

Validation of the Danish National Diabetes Register

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Abstract: The Danish National Diabetes Register (NDR) was established in 2006 and builds on data from Danish health registers. We validated the content of NDR, using full information from the Danish National Patient Register and data from the literature. Our study indicates that the completeness in NDR is $\geq 95\%$ concerning ascertainment from data sources specific for diabetes, ie, prescriptions with antidiabetic drugs and diagnoses of diabetes in the National Patient Register. Since the NDR algorithm ignores diabetes-related hospital contacts terminated before 1990, the establishment of the date of inclusion is systematically delayed for $\geq 10\%$ of the registrants in general and for $\geq 30\%$ of the inclusions before 1997 in particular. This bias is enhanced for ascertainment by chiropody services and by frequent measurements of blood glucose because the date of reimbursement of services, rather than the date of encounter, has been taken as the date of inclusion in NDR. We also find that some 20% of the registrations in NDR may represent false positive inclusions of persons with frequent measurements of blood glucose without having diabetes. We conclude that NDR is a novel initiative to support research in the epidemiological and public health aspects of diabetes in Denmark, but we also present a list of recommended changes for improving validity, by reducing the impact of current sources of bias and misclassifications.

Keywords: diabetes mellitus, epidemiology, ascertainment, validity

Introduction

Like the other Nordic countries, Denmark offers unique opportunities for register-based research in epidemiology and health care.^{1,2} Denmark has a particularly strong tradition of epidemiological research in diabetes,³ which has been further enhanced with the establishment of the Danish National Diabetes Register (NDR) and the utilization of NDR for research purposes.^{4,5}

We have previously used Danish health registers and population-based samples for studies of the epidemiological and public health aspects of diabetes.^{6,7} The establishment of NDR represents a new source for access to individual patient data that we are currently making use of, within the framework of a project called the Diabetes Impact Study 2013, to update our previous studies.

A study that used supplementary methods to ascertain persons with diabetes was performed recently in parts of the former county of Vejle and found that clinical characteristics and mortality differed according to the ascertainment methods applied.⁸ We present here a comprehensive validation analysis of NDR, as part of the Diabetes Impact Study 2013.

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Materials and methods

Sources and content of NDR

NDR was established by the Danish National Board of Health in 2006 to provide a resource for monitoring the clinical course in patients with diabetes.⁴ NDR is built upon information from already existing Danish health registers, including the Danish National Patient Register (DNPaR),⁹ the Danish National Prescription Registry,¹⁰ and the Danish National Health Service Register¹¹ as well as the Danish Civil Registration System (DCRS).¹² In addition, NDR extracts information from DCRS on residence (municipality code) for each registrant as well as recorded survival status. Linkage of person-specific data between the registers is possible using the person identification number (PIN) that is assigned to each Danish citizen at birth or when granted Danish citizenship. For persons without Danish citizenship but with a permission to stay in Denmark, a temporary PIN is assigned. The PIN is unique and used for administrative purposes throughout the public and private sectors in Denmark, and is saved in the central health registers even after the death of the citizen.

Table 1 provides an overview, based on the account of Carstensen et al,⁴ of the main content of NDR, together with the data sources. The date that a given person is qualified for inclusion in NDR is known for each of the

data sources (Table 1). The date of inclusion in NDR for a given person occurs when the first of any qualifying events (Table 1) for that person is fulfilled.

It should be noted that according to the detailed documentation, NDR ignores all hospital contacts terminated before 1990 in the search for diabetes-related activities in the DNPaR.¹³ Overall, the algorithms employed are assumed to identify persons with diabetes with a high level of probability and to reduce false positive inclusion of young women with polycystic ovarian syndrome and women with gestational diabetes. NDR is supposed to include all persons with inclusion date since the start of 1997, as well as all persons who were alive at the end of 1996 and identified in the health registers with an inclusion date before 1997.

NDR is operated by the Statens Serum Institut and is updated on an annual basis, most recently by the inclusion of data for the year 2012. The content of NDR is available for research, provided permission is obtained from the Danish Data Protection Agency and the Danish National Board of Health.

Supplementary data

The present analysis is based on the content of NDR, as extracted on July 3, 2013, containing the data on all inclusions in NDR until 2012. For all registered persons (n=497,232), data were extracted on all contacts (regardless of date) registered in DNPaR, containing information on all interventions performed, primary and secondary diagnoses underlying the contact, hospitalization, and all relevant dates.⁹ The supplementary material presents a list of all diagnoses (according to International Classification of Disease [ICD]8 and ICD10 classifications) and intervention codes (according to the official Danish classification of codes in healthcare) used for this extraction. It should be noted that the codes for diabetes related to pregnancy (ICD10 code DO24) were not used for the purpose of obtaining supplementary data. In addition, all available data on moves, migrations, and deaths recorded on the population of NDR registrants were obtained from DCRS.¹²

Methods of analysis

All inclusions in NDR were described by ascertainment source. For this purpose, ascertainment by means of a diagnosis of diabetes in DNPaR was labeled “DMdiag”. Ascertainment by frequency of blood glucose measurements (“blod2i5” and “blod5i1”, respectively) was grouped to the ascertainment category “BSfreq” (Table 1). Similarly, ascertainment by means of purchased antidiabetic drugs

