophan Metabolism Confers Immunosuppressive Properties on Dendritic Cells

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Figure S1. CD11b⁺ but not CD11c⁺ cells from Itgax-cre;Arg¹⁰/¹⁰ upregulate Arg1 expression in response to IL-4 (related to Figure 2). Real-time PCR analysis of Arg1 transcripts in CD11b⁺ and CD11c⁺ cells purified from the spleen of wild-type (WT) or Itgax-cre;Arg¹⁰/¹⁰ mice and stimulated for 3 hr with IL-4. Data were normalized to the expression of Gapdh (encoding glyceraldehyde phosphate dehydrogenase) and are presented relative to results in untreated cells. One experiment representative of three,
L-Arginine $\xrightarrow{\text{O}_2, \text{NO}}$ L-Citrulline

$\xrightarrow{\text{H}_2\text{O}, \text{Urea}}$ Arg1

L-Ornithine $\xrightarrow{\text{OAT}}$ L-Proline

$\xrightarrow{\text{ODC}}$ Putrescine

$\xrightarrow{\text{SRM}}$ Spermidine

$\xrightarrow{\text{SMS}}$ Spermine

Figure S2. Scheme of the Arg1 Pathway (related to Figure 5). The reaction catalyzed by nitric oxide synthases (NOS) is also shown. OAT, ornithine aminotransferase. ODC, ornithine decarboxylase. SRM, spermidine synthase. SMS, spermine synthase.
Figure S3. Orn Upregulates IDO1 Expression in Human DCs via ODC Catalytic Activity (related to Figure 5). DCs (5 × 10^5), obtained as described in Experimental Procedures, were exposed to 1 mM DFMO or medium alone for 1 hr prior to incubation with Orn at 100 µM or medium for 24 hr. Expression of IDO1 transcripts was evaluated by Real-Time PCR, normalized to the expression of ACTB (i.e., the human β-actin encoding gene), and presented relative to values in untreated cells (fold change = 1; dotted line). **, p < 0.01 (unpaired Student’s t test; DCs treated with DFMO, Orn, or a combination of both vs. untreated samples). One experiment representative of three.
Figure S4. Effect of Polyamines on IDO1 Protein Expression and Activation of Kinases in DCs (related to Figure 6).

(A) Real-time PCR analysis of *Srm* (coding for spermidine synthase) and *Sms* (spermine synthase) gene expression in DCs stimulated with TGF-β or IL-4 for 18 hr, normalized to the expression of *Gapdh* (encoding glyceraldehyde phosphate dehydrogenase) and presented relative to results in untreated cells (dotted line, one fold).

(B) Splenic DCs from WT mice were incubated with 20 µM putrescine (Put), spermidine (Spd), or spermine (Spm) and, after 24 hr, cell lysates were examined for IDO1 expression by means of a specific anti-mouse IDO1 antibody. β-tubulin expression was used as a loading control.

(C) Real-time PCR analysis of *Psm* 8, *Psm* 9, and *Psm* 10 (coding for β5i, β1i, and β2i immunoproteasome subunits, respectively) gene expression in DCs incubated with putrescine or spermidine for 18 hr, normalized as in (A).

(D) Splenic DCs from WT mice were incubated with 5 µg/ml cycloheximide (an inhibitor of protein synthesis) for 30 min prior to addition of 10 µM MG132 (a proteasome inhibitor), 20 µM putrescine (Put), or a combination of both. Control cells (Unt) were treated with cycloheximide and then incubated with medium alone. After 3 hr, cell lysates were examined for IDO1 expression by means of a specific anti-mouse IDO1 antibody. β-tubulin expression was used as a loading control.

(E) Average of serine, threonine, and tyrosine phosphorylation of Chip peptides in DCs incubated with putrescine (Put) or spermidine (Spd) for 15 or 30 min. Untreated cells (vehicle) were used control. One experiment representative of two (D and E) or three (A–C) is shown.
Figure S5. Co-expression of Arg1 and IDO1 in DCs and upregulation of both Arg1 and Ido1 transcripts by TGF-β in plasmacytoid DCs (related to Figure 7 and Experimental Procedures).

(A) Immunofluorescence analysis of DCs, incubated with medium alone (Unt) or stimulated with TGF-β for 24 hr and stained with anti-Arg1 (red) and anti-IDO1 (green) antibodies. Nuclei were stained with DAPI (blue).

