Proteomics reveals unique identities of human TGF-β-induced and thymus-derived CD4+ regulatory T cells

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The CD4+ regulatory T (Treg) cell lineage, defined by FOXP3 expression, comprises thymus-derived (t)Treg cells and peripherally induced (p)Treg cells. As a model for Treg cells, studies employ TGF-β-induced (i)Treg cells generated from CD4+ conventional T (Tconv) cells in vitro. Here, we describe how human iTreg cells relate to human blood-derived tTreg and Tconv cells according to proteomic analysis. Each of these cell populations had a unique protein expression pattern. iTreg cells had very limited overlap in protein expression with tTreg cells, regardless of cell activation status and instead shared signaling and metabolic proteins with Tconv cells. tTreg cells had a uniquely modest response to CD3/CD28-mediated stimulation. As a benchmark, we used a previously defined proteomic signature that discerns ex vivo naïve and effector Treg cells from Tconv cells and includes conserved Treg cell properties. iTreg cells largely lacked this Treg cell core signature and highly expressed e.g. STAT4 and NFATC2, which may contribute to inflammatory responses. We also used a proteomic signature that distinguishes ex vivo effector Treg cells from Tconv cells and naïve Treg cells. iTreg cells contained part of this effector Treg cell signature, suggesting acquisition of pTreg cell features. In conclusion, iTreg cells are distinct from tTreg cells and share limited features with ex vivo Treg cells at the proteomic level.

Immune responses depend on a balance between the opposing activities of conventional (Tconv) and regulatory T (Treg) cells1. While Tconv cells are key mediators of adaptive immunity, Treg cells are specialized in maintaining immune tolerance and controlling inflammation1. Based on their origin, Treg cells can be classified into two main groups. Thymus-derived (t)Treg cells are primarily directed at self-antigens and stably express the Treg cell master transcription factor FOXP3, which is indispensable for their identity and suppressive function2–6. Based on their origin, Treg cells can be classified into two main groups. Thymus-derived (t)Treg cells are primarily directed at self-antigens and stably express the Treg cell master transcription factor FOXP3, which is indispensable for their identity and suppressive function2–6. In addition, they express the transcription factor Helios (IKZF2), which is consequently used as a marker of stable Treg cells7,8. Peripherally induced (p)Treg cells are converted from Tconv cells in response to foreign antigens9 and reportedly have less stable FOXP3 expression10–11, in particular recently after their development14. Stable expression of FOXP3 is established epigenetically by demethylation of several regions within the FOXP3 gene locus, including the so-called Treg-specific demethylated region (TSDR) in the CNS2 enhancer5,15.

Based on expression of CD45RA, human Treg cells from peripheral blood can be divided into naïve (CD45RA+) and effector (CD45RA–) Treg cells16. Since conversion of Tconv cells into pTreg cells requires their activation, it stands to reason that the naïve Treg cell population is mainly composed of tTreg cells17,18. Such naïve Treg cells can differentiate into effector Treg cells in response to inflammatory signals19. Therefore, the effector Treg cell population arguably consists of a mixture of pTreg cells and activated tTreg cells. Amongst human effector Treg cells, it is currently not possible to definitively discriminate Treg cells from Tconv cells for a lack of cell surface markers20.

Naïve Treg cells occur in low abundance in the blood21,22 and effector Treg cells from the blood are not amenable to expansion in vitro19,23. For these and other reasons, studies that investigate Treg cell differentiation, function or metabolism in vitro often employ TGF-β-induced (i)Treg cells that are generated from Tconv cells26–27. Such iTreg cells resemble Treg cells by their expression of FOXP3 and their suppressive capacity28,29. However,
in terms of FOXP3 expression and suppressive function, iTreg cells are less stable than tTreg cells. For example, adoptively transferred iTreg cells in mice lost Foxp3 expression and failed to prevent graft-versus-host disease, in contrast to tTreg cells26. Similarly, others showed that iTreg cells had reduced Foxp3 expression and suppressive capacity after removal of TGF-β in vitro and that the majority of iTreg cells rapidly lost Foxp3 expression following adoptive transfer in wild-type mice30.

Despite the acquisition of Treg cell-like features, it is unclear whether iTreg cells represent an equivalent of ex vivo Treg cells. We recently described that ex vivo naive and effector Treg cells from human peripheral blood have a common proteomic signature that sets them apart from other CD4+ T cell subsets22. This Treg cell core signature reflects unique Treg cell properties associated with signal transduction, transcriptional regulation and cell metabolism. In addition, we published a proteomic signature that is unique to ex vivo effector Treg cells as opposed to Tconv cells and naive Treg cells22. Using these proteomic signatures as benchmarks, we aimed to determine to what extent in vitro-generated iTreg cells resemble ex vivo Treg cells. The recently identified cell surface molecule GPA33, which marks naïve Treg cells in human blood, enabled us to include in our comparative study highly pure iTreg cells, without using the mTOR inhibitor rapamycin11,22. In brief, our proteomic analysis shows that iTreg cells have minimal features in common with ex vivo Treg cells. We report large differences between human iTreg, tTreg and Tconv cells in terms of protein expression and biological processes, providing a proteomic resource for further research into human Treg cells.