Table 1 Source registers and content of NDR

Source register	Item in NDR (label)	Inclusion algorithm
DNPaR	lpr (“DMdiag”)	Registration in DNPaR with a diagnosis of diabetes, defined as ICD-10: DE10-14, DH36.0, DO24 (excluding DO24.4), and ICD-8 (prior to 1994): 249, 250
DNPrR	ins (“AntiDiab”)	Purchase of insulin recorded at least twice in DNPrR ^a
	oad (“AntiDiab”)	Purchase of oad at least twice recorded in DNPrR. ^a An exception is females aged 20–39 prescribed metformin exclusively
DNHSR	blod5i1 (“BSfreq”)	Five blood glucose measurements in a 1-year period in DNHSR
	blod2i5 (“BSfreq”)	Two blood glucose measurements per year in 5 consecutive years in DNHSR
	fodt (“Chiro”)	Registration of chiropody (as a service for a diabetic patient) in DNHSR

Note: ^aInclusion criteria only fulfilled at purchase of second prescription of the drug.
Abbreviations: AntiDiab, antidiabetic drug; BSfreq, frequency of blood sugar measurement; Chiro, chiropody; DMdiag, diagnoses of diabetes; DNHSR, Danish National Health Service Register; DNPaR, Danish National Patient Register; DNPrR, Danish National Prescription Registry; ICD, International Classification of Diseases; ins, insulin; NDR, National Diabetes Register; oad, oral antidiabetic drug.

(“ins” and “oad”) was grouped to the ascertainment category “AntiDiab”. Ascertainment by receiving a service as a person with diabetes by a registered chiroprapist was categorized as “Chiro”.

Using the access to the full information from DNPaR, we identified for each registrant in NDR, all hospital contacts where diabetes formed part of the primary diagnosis or where interventions performed during the contact were specific for diabetes. The supplementary material contains the list of diagnostic codes (ICD8 classification for contacts before 1995; ICD10 classification for contacts for 2005 and forward) and the procedure codes specific for intervention in diabetes as used in all Danish hospitals. If such contacts were later than the inclusion date registered in NDR, the inclusion date in NDR was set operationally as the date of diagnosis of diabetes. If such contacts were earlier than the inclusion date registered in NDR, the earliest admission date was set operationally as the date of diagnosis.

Secondary ascertainment for a given patient occurred by the fulfillment of inclusion criteria by any other data sources than the first one. For each ascertainment, source methods for survival analysis¹⁴ were used to estimate the cumulative

probability of being ascertained by the first occurring secondary source. Patients were censored if death occurred before experiencing secondary ascertainment.

All data were analyzed using anonymized PINs. Since this was a purely descriptive study without any prior specified hypotheses to be tested, only descriptive analyses were performed.

Ethical aspects

The Diabetes Impact Study 2013 is observational, with no contact to registered persons. Permission to extract and analyze data was obtained from the Danish Data Protection Agency (permit number Jnr 2013-21-1749) and the Danish National Board of Health (permit number FSEID-00000440).

Results

Ascertainment by source in NDR

Figure 1 provides an overview of the population of persons registered in NDR. In total, 688 persons (0.1%) have been excluded for various reasons. In particular, 530 persons with inclusion after death represent persons who died

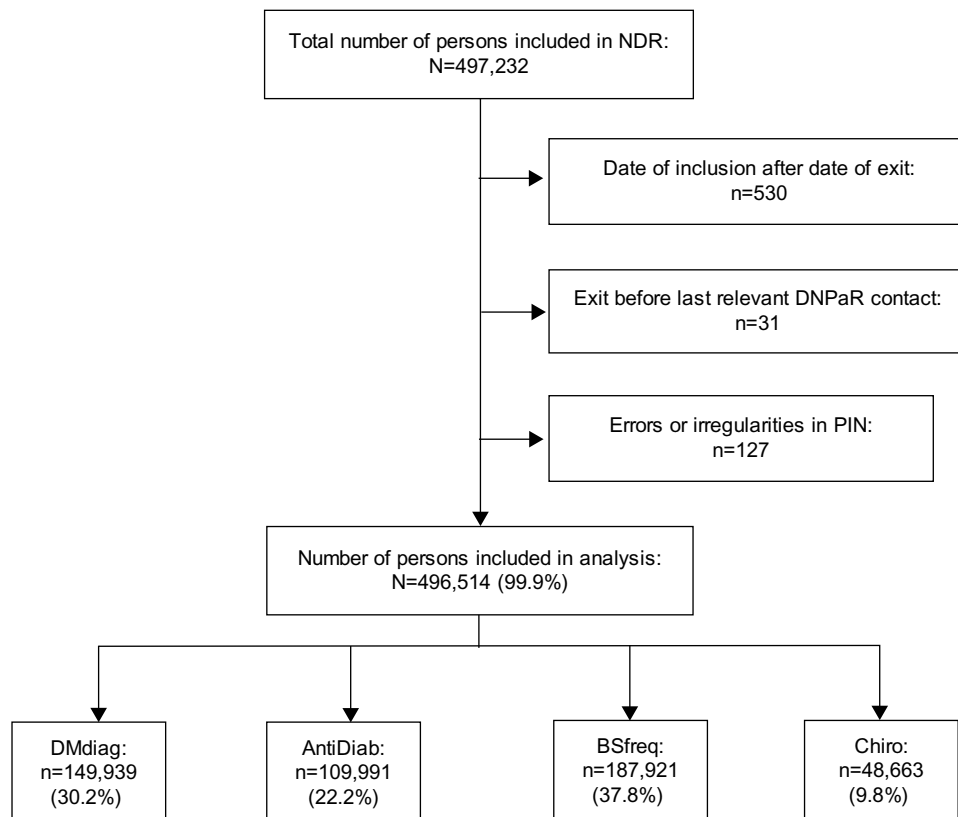


Figure 1 Flowchart of the population of persons registered as of July 3, 2013 in NDR.

