

Review Article

Risk of Cancer in Diabetes: The Effect of Metformin

Mojtaba Malek, Rokhsareh Aghili, Zahra Emami, and Mohammad E. Khamseh

*Endocrine Research Center (Firouzgar), Institute of Endocrinology and Metabolism,
Tehran University of Medical Sciences, Tehran 15937-48711, Iran*

Correspondence should be addressed to Mohammad E. Khamseh; m-khamseh@tums.ac.ir

Received 20 July 2013; Accepted 16 August 2013

Academic Editors: A. Kautzky-Willer and N. Tentolouris

Copyright © 2013 Mojtaba Malek et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Cancer is the second cause of death. Association of diabetes as a growing and costly disease with cancer is a major health concern. Meanwhile, preexisting diabetes is associated with an increased risk of all-cause and cancer-specific mortalities. Presence of diabetes related comorbidities, poorer response to cancer treatment, and excess mortality related to diabetes are among the most important explanations. Although diabetes appear to be a risk factor for cancer and is associated with the mortality risk in cancer patients, several factors such as diabetes duration, multiple drug therapy, and the presence of diabetes comorbidities make the assessment of the effect of diabetes treatment on cancer risk and mortality difficult. Metformin is the drug of choice for the treatment of type 2 diabetes. The available evidence from basic science, clinical, and population-based research supports the anticancer effect of metformin. However, randomized controlled clinical trials do not provide enough evidence for a strong protective effect of metformin on cancer incidence or mortality. One of the most important limitations of these trials is the short duration of the followup. Further long-term randomized controlled clinical trials specifically designed to determine metformin effect on cancer risk are needed to provide the best answer to this challenge.

1. Diabetes and Cancer

The prevalence of diabetes in newly diagnosed cancer patients is reported to be from 8% to 18% [1]. A meta-analysis of 12 cohort studies reported a significant pooled adjusted risk ratio (RR) of all-cancer incidence (RR = 1.10, 95% CI, and 1.04 to 1.17). The increased risk was observed for both men (RR = 1.14, CI, and 1.06 to 1.23) and women (RR = 1.18, CI, and 1.08 to 1.28) [2].

An increase for site-specific cancer incidence has also been described in many systematic reviews and meta-analyses. Deng et al. reported a 26% increase in the incidence of colorectal cancer. The rate was similar in both women and men. These results were obtained from eight case-control and 16 cohort studies without heterogeneity between studies ($P = 0.296$) [3]. Parallel to these results, Jiang et al. also described an increased incidence of colorectal cancer in a systematic review of 41 cohort studies (summary of relative risk 1.27, 95% CI: 1.21–1.34) [4]. Although the studies showed significant heterogeneity ($P = 0.002$, $I^2 = 48.4\%$), the incidence seems not to be affected by sex (RR = 1.25, 95% CI: 1.19–1.14 in men, and RR = 1.34, 95% CI: 1.22–1.47 in women) [5].

Breast cancer has also been shown to be more prevalent in women with type 2 diabetes (SRR = 1.27, 95% CI: 1.16–1.39). In contrast, type 1 diabetes seems not to be associated with the risk of breast cancer (SRR 1.00, 95% CI: 0.74–1.3). In type 2 diabetes, the risk was lower in prospective (23%) than in retrospective studies (36%). Adjustment for BMI showed a lower risk (16%), compared with nonadjustment (33%) [6].

Another meta-analysis included 38 studies investigating breast cancer incidence in women, and 5 studies investigating the incidence in men found a significantly increased risk in both women (RR 1.24, 95% CI: 1.15–1.35) and men (RR 1.40, 95% CI: 1.10–1.79) [7].

Two meta-analyses explored the incidence of gastric cancer in people with type 2 diabetes in case-control and cohort studies [8, 9]. A slight increase in the incidence of gastric cancer was reported, although the risk seems to be affected by sex. Women with diabetes showed an 18% increase in risk compared with nondiabetic individuals, while the risk of gastric cancer was similar for diabetic men compared to nondiabetic men [8].

Stevens et al. conducted a systematic review of three cohort and six case-control studies and reported pancreatic

TABLE 1: Cancer risks in diabetes.

First author, year of publication	Study method [reference]	Sample	Risk of cancer among DM participants (95% CI)
Noto, 2011	Meta-analysis [2]	12 cohort studies	Men RR = 1.14 (1.06–1.23) Women RR = 1.18 (1.08–1.28)
Deng, 2012	Systematic review and meta-analysis [3]	8 case-control studies 16 cohort studies	RR = 1.26 (1.20–1.31)
Jiang, 2011	Systematic review and meta-analysis [4]	30 cohort studies	Summary RR = 1.27 (1.21–1.34)
Krämer, 2012	Meta-analysis [5]	8 case-control studies 21 cohort studies	Men RR = 1.29 (1.19–1.40) Women RR = 1.34 (1.22–1.47).
Boyle, 2012	Meta-analysis [6]		Summary RR = 1.27 (1.16–1.39)
Hardefeldt, 2012	Meta-analysis [7]	43 studies	Men RR = 1.40 (1.10–1.79) Women RR = 1.24 (1.15–1.35)
Tian, 2012	Meta-analysis [8]	7 case-control studies 18 cohort studies	Summary RR = 1.11 (1.00–1.24)
Ge, 2011	Meta-analysis [9]	4 case-control studies 17 cohort studies	Summary RR = 1.09 (0.98–1.22) Women Summary RR = 1.18 (1.01–1.39)
Stevens, 2007	Systematic review and meta-analysis [10]	6 case-control studies 3 cohort studies	RR = 2.00 (1.37–3.01)
Ben, 2011	Meta-analysis [11]	35 cohort studies	Summary RR = 1.94 (1.66–2.27)
Larsson, 2011	Meta-analysis [12]	9 cohort studies	Men RR = 1.26 (1.06–1.49) Women RR = 1.70 (1.47–1.97)
Bansal, 2013	Meta-analysis [13]	16 case-control studies 29 cohort studies	RR = 0.86 (0.80–0.92)
Peairs, 2011	Systematic review and meta-analysis [14]	8 studies	HR = 1.49 (1.35–1.65)
Noto, 2010	Systematic review and meta-analysis [15]	1 case-control study 4 cohort studies	Men RR = 1.25 (1.06–1.46) Women RR = 1.23 (0.97–1.56)

cancer by diabetes subtype. The overall RR for pancreatic cancer in type 1 or young-onset diabetes was significantly higher compared to those with no diabetes (RR 2.00, 95% CI: 1.37–3.01) [10]. Furthermore, in a meta-analysis of 35 cohort studies, diabetes was associated with an increased risk of pancreatic cancer (SRR 1.94, 95% CI: 1.66–2.27). However, there was a significant heterogeneity among the studies ($P < 0.001$, $I^2 = 93.6\%$) [11]. In this meta-analysis, the risk of pancreatic cancer was independent of body mass index, alcohol consumption, and smoking status.

