

RESEARCH ARTICLE

Inhibition of Phosphoinositide 3-Kinase p110delta Does Not Affect T Cell Driven Development of Type 1 Diabetes Despite Significant Effects on Cytokine Production

Ariana Barbera Betancourt¹, Juliet L. Emery², Asha Recino¹, F. Susan Wong³, Anne Cooke¹, Klaus Okkenhaug², Maja Wallberg^{1*}

1 Department of Pathology, University of Cambridge, Tennis Court Road, CB2 1QP Cambridge, United Kingdom, **2** Laboratory of Lymphocyte Signalling and Development, Babraham Institute, Cambridge CB22 3AT, United Kingdom, **3** Diabetes Research Group, Institute of Molecular and Experimental Medicine, Cardiff School of Medicine, Cardiff University, Cardiff CF14 4XN, United Kingdom

* mw394@cam.ac.uk



OPEN ACCESS

Citation: Barbera Betancourt A, Emery JL, Recino A, Wong FS, Cooke A, Okkenhaug K, et al. (2016) Inhibition of Phosphoinositide 3-Kinase p110delta Does Not Affect T Cell Driven Development of Type 1 Diabetes Despite Significant Effects on Cytokine Production. PLoS ONE 11(1): e0146516. doi:10.1371/journal.pone.0146516

Editor: Dongmei Li, University of Rochester, UNITED STATES

Received: October 13, 2015

Accepted: December 18, 2015

Published: January 19, 2016

Copyright: © 2016 Barbera Betancourt et al. This is an open access article distributed under the terms of the [Creative Commons Attribution License](https://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Data Availability Statement: All relevant data are within the paper and its Supporting Information files.

Funding: This work was supported by Grant number 09/0003840, Diabetes UK <https://www.diabetes.org.uk/> (MW); Grant number 5-2006-229, Juvenile Diabetes Research Foundation <https://www.jdrf.org.uk/> (KO); Grant number BBS/E/B/0000C236 Biotechnology and Biological Sciences Research Council (KO); Grant number health-f5-2009-241883 European Research Council 7th Frame Programme http://ec.europa.eu/research/fp7/index_en.cfm (AC);

Abstract

Type 1 diabetes is caused by the destruction of insulin producing beta cells by the immune system. The p110δ isoform of PI3K is expressed primarily in cells of haematopoietic origin and the catalytic activity of p110δ is important for the activation of these cells. Targeting of this pathway offers an opportunity to reduce immune cell activity without unwanted side effects. We have explored the effects of a specific p110δ isoform inhibitor, IC87114, on diabetogenic T cells both *in vitro* and *in vivo*, and find that although pharmacological inhibition of p110δ has a considerable impact on the production of pro-inflammatory cytokines, it does not delay the onset of diabetes after adoptive transfer of diabetogenic cells. Further, we demonstrate that combination treatment with CTLA4-Ig does not improve the efficacy of treatment, but instead attenuates the protective effects seen with CTLA4-Ig treatment alone. Our results suggest that decreased IL-10 production by Foxp3⁺ CD4⁺ T cells in the presence of IC87114 negates individual anti-inflammatory effects of IC87114 and CTLA4-Ig.

Introduction

Type 1 diabetes is an autoimmune inflammatory disease that is caused by immune cell mediated destruction of the insulin-producing beta cells in the pancreas. T cells are assumed to play a considerable role in the pathogenesis of this disease which comes from the demonstration that its strongest genetic risk is conferred by the HLA locus and other loci affecting the biology of these cells [1, 2]. Islet specific CD4⁺ and CD8⁺ cells mediate diabetogenesis in NOD mice [3] and have been identified in human type 1 diabetes patients [4–6]. Other cell types including B cells, dendritic cells and macrophages are also necessary for the initiation of the anti-islet immune responses (reviewed in [7]). Once the beta cell pool is destroyed by the immune system, insufficient amounts of insulin are produced to sustain glucose uptake into insulin

and Grant number 02BX12ACYD, the Britain Israel Research and Academic Exchange Partnership (BIRAX) <http://www.britishcouncil.org/en/programmes/science/birax>. The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

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

dependent cells, notably muscle cells, causing wasting due to protein breakdown, even though high levels of glucose are present in the blood. The disease was fatal until Banting and Best discovered insulin in 1922 [8]. Although much has been learned about type 1 diabetes since then, no protocols for sustained cure or prevention have been discovered. Strategies attempting to abrogate immune responses have been successful in mouse models [7], and some have also resulted in delay of beta cell destruction in patients, such as the anti-CD3 treatment clinical trials [9, 10] and CTLA-4 Ig [11].

CTLA-4 Ig prevents the activation of T cells by inhibiting interaction with the co-stimulatory molecule CD28. However, CD28 blockade alone is unlikely to be sufficient to prevent autoimmunity due to the fact that memory and CD8⁺ T cells are not as dependent as naïve CD4⁺ T cells on CD28 costimulation. As none of the clinical trials have resulted in long term reversal of diabetes, it has been suggested that a combination of treatments targeting various stages of immune activation may be more successful [12].

A widely explored approach is the inhibition of key signalling enzymes involved in the activation and metabolism of immune cells. PI3Ks constitute a family of enzymes involved in cellular functions such as cell growth, proliferation, differentiation, motility, survival and intracellular trafficking [13]. The class I PI3Ks catalyse the final step to PIP3 by phosphorylating PI(4,5)P2 to become PI(3,4,5)P3. PIP3 activates Akt, which in turn leads to activation of mTOR and inhibition of Foxo. Of the four class I PI3K subunits, PI3K p110 δ is uniquely expressed in immune cells [14]. Specific inhibition of this subunit therefore primarily affects cells of the immune system, leaving other cell types unchanged. p110 δ inhibitors such as IC87114 have been used to inhibit p110 δ activity *in vitro* and *in vivo*. It has been shown that IC87114 inhibits TCR-induced cytokine production by both naïve and effector T cells [15]. In addition to its effects on T cells, p110 δ inhibitors also target other immune cells such as B cells, NK and myeloid cells (reviewed in [14, 16]). The typical outcome of PI3K δ inactivation is a reduction but not a complete abrogation of leukocyte functions, probably due to the overlapping functions with other PI3K members that are also present in leukocytes. IC87114 reduced disease severity in preclinical rodent models of rheumatoid arthritis, asthma, and allergy [17–19]. The PI3K δ inhibitor idelalisib was approved in 2014 for the treatment of chronic lymphocytic leukaemia.

CD28 and p110 δ act synergistically to provide full T-cell stimulation [20], and we hypothesised that dual inhibition of CD28 and p110 δ may achieve more potent alleviation of pathologic immune responses than can be achieved with either inhibitor alone. To test this hypothesis, we used IC and CTLA-4 Ig, either alone or in combination, to inhibit signalling via p110 δ , CD28 or both in naïve and activated T cells from non obese diabetic (NOD) mice and TCR transgenic mice on a NOD background, and monitored proliferation and cytokine production. In addition, we have investigated whether such a combination could limit the development of autoimmune diabetes in the NOD mouse model after the transfer of either naïve or activated islet reactive BDC2.5 CD4⁺ T cells.

Materials and Methods

Mice

Female NOD, NOD-*scid*, BDC2.5 NOD mice [21], NOD-CD2-GFP mice [22], G9C8 TCR transgenic NOD mice [23, 24] and NOD-Foxp3-GFP [25] were bred in the Department of Pathology, University of Cambridge and maintained under specific pathogen-free conditions. The mice are housed in individually ventilated cages with free access to standard chow and water. The facility is kept on a 12 hour light, 12 hour dark cycle. The humane endpoints for these experiments specify that any mouse that loses more than 15% of its body weight (compared to healthy littermates), or in other ways looks unwell and likely to exceed the Home

Office standard of moderate severity must be culled. However, no mice used in this study required early culling. At the end the experiments, mice were culled using a CO₂-chamber followed by dislocation of the neck. In cases where we were harvesting islets for transplantation it was important to maintain an intact bile duct, and death was instead confirmed through palpation of the chest to assess the absence of a heart beat.

Ethics statement

This study was carried out in strict accordance with U.K. Home Office project licence regulations (Project Licence number 80/2442) after approval by the Ethical Review Committee of the University of Cambridge.

PI3K- δ inhibitor

The PI3K p110 δ inhibitor IC87114 was synthesized by Jonathan Clark (Babraham Institute) as described (D030 from patent WO 01/81346) [15]. For *in vivo* administration, IC87114 was dissolved in methyl cellulose 400 cps (Sigma) using a sonicator (Heat Systems Ultrasonics), and administered through oral gavage twice daily in 100 μ l at a dose of 30mg/kg body weight. This dose was chosen based on previous reports of its efficacy *in vivo* [17]. In our hands, a 30 mg/kg by gavage achieves \sim 2 μ M 90 min post-administration and the drug is cleared from the blood 4–7 hours post administration. IC87114 is selective for p110 δ at plasma concentrations of 5 μ M [17].

CTLA4-Ig

CTLA4-Ig (Abatacept) was provided by Bristol Myers Squibb (BMS). CTLA4-Ig was administered by intraperitoneal (ip) injection starting on day 0 with 500 μ g, then 250 μ g every other day [26]. For *in vitro* assays, CTLA4-Ig was added to cultures at 100 ng/ml.

Th1 differentiation for *in vitro* studies and adoptive transfer

CD4⁺CD25⁻ T cells (for *in vitro* studies) or CD4⁺CD62L^{hi} CD25⁻B220⁻ T cells (for adoptive transfer) were isolated by cell sorter from 5-week-old BDC2.5 TCR transgenic NOD mice and differentiated into Th1 cells by culturing them with plate bound anti-CD3 (2 μ g/mL), soluble anti-CD28 (10 μ g/mL), IL-2 (100u/ml), IL-12 (10ng/ml) and IFN- γ (100u/ml) for 4 days at 37°C with 5% CO₂. Afterwards, the production of IFN- γ was checked by specific ELISA (R&D).

T cell activation for functional assays

Cells were isolated from spleen and lymph nodes and cultured in IMDM with 10% fetal calf serum, 1% penicillin-streptomycin, and β -mercaptoethanol. 2x10⁵ total lymphocytes were stimulated as appropriate (see below) for 3 days in the presence or absence of rising concentrations of IC (0.6, 1.25, 2.5, 5 and 10 μ g/mL) at 37°C with 5% CO₂. NOD mouse cells were stimulated with plate bound anti-CD3 (2 μ g/mL) and soluble anti-CD28 (10 μ g/mL), whereas cells isolated from BDC2.5 or G9C8TCR transgenic NOD mice were stimulated with BDC2.5 mimotope or insulin peptide insB 15–23, respectively. In other experiments, 2.5x10⁵ Th1 cells were cultured with 1x10⁴ APCs and BDC2.5 mimotope (0.5 μ g/mL) with or without increasing concentrations of IC as previously described for 72 hours. Cells cultured in the presence of the proliferative stimulus but without IC87114 were positive controls, whereas non-stimulated cells were negative controls.

