

# PUBLISHED VERSION

Wen-Peng You, Maciej Henneberg

**Type 1 diabetes prevalence increasing globally and regionally: the role of natural selection and life expectancy at birth**

BMJ Open Diabetes Research & Care, 2016; 4(1):e000161-1-e000161-8

This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

Originally published at:

<http://doi.org/10.1136/bmjdr-2015-000161>

## PERMISSIONS

<http://creativecommons.org/licenses/by-nc/4.0/>



**Attribution-NonCommercial 4.0 International** (CC BY-NC 4.0)

This is a human-readable summary of (and not a substitute for) the [license](#).

[Disclaimer](#)

### You are free to:

**Share** — copy and redistribute the material in any medium or format

**Adapt** — remix, transform, and build upon the material

The licensor cannot revoke these freedoms as long as you follow the license terms.

### Under the following terms:



**Attribution** — You must give [appropriate credit](#), provide a link to the license, and [indicate if changes were made](#). You may do so in any reasonable manner, but not in any way that suggests the licensor endorses you or your use.



**NonCommercial** — You may not use the material for [commercial purposes](#).

**No additional restrictions** — You may not apply legal terms or [technological measures](#) that legally restrict others from doing anything the license permits.

<http://hdl.handle.net/2440/98907>

# Type 1 diabetes prevalence increasing globally and regionally: the role of natural selection and life expectancy at birth

Wen-Peng You,<sup>1</sup> Maciej Henneberg<sup>1,2</sup>

**To cite:** You W-P, Henneberg M. Type 1 diabetes prevalence increasing globally and regionally: the role of natural selection and life expectancy at birth. *BMJ Open Diabetes Research and Care* 2016;**4**: e000161. doi:10.1136/bmjdr-2015-000161

► Additional material is available. To view please visit the journal online (<http://dx.doi.org/10.1136/bmjdr-2015-000161>).

Received 8 October 2015  
Revised 18 January 2016  
Accepted 21 January 2016



CrossMark

<sup>1</sup>School of Medicine, The University of Adelaide, Adelaide, South Australia, Australia

<sup>2</sup>Institute of Evolutionary Medicine, University of Zurich, Zurich, Switzerland

**Correspondence to**  
Dr Wen-Peng You; Wenpeng.you@adelaide.edu.au

## ABSTRACT

**Objective:** Prevalence of type 1 diabetes (T1D) disease is increasing worldwide. We aim to test correlation of T1D prevalence to the reduced natural selection measured by Biological State Index ( $I_{bs}$ ).  
**Research design and methods:** Country-specific estimates of T1D prevalence, life expectancy, obesity prevalence rate, urbanization rates, per capita sugars consumption and per capita gross domestic product (GDP) were obtained.  $I_{bs}$  and country-specific longevity ( $e_{50}$ ) increase for each country were self-calculated. These data were then matched to T1D prevalence by country for our ecological study among 118 countries. Countries were also grouped to study the associations in different regions. SPSS V.22 was used for correlation analysis.

**Results:** Worldwide, both  $I_{bs}$  and life expectancy at birth ( $I_{bs}$  proxy) were significantly correlated to T1D prevalence in Pearson  $r$  ( $r=0.713$ ,  $p<0.001$  and  $r=0.722$ ,  $p<0.001$ , respectively) and Spearman's  $r$  ( $r=0.724$ ,  $p<0.001$  and  $r=0.689$ ,  $p<0.001$ , respectively). T1D prevalence was not correlated to longevity increase measured as life expectancy at 50 years old. T1D prevalence was significantly associated with  $I_{bs}$  ( $r=0.307$ ,  $p<0.001$ ) and newborn life expectancy ( $r=0.349$ ,  $p<0.001$ ) independent of per capita total sugar consumption, per capita GDP, urbanization and obesity prevalence in partial correlation. Globally, both life expectancy at birth and  $I_{bs}$  exponentially correlated to T1D prevalence. Pearson correlations generally existed in different country categorizations by geographic region, culture background and economic status.

**Conclusions:** Reduced natural selection may have contributed to the increasing T1D prevalence worldwide. T1D epidemiology study in total population may be the practical solution to identify the causes of increasing T1D prevalence.

## BACKGROUND

Type 1 diabetes (T1D) is an autoimmune disease with a strong genetic component.<sup>1 2</sup> It can occur at any age, but tends to develop in childhood,<sup>3</sup> so it has long been called 'juvenile diabetes'. T1D is characterized by

## Key messages

- Prevalence of type 1 diabetes (T1D) is increasing worldwide.
- T1D prevalence may reflect accumulation of faulty genes in gene pools of populations because of its partly genetic background.
- Relaxation of the opportunity for natural selection correlates with T1D prevalence.