The risk for kidney cancer was investigated in a meta-analysis of nine cohort studies. The results showed a 42% increased risk of kidney cancer, although the studies were heterogenous ($P < 0.001$). The risk was higher in women (RR 1.70, 95% CI: 1.47–1.9) than in men (RR 1.26, 95% CI: 1.06–1.49). Adjustment for BMI lowered the association significantly (RR 1.12, 95% CI: 0.99–1.27) [12].

The available evidence for the risk of prostate cancer in type 2 diabetes is conflicting. A recent meta-analysis, of 29 cohorts and 16 case-control studies, examined the association between type 2 diabetes and risk of prostate cancer and showed a significant inverse association (RR 0.86, 95% CI: 0.80–0.92) [13]. Table 1 summarizes the cancer risks in diabetes.

Considering cancer mortality, preexisting diabetes is associated with the risk of all-cause and site-specific mortality [2, 4, 8, 14].

Noto et al. described in a qualitative review and meta-analysis that diabetes was associated with an increased relative risk of all-cancer mortality (RR 1.16, 95% CI: 1.03–1.30) which remained significant for both sexes in a subgroup analysis [2].

Site-specific cancer mortality for colorectal cancer was evaluated in a systematic review and a meta-analysis of cohort studies by Jiang et al. They reported that diabetes was associated with a 20% increase in the incidence of colorectal cancer that was independent of sex, body mass index, family history of colorectal cancer, smoking, geographic location, and physical activity [4].

Similarly, mortality from gastric cancer in a patient with diabetes was reported to be higher than those individuals without diabetes (SRR 1.29, 95% CI: 1.04–1.59) [8].

Breast cancer outcomes in people with preexisting diabetes were reported in a systematic review and a meta-analysis by Peairs et al. In this study preexisting diabetes was associated with 49% increase in all-cause mortality in patients with breast cancer. In addition, these people had a more advanced stage on presentation and experienced more toxicity from chemotherapy [14].

TABLE 2: The potential explanations for the associations between the risk of all-cause and site-specific mortality in diabetes.

(1) Increase in cell proliferation rate in the presence of hyperglycemia and hyperinsulinemia
(2) Increase in cell permeability and changes in basement membranes caused by an increased level of reactive oxygen species (ROS)
(3) Presence of diabetes related morbidities which may alter clinical decisions regarding the treatment for the cancer
(4) Poorer response to the cancer treatment
(5) Suboptimal cancer screening programs in diabetic patients
(6) Excess mortality risk related to diabetes

Table 2 summarizes the potential explanations for the observed association of diabetes and mortality risk.

2. Hyperinsulinemia and Cancer

One of the most important metabolic abnormalities that might explain the relationship between type 2 diabetes and cancer is hyperinsulinemia [16].

Hyperinsulinemia is the result of insulin resistance in peripheral tissues.

Nocturnal lipolysis plays an important role in causing insulin resistance and compensatory hyperinsulinemia [17]. Insulin can have direct and indirect effect on tumor cells. Direct mitogenic effect of insulin on tumor cells is mediated through the insulin receptors that are expressed on tumor cells.

On the other hand, insulin, in high concentrations, is capable of activating the related growth factors receptors such as insulin-like growth factor one receptors. These receptors are also expressed on all tumor cells.

Furthermore, insulin can stimulate the synthesis of insulin growth factor one (IGF-1) and promote steroidogenesis.

In addition, insulin has an inhibitory effect on the expression of sex hormone binding globulins.

So, insulin can also promote tumor cell growth indirectly through elevated levels of circulating IGF-1 and sex steroids [18].

In 2004, Renehan et al. presented the idea of the direct link of insulin-like growth factors (IGFs) and insulin-like growth factor binding proteins (IGFBPs) and cancer risk [19].

Some studies described the association of circulating C-peptide, as the marker of insulin level, and colorectal cancer [19–34]. Similar results have been reported for breast cancer, and it has been shown that the overall risk of breast cancer is higher in the upper levels of C-peptide.

In summary, the proposed mechanisms explaining the possible positive link between hyperinsulinemia and cancer risk are

- (1) direct promotion of tumor proliferation [35],
- (2) modulation of circulating growth factors and their binding proteins [36],
- (3) binding to growth factor specific receptors [37].

Hyperinsulinemia has been shown to be associated with an increased risk of cancer mortality. Cremona study described a higher risk of cancer mortality (HR = 1.62, 95% CI: 1.19–2.20; $P < 0.002$) in general and a higher risk of mortality for gastrointestinal cancer (HR = 2.61, 95% CI: 1.73–3.94; $P < 0.0001$) [38]. The effect of hyperinsulinemia on cancer mortality was independent of diabetes, obesity, and the metabolic syndrome. In another study on endometrial cancer patients, hyperinsulinemia was associated with a more aggressive course in patients with well or moderately differentiated endometrial adenocarcinoma [39].

3. Metformin and Cancer

3.1. Metformin-Historical Perspective. *Galega officinalis* (French lilac or goat's rue) is the plant which biguanides are derived from. In ancient Egypt and medieval Europe, it was used for the relief of polyuria [40–42].

In the 1920s, biguanides were identified as the active ingredient of *Galega officinalis* and in the 1950s, they were developed as therapeutics. Since then, metformin, phenformin, and buformin were developed and used for treatment of type 2 diabetes [43]. The problem with the latter two, that is, phenformin and buformin, was toxicity related to lactic acidosis. Hence, they were withdrawn from the market by the 1970s.

However, metformin proved to be safe and was recognized as one of the most effective and safe therapeutics for treatment of type 2 diabetes. In the UK, it got the approval for the treatment of hyperglycemia in 1958, and three decades later it was approved in the US.

Several years later, “a new life for an old drug” was begun [44]. The role of biguanides in metabolic immunotherapy and metabolic rehabilitation opened a new window towards the future: the use of metformin beyond diabetes [44–47].