Proliferation and cytokine analysis

In all experiments proliferation was assessed by CFSE staining (5 μ M). After gating on CD4⁺ and/or CD8⁺ T cells, the percentage of proliferating cells in each population was determined. For cytokine analysis, supernatants were taken at the end of the time cultures and IFN- γ production or IL-10 was assessed by specific ELISA (R&D Systems). Levels of other cytokines were detected using a cytometric bead array (eBioscience). For intracellular cytokine staining, cells were washed and stimulated with PMA (50 ng/mL) and ionomycin (2000 ng/mL) for 5 hours. BFA (5 μ g/mL) was added for the last 3 hours. Afterwards, the staining of cell surface markers was performed. Cells were washed, fixed, permeabilized (intracellular cytokine staining kit, eBioscience), and stained for detection of IFN- γ .

In vitro assessment of regulatory T cells

CD4⁺ CD25⁺ GFP⁺ cells from spleens and lymph nodes of 5-week-old Foxp3/GFP⁺ BDC2.5 TCR transgenic NOD mice were isolated by cell sorter (MoFlo, BD). Tregs were cultured with anti-CD3 (5 μ g/mL), anti-CD28 (20 μ g/mL) and IL-2 (1000 u/ml) with or without IC87114 (5 and 10 μ M) for 72 hours. Proliferation was assessed by dilution of CFSE staining (5 μ M, Invitrogen) after gating on CD4⁺CD25⁺ cells. Supernatants were assessed for IL-10 by specific ELISA (R&D). Positive control cells were stimulated with anti-CD3/28 antibodies and IL-2 whereas negative control cells were not stimulated at all.

Islet transplantation

Diabetes was induced in recipient mice (normal (WT), CD28^{-/-}, p110 δ ^{D910A/D910A} and CD28^{-/-}; p110 δ ^{D910A/D910A} double knockouts (DKOs)) by streptozotocin injection. All strains were backcrossed on the C57BL/6 background. Islets were prepared from MHC-mismatched donors (Cba1-C57BL/6 F1). Pancreatic islets were isolated through inflation of the pancreas via the bile duct [27], and islet transplantation was performed according to standard protocols [28]. Confirmed diabetic recipient mice received between 300 and 500 islets, giving approximately 15 islets per gram of body weight. We anaesthetised the recipient mice with isoflurane inhalation anaesthesia, and gave them sc temgesic for post-surgery analgesia. Islets were injected beneath the kidney capsule of female recipients that have been confirmed to be diabetic after the streptozotocin injection (Blood glucose level > 20 mM at two consecutive readings). Blood glucose was assessed three times per week for the period of graft survival, and daily at the onset of graft rejection—for a maximum of three days. Some mice which retained their grafts for longer than 100 days had the engrafted kidney removed at the end of the experiment to ascertain that the graft was responsible for the restoration of euglycemia.

Adoptive transfer models

CD4⁺ T cells were isolated by cell sorting from spleen and lymph nodes of BDC2.5 TCR transgenic mice. 1x10⁶ CD4⁺ T cells were transferred by the ip route to 5-week old NOD-*scid* mice. Mice received IC87114 treatment (30mg/Kg) by oral gavage twice per day and/or CTLA-4 Ig administered by the ip route at a concentration of 500 μ g/mL (the first time) and then 250 μ g/mL every other day from day 0 to 10. Positive control mice did not receive any treatment. After day 7, mice were monitored for the development of diabetes by measuring blood glucose or urine glucose levels with Diastix strips (Bayer) every day. In other experiments, 5x10⁵ Th1 cells (see above) were transferred by the ip route to 5-week old NOD mice. Mice received the same treatment as described above, but from day 0 to 5. After day 3, mice were monitored for the development of diabetes as previously described.

Statistical analysis

Differences between groups were tested using the student t-test, significant p-values are indicated with * ($p \leq 0.05$), ** ($p \leq 0.001$) or *** ($p \leq 0.0001$). Differences between animals regarding diabetes incidence or recurrence after islet transplant were tested using the Log rank survival test, with actual p-values displayed within the relevant figure or legend. All analysis was performed using GraphPad Prism software.

Results

IC87114 inhibition of the PI3K p110 δ subunit does not diminish T cell proliferation but has profound effects on cytokine production

We assessed the effects of increasing levels of IC87114 on T cells from diabetes-prone NOD mice in *in vitro* culture, looking at anti-CD3 and anti-CD28 induced proliferation of CD4⁺ T cells (Fig 1A, left hand panel) and CD8⁺ T cells (Fig 1A, right hand panel) and production of IFN- γ (Fig 1B). We found that the presence of IC87114 did not suppress proliferation of either CD4⁺ or CD8⁺ T cells (Fig 1A) but that the production of IFN- γ was severely impaired even at the lowest concentration of IC87114 tested (Fig 1A). Unaffected proliferation (as measured by % of cells divided ≥ 1 time) and suppressed IFN- γ production were seen both in islet specific TCR transgenic BDC2.5 CD4⁺ T cells [21] (Fig 1C and 1D), which recognise a posttranslationally modified peptide of chromogranin A [29], and in TCR transgenic G9C8 CD8⁺ T cells which recognise an insulin peptide [23, 24] (Fig 1E and 1F). We performed an extended analysis of the effects on cytokine production by IC87114, and found that increasing concentration of IC87114 suppressed all the pro-inflammatory cytokines assessed in supernatants from peptide-stimulated BDC2.5 CD4⁺ T cells, including IFN- γ , IL-2, GM-CSF, and IL-6, while there was a trend towards increased production of anti-inflammatory cytokine IL-10 and Th2 associated cytokines IL-4 and IL-5 (S1 Fig). Production of the cytokines that could be detected in supernatants from insulin peptide stimulated G9C8 CD8⁺ T cells was also decreased, with lower levels of IFN- γ , IL-2, TNF, IL-17, GM-CSF, IL-10 and IL-6 recorded even at the lowest concentrations of added IC87114 (S2 Fig). An interesting finding was that T cell proliferation was if anything increased in the presence of higher concentrations of IC87114, with more cells dividing several times (S3A Fig) and with BDC2.5 CD4⁺ T cells upregulating CD25 in response to increasing concentration of IC87114 (S3B Fig).

Administration of IC87114 via oral gavage does not delay onset of diabetes after BDC2.5 cell transfer into NOD-*scid* recipients

As production of proinflammatory cytokines is central to the process whereby islet-specific T cells cause type 1 diabetes through incapacitating and killing beta cells [30], we wanted to test whether oral administration of IC87114 could prevent or delay onset of diabetes in NOD-*scid* mice after transfer of potentially diabetogenic islet-specific BDC2.5 CD4⁺ T cells. We found that oral administration of 30mg/kg body weight twice per day (Fig 2A) had no effect on the development of diabetes after cell transfer, as all mice in both groups had developed diabetes by day 13 after transfer (Fig 2B).

IC87114 suppresses IFN- γ production from Th1-differentiated islet specific effector T cells but does not stop them from causing diabetes after transfer into wt NOD recipients

As IC87114 did not have an effect on development of diabetes after transfer of BDC2.5 cells we wished to elucidate whether it could decrease IFN- γ production from already Th1