(B) Cytofluorimetric analysis of DCs purified from the mouse spleen. Cells were stained with anti-CD11c and –MHC II (to evaluate cDC purity) or anti-B220 and –PDCA (to evaluate the contamination of pDCs) antibodies. Numbers show percentages of positive cells in the indicate gate. The fact that PDCA alone but not in combination with B220 was highly expressed in the DC population may derive from nonspecific effects of cell manipulation in vitro, as previously reported (Blasius et al., 2006).

(C) Kinetic analysis of Arg1 and Ido1 transcripts in purified pDCs from WT mice after incubation with TGF-β for different times (indicated). Data are normalized to the expression of Gapdh and presented relative to results in untreated cells (dotted line, one fold). In (A-C), one representative experiment out of four (A and B) or three (C) is shown.
Table S1. Modulation of Src-like Kinase Activity Following Stimulation with Putrescine or Spermidine (related to Fig. 6)

Kinase activity in vehicle control cultures and those stimulated with putrescine or spermidine towards Src-like kinase signal-transduction relevant undeca peptides. Cells were lysed and analysed by in vitro kinase assays as described in the Experimental Procedures section (see also Figure S4B). Results (in artificial phosphoimager units) represent the average value and standard deviation obtained from three technical replicas (three separate reactions carried out in parallel). Values are STORM phosphoimager units (24 hr exposure).