Abbreviations: AntiDiab, antidiabetic drug; BSfreq, frequency of blood sugar measurement; Chiro, chiroprapist; DMdiag, diagnoses of diabetes; DNPaR, Danish National Patient Register; NDR, National Diabetes Register; PIN, personal identification number.

before inclusion. Further, 31 persons registered with a relevant contact in DNPaR after exit represent persons with date of emigration registered in DCRS before the event registered in DNPaR. Almost 40% of registrants are ascertained by “BSfreq” as the primary source, whereas less than 10% of the registrants are ascertained by “Chiro” as the primary source.

Trends over time in inclusion by primary ascertainment source are illustrated from 1997 and onward in Figure 2. The annual number of inclusions, representing registration incidence, has increased steadily, from about 15,000 to more than 30,000, over the period 1997 through 2011. A relatively large increase has been seen for primary ascertainment by “AntiDiab”, while the contribution from “DMdiag” has been rather constant and that for “BSfreq” has been modestly increasing. The number of inclusions by “Chiro” as the primary source is limited but shows relative increases during 2002–2004 and again for the year 2011.

Secondary ascertainment in NDR

Inclusion in NDR may be qualified in several ways, and registrants may have been ascertained subsequently by other sources than by the primary ascertainment source with the earliest qualification fulfilled. Figure 3 shows the cumulative probability of ascertainment by at least one additional source, by follow-up time from the date of primary ascertainment. This analysis uses inclusions from 1997 throughout 2011. The patterns of cumulative probability vary by primary ascertainment source. For primary ascertainment by both

“DMdiag” and “AntiDiab”, some 80% of the persons will be ascertained by at least one additional source within a period of 10 years, however, with a sharper initial rise for persons primarily ascertained by “DMdiag”. Primary ascertainment by “Chiro” is associated with a cumulative probability at a lower level and with a more slow increase. For persons primarily ascertained by “BSfreq”, only about 50% will be ascertained by a second source and with only a modest initial sharp increase in the probability curve.

Operationally defined date of diagnosis of diabetes

Correspondence between the date of inclusion in NDR and the operationally defined date of diagnosis of diabetes, when using the full historical information in DNPaR, is shown in Table 2, grouped by period of inclusion in NDR. Overall, the operational date of diagnosis equals the date of inclusion in NDR for almost 89% of persons. However, the percentage with agreement is only about 68% for persons with inclusion before 1997 against about 98% for persons with inclusion from 1997 and onward. The proportion of persons with more than 10 years of difference from operationally defined date of diagnosis until inclusion in NDR is about 10% for inclusion before 1997 against 0.4% for inclusion from 1997 and onward.

Discussion

NDR is a nationwide register that provides a unique platform for comprehensive research in the epidemiology and public health aspects of diabetes. As a main advantage, NDR makes

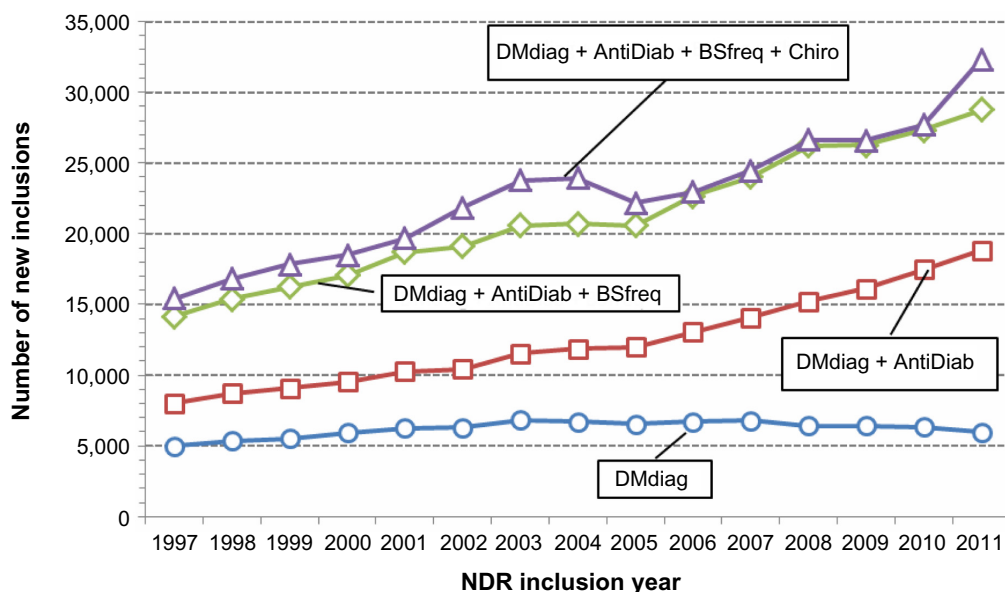


Figure 2 Ascertainment in NDR by source and calendar time, for inclusions in NDR from 1997 through 2011.