3.2. Mechanism of Action of Metformin. Adenosine monophosphate kinase (AMPK) is the cellular energy sensor located within the cytoplasm. It is involved in regulating metabolism within the cells (Figure 1).

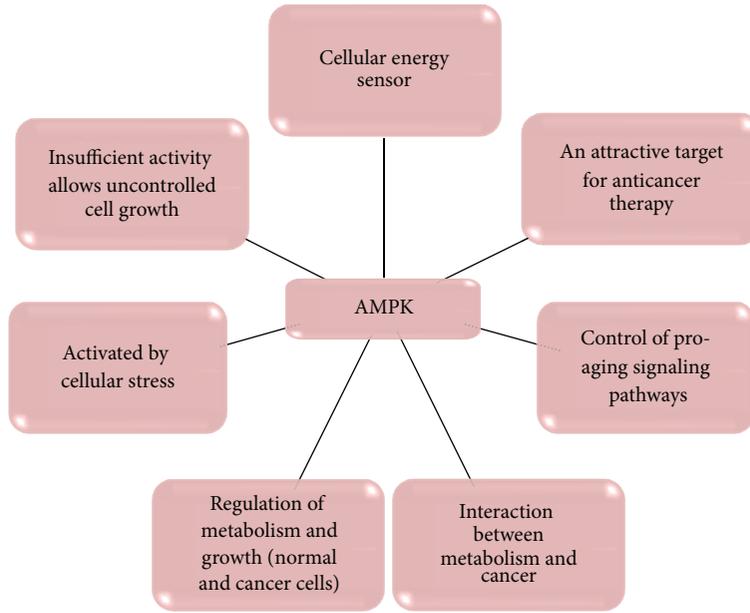
A protein called liver kinase B₁ (LKB₁) is the product of the tumor suppressor gene. Phosphorylation of AMPK catalytic subunit takes place in the presence of LKB1 and is facilitated by AMP. Increasing intracellular levels of adenosine monophosphate (AMP) activates AMPK [48, 49].

AMPK activation leads to the inhibition of mammalian target of rapamycin (mTOR) signaling, hence, downregulates gluconeogenesis by the liver.

Figure 2 illustrates that the energy sensing pathways converge on the coactivator TORC2.

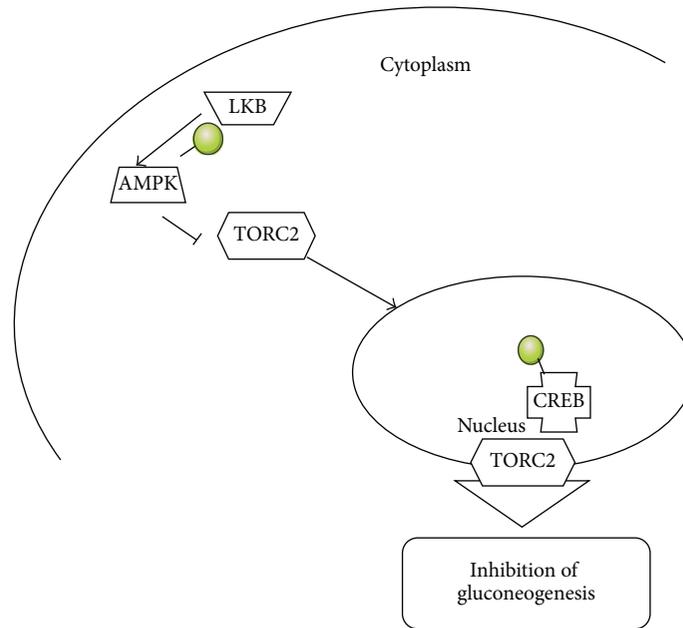
The role of AMPK in regulating energy balance is illustrated in Table 3.

Metformin exerts both indirect (insulin dependent) and direct (insulin independent) actions at the cellular level. Its direct effect is mediated via AMPK activation and reduction of mTOR signaling pathway which leads to inhibition of gluconeogenesis in the liver, protein synthesis and cell proliferation in cancer cells [50–53].



AMPK: adenosine monophosphate kinase

FIGURE 1: The role of AMPK* at cellular level.



Phosphor

AMPK: adenosine monophosphate kinase

AMP: adenosine monophosphate

LKB₁: liver kinase B₁

mTOR: mammalian target of rapamycin

TORC2: transducer of regulated CREB activity 2

CREB: cyclic AMP response element binding

FIGURE 2: Energy sensing pathways converge on the coactivator TORC2.

TABLE 3: Role of AMPK in regulating energy balance at the whole body.

Pancreas	Modulation of insulin secretion
	Inhibition of gluconeogenesis
Liver	Inhibition of cholesterol synthesis
Adipose tissue	Inhibition of lipolysis
Skeletal muscle	Increase glucose uptake
	Increase fatty acid oxidation
Hypothalamus	Increase food intake
Heart	Increase glucose uptake
	Increase fatty acid oxidation

AMPK: adenosine monophosphate kinase.

The indirect effects of metformin are mediated through its blood glucose lowering ability and subsequent reduction of the circulating insulin level.

Metformin activates AMPK in the liver and skeletal muscles. In this way, it reduces gluconeogenesis in the liver and fosters glucose uptake by the peripheral tissues, leading to a lower blood glucose and insulin levels [40, 54].

Insulin has been known to have a mitogenic activity and theoretically could have a promoting effect on the growth of tumor cells [55, 56].

Although various anticancer effects for metformin have been described [57, 58], the LKB₁ mediated mTOR signaling suppression seems to be the fundamental mechanism of anticancer effect of metformin. Table 4 shows the possible anticancerogenic effects of metformin.

3.3. Metformin, Cancer Risk, and Cancer Mortality. Metformin is the drug of choice for the treatment of type 2 diabetes [59].

It is a safe drug, and its cost is very low. So, it remains one of the most common prescribed drugs worldwide [60]. In addition, metformin can inhibit cancer cell growth in vitro [60–62].

Hirsch et al. [63] explained that metformin selectively kills cancer stem cells and blocks tumor growth. They also observed a synergistic action of metformin with chemotherapeutic drugs in order to reduce tumor mass and prolongation of remission in nude mice.

They showed that a combination treatment (metformin plus doxorubicin) can induce remission and maintained it for at least 60 days after treatment withdrawal. Relapse of the tumor growth was observed after 20 days in mice treated with doxorubicin alone. On the other hand, combination therapy was associated with a prolong remission that might represents a cure.