| Peptide used for kinase reaction | Phosphorylation site in protein | Upstream kinase | Vehicle | 15’putrescine | 30’putrescine | 15’spermidine | 30’spermidine |
|----------------------------------|---------------------------------|-----------------|---------|---------------|---------------|---------------|---------------|
| PCTTIYVAATE                      | SLAM<sub>Tyr307</sub>          | Fyn             | 1.0 ± 0.0·10<sup>0</sup> | 0.0           | 3.2 ± 0.0·10<sup>1</sup> | 0.0           | 1.0 ± 0.0·10<sup>0</sup> |
| EELAAYAEIRV                      | SIGLEC4A<sub>Tyr620</sub>      | Fyn             | 2.0 ± 0.0·10<sup>0</sup> | 3.0 ± 0.0·10<sup>1</sup> | 0.0           | 1.0 ± 0.0·10<sup>0</sup> | 0.0           |
| SLESLYSACSM                      | Grb10<sub>Tyr61</sub>          | Fyn             | 2.0 ± 1.0·10<sup>0</sup> | 9.9 ± 0.0·10<sup>1</sup> | 0.0           | 2.0 ± 0.0·10<sup>0</sup> | 0.0           |
| DDQEYDDVAE                       | F-YB<sub>Tyr295</sub>          | Fyn             | 1.7 ± 0.6·10<sup>0</sup> | 1.3 ± 0.7·10<sup>2</sup> | 1.5 ± 0.2·10<sup>2</sup> | 3.0 ± 0.0·10<sup>0</sup> | 2.0 ± 0.0·10<sup>0</sup> |
| KAGNYDISED                       | NR2B<sub>Tyr1259</sub>         | Fyn             | 0.0           | 1.5 ± 0.0·10<sup>2</sup> | 3.3 ± 0.0·10<sup>1</sup> | 1.0 ± 0.0·10<sup>0</sup> | 1.0 ± 0.0·10<sup>0</sup> |
| VNLINYQDDAE                      | CTNNB1<sub>Tyr142</sub>        | Fyn             | 0.0           | 0.0           | 0.0           | 0.0           | 8.0 ± 0.0·10<sup>0</sup> |
| DVLKFXDSNTV                      | PAK2<sub>Tyr130</sub>          | Hck             | 2.3 ± 0.6·10<sup>0</sup> | 3.0 ± 0.0·10<sup>2</sup> | 2.6 ± 0.4·10<sup>2</sup> | 4.0 ± 0.0·10<sup>0</sup> | 3.0 ± 1.4·10<sup>0</sup> |
| IPSVPYPAFFA                      | GRNF2<sub>Tyr322</sub>         | Hck             | 2.0 ± 0.0·10<sup>0</sup> | 1.2 ± 0.0·10<sup>2</sup> | 1.8 ± 0.2·10<sup>2</sup> | 2.0 ± 0.0·10<sup>0</sup> | 0.0           |
| DEGDEIYEDLMR                     | Va<sub>Tyr174</sub>            | Lck             | 1.0 ± 0.0·10<sup>0</sup> | 7.8 ± 0.5·10<sup>1</sup> | 8.2 ± 4.7·10<sup>1</sup> | 0.0           | 1.0 ± 0.0·10<sup>0</sup> |
| FDDPSYVQNQ                       | Shc<sub>Tyr427</sub>           | Lck             | 4.3 ± 1.5·10<sup>0</sup> | 3.2 ± 1.2·10<sup>2</sup> | 7.9 ± 0.0·10<sup>1</sup> | 2.6 ± 0.6·10<sup>0</sup> | 3.5 ± 0.7·10<sup>0</sup> |
| GGGDDYEDIIK                      | Va<sub>Tyr172</sub>            | Lck             | 2.0 ± 1.4·10<sup>0</sup> | 3.3 ± 3.8·10<sup>2</sup> | 1.3 ± 0.7·10<sup>2</sup> | 2.3 ± 0.6·10<sup>0</sup> | 2.0 ± 0.0·10<sup>0</sup> |
| TTVELYSLAER                      | PKC0<sub>Tyr90</sub>           | Lck             | 0.0           | 0.0           | 0.0           | 2.0 ± 0.0·10<sup>0</sup> | 1.0 ± 0.0·10<sup>0</sup> |
| IKEDVLYLDHH                      | PTK6<sub>Tyr342</sub>          | Lck             | 0.0           | 2.8 ± 1.2·10<sup>2</sup> | 2.4 ± 0.4·10<sup>2</sup> | 3.5 ± 0.7·10<sup>0</sup> | 3.0 ± 0.7·10<sup>0</sup> |
| LNEEWWVSYIT                      | SKP76<sub>Tyr423</sub>         | Lck             | 4.3 ± 1.0·10<sup>0</sup> | 3.1 ± 0.2·10<sup>2</sup> | 3.6 ± 0.4·10<sup>2</sup> | 4.0 ± 1.0·10<sup>0</sup> | 3.7 ± 0.6·10<sup>0</sup> |
| SRLSAYPALEG                      | CD5<sub>Tyr487</sub>           | Lck             | 1.5 ± 0.7·10<sup>0</sup> | 1.6 ± 0.4·10<sup>2</sup> | 1.5 ± 0.6·10<sup>2</sup> | 1.0 ± 0.0·10<sup>0</sup> | 2.3 ± 2.3·10<sup>0</sup> |
| ADDSSYTARSA                      | ZAP70<sub>Tyr493</sub>         | Lck             | 3.0 ± 0.0·10<sup>0</sup> | 1.5 ± 0.0·10<sup>2</sup> | 2.1 ± 0.5·10<sup>2</sup> | 3.5 ± 2.1·10<sup>0</sup> | 1.7 ± 0.6·10<sup>0</sup> |
| ASEPFFYNVFEF                     | BCR<sub>Tyr177</sub>           | Lyn             | 0.0           | 1.6 ± 0.7·10<sup>2</sup> | 1.6 ± 0.9·10<sup>2</sup> | 1.6 ± 0.6·10<sup>0</sup> | 2.0 ± 0.0·10<sup>0</sup> |
| VLDDEYVSSFG                      | TXK<sub>Tyr420</sub>           | Lyn             | 1.0 ± 0.0·10<sup>0</sup> | 1.9 ± 0.0·10<sup>2</sup> | 1.4 ± 0.0·10<sup>2</sup> | 0.0           | 1.0 ± 0.0·10<sup>0</sup> |
| FNNPASYCSVGL                     | IPPL1<sub>Tyr986</sub>         | Lyn             | 0.0           | 0.0           | 5.3 ± 0.0·10<sup>1</sup> | 1.0 ± 0.0·10<sup>0</sup> | 2.0 ± 1.0·10<sup>0</sup> |
| LGSQSYEDMRG                      | CD19<sub>Tyr531</sub>          | Lyn             | 1.5 ± 0.7·10<sup>0</sup> | 1.8 ± 1.2·10<sup>2</sup> | 1.1 ± 0.0·10<sup>2</sup> | 2.0 ± 0.0·10<sup>0</sup> | 1.5 ± 0.7·10<sup>0</sup> |
| ETNNYETAGD                       | CD32<sub>Tyr279</sub>          | Lyn             | 0.0           | 1.1 ± 0.5·10<sup>2</sup> | 0.0           | 1.5 ± 0.7·10<sup>0</sup> | 1.0 ± 0.0·10<sup>0</sup> |
| RRCKHYELLV                       | VRLC<sub>2</sub><sub>Tyr253</sub> | Lyn             | 3.7 ± 0.6·10<sup>0</sup> | 1.2 ± 0.1·10<sup>2</sup> | 1.7 ± 0.1·10<sup>2</sup> | 3.0 ± 0.7·10<sup>0</sup> | 1.7 ± 0.6·10<sup>0</sup> |
| FSSGELYGGLPP                     | NIP1<sub>Tyr264</sub>          | Lyn             | 2.0 ± 0.0·10<sup>0</sup> | 7.6 ± 3.2·10<sup>1</sup> | 1.1 ± 0.0·10<sup>2</sup> | 1.5 ± 0.7·10<sup>0</sup> | 1.0 ± 0.0·10<sup>0</sup> |
| Peptide          | Tyrosine Mutation | Src   | CD19 Ty409 | Lyn | CD19 Ty409 | Lyn | CD19 Ty409 | Lyn | CD19 Ty409 | Lyn | CD19 Ty409 | Lyn | CD19 Ty409 | Lyn | CD19 Ty409 | Lyn | CD19 Ty409 | Lyn |
|------------------|-------------------|-------|------------|-----|------------|-----|------------|-----|------------|-----|------------|-----|------------|-----|------------|-----|------------|-----|
| EQRNVYKDYRQ      |                   |       |            |     |            |     |            |     |            |     |            |     |            |     |            |     |            |     |
| FTNPVYATLYM      |                   |       |            |     |            |     |            |     |            |     |            |     |            |     |            |     |            |     |
| PKDevSkYyT       |                   |       |            |     |            |     |            |     |            |     |            |     |            |     |            |     |            |     |
| CD33 Ty340       |                   |       |            |     |            |     |            |     |            |     |            |     |            |     |            |     |            |     |
| PDK1 Ty373       |                   |       |            |     |            |     |            |     |            |     |            |     |            |     |            |     |            |     |
| TTSQYDVPi        |                   |       |            |     |            |     |            |     |            |     |            |     |            |     |            |     |            |     |
| Clathrin1 Ty1477 |                   |       |            |     |            |     |            |     |            |     |            |     |            |     |            |     |            |     |