Results

Validation of Tconv, tTreg and iTreg cell populations used in this study. To obtain the cell populations of interest, we first isolated CD4+ T cells from human peripheral blood. These cells were flow cytometrically sorted based on a CD25loCD127hiCD45RA+GPA33int phenotype to obtain naïve Tconv cells, or a CD25loCD127loCD45RA+GPA33hi phenotype to obtain naïve Treg cells, as described before11,12 (Supplementary Fig. 1). After sorting, the cells were expanded using agonistic monoclonal antibodies (mAbs) against CD3 and CD28 in the presence of IL-2. Cells derived from naïve Treg cells are hereinafter referred to as tTreg cells. To generate iTreg cells, TGF-β was added to Tconv cell cultures. After 2 weeks of expansion in vitro, cells "rested" for 4 days prior to restimulation with anti-CD3/CD28 antibodies or not for 24 h and analysis (Fig. 1a).

At this point, Tconv cells were FOXP3−Helios− and largely retained this phenotype after CD3/CD28-mediated restimulation (Fig. 1b,c). iTreg cells showed a uniform expression of FOXP3 and Helios (IKFZ2), both before and after restimulation, suggesting that the isolation and cell culture method generated pure iTreg cells (Fig. 1b,c). The iTreg cells showed bimodal FOXP3 expression, with an upregulation of FOXP3 upon CD3/CD28-mediated restimulation, while Helios was minimally expressed, as described11 (Fig. 1b,c).

The TSDR of the FOXP3 gene locus had the highly demethylated configuration in tTreg cells and was methylated in Tconv and iTreg cells, in agreement with published data32 (Fig. 1d). Next, we functionally characterized the cell populations in a standard assay, testing their ability to suppress CD3-induced proliferation of CellTrace Violet-labeled Tconv cells from peripheral blood. In this assay, tTreg cells were suppressive and more so than iTreg cells, while Tconv cells were not, in agreement with earlier research33 (Fig. 1e). Together, these data validate the purification and cell culture protocols to generate iTreg, tTreg and Tconv cell populations for further characterization.

Expression of inhibitory and stimulatory receptors on tTreg and iTreg cells. We next examined the expression of a number of important inhibitory and stimulatory cell surface receptors. Among these, CTLA-4 is a key mediator of suppression by tTreg cells13 by reducing the availability of CD80/86 for the costimulatory receptor CD28 on Tconv cells35,36. As CTLA-4 recycles between the cell surface and endosomes37, we measured both total (cell surface plus intracellular) CTLA-4 levels, as well as specifically the CTLA-4 levels on the cell surface only. Both tTreg and iTreg cells showed higher total CTLA-4 expression than Tconv cells (Fig. 2a, Supplementary Fig. 2). All three cell types upregulated total CTLA-4 protein levels upon CD3/CD28-mediated restimulation, but levels were still higher in tTreg and iTreg cells than in Tconv cells. Cell surface expression of CTLA-4 was only found on all three cell types after CD3/CD28-mediated restimulation, with iTreg cells showing the highest levels (Fig. 2b, Supplementary Fig. 2). The costimulatory receptors OX40 and GITR have received attention as regulators of Treg cell responses38–40. They were only detected on the cell surface following CD3/CD28-mediated restimulation, on all three cell types, with highest levels on iTreg cells (Fig. 2c,d, Supplementary Fig. 2). PD-1, which is described as a co-inhibitory receptor for both Tconv and tTreg cells41, was induced by CD3/CD28-mediated restimulation on both iTreg and Tconv cells, but not on tTreg cells (Fig. 2e, Supplementary Fig. 2). The cell surface enzyme CD39 (ENTPD1) facilitates extracellular ATP hydrolysis, thereby generating adenosine that has immunosuppressive effects on Tconv cells42. CD39 protein levels were highest on iTreg cells both before and after restimulation (Fig. 2f, Supplementary Fig. 2). This analysis shows that iTreg cells differ from Tconv and iTreg cells particularly in high CTLA-4 cell surface expression and lack of PD-1 expression after CD3/CD28-mediated activation, while iTreg cells are unique in high CD39 expression (Fig. 2g).

Proteomic analysis reveals that iTreg cells are distinct from tTreg and Tconv cells. We performed label-free quantitative proteomics to compare overall protein expression profiles in resting Tconv, tTreg and iTreg cells in an unbiased manner. Based on the levels of the 4285 proteins that were identified (Supplementary Table 1), principal component analysis (PCA) showed distinct grouping of Tconv, tTreg and iTreg cells (Fig. 3a). PCA1 explained most of the differences between the populations (33%). It revealed that iTreg and tTreg cells are distinct from one another and each more similar to Tconv cells. In contrast, PC2 (16%) indicated commonalities between iTreg and tTreg cells that set them apart from Tconv cells. Hierarchical clustering, based on the 475 proteins that were differentially expressed between the three cell types, also revealed marked differences
between all populations (Fig. 3b, Supplementary Table 2). Each cell population exhibited a unique pattern of protein clusters with relatively high or low expression. This analysis clustered iTreg and Tconv cells together in one branch of the dendrogram, whereas tTreg cells clustered in a different branch, showing that iTreg cells are overall more related to Tconv cells than to tTreg cells. In fact, only two small protein clusters were shared by
iTreg and tTreg cells but not Tconv cells: cluster 5 and cluster 10, which represented low or high expression of proteins compared to Tconv cells, respectively. Gene Ontology (GO) analysis indicated an overrepresentation of proteins associated with protein folding and endoplasmic reticulum stress in these two clusters (Fig. 3c, Supplementary Fig. 3). Notable was furthermore shared low expression of GIMAP proteins that are implicated in T cell homeostasis and shared low expression of YY1, which inhibits FOXP3 expression and function.