Meta-analyses of metformin and cancer risk in diabetic patients described around one-third reduction in overall cancer risk and cancer mortality in metformin users compared with other antidiabetic drugs [14, 64].

Moreover, the pooled risk ratios for the incidence of specific cancer sites were lower in metformin users: 0.68 (95% CI, 0.53–0.88) for colorectal cancer, 0.2 (95% CI, 0.07–0.59) for

TABLE 4: The putative anticarcinogenic effects of metformin.

- (1) Activation of LKB₁/AMPK pathway
- (2) Induction of cell cycle arrest and/or apoptosis
- (3) Inhibition of protein synthesis
- (4) Reduction in circulating insulin levels
- (5) Activation of the immune system
- (6) Eradication of cancer stem cells
- (7) Reduced IGF1, insulin, and HER2-mediated signaling
- (8) Inhibition of angiogenesis

AMPK: adenosine monophosphate kinase.

LKB₁: liver kinase B₁.

IGF1: insulin-like growth factor 1.

HER2: human epidermal growth factor receptor 2.

hepatocellular cancer, and 0.67 (95% CI, 0.45–0.99) for lung cancer [14].

The potential role of metformin in chemoprevention for liver cancer is clearly described in a recent meta-analysis [65]. The study indicated that among patients with type 2 diabetes, metformin is associated with a 62% reduction in the estimated risk of liver cancer and 70% risk reduction for hepatocellular carcinoma (HCC).

The finding that the risk reduction for cancer incidence varies among certain sites might be due to the difference in carcinogenesis at different sites. Growth of the epithelial malignant tumors such as colon, pancreas, and breast is affected by hyperglycemia and hyperinsulinemia [66–70]. These findings support the anticancer effect of metformin especially because the analysis was based on large population-based data obtained from multiple nations, including Asians who are generally lean and insulinopenic [15].

Although meta-analyses of observational studies suggest a cancer risk reduction around one third in metformin users, results of a recently published systematic review and collaborative meta-analyses of randomized clinical trials do not support this concept [71]. This meta-analysis obtained the data for cancer incidence from 11 RCTs and for all-cause mortality from 13 RCTs. The summary RR for the incident of cancer in people randomized to metformin compared with comparators was 1.03 (95% CI: 0.82–1.28, $I^2 = 15%$).

In addition, the summary RR for all-cause mortality in metformin users compared with comparators was 0.94 (95% CI: 0.79–1.12).

The analyses of the trials comparing metformin to placebo or usual care and trials with periods longer than one year were not in favor of metformin. However, the confidence intervals were wide, and there was a high clinical heterogeneity between the trials considering the comparators. Furthermore, there was not sufficient data to examine individual cancer endpoints. Another important limitation was the short follow-up period (average 4.1 years). In most observational studies, the beneficial effect of metformin to reduce the cancer risk was evident when the drug was used for more than five years [64].

As most patients with diabetes need multiple drugs for optimal metabolic control [72], the possibility of drug interactions is an important limitation for the evaluation of cancer risk of specific glucose lowering drugs in both observational studies and clinical trials.

3.4. Glucose Lowering Drugs, Cancer Risk, and Cancer Mortality. Although diabetes appears to be a risk factor for some types of cancer and is associated with the risk of mortality in cancer patients, the impact of intensive glycemic control on the cancer risk reduction and improving cancer outcomes is controversial.

The majority of the evidence for the association between glycemic control and cancer outcomes is based on the epidemiologic studies which harbor considerable confounds [73–76].

Several factors such as the duration of diabetes, the need for combination drug therapy to reach metabolic targets, and the presence of diabetes comorbidities make the assessment of the effect of diabetes treatment on cancer risk and cancer mortality difficult.

One clinical retrospective study examined the pathologic complete response rate (PCR) in women with early stage breast cancer and compared the PCR rate in diabetic women receiving metformin with non-metformin users and nondiabetic controls. The PCR was 24% in the metformin group, 8.0% in the nonmetformin group, and 16% in nondiabetic group [77, 78]. Other studies found similar effects for metformin among patients with nonsmall lung cancer and colorectal cancer with diabetes [79, 80]. However, there was no significant improvement in estimated 3-year relapse-free survival rate.

In a case control study [81], metformin was found to be associated with a risk reduction for cancer in general. UKPDS metformin study compared metformin-based intensified treatment against dietary management in overweight people with type 2 diabetes. It supports the beneficial effect of metformin on cancer mortality [82]. However, the sample size was small in this study.

An epidemiologic study found that metformin use was associated with a lower risk of cancer-related mortality compared to sulfonylureas users [61]. A meta-analysis of seventeen observational studies investigated the risk of all cancer and site-specific cancer in people with type 2 diabetes and compared metformin with sulfonylurea [83]. The meta-analysis showed that metformin was significantly associated with decreased RR for all cancer (SRR 0.61, 95% CI: 0.54–0.7), although the studies included in the final analysis were significantly heterogeneous. Furthermore, except for colorectal cancer, metformin was not associated with any significant effect on the incidence of other cancers, for example, prostate and breast cancers.

In another study, Currie described the cancer risk according to treatment of type 2 diabetes. It was nadir for metformin monotherapy compared to sulfonylurea monotherapy, sulfonylurea plus metformin, and insulin-based therapy. The risk for insulin secretagogues was similar to that of insulin

[84]. However, it should be mentioned that patients on combination or insulin therapies might be expected to have diabetes for a longer period of time.

In a recent large population-based study [85], a lower risk of cancer in general and specific cancers was described in patients treated with metformin in comparison with those received sulfonylurea derivatives. Both groups had a similar duration of diabetes; however, the cause of death was not identified. So, they could not be able to compare the association of the risk of cancer death between metformin users and those who had been treated with sulfonylurea derivatives.

The evidence from randomized trials differs from those based on observational studies. Home et al. [86] collected data for malignancies as an adverse event from a Diabetes Outcome Progression Trial (ADOPT) and Rosiglitazone Evaluated for Cardiovascular Outcomes and Regulation of Glycemia in Diabetes (RECORD).

Incident cancer was considered as an adverse event in these two randomized controlled clinical trials. The results suggest that metformin has no advantage over rosiglitazone for cancer protection, although there is a possibility that sulfonylureas may have a small disadvantage over the other two drugs. However, it should be emphasized that the number of malignancies was limited in these trials, and the use of insulin for long-term glucose control might dilute the differences between the medications.