destruction of pancreatic  $\beta$ -cells, culminating in absolute insulin deficiency.<sup>4</sup> As of 2014, an estimated 387 million people have diabetes worldwide,<sup>5</sup> of which T1D accounts for between 5% and 10%.<sup>6</sup> Diabetic complications continue to be a major cause of morbidity and mortality in persons with T1D.<sup>7</sup> Great efforts have been made to assess the incidence and prevalence of T1D. Unfortunately, the exact etiology and pathogenesis of T1D is still unknown. Generally, longitudinal or cross-sectional studies are often locally or regionally performed. Consequently, it is difficult to access generalizable results because the epidemiology of T1D is known to be heterogeneous regarding geography and ethnicity. Genetic predisposition to T1D is only alleged to explain some of the geographic variability in T1D occurrence, but it cannot account for its rapidly increasing frequency.<sup>8</sup> A number of studies have associated gross domestic product (GDP) level with T1D prevalence or incidence,<sup>8–11</sup> but GDP does not fully explain variations and trends in T1D prevalence rates observed in many countries, for example Japan. It has been postulated that environmental factors may be able to trigger an autoimmune destruction of the  $\beta$ -cells leading to absolute dependence on insulin treatment,<sup>8 9 12–17</sup> however, these environmental factors are circumstantial.<sup>13</sup>

Natural selection, as one of the basic mechanisms of evolution, is the differential survival and fertility of individuals due to

differences in phenotype that reflect genetic differences. In our modern society, natural selection still acts on all members of a population, selecting those individuals that have an increased reproductive success (survival and/or fertility).<sup>18</sup> The 'Biological State Index ( $I_{bs}$ )' has been proposed to measure the populational reproductive success by taking into account potential loss of reproductive success by dying at age  $x$ , summed over all age categories.<sup>19 20</sup> The  $I_{bs}$  is calculated by combining age-specific death frequency ( $dx$  variable of a life table) with an age-specific reproductive loss ( $s_x$ ):

$$I_{bs} = 1 - \sum_{x=0}^{x=\omega} dx \cdot Sx$$

where  $dx$  is the frequency of death at age  $x$  or represents the mortality rate,  $s_x$  the reproductive loss from dying at age  $x$ , that is, the estimated probability of not possessing the complete number of births at age  $x$ .  $s_x$  is based on the cumulative number of births at specific ages.<sup>20 21</sup> The construction and interpretation of the  $I_{bs}$  was predicated on the assumption that heritability of human fertility variance is negligible.<sup>22</sup>

An  $I_{bs}$  value of 1 indicates total adaptation of the population to their environment (ability to overcome selection pressures that are present). An  $I_{bs}$  value of 0 signifies a total lack of adaptation (inability to overcome selection pressures that are present), and an impossibility to give life to the next generation. An  $I_{bs}$  value close to 0 indicates large effective natural selection pressures acting on a population, since few individuals are surviving to produce offspring. In such a scenario there is a possibility for fast evolution, since many genes may not be passed to the next generation. An  $I_{bs}$  value close to 1 indicates that natural selection is not having much effect on the population since many individuals are able to maximally contribute to producing the next generation. Thus, the  $I_{bs}$  permits the estimation of the magnitude of the successful reproduction of a population.

The genetic trait of T1D may allow individuals from a population to pass their T1D genes on to their next generation. What fraction of a population had a chance to fully participate in reproducing under a given set of mortality conditions may be associated with the proportion of population carrying T1D genes in the next generation. Previously, Stephan and Henneberg<sup>23</sup> raised a concern that the developed populations may have accumulated more unfavorable genes, such as T1D genes because natural selection (measured by the  $I_{bs}$ ) has been greatly reduced. Recently, a systematic review concluded that the T1D prevalence rate was associated with age increase in population.<sup>3</sup> Therefore, the objective of the current paper was to use country-specific data to test, from a global perspective, that population with greater  $I_{bs}$  value, fuelled by life expectancy at birth, may have higher T1D prevalence, using empirical, macro-level data collected from the major international organizations.

## MATERIAL AND METHOD

The dependent variable in the analysis was the country-specific estimate of T1D prevalence (sum of rates in 0–14 and 15+years old groups) which was published by the International Diabetes Federation (IDF) in 2000.<sup>24</sup> We used  $I_{bs}$  and life expectancy at birth of each country as the independent variables. The  $I_{bs}$  was calculated as proposed by Henneberg<sup>20</sup> and Henneberg and Piontek<sup>19</sup> with the fertility data of each country published by United Nations in 2008<sup>25</sup> and the mortality data of life tables (2009) published by WHO in 2012.<sup>26</sup> Life expectancy at birth is a proxy for  $I_{bs}$  since it expresses an opportunity of a newborn to survive to a specific age, which is usually an age falling into reproductive life span or above it. We extracted the country-specific life expectancy (years) at birth by country published by WHO in 2013.<sup>27</sup>

GDP,<sup>8–11</sup> urbanization<sup>8</sup> and body weight status<sup>17</sup> have been associated with T1D prevalence. It has been suggested that a population with greater sugar consumption may have greater diabetes (total) prevalence<sup>28</sup> and that sugar consumption may affect health of patients with T1D.<sup>29</sup> Therefore, we controlled for market availability of sugars and sweeteners (sugars in short) in g/capita/day in each country for year 2010 from United Nations Food and Agricultural Organization (FAO) Food Balance Sheet, per capita GDP (expressed in purchasing power parity in 2010 US dollars for comparability among countries) and urbanization (percentage of population living in urban areas in each country in 2010) from the World Bank's World Development Indicators Database, and obesity prevalence rate (percentage of the population aged 18+ years old with body mass index (BMI)  $\geq 30$  kg/m<sup>2</sup> in each country during 2010) from the WHO.