Apart from this limited overlap in protein expression between iTreg and tTreg cells, a far greater number of proteins (clusters 1–4 and 6–9) was differentially expressed between these cell types (Fig. 3b). Proteins in clusters 1–4 were expressed at lower levels in iTreg cells than in tTreg cells and were mainly associated with metabolic processes (Fig. 3d). Part of these clusters showed relatively low protein expression in both iTreg and Tconv cells (clusters 1–2), whereas the other part showed low protein expression uniquely in iTreg cells (clusters 3–4) (Supplementary Fig. 3). Noteworthy was the lower expression of 26 proteins involved in mitochondrial organization in iTreg cells compared to tTreg cells (Fig. 3d). Of these, 18 proteins were located in cluster 2, indicating that iTreg cells largely shared low expression of these proteins with Tconv cells. Clusters 6–9 consisted of proteins that were present at higher levels in iTreg cells than in tTreg cells. These proteins are mainly involved in cell cycling, immune effector processes, vesicle-mediated transport and cell adhesion (Fig. 3e). These clusters included proteins related to lymphocyte activation, such as MHC class II, STAT transcription factors, and signaling molecules.

**Figure 2.** Expression of characteristic cell surface molecules associated with Treg cells. (a–f) Quantification of flow cytometric analysis of the protein expression of total CTLA-4 (cell surface plus intracellular) (n = 13) (a), surface CTLA-4 (n = 8) (b), surface OX40 (n = 8) (c), surface GITR (n = 8) (d), surface PD-1 (n = 8) (e) and surface CD39 (ENTPD1) (n = 8) (f) using indicated cell populations. Mean fluorescence intensity (MFI) is depicted on the y-axis. Two-way ANOVA with Tukey’s post hoc test was used for statistical analysis. Data are presented as mean ± SEM. Sample size (n) represents cells from individual donors, analyzed in independent experiments. Asterisks without lines indicate comparisons within cell types following restimulation. *p < 0.05, **p < 0.01, ***p < 0.001, ****p < 0.0001. (g) Table summarizing the data in (a–f), indicating relatively low (−), intermediate (+/−) or high (+) expression levels.
Figure 3. The iTreg, tTreg and Tconv cell proteomes are distinct. (a) PCA plot of the proteome of unstimulated Tconv, iTreg and tTreg cells (n = 3). (b) Heat map showing hierarchical clustering of the 475 proteins that were differentially expressed between unstimulated Tconv, iTreg and tTreg cells (ANOVA, FDR < 0.05). Samples are colored according to cell type as in (a). Z-scores showing relative protein expression values are color-coded. Numbered boxes (1–10) indicate clusters of proteins. (c) STRING network of proteins that have shared low (cluster 5) or high (cluster 10) expression in iTreg and tTreg cells, relative to Tconv cells. Proteins with high or low expression are depicted as yellow or blue nodes, respectively. Protein associations according to STRING are connected with lines. (d) GO enrichment analysis of clusters 1–4 containing proteins with low expression in iTreg cells compared to tTreg cells, showing ten significantly enriched GO biological processes. For example, proteins associated with mitochondrion organization are shown in a STRING network. Blue nodes indicate low expression in iTreg cells relative to tTreg cells. (e) GO enrichment analysis of clusters 6–9 containing proteins with high expression in iTreg cells compared to tTreg cells, showing ten significantly enriched GO biological processes. For example, proteins involved in immune effector processes are displayed in a STRING network. Yellow nodes indicate high expression in iTreg cells relative to tTreg cells.
such as CBL, VAV1, GRB2, MAPK1 and PAK1 (Fig. 3e). High expression of these proteins was partly shared by iTreg and Tconv cells (clusters 6–8) and partly unique for iTreg cells (cluster 9) (Supplementary Fig. 3). This analysis indicates that iTreg cells display a protein expression profile that largely sets them apart from tTreg cells and indicates similarity with Tconv cells particularly in metabolic features and potential responses to activation signals. However, iTreg cells do share some features with tTreg cells, in particular regarding endoplasmic reticulum-associated processes.

We also analyzed the proteomes of Tconv, tTreg and iTreg cells following CD3/CD28-mediated restimulation (Supplementary Table 1). Distinct clusters were identified by PCA, showing that particularly Tconv and iTreg cells altered their protein expression upon activation (Supplementary Fig. 4a). A much smaller response was observed in tTreg cells, as these samples clustered adjacent to each other regardless of stimulation. We performed
A core signature reflecting unique Treg cell properties is present in tTreg but not iTreg cells. Recently, we performed proteomic analysis of Treg cells that were isolated to high purity in the naive state from human peripheral blood and therefore very likely represent Treg cells. In that study, we defined a proteomic signature shared between these naïve Treg cells and effector phenotype Treg cells that distinguishes them from CD4+ Tconv cells, which we termed the Treg cell core signature22. This signature defines characteristic protein expression in expanded Tconv, tTreg and iTreg cells. Samples are colored according to cell type as in (c). Z-scores depicting relative protein expression values are color-coded. Numbered boxes (1–5) indicate clusters of proteins. For reference, protein names are colored according to high (yellow) or low (blue) expression in the Treg cell core signature as shown in (a). Significantly differential expression (FDR < 0.05) is indicated by asterisks and bold letters. Proteins are marked (#) when expression in in vitro-expanded Treg cells from the present dataset deviates from the core signature derived from ex vivo Treg cells22. (e) Quantification of flow cytometric analysis of intracellular IFN-γ and IL-2 protein expression in indicated PMA/ionomycin-treated cell populations from the same donors (n = 6). Percentage of IFN-γ+ or IL-2+ cells is shown on the y-axis. One-way ANOVA with Tukey’s post hoc test was used for statistical analysis. (f) Quantification of flow cytometric analysis of the protein expression of total NFATC2 (n = 4), surface ITGA4 (CD49d) (n = 5) and surface ICOS (n = 5) on indicated cell populations. MFI is depicted on the y-axis. Two-way ANOVA with Tukey’s post hoc test was used for statistical analysis. Data are presented as mean ± SEM. Sample size (n) represents cells from individual donors, analyzed in independent experiments.

