Metformin in obesity, cancer and aging: addressing controversies

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Abstract: Metformin, an oral anti-diabetic drug, is being considered increasingly for treatment and prevention of cancer, obesity as well as for the extension of healthy lifespan. Gradually accumulating discrepancies about its effect on cancer and obesity can be explained by the shortage of randomized clinical trials, differences between control groups (reference points), gender- and age-associated effects and pharmacogenetic factors. Studies of the potential antiaging effects of antidiabetic biguanides, such as metformin, are still experimental for obvious reasons and their results are currently ambiguous. Here we discuss whether the discrepancies in different studies are merely methodological or inherently related to individual differences in responsiveness to the drug.

The epidemics of obesity and diabetes, which has been the matter of major concern for endocrinologists over the last two decades, has lead to an apparently unexpected, yet actually long foretold [1-3] result: the concern spread not only to cardiologists but, also, to oncologists and specialists engaged in tackling of more general problems, including aging and age-related pathology [4-7]. The wave of interest, with periodical decays and increasing surges, was associated with the attempts to use antidiabetic biguanides to control body weight and tumor growth [8-13]. Another facet of the situation is that almost 45 years ago these drugs were suggested to promote longevity [14]. Over the last years, the expanding bodies of relevant evidence, which mainly related to metformin, started to merge and occupy increasing place in current literature. The objective of the present essay is to attract more attention to accumulating inconsistencies. The first two sections of the essay, which are related to obesity and cancer, are based mostly on clinical data. The third section, which is related to aging or, rather, antiaging, is based predominately on experimental evidence obtained in rodents. Clearly, obesity and cancer have numerous interrelationships with aging (see details in [3, 15, 16]); however, we will separate these aspects for the sake of clarity in discussing the relevant effects of metformin.

Metformin and obesity

Weight-reducing effects of biguanides may explain in part their antitumor activity [13, 17]. Obesity is associated with increased mortality and, hence, decreased lifespan [2, 4]. Therefore, first of all, we will discuss whether biguanides reduce weight and affect body composition (fat vs. lean mass and visceral vs. subcutaneous fat). Two meta-analyses published in 2005 and 2011 summarized the use of metformin to treat adults having excessive body mass (fat) and showed that, among more than fifty potentially relevant studies, only less than ten satisfied all criteria. Only two or three of the latter confirmed that metformin had moderate weight-reducing effects, which, however, were inferior to the effects of behavioral interventions and gastrointestinal fat absorption inhibitor orlistat [18, 19]. Nevertheless, more scrutiny in treating the data accumulated so far, a part of which reproduces earlier results, suggests that it is reasonable to dissect the evidence into several subsections.

Obesity in type 2 diabetes mellitus

For obese diabetics and people at risk of diabetes, metformin remains a treatment able to moderately reduce body weight (by 5% on average). This is believed to be an additional benefit in treating diabetes
and is suggested to be caused by reducing of insulin resistance and hyperinsulinemia rather than by anorexic or other effects [20, 21].

Polycystic ovary syndrome (PCOS) in women with excess body weight
Only in a part of the studies in PCOS patients with obesity, metformin treatment without behavioral interventions decreased body mass index (BMI). It is not clear whether this can be explained by metformin underdosage [22], although a recent meta-analysis indicates that both drug dose and duration of treatment are highly relevant [23].

Age, genetic factors and oxidative stress
According to a recent randomised study [24], confirmed by other publications [25], children and adolescents may be more responsive to metformin-induced weight reduction compared to adults. This is reflected by body weight dynamics and by changes in the waist/hip ratio and fat deposits. Metformin has been shown to upregulate the expression of the organic cation transporter OCT1 in adipose tissue. Therefore, OCT1 may be a potential marker of metformin accumulation in the tissue [26]. On the other hand, bearing of one of the polymorphisms of the neuron growth regulator NEGR1 predicts the magnitude of the weight-reducing effect of metformin and, ironically, the propensity to restore body weight after the discontinuation of treatment with metformin [27]. The significance of oxidative stress also is difficult to characterize. This process is clearly increased in the adipose tissue in obese patients [28], and although metformin is known to have an antioxidant activity, especially in diabetes [29], this drug can enhance free radical generation in differentiating adipocytes [30].

