Anti-proliferative activity of oral anti-hyperglycemic agents on human vascular smooth muscle cells: thiazolidinediones (glitazones) have enhanced activity under high glucose conditions

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Abstract

Background: Inhibition of vascular smooth muscle cell (vSMC) proliferation by oral anti-hyperglycemic agents may have a role to play in the amelioration of vascular disease in diabetes. Thiazolidinediones (TZDs) inhibit vSMC proliferation but it has been reported that they anomalously stimulate [3H]-thymidine incorporation. We investigated three TZDs, two biguanides and two sulfonylureas for their ability of inhibit vSMC proliferation. People with diabetes obviously have fluctuating blood glucose levels thus we determined the effect of media glucose concentration on the inhibitory activity of TZDs in a vSMC preparation that grew considerably more rapidly under high glucose conditions. We further explored the mechanisms by which TZDs increase [3H]-thymidine incorporation.

Methods: VSMC proliferation was investigated by [3H]-thymidine incorporation into DNA and cell counting. Activation and inhibition of thymidine kinase utilized short term [3H]-thymidine uptake. Cell cycle events were analyzed by FACS.

Results: VSMC cells grown for 3 days in DMEM with 5% fetal calf serum under low (5 mM glucose) and high (25 mM glucose) increased in number by 2.5 and 4.7 fold, respectively. Rosiglitazone and pioglitazone showed modest but statistically significantly greater inhibitory activity under high versus low glucose conditions (P < 0.05 and P < 0.001, respectively). We confirmed an earlier report that troglitazone (at low concentrations) causes enhanced incorporation of [3H]-thymidine into DNA but did not increase cell numbers. Troglitazone inhibited serum mediated thymidine kinase induction in a concentration dependent manner. FACS analysis showed that troglitazone and rosiglitazone but not pioglitazone placed a slightly higher percentage of cells in the S phase of a growing culture. Of the biguanides, metformin had no effect on proliferation assessed as [3H]-thymidine incorporation or cell numbers whereas phenformin was inhibitory in both assays albeit at high concentrations. The sulfonylureas chlorpropamide and gliclazide had no inhibitory effect on vSMC proliferation assessed by either [3H]-thymidine incorporation or cell numbers.

Conclusion: TZDs but not sulfonylureas nor biguanides (except phenformin at high concentrations) show favorable vascular actions assessed as inhibition of vSMC proliferation. The activity of rosiglitazone and pioglitazone is enhanced under high glucose conditions. These data provide further in vitro evidence for the potential efficacy of TZDs in preventing multiple cardiovascular diseases. However, the plethora of potentially beneficial actions of TZDs in cell and animal models have not been reflected in the results of major clinical trials and a greater understanding of these complex drugs is required to delineate their ultimate clinical utility in preventing macrovascular disease in diabetes.

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Background
The role of vascular smooth muscle cell (vSMC) proliferation in vascular disease, particularly atherosclerosis, is controversial and unresolved [1]. However, emerging information is identifying the situations such as post-angioplasty restenosis in people with diabetes in which hyperproliferation is clearly critical in determining the clinical outcome [2]. Although coronary artery by-pass grafting (CABG) was initially the preferred intervention over angioplasty in people with diabetes and coronary artery disease [3] the introduction of coronary artery stents and drug coated stents and possibly supplemented with systemic therapy has raised the possibility that this less invasive treatment may be suitable for this population [4,5]. Although factors such as proteoglycan mediated lipid deposition [6,7] and inflammation [8,9] are clearly important in the process of atherosclerosis and restenosis, in the setting of diabetes vSMC proliferation is clearly critical and thus a target for therapy [2]. As people with diabetes clearly have ongoing hyperglycaemia after a clinical intervention for coronary artery disease (CAD), the role of the anti-hyperglycemic therapy in providing a complementary action to prevent vSMC cell proliferation is of potential therapeutic interest [10]. It is further possible that an oral anti-proliferative agent may also be useful as adjunct therapy following vascular intervention even in the absence of diabetes [11].