\[ p < 0.05, \ * p < 0.01, \ ** p < 0.001, \ *** p < 0.0001. \]

For instance, iTreg cells expressed high levels of the transcription factor STAT4, while low levels in Treg cells are critical to conserve Treg cell function23. Low expression of transcription factor NFATC2 is another feature of Treg cells22, but iTreg cells shared relatively high expression of this protein with Tconv cells. STAT4 and NFATC2 are mediators of pro-inflammatory cytokine signaling and regulate IFNG and IL2 gene expression, respectively24,25. We observed that iTreg cells but not Treg cells were able to produce IFN-γ and IL-2 following PMA/ionomycin-mediated stimulation, suggesting that iTreg cells lack anti-inflammatory characteristics of Treg cells (Fig. 4e, Supplementary Fig. 5a). We furthermore confirmed differential expression of core signature proteins NFATC2, ITGA4 and ICOS between iTreg and Treg cells by flow cytometry, both before and after CD3/CD28-mediated restimulation (Fig. 4f, Supplementary Fig. 5b).

Since iTreg cells responded strongly to CD3/CD28-mediated restimulation in terms of altered protein expression (Supplementary Fig. 4a,b), we examined whether they obtained more of the Treg cell core signature under these conditions. The core signature is present in naïve and effector iTreg cells ex vivo, and therefore, the expression of core signature proteins remains rather stable in iTreg cells upon activation22. This was indeed observed in restimulated iTreg cells, and interestingly, levels of their own core signature proteins were also largely maintained in iTreg or Tconv cells upon restimulation (Supplementary Fig. 4c, Supplementary Table 3). Thus, iTreg cells did not have characteristic expression of many Treg cell core signature proteins, regardless of whether the cells were resting or had been restimulated via CD3/CD28. These data suggest that iTreg cells share only a part of the unique characteristics of ex vivo Treg cells.

iTreg cells have aspects of an effector Treg cell signature. We previously also defined an effector Treg cell signature, composed of proteins that set ex vivo effector Treg cells apart from naïve Treg cells and Tconv cells22. The 39 lowly expressed proteins of this signature (effector Treg) primarily include those associ-
phenotype that was not majorly affected by ATRA or butyrate. Addition of rapamycin to TGF-β and ATRA cells generated by the more traditional protocol used here.

Fig. 6). Altogether, the majority of effector Treghi signature proteins was expressed at relatively high levels in iTreg cells. Thus, the effector Treg cell signature is partially present in iTreg cells, indicating that iTreg cells resemble ex vivo effector Treg cells to some extent.

Discussion

The Treg cell lineage can be subdivided into tTreg and pTreg cells based on developmental origin. iTreg cells have escaped negative selection in the thymus and are mainly geared towards self-antigen recognition. They comprise the majority of Treg cells in vivo and control systemic and tissue-specific autoimmunity. pTreg cells recognize foreign antigen and are described to reside at mucosal surfaces and the maternal–fetal interface.

Using the cell surface marker Nrp1, pTreg cells can be distinguished from Treg cells in mice, where common and unique features in their phenotypes and transcriptome profiles have been observed. tTreg and pTreg cells complement each other in immune homeostasis, which may in part be due to their distinct TCR repertoires and differential expression of transcriptional regulators. In human, effector phenotype tTreg or pTreg cells cannot be discriminated for lack of surface markers. Because pTreg cells cannot be isolated ex vivo, TGF-β-driven iTreg cells are often employed as an in vitro model for Treg cell function, differentiation or metabolism.

Upon their generation from Tconv cells, iTreg cells acquire Treg cell-like features such as expression of FOXP3 and CTLA-4. However, FOXP3 expression in iTreg cells is heterogeneous, as observed by us and others. Stable FOXP3 expression is established at the epigenetic level by demethylation of several regions within the FOXP3 gene locus. A critical region for sustained FOXP3 expression is the TSDR in the CNS2 enhancer, which we confirmed to be highly demethylated in iTreg cells but not in tTreg cells, as described. TGF-β can induce FOXP3 expression in activated Tconv cells via the CNS1 enhancer, which is reportedly less stable. Natural compounds including all-trans retinoic acid (ATRA), a vitamin A metabolite, and microbial product butyrate can enhance FOXP3 expression. In vivo, these compounds are involved in pTreg cell generation, particularly in the gut. Schmidt et al. reported that addition of ATRA or butyrate to TGF-β increased the frequency of human FOXP3-expressing cells, but nevertheless did not improve CTLA-4 expression and suppressive function. It was recently reported for murine Treg cells that deprivation of CD28 costimulation led to increased DNA demethylation within Treg cell-specific genes, even including Foxp3 CNS2 demethylation, giving rise to more stable Foxp3 expression and enhanced suppressive function. It will therefore be of interest to generate human iTreg cells in absence of CD28 costimulation and to determine whether such cells resemble tTreg cells better than the iTreg cells generated by the more traditional protocol used here.