Abbreviations: AntiDiab, antidiabetic drug; BSfreq, frequency of blood sugar measurement; Chiro, chiropody; DMdiag, diagnoses of diabetes; NDR, National Diabetes Register.

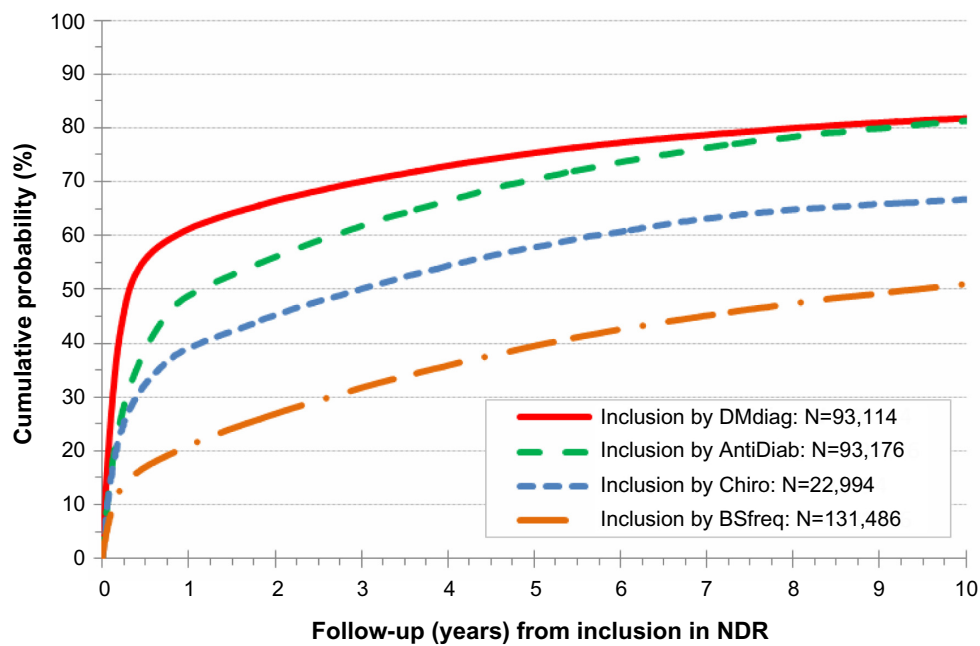


Figure 3 Secondary ascertainment in NDR by primary ascertainment source, for inclusions in NDR from 1997 through 2011.

Abbreviations: AntiDiab, antidiabetic drug; BSfreq, frequency of blood sugar measurement; Chiro, chiropody; DMdiag, diagnoses of diabetes; NDR, National Diabetes Register.

use of already existing data that are routinely collected and maintained at the national level. As a further advantage, NDR attempts to cover both the secondary and the primary health care sector. In the attempt to validate important elements of NDR, the present study has identified several areas of concern about the validity of NDR, some of which may be alleviated by simple measures.

Validity of ascertainment in NDR

Ascertainment by purchase of antidiabetic drugs and by a primary hospital discharge diagnosis of diabetes as well as by diabetes-related chiropody may be characterized as specific for diabetes. However, the present list of diagnoses used for the identification of diabetes-related contacts in DNPaR is not complete and should be supplemented with diagnostic codes for diabetic eye complications. Furthermore, additional relevant contacts may be identified by codes of

interventions specific for treatment of diabetes not included in the present NDR algorithm. Such codes may be identified by comparing the codes specified in Table 1 with the codes listed in the supplementary material. The use of such supplementary codes may not increase the ascertainment in NDR substantially but may optimize the establishment of date of inclusion in NDR.

The inclusion of a substantial proportion of persons by means of the frequency of blood sugar measurements in the primary health care sector gives rise to concern about the validity of ascertainment, particularly because currently it is not possible to access information on the results of blood glucose measurements at the national level. Even though validation performed prior to the establishment of NDR indicated a high predictive value of diabetes for the “BSfreq” criteria currently used by NDR,⁴ increased awareness in diabetes may have changed clinical practice and thereby changed

Table 2 Delay (in years) between date of inclusion in NDR and the operational date of diagnosis

Delay (in years)	NDR inclusion before 1997		NDR inclusion 1997–2011 (incl)		Total	
	Number	%	Number	%	Number	%
No delay	106,142	68.2	335,306	98.4	441,448	88.9
0 < delay < 1	2,551	1.6	2,489	0.7	5,040	1.0
1 ≤ delay < 5	13,860	8.9	1,140	0.3	15,000	3.0
5 ≤ delay < 10	17,470	11.2	383	0.1	17,853	3.6
10 ≤ delay	15,717	10.1	1,456	0.4	17,173	3.5
All persons	155,740	100.0	340,774	100.0	496,514	100.0

Abbreviation: NDR, National Diabetes Register.

the predictive performance of the “BSfreq” algorithms. Our results show that the segment of registrants ascertained by “BSfreq” has a substantially lower probability, at about 50%, of being ascertained subsequently by a secondary source. This finding suggests that the group of persons ascertained exclusively by “BSfreq”, to some extent, contains persons with blood sugar examined without having clinically diagnosed diabetes. According to a “worst-case” scenario, 50% of those 40% of registrants that are ascertained by “BSfreq” do not have diabetes; this corresponds to about 20% of the total population registered in NDR.