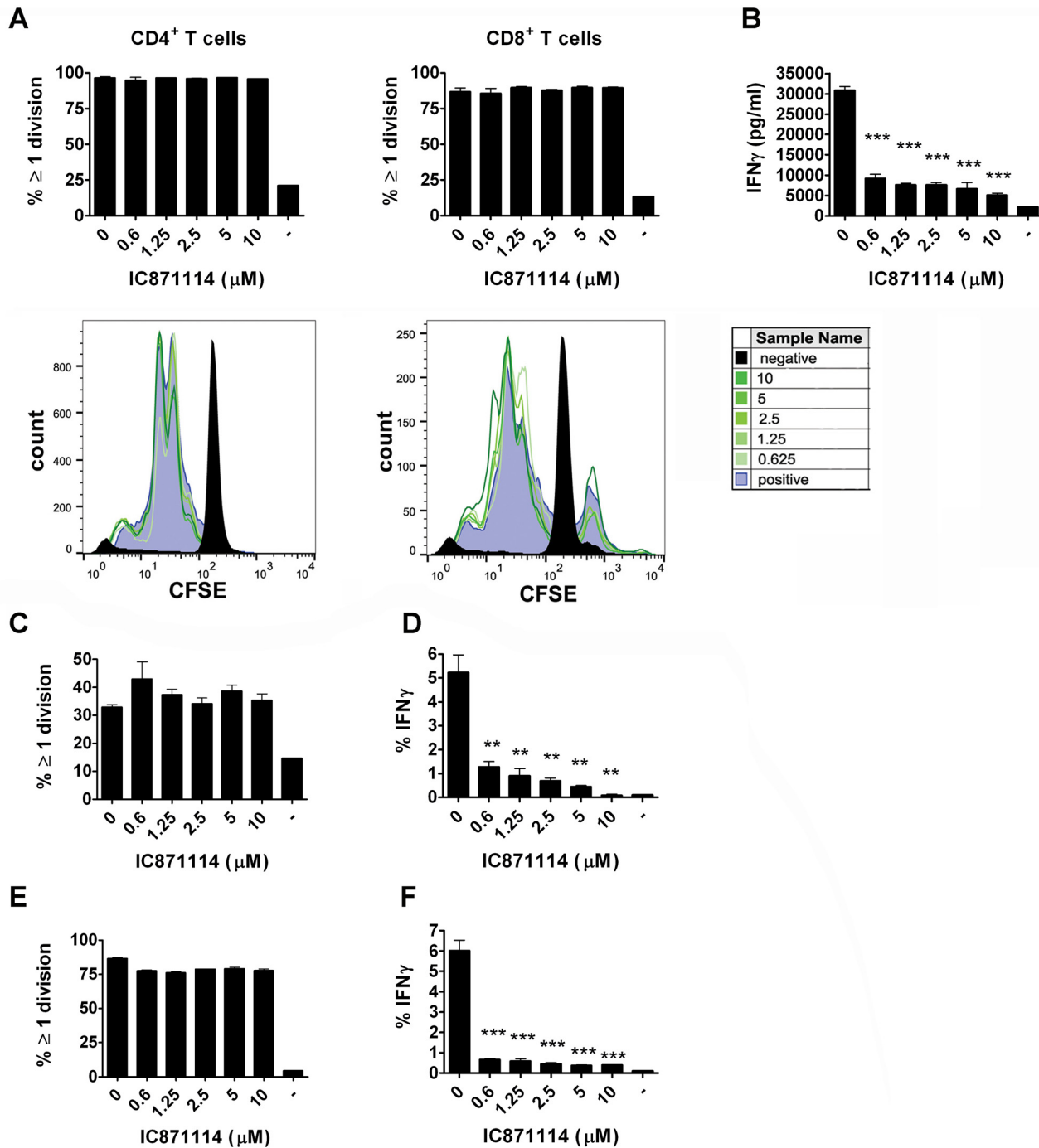


Fig 1. IC871114 blocks IFN- γ production by activated NOD cells. Cells isolated from the spleens and lymph nodes of regular NOD mice (A,B); BDC2.5 TCR transgenic NOD mice (C,D) or G9C8 TCR transgenic NOD mice (E,F) were stimulated with anti-CD3/28 antibodies (A, B), BDC2.5 mimotope (C, D) or insulin peptide (E, F), respectively, with or without increasing concentrations of IC871114 (0.6–10 μ M) for 72 hours. (A,C,E) Cells were stained with CFSE and after gating on CD4⁺ and/or CD8⁺ T cells, the percent of proliferating cells in each population was determined. (B) Supernatants were collected and IFN- γ production was assessed by specific ELISA (B) or intracellular staining (D, F). All data were expressed as the mean \pm SD for triplicate samples, differences between groups were tested using the student t-test. The data is representative of at least three independent experiments.

doi:10.1371/journal.pone.0146516.g001

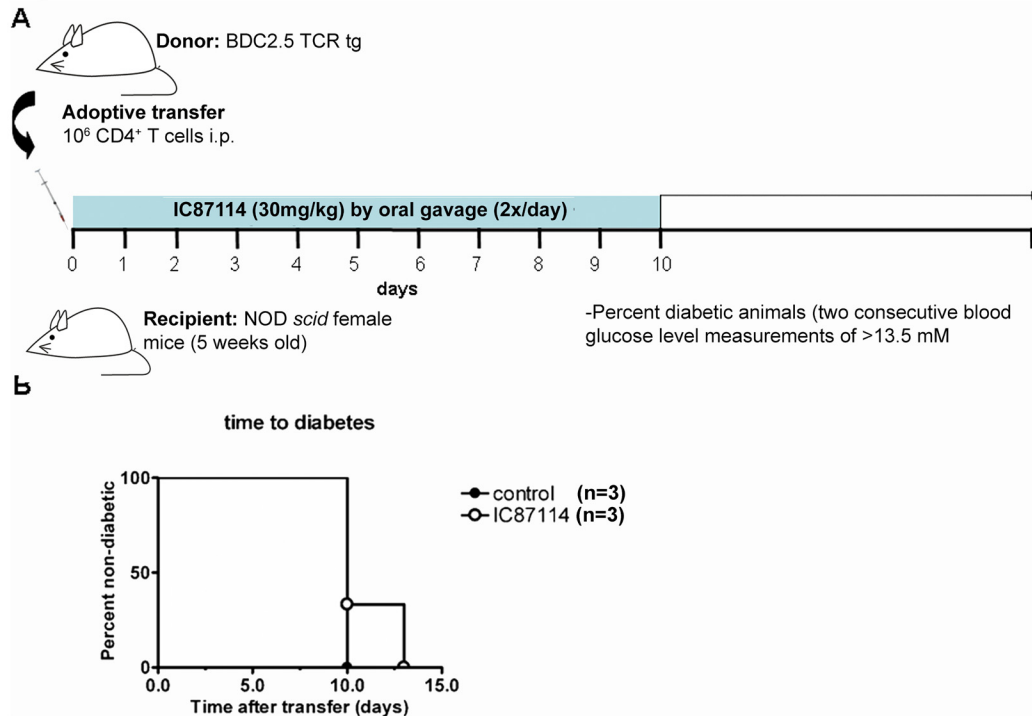


Fig 2. Administration of IC87114 does not prevent diabetes after adoptive transfer of naïve diabetogenic cells. CD4⁺ T cells were isolated by cell sorting from spleen and lymph nodes of BDC2.5 TCR transgenic mice. 1x10⁶ CD4⁺ T cells were transferred i.p. route to NOD-*scid* mice. Mice received IC87114 treatment (30mg/Kg) by oral gavage (2x/day) from day 0 to 10 (A). Positive controls are mice that received CD4⁺ T cells from BDC2.5 mice without any other treatment. After this time, blood glucose levels were checked every day, and mice were considered irrevocably diabetic and sacrificed when they reached two consecutive blood glucose levels >13mM (B).

doi:10.1371/journal.pone.0146516.g002

differentiated effector CD4⁺ T cells. We found that, just as seen in *in vitro* cultures with naïve cells, the Th1 differentiated effector cells proliferated equally well in the presence of IC87114 (Fig 3A), but that their IFN- γ production, although more robust than from naïve T cells, was decreased by IC87114 in a dose-dependent way (Fig 3B). Th1 differentiated effector BDC2.5 CD4⁺ T cells cause diabetes after transfer into wildtype NOD mice in a process heavily dependent on production of IFN- γ (Fig 4A) [30], and we hypothesised that administration of IC87114 via oral gavage which can stop IFN- γ production from these cells *in vitro*, would inhibit or delay onset of diabetes in the IFN- γ dependent model of disease. However, oral administration of 30mg/kg body weight twice per day from the day of cell transfer did not affect disease development, and both the IC87114-treated and the vehicle treated group developed diabetes on day 5 after adoptive transfer (Fig 4B). Oral administration did not appear to affect IFN- γ production *in vivo* as cells isolated from pancreas (Fig 4C) or pancreatic lymph nodes (Fig 4D) of IC87114 or vehicle-treated mice produced equal amounts of IFN- γ .

Combination with CTLA4-Ig does not alter or potentiate the *in vitro* effects of IC87114

As IC87114 showed no effect on *in vivo* diabetes development despite its dramatic effects on cytokine production *in vitro*, we hypothesised that a combination with another agent that targets a different pathway leading to the activation of T cells might synergise with the IC87114. Indeed, in work leading up to the current study, we found that p110 δ -CD28 double knockout mice are immune suppressed and failed to reject allogeneic islet grafts ([20] and S4 Fig). Since

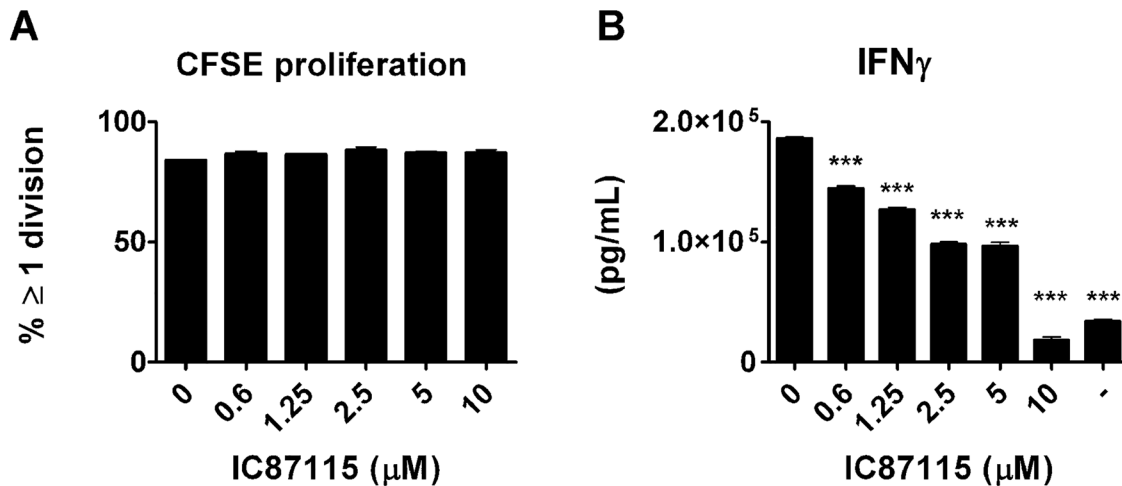


Fig 3. IC87114 reduces IFN- γ production by differentiated Th1 cells. Cells isolated from the spleens and lymph nodes of BDC2.5 TCR transgenic NOD mice were differentiated into Th1 cells by culturing them in the presence of anti-CD3/CD28 antibodies, IL-2, IL-12 and IFN- γ for 4 days. 2.5×10^5 Th1 cells were then cultured with 1×10^4 APCs and BDC2.5 mimotope (0.5 $\mu\text{g}/\text{mL}$) with or without increasing concentrations of IC87114 (0.6–10 μM) for 72 hours. Cells were stained with CFSE and after gating on CD4⁺ T cells, the percent of proliferating cells was determined (A). Supernatants were collected and IFN- γ production was assessed by specific ELISA (B). All data were expressed as the mean \pm SD for triplicate samples. The results are representative of at least three independent experiments, differences between groups were tested using the student t-test.

doi:10.1371/journal.pone.0146516.g003

CTLA-4 Ig inhibits the ability of T cells to receive signals via CD28, we assessed the effects of combining increasing levels of IC87114 with CTLA4-Ig on T cells from NOD mice in *in vitro* culture, looking at anti-CD3 induced proliferation of CD4⁺ T cells (Fig 1A, left hand panel) and CD8⁺ T cells (Fig 1A, right hand panel) and production of IFN- γ (Fig 1B). We found that the combination of CTLA4-Ig with IC87114 did not suppress proliferation of either CD4⁺ or CD8⁺ T cells any differently than IC87114 on its own (Fig 5A) and that the production of IFN- γ was similarly impaired in the presence of CTLA4-Ig (Fig 5B) compared with IC87114 alone (Fig 1B). The presence of CTLA4-Ig modestly impaired proliferation of BDC2.5 CD4⁺ T cells, but this was not potentiated by increasing concentrations of IC87114 (Fig 5C). However, CTLA4-Ig did have a significant effect on the production of IFN- γ from the BDC2.5 CD4⁺ T cells, and combination with IC87114 decreased IFN- γ production even further (Fig 5D). The proliferation and IFN- γ production of G9C8 CD8⁺ T cells was not affected by the addition of CTLA4-Ig (Fig 5D and 5E), demonstrating the same reduction of IFN- γ production in response to IC87114 as when no CTLA4-Ig was present (Fig 1F). We performed an extended analysis of the effects on cytokine production by combining CTLA4-Ig and IC87114, and found that addition of CTLA4-Ig on its own decreased the production of IL-2 in supernatants from peptide-stimulated BDC2.5 CD4⁺ T cells, but that other cytokines assessed demonstrated no additional downregulation in response to the additional presence of CLTA4-Ig to the increasing concentrations of IC87114 in the culture (S5 Fig). We did not detect an effect of CTLA-4 Ig with the bead array, which could be due to the fact that the high concentration of IFN- γ in the samples exceeding the top limit of the assay (S5 Fig).