Peptides in bold are those mentioned in the main text.
Table S2. Sequence of Primers Used in Real-Time Experiments (related to Experimental Procedures)

| Gene   | Forward primer (5’- 3’)                  | Reverse primer (5’- 3’)            |
|--------|-----------------------------------------|-----------------------------------|
| Ido1   | CGATGTTCGAAAGGTGCTGC                    | GCAGGAGAAGCTGCGATTTGC             |
| Arg1   | CAGAAGAATGGAAGAGTCAG                    | CAGATATGCAGGGAGTCACC              |
| Ptpn6  | CAGCTGCTAGGTCCAGATGAGA                  | CAGCTCAGGTACTGGTAGTGC             |
| Tgfb1  | CACAGAGAAGAACTGCTGTG                    | AGGAGCGCACAATCATGTTG              |
| Gapdh  | CTGCCAGAACATCATCCCT                     | ACTTGGCAGGTTTCTCCAGG              |
| IDO1   | TCACAGACCACAAGTCACAG                    | GCAAGACCTTACGGACATCT              |
| ACTB   | TCAGGGTGAGGATGCCTCCT                    | CTCGTCGTCGACACGGCT               |

Table S3. Complete List of Arrays Used in This Study (related to Fig. 5)
Gene expression data of DCs, obtained using Mouse Gene 1.0 ST and Mouse Genome 430 2.0 arrays, were downloaded from Gene Expression Omnibus (http://www.ncbi.nlm.nih.gov/geo) and ArrayExpress (https://www.ebi.ac.uk/arrayexpress/).