In a comprehensive proteomic analysis, Schmidt et al. found TGF-β to be the key driver of the iTreg cell phenotype that was not majorly affected by ATRA or butyrate. Addition of rapamycin to TGF-β and ATRA generated iTreg cells with a unique protein expression profile and enhanced suppressive activity in vitro, but these iTreg cells were not suppressive in vivo. Given these data, we here used iTreg cells induced by TGF-β only and compared them to highly pure GPA33hi tTreg cells and Tconv cells, using several benchmarks of ex vivo Treg cells as previously defined by proteomics. Proteomics gives more direct insight into the functional cellular programs compared to transcriptomics due to differential regulation of protein and mRNA levels, as also shown for Treg cells.

iTreg cells proved to display a protein expression profile that markedly sets them apart from Treg cells. The proteins that were differentially expressed between iTreg and Treg cells were enriched in metabolic regulators (Fig. 3d). Studies on T-cell metabolism have used iTreg cells as representative of ex vivo Treg cells and described certain metabolic features that would be unique to Treg cells as compared to CD4+ Tconv cells. The clear distinction in expression of metabolic regulators between iTreg and Treg cells underlines that these cell populations should not be equated in metabolic studies. iTreg cells furthermore differed from tTreg cells in high expression of signaling proteins involved in lymphocyte activation. This result predicts a divergent response of iTreg versus tTreg cells to extracellular stimuli. Based on the proteomic Treg cell core signature, we have previously
Figure 5. Aspects of an effector Treg cell signature are present in iTreg cells. (a,b) The effector Treg cell signature represents 39 lowly and 41 highly expressed proteins in ex vivo effector Treg cells compared to naïve Treg cells and other CD4+ T cells from human blood. Shown are STRING networks displaying the lowly and highly expressed proteins as blue and yellow nodes, respectively. (c,d) Heat maps showing hierarchical clustering of the lowly (c) or highly (d) expressed proteins of the effector Treg cell signature based on their expression in unstimulated Tconv, tTreg and iTreg cells (n=3). Samples are colored according to cell type. Z-scores showing relative protein expression values are color-coded. Numbered boxes indicate clusters of proteins. Significantly differential expression is indicated by asterisks and bold letters (FDR < 0.05). CCR4, HLA-DRA and FAS (CD95) are marked in (d), as these molecules are also analyzed by flow cytometry in (e). (e) Quantification of flow cytometric analysis of the protein expression of surface CCR4 (n=4), surface HLA-DR (n=4) and surface FAS (n=4) on indicated cell populations. MFI is depicted on the y-axis. Two-way ANOVA with Tukey’s post hoc test was used for statistical analysis. Data are presented as mean ± SEM. Sample size (n) represents cells from individual donors, analyzed in independent experiments. ***p < 0.001, ****p < 0.0001.
reported that iTreg cells have adaptations in signaling pathways downstream of immune receptors such as the TCR, CD28 and cytokine receptors. In particular, low expression of signaling molecules in iTreg cells may lead to attenuated responses to pro-inflammatory cytokines, as we demonstrated before for STAT4. The high expression of multiple signaling molecules in iTreg cells, together with proteins involved in the cell cycle indicates a more activated and responsive state of iTreg cells compared to tTreg cells. Accordingly, we observed that CD3/CD28-mediated restimulation induced a strong response in iTreg and Tconv cells. In contrast, tTreg cells had a strikingly modest response to this stimulus. The proteomic similarities between iTreg and tTreg cells were limited. GO analysis primarily indicated a similar enrichment of proteins involved in endoplasmic reticulum-associated processes. The transcription factor Eos (Ikzf4) is associated with controlling Treg cell function and phenotype and interacts directly with Foxp3 to mediate gene silencing in murine Treg cells. Eos is also reported to be highly expressed in human iTreg cells, which we confirmed by proteomics and flow cytometry (Supplementary Fig. 7). We furthermore observed shared low expression of transcription factor YY1 by proteins, which can also interact with FOXP3 to regulate transcription. Helios was not expressed at similar high levels in iTreg cells as observed in tTreg cells, but minor upregulation was found compared to Tconv cells. It has been demonstrated in mice that, after recent egress from the thymus, a small proportion of naïve CD4+ Tconv cells can develop into Helios+ pTreg cells. Nevertheless, Helios is preferentially expressed in iTreg cells and marks their high expression of involved mediators, such as STAT4 and NFATC2.