In a recent study, a regional algorithm based on information on glycosylated hemoglobin (HbA_{1c}) measurements as a supplementary ascertainment source for the identification of persons with diabetes was compared with the algorithm used by NDR.⁸ Based on an analysis of ascertainment patterns in about 14,000 persons ascertained jointly by the two algorithms, it was estimated that about 21% of persons ascertained by means of “BSfreq” in NDR may not have diagnosed diabetes but have their blood glucose examined frequently for other reasons. This estimate is close to the one presented above, which was obtained by a completely different approach. Furthermore, almost 700 persons were identified by the regional algorithm as having at least one elevated measurement of HbA_{1c} diagnostic of diabetes but without being registered with NDR.⁸ Taking these findings as representing ascertainment failure in NDR, a proportionate analysis suggests the estimated completeness of diagnosed diabetes in NDR to be 93%–95%. Although a full systematic analysis of the completeness of ascertainment has been difficult because of differences in the definitions of the underlying patient populations, it is possible to perform formal analysis for specific segments. Thus, applying capture–recapture methods of analysis to the data in Table 1 of Nielsen et al⁸ for ascertainment by “AntiDiab” and “DMdiag”, the completeness of ascertainment in NDR is estimated at 95.8% (95% confidence interval [CI] 95.4%–96.3%) and 97.8% (95% CI 97.4%–98.6%), respectively.

Whereas the completeness of ascertained diagnosed diabetes in NDR may be 95% or higher, a substantial proportion, possibly amounting to about 20%, of the persons registered with NDR may have been falsely included by means of frequent measurement of blood glucose. In June 2012, the Danish National Board of Health decided to accept an HbA_{1c} level above 6.5% as a diagnostic criterion for diabetes. If readings of HbA_{1c} measurements will be accessible at the national level in the future, it should be considered to revise the algorithm for inclusion in NDR accordingly.

In terms of a full assessment of the performance of the algorithm of inclusion in NDR, the completeness of ascertainment is interpretable as the sensitivity, which we estimate to be 95% or higher. If further assumed that some 20% of the registrants in NDR do not have diabetes, the positive predictive value of ascertainment is about 80%. Applying a sensitivity at 95% and a positive predictive value at 80% in a scenario with a total population like the Danish, with some six million inhabitants and an annual number of new inclusions in NDR at 30,000, yields an estimated specificity (ie, the proportion of the total population without diabetes and that not ascertained by NDR) at 99.9%.

Date of inclusion in NDR versus operationally defined date of diagnosis of diabetes

Since the date of clinical diagnosis of diabetes cannot be recorded in NDR, it is necessary to use the date of inclusion as a proxy. This must be done with due reservation because in most cases, the clinical diagnosis may have been established before inclusion in NDR.⁴ The impact of this form of bias may be reduced by searching for the earliest possible registration of diabetes-related contacts in the source registers, notably DNPdR. As per the current NDR algorithm, all hospital contacts closed before 1990 are ignored.¹³ The reasons for this are not obvious, but the ignorance will enhance bias when using the date of inclusion in NDR as a proxy for the date of diagnosis of diabetes. Since our study takes account of all hospital activities, regardless of time, we can show that by ignoring contacts in DNPdR before 1990, more than 10% of the registrants have a delay in the operationally defined date of diagnosis. The delay affects more than 30% of the registrants with inclusion date before 1997 and is substantial, representing 10 years or even more for about 10% of the persons included in NDR before 1997. While this may not affect the completeness of ascertainment in NDR, it introduces a bias, particularly in investigations that involve the study of diabetes duration and the impact of diabetes duration over time.

Our study also has identified that persons may be included in NDR after death. To explore on this finding, we investigated further the timing of inclusion in NDR. It appears that for patients primarily ascertained by “DMdiag” and by “AntiDiab”, the inclusion dates seem distributed by random, albeit with relatively fewer, inclusions during weekends. In contrast, primary ascertainment by “Chiro” is systematically associated with inclusion on Wednesdays, whereas primary ascertainment by “BSfreq” is associated

with inclusion also on Wednesday, however with possible spacing between Wednesdays in intervals of up to 4 weeks or even more. Thus, it is likely that for inclusion by methods using contacts with health care providers in the primary health care sector, the inclusion date in NDR represents the date of reimbursement from the National Health Service to the health care provider, rather than the date of actual provision of service. Accordingly, a limited number of patients have died in the interval between the date of contact and the date of reimbursement. This element of the NDR algorithm enhances the bias when using date of inclusion in NDR as a proxy for date of diagnosis.

The present study has demonstrated how administrative changes may influence inclusion in NDR. The number of inclusions by “Chiro” as the primary source is limited but shows a distinct peak for the period 2002–2004 and again for the year 2011. These peaks coincide with periods during which agreements between the chiropractors and the National Health Service were in effect concerning reimbursement of expenses of chiropractic for diabetic patients. In fact, the most recent summary report from NDR¹⁵ shows a relative decrease in the number of new inclusions for the year 2012, which may be interpreted as a compensation for the relatively steep increase in 2011. Such fluctuations must be interpreted with appropriate caution to the extent that the inclusion rate in NDR is used as a proxy for the incidence rate of diabetes.