We have made a direct comparison of the inhibitory activity of the three major classes of oral anti-hyperglycemic agents thiazolidinediones (TZDs) also known as glitazones, biguanides and sulphonylureas towards vSMC proliferation. Further, we used multiple assays to evaluate the mechanism of inhibition and addressed the clinically relevant question of the effect of glucose concentration on the inhibitory activity of the TZDs.

The data shows that only TZDs show appreciable inhibitory activity towards vSMC proliferation amongst currently used oral anti-hyperglycemic agents. Furthermore, under high glucose conditions in which vSMC proliferation is markedly enhanced, the inhibitory potency of the clinical TZDs, rosiglitazone and pioglitazone, is increased not diminished. We also reveal an action of TZDs to stimulate [3H]-thymidine incorporation secondary to stimulation of uptake suggesting that other assays of proliferation are more suitable for studies with this class of drug.

Methods

Materials
Phenformin, metformin, chlorpropamide, dimethylsulfoxide (DMSO), platelet-derived growth factor (PDGF), Whatman 3 MM chromatography paper and DEAE-Sephacel were obtained from Sigma Aldrich (St Louis, MO). Gliclazide was provided by Institute de Recherches Internationale Servier (France). Troglitazone (Rezulin, Parke-Davis/Warner Lambert), rosiglitazone (Avandia, GlaxoSmithKline) and pioglitazone (Actos, Takeda) were provided as gifts by Parke Davis Pharmaceutical Research (Ann Arbor, MI), GlaxoSmithKline (Boronia, Australia) and Eli Lilly (West Ryde, Australia), respectively. Foetal bovine serum (FBS) was obtained from CSL (Parkville, Australia). [3H]-Thymidine and cetylpyridinium chloride (CPC) were from ICN Biomedicals Inc. (Irvine, CA). Scintillation fluid Instagel was from Packard (Groningen, The Netherlands).

Cell culture preparations
Human vSMCs were isolated using the explant technique from discarded segments of the saphenous veins and internal mammary arteries from a variety of patient donors undergoing surgery at the Alfred Hospital (Melbourne, Australia), the acquisition of which was approved by the Alfred Hospital Ethics committee. Cells from both sources were used for these experiments and in accord with our recent data [12] there were no systematic differences in the results obtained with cells from either vascular origin.

Analysis of mitogenic response by 3H-thymidine incorporation
Cells were seeded at 4.5 × 10^4 cells per well in a 24 well plate in DMEM containing 5 mM glucose with 10% FBS for 24 h. Cells were serum deprived for 48 h followed by treatment with control media (0.1% DMSO) or treatment media containing the anti-hyperglycaemic agents in the presence of PDGF (50 ng/ml) and incubated overnight at 37°C in 5% CO₂. Cells were labelled with [3H]-thymidine (1 μCi/ml) or 3 h and assessed for [3H] incorporation into newly synthesized DNA as previously described [13].

Analysis of cell proliferation by assessment of cell number
Human vSMCs in the log growth phase were treated with anti-hyperglycaemic agents (10–300 μM) in the presence of 5% serum for 72 h and assessed for cell number as previously described [12].

Assessment of thymidine kinase induction by acute [3H]-thymidine uptake
Acute [3H]-thymidine uptake was assessed as previously described [14]. Briefly, vSMCs were grown to confluency in 24 well plates and serum deprived for 24 h in DMEM (5 mM glucose) and 0.1% FCS. Growth factors and drugs as indicated were added for 18 h. The cells were washed once with DMEM (5 mM glucose) prior to an equilibrating 30 min incubation in DMEM (5 mM glucose). [3H]-Thymidine (1 μCi/ml) was added and uptake proceeded for 4 mins at 37°C. At the end of the incubation period, the cells were immediately placed on ice and rapidly washed with ice cold PBS. The radioactivity was released
into solution by 1 M NaOH. The amount of [3H]-thymidine was quantitated by liquid scintillation counting.