Upon activation, iTreg cells can travel from secondary lymphoid organs through the blood to peripheral tissues, where they experience environmental signals. They can adapt themselves to locally resident effector T cells by responding to specific cytokines. This involves the gain of lineage-determining transcription factors such as GATA-3 or T-bet in addition to FOXP3, which alters their functionality. Regardless of their activation status, tTreg cells retain core features that safeguard their suppressive nature and prevent them from exerting pro-inflammatory functions. To establish whether in vitro generated iTreg cells acquired these features, we examined to what extent they exhibited the characteristic proteomic signatures from human Treg cells that were analyzed directly from peripheral blood. The first benchmark we used was the Treg cell core signature, a conserved set of proteins between naïve and effector Treg cells compared to other CD4+ T cell subsets that emphasizes critical Treg cell aspects. This signature strongly suggests differential wiring of signaling pathways in Treg and Tconv cells, given the constitutive differential expression of key signal transduction components between these cell types. We observed that iTreg cells only share a limited part of these conserved characteristics, while the signature was largely present in the in vitro-expanded tTreg cells studied. Noteworthy, expression of the core signature proteins remained comparable upon CD3/CD28-mediated restimulation in all cell types studied, supporting that the signature reflects essential Treg cell properties. Thus, iTreg cells do not resemble Treg cells found in peripheral blood. Since pTreg cells cannot be isolated ex vivo due to lack of cell surface markers, it cannot be directly assessed to what extent iTreg cells share features with pTreg cells, whose TGF-β-induced features they are thought to reproduce.

Based on the differential expression of signaling molecules, it is likely that iTreg cells show an enhanced response to pro-inflammatory cytokines compared to tTreg cells. For example, the Treg cell core signature includes low expression of STAT4 and NFATC2, which are mediators of pro-inflammatory cytokine signaling. Low expression of these molecules may aid iTreg cells to protect their anti-inflammatory nature in a pro-inflammatory environment, while they can still respond to pro-inflammatory cytokines through other pathways. We have previously shown that deliberate elevation of STAT4 expression in iTreg cells increases their susceptibility to destabilization by pro-inflammatory cytokines, provoking effector cytokine production and loss of FOXP3 expression. Both STAT4 and NFATC2 had relatively high expression in iTreg cells compared to tTreg cells, suggesting that iTreg cells may respond to pro-inflammatory cues in a pro-inflammatory manner, e.g. by IFN-γ production. iTreg cells were shown to be less stable after transfer into inflamed mice than iTreg cells. This has been attributed to their fragile epigenetic stability, but our findings now suggest that this may also be caused by their high expression of involved mediators, such as STAT4 and NFATC2.

The second benchmark we examined was an effector Treg cell signature, which sets effector Treg cells apart from naïve Treg cells and other CD4+ T cell subsets. Relative to protein expression in iTreg and Tconv cells, iTreg cells acquired aspects of the effector Treg cell signature, indicating that iTreg cells may resemble ex vivo effector Treg cells to some extent. Since iTreg cells are thought to mimic pTreg cells because of TGF-β-mediated induction, the enrichment of the effector Treg cell signature in iTreg cells may fit with the idea that the effector Treg cell population in peripheral blood contains both activated iTreg cells and pTreg cells.

In conclusion, our global proteomic analysis showed that iTreg and tTreg cells are distinct and each share more protein expression with Tconv cells than with one another, providing a resource for further research into human Treg cells. Moreover, iTreg cells only partially have the protein expression features of ex vivo effector Treg cells. These aspects limit their utility as an in vitro model for Treg cells.

**Methods**

### Cell isolation and flow cytometric sorting

Human materials were obtained in compliance with the Declaration of Helsinki and the Dutch rules regarding the use of human materials from voluntary donors. Buffy coats from healthy anonymized male donors were obtained after their written informed consent, as approved by the Ethical Advisory Council of Sanquin Blood Supply Foundation, Plesmanlaan 125, Amsterdam, the Netherlands. Total CD4+ T cells were collected from buffy coats by first performing a Ficoll-Paque Plus (GE Healthcare) density gradient centrifugation to isolate PBMCs and subsequently using CD4 magnetic MicroBeads (Miltenyi Biotec) according to the manufacturers’ protocols. Alternatively, total CD4+ T cells were directly isolated from buffy coats using the StraightFrom Buffy Coat CD4 MicroBead kit (Miltenyi Biotec) according to the manufacturer’s instructions. To sort naïve Tconv and (Treg cells, CD4+ T cells were stained with CD4-PE-Cy7, CD127-
added. From day 7 to day 14, cells were cultured in 24-well plates (Greiner) at 5 × 10^5 cells per ml, while fresh mAbs against CD3 (clone CLB-T3/4.E, IgE isotype, Sanquin, 0.1 μg/ml) and CD28 (clone CLB-CD28/1, Sanquin, 0.2 μg/ml) in solution, added on day 0 and day 7 during expansion cultures. On day 4, fresh T-cell medium was added. From day 7 to day 14, cells were cultured in 24-well plates (Greiner) at 5 × 10^5 cells per ml, while fresh T-cell medium was added on day 11. On day 14, cells were cultured in 6-well plates (Corning) at 1 × 10^6 cells per ml for 4 days using fresh T-cell medium in the absence of agonistic mAbs. To generate iTreg cells, naïve Tconv cells were cultured as described above in the presence of TGF-β (Peprotech, 10 ng/ml). Prior to restimulation experiments, dead cells were removed by Ficolc-Paque Plus density gradient centrifugation. T cell restimulation was performed where indicated, using agonistic mAbs in the presence of agonistic mAbs against CD3 (0.1 μg/ml) and CD28 (0.2 μg/ml), but without TGF-β. For cytokine production assays, resting T cells were cultured for 5 h in presence of GolgiPlug (BD Biosciences, 1:1000 dilution), with or without PMA (5 ng/ml) and ionomycin (750 ng/ml).