Metformin and the heterogeneity of obesity
Possible associations between differences in metabolic phenotypes could account for differences in metformin action. The list of conditions is long and includes among others such varieties as ‘standard’ obesity, metabolically healthy obese state (MHO), sarcopenic obesity, and metabolically obese normal weight state (MONW) (see [31]). The incidences of major non-infectious age-dependent diseases, the ways of treating them, and treatment outcomes, especially at advanced ages, are important as well known from both, basic science and, especially, applied science standpoints. With regard to treatment, it should be noted that, although there are still many proponents of the idea that obesity should be treated irrespective of its type, it is also suggested that such a straightforward approach may be inadequate, in particular, when applied to MHO [32]. There is still no evidence about metformin use with regards to the some aforementioned subtypes of obesity, in contrast to numerous studies of its use in the visceral/abdominal obesity (e.g., see [33] and many others). The sarcopenic obesity, i.e. body fat gain associated with skeletal muscle loss, appears in cancer patients to lead to poorer outcomes of endocrine therapy and chemotherapy, treatments that tend to increase body fat by themselves.

Metformin and body weight in cancer patients
This issue is addressed implicitly in many studies of polycystic ovary syndrome (see above), which is believed to be a risk factor of several hormone-dependent malignancies, first of all, endometrial cancer. Several relevant observations relate to prostate cancer patients exposed to androgen deprivation, which is associated often with insulin resistance and other manifestations of the metabolic syndrome, including body weight gain. In one study of such patients, metformin treatment during 6 months was associated with decreased BMI and waist circumference [34]. The important question of the influence of body weight and body fat on the ability of metformin to affect cancer incidence in diabetic patients (see the next section) is unanswered as of yet, although BMI as potential risk modulator was taken in consideration in some of such studies, e.g. [35].

Cancer incidence in diabetic patients treated with metformin
As noted earlier [36] based on evidence accumulated by mid-2010ies, the emerging picture is rather vague. In particular, metformin effects are rather variable with regard to specific cancer locations, except for two notable cancers where the incidence decreased with metformin (colon cancer and hepatoblastoma). These data suggest that the possible anticancer effects of metformin are tissue-specific. It was reasoned by some authors that it is not that metformin is ‘thus good’, but the reference therapies, including sulfonylureas and insulin, may be ‘thus bad’ [35, 36]. Also, no data were available to judge whether glucose intolerance reduction/compensation by biguanides is important for the total and site-specific cancer incidence shifts.

Much of relevant evidence was gained over the last two years. The main findings may be categorized according to the following subtitles:

Cancer location. Reduced colon cancer incidence in diabetic patients treated with metformin was confirmed by meta-analysis of 5 observational studies where total odds ratio (OR) for treated vs. untreated subjects was 0.68 [37]; however, this conclusion was questioned later [38]. Some controversy remains in regards to prostate
cancer because in a case-control study of diabetics treated with metformin the OR for prostate cancer was found to increase to 1.23 [39]. In a study based on data obtained from Danish Cancer Register, the OR for breast cancer incidence was found to decrease to 0.77 upon the use of metformin in patients with type 2 diabetes [40]. Despite this positive result, the situation with breast cancer is somewhat debatable, in particular, because several known molecular-biological subtypes of this tumor are significantly different in their risk factors and responses to therapy. Accordingly, some studies found no association between metformin therapy and breast cancer incidence in diabetic women [41, 42]. In a recent study performed in Taiwan, metformin intake by diabetics was found to be associated with reduced risks of colorectal carcinomas (the effect was more expressed in women) and liver cancer (only in men) but not of oesophagus, stomach and pancreas cancer [43].