**Cell cycle analysis by Fluorescence Activated Cell Sorting (FACS)**

Cells were seeded at 2.5 × 10^5 cells per 90 mm plate in duplicate in DMEM containing 5 mM glucose with 10% FBS for 24 h. Cells were serum deprived for 48 h followed by treatment with control media (0.1% DMSO) or treatment media containing the anti-hyperglycaemic agents in the presence of PDGF (50 ng/ml) in the presence of 5% serum and incubated for 72 h at 37°C in 5% CO₂. For analysis of S phase entry and cells were labelled with 10 μM 5-bromo-2’-deoxyuridine (BrdU, Sigma, St. Louis, MO) for 4 hr at 37°C in the culture medium. BrdU incorporation was detected using FITC-conjugated anti-BrdU (Becton Dickinson) and DNA was stained with propidium iodide (5 μg/ml) 30 min at room temperature in the dark. Flow cytometry and cell cycle analysis was carried out with a FACS Calibur (Becton Dickinson) with 488 nm excitation.

**Statistical analyses**

Data are presented as mean ± standard error of the mean (SEM) from 2–5 experiments as indicated in each individual figure. Data were statistically analysed using a Student’s t test or 1-way ANOVA.

**Results**

**Inhibition of vSMC proliferation by TZDs – dependence of the inhibitory activity of TZDs on the culture media glucose concentration**

People with diabetes have blood glucose concentrations that may range at times from 2 to 30 mM and thus the effect of glucose concentration on the actions of antihyperglycaemic agents is clinically relevant. We identified a human vSMC preparation that showed enhanced proliferation in response to high glucose media. Under “low” glucose conditions which actually equate to normoglycaemia (i.e. DMEM with 5 mM glucose) and in the presence of 5 per cent FBS, cell numbers over 3 days increased from 36,800 ± 3,800 to 91,900 ± 8,500 cells/well (mean ± sem, n = 5 experiments in triplicate). Under high glucose conditions (DMEM containing 25 mM glucose) and in the presence of 5 per cent FBS, vSMC numbers increased from 54,400 ± 5,500 to 250,500 ± 28,000 cells/well (mean ± sem, n = 5 experiments in triplicate). We used this cell model to assess the inhibitory activity of the two clinical TZDs, rosiglitazone and pioglitazone. Rosiglitazone was evaluated at 3–100 μM and pioglitazone 0.3–30 μM noting that pioglitazone precipitates in the culture media at concentration above 30 μM. The inhibitory effects of rosiglitazone in low and high glucose media were normalised to the control which was set as 100% (Fig. 1A and 1B, respectively). The data is presented as a dose response curve with statistical comparison of the curves (Fig. 1C). The data shows that the inhibitory effect of rosiglitazone is enhanced (P < 0.001) under high glucose condition (Fig. 1C). Similar data for the effect of pioglitazone is shown in (Figs 1D, E and 1F). Pioglitazone showed a greater inhibitory potency (P < 0.001) in high compared to low glucose DMEM.

**Evaluation of the effect of TZDs as assessed by [3H]-thymidine incorporation**

Multiple assays have been used to evaluate the inhibitory effects of TZDs on vSMC proliferation. Takeda et al. [15] have reported that TZDs cause a marked but anomalous increase in [3H]-thymidine incorporation in vSMCs and it was further suggested that this is associated with a hyperproliferative response [15]. We investigated the effects of troglitazone, the TZD used by those authors, on the [3H]-thymidine incorporation by human vSMCs. Troglitazone had a bi-phasic effect on [3H]-thymidine incorporation into DNA with stimulation at low concentrations and inhibition at higher concentrations (Fig. 2). We investigated the mechanism of this increase in [3H]-thymidine incorporation which is at variance with the inhibition observed when inhibition of proliferation is assessed by cell counting as we previously described [12].