No ethical approval or written informed consent for participation was required in this study as all the aforementioned data were freely downloaded from the United Nations agents' websites.

We aligned all independent variables and confounding factors with the country-specific T1D prevalence, and obtained a set of data for 118 countries. All country-specific data were put in a uniform format. Each country was treated individually and all of their available information was analysed. Data of calculated  $I_{bs}$  and summary statistics are further described in the supporting information (see online supplementary tables S1 and S2).

In order to demonstrate the universal association between T1D prevalence and  $I_{bs}$  and life expectancy at birth respectively in different country groupings, we categorized the countries for correlation analyses based on (1) the WHO regional classifications;<sup>30</sup> (2) the strong contrast in terms of geographic distributions, per capita GDP levels and/or cultural backgrounds. We analysed the correlation in the six country groupings: Latin America and the Caribbean (LAC),<sup>31</sup> the Arab World,<sup>32</sup> the Organisation for Economic Co-operation and Development (OECD),<sup>33</sup>

**Table 1** Global associations between T1D prevalence rate and  $I_{bs}$  and life expectancy (years) at birth respectively\*

	Pearson's r		Spearman's r		Partial correlation†	
	r	n	r	n	r	df
Log $I_{bs}$	0.713	118	0.724	118	0.307	103
Log life expectancy at birth	0.722	118	0.689	118	0.349	103
Log sugars per capita	0.666	109	0.534	109	—	—
Log GDP per capita	0.720	116	0.749	116	—	—
Log BMI $\geq 30$ prevalence	0.636	109	0.538	118	—	—
Log urbanization	0.507	118	0.567	118	—	—

\*All correlations are significant at the 0.001 level (two-tailed).

†Keeping GDP, BMI, urbanization and sugars intake constant.

BMI, body mass index; GDP, gross domestic product;  $I_{bs}$ , Biological State Index; T1D, type 1 diabetes.

European Economic Area (EEA),<sup>34</sup> Asia Cooperation Dialogue (ACD)<sup>35</sup> and the Asia-Pacific Economic Cooperation (APEC).<sup>36</sup> In our analysis, we only included those countries for which we could access their data for the specific groupings.

To a large extent, grouping countries for analysis may also allow us to align our findings against previous local or regional studies regarding heterogeneous T1D epidemiology due to various geographic location and ethnicity.

It might be considered that T1D prevalence is a result of the increase in longevity rather than relaxed natural selection of the genetic background of T1D since general health and advances in medical care improve survival of patients with T1D. This consideration is clarified by much stronger correlation between T1D prevalence and  $I_{bs}$  than that between T1D and life expectancy ( $e_{50}$ ) increase (across two periods of 1950–1955 and 2005–2010). The rationale to use country-specific life expectancy ( $e_{50}$ ) is that (1) the estimate of life expectancy based on this segment of the population may not be biased with child (0–15 years) mortality, in particular due to deaths caused by T1D disease, and (2) deaths of adults, especially females, during the reproductive life span (15–50 years) that may differentiate numbers of T1D genes passed on to new generations. Thus, we obtained country-specific life expectancies ( $e_{50}$ ) for the periods of 1950–1955 and 2005–2010 respectively from the WHO life tables.<sup>37</sup> And then we calculated the life expectancy increase from the period of 1950–1955 to the period of 2005–2010 for each country producing a new variable, which is 'life expectancy increase ( $e_{50}$ , 1950–2010)' for each country across the 55 years. We repeated the above correlation analysis after we replaced the variable of 'life expectancy at birth' with life expectancy increase ( $e_{50}$ , 1950–2010).

Pearson's correlation coefficient, Spearman's  $r$  and partial correlation analyses were conducted using SPSS V. 22 (SPSS Inc, Chicago Illinois, USA). In this study, the data were log transformed for correlation analysis in SPSS.

## RESULTS

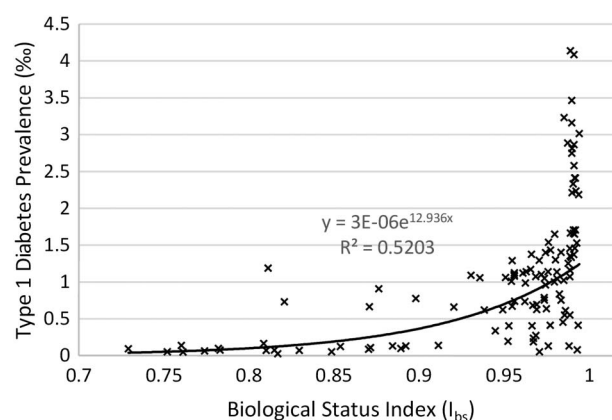
Worldwide T1D prevalence was in significant associations with  $I_{bs}$  using Pearson  $r$  ( $r=0.713$ ,  $p<0.001$ ) and

Spearman's  $r$  ( $r=0.724$ ,  $p<0.001$ ), respectively. The similar associations were also observed between T1D and life expectancy in Pearson model ( $r=0.722$ ,  $p<0.001$ ) and Spearman's model ( $r=0.689$ ,  $p<0.001$ ), respectively (table 1). Further investigation with partial correlation analysis showed that worldwide the association between T1D prevalence was still strongly associated with  $I_{bs}$  ( $r=0.307$ ,  $p<0.001$ ) and life expectancy ( $r=0.349$ ,  $p<0.001$ ) when we controlled for per capita total sugars availability, per capita GDP, urbanization and obesity prevalence (table 1). All confounders were in significant associations with T1D prevalence rate in both Pearson  $r$  and Spearman's  $r$ .