Table S4. Peptide array raw data (related to Figure 6 and Figure S4)
For peptide array analysis, we employed the Pepchip kinomics array, featuring 960 different human-only kinase substrate peptides in addition to 70 positive and negative controls, each spotted in triplicate. The chips were exposed to a phosphor screen for 72 hours, and the density of the spots was measured and analyzed by means of array software (ScanAnalyze).
Supplemental Experimental Procedures

Isolation and Treatments of DCs and MDSCs

Splenic DCs were purified using CD11c MicroBeads (Miltenyi Biotec) in the presence of EDTA to disrupt DC-T cell complexes. Cells were 90-95% CD11c+, >95% MHC I-A+, >95% B7-2+, <0.1% CD3+, and appeared to consist of 90–95% CD8−, 5–10% CD8+, and 1–5% B220+PDCA+ (i.e., plasmacytoid DCs or pDCs) cells (Grohmann et al., 2002) (Figure S5B and data not shown). Although the presence of a small percentage of pDCs (previously demonstrated to activate the IDO1 signaling in response to TGF-β (Pallotta et al., 2011)) may raise the possibility that Arg1 and IDO1 pathways are occurring in distinct DC subsets, we demonstrated by immunofluorescence that cDCs do upregulate both Arg1 and IDO1 proteins following TGF-β stimulation (Figure S5A) and, by Real-Time PCR, that also purified pDCs can upregulate the gene expression of both amino acid catabolizing enzymes (Figure S5C). DC populations were further fractionated according to CD8 expression to obtain purified CD8− DCs by means of selective MicroBeads (Miltenyi Biotec). The CD8− fraction was 45% CD4+ and typically contained <0.5% contaminating CD8+ cells.

MDSCs were obtained from bone marrow cells, harvested and plated at 1 × 10^6 cells/ml in RPMI 1640 medium (Thermo Fisher Scientific) supplemented with 10% FCS, 10 ng/ml recombinant mouse (rm) GM-CSF (Peprotech), 10 ng/ml rmIL-4 (Peprotech) plus 30% v/v of tumor cell conditioned medium (TCCM) from B16F10 tumor cells, as previously described (Youn et al., 2008). Cultures were incubated at 37°C for five days. On day 3, fresh medium with cytokines and TCCM was replaced. Cells were collected on day 5 and analyzed by flow cytometry for CD11b and Gr1 expression. Recovered cells routinely resulted > 70 % co-expressing CD11b and Gr1.

Peripheral blood mononuclear cells (PBMCs) were cultured to produce human monocyte-derived DCs as previously described (Woltman et al., 2003). Briefly, PBMCs were isolated from a buffy coat obtained from healthy donors using Ficoll-Hypaque (GE Healthcare Life Science) density gradient centrifugation and were plated at density of 2 x 10^6 cells/ml in 6-well tissue culture dishes. After 2 hr at 37°C, non-adherent cells were removed by washing twice with cold PBS. The adherent fraction was cultured in RPMI 1640 medium (Thermo Fisher Scientific) supplemented with 10% FCS, 20 ng/ml recombinant human (rh) GM-CSF (Peprotech) and 20 ng/ml rhIL-4 (Peprotech). Cultures were incubated at 37°C for seven days. On day 4, fresh medium with cytokines at the same dosages was replaced.

For all in vitro studies, CD11c+ or CD8− DCs were cultured at 1 × 10^6 cells per well in 24-well plates in Iscove’s Modified Dulbecco’s medium (IMDM, Thermo Fisher Scientific) or, in selected experiments, in
Dulbecco’s Modified Eagle Medium (DMEM, Thermo Fisher Scientific), containing low or standard Arg levels (4, 40, or standard 400 μM) and completed by adding L-asparagine (120 μM) and L-lysine (790 μM). rmIL-4 and rmIFN-γ (Peprotech) were used to stimulate DCs at the final concentration of 10 ng/ml and 100 U/ml, respectively. Arg, Orn, and polyamines (putrescine, spermidine and spermine) were from Sigma-Aldrich. The Arg1 inhibitor nor-NOHA (Bachem) was used at a final concentration of 50 or 100 μM. DFMO (Sigma-Aldrich) was used at the final concentration of 1 mM. PP2 (an Src inhibitor), and PP3 (negative control for the Src inhibitor; both from Tocris Bioscience), were used at the final concentration of 5 μM. The TGF-β receptor signaling inhibitors SB-431542, LY-2109761, and LY2157299 were purchased from Cayman Chemical and used at the final concentration of 10 μM. In specific experiments, DCs were conditioned by co-culture for 24 hr with MDSCs (either such or pretreated for 1 hr with Nor-NOHA or DFMO using transwell cell culture inserts (Nunc).