**Flow cytometry.** For expression analysis of cell surface molecules, T cells were washed in PBS/1% FCS and stained using the following mAbs in appropriate combinations: anti-CD4-FITC (Beckman Coulter, hereinafter referred to as T-cell medium, at 37 °C and 5% CO₂. Cells were expanded using agonistic mAbs against CD3 (clone CLB-T3/4.E, IgE isotype, Sanquin, 0.1 μg/ml) and CD28 (clone CLB-CD28/1, Sanquin, 0.2 μg/ml) in solution. Fresh T-cell medium was added. From day 7 to day 14, cells were cultured in 24-well plates (Greiner) at 5 × 10^5 cells per ml, while fresh T-cell medium was added on day 11. On day 14, cells were cultured in 6-well plates (Corning) at 1 × 10^6 cells per ml for 4 days using fresh T-cell medium in the absence of agonistic mAbs. To generate iTreg cells, naïve Tconv cells were cultured as described above in the presence of TGF-β (Peprotech, 10 ng/ml). Prior to restimulation experiments, dead cells were removed by Ficolc-Paque Plus density gradient centrifugation. T cell restimulation was performed where indicated, using agonistic mAbs in the presence of agonistic mAbs against CD3 (0.1 μg/ml) and CD28 (0.2 μg/ml), but without TGF-β. For cytokine production assays, resting T cells were cultured for 5 h in presence of GolgiPlug (BD Biosciences, 1:1000 dilution), with or without PMA (5 ng/ml) and ionomycin (750 ng/ml).

**Flow cytometry.** For expression analysis of cell surface molecules, T cells were washed in PBS/1% FCS and stained using the following mAbs in appropriate combinations: anti-CD340-Pe-Cy7 (BioLegend), anti-CD3-APC-Cy7 (BioLegend), anti-CD399-BV510 (BioLegend), anti-CD44-PE-Cy7 (BioLegend), anti-CD45RA-FITC (Immunotools) and GPA33-AF647 mAbs (BioLegend) in culture during restimulation for 24 h. Near-IR dead cell stain kit (Invitrogen) was used as a live/dead marker. For intracellular staining, cells were fixed and permeabilized using the FOXP3 transcription factor staining buffer set (Invitrogen) according to the manufacturer’s instructions. Next, cells were stained with combinations of anti-FOX3-APC (Invitrogen), anti-Helixos-PE-Cy7, anti-CTLA-4-PE-Dazzle594, anti-EOS-PE-DR-APC-eFluor 780 (Invitrogen), anti-HLA-DR-BV605 (BD Biosciences) and anti-FAS-BB700 (CD95) (BD Biosciences), anti-ICOS-PerCP-Cy5.5 (BioLegend), anti-CCR4-PE-Cy7 (BioLegend), anti-HLA-DR-APC-eFluor 780 (Invitrogen), anti-HLA-DR-BV605 (BD Biosciences) and anti-FAS-BB700 (CD95) (BD Biosciences). Cell surface expression of CTLA-4 was measured by adding anti-CTLA-4-PE-Dazze594 mAb (BioLegend) in culture during restimulation for 24 h. Near-IR dead cell stain kit (Invitrogen) was used as a live/dead marker. For intracellular staining, cells were fixed and permeabilized using the FOXP3 transcription factor staining buffer set (Invitrogen) according to the manufacturer’s instructions. Next, cells were stained with combinations of anti-FOX3-APC (Invitrogen), anti-Helixos-PE-Cy7, anti-CTLA-4-PE-Dazzle594, anti-EOS-PE (BioLegend) and anti-NEATC2-AF488 (Cell Signaling Technology) mAbs. Anti-IL-2-PerCP-Cy5.5 (BioLegend) and anti-IFN-γ-BV421 (BD Biosciences) mAbs were used for intracellular cytokine staining. Flow cytometry was performed using a BD LSR Fortessa or BD LSR II cell analyzer (BD Biosciences). Sample acquisition was performed using FACSDiva software version 8 and data were analyzed using FlowJo software version 10.6.0.

**Methylation assay of the TSDR.** The methylation status of the TSDR within the FOXP3 locus was determined in Tconv, tTreg and iTreg cells as described previously. Only male donors were used for these analyses, given the X-chromosomal location of the FOXP3 locus. In brief, expanded T cells were resuspended in PBS for proteinase K digestion, followed by bisulfite conversion of DNA using the EZ DNA Methylation-Direct kit (Zymo Research) and methylation-specific quantitative PCR using iQTM SYBR Green Supermix (Bio-Rad). Methylation-specific primers were 5'-CGATAGGGTAGTTAGTTGAGCC-3' and 5'-GGTACTAAGCCCAGGA-3'. Demethylation-specific primers were 5'-TAGGTTAGTTGAGCCGAGTA-3' and 5'-CCATTTACATCAACACACGAA-3'. Melt curve analysis was performed on a LightCycler® 480-II (Roche). Methylation of the TSDR (%) was calculated using the following formula: 100/(1 + 2^Ct[CG] − Ct[TG]), where Ct[CG] is defined as Ct values obtained using methylation-specific primers and Ct[TG] is defined as Ct values obtained using demethylation-specific primers.

**Suppression assay.** Total human PBMCs were washed and resuspended in PBS, followed by incubation for 8 min using 5 μM CellTrace Violet dye (Invitrogen). Following fluorescent labeling, an equal volume of FCS was added and cells were subsequently washed twice in IMDM supplemented with 8% FCS. CellTrace Violet-labeled PBMCs were cocultured with expanded Tconv, tTreg or iTreg cells for 96 h at indicated ratios, in the presence of anti-CD3 mAb (0.05 μg/ml). Proliferation of PBMCs was examined by measuring CellTrace Violet dye dilution by flow cytometry on a BD LSR Fortessa or BD LSR II cell analyzer. Data were analyzed using FlowJo software (version 10.6.0).