Conclusion and recommendations

NDR represents a novel initiative by which unique opportunities have been created for the monitoring of the basic epidemiological characteristics of diabetes as well as for further in-depth studies on the epidemiological and public health aspects of diabetes in Denmark. Comparisons with other samples of diabetic patients have suggested that NDR may perform with a high level of completeness of ascertainment.⁸ Even though, our study has identified a range of issues that may question other aspects of the validity of NDR, with corresponding consequences for the use of NDR for research purposes. Some of these issues may be resolved by relatively simple changes in the NDR algorithm. Specifically, we recommend the following: 1) In the search for qualifying contacts in DNPaR, the NDR algorithm should include the content of the total DNPaR, rather than being limited to contacts after 1990. This simple measure will reduce substantially the bias in the operationally defined start of the diabetes course that preferentially exists for persons with inclusion date before 1997. 2) The set of diagnostic codes used in the NDR should be reviewed carefully and updated on a periodic basis with

current and new relevant diagnostic codes and intervention codes specific for diabetes, as suggested in the supplementary material of this paper. While it may have limited impact only concerning the number of persons identified, such an improvement facilitates the establishment of a more qualified date of inclusion. 3) For inclusion by means of “BSfreq” and “Chiro”, the true date of encounter, if available, in the Danish National Health Service Register should be used, rather than using the date of reimbursement. This will further support the establishment of a more qualified date of inclusion, and at the same time, the strikingly invalid feature of including persons after their death will be avoided. 4) As soon as results of HbA_{1c} measurements may become available as part of a future nationwide Danish laboratory database, the NDR algorithm should switch to rely upon the identification of measurements of elevated HbA_{1c}, instead of the frequency of blood glucose measurements. Such a switch will significantly reduce the false inclusion in NDR of persons with frequent blood glucose measurements for other clinical reasons than diabetes control. 5) Until the NDR algorithm has been further improved according to the recommendations presented here, we suggest performing sensitivity analyses with respect to the segment of persons ascertained by “BSfreq”, in studies that make use of the content of NDR.

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Disclosure

The authors report no conflicts of interest in this work.

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Supplementary material

Table S1 Diagnosis codes and intervention codes used for the identification of diabetes-related hospital contacts

Code	Code category	Code text
24900	ICD8 – disease classification	Insulin-dependent diabetes mellitus, without complications
24901	ICD8 – disease classification	Cataract, retinopathy in insulin-dependent diabetes
24902	ICD8 – disease classification	Diabetic nephropathy, Kimmelstiel–Wilson syndrome, insulin-dependent diabetes
24903	ICD8 – disease classification	Neuropathy, diabetic polyneuritis, insulin-dependent diabetes
24904	ICD8 – disease classification	Diabetic angiopathy in extremities, insulin-dependent diabetes
24905	ICD8 – disease classification	Diabetic gangrene, insulin-dependent diabetes
24906	ICD8 – disease classification	Diabetic coma without ketonuria, insulin-dependent diabetes
24907	ICD8 – disease classification	Diabetic coma (incl precoma), insulin-dependent diabetes
24908	ICD8 – disease classification	Insulin-dependent diabetes mellitus, with complications not otherwise specified
24909	ICD8 – disease classification	Insulin-dependent diabetes mellitus
25000	ICD8 – disease classification	Insulin independent diabetes mellitus, without complications
25001	ICD8 – disease classification	Cataract, retinopathy in insulin independent diabetes
25002	ICD8 – disease classification	Diabetic nephropathy, Kimmelstiel–Wilson syndrome, insulin independent diabetes
25003	ICD8 – disease classification	Neuropathy, diabetic polyneuritis, insulin independent diabetes
25004	ICD8 – disease classification	Diabetic angiopathy in extremities, insulin independent diabetes
25005	ICD8 – disease classification	Diabetic gangrene, insulin independent diabetes
25006	ICD8 – disease classification	Diabetic coma without ketonuria, insulin independent diabetes
25007	ICD8 – disease classification	Diabetic coma (incl precoma), insulin independent diabetes
25008	ICD8 – disease classification	Insulin independent diabetes mellitus, with complications not otherwise specified
25009	ICD8 – disease classification	Insulin independent diabetes mellitus
BBHA	National Intervention Classification	Peroral diabetes treatment
BBHA0	National Intervention Classification	Treatment with peroral antidiabetic agents
BBHA00	National Intervention Classification	Treatment with agents stimulating insulin production
BBHA01	National Intervention Classification	Treatment with glucose-lowering antidiabetic agents
BBHF0	National Intervention Classification	Insulin treatment
BBHF00	National Intervention Classification	Insulin treatment, mono- or dual therapy
BBHF01	National Intervention Classification	Insulin treatment, basal and bolus therapy
BBHF02	National Intervention Classification	Insulin pump treatment
BBKB	National Intervention Classification	Instruction in injecting insulin
BUBE0	National Intervention Classification	Diabetes diet
BUBE00	National Intervention Classification	Diabetes diet, type 1 diabetes
BUBE01	National Intervention Classification	Diabetes diet, type 2 diabetes
BUFA	National Intervention Classification	Dietary instruction in relation to diabetes
BUFA0	National Intervention Classification	Dietary instruction in relation to type 1 diabetes
BUFA00	National Intervention Classification	Initial dietary instruction in relation to type 1 diabetes
BUFA01	National Intervention Classification	Follow-up dietary instruction in relation to type 1 diabetes
BUFA1	National Intervention Classification	Dietary instruction in relation to type 2 diabetes
BUFA10	National Intervention Classification	Initial dietary instruction in relation to type 2 diabetes
BUFA11	National Intervention Classification	Follow-up dietary instruction in relation to type 2 diabetes
BUFA3	National Intervention Classification	Dietary instruction in relation to pregnant women with type 1 diabetes
BUFA30	National Intervention Classification	Initial dietary instruction in relation to pregnant women with type 1 diabetes
BUFA31	National Intervention Classification	Follow-up dietary instruction in relation to pregnant women with type 1 diabetes
DE10	ICD10 – disease classification	Type 1 diabetes
DE100	ICD10 – disease classification	Type 1 diabetes with coma
DE100A	ICD10 – disease classification	Diabetic coma, hyperosmolar in IDDM
DE100B	ICD10 – disease classification	Diabetic coma in IDDM with ketoacidosis
DE100C	ICD10 – disease classification	Diabetic coma in IDDM without ketoacidosis
DE100D	ICD10 – disease classification	Diabetic coma, hyperglycemic in IDDM
DE100E	ICD10 – disease classification	Diabetic coma, hypoglycemic in IDDM
DE100F	ICD10 – disease classification	Insulin-dependent diabetes with diabetic coma
DE101	ICD10 – disease classification	Type 1 diabetes with ketoacidosis
DE102	ICD10 – disease classification	Type 1 diabetes with renal complication
DE103	ICD10 – disease classification	Type 1 diabetes with eye complication
DE104	ICD10 – disease classification	Type 1 diabetes with neurological complication