In conclusion, metformin is a safe and effective drug for treatment of type 2 diabetes with a low cost. The current evidence is not sufficient to support the anticancer effect of metformin. The protective effect of metformin might be evident over a longer period of time or in certain treatment groups, for example, insulin users.

Long-term randomized clinical trials specifically designed to determine metformin effect on cancer risk are needed to examine the hypothesis that metformin has an anticancer effect.

Key Messages

- (i) Type 2 diabetes is associated with an increased risk of incident cancer.
- (ii) Type 1 diabetes is associated with pancreatic cancer.
- (iii) Preexisting diabetes is associated with the risk of all-cause and cancer-specific mortality.
- (iv) Available evidence from observational studies supports the hypothesis that metformin has an anticancer effect.
- (v) Randomized controlled clinical trials do not provide strong evidence supporting the protective effect of metformin on cancer incidence and mortality.
- (vi) Long-term RCTs are needed to specifically target the effect of metformin on cancer risk and cancer mortality.

Abbreviations

UK: United Kingdom
US: United States

AMPK: Adenosine monophosphate kinase
 LKB₁: Liver kinase B₁
 AMP: Adenosine monophosphate
 mTOR: Mammalian target of rapamycin
 TORC2: Transducer of Regulated CREB activity 2
 CREB: Cyclic AMP response element binding
 IGF1: Insulin-like growth factor 1
 HER2: Human epidermal growth factor receptor 2
 IGFBPs: Insulin-like growth factor binding proteins
 ROS: Reactive oxygen species
 PCR: Pathologic complete response rate
 UKPDS: United Kingdom Prospective Diabetes Study
 EASD: European Association for the Study of Diabetes
 ADA: American Diabetes Association
 SRRs: Standardised registration ratios
 HCC: Hepatocellular carcinoma
 RRs: Relative risks.

Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

References

- [1] C. Ko and S. Chaudhry, "The need for a multidisciplinary approach to cancer care," *Journal of Surgical Research*, vol. 105, no. 1, pp. 53–57, 2002.
- [2] H. Noto, T. Tsujimoto, T. Sasazuki, and M. Noda, "Significantly increased risk of cancer in patients with diabetes mellitus: a systematic review and meta-analysis," *Endocrine Practice*, vol. 17, no. 4, pp. 616–628, 2011.
- [3] L. Deng, Z. Gui, L. Zhao, J. Wang, and L. Shen, "Diabetes mellitus and the incidence of colorectal cancer: an updated systematic review and meta-analysis," *Digestive Diseases and Sciences*, vol. 57, no. 6, pp. 1576–1585, 2012.
- [4] Y. Jiang, Q. Ben, H. Shen, W. Lu, Y. Zhang, and J. Zhu, "Diabetes mellitus and incidence and mortality of colorectal cancer: a systematic review and meta-analysis of cohort studies," *European Journal of Epidemiology*, vol. 26, no. 11, pp. 863–876, 2011.
- [5] H. U. Krämer, B. Schöttker, E. Raum, and H. Brenner, "Type 2 diabetes mellitus and colorectal cancer: meta-analysis on sex-specific differences," *European Journal of Cancer*, vol. 48, pp. 1269–1282, 2012.
- [6] P. Boyle, M. Boniol, A. Koechlin et al., "Diabetes and breast cancer risk: a meta-analysis," *British Journal of Cancer*, vol. 107, pp. 1608–1617, 2012.
- [7] P. J. Hardefeldt, G. D. Eslick, and S. Edirimanne, "Benign thyroid disease is associated with breast cancer: a meta-analysis," *Breast Cancer Research and Treatment*, vol. 133, no. 3, pp. 1169–1177, 2012.
- [8] T. Tian, L. Q. Zhang, X. H. Ma, J. N. Zhou, and J. Shen, "Diabetes mellitus and incidence and mortality of gastric cancer: a meta-analysis," *Experimental and Clinical Endocrinology and Diabetes*, vol. 120, no. 4, pp. 217–223, 2012.
- [9] Z. Ge, Q. Ben, J. Qian, Y. Wang, and Y. Li, "Diabetes mellitus and risk of gastric cancer: a systematic review and meta-analysis of observational studies," *European Journal of Gastroenterology and Hepatology*, vol. 23, no. 12, pp. 1127–1135, 2011.
- [10] R. J. Stevens, A. W. Roddam, and V. Beral, "Pancreatic cancer in type 1 and young-onset diabetes: systematic review and meta-analysis," *British Journal of Cancer*, vol. 96, no. 3, pp. 507–509, 2007.
- [11] Q. Ben, M. Xu, X. Ning et al., "Diabetes mellitus and risk of pancreatic cancer: a meta-analysis of cohort studies," *European Journal of Cancer*, vol. 47, no. 13, pp. 1928–1937, 2011.
- [12] S. C. Larsson and A. Wolk, "Diabetes mellitus and incidence of kidney cancer: a meta-analysis of cohort studies," *Diabetologia*, vol. 54, no. 5, pp. 1013–1018, 2011.
- [13] D. Bansal, A. Bhansali, G. Kapil et al., "Type 2 diabetes and risk of prostate cancer: a meta-analysis of observational studies," *Prostate Cancer and Prostatic Diseases*, vol. 16, no. 2, pp. 151–158, 2013.
- [14] K. S. Peairs, B. B. Barone, C. F. Snyder et al., "Diabetes mellitus and breast cancer outcomes: a systematic review and meta-analysis," *Journal of Clinical Oncology*, vol. 29, no. 1, pp. 40–46, 2011.
- [15] H. Noto, K. Osame, T. Sasazuki, and M. Noda, "Substantially increased risk of cancer in patients with diabetes mellitus: a systematic review and meta-analysis of epidemiologic evidence in Japan," *Journal of Diabetes and its Complications*, vol. 24, no. 5, pp. 345–353, 2010.
- [16] D. Lann and D. LeRoith, "The role of endocrine insulin-like growth factor-I and insulin in breast cancer," *Journal of Mammary Gland Biology and Neoplasia*, vol. 13, no. 4, pp. 371–379, 2008.
- [17] I. R. Hsu, S. P. Kim, M. Kabir, and R. N. Bergman, "Metabolic syndrome, hyperinsulinemia, and cancer," *The American Journal of Clinical Nutrition*, vol. 86, no. 3, pp. S867–S871, 2007.
- [18] R. Novosyadlyy and D. LeRoith, "Hyperinsulinemia and type 2 diabetes: impact on cancer," *Cell Cycle*, vol. 9, no. 8, pp. 1449–1450, 2010.
- [19] A. G. Renehan, M. Zwahlen, C. Minder, S. T. O'Dwyer, S. M. Shalet, and M. Egger, "Insulin-like growth factor (IGF)-I, IGF binding protein-3, and cancer risk: systematic review and meta-regression analysis," *The Lancet*, vol. 363, no. 9418, pp. 1346–1353, 2004.
- [20] E. Giovannucci, "Insulin, insulin-like growth factors and colon cancer: a review of the evidence," *Journal of Nutrition*, vol. 131, no. 11, pp. 3109S–3120S, 2001.
- [21] I. Tripkovic, A. Tripkovic, M. Strnad, V. Capkun, and L. Zekan, "Role of insulin-like growth factor-1 in colon cancerogenesis: a Case-Control Study," *Archives of Medical Research*, vol. 38, no. 5, pp. 519–525, 2007.
- [22] R. E. Schoen, C. M. Tangen, L. H. Kuller et al., "Increased blood glucose and insulin, body size, and incident colorectal cancer," *Journal of the National Cancer Institute*, vol. 91, no. 13, pp. 1147–1154, 1999.
- [23] R. Kaaks, P. Toniolo, A. Akhmedkhanov et al., "Serum C-peptide, insulin-like growth factor (IGF)-I, IGF-binding proteins, and colorectal cancer risk in women," *Journal of the National Cancer Institute*, vol. 92, no. 19, pp. 1592–1600, 2000.
- [24] S. H. Saydah, E. A. Platz, N. Rifai, M. N. Pollak, F. L. Brancati, and K. J. Helzlsouer, "Association of markers of insulin and glucose control with subsequent colorectal cancer risk," *Cancer Epidemiology Biomarkers and Prevention*, vol. 12, no. 5, pp. 412–418, 2003.