### Real-Time RT-PCR and Western Blotting

Real-Time RT-PCR (for mouse Ido1, Arg1, Ptpn6, Tgfb1, and Gapdh) analyses were carried out as described (Pallotta et al., 2011), using primers listed in Table S2. Data were calculated as the ratio of gene to Gapdh expression by the relative quantification method (ΔΔCT; means ± SD of triplicate determination), and data are presented as normalized transcript expression in the samples relative to normalized transcript expression in control cultures (in which fold change = 1; dotted line). Anti-Src, and -pSrc (Tyr416), antibodies were from Cell Signaling Technology, whereas anti–β-tubulin was from Sigma-Aldrich.

### Determination of Arg1 and IDO1 Catalytic Activity

Arg1 activity was measured in cell lysates in terms of urea production, as described (Bronte et al., 2003). The urea concentration was determined by measuring absorbance at 430 nm by means of a spectrophotometer, and then normalized for the total protein content. IDO1 functional activity was measured in culture supernatants in terms of the ability to metabolize L-tryptophan to Kyn, whose concentrations were measured by high performance liquid chromatography in 24 hr–culture supernatants after the addition of 100 μM L-tryptophan for the final 8 hr (Fallarino et al., 2003; Grohmann et al., 2007).

### Meta-analysis of Dendritic Cell Gene Expression Data

*Microarray data collection and processing.* Gene expression data of DCs, obtained using Mouse Gene 1.0 ST and Mouse Genome 430 2.0 arrays, were downloaded from Gene Expression Omnibus.
Accession numbers of samples used in this study are listed in Table S3. Microarray probe fluorescence signals of samples belonging to the same type of array were converted to expression values using the Robust Multiarray Average procedure (Irizarry et al., 2003) coded in the affy Bioconductor package. Briefly, fluorescence intensities were background-adjusted and normalized using quantile normalization, and expression values were calculated using median polish summarization and Entrez-based chip definition files for Mouse Gene 1.0 ST Array (mogene10stv1_Mm_ENTREZG version 15.0.0; 21225 custom probe sets) and for Mouse Genome 430 2.0 Array (Mouse4302_Mm_ENTREZG version 15.0.0; 17306 custom probe sets), respectively (Dai et al., 2005).

All data analyses were performed in R version 2.14.2 using Bioconductor libraries and R statistical packages.

Merging of Gene 1.0 ST and Mouse Genome 430 transcriptional data. Transcriptional data of samples hybridized on Mouse Gene 1.0 ST arrays were merged with those obtained with Mouse Genome 430 2.0 arrays matching 16547 common Entrez gene IDs, i.e., the common identifier of custom probe sets in both data sets. A direct merging of raw fluorescence signals (i.e., of CEL files), although desirable for an optimal removal of batch effects, was unfeasible due to the different probe sequences synthesized on Mouse Gene 1.0 ST and Mouse Genome 430 arrays. Thus, batch effects were removed applying ComBat to the merged matrix. ComBat was used with default parameters with the exception of the adjustment variables that were imputed as a vector of array type labels (Johnson et al., 2007).

Metabolomic Analyses

For metabolomic analysis, DCs (1 x 10⁶ cells/sample; in triplicate) were incubated in the presence or absence of Orn at 100 µM. After 4 and 16 hr, supernatants were collected to perform extraction of metabolites as described (D'Alessandro et al., 2014; Viticchio et al., 2015). Twenty µl of extract for each sample were injected into a system of ultraperformance liquid chromatography-high resolution mass spectrometry (UPLC-HRMS) (Ultimate 3000, Thermo) and running on a positive mode. Samples were loaded onto a ReproSil C18 column (2.0 mm × 150 mm, 2.5 µm — Dr. Maisch GmbH) and chromatographic separations were achieved at a temperature of 30 °C and flow rate of 0.2 ml/min. The UPLC system was coupled online with a mass spectrometer Q Exactive (Thermo) scanning in full MS mode (2 µscans) at 70,000 resolution. Calibration was performed before each analysis against positive ion mode calibration mixes (Piercenet, Thermo Fisher) to ensure sub ppm error of the intact mass. Metabolite assignments were performed using computer software (Maven), upon conversion of .raw files into .mzXML format through MassMatrix.
Kinome Profiling Analyses