Activation of [3H]-thymidine incorporation into DNA of proliferating cells arises from growth factor mediated activation of thymidine kinase activity and enhanced [3H]-thymidine uptake into the salvage pathway of thymidylate synthase which provides adequate precursors for DNA synthesis in vSMC [14]. We hypothesised that troglitazone may have a growth factor like effect (gene induction) on thymidine kinase leading to enhanced uptake and consequently increased incorporation of [3H]-thymidine into DNA. Thymidine kinase activity can be assessed as the uptake by the cell of [3H]-thymidine over a period of several minutes [14]. We treated vSMCs with low (0.3 μM) and high (10 μM) concentrations of troglitazone which stimulate and inhibit, respectively, [3H]-thymidine incorporation into DNA precipitable material (Fig. 2). We undertook the acute (3 min) uptake experiment concomitant with an examination of the effect of troglitazone on incorporation of [3H]-thymidine into DNA in a parallel experiment as described above. Troglitazone caused concentration dependent inhibition of the acute uptake of [3H]-thymidine (Fig. 3A) with no evidence of stimulation. The parallel experiment for assessing [3H]-thymidine incorporation into DNA produced the expected result of enhanced [3H]-thymidine incorporation at low concentration of troglitazone (0 – 3 μM) and inhibition of incorporation at high troglitazone concentrations (10 nM) (Fig. 3B).
Figure 1
Effect of media glucose concentration on the inhibitory activity of TZDs. Panels A and B show the effect of rosiglitazone under low (5 mM) glucose (A) and high glucose (25 mM) and the data shown as a dose response curve (C). Data for pioglitazone is shown in panels D, E and F. Data represent the mean ± SEM from 3 experiments in triplicate *P < 0.05, ***,P < 0.001
Troglitazone treatment of vSMC increases [\(^3\)H]-thymidine incorporation into newly synthesized DNA which implies that more cells are in the S phase of the cell cycle. We used FACS to investigate the effects of the three TZDs on cell cycle progression in vSMCs. The treatment of vSMCs with troglitazone and rosiglitazone resulted in a small increase in cells in the S phase (Fig. 4). In contrast, treatment of cells with pioglitazone reduced the proportion of cells in the S phase of the cell cycle (Fig. 4). The data indicates that the [\(^3\)H]-thymidine incorporation truly reflects an increased number of cells in the S phase in the presence of rosiglitazone and troglitazone however this does not result in cell cycle progression as no increase in cell numbers is observed [12].

**Effect of biguanides on vSMC proliferation**

The effects of metformin and phenformin were assessed as described above. Although phenformin has been withdrawn from clinical use, its use also allowed us to assess two compounds in the biguanide class. Metformin had no inhibitory effect on [\(^3\)H]-thymidine incorporation stimulated by PDGF and indeed a small anomalous stimulatory effect at 30 \(\mu\)M that attained statistical significance (Fig. 5A). Phenformin (10, 30, 100 \(\mu\)M) caused a partially concentration dependent inhibitory response with a half maximal inhibitory concentration of approximately 30 \(\mu\)M (Fig. 5A). In terms of vascular pathology the response of most interest is that of changes in cell number. Thus we examined the effect of metformin and phenformin on serum-stimulated increases in cell number. Metformin had no effect on cell numbers but phenformin caused a concentration dependent reduction in cell proliferation with a half maximally inhibitory concentration between 30 and 100 \(\mu\)M (Fig. 5B). There were no apparent toxic effects of phenformin on the cells as assessed by lactate...
Assessment of the cell cycle state of human vSMCs following treatment with clinical TZDs. S phase fraction in cultures treated with TZDs either Troglitazone, Rosiglitazone or Pioglitazone (1 μM each) in the presence of PDGF (50 ng/mL) for 24 h as compared with untreated controls. S phase fraction was analysed by flow cytometry using DNA staining with propidium iodide.

dehydrogenase release and phase contrast microscopy (data not shown).

Effect of sulphonylureas on vSMC proliferation

We examined the effect of two representative sulfonylureas on the inhibition of cell cycle S-phase [3H]-thymidine incorporation into human vSMCs stimulated by PDGF [16]. Neither chlorpropamide (10, 30, 100 μM) nor glipizide (10, 30, 100 μM) inhibited the ability of PDGF to stimulate [3H]-thymidine incorporation even at high concentrations (data not shown). We also assessed the effect of chlorpropamide (3–100 μM) and gliclazide (3–100 μM) to inhibit proliferation assessed by cell counting. Chlorpropamide and gliclazide had no effect on cell proliferation stimulated by 5 per cent fetal bovine serum (FBS) assessed after 3 days proliferation (data not shown).