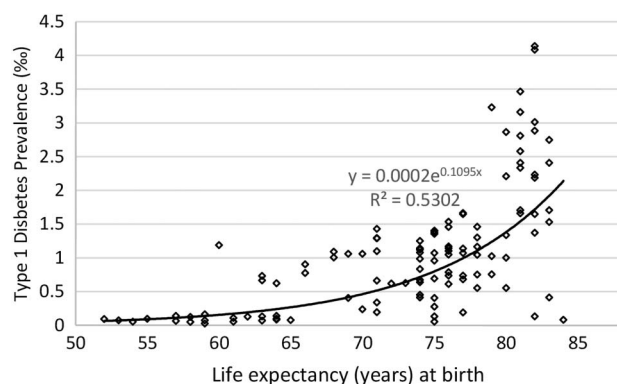
Globally, T1D is noted to be exponentially related with both  $I_{bs}$  ( $R^2=0.5203$ ) and life expectancy ( $R^2=0.5302$ ) (figures 1 and 2) after we removed two outliers of extremely high T1D prevalence in Finland and Sweden, respectively.

Worldwide, strong positive Pearson  $r$  coefficients were observed in the correlation of  $I_{bs}$  to life expectancy (proxy of  $I_{bs}$ ,  $r=0.908$ ,  $p<0.001$ ) and to GDP ( $r=0.781$ ,  $p<0.001$ ).

Table 2 presented that Europe had the significant associations with  $I_{bs}$  ( $r=0.502$ ,  $p=0.001$ ) and life expectancy ( $r=0.610$ ,  $p<0.001$ ), respectively. The sub-Europe (EEA) also had the very strong associations of T1D prevalence

**Figure 1** Relationship between  $I_{bs}$  and type 1 diabetes prevalence worldwide.





**Figure 2** Relationship between life expectancy (years) at birth and type 1 diabetes prevalence worldwide.

to both  $I_{bs}$  ( $r=0.479$ ,  $p=0.009$ ) and life expectancy ( $r=0.574$ ,  $p=0.001$ ) (table 3). We observed the slightly negative correlations of T1D prevalence rate to the  $I_{bs}$  and life expectancy, respectively, in two Asian country groupings, SEARO (South-East Asia) and the ACD (table 3). Both  $I_{bs}$  and life expectancy were insignificantly associated with T1D prevalence rate in other four WHO regions, AFRO, AMRO, EMRO and WPRO (table 2).

Table 3 showed that the T1D prevalence rate was associated with both the  $I_{bs}$  and newborn life expectancy in the groupings consisting of countries with similar cultures (Arab World and EEA) and different cultures (APEC and OECD), with similar economy status (OECD) and those with economy status in disparity (APEC), and with the heterogeneous region (LAC) and the homogeneous area (EEA and ACD).

Table 4 showed that in general T1D prevalence is not correlated to longevity increase, but strongly correlated to  $I_{bs}$  at country level. Life expectancy increase (e50, 1950–2010) does not correlate significantly with T1D prevalence in Pearson correlation ( $r=0.165$ ,  $p=0.079$ ) or Spearman's  $r$  ( $r=0.166$ ,  $p=0.077$ ). These contrasted with the correlation between  $I_{bs}$  and T1D prevalence in Pearson correlation ( $r=0.713$ ,  $p<0.001$ ) and in Spearman  $r$  ( $r=0.724$ ,  $p<0.001$ ), respectively. In partial correlation analysis, when we controlled for GDP, BMI $\geq 30$ , urbanization and sugars intake, T1D was negatively and insignificantly correlated to life expectancy increase (e50, 1950–2010) ( $r=-0.070$ ,

$p=0.487$ ), but it was in strong and significant correlation to  $I_{bs}$  ( $r=0.276$ ,  $p=0.005$ ).

## DISCUSSION

The worldwide trend of increased T1D prevalence likelihood has multiple etiologies, which may act through multiple mechanisms. By assessing the T1D prevalence rate data for 118 countries we have shown that globally and regionally population which had greater value of  $I_{bs}$  (less opportunity for natural selection) may have greater T1D prevalence and secondly, that newborn life expectancy was significantly associated with T1D prevalence rate at population level.