Study design. The aforementioned study from Taiwan [43] may be referred to as prospective, since study subjects, which were recruited in 2000, where followed for 7 to 8 years. Unfortunately, prospective studies related to the ‘metformin-diabetes-cancer risk problem’ are rare and are not designed specifically to tackle this problem. Available publications are largely based on observational or retrospective case-control studies or, rarely, cohort follow-up [35]. To the best of our knowledge, there is only one meta-analysis of randomized controlled trials (RCTs) that was reported at the Annual American Diabetes Association Conference in June 2011 [44] and cited in the editorial article [45]. The analysis was based on 4 published trials and 3 datasets obtained directly from Principal Investigators. Inclusion criteria for the trials stipulated that there must be ≥ 500 diabetes mellitus patients followed for ≥ 1 year. Output parameters were cancer incidence and all-cause mortality. The OR for cancer risks upon metformin intake vs. other therapies varied from 1.03 to 1.47 arguing against the ability of metformin to reduce cancer incidence in diabetic patients [44]. It cannot be excluded, however, that the above meta-analysis included the data of an independently published study [46] based on ADAPT and RECORD RCTs that did not show differences between metformin and roziglitazone with regard to their effects on cancer risk, but did demonstrate and confirm that metformin affords some benefit compared with sulfonylurea. To tell the truth, both these trials mentioned in the paper of P.D. Home et al. [46] were not designed specifically to address cancer incidence (as well as many previous observational studies that suggested a reduction of cancer risk by metformin therapy).

Reference points. The importance of ‘reference therapy’ or its absence (placebo) for judging the effects of metformin follows from the preceding paragraph and was mentioned above. Studies where data on metformin-treated diabetes were compared with untreated diabetes are few for obvious ethical and morbidity reasons. One such study is the aforementioned Taiwanese study [43] where total cancer incidence per 10 000 person×years were reported for “no diabetes” (46.0), “diabetes treated with metformin” (44.8), “diabetes treated with other antidiabetic therapies without metformin” (91.7), and “diabetes without any treatment” (97.6). The last two figures are very close suggesting that differences between metformin and other drugs may be associated with the adverse effects of the latter. Such proposals, in particular related to sulfonylurea derivatives, have been put forward several times before and are exemplified by the recent quotation: “However, whether this should indeed be seen as a decreased risk of cancer for the use of metformin compared with the use of sulfonylurea derivatives or as an increased risk of cancer for the use of sulfonylurea derivatives compared with the use of metformin remains to be elucidated” [47].

It should be noted that when cancer-related mortality, rather than cancer risk, is assessed in diabetics (once again, at ages mainly above 50), the belief that their survival is better in metformin-treated groups is not always confirmed for prostate and breast cancer [48, 49], although one study does suggest better survival for these cancers as well [50]. Metformin was found to be relatively more beneficial with regard to colon and ovarian cancer survival in diabetics [51, 52]. However, in these studies, metformin was compared with other antidiabetic drugs, whose less favorable effects cannot be ruled out.

Effectiveness of biguanides in cancer patients without diabetes

Past- vs. present-time evidence. The currently discussed prospects for using antidiabetic biguanides as potential antitumor drugs in cancer patients without diabetes [13, 53] attracted a great deal of attention in the past. In particular, it was shown in studies initiated by Prof. V.M. Dilman more than 40 years ago that the inclusion of phenformin as a long-term adjuvant modality into programs of metabolic rehabilitation of postoperative colorectal and breast cancer patients was associated with increased total and relapse-free survival [10, 54, 55]. In recent study, metformin was administered at a dose of 500 to 1000 mg daily for 2 to 4 weeks before surgery (that is, in a neoadjuvant manner) to breast cancer patients. Most of the patients, whose mean age was 59.9 years, were postmenopausal. This treatment
Clinical trials. Randomized clinical trials are now underway in several countries, mostly under auspices of the National Cancer Institute (http://www.clinicaltrials.gov/), to clarify the issue of metformin usability in nondiabetic patients. However, one of the initiators of these trials, Prof. Pamela Goodwin (University of Toronto, Canada), who, in particular, is directing a large multicenter, 5 year study of adjuvant metformin in nondiabetic breast cancer patients, is rather sober in her recent comment published in Science [59]: “We anticipate that it will take another 2 years to enroll everyone and maybe 2 to 3 years after that before we have results. All the preclinical and epidemiological evidence is pretty consistent and compelling, but all it's done is help us form a hypothesis... Until then, the best we can say is that metformin may be beneficial for cancer. We need to proceed from here very, very carefully.”