Discussion

We have systematically evaluated the effect of the three major classes of oral anti-hyperglycaemic agents for their ability to inhibit vSMC proliferation using the techniques of [3H]-thymidine incorporation into DNA and cell counting plus supporting mechanistic studies. We confirmed our earlier report that all three clinically used TZDs cause concentration dependent inhibition of vSMC proliferation assessed by cell counting [12]. Utilizing human vSMCs that showed glucose dependent cell proliferation (accelerated growth under high glucose conditions) we demonstrated that the inhibitory potency of rosiglitazone and pioglitazone was slightly but statistically significantly greater under high glucose conditions. A representative TZD, troglitazone showed an unexpected but previously observed effect to stimulate [3H]-thymidine incorporation into DNA at low concentrations with clear and marked inhibition at high concentrations. We investigated the possibility that the stimulation was secondary to induction of thymidine kinase activity and thus acute stimulation of [3H]-thymidine uptake but this was not the mechanism. However, cell cycle analysis by FACS confirmed that the increase in [3H]-thymidine incorporation correlated with the increase in cells in the S phase of the
cell cycle. Two sulfonylureas showed no inhibitory activity in the former assay and were not further investigated. Of the two biguanides only phenformin, which is not now clinically used, showed inhibitory activity and this was expressed in both the assay of \(^{3}\text{H}\)-thymidine incorporation into DNA and cell counting. Thus of the clinically relevant oral hypoglycemic agents only the TZDs showed inhibition of vSMC proliferation.

The major clinical implication of these studies relates to the role of vSMC proliferation in cardiovascular disease. The most prominent finding is that the diabetic milieu leads to hyper proliferation of vSMCs such that in people with diabetes proliferation limits the efficacy of vascular interventions due to hyperplasia of vSMCs [2]. Such proliferation also occurs in animal models such as the streptozotocin treated mini pigs in which a clear development of vascular hyperplasia occurs in association with increased expression of inflammatory markers [17]. Thus, there may be a role for choosing an oral hypoglycemic agent with anti-proliferative actions to provide additional potential benefit for people with diabetes. Indeed rosiglitazone has been shown to limit the increase in intimal medial thickening and restenosis in people with diabetes [4,11]. Our demonstration that the anti-proliferative action of TZDs is maintained even enhanced in a high glucose environment favors the use of these agents for glucose lowering in people with diabetes although the overall impact of these agents on cardiovascular disease is the subject of considerable controversy as addressed later [18,19].

The impact of hyperglycemia on the actions of TZDs on cardiovascular cells is an important aspect of the action of anti-hyperglycemic agents, particularly for TZDs which are being considered for wider use in the cardiovascular setting. We are unaware of any reports of the effect of glucose concentration on the cellular action of clinical TZDs in any in vitro setting. There are many examples where high glucose stimulates responses relevant to diabetes proliferation limits the efficacy of vascular interventions due to hyperplasia of vSMCs [2]. Such proliferation also occurs in animal models such as the streptozotocin treated mini pigs in which a clear development of vascular hyperplasia occurs in association with increased expression of inflammatory markers [17]. Thus, there may be a role for choosing an oral hypoglycemic agent with anti-proliferative actions to provide additional potential benefit for people with diabetes. Indeed rosiglitazone has been shown to limit the increase in intimal medial thickening and restenosis in people with diabetes [4,11]. Our demonstration that the anti-proliferative action of TZDs is maintained even enhanced in a high glucose environment favors the use of these agents for glucose lowering in people with diabetes although the overall impact of these agents on cardiovascular disease is the subject of considerable controversy as addressed later [18,19].