Combination treatment with CTLA4-Ig does not increase protection from diabetes by IC87114

As production of IFN- γ was further decreased in BDC2.5 CD4⁺ T cells *in vitro* when CTLA4-Ig was added to IC87114 in culture, we tested whether oral administration of IC87114 in combination with CTLA4-Ig could prevent or delay onset of diabetes in NOD *scid* mice after transfer

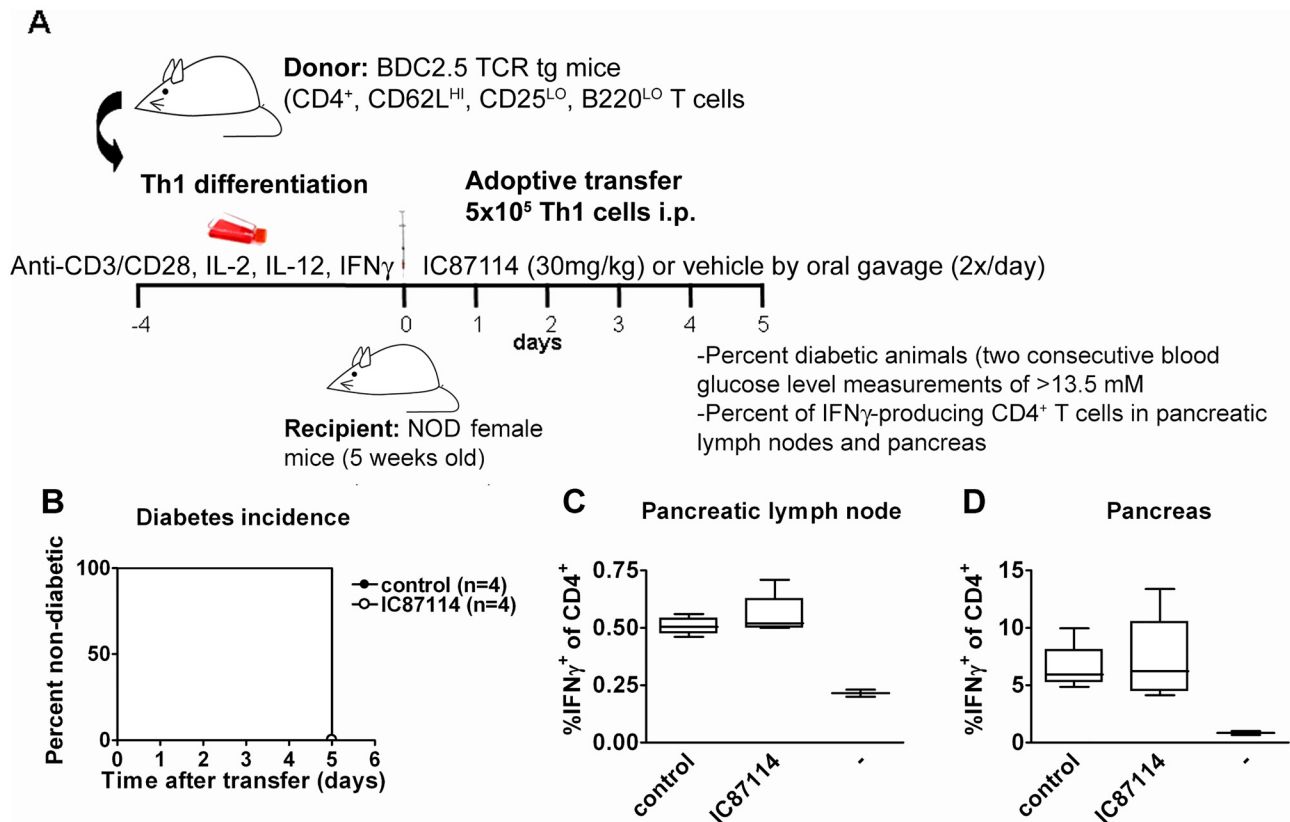


Fig 4. IC87114 treatment does not reduce the incidence of diabetes after adoptive transfer of Th1 cells to wt NOD mice. CD4⁺, CD62L^{hi}, CD25^{lo} B220^{lo} T cells were isolated by cell sorting from spleen and lymph nodes of BDC2.5 TCR transgenic mice and differentiated into Th1 cells by culturing them in the presence of anti-CD3/CD28 antibodies, IL-2, IL-12 and IFN- γ for 4 days. 5x10⁵ Th1 cells were transferred i.p. to wt NOD mice and IC87114 (30mg/Kg) or vehicle were administered by oral gavage twice daily from day 0 to 5 (A). Mice were considered diabetic after 2 consecutive blood glucose readings > 13mM (B). We assessed the percentages of IFN- γ -producing CD4⁺T cells from pancreatic lymph nodes (C) and pancreas (D) in vehicle treated mice, IC87114 treated mice and negative control mice (4 mice per group). Negative control NOD mice did not receive Th1 cells nor any treatment.

doi:10.1371/journal.pone.0146516.g004

of BDC2.5 CD4⁺ T cells (Fig 6A). We found that intraperitoneal injections of CTLA4-Ig every other day for the first 10 days after transfer significantly delayed onset of diabetes in recipient mice (Fig 6B), but that combination with twice daily administration of IC87114 did not enhance the protective effect, but rather appeared to attenuate it (Fig 6B).

Combination with CTLA4-Ig does not alter or potentiate the *in vitro* or *in vivo* effects of IC87114 on Th1 differentiated effector T cells

We also assessed whether addition of CTLA4-Ig had an additional inhibitory affect on IFN- γ production from already Th1 differentiated effector CD4⁺ T cells. We found that, just as seen in Fig 3, that the Th1 differentiated effector cells proliferated equally well in the presence of IC87114 (Fig 3A), but that their IFN- γ production, although more robust than from naïve T cells, was decreased by IC87114 in a dose dependent way. Addition of CTLA4-Ig to the culture had no effect on proliferation (Fig 7A), and had some effect on IFN- γ production, not appearing to result in synergistic suppression (Fig 7B). Neither treatment with CTLA4-Ig on its own nor in combination with IC87114 affected development of diabetes after transfer of Th1 differentiated effector BDC2.5 CD4⁺ T cells into wt NOD mice (Fig 7C), as all mice regardless of treatment developed diabetes 5 to 6 days after injection of the diabetogenic cells.

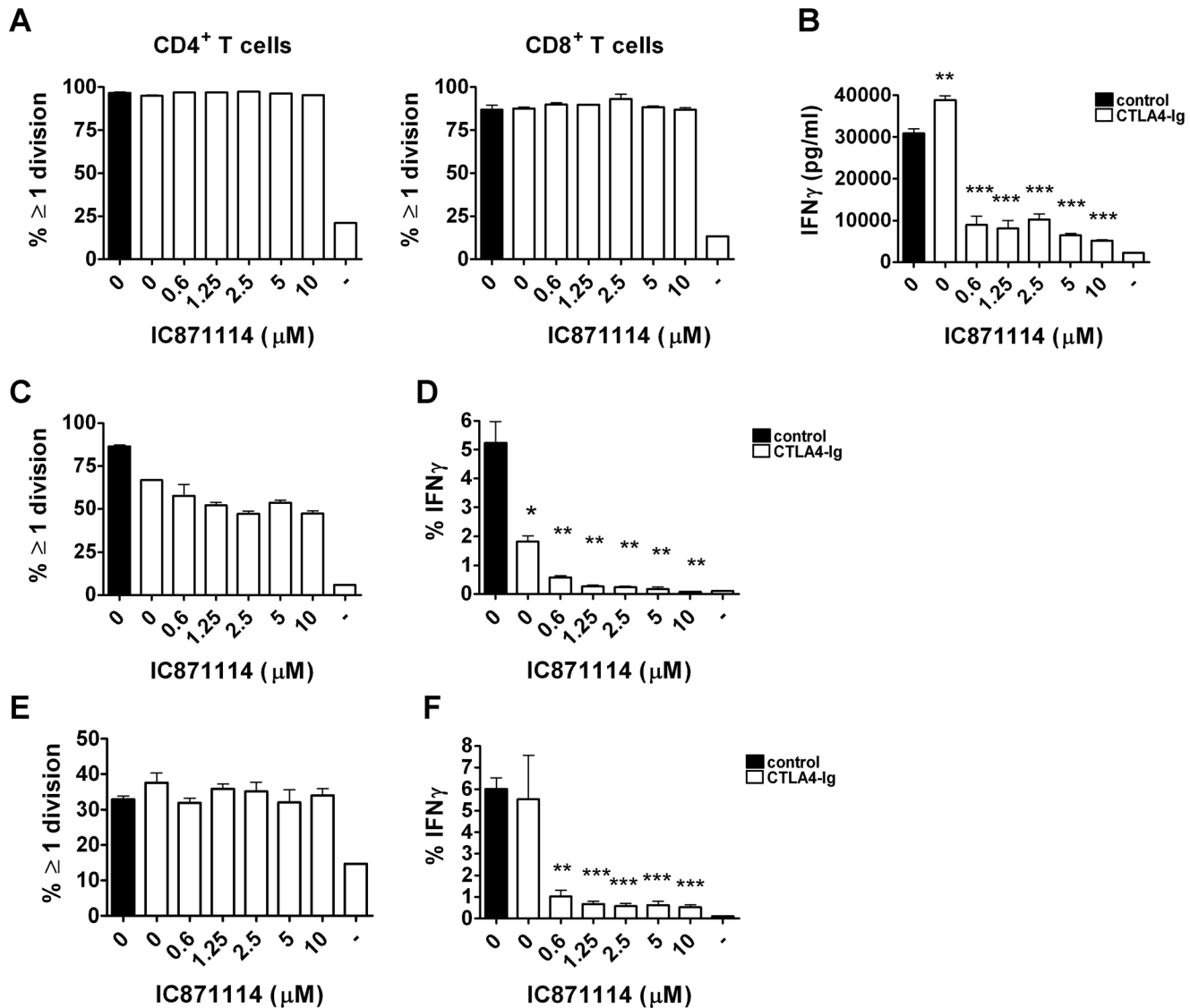


Fig 5. Combination of CTLA4-Ig with IC87114 has no major effect on diabetogenic cells. Cells isolated from the spleens and lymph nodes of wild-type NOD mice (A,B); BDC2.5 TCR transgenic NOD mice (C,D) or G9C8 TCR transgenic NOD mice (E,F) were stimulated with anti-CD3/28 antibodies (A, B), BDC2.5 mimotope (C, D) or insulin peptide (E, F), respectively, with or without increasing concentrations of IC87114 (0.6–10μM), with or without CTLA4-Ig (100 ng/ml) as indicated, for 72 hours. Cells were stained with CFSE and after gating on CD4⁺ and/or CD8⁺ T cells, the percent of proliferating cells in each population was determined (A,C,E). IFN-γ production was assessed either through ELISA (B) or intracellular staining (D, F). All data were expressed as the mean ± SD for triplicate samples, differences between groups were tested using the student t-test. The data is representative of at least three independent experiments.