Overall, the operation of natural selection on contemporary populations is declining due to modern medicine,<sup>23</sup> but the magnitude of the decline may differ between countries due to their specific level of sanitation, medical interventions and public health measures. Natural selection is still one of the major evolutionary forces that informs changes in gene frequencies in a population through the action of differential fertility and mortality over generations.<sup>38</sup> For example, studies have shown that the increasing prevalence rates of a partially heritable disease, nasal septa and lacrimal bone defects may be attributed to the decreasing effect of natural selection.<sup>39</sup> More than 40 genetic loci located in different chromosomes have been associated with T1D in multiple studies.<sup>1–2</sup> Although T1D can be fatal, the majority of genetically predisposed people do not develop T1D.<sup>40</sup> This allows for accumulation of genetic predisposition in human populations. This accumulation will increase when fewer persons who developed a disease would die. Differential fertility and mortality are the basic events of natural selection, which operate singly or jointly to determine the fitness (reproductive success) of a particular population in a given environment.<sup>38</sup> The country-specific fertility-based and mortality-based  $I_{bs}$  at different levels may indicate their different successful reproduction opportunities of individuals in the succeeding countries.<sup>23</sup> The reproduction success opportunity of each population may determine their magnitude of T1D genes accumulation, thus influenced prevalence rate of patients with T1D in their next generations. In the present study, the correlation of  $I_{bs}$

**Table 2** Associations between T1D prevalence rate and  $I_{bs}$  and newborn life expectancy (years) at birth respectively in the WHO regions

WHO region	$I_{bs}$		Newborn life expectancy	
	Pearson's r	Significance	Pearson's r	Significance
AFRO (n=21)	0.343	0.128	0.214	0.351
AMRO (n=28)	0.145	0.461	0.173	0.380
EMRO (n=15)	0.783	0.001	0.541	0.037
EURO (n=38)	0.502	0.001	0.610	<0.001
SEARO (n=5)	−0.479	0.415	−0.436	0.463
WPRO (n=11)	0.345	0.298	0.330	0.322

$I_{bs}$ , Biological State Index; T1D, type 1 diabetes.

**Table 3** Associations between T1D prevalence rate and both the  $I_{bs}$  and life expectancy (years) at birth in different country groupings categorized considering their socioeconomic, geographic location and cultural backgrounds

Country grouping	$I_{bs}$		Life expectancy at birth	
	Pearson's r	Significance	Pearson's r	Significance
APEC (n=16)	0.340	0.197	0.369	0.160
Arab World (n=13)	0.748	0.003	0.469	0.106
EEA (n=29)	0.479	0.009	0.574	0.001
ACD (n=20)	-0.268	0.253	-0.392	0.087
OECD (n=34)	0.365	0.034	0.155	0.382
LAC (n=26)	0.524	0.006	0.044	0.831

ACD, Asia Cooperation Dialogue; APEC, Asia-Pacific Economic Cooperation; EEA, European Economic Area;  $I_{bs}$ , Biological State Index; LAC, Latin America and the Caribbean; OECD, Organisation for Economic Co-operation and Development; T1D, type 1 diabetes.

to the T1D prevalence rate has been observed, which was compatible with suggestion that lower opportunity for selection allows accumulation of unfavorable genes.<sup>23 41</sup> Our analysis of correlations between T1D prevalence and  $I_{bs}$  by region or by WHO grouped countries seem to indicate that in regions where insulin was available earlier and that had better availability of healthcare the relationship is stronger. This provides the analog of a snapshot what could happen at different times in the same region as time from insulin introduction and improvement of healthcare increased. Thus, the distribution across different populations could be interpreted as a surrogate measure of the evolution in time of T1D prevalence after the introduction of insulin. Artificial insulin introduced for T1D treatment and increasing insulin availability may have played a key role in reducing natural selection as insulin enables countless people with onset of T1D to survive<sup>3</sup> and maintain normal reproductive capacity.<sup>42</sup> This may have been boosting T1D genes accumulation and prevalence of T1D. T1D can affect people of any age, but usually occurs in children or sexually mature young adults<sup>3</sup> who have greater potential to reproduce than older adults. T1D has been historically, and continues to be, the most common type of diabetes in children and adolescents.<sup>43</sup> Insulin is the priority for T1D treatment. Otherwise, patient with T1D may only live up to 1 year, some only a week. Several human generations have benefited from insulin since it was discovered and became available in early 1920s.<sup>44</sup> Reduced natural selection boosted by

insulin treatment of several generations may have enabled cumulative effect of T1D genes frequency in human population to occur quickly and to be noticeable for a couple of decades.<sup>9</sup> Studies have shown that a partially heritable disease, phenylketonuria was only noticeable after being accumulated for several generations<sup>23</sup> with about 2% increase in each.<sup>45</sup>

T1D prevalence/incidence is increasing worldwide<sup>46</sup> with special regard to the developed countries.<sup>9 10 47</sup> This may be partially attributable to earlier and greater affordability of insulin, in addition to relative more reduced natural selection (greater  $I_{bs}$  values) in those developed countries. Although exogenous insulin can be obtained from animals (bovine and porcine),<sup>44</sup> production, transportation, storage and administration of such insulin was extremely expensive,<sup>48</sup> which may be beyond the affordability of many patients with T1D, especially those from developing countries. Biosynthetic insulin based on DNA technology has been commercially available since 1982<sup>44</sup> and it has been thought that it can continue to accommodate global demand<sup>44</sup> because of low cost from the production to administration. However, unfortunately life-saving insulin is still less accessible, affordable, or both to people diagnosed with diabetes in a developing country than their counterparts in the developed world.<sup>49</sup> This lower survivorship of patients with T1D may contribute to lower prevalence figures directly, besides the fact that less predisposing genes have accumulated in the gene pools of those countries.