- [25] P. Stattin, A. Lukanova, C. Biessy et al., "Obesity and colon cancer: does leptin provide a link?" *International Journal of Cancer*, vol. 109, no. 1, pp. 149–152, 2004.
- [26] E. K. Wei, J. Ma, M. N. Pollak et al., "A prospective study of C-peptide, insulin-like growth factor-I, insulin-like growth factor binding protein-1, and the risk of colorectal cancer in women," *Cancer Epidemiology Biomarkers and Prevention*, vol. 14, no. 4, pp. 850–855, 2005.
- [27] P. Toniolo, P. F. Bruning, A. Akhmedkhanov et al., "Serum insulin-like growth factor-I and breast cancer," *International Journal of Cancer*, vol. 88, pp. 828–832, 2000.
- [28] P. Muti, T. Quattrin, B. J. B. Grant et al., "Fasting glucose is a risk factor for breast cancer: a prospective study," *Cancer Epidemiology Biomarkers and Prevention*, vol. 11, no. 11, pp. 1361–1368, 2002.
- [29] M. Verheus, P. H. M. Peeters, S. Rinaldi et al., "Serum C-peptide levels and breast cancer risk: results from the European Prospective Investigation into Cancer and Nutrition (EPIC)," *International Journal of Cancer*, vol. 119, no. 3, pp. 659–667, 2006.
- [30] A. H. Eliassen, S. S. Tworoger, C. S. Mantzoros, M. N. Pollak, and S. E. Hankinson, "Circulating insulin and C-peptide levels and risk of breast cancer among predominately premenopausal women," *Cancer Epidemiology Biomarkers and Prevention*, vol. 16, no. 1, pp. 161–164, 2007.
- [31] P. F. Bruning, J. M. G. Bonfrer, P. A. H. Van Noord, A. A. M. Hart, M. De Jong-Bakker, and W. J. Nooljen, "Insulin resistance and breast-cancer risk," *International Journal of Cancer*, vol. 52, no. 4, pp. 511–516, 1992.
- [32] M. E. Del Giudice, I. G. Fantus, S. Ezzat, G. McKeown-Eyssen, D. Page, and P. J. Goodwin, "Insulin and related factors in premenopausal breast cancer risk," *Breast Cancer Research and Treatment*, vol. 47, no. 2, pp. 111–120, 1998.
- [33] A. Malin, Q. Dai, H. Yu et al., "Evaluation of the synergistic effect of insulin resistance and insulin-like growth factors on the risk of breast carcinoma," *Cancer*, vol. 100, no. 4, pp. 694–700, 2004.
- [34] C. Schairer, D. Hill, S. R. Sturgeon et al., "Serum concentrations of IGF-I, IGFBP-3 and c-peptide and risk of hyperplasia and cancer of the breast in postmenopausal women," *International Journal of Cancer*, vol. 108, no. 5, pp. 773–779, 2004.
- [35] M. Bontenbal, A. M. Sieuwerts, J. G. M. Klijn et al., "Effect of hormonal manipulation and doxorubicin administration on cell cycle kinetics of human breast cancer cells," *British Journal of Cancer*, vol. 60, no. 5, pp. 688–692, 1989.
- [36] A. G. Renehan, J. Frystyk, and A. Flyvbjerg, "Obesity and cancer risk: the role of the insulin-IGF axis," *Trends in Endocrinology and Metabolism*, vol. 17, no. 8, pp. 328–336, 2006.
- [37] G. Pandini, F. Frasca, R. Mineo, L. Sciacca, R. Vigneri, and A. Belfiore, "Insulin/insulin-like growth factor I hybrid receptors have different biological characteristics depending on the insulin receptor isoform involved," *Journal of Biological Chemistry*, vol. 277, no. 42, pp. 39684–39695, 2002.
- [38] G. Perseghin, G. Calori, G. Lattuada et al., "Insulin resistance/hyperinsulinemia and cancer mortality: the Cremona study at the 15th year of follow-up," *Acta Diabetologica*, vol. 49, no. 6, pp. 421–428, 2012.
- [39] L. M. Berstein, J. O. Kvatchevskaya, T. E. Poroshina et al., "Insulin resistance, its consequences for the clinical course of the disease, and possibilities of correction in endometrial cancer," *Journal of Cancer Research and Clinical Oncology*, vol. 130, no. 11, pp. 687–693, 2004.
- [40] L. A. Witters, "The blooming of the French lilac," *Journal of Clinical Investigation*, vol. 108, no. 8, pp. 1105–1107, 2001.