doi:10.1371/journal.pone.0146516.g005

IC87114 does not alter the percentage of regulatory T cells, but reduces their production of IL-10

Treatment with IC87114 appeared to break the temporary tolerance afforded by CTLA4-Ig treatment after islet specific T cell transfer (Fig 6B). As the effects of IC87114 on the cytokine production of these cells were so striking, we wondered if effects of IC87114 on the regulatory T cell pool could be even more important, and thus cancel out and even override any anti-inflammatory effects. We found that oral administration of IC87114 had no effect on the percentages of Treg present in the spleen, inguinal or pancreatic lymph nodes of treated mice (Fig

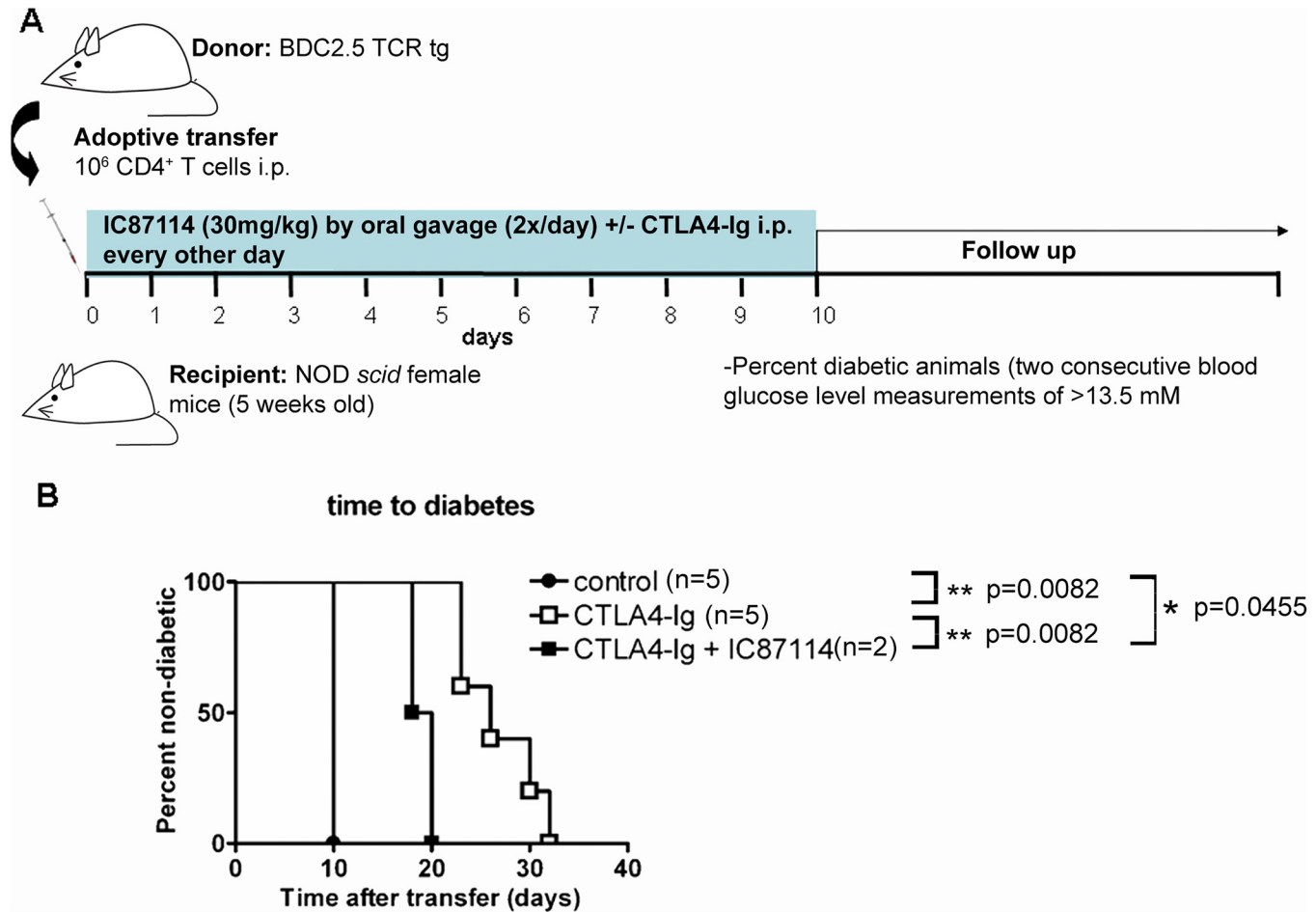


Fig 6. Combination treatment with CTLA4-Ig and IC87114 does not prevent diabetes after adoptive transfer of naïve diabetogenic cells. CD4⁺ T cells were isolated by cell sorting from spleen and lymph nodes of BDC2.5 TCR transgenic mice. 1x10⁶ CD4⁺ T cells were transferred i.p. route to NOD-*scid* mice. Mice received IC87114 treatment (30mg/Kg) or vehicle by oral gavage (2x/day) from day 0 to 10, and/or ip injections of CTLA4-Ig on day 0, 3 and 5 (A). Positive control mice received CD4⁺ T cells from BDC2.5 mice without any other treatment. Blood glucose levels were checked every day from day 7, and mice were considered irrevocably diabetic and sacrificed when they reached two consecutive blood glucose levels >13mM (B). Differences between treatment groups were determined using the Log Rank survival test.

doi:10.1371/journal.pone.0146516.g006

8A) or on their absolute numbers (data not shown), and that presence of IC87114 in culture medium did not affect proliferation of sorted Treg from Foxp3-GFP reporter NOD mice *in vitro* (Fig 8B). However, we found that cytokine production was strongly affected by IC87114, demonstrating a ~65% drop in IL-10 production in the presence of 5mM IC87114 (Fig 8C).

Discussion

IC87114 has a profound suppressive effect on the production of proinflammatory cytokines such as IFN-γ and IL-17 as well as IL-2 from islet-specific cells activated *in vitro*, while increasing the production of anti-inflammatory cytokines such as IL-4 and IL-10. As the development of diabetes in NOD mice depends on the production of IFN-γ from CD4⁺ T cells [30], and IFN-γ production and IL-17 production is elevated in islet-reactive CD4⁺ T cells in recent onset type 1 diabetes patients [5, 31], we hypothesised that administration of IC87114 could have beneficial effects on anti-islet inflammation and be a feasible option of type 1 diabetes therapy. A previous study has demonstrated modest positive effects of IC87114 on the