Resistance and responsiveness to metformin, and its tolerability and pharmacogenetics. The aspects listed in the subtitle are relevant directly to metformin use in obesity clinics (as was mentioned already above with regard to resistance), oncology [36, 60] and, undoubtedly, aging research (see [7] and below) and therefore need further scrutiny. Unresponsiveness to metformin, which was displayed by a number of normal and transformed cell lines, is probably caused by specific features of mitochondrial function as they relate to apoptosis [61]. In the field of pharmacogenetics, the relatively long known polymorphism of the organic cation transporter OCT1 [62] has been added to the ever increasing number of other markers associated with differences in the metabolism of biguanides and, thereby, in their effects [63, 64]. It is well known that, mainly because of gastrointestinal discomfort, metformin treatment is cancelled or interrupted in every fifth to sixth diabetic patient, and the rate of such adverse effects is increased in elderly subjects [65]. Such effects also are observed in nondiabetic cancer patients treated with metformin [56]. According to the latest Cochrane Collaboration estimates, the risk of lactic acidosis resulting from metformin intake (4-5 cases per 100000 subjects×year) is lower than previously thought [66]. In this regard, metformin is 7-10 times better than phenformin. Moreover, there is no evidence that antidiabetic biguanides can induce lactic acidosis in nondiabetics, even at older or advanced ages [13, 56]; therefore, the gastrointestinal side effects, especially in the elderly, seem to be the primary concern associated with metformin usage.

Metformin in the antiaging research agenda

Potential antiaging drugs are expected to prevent or eliminate age-related diseases [7]. Evidence that metformin is more beneficial that other antidiabetic drugs in reducing all-cause mortality and, therefore, increasing life expectancy in diabetic patients was presented earlier. This important feature is believed to be associated with the ability of metformin to influence the rate of macrovascular complications of diabetes [67, 68] rather than the basic mechanisms of aging. Such mechanisms as potential targets of metformin are under increasing scrutiny in the recent years. Among proximal targets under discussion are those involved in insulin resistance, insulin/IGF-1 system, and fatty acid oxidation and utilization [7, 69-71], which were considered earlier with regard to the antiaging effects of phenformin [3, 14, 72]. Among the most discussed targets of metformin are AMPK activity and AMP-related signaling, glycation reactions and glycation end-products, mitochondrial membranes, reactive oxygen species generation, epigenetic mechanisms, pluripotent stem cells, cell proliferative senescence and mTOR pathway [7, 71, 73-77]. Without digging into all possible mechanistic details, the only endpoints used to assess metformin as an antiaging agent will be considered below.

Metformin has been shown to slow-down lipofuscin accumulation, enhance locomotor activity and increase mean lifespan in Caenorhabditis elegans nematodes in a dose-dependent manner within concentration range of 1 to 50 mM in culture medium [78]. In R6/2 mice, used to model Huntington’s disease, metformin increased the lifespan of males, but not of females at a concentration of 2 mg/mL in drinking water but not at 5 mg/mL [79]. However, in order to differentiate changes in rodent lifespan resulting from influences on the basic mechanisms of aging rather than on specific disease-
related mortality, it is more appropriate, according to S.R. Spindler [80], to use genetically heterogeneous long-lived healthy populations, because short-lived or weakened animals have not been shown to predict longevity effects observed in long-lived ones. It is also mandatory to report the data with regards to monitored food consumption and body weight, thereby excluding the potential effects of caloric restriction; more than that, a positive control (e.g., a calorically restricted group) is highly desirable too [80]. Of note, rodent species, such as mice and rats, as well as nematodes and fruit flies, originated as a consequence of r-selection with an emphasis on a high growth rate resulting in numerous offspring, each of which has a relatively low probability of surviving to adulthood. In contrast to that, higher primate species including humans were molded in evolution primarily by K-selection resulting from living in crowded niches and having fewer offspring, each of which has a relatively high probability of surviving to adulthood [81, 82]. Nevertheless, for a number of practical reasons, properly chosen rodents remain the best choice for the selection of lifespan expanding drugs [80].