**Sample preparation for mass spectrometry.** Following T-cell expansion, at least 1 × 10^6 resting or restimulated Tconv, tTreg and iTreg cells were washed in PBS and lysed in 100 mM Tris HCl pH 8.0 with 4% SDS and 100 mM DTT, as described previously. Samples were heated at 95 °C, sonicated (Bioruptor) and centrifuged (16,000 g), followed by isolation of cell lysates and storage at −80 °C. Cell lysates were processed using filter-aided sample preparation (FASP) as described. In short, proteins were alkylated and digested into peptides using sequencing-grade Trypsin (Promega). Samples were desalted using StageTips, the aqueous phase was evaporated in a speedvac, and proteins were reconstituted in 2% acetonitrile in 0.1% TFA in water before analysis by mass spectrometry.

**Mass spectrometry data acquisition.** As described previously, tryptic peptides were separated by nanoscale C18 reverse chromatography coupled on line to an Orbitrap Fusion Tribrid mass spectrometer...
Cytoscape. The largest gene set clusters were displayed using manually assigned annotations to summarize the differentially regulated following restimulation of T cells, the EnrichmentMap app (version 3.3.2) was used in using default parameters and gene set permutations. To identify related groups of biological processes that were analyzed for each cell type using GO biological processes in GSEA software (version 4.1.0). GSEA was performed enriched genes. STRING was used to visualize protein networks. The proteomic response to restimulation was heat maps. Protein clusters were subjected to overrepresentation analysis to assess enrichment of GO biological processes, a redundancy cutoff was applied when significant GO terms contained large overlap based on statistical heat maps. Protein clusters were subjected to overrepresentation analysis to assess enrichment of GO biological processes, a redundancy cutoff was applied when significant GO terms contained large overlap based on statistical analyses were performed as indicated in the figure legends.  Log2 transformation was performed when data were not normally distributed. Data are presented as mean ± SEM and a two-sided p<0.05 was considered statistically significant.

Mass spectrometry data processing and analysis. RAW mass spectrometry files were processed as previously described25 with the MaxQuant computational platform25 (version 1.5.2.8) using label-free quantitation (LFQ). Peptides were identified using the Andromeda search engine by querying the human UniProt database with a 1% false discovery rate (FDR) cutoff both at peptide and protein level. Potential contaminants and reverse hits were removed using Perseus (version 1.6.12). LFQ values were log2-transformed and the biological replicates were grouped based on cell type and treatment. For analysis of unstimulated samples, proteins with a minimum of 3 valid values in at least one cell type were included for further analysis (Supplementary Table 1). For analysis of samples with or without CD3/CD28-mediated restimulation, a minimum of 3 valid values in at least one sample group was required to include proteins for further analysis (Supplementary Table 1). Missing value imputation was performed separately for each sample, replacing missing values by random numbers drawn from the lower part of the normal distribution (width = 0.3, shift = 1.8). The proteomics experiment was conducted with three biological replicates per sample group, but one sample, i.e. restimulated iTreg cells from donor 1, was excluded from the analysis due to lower protein content that led to fewer detected proteins. PCA revealed that this sample was an outlier.

Proteomics data were analyzed using Qlucore Omics Explorer (version 3.8) to perform PCA and ANOVA test followed by a Benjamini–Hochberg multiple testing correction with a 5% FDR. After z-score normalization, clusters of proteins with similar expression patterns were identified using hierarchical clustering and visualized in heat maps. Protein clusters were subjected to overrepresentation analysis to assess enrichment of GO biological processes using the STRING app (version 1.6.0) in Cytoscape (version 3.9.0). To refine the analysis of biological processes, a redundancy cutoff was applied when significant GO terms contained large overlap based on enriched genes. STRING was used to visualize protein networks. The proteomic response to restimulation was analyzed for each cell type using GO biological processes in GSEA software (version 4.1.0). GSEA was performed using default parameters and gene set permutations. To identify related groups of biological processes that were differentially regulated following restimulation of T cells, the EnrichmentMap app (version 3.3.2) was used in Cytoscape. The largest gene set clusters were displayed using manually assigned annotations to summarize the biological processes represented by the clusters.

Statistical analysis. Data were analyzed using GraphPad Prism (version 9.0.1), except for proteomics data. Statistical analyses were performed as indicated in the figure legends. Log2 transformation was performed when data were not normally distributed. Data are presented as mean ± SEM and a two-sided p < 0.05 was considered statistically significant.

Data availability Proteomics data on iTreg and Tconv cells have been deposited to the ProteomeXchange Consortium (http://proteomecentral.proteomexchange.org) via the PRIDE repository23 with the dataset identifier PXD007745, as outlined before23. Proteomics data on iTreg cells are also available via ProteomeXchange with the dataset identifier PXD038302.

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M.M., S.d.K. and J.B. designed the study and wrote the manuscript, M.M. and S.d.K. performed experiments and analyzed the data, E.S. provided technical assistance, E.C. provided data and insights, D.A. provided conceptual advice in the study design and contributed to the manuscript.

Competing interests
The authors declare no competing interests.

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