**Table 4** Comparing correlations between type 1 diabetes prevalence and  $I_{bs}$  and life expectancies increase (e50)

	Pearson's r			Spearman's r			Partial correlation*		
	r	p Value	n	r	p Value	n	r	p Value	df
Log $I_{bs}$	0.713	<0.001	118	0.724	<0.001	118	0.276	0.005	98
Log life expectancy (e50) increase	0.165	0.079	114	0.166	0.077	114	-0.070	0.487	98
Log sugars per capita	0.666	<0.001	109	0.534	<0.001	109	—	—	—
Log GDP per capita	0.720	0.720	116	0.749	<0.001	116	—	—	—
Log BMI $\geq$ 30 prevalence	0.636	<0.001	109	0.538	<0.001	118	—	—	—
Log urbanization	0.507	<0.001	118	0.567	<0.001	118	—	—	—

\*Keeping GDP, BMI, urbanization and sugars intake constant.

BMI, body mass index; GDP, gross domestic product;  $I_{bs}$ , Biological State Index.

Our study showed that the relationship between life expectancy and T1D prevalence rate was exponential (figure 2,  $R^2=0.5266$ ). The Australian Institute of Health and Welfare (AIHW) also indicated the exponential relationship between T1D prevalence rate between age increase of Australian population through the Australian National Diabetes Register.<sup>50</sup> Additionally, Neville *et al*<sup>51</sup> reported that the increased longevity of diabetic patients contributed to the increasing prevalence of diabetes in Japanese population. The life expectancy gap between patients with T1D and non-diabetic people has reduced significantly<sup>52</sup> due to developments in sanitation, medical interventions and public health measures. Therefore, the underlying reason for the exponential relationship in our study may be because the number of individual patients with T1D have increased in the human population.<sup>3</sup> The American Diabetes Association has also stated that the majority of individuals with T1D are adults even though T1D has been more frequent and a relatively straightforward diagnosis in children.<sup>53</sup>

The correlations of T1D prevalence rates to both  $I_{bs}$  and life expectancy were not only observed worldwide, but also in different country groupings sharing specific characteristics such as geographic locations (table 2), culture backgrounds (table 3) and affiliations to international functional organizations (table 3). Results' highlights indicated that the correlations of  $I_{bs}$  and life expectancy to T1D were significant or very strong in European country groupings (WHO-Europe in table 2 and EEA in table 3), but very weak in Asian country grouping (WHO-SEA in table 1 and ACD in table 3). This may be attributable to high-genetic predispositions<sup>13–15 46</sup> in Europe, but low-genetic predisposition in Asia.<sup>46 53</sup>

Although we found that the correlations of  $I_{bs}$  and life expectancy to T1D prevalence rate existed globally and in different country groupings categorized with a variety of criteria, there are several limitations, including the intrinsic limitations (conceptualized as ecological fallacy) to this study.

First, the data analyzed were calculated for per capita in each country, so we could only demonstrate the relationships between T1D prevalence rate and  $I_{bs}$  and life expectancy at country/population level, which does not necessarily correspond to the same relationships holding true at the individual level. We also need to point out that it would be difficult to test the relationships at the individual level due to very rare T1D occurrence rate.

Second, the slow changes in the genetic code of the human populations may not fully explain the increasing T1D prevalence. Non-genetic (environmental) factors partially determine whether, and how risk-associated genotypes may lead to overt T1D disease. Unfortunately, our  $I_{bs}$  does not indicate if fitness change at population level is due to evolution of individuals or change/s. It may also be that altered lower natural immunity to infections following decades of using antibiotics may influence increased rates of autoimmune diseases including T1D.

Third, the data compiled and/or collected by the major international agencies (IDF, WHO, FAO and the World Bank) are fairly crude, and may contain some random errors.

Finally, current evidence of the increasing frequency of many heritable genetic disorders, including T1D does not appear to be available. To the best of our knowledge, the T1D prevalence rate for all age groups at country level published by IDF may be the only version to single out T1D prevalence worldwide after consulting the major diabetes research or data collecting institutions. This may be because clinically, adult T1D is difficult to discriminate from certain forms of type 2 diabetes and from latent autoimmune diabetes in adults (LADA).<sup>53</sup> Therefore, we do not know how much this set of data was confounded by other forms of diabetes.

The current prevailing paradigm on the increasing prevalence of T1D is that environmental pressures are now able to trigger genotypes.<sup>8 9 12–17</sup> Currently, medical gene intervention in modern medicine at this stage cannot remove T1D genes, and eugenics (improvement in the genetic stock) can offer no direction due to ethics issue. Therefore, study of T1D epidemiology based on prevalence/incidence T1D data of all age groups has become imperative as it may offer optimal solution to address or at least slow down T1D genetic load increases in different populations.