### Table 1. Data on lifespan in rodents receiving metformin

| Species, gender and strain | Drug administration mode and the age of its onset | Food consumption vs. control | Body weight vs. control | Lifespan, days (control/experiment, % change) | Ref. |
|----------------------------|--------------------------------------------------|-----------------------------|------------------------|---------------------------------------------|------|
| Mice, F Her-2/neu transgenic (FVB/N) | Drinking water, 100 mg/kg, 2 months | Decrease on months 4 & 6 (p <0.05) | No changes | 264±4/285±5 +8.0% | [83] |
| Mice, F Her-2/neu transgenic (FVB/N) | Drinking water, 100 mg/kg, 2 months | Decrease on months 7 & 8 | No changes | 285±12/304±10 +6.7% p for log-rank test 0.21 | C 396/359 -9.3% 396/352 -11.1%* p<0.001 [84]* |
| Mice, F SHR | Drinking water, 100 mg/kg, 3 months | Increase on months 12-16 (p<0.05) and decrease on month 22 (p<0.01) | Tends to decrease after 20 months | 388±29/535±32 +37.9%* p<0.01 | 814/898 +10.3% 727±23/878±7 +20.8% [85] |
| Mice, F SHR | Drinking water, 100 mg/kg, 2 months | No difference | No changes | 559±22/583±27 +4.1% | 941/972 +3.3% 892±12/897±28 +0.6% [86]** |
| Mice, F SHR | Drinking water, 100 mg/kg | No difference | No changes | 511±20/583±27 +14.1%, p= 0.17 | 941/972 +3.3% 881±13/897±28 +2.0% [88]** |
| Mice, F SHR | Drinking water, 100 mg/kg | 9 months | No changes | 583±18/619±20 +6.2% | 941/855 -9.1% 892±12/820±14 -8.8% [87] |
| Mice, F SHR | Drinking water, 100 mg/kg | 15 months | Decreases after 20 months | 668±16/647±21 -4.2% | 941/966 +2.7% 913±9/892±47 -2.3% [87] |
| Mice, F 129/Sv | Drinking water, 100 mg/kg, 3 months | Decrease on months 15-21 | Decreases after 25 months | 706±21/742±16 +5.1% | 930/966 +3.9% 910±9/913±19 +0.3% [87] |
| Mice, M 129/Sv | Drinking water, 100 mg/kg, 3 months | Decrease on months 15-21 | Decreases after 22 months | 662±28/573±27 -13.4%* | 1029/1044 +1.5% 951±32/931±30 -2.1% [87] |
| Rats, M Fisher F344 | Chow, 300 mg/kg, 6 months | No difference | Decreases at 48-74 months | 796±170/815±186 +2.4% | 1065/1062 -0.3% 1039±30/1061±3 +2.1% [89] |

Notes: * differences are statistically significant
# Data in [84] are not fully consistent with [83]
## Data in [86] and partly in [88] do not reproduce data in [85]; the mean lifespan in control was much shorter in [85] than in [86, 88]

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The aforesaid is of importance in considering experimental data about metformin, which are presented in Table 1. Most experiments were performed by the group lead by Prof. V.N. Anisimov [70, 83-88] with involvement, in some cases, of our laboratory. It is necessary to note, that unfortunately not all of these experiments fully satisfy the above requirements [80]. For example, the transgenic HER-2/neu mice are short-living and die mainly of mammary carcinomas. In some experiments, metformin-treated animals exhibited changes in food-consumption and body weight (see Table 1 and [83-85, 87-88]). Nevertheless, their representation, supplemented by the results reported by D.L. Smith et al. [89], helps to see the whole picture. Certain sex-related differences in effects observed in mice [87] were discussed earlier [90]. Noteworthy is that metformin tends to be more efficient upon earlier onset of its application ([88] and Table 1). This data corroborates previous experimental observations related to newborn macrosomy [91] and suggests that early-onset usage of biguanides, including during pregnancy and even preceding pregnancy, may be more beneficial than the late-onset, notwithstanding all difficulties in the practical realization of such recommendations.