(Continued)

Table S1 (Continued)

Code	Code category	Code text
DE105	ICD10 – disease classification	Type 1 diabetes with complications in peripheral vascular system
DE105A	ICD10 – disease classification	Type 1 diabetes with peripheral angiopathy
DE105B	ICD10 – disease classification	Type 1 diabetes with foot ulcer
DE105C	ICD10 – disease classification	Type 1 diabetes with gangrene
DE105D	ICD10 – disease classification	Type 1 diabetes with microangiopathy
DE106	ICD10 – disease classification	Type 1 diabetes with other complication
DE107	ICD10 – disease classification	Type 1 diabetes with multiple complications
DE108	ICD10 – disease classification	Type 1 diabetes with complication UNS
DE109	ICD10 – disease classification	Type 1 diabetes without complications
DE109A	ICD10 – disease classification	Type 1 diabetes UNS
DE11	ICD10 – disease classification	Type 2 diabetes
DE110	ICD10 – disease classification	Type 2 diabetes with coma
DE110A	ICD10 – disease classification	Diabetic coma in NIDDM without ketoacidosis
DE110B	ICD10 – disease classification	Diabetic coma, hypoglycemic in NIDDM
DE110C	ICD10 – disease classification	Diabetic coma, hyperosmolar in NIDDM
DE110D	ICD10 – disease classification	Diabetic coma, hyperglycemic in NIDDM
DE110E	ICD10 – disease classification	Diabetic coma in NIDDM with ketoacidosis
DE111	ICD10 – disease classification	Type 2 diabetes with ketoacidosis
DE112	ICD10 – disease classification	Type 2 diabetes with renal complication
DE113	ICD10 – disease classification	Type 2 diabetes with eye complication
DE114	ICD10 – disease classification	Type 2 diabetes with neurological complication
DE115	ICD10 – disease classification	Type 2 diabetes with complications in peripheral vascular system
DE115A	ICD10 – disease classification	Type 2 diabetes with peripheral angiopathy
DE115B	ICD10 – disease classification	Type 2 diabetes with foot ulcer
DE115C	ICD10 – disease classification	Type 2 diabetes with gangrene
DE115D	ICD10 – disease classification	Type 2 diabetes with microangiopathy
DE116	ICD10 – disease classification	Type 2 diabetes with other complication
DE117	ICD10 – disease classification	Type 2 diabetes with multiple complications
DE118	ICD10 – disease classification	Type 2 diabetes with complication UNS
DE119	ICD10 – disease classification	Type 2 diabetes without complications
DE119A	ICD10 – disease classification	Type 2 diabetes UNS
DE12	ICD10 – disease classification	Malnutrition-related diabetes
DE120	ICD10 – disease classification	Malnutrition-related diabetes with coma
DE120A	ICD10 – disease classification	Diabetic coma, hyperglycemic in malnutrition-related diabetes
DE120B	ICD10 – disease classification	Diabetic coma, hypoglycemic in malnutrition-related diabetes
DE120C	ICD10 – disease classification	Diabetic coma in malnutrition-related diabetes with ketoacidosis
DE120D	ICD10 – disease classification	Diabetic coma in malnutrition-related diabetes without ketoacidosis
DE120E	ICD10 – disease classification	Diabetic coma, hyperosmolar in malnutrition-related diabetes
DE121	ICD10 – disease classification	Malnutrition-related diabetes with ketoacidosis
DE122	ICD10 – disease classification	Malnutrition-related diabetes with renal complication
DE123	ICD10 – disease classification	Malnutrition-related diabetes with eye complication
DE124	ICD10 – disease classification	Malnutrition-related diabetes with neurological complication
DE125	ICD10 – disease classification	Malnutrition-related diabetes with complications in peripheral vascular system
DE125A	ICD10 – disease classification	Malnutrition-related diabetes with peripheral angiopathy
DE125B	ICD10 – disease classification	Malnutrition-related diabetes with foot ulcer
DE125C	ICD10 – disease classification	Malnutrition-related diabetes with gangrene
DE125D	ICD10 – disease classification	Malnutrition-related diabetes with microangiopathy
DE126	ICD10 – disease classification	Malnutrition-related diabetes with other complication
DE127	ICD10 – disease classification	Malnutrition-related diabetes with multiple complications
DE128	ICD10 – disease classification	Malnutrition-related diabetes with complication UNS
DE129	ICD10 – disease classification	Malnutrition-related diabetes without complications
DE13	ICD10 – disease classification	Other forms of diabetes
DE130	ICD10 – disease classification	Other form of diabetes with coma
DE131	ICD10 – disease classification	Other form of diabetes with ketoacidosis
DE132	ICD10 – disease classification	Other form of diabetes with renal complication
DE133	ICD10 – disease classification	Other form of diabetes with eye complication
DE134	ICD10 – disease classification	Other form of diabetes with neurological complication