## CONCLUSIONS

Our study suggested that reduced natural selection ( $I_{bs}$ ) may be the major contributor to the increasing prevalence of T1D worldwide with special regard to European countries. It seems that T1D epidemiology study based on all age groups may be the practical solution to identify the causes of increasing T1D prevalence and to address, or at least slow down, T1D genetic load increases in different populations as modern medicine cannot operate effectively at the gene level yet.

**Acknowledgements** The authors express appreciation to Dr Arthur Saniotis for his editorial assistance.

**Contributors** MH conceived the idea for this study. W-PY extracted the data and W-PY took responsibility of data integrity. MH and W-PY interpreted the data. W-PY wrote the manuscript. W-PY and MH edited and approved the manuscript for submission to the journal.

**Competing interests** Financial support to MH was provided by the Wood Jones Bequest to the University of Adelaide.

**Provenance and peer review** Not commissioned; externally peer reviewed.

**Data sharing statement** Not relevant as all data for this research were freely downloaded from the United Nations agents' websites. However, the corresponding authors could supply the country-specific data used in this research on request.

**Open Access** This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

## REFERENCES

- Noble JA, Erlich HA. Genetics of type 1 diabetes. *Cold Spring Harb Perspect Med* 2012;2:1–15.
- Steck AK, Rewers MJ. Genetics of type 1 diabetes. *Clin Chem* 2011;57:176–85.
- Frese T, Sandholzer H. The epidemiology of type 1 diabetes mellitus. In: Escher S, Li A, eds. *type 1 diabetes*. InTech, 2013:1–22.
- Maahs DM, West NA, Lawrence JM, et al. Epidemiology of type 1 diabetes. *Endocrinol Metab Clin North Am* 2010;39:481–97.
- International Diabetes Federation. *Key findings 2014*. <http://www.idf.org/diabetesatlas/update-2014>
- Melmed S, Polonsky SK, et al. *Williams textbook of endocrinology*. 12th edn. Philadelphia: Elsevier/Saunders, 2011.
- Libby P, Nathan DM, Abraham K, et al. Report of The National Heart, Lung, and Blood Institute—National Institute of Diabetes and Digestive and Kidney diseases working group on cardiovascular complications of type 1 diabetes mellitus. *Circulation* 2005;111:3489–93.
- Borchers AT, Uibo R, Gershwin ME. The geoepidemiology of type 1 diabetes. *Autoimmun Rev* 2010;9:A355–65.
- Gale EA. The rise of childhood type 1 diabetes in the 20th century. *Diabetes* 2002;51:3353–61.
- International Diabetes Federation. *IDF diabetes atlas*. 3rd edn. Brussels, Belgium: International Diabetes Federation, 2006.
- Patterson CC, Dahlquist G, Soltész G, et al. Is childhood-onset type 1 diabetes a wealth-related disease? An ecological analysis of European incidence rates. *Diabetologia* 2001;44(Suppl 3):B9–16.
- Eisenbarth GS, Naylor RC, Rabinow SL. Type 1 diabetes as a chronic autoimmune disease. *J Diabet Complications* 1988;2:54–8.
- Patterson C, Guariguata L, Dahlquist G, et al. Diabetes in the young—a global view and worldwide estimates of numbers of children with type 1 diabetes. *Diabetes Res Clin Pract* 2014;103:161–75.
- Kaprio J, Tuomilehto J, Koskenvuo M, et al. Concordance for type 1 (insulin-dependent) and type 2 (non-insulin-dependent) diabetes mellitus in a population-based cohort of twins in Finland. *Diabetologia* 1992;35:1060–7.
- Kyvik KO, Green A, Beck-Nielsen H. Concordance rates of insulin dependent diabetes mellitus: a population based study of young Danish twins. *BMJ* 1995;311:913–17.
- Gale EA. Declassifying diabetes. *Diabetologia* 2006;49:1989–95.
- Wilkin TJ. The accelerator hypothesis: weight gain as the missing link between type I and type II diabetes. *Diabetologia* 2001;44:914–22.
- Byars SG, Ewbank D, Govindaraju DR et al. Colloquium papers: natural selection in a contemporary human population. *Proc Natl Acad Sci USA* 2010;107(1):1787–92.
- Henneberg M, Piontek J. Biological state index of human groups. *Przegl Anthropologiczny* 1975;XLI:191–201.
- Henneberg M. Reproductive possibilities and estimations of the biological dynamics of earlier human populations. *J Hum Evol* 1976;5:41–8.
- Henneberg M. Notes on the reproduction possibilities of human prehistorical populations. *Przegl Anthropologiczny* 1975;41:75–89.
- Henneberg M. Quantitative evaluation of actual intensity of natural selection through differential fertility in human populations. *Am J Phys Anthropol* 1985;66:181.
- Stephan CN, Henneberg M. Medicine may be reducing the human capacity to survive. *Med Hypotheses* 2001;57:633–7.
- International Diabetes Federation. *IDF diabetes atlas*. 1st edn. Brussels, Belgium: International Diabetes Federation, 2000.
- The United Nations. *World Fertility Data 2008*. 2012 (29.07.2015). <http://www.un.org>.