The cardiovascular effects and potential negative outcomes associated with rosiglitazone therapy have been the subject of recent controversy [18]. A meta analysis showed that rosiglitazone therapy was associated with statistically significantly higher rates of myocardial infarction and cardiovascular death [18]. Notably a similar analysis showed that this apparently higher risk of cardiovascular misadventure did not apply to the other clinical TZD, pioglitazone, for which a lower risk of cardiovascular events was calculated [24]. However, an alternative analysis of the rosiglitazone data, an analysis considering for example the role of omitting studies in which there were zero cardiovascular events has shown that the odds ratio for excess cardiovascular events associated with rosiglitazone therapy is much lower and not statistically significant [19]. The blood glucose lowering action of TZDs are beneficial in terms of reductions in microvascular complications which are very closely linked to long term estimates of glycaemia [25-27]. However, more than glucose lowering actions are required to manifest beneficial outcomes on cardiovascular disease such as strokes and heart attacks [28]. The lack of human efficacy of TZDs towards cardiovascular disease in initial trials [29] is surprising in view of the overwhelmingly positive occurrence of beneficial actions in cell and animal models for these compounds [30-33]. Nevertheless, human therapeutic experience is
the definitive factor and further results from clinical trials will provide information and guidance to the ultimate usefulness of this class of compound [34]. In terms of macrovascular disease the major contributing factors are inflammation, oxidation and the retention of atherogenic lipoproteins by extracellular matrix molecules particularly the proteoglycan, biglycan [35,36]. The role of proteoglycans was demonstrated recently in a human pathology study in coronary arteries in which it was shown that atherosclerosis commences with the deposition lipoproteins associated with the expression of biglycan in the outer layer of the diffuse intimal thickenings [7]. We recently reported on the action of oral anti-hyperglycaemic agents to modify the synthesis and structure of proteoglycans and in accord with the present study it was shown that TZDs are the only group with direct vascular actions [37]. In the present study we observed that the biguanide phenformin inhibited vSMC proliferation and that finding is in accord with our earlier observation that it inhibits protein synthesis in these cells [37]. Metformin appears to be the only non-TZD that shows favourable outcomes on macrovascular disease UKPDS [38] and which may be associated with the important role of insulin resistance in macrovascular disease [39].

Conclusion
This data supports numerous studies in which TZDs have appreciable direct actions on vSMCs that if expressed in vivo would be beneficial in reducing cardiovascular disease. Our finding that the action of TZDs is enhanced under high glucose conditions is also positive for the use of these compounds in the setting of diabetes. Further understanding is required to match the clinical utility of these agents with the multitude of in vitro and in vivo data which supports a role in the prevention of cardiovascular disease associated with diabetes.

Abbreviations
CABG: Coronary artery bypass grafting
CAD: Coronary artery disease
DMSO: Dimethylsulfoxide
TZDs: Thiazolidinediones
vSMCs: Vascular smooth muscle cells

Competing interests
PJL has received honoraria and expenses in relation to presentations at meetings sponsored by GSK. PJL received a GSK post-graduate support award (226/07). JN received travel support from GSK to attend and present related data at scientific meetings.

Authors’ contributions
PJL conceived the study and wrote the manuscript. NO and MB undertook the FACS experiments and assisted with the preparation of the manuscript. SD, NC and JN undertook various experiments.