- [41] D. R. Hadden, "Goat's rue—French lilac—Italian fitch—Spanish sainfoin: gallega officinalis and metformin: the Edinburgh connection," *The Journal of the Royal College of Physicians of Edinburgh*, vol. 35, no. 3, pp. 258–260, 2005.
- [42] C. J. Bailey and C. Day, "Metformin: its botanical background," *Practical Diabetes International*, vol. 21, no. 3, pp. 115–117, 2004.
- [43] C. J. Bailey and R. C. Turner, "Metformin," *The New England Journal of Medicine*, vol. 334, no. 9, pp. 574–579, 1996.
- [44] C. M. Rotella, M. Monami, and E. Mannucci, "Metformin beyond diabetes: new life for an old drug," *Current Diabetes Reviews*, vol. 2, no. 3, pp. 307–315, 2006.
- [45] V. M. Dilman, L. M. Berstein, and M. N. Ostroumova, "Metabolic immunodepression and metabolic immunotherapy. An attempt at improvement in immunologic response in breast cancer patients by correction of metabolic disturbances," *Oncology*, vol. 39, no. 1, pp. 13–19, 1982.
- [46] V. M. Dilman, L. M. Berstein, T. P. Yevtushenko et al., "Preliminary evidence on metabolic rehabilitation of cancer patients," *Archiv fur Geschwulstforschung*, vol. 58, no. 3, pp. 175–183, 1988.
- [47] L. M. Berstein, "Modern approach to metabolic rehabilitation of cancer patients: biguanides (phenformin and metformin) and beyond," *Future Oncology*, vol. 6, no. 8, pp. 1313–1323, 2010.
- [48] B. B. Kahn, T. Alquier, D. Carling, and D. G. Hardie, "AMP-activated protein kinase: ancient energy gauge provides clues to modern understanding of metabolism," *Cell Metabolism*, vol. 1, no. 1, pp. 15–25, 2005.
- [49] G. Zhou, R. Myers, Y. Li et al., "Role of AMP-activated protein kinase in mechanism of metformin action," *Journal of Clinical Investigation*, vol. 108, no. 8, pp. 1167–1174, 2001.
- [50] R. J. O. Dowling, M. Zakikhani, I. G. Fantus, M. Pollak, and N. Sonenberg, "Metformin inhibits mammalian target of rapamycin-dependent translation initiation in breast cancer cells," *Cancer Research*, vol. 67, no. 22, pp. 10804–10812, 2007.
- [51] M. Zakikhani, R. Dowling, I. G. Fantus, N. Sonenberg, and M. Pollak, "Metformin is an AMP kinase-dependent growth inhibitor for breast cancer cells," *Cancer Research*, vol. 66, no. 21, pp. 10269–10273, 2006.
- [52] W. H. Gotlieb, J. Saumet, M.-C. Beauchamp et al., "In vitro metformin anti-neoplastic activity in epithelial ovarian cancer," *Gynecologic Oncology*, vol. 110, no. 2, pp. 246–250, 2008.
- [53] I. N. Alimova, B. Liu, Z. Fan et al., "Metformin inhibits breast cancer cell growth, colony formation and induces cell cycle arrest in vitro," *Cell Cycle*, vol. 8, no. 6, pp. 909–915, 2009.
- [54] K. Cusi, A. Consoli, and R. A. Defronzo, "Metabolic effects of metformin on glucose and lactate metabolism in noninsulin-dependent diabetes mellitus," *Journal of Clinical Endocrinology and Metabolism*, vol. 81, no. 11, pp. 4059–4067, 1996.
- [55] A. Belfiore and F. Frasca, "IGF and insulin receptor signaling in breast cancer," *Journal of Mammary Gland Biology and Neoplasia*, vol. 13, no. 4, pp. 381–406, 2008.
- [56] A. M. Mulligan, F. P. O'Malley, M. Ennis, I. G. Fantus, and P. J. Goodwin, "Insulin receptor is an independent predictor of a favorable outcome in early stage breast cancer," *Breast Cancer Research and Treatment*, vol. 106, no. 1, pp. 39–47, 2007.
- [57] A. Kalender, A. Selvaraj, S. Y. Kim et al., "Metformin, independent of AMPK, inhibits mTORC1 in a rag GTPase-dependent manner," *Cell Metabolism*, vol. 11, no. 5, pp. 390–401, 2010.
- [58] M. Foretz, S. Hébrard, J. Leclerc et al., "Metformin inhibits hepatic gluconeogenesis in mice independently of the LKB1/AMPK pathway via a decrease in hepatic energy state," *Journal of Clinical Investigation*, vol. 120, no. 7, pp. 2355–2369, 2010.