DCs were lysed in ice-cold Pepchip cell lysis buffer (20 mM Tris-HCl, pH 7.5, 150 mM NaCl, 1 mM Na₂EDTA, 1 mM EGTA, 1% Triton X-100, 2.5 mM sodium pyrophosphate, 1 mM MgCl₂, 1 mM µ-glycerophosphate, 1 mM Na₃VO₄, 1 mM NaF, 1 µg/ml leupeptin, 1 µg/ml aprotinin, 1 mM PMSF). Samples were sonicated four times for 5 s each time on ice and centrifuged at 7000 × g for 10 min at 4°C. Protein content in the clear supernatant was determined with a bicinchoninic acid protein assay kit (Pierce, Rockford, IL), using bovine serum albumin (BSA) as standard, and supernatants were stored at −80°C until peptide array analysis. For peptide array analysis, we employed the Pepchip kinomics array, featuring 960 different human-only kinase substrate peptides in addition to 70 positive and negative controls, each spotted in triplicate. Cell lysates were cleared by centrifugation and peptide array incubation mix was produced by adding 10 µl of activation mix (50% glycerol, 50 µM ATP, 0.05% v/v Brij-35, and 0.25 mg/ml BSA) and 2 µl [γ-³²P] ATP (approx. 1000 kBq (Amersham AH9968). Next, the peptide array mix was added onto the chip, which was kept at 37°C in a humidified stove for 90 min. Subsequently, the peptide array was washed twice with Tris-buffered saline with Tween 20, twice in 2 M NaCl, and twice in demineralised H₂O and then air-dried. The chips were exposed to a phosphor screen for 72 hours, and the density of the spots was measured and analyzed by means of array software (ScanAnalyze). Using grid tools, spot density and individual background were corrected and spot intensities and background intensities were analyzed. Data were exported to an excel sheet (Table S4) for further analysis.
Supplementary Text

Role of TGF-βR Subunits in Arg1-Mediated IDO1 Signaling

Activation of the TGF-β receptor type II subunit (TGF-βRII) by TGF-β binding is known to be absolutely required for activation of the TGF-βRI kinase and subsequent triggering of the TGF-βR signaling, either Smad-dependent or Smad-independent, including that mediated by PI3K (Gorelik and Flavell, 2002; Pickup et al., 2013; Tu et al., 2014). We previously demonstrated that the mechanisms involved in the activation of IDO1 signaling by TGF-β are rather complex, being IDO1 phosphorylation (which happens early) Smad-independent but PI3K-dependent, whereas the later induction of SHP phosphatases is contingent on both Smad and PI3K (Pallotta et al., 2011). In order to evaluate whether autocrine/paracrine TGF-β could play a role in the effects of ornithine (Orn) in DCs, wild-type (WT) DCs were incubated with the Arg metabolite for different times after a 1-hr pretreatment with an inhibitor of TGF-β receptor signaling, namely, SB-431542 (selective for TGF-βRI or ALK5), LY2109761 (inhibiting both TGF-βRI and TGF-βRII), or LY2157299/galunisertib (targeting TGF-βRI) (Figure 3D). DCs purified from transgenic CD11<sup>cdnR</sup> mice (expressing a truncated form of TGF-βRII in CD11c<sup>+</sup> cells) (Laouar et al., 2005) were also assayed (Figure 3E). Our findings showed that inhibition of TGF-βRI (as provided by all tested TGF-βR inhibitors) is apparently more effective than functional inactivation of TGF-βRII alone (as occurs in CD11<sup>cdnR</sup> DCs). Although difficult to reconcile with the widely accepted view of TGF-βR signaling, our data may be explained by data provided by Ishigame et al. in a Listeria infection model, which suggested that CD11<sup>cdnR</sup> mice perhaps still express a functional TGF-βRII that can induce some level of TGF-β signaling (Ishigame et al., 2013). Moreover, Xavier et al reported that TGF-βRI may trigger TGF-βR signaling (both Smad and PI3K-dependent) independently from the other subunit (Xavier et al., 2009). Whatever the true mechanism/s will be, our current data do indicate that the TGF-βR signaling is involved in the Idol upregulating effects of Orn in DCs.
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