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References
1. Ross R: The pathogenesis of atherosclerosis. N Engl J Med 1986, 314:488-500.
2. Kornowski R, Mintz GS, Kent KM, Pichard AD, Satler F, Bucher TA, Hong MK, Popma JJ, Leon MB: Increased restenosis in diabetes mellitus after coronary interventions is due to exaggerated intimal hyperplasia. Circulation 1997, 95:1366-1369.
3. Ferguson JJ: NHLBI BARI clinical alert on diabetes treated with angioplasty. Circulation 1995, 92:3371.
4. Choi D, Kim SK, Choi SH, Ko YG, Ahn CW, Jang Y, Lim SK, Lee HC, Cha BS: Preventative effects of rosiglitazone on restenosis after coronary stent implantation in patients with type 2 diabetes. Diabetes Care 2004, 27:2654-60.
5. Takagi T, Akasaka T, Yamamuro A, Honda Y, Hozumi T, Morokusa K, Yoshida K: Troglitazone reduces neointimal tissue proliferation after coronary stent implantation in patients with non-insulin dependent diabetes mellitus: a serial intravascular ultrasound study. J Am Coll Cardiol 2000, 36:529-35.
6. Williams KJ, Tabas I: The response-to-retention hypothesis of early atherosclerosis. Arterioscler Thromb Vasc Biol 1995, 15:531-61.
7. Nakashima Y, Fuji H, Sumiyoshi S, Wight TN, Sueishi K: Early human atherosclerosis: accumulation of lipid and proteoglycans in intimal thickenings followed by macrophage infiltration. Arterioscler Thromb Vasc Biol 2007, 27:1159-65.
8. Libby P: Inflammation in atherosclerosis. Nature 2002, 420:868-74.
9. Ross R: Atherosclerosis – an inflammatory disease. N Engl J Med 1999, 340:115-26.
10. Weissman P: Reappraisal of the pharmacologic approach to treatment of type 2 diabetes mellitus. Am J Cardiol 2002, 90:42G-50G.
11. Sidhu JS, Kaposzta Z, Markus HS, Kaski JC: Effect of Rosiglitazone on Common Carotid Intima-Media Thickness Progression in Coronary Artery Disease Patients Without Diabetes Mellitus. Arterioscler Thromb Vasc Biol 2004, 24:930-934.
12. de Dios ST, Bremmer D, Dilley RJ, Ivey ME, Jennings GL, Law RE, Little PJ: Inhibitory activity of clinical thiazolidinedione peroxi-some proliferator activating receptor-gamma ligands toward internal mammary artery, radial artery, and saphenous vein smooth muscle cell proliferation. Circulation 2003, 107:2548-50.
13. Little PJ, Allen TJ, Hashimura K, Nigro J, Farrellly CA, Dilley RJ: High glucose potentiates mitogenic responses of cultured ovine coronary smooth muscle cells to platelet derived growth fac-tor and transforming growth factor-beta1. Diabetes Res Clin Pract 2003, 59:93-101.
14. Agrotis A, Little PJ, Salits J, Bobik A: Dihydropyridine Ca2+ channel antagonists inhibit the salvage pathway for DNA synthesis in human vascular smooth muscle cells. Eur J Pharmacol 1993, 244:269-75.
15. Takeda K, Ichiki T, Tokunou T, Iino N, Takeshita A: 15-Deoxy-delta 12,14-prostaglandin J2 and thiazolidinediones activate the...
MEK/ERK pathway through phosphatidylinositol 3-kinase in vascular smooth muscle cells. J Biol Chem 2001, 276:48950-5.

Ross R, JA G, Kariya B, Harker L: A platelet-dependent serum factor that stimulates the proliferation of arterial smooth muscle cells in vitro. Proc Natl Acad Sci USA 1974, 71:1207-1210.

Zhang Q, Lu L, Pu L, Zhang R, Shen J, Zhu Z, Hu J, Yang Z, Chen Q, Shen W: Neointimal hyperplasia persists at six months after sirolimus-eluting stent implantation in diabetic porcine. Cardiovasc Diabetology 2007, 6:16.

Nissen SE, Wolski K: Effect of rosiglitazone on the risk of myocardial infarction and death from cardiovascular causes. N Engl J Med 2007, 356:2457-71.

Diamond GA, Bax L, Kaul S: Uncertain Effects of Rosiglitazone on the Risk for Myocardial Infarction and Cardiovascular Death. Ann Intern Med 2007.

Sato A, Sasaoka T, Yamazaki K, Nakamura N, Temaru R, Ishiki M, Takata M, Kishida M, Wada T, Ishihara H, et al.: Glucosamine enhances platelet-derived growth factor-induced DNA synthesis via phosphatidylinositol 3-kinase pathway in rat aortic smooth muscle cells. Atherosclerosis 2001, 157:341-52.

Ling S, Little PJ, Williams MR, Dai A, Hashimura K, Liu JP, Komesaroff P, et al.: Effect of 17beta-estradiol in human vascular smooth muscle cells. Am J Physiol Endocrinol Metab 2002, 282:E746-51.