- WHO. *World Health Statistics 2012. Life tables for WHO member states*. Geneva: World Health Organization, 2012.
- WHO. *Life expectancy at birth (years)*. 2013. (29.07.2015). <http://apps.who.int>
- Basu S, Stuckler D, McKee M, et al. Nutritional determinants of worldwide diabetes: an econometric study of food markets and diabetes prevalence in 173 countries. *Public Health Nutr* 2013;16:179–86.
- Lamb MM, Frederiksen B, Seifert JA, et al. Sugar intake is associated with progression from islet autoimmunity to type 1 diabetes: the Diabetes Autoimmunity Study in the Young. *Diabetologia* 2015;58:2027–34.
- WHO. *WHO regional offices*. (11.26.2015). <http://www.who.int>
- The United Nations Educational Scientific and Cultural Organization. *UNESCO Regions-Latin America and the Caribbean*. 2014. <http://www.unesco.org>
- The World Bank. *Arab World I Data*. 2015. <http://data.worldbank.org>
- The OECD. *List of OECD member countries*. 2015. <http://www.oecd.org>
- The European Free Trade Association. *What is the European Economic Area?* <http://www.efta.int/eea>
- Asia Cooperation Dialogue. *Member countries*. <http://www.acddialogue.com>
- Asia-Pacific Economic Cooperation. *Member Economies-Asia-Pacific Economic Cooperation*. <http://www.apec.org>
- United Nations, Department of Economic and Social Affairs, and Population Division. *World Population Prospects: The 2012 Revision*. 2013: DVD Edition.
- Livingstone FB, Sphuler JN. Cultural determinants of natural selection. *United Nations Educational, Scientific and Cultural Organization: Expert Meeting on the Biological Aspect of Race*; Moscow, 1964:1–3.
- Post RH. Deformed nasal septa and relaxed selection. *Eugen Q* 1966;13:101–12.
- Tuomilehto J. The emerging global epidemic of type 1 diabetes. *Curr Diab Rep* 2013;13:795–804.
- Saniotis A, Henneberg M. Medicine could be constructing human bodies in the future. *Med Hypotheses* 2011;77:560–4.
- Moroni L, Bianchi I, Lleo A. Geoepidemiology, gender and autoimmune disease. *Autoimmun Rev* 2012;11:A386–92.
- Dabelea D, Bell RA, D'Agostino RB Jr, et al. The Writing Group for the SEARCH for Diabetes in Youth Study Group. Incidence of diabetes in youth in the United States. *JAMA* 2007;297:2716–25.
- King KM. A history of insulin: from discovery to modern alternatives. *Br J Nurs* 2003;12:1137–41.
- Medawar PB. Do advances in medicine lead to genetic deterioration? In: Bajema CJ, ed. *Natural selection in human populations*. New York: Robert E. Krieger Publishing Co, 1971:300–8.
- Onkamo P, Väänänen S, Karvonen M, et al. Worldwide increase in incidence of Type I diabetes—the analysis of the data on published incidence trends. *Diabetologia* 1999;42:1395–403.
- Kawasaki E, Eguchi K. Is type 1 diabetes in the Japanese population the same as among Caucasians? *Ann N Y Acad Sci* 2004;1037:96–103.
- Suleyman F. Landmarks in diabetes care: a historical perspective. *Community Nurse* 1998;4:13–16.
- Ogle G, Middlehurst A, Short-Hobbs R, et al. *Children and diabetes: success and challenge in the developing world*. 2013. <http://www.idf.org>
- Australian Institute of Health and Welfare. *Prevalence of type 1 diabetes in Australian children, 2008. Diabetes series Number15. Cat. no. CVD 54*. Canberra: AIHW, 2011.
- Neville SE, Boye KS, Montgomery WS, et al. Diabetes in Japan: a review of disease burden and approaches to treatment. *Diabetes Metab Res Rev* 2009;25:705–16.
- Miller RG, Secrest AM, Sharma RK, et al. Improvements in the life expectancy of type 1 diabetes: the Pittsburgh epidemiology of diabetes complications study cohort. *Diabetes* 2012;61:2987–92.
- Chiang JL, Kirkman MS, Laffel LM, et al. Type 1 diabetes through the life span: a position statement of the American Diabetes Association. *Diabetes Care* 2014;37:2034–54.



# Type 1 diabetes prevalence increasing globally and regionally: the role of natural selection and life expectancy at birth

Wen-Peng You and Maciej Henneberg

*BMJ Open Diab Res Care* 2016 4:  
doi: 10.1136/bmjdr-2015-000161

---

Updated information and services can be found at:  
<http://drc.bmj.com/content/4/1/e000161>

---

*These include:*

## References

This article cites 32 articles, 7 of which you can access for free at:  
<http://drc.bmj.com/content/4/1/e000161#BIBL>

## Open Access

This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

## Email alerting service

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

---

## Topic Collections

Articles on similar topics can be found in the following collections  
[Epidemiology/health services research](#) (36)

---

## Notes

---

To request permissions go to:  
<http://group.bmj.com/group/rights-licensing/permissions>

To order reprints go to:  
<http://journals.bmj.com/cgi/reprintform>

To subscribe to BMJ go to:  
<http://group.bmj.com/subscribe/>