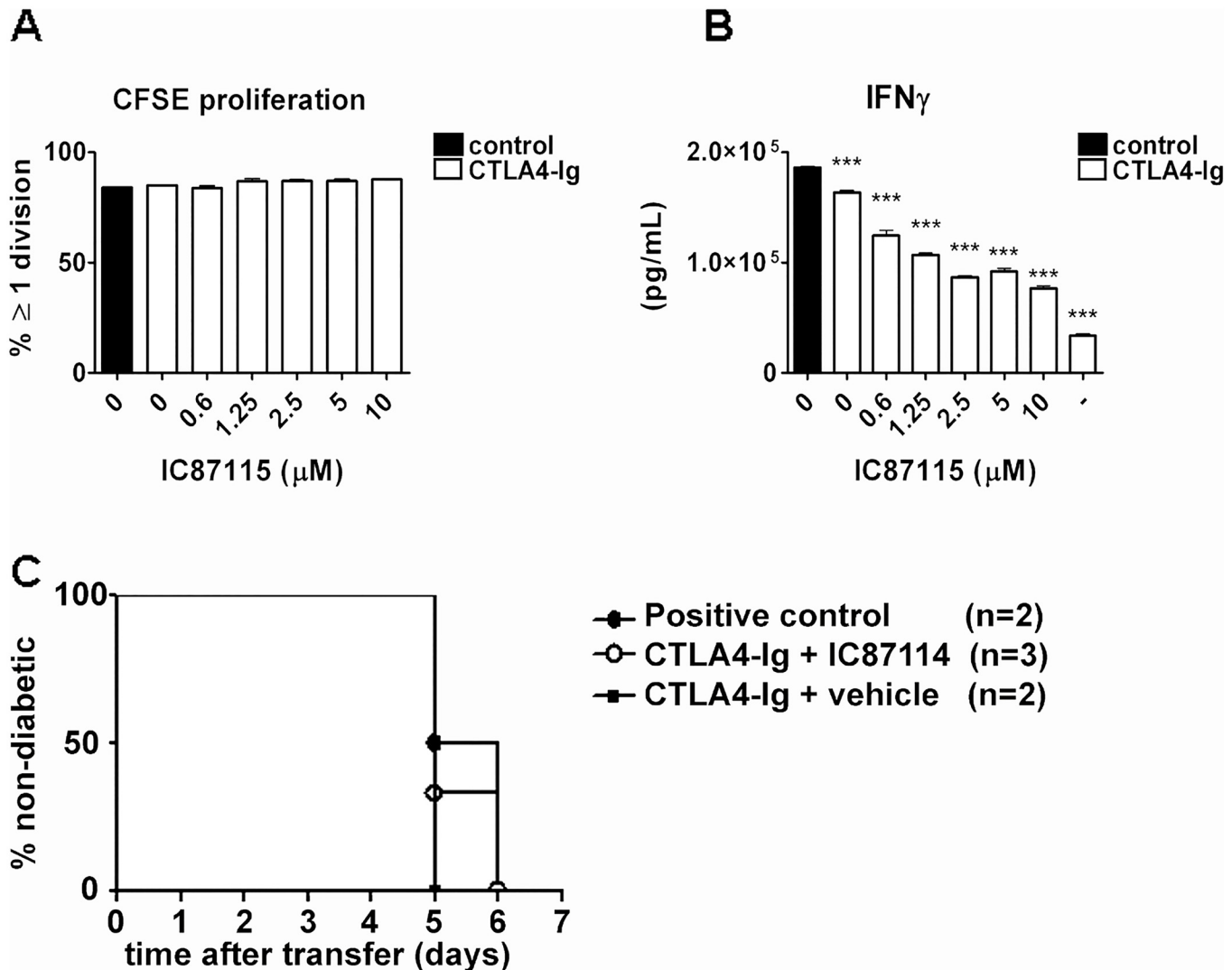


Fig 7. Combined treatment of IC87114 and CTLA-4 Ig does not reduce the incidence of diabetes in the adoptive transfer model of *Th1* cells from BDC2.5 transgenic mice to NOD mice. CD4⁺, CD62L^{hi}, CD25^{lo} B220^{lo} T cells were isolated by cell sorting from spleen and lymph nodes of BDC2.5 TCR transgenic mice and differentiated into Th1 cells by culturing them in the presence of anti-CD3/CD28 antibodies, IL-2, IL-12 and IFN- γ for 4 days. Th1 cells were then cultured with 1×10^4 APCs and BDC2.5 mimotope (0.5 μ g/mL) with or without increasing concentrations of IC87114 (0.6–10 μ M)), with or without CTLA4-Ig (100 ng/ml) as indicated, for 72 hours. Cells were stained with CFSE and after gating on CD4⁺ T cells, the percentage of proliferating cells was determined (A). Supernatant IFN- γ assessed by specific ELISA (B). All data expressed as the mean \pm SD for triplicate samples, differences between groups were tested using the student t-test. The results are representative of at least three independent experiments. 5×10^5 Th1 cells were transferred i.p. to wt NOD mice. IC87114 (30mg/Kg) or vehicle were administrated by oral gavage twice daily from day 0 to 5, and/ CTLA4-Ig injected ip on day 0, 3 and 5. The number of mice per group is indicated within the Fig Mice were considered diabetic after 2 consecutive blood glucose readings > 13mM (C).

doi:10.1371/journal.pone.0146516.g007

development of diabetes in NOD mice [32], and we hypothesised that combination of this agent which has such dramatic effects on production of pro-inflammatory cytokines even from already activated cells [15] with another well tolerated treatment targeting the activation of T cells [11, 26] would have a chance of preventing disease completely. Indeed, preliminary studies showed that genetic inactivation of PI3K δ and CD28 resulted in indefinite acceptance of islet allografts.

However, we found that, despite seeing considerable inhibition of pro-inflammatory cytokine production even at the lowest concentrations of IC87114 tested, treatment with IC87114

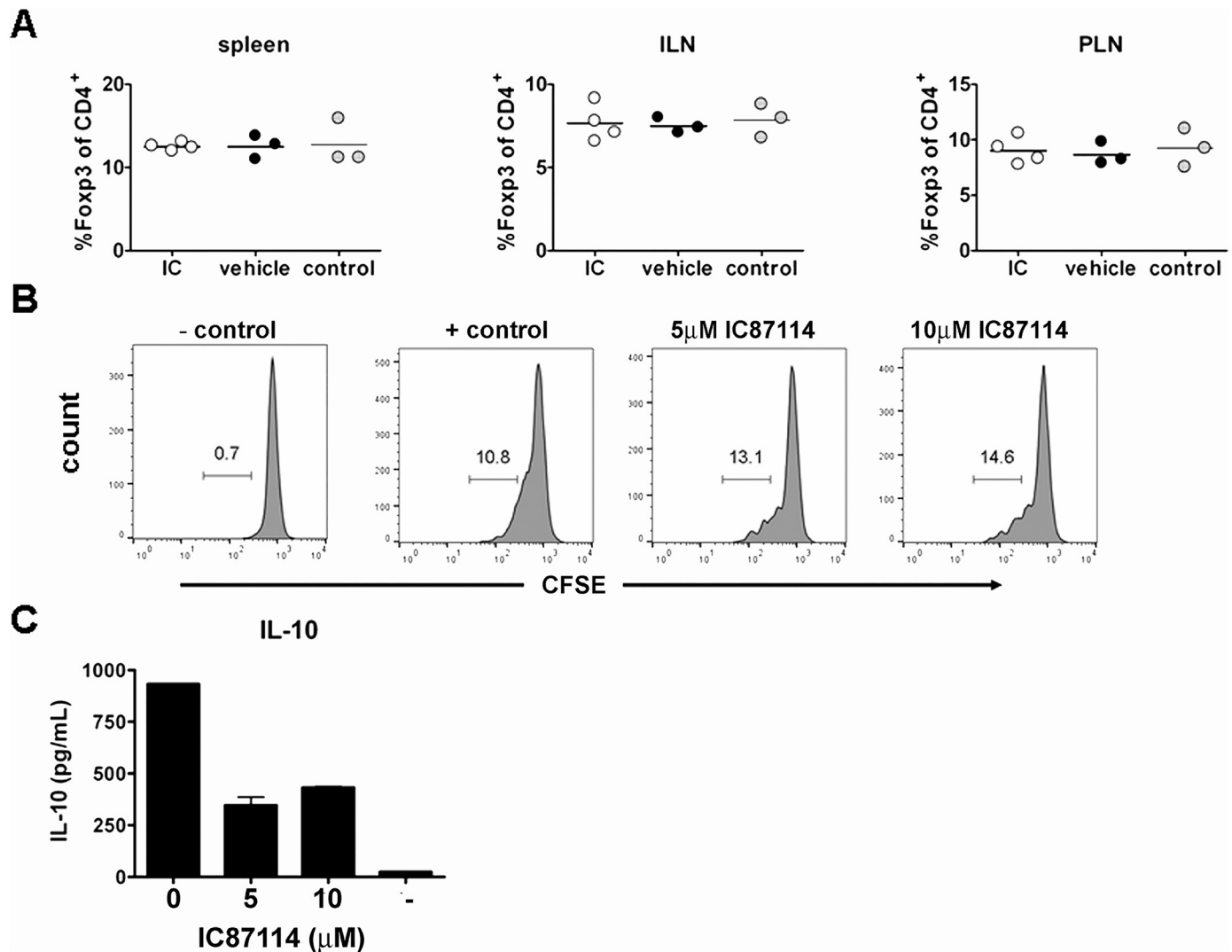


Fig 8. IC87114 does not affect percentages of Foxp3⁺ Treg in treated mice, but reduces IL-10 production from Treg. Percentages of Foxp3⁺ cells were assessed in spleen, inguinal lymph nodes and pancreatic lymph nodes of mice treated with IC87114 twice daily for 10 days. Each data point represents one mouse (A). Proliferation of sorted CFSE labeled BDC2.5 Foxp3-GFP cells in response to anti-CD3/CD28 antibodies in the presence of IC87114 (B). Production of IL-10 from stimulated, sorted, BDC2.5 Foxp3-GFP in presence of IC87114 measured by supernatant ELISA (C). Results are representative of at least two experiments.

doi:10.1371/journal.pone.0146516.g008

had no effect on the development of diabetes either after adoptive transfer of naïve BDC2.5 CD4⁺ T cells into NOD *scid* mice or after transfer of Th1 differentiated effector BDC2.5 CD4⁺ T cells into wild type NOD mice. The group sizes we used for *in vivo* experiments were small, and it cannot be excluded that differences between groups could have been detected had the sample size been larger. However, as no positive effects of IC87114 were detected in the preliminary *in vivo* experiments, we decided against using larger numbers of mice which might have allowed us to detect subtle differences between groups.

Adding CTLA4-Ig to cultures had specific effects, targeting CD4⁺ T cell proliferation and production of IL-2 from these cells, and combination of CTLA4-Ig with IC87114 had a major effect on IFN-γ production from CD4⁺ T cells. This was considered promising as IFN-γ production from CD4⁺ T cells that has been shown to be particularly important in the pathology

of diabetes in mice as well as men [5, 30]. However, combination treatment with twice daily oral administration of IC87114 and intraperitoneal injections with CTLA4-Ig every other day for 10 days did not lead to protection from diabetes either after transfer of naïve cells or pre-activated Th1 cells. In fact, treatment with IC87114 appeared to attenuate the protective effect afforded by CTLA4-Ig injection, raising the question of whether effects on the regulatory T cell subset may be greater than any effects on anti-inflammatory cytokine production. We found that although IC87114 treatment did not change the percentage of Foxp3⁺ regulatory CD4⁺ T cells present in secondary lymphoid organs, these cells produced less anti-inflammatory IL-10 when activated in the presence of IC87114, and are thus also a feasible targets of IC87114 treatment. Production of anti-inflammatory cytokines such as IL-10 [33] and TGF- β is known to suppress development of diabetes in NOD mice [34, 35], and it is not surprising that a reduction in the production of anti-inflammatory cytokines could precipitate disease.

Regulatory T cells have been identified as a target of PI3K p110 δ inhibitors, and targeting p110 δ signalling in these cells increases anti-tumour immune responses by releasing cytotoxic CD8⁺ T cells from Treg control [36, 37] even though lack of p110 δ signalling also resulted in decreased CD8⁺ T cell expression of activation marker CD44 as well as granzyme B and perforin. Our findings demonstrate major effects of IC87114 on the production of pro-inflammatory cytokines from diabetogenic T cell clones *in vitro*, but no decrease in their diabetogenicity *in vivo*. In fact, we see that treatment with IC87114 abrogates the protective effects of CTLA4-Ig treatment and causes a quicker progression to diabetes after cell transfer. Our results indicate that IC87114 is not a promising candidate for treatment of type 1 diabetes, even in combination with other drugs such as CTLA4-Ig, but that it leaves the *in vivo* inflammatory response unchanged. Instead, our data adds weight to the proposal that PI3K p110 δ inhibition, as a means to reduce the effect of Treg, is a feasible strategy for breaking tumour-specific immune tolerance to achieve improved cancer immunotherapy.