Gomperts BD, Kramer IM, Tatham PER: Signal Transduction. San Diego: Elsevier Academic Press; 2004.

Sperti G, Colucci WS: Calcium influx modulates DNA synthesis and proliferation in A7r5 vascular smooth muscle cells. Eur J Pharmacol 1991, 206:279-84.

Lincoff AM, Wolski K, Nicholls SJ, Nissen SE: Pioglitazone and risk of cardiovascular events in patients with type 2 diabetes mellitus: a meta-analysis of randomized trials. JAMA 2007, 298:1180-8.

DCCT: The absence of a glycemic threshold for the development of long-term complications: the perspective of the Diabetes Control and Complications Trial. Diabetes 1996, 45:1289-1298.

Wei M, Haffner SM, Gaskill SP, Stern MP: Effects of diabetes and level of glycemia on all-cause and cardiovascular mortality. Diabetes Care 1998, 21:1167-1172.

Vasudevan AR, Burns A, Fonseca VA: The effectiveness of intensive glycemic control for the prevention of vascular complications in diabetes mellitus. Treat Endocrinol 2006, 5:273-86.

UK Prospective Diabetes Study Group: Tight blood pressure control and risk of macrovascular and microvascular complications in type 2 diabetes: UKPDS 38. Br Med J 1998, 317:703-13.

Berrimandy JA, Charbonnel B, Eckland DJ, Erdmann E, Massi-Benedetti M, Moules IK, Skene AM, Tan MH, Lefebvre PJ, Murray GD, et al.: Secondary prevention of macrovascular events in patients with type 2 diabetes in the PROactive Study (PROspective pioglitAzone Clinical Trial In macroVascular Events): a randomised controlled trial. Lancet 2005, 366:1279-89.

Plutzky J: Medicine. PPARs as therapeutic targets: reverse cardiology? Science 2003, 302:406-7.

Marx N, Schonbeck U, Lazar MA, Libby P, Plutzky J: Peroxisome proliferator-activated receptor gamma activators inhibit gene expression and migration in human vascular smooth muscle cells. Circulation Research 1998, 83:1097-1103.

Collins AR, Meehan WP, Kintscher U, Jackson S, Wakino S, Noh G, Palinski W, Hsueh WA, Law RE: Troglitazone inhibits formation of early atherosclerotic lesions in diabetic and nondiabetic low density lipoprotein receptor-deficient mice. Arterioscler Thromb Vasc Biol 2001, 21:365-71.

Calkin AC, Forbes JM, Smith CM, Lassila M, Cooper ME, Jandeleit-Dahm KA, Allen TJ: Rosiglitazone attenuates atherosclerosis in a model of insulin insufficiency independent of its metabolic effects. Arterioscler Thromb Vasc Biol 2005, 25:1903-9.

Nigro J, Ballinger ML, Osman N, Dart AM, Little PJ: Anti-atherogenic role of peroxisome proliferator activated receptor ligands. Current Cardiology Reviews 2005, 1:89-102.

Libby P: Inflammation and cardiovascular disease mechanisms. Am J Clin Nutr 2006, 83:4565-4605.

Little PJ, Ballinger ML, Osman N: Vascular wall proteoglycan synthesis and structure as a target for the prevention of atherosclerosis. Vascular Health Risk Management 2007, 3:1-8.

37. de Dios ST, Frontanilla KV, Nigro J, Ballinger ML, Ivey ME, Cawson EA, Little PJ: Regulation of the atherogenic properties of vascular smooth muscle proteoglycans by oral anti-hyperglycemic agents. J Diabetes Complications 2007, 21:108-17.

38. UK Prospective Diabetes Study (UKPDS) Group: Effect of intensive blood-glucose control with metformin on complications in overweight patients with type 2 diabetes (UKPDS 34). Lancet 1998, 352:854-65.

39. Nigro J, Osman N, Dart AM, Little PJ: Insulin Resistance and Atherosclerosis. Endocr Rev 2006, 27:242-259.
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Little, PJ; Osman, N; de Dios, ST; Cemerlang, N; Ballinger, M; Nigro, J

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