On the whole, the data collected till present unfortunately are not always reproduced in the same experimental settings (Table 1), and demonstrate mixed results as has been already pointed out (see [89, 93] and in part [71, 92]). This conclusion relates as well to slowing down of aging rate assessed by the Gompertz model [83] since shifts of α parameter value in this model into the favorable side were not found in all the experiments mentioned above.

Conclusions

The main and uniform conclusion presented herein is that metformin acts ‘selectively’ and its effects on obesity, cancer, and lifespan experiments vary depending on gender [90], age [24, 25, 51, 88] and other factors, including tumor location (tissue specificity). It has been pointed out earlier [36, 55] that, in order to make use of the potential antitumor and/or weight-reducing activities of metformin, it is necessary to consider its direct and indirect effects, the presence or absence of glucose intolerance, and impairments in the insulin/IGF-1 system and tissue responsiveness, which are determined, among others, by pharmacogenetic factors (see also [53, 63]). This can be true as well for antiaging applications where, in particular, the evaluation of mTOR-related pathways and gene-expression markers is recommended as a means for the choice of geroprotectors [77, 94]. With all the known benefits of metformin in different areas, including reducing the rate of certain complications in diabetic patients [67, 68], only further studies will allow the specific molecular targets of metformin to be elucidated fully with respect to treatment and prevention of obesity, cancer, other age-related pathology and lifespan as discussed above.

Of special interest is the question of whether metformin is the most appropriate biguanide for oncology [36, 53, 95] and beyond. The authors of publications cited herein, while recognizing that only metformin is authorized by currently valid pharmacopeias for clinical use, note that phenformin, which is more associated with lactic acidosis in diabetic patients, may be more potent in anticancer applications, as follows from several preclinical studies [95, 96]. The potential antiaging activity of phenformin was reported long ago [97]. Therefore, it makes sense to compare directly metformin and phenformin for their ability to influence lifespan under identical experimental settings and in the same experiment and thus to make an additional step to developing of approaches to the “…recommendation for healthy life, which may help to bring one's lifespan several years closer to the reliably recorded maximum” [98].

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CONFLICT OF INTERESTS STATEMENT

The authors of this manuscript have no conflict of interest to declare.

Note added after submitting of this paper

Recently, R.Boussageon et al. performed the meta-analysis of randomised controlled trials of metformin efficacy against morbidity or mortality in patients with type 2 diabetes. Metformin did not significantly affect the primary outcomes: all-cause mortality, risk ratio (RR) = 0.99 (95% CI: 0.75 to 1.31), and cardiovascular mortality. The secondary cardiovascular outcomes were also unaffected by metformin treatment. The authors concluded that although metformin is considered the gold standard, its benefit/risk ratio remains uncertain, and needs further study [Boussageon R, Supper I, Bejan-Angoulvant T, Kellou N, Cucherat M, Boissel JP, et al. (2012) Reappraisal of Metformin Efficacy in the Treatment of Type 2 Diabetes: A Meta-Analysis of
Randomised Controlled Trials. PLoS Med 9(4): e1001204. Epub 2012 Apr 10.

Also, H.Noto et al. performed a search for pertinent articles published as of October 12, 2011 and calculated pooled risk ratios (RRs) for overall cancer mortality and cancer incidence in patients with diabetes treated with metformin. The use of metformin was associated with significantly lower risks of cancer mortality and incidence. However, as mentioned by the authors, this analysis was mainly based on observational studies and the findings underscore the more need for long-term randomized clinical trials to confirm potential benefit for individuals with diabetes [Noto H, Goto A, Tsujimoto T, Noda M. (2012) Cancer Risk in Diabetic Patients Treated with Metformin: A Systematic Review and Meta-analysis. PLoS One. 7(3):e33411. Epub 2012 Mar 20].

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