- [59] S. E. Inzucchi, R. M. Bergenstal, J. B. Buse et al., "Management of hyperglycaemia in type 2 diabetes: a patient-centered approach. Position statement of the American Diabetes Association (ADA) and the European Association for the Study of Diabetes (EASD)," *Diabetologia*, vol. 55, no. 6, pp. 1577–1596, 2012.
- [60] M. Monami, C. Lamanna, D. Balzi, N. Marchionni, and E. Mannucci, "Sulphonylureas and cancer: a case-control study," *Acta Diabetologica*, vol. 46, no. 4, pp. 279–284, 2009.
- [61] S. L. Bowker, S. R. Majumdar, P. Veugelers, and J. A. Johnson, "Increased cancer-related mortality for patients with type 2 diabetes who use sulfonylureas or insulin," *Diabetes Care*, vol. 29, no. 2, pp. 254–258, 2006.
- [62] T. Patel, G. Hruby, K. Badani, C. Abate-Shen, and J. M. McKiernan, "Clinical outcomes after radical prostatectomy in diabetic patients treated with metformin," *Urology*, vol. 76, no. 5, pp. 1240–1244, 2010.
- [63] H. A. Hirsch, D. Iliopoulos, P. N. Tsiachlis, and K. Struhl, "Metformin selectively targets cancer stem cells, and acts together with chemotherapy to block tumor growth and prolong remission," *Cancer Research*, vol. 69, no. 22, article 8832, 2009.
- [64] A. DeCensi, M. Puntoni, P. Goodwin et al., "Metformin and cancer risk in diabetic patients: a systematic review and meta-analysis," *Cancer Prevention Research*, vol. 3, no. 11, pp. 1451–1461, 2010.
- [65] Z. J. Zhang, Z. J. Zheng, R. Shi et al., "Metformin for liver cancer prevention in patients with type 2 diabetes: a systematic review and meta-analysis," *Journal of Clinical Endocrinology and Metabolism*, vol. 97, pp. 2347–2353, 2012.
- [66] E. Giovannucci, "Metabolic syndrome, hyperinsulinemia, and colon cancer: a review," *The American Journal of Clinical Nutrition*, vol. 86, no. 3, pp. 836S–842S, 2007.
- [67] C. Algire, L. Amrein, M. Zakikhani, L. Panasci, and M. Pollak, "Metformin blocks the stimulative effect of a high-energy diet on colon carcinoma growth in vivo and is associated with reduced expression of fatty acid synthase," *Endocrine-Related Cancer*, vol. 17, no. 2, pp. 351–360, 2010.
- [68] S. Raimondi, P. Maisonneuve, and A. B. Lowenfels, "Epidemiology of pancreatic cancer: an overview," *Nature Reviews Gastroenterology and Hepatology*, vol. 6, no. 12, pp. 699–708, 2009.
- [69] M. B. Schneider, H. Matsuzaki, J. Haorah et al., "Prevention of pancreatic cancer induction in hamsters by metformin," *Gastroenterology*, vol. 120, no. 5, pp. 1263–1270, 2001.
- [70] F. Xue and K. B. Michels, "Diabetes, metabolic syndrome, and breast cancer: a review of the current evidence," *The American Journal of Clinical Nutrition*, vol. 86, no. 3, pp. S823–S835, 2007.
- [71] R. J. Stevens, R. Ali, C. R. Bankhead et al., "Cancer outcomes and all-cause mortality in adults allocated to metformin: systematic review and collaborative meta-analysis of randomised clinical trials," *Diabetologia*, vol. 55, pp. 2593–2603, 2012.
- [72] R. C. Turner, C. A. Cull, V. Frighi, and R. R. Holman, "Glycemic control with diet, sulfonylurea, metformin, or insulin in patients with type 2 diabetes mellitus. Progressive requirement for multiple therapies (UKPDS 49)," *Journal of the American Medical Association*, vol. 281, no. 21, pp. 2005–2012, 1999.
- [73] U. Smith and E. A. M. Gale, "Editorial: does diabetes therapy influence the risk of cancer?" *Diabetologia*, vol. 52, no. 9, pp. 1699–1708, 2009.
- [74] S. K. Garg, I. B. Hirsch, and J. S. Skyler, "Editorial: insulin glargine and cancer—an unsubstantiated allegation," *Diabetes Technology and Therapeutics*, vol. 11, no. 8, pp. 473–476, 2009.
- [75] B. B. Duncan and M. I. Schmidt, "Metformin, cancer, alphabet soup, and the role of epidemiology in etiologic research," *Diabetes Care*, vol. 32, no. 9, pp. 1748–1750, 2009.
- [76] S. J. Pocock and L. Smeeth, "Insulin glargine and malignancy: an unwarranted alarm," *The Lancet*, vol. 374, no. 9689, pp. 511–513, 2009.
- [77] A. D. Hanchate, K. M. Clough-Gorr, A. S. Ash, S. S. Thwin, and R. A. Silliman, "Longitudinal patterns in survival, comorbidity, healthcare utilization and quality of care among older women following breast cancer diagnosis," *Journal of General Internal Medicine*, vol. 25, no. 10, pp. 1045–1050, 2010.
- [78] S. Jiralerspong, S. L. Palla, S. H. Giordano et al., "Metformin and pathologic complete responses to neoadjuvant chemotherapy in diabetic patients with breast cancer," *Journal of Clinical Oncology*, vol. 27, no. 20, pp. 3297–3302, 2009.
- [79] B.-X. Tan, W.-X. Yao, J. Ge et al., "Prognostic influence of metformin as first-line chemotherapy for advanced nonsmall cell lung cancer in patients with type 2 diabetes," *Cancer*, vol. 117, no. 22, pp. 5103–5111, 2011.
- [80] J. H. Lee, T. I. Kim, S. M. Jeon, S. P. Hong, J. H. Cheon, and W. H. Kim, "The effects of metformin on the survival of colorectal cancer patients with diabetes mellitus," *International Journal of Cancer*, vol. 7, no. 6, pp. 752–759, 2011.
- [81] J. M. M. Evans, L. A. Donnelly, A. M. Emslie-Smith, D. R. Alessi, and A. D. Morris, "Metformin and reduced risk of cancer in diabetic patients," *British Medical Journal*, vol. 330, no. 7503, pp. 1304–1305, 2005.
- [82] J. A. Johnson and S. L. Bowker, "Intensive glycaemic control and cancer risk in type 2 diabetes: a meta-analysis of major trials," *Diabetologia*, vol. 54, no. 1, pp. 25–31, 2011.
- [83] D. Soranna, L. Scotti, A. Zamboni et al., "Cancer risk associated with use of metformin and sulfonylurea in type 2 diabetes: a meta-analysis," *Oncologist*, vol. 17, pp. 813–822, 2012.
- [84] C. J. Currie, C. D. Poole, and E. A. M. Gale, "The influence of glucose-lowering therapies on cancer risk in type 2 diabetes," *Diabetologia*, vol. 52, no. 9, pp. 1766–1777, 2009.
- [85] R. Ruiters, L. E. Visser, M. P. P. Van Herk-Sukel et al., "Lower risk of cancer in patients on metformin in comparison with those on sulfonylurea derivatives: results from a large population-based follow-up study," *Diabetes Care*, vol. 35, no. 1, pp. 119–124, 2012.
- [86] P. D. Home, S. E. Kahn, N. P. Jones, D. Noronha, H. Beck-Nielsen, and G. Viberti, "Experience of malignancies with oral glucose-lowering drugs in the randomised controlled ADOPT (A Diabetes Outcome Progression Trial) and RECORD (Rosiglitazone Evaluated for Cardiovascular Outcomes and Regulation of Glycaemia in Diabetes) clinical trials," *Diabetologia*, vol. 53, no. 9, pp. 1838–1845, 2010.



Hindawi

Submit your manuscripts at
<http://www.hindawi.com>