Supporting Information

S1 Fig. Effects of IC87114 on cytokine production in BDC2.5 CD4⁺ T cells. Cells isolated from the spleens and lymph nodes of BDC2.5 TCR transgenic NOD mice were stimulated with the BDC2.5 mimotope (0.5 μ g/mL) with or without increasing concentrations of IC87114 (0.6–10 μ M) for 48 hours. Cytokines from supernatants were assessed in duplicate using a bead cytokine array, differences between groups were tested using the student t-test. (TIF)

S2 Fig. Effects of IC87114 on cytokine production in G9C8 CD8⁺ T cells. Cells isolated from the spleens and lymph nodes of G9C8 TCR transgenic NOD mice were stimulated with the insulinB 15–23 peptide (0.5 μ g/mL) with or without increasing concentrations of IC87114 (0.6–10 μ M) for 48 hours. Cytokines from supernatants were assessed in duplicate using a bead cytokine array, differences between groups were tested using the student t-test. (TIF)

S3 Fig. Effects of IC87114 on the distribution of divisions and activation status. Cells isolated from the spleens and lymph nodes of BDC2.5 TCR transgenic NOD mice were stimulated with the BDC2.5 mimotope (0.5 μ g/mL) with or without increasing concentrations of IC87114 (0.6–10 μ M) for 72 hours, and then stained for CD25 A histogram overlay of representative cultures gated on CD4⁺ cells (A, left) and a graph showing all data (A, right). Cells isolated from the spleens and lymph nodes of G9C8 TCR transgenic NOD mice were stained with CFSE and stimulated with the insulinB 15–23 peptide (0.5 μ g/mL) with or without increasing concentrations of IC87114 (0.6–10 μ M) for 72 hours. A histogram overlay of representative cultures

gated on CD8⁺ cells (B, left), and a graph showing all data (B, right). Differences between groups were tested using the student t-test.

(TIF)

S4 Fig. Survival of MHC mis-matched islets in streptozotocin induced diabetic recipients.

Wt C57BL/6 mice, CD28 KO, PI3K p110D910A (D910A) and CD28-D910A double deficient mice (DKO) were rendered diabetic through injection of streptozotocin. Diabetic mice received a MHC mis-matched (Cba1-C57BL/6 F1 donor) islet graft under the kidney capsule. Blood glucose was monitored in the recipient mice for up to 215 days. Some DKO mice that remained euglycemic for a long time underwent nephrectomy at the end of the experiment to ascertain that the graft was the cause of the restored euglycemia. The difference in euglycemic survival between wt recipient mice and DKO recipient mice was assessed using the Log Rank survival test, resulting in a p-value of 0.0027 (**).

(TIF)

S5 Fig. Effects of combination of CTLA4-Ig and IC87114 on cytokine production in BDC2.5 CD4⁺ T cells.

Cells isolated from the spleens and lymph nodes of BDC2.5 TCR transgenic NOD mice were stimulated with the BDC2.5 mimotope (0.5 μ g/mL) in the presence of CTLA4-Ig (100 ng/mL) with or without increasing concentrations of IC87114 (0.6–10 μ M) for 48 hours. Cytokines from supernatants were assessed in duplicate using a bead cytokine array, differences between groups were tested using the student t-test.

(TIF)

Acknowledgments

The authors wish to thank Yvonne Sawyer, Nigel Miller, Lynn Broom and Jonathan Clark for excellent technical assistance, and Nick Holmes for helpful discussion of the data.

Author Contributions

Conceived and designed the experiments: ABB KO MW. Performed the experiments: ABB JE AR MW. Analyzed the data: ABB AR AC MW. Contributed reagents/materials/analysis tools: FSW AC. Wrote the paper: ABB FSW AC KO MW.

References

1. Barrett JC, Clayton DG, Concannon P, Akolkar B, Cooper JD, Erlich HA, et al. Genome-wide association study and meta-analysis find that over 40 loci affect risk of type 1 diabetes. *Nat Genet.* 2009; 41(6):703–7. Epub 2009/05/12. ng.381 [pii] doi: [10.1038/ng.381](https://doi.org/10.1038/ng.381) PMID: [19430480](https://pubmed.ncbi.nlm.nih.gov/19430480/).
2. Steck AK, Rewers MJ. Genetics of type 1 diabetes. *Clin Chem.* 2011; 57(2):176–85. Epub 2011/01/06. clinchem.2010.148221 [pii] doi: [10.1373/clinchem.2010.148221](https://doi.org/10.1373/clinchem.2010.148221) PMID: [21205883](https://pubmed.ncbi.nlm.nih.gov/21205883/).
3. Phillips JM, Parish NM, Bland C, Sawyer Y, De La Pena H, Cooke A. Type 1 Diabetes Development Requires Both CD4⁺ and CD8⁺ T cells and Can Be Reversed by Non-Depleting Antibodies Targeting Both T Cell Populations. *Rev Diabet Stud.* 2009; 6(2):97–103. Epub 2009/10/07. doi: [10.1900/RDS.2009.6.97](https://doi.org/10.1900/RDS.2009.6.97) PMID: [19806239](https://pubmed.ncbi.nlm.nih.gov/19806239/).
4. Willcox A, Richardson SJ, Bone AJ, Foulis AK, Morgan NG. Analysis of islet inflammation in human type 1 diabetes. *Clin Exp Immunol.* 2009; 155(2):173–81. Epub 2009/01/09. CEI3860 [pii] doi: [10.1111/j.1365-2249.2008.03860.x](https://doi.org/10.1111/j.1365-2249.2008.03860.x) PMID: [19128359](https://pubmed.ncbi.nlm.nih.gov/19128359/); PubMed Central PMCID: [PMC2675247](https://pubmed.ncbi.nlm.nih.gov/PMC2675247/).
5. Arif S, Tree TI, Astill TP, Tremble JM, Bishop AJ, Dayan CM, et al. Autoreactive T cell responses show proinflammatory polarization in diabetes but a regulatory phenotype in health. *J Clin Invest.* 2004; 113(3):451–63. Epub 2004/02/03. doi: [10.1172/JCI19585](https://doi.org/10.1172/JCI19585) PMID: [14755342](https://pubmed.ncbi.nlm.nih.gov/14755342/).
6. Skowera A, Ellis RJ, Varela-Calvino R, Arif S, Huang GC, Van-Krinks C, et al. CTLs are targeted to kill beta cells in patients with type 1 diabetes through recognition of a glucose-regulated preproinsulin epitope. *J Clin Invest.* 2008; 118(10):3390–402. Epub 2008/09/20. doi: [10.1172/JCI35449](https://doi.org/10.1172/JCI35449) PMID: [18802479](https://pubmed.ncbi.nlm.nih.gov/18802479/); PubMed Central PMCID: [PMC2542849](https://pubmed.ncbi.nlm.nih.gov/PMC2542849/).

7. Wallberg M, Cooke A. Immune mechanisms in type 1 diabetes. *Trends Immunol.* 2013; 34(12):583–91. Epub 2013/09/24. S1471-4906(13)00131-2 [pii] doi: [10.1016/j.it.2013.08.005](https://doi.org/10.1016/j.it.2013.08.005) PMID: [24054837](https://pubmed.ncbi.nlm.nih.gov/24054837/).
8. Banting FG, Best CH, Collip JB, Campbell WR, Fletcher AA. Pancreatic Extracts in the Treatment of Diabetes Mellitus. *Can Med Assoc J.* 1922; 12(3):141–6. Epub 1922/03/01. PMID: [20314060](https://pubmed.ncbi.nlm.nih.gov/20314060/).
9. Sherry N, Hagopian W, Ludvigsson J, Jain SM, Wahlen J, Ferry RJ Jr, et al. Teplizumab for treatment of type 1 diabetes (Protege study): 1-year results from a randomised, placebo-controlled trial. *Lancet.* 2011; 378(9790):487–97. Epub 2011/07/02. S0140-6736(11)60931-8 [pii] doi: [10.1016/S0140-6736\(11\)60931-8](https://doi.org/10.1016/S0140-6736(11)60931-8) PMID: [21719095](https://pubmed.ncbi.nlm.nih.gov/21719095/).
10. Ambery P, Donner TW, Biswas N, Donaldson J, Parkin J, Dayan CM. Efficacy and safety of low-dose otelexizumab anti-CD3 monoclonal antibody in preserving C-peptide secretion in adolescent type 1 diabetes: DEFEND-2, a randomized, placebo-controlled, double-blind, multi-centre study. *Diabet Med.* 2014. Epub 2013/11/19. doi: [10.1111/dme.12361](https://doi.org/10.1111/dme.12361) PMID: [24236828](https://pubmed.ncbi.nlm.nih.gov/24236828/).
11. Orban T, Bundy B, Becker DJ, DiMeglio LA, Gitelman SE, Goland R, et al. Co-stimulation modulation with abatacept in patients with recent-onset type 1 diabetes: a randomised, double-blind, placebo-controlled trial. *Lancet.* 2011; 378(9789):412–9. Epub 2011/07/02. S0140-6736(11)60886-6 [pii] doi: [10.1016/S0140-6736\(11\)60886-6](https://doi.org/10.1016/S0140-6736(11)60886-6) PMID: [21719096](https://pubmed.ncbi.nlm.nih.gov/21719096/).
12. von Herrath MG, Korsgren O, Atkinson MA. Factors Impeding the Discovery of an Intervention-based Treatment for Type 1 Diabetes. *Clin Exp Immunol.* 2015. Epub 2015/05/20. doi: [10.1111/cei.12656](https://doi.org/10.1111/cei.12656) PMID: [25989477](https://pubmed.ncbi.nlm.nih.gov/25989477/).
13. Okkenhaug K, Vanhaesebroeck B. PI3K in lymphocyte development, differentiation and activation. *Nat Rev Immunol.* 2003; 3(4):317–30. Epub 2003/04/02. doi: [10.1038/nri1056](https://doi.org/10.1038/nri1056) nri1056 [pii]. PMID: [12669022](https://pubmed.ncbi.nlm.nih.gov/12669022/).
14. Okkenhaug K. Signaling by the phosphoinositide 3-kinase family in immune cells. *Annu Rev Immunol.* 2013; 31:675–704. Epub 2013/01/22. doi: [10.1146/annurev-immunol-032712-095946](https://doi.org/10.1146/annurev-immunol-032712-095946) PMID: [23330955](https://pubmed.ncbi.nlm.nih.gov/23330955/).
15. Soond DR, Bjorgo E, Moltu K, Dale VQ, Patton DT, Torgersen KM, et al. PI3K p110 delta regulates T-cell cytokine production during primary and secondary immune responses in mice and humans. *Blood.* 2010; 115(11):2203–13. Epub 2010/01/19. blood-2009-07-232330 [pii] doi: [10.1182/blood-2009-07-232330](https://doi.org/10.1182/blood-2009-07-232330) PMID: [20081091](https://pubmed.ncbi.nlm.nih.gov/20081091/).
16. Hawkins PT, Stephens LR. PI3K signalling in inflammation. *Biochim Biophys Acta.* 2015; 1851(6):882–97. Epub 2014/12/17. S1388-1981(14)00258-3 [pii] doi: [10.1016/j.bbali.2014.12.006](https://doi.org/10.1016/j.bbali.2014.12.006) PMID: [25514767](https://pubmed.ncbi.nlm.nih.gov/25514767/).
17. Ali K, Bilancio A, Thomas M, Pearce W, Gilfillan AM, Tkaczyk C, et al. Essential role for the p110delta phosphoinositide 3-kinase in the allergic response. *Nature.* 2004; 431(7011):1007–11. Epub 2004/10/22. nature02991 [pii] doi: [10.1038/nature02991](https://doi.org/10.1038/nature02991) PMID: [15496927](https://pubmed.ncbi.nlm.nih.gov/15496927/).
18. Nashed BF, Zhang T, Al-Alwan M, Srinivasan G, Halayko AJ, Okkenhaug K, et al. Role of the phosphoinositide 3-kinase p110delta in generation of type 2 cytokine responses and allergic airway inflammation. *Eur J Immunol.* 2007; 37(2):416–24. Epub 2007/01/20. doi: [10.1002/eji.200636401](https://doi.org/10.1002/eji.200636401) PMID: [17236236](https://pubmed.ncbi.nlm.nih.gov/17236236/).
19. Randis TM, Puri KD, Zhou H, Diacovo TG. Role of PI3Kdelta and PI3Kgamma in inflammatory arthritis and tissue localization of neutrophils. *Eur J Immunol.* 2008; 38(5):1215–24. Epub 2008/04/17. doi: [10.1002/eji.200838266](https://doi.org/10.1002/eji.200838266) PMID: [18412166](https://pubmed.ncbi.nlm.nih.gov/18412166/).
20. Garcon F, Patton DT, Emery JL, Hirsch E, Rottapel R, Sasaki T, et al. CD28 provides T-cell costimulation and enhances PI3K activity at the immune synapse independently of its capacity to interact with the p85/p110 heterodimer. *Blood.* 2008; 111(3):1464–71. Epub 2007/11/17. blood-2007-08-108050 [pii] doi: [10.1182/blood-2007-08-108050](https://doi.org/10.1182/blood-2007-08-108050) PMID: [18006698](https://pubmed.ncbi.nlm.nih.gov/18006698/).
21. Katz JD, Wang B, Haskins K, Benoist C, Mathis D. Following a diabetogenic T cell from genesis through pathogenesis. *Cell.* 1993; 74(6):1089–100. Epub 1993/09/24. 0092-8674(93)90730-E [pii]. PMID: [8402882](https://pubmed.ncbi.nlm.nih.gov/8402882/).
22. Raine T, Zaccane P, Mastroeni P, Cooke A. Salmonella typhimurium infection in nonobese diabetic mice generates immunomodulatory dendritic cells able to prevent type 1 diabetes. *J Immunol.* 2006; 177(4):2224–33. Epub 2006/08/05. 177/4/2224 [pii]. PMID: [16887982](https://pubmed.ncbi.nlm.nih.gov/16887982/).
23. Wong FS, Karttunen J, Dumont C, Wen L, Visintin I, Pilip IM, et al. Identification of an MHC class I-restricted autoantigen in type 1 diabetes by screening an organ-specific cDNA library. *Nat Med.* 1999; 5(9):1026–31. PMID: [10470079](https://pubmed.ncbi.nlm.nih.gov/10470079/).
24. Wong FS, Siew LK, Scott G, Thomas IJ, Chapman S, Viret C, et al. Activation of insulin-reactive CD8 T-cells for development of autoimmune diabetes. *Diabetes.* 2009; 58(5):1156–64. Epub 2009/02/12. db08-0800 [pii] doi: [10.2337/db08-0800](https://doi.org/10.2337/db08-0800) PMID: [19208910](https://pubmed.ncbi.nlm.nih.gov/19208910/); PubMed Central PMCID: [PMC2671054](https://pubmed.ncbi.nlm.nih.gov/PMC2671054/).