(Continued)

Table S1 (Continued)

Code	Code category	Code text
DE135	ICD10 – disease classification	Other form of diabetes with complications in peripheral vascular system
DE135A	ICD10 – disease classification	Other form of diabetes with peripheral angiopathy
DE135B	ICD10 – disease classification	Other form of diabetes with foot ulcer
DE135C	ICD10 – disease classification	Other form of diabetes with gangrene
DE135D	ICD10 – disease classification	Other form of diabetes with microangiopathy
DE136	ICD10 – disease classification	Other form of diabetes with other complication
DE137	ICD10 – disease classification	Other form of diabetes with multiple complications
DE138	ICD10 – disease classification	Other form of diabetes with complication UNS
DE139	ICD10 – disease classification	Other form of diabetes without complications
DE14	ICD10 – disease classification	Unspecified diabetes
DE140	ICD10 – disease classification	Diabetes UNS with coma
DE140A	ICD10 – disease classification	Diabetic coma in malnutrition-related diabetes
DE140B	ICD10 – disease classification	Diabetic coma, hyperglycemic in unspecified diabetes
DE140C	ICD10 – disease classification	Diabetic coma, hyperosmolar in unspecified diabetes
DE140D	ICD10 – disease classification	Diabetic coma, hypoglycemic in unspecified diabetes
DE141	ICD10 – disease classification	Diabetes UNS with ketoacidosis
DE142	ICD10 – disease classification	Diabetes UNS with renal complication
DE143	ICD10 – disease classification	Diabetes UNS with eye complication
DE144	ICD10 – disease classification	Diabetes UNS with neurological complication
DE145	ICD10 – disease classification	Diabetes UNS with complications in peripheral vascular system
DE145A	ICD10 – disease classification	Diabetes UNS with peripheral angiopathy
DE145B	ICD10 – disease classification	Diabetes UNS with foot ulcer
DE145C	ICD10 – disease classification	Diabetes UNS with gangrene
DE145D	ICD10 – disease classification	Diabetes UNS with microangiopathy
DE146	ICD10 – disease classification	Diabetes UNS with other complication
DE147	ICD10 – disease classification	Diabetes UNS with multiple complications
DE148	ICD10 – disease classification	Diabetes UNS with complication UNS
DE149	ICD10 – disease classification	Diabetes UNS without complications
DE159	ICD10 – disease classification	Hypoglycemic coma UNS
DE159A	ICD10 – disease classification	Insulin shock
DG632	ICD10 – disease classification	Diabetic polyneuropathy
DH280	ICD10 – disease classification	Diabetic cataract
DH360	ICD10 – disease classification	Diabetic retinopathy UNS
DH360A	ICD10 – disease classification	Simplex retinopathy IDDM
DH360B	ICD10 – disease classification	Proliferative retinopathy IDDM
DH360C	ICD10 – disease classification	Simplex retinopathy NIDDM
DH360D	ICD10 – disease classification	Proliferative retinopathy NIDDM
DH360E	ICD10 – disease classification	Diabetic maculopathy IDDM
DH360F	ICD10 – disease classification	Diabetic maculopathy NIDDM
DH360H	ICD10 – disease classification	Simplex diabetic retinopathy
DH360J	ICD10 – disease classification	Proliferative diabetic retinopathy
DH360K	ICD10 – disease classification	Diabetic maculopathy
DZ135AI	ICD10 – disease classification	Screening for diabetic retinopathy

Abbreviations: ICD, International Classification of Disease; IDDM, insulin-dependent diabetes mellitus; NIDDM, non-insulin-dependent diabetes mellitus; UNS, unspecified.

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