25. Bettelli E, Carrier Y, Gao W, Korn T, Strom TB, Oukka M, et al. Reciprocal developmental pathways for the generation of pathogenic effector TH17 and regulatory T cells. *Nature*. 2006; 441(7090):235–8. Epub 2006/05/02. [nature04753](https://doi.org/10.1038/nature04753) [pii] doi: [10.1038/nature04753](https://doi.org/10.1038/nature04753) PMID: [16648838](https://pubmed.ncbi.nlm.nih.gov/16648838/).
26. Vergani A, D'Addio F, Jurewicz M, Petrelli A, Watanabe T, Liu K, et al. A novel clinically relevant strategy to abrogate autoimmunity and regulate alloimmunity in NOD mice. *Diabetes*. 2010; 59(9):2253–64. Epub 2010/09/02. [db09-1264](https://doi.org/10.2337/db09-1264) [pii] doi: [10.2337/db09-1264](https://doi.org/10.2337/db09-1264) PMID: [20805386](https://pubmed.ncbi.nlm.nih.gov/20805386/).
27. Szot GL, Koudria P, Bluestone JA. Murine pancreatic islet isolation. *J Vis Exp*. 2007;(7):255. Epub 2008/11/08. doi: [10.3791/255](https://doi.org/10.3791/255) PMID: [18989427](https://pubmed.ncbi.nlm.nih.gov/18989427/); PubMed Central PMCID: [PMC2565847](https://pubmed.ncbi.nlm.nih.gov/PMC2565847/).
28. Szot GL, Koudria P, Bluestone JA. Transplantation of pancreatic islets into the kidney capsule of diabetic mice. *J Vis Exp*. 2007;(9):404. Epub 2008/11/08. doi: [10.3791/404](https://doi.org/10.3791/404) PMID: [18989445](https://pubmed.ncbi.nlm.nih.gov/18989445/); PubMed Central PMCID: [PMC2566322](https://pubmed.ncbi.nlm.nih.gov/PMC2566322/).
29. Delong T, Baker RL, He J, Barbour G, Bradley B, Haskins K. Diabetogenic T-cell clones recognize an altered peptide of chromogranin A. *Diabetes*. 2012; 61(12):3239–46. Epub 2012/08/23. [db12-0112](https://doi.org/10.2337/db12-0112) [pii] doi: [10.2337/db12-0112](https://doi.org/10.2337/db12-0112) PMID: [22912420](https://pubmed.ncbi.nlm.nih.gov/22912420/).
30. Bending D, De La Pena H, Veldhoen M, Phillips JM, Uytendhoeve C, Stockinger B, et al. Highly purified Th17 cells from BDC2.5NOD mice convert into Th1-like cells in NOD/SCID recipient mice. *J Clin Invest*. 2009. Epub 2009/02/04. 37865 [pii] doi: [10.1172/JCI37865](https://doi.org/10.1172/JCI37865) PMID: [19188681](https://pubmed.ncbi.nlm.nih.gov/19188681/).
31. Arif S, Moore F, Marks K, Bouckenooghe T, Dayan CM, Planas R, et al. Peripheral and islet interleukin-17 pathway activation characterizes human autoimmune diabetes and promotes cytokine-mediated beta-cell death. *Diabetes*. 2011; 60(8):2112–9. Epub 2011/06/11. [db10-1643](https://doi.org/10.2337/db10-1643) [pii] doi: [10.2337/db10-1643](https://doi.org/10.2337/db10-1643) PMID: [21659501](https://pubmed.ncbi.nlm.nih.gov/21659501/).
32. Durand CA, Richer MJ, Brenker K, Graves M, Shanina I, Choi K, et al. Selective pharmacological inhibition of phosphoinositide 3-kinase p110delta opposes the progression of autoimmune diabetes in non-obese diabetic (NOD) mice. *Autoimmunity*. 2013; 46(1):62–73. Epub 2012/10/09. doi: [10.3109/08916934.2012.732130](https://doi.org/10.3109/08916934.2012.732130) PMID: [23039284](https://pubmed.ncbi.nlm.nih.gov/23039284/).
33. Tai N, Yasuda H, Xiang Y, Zhang L, Rodriguez-Pinto D, Yokono K, et al. IL-10-conditioned dendritic cells prevent autoimmune diabetes in NOD and humanized HLA-DQ8/RIP-B7.1 mice. *Clin Immunol*. 2011; 139(3):336–49. Epub 2011/04/05. [S1521-6616\(11\)00073-8](https://doi.org/10.1016/j.clim.2011.03.003) [pii] doi: [10.1016/j.clim.2011.03.003](https://doi.org/10.1016/j.clim.2011.03.003) PMID: [21458378](https://pubmed.ncbi.nlm.nih.gov/21458378/).
34. Thomas DC, Wong FS, Zaccane P, Green EA, Wallberg M. Protection of islet grafts through transforming growth factor-beta-induced tolerogenic dendritic cells. *Diabetes*. 2013; 62(9):3132–42. Epub 2013/05/30. [db12-1740](https://doi.org/10.2337/db12-1740) [pii] doi: [10.2337/db12-1740](https://doi.org/10.2337/db12-1740) PMID: [23715623](https://pubmed.ncbi.nlm.nih.gov/23715623/).
35. Wallberg M, Wong FS, Green EA. An islet-specific pulse of TGF-beta abrogates CTL function and promotes beta cell survival independent of Foxp3+ T cells. *J Immunol*. 2011; 186(4):2543–51. Epub 2011/01/11. [jimmunol.1002098](https://doi.org/10.4049/jimmunol.1002098) [pii] doi: [10.4049/jimmunol.1002098](https://doi.org/10.4049/jimmunol.1002098) PMID: [21217013](https://pubmed.ncbi.nlm.nih.gov/21217013/).
36. Ali K, Soond DR, Pineiro R, Hagemann T, Pearce W, Lim EL, et al. Inactivation of PI(3)K p110delta breaks regulatory T-cell-mediated immune tolerance to cancer. *Nature*. 2014; 510(7505):407–11. Epub 2014/06/12. [nature13444](https://doi.org/10.1038/nature13444) [pii] doi: [10.1038/nature13444](https://doi.org/10.1038/nature13444) PMID: [24919154](https://pubmed.ncbi.nlm.nih.gov/24919154/).
37. Patton DT, Garden OA, Pearce WP, Clough LE, Monk CR, Leung E, et al. Cutting edge: the phosphoinositide 3-kinase p110 delta is critical for the function of CD4+CD25+Foxp3+ regulatory T cells. *J Immunol*. 2006; 177(10):6598–602. Epub 2006/11/04. [177/10/6598](https://doi.org/10.1093/infdis/jih257) [pii]. PMID: [17082571](https://pubmed.ncbi.nlm.nih.gov/17082571/).