The impact of consanguinity on the frequency of inborn errors of metabolism

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ABSTRACT
Inborn errors of metabolism (IEM) are a heterogeneous group of genetic disorders present in all ethnic groups. We investigated the frequency of consanguinity among parents of newborns with IEM diagnosed by neonatal screening.

Data were obtained from 15 years of expanded newborn screening for selected IEM with autosomal recessive mode of inheritance, a national screening program of newborns covering the period from 2002 until April 2017. Among the 838,675 newborns from Denmark, the Faroe Islands and Greenland, a total of 196 newborns had an IEM of whom 155 from Denmark were included in this study. These results were crosschecked against medical records. Information on consanguinity was extracted from medical records and telephone contact with the families.

Among ethnic Danes, two cases of consanguinity were identified in 93 families (2.15%). Among ethnic minorities there were 20 cases of consanguinity among 33 families (60.6%). Consequently, consanguinity was 28.2 times more frequent among descendants of other geographic place of origin than Denmark. The frequency of consanguinity was conspicuously high among children of Pakistani, Afghan, Turkish and Arab origin (71.4%). The overall frequency of IEM was 25.5 times higher among children of Pakistani, Turkish, Afghan and Arab origin compared to ethnic Danish children (5.35:10,000 v 0.21:10,000). The frequency of IEM was 30-fold and 50-fold higher among Pakistanis (6.5:10,000) and Afghans (10.6:10,000), respectively, compared to ethnic Danish children.

The data indicate a strong association between consanguinity and IEM. These figures could be useful to health professionals providing antenatal, pediatric, and clinical genetic services.

1. Introduction
Inborn errors of metabolism (IEM) are a heterogeneous group of rare genetic disorders that are an important cause of morbidity and mortality in children and adults. The outcome of many IEM is directly related to how early correct diagnoses are made and appropriate treatment instituted. Most of the IEM are inherited in an autosomal recessive manner, and the majority of cases are due to enzymatic defects arising from aberrations in single genes. Autosomal recessive IEM are characterized by their varying frequencies in different ethnic groups, as a result of natural selection, genetic drift, and large-scale migration from countries where consanguinity is favored [1]. This is apparent in a somewhat ethnically diverse country as Denmark and other Western societies, where migration from countries with tradition for consanguineous unions has been a significant demographic feature [2], thus, having an impact on the patient profile of medical genetics clinics in recipient countries [3,4].

To determine the genetic basis of IEM in Denmark, we undertook a national study and investigated the frequency of IEM among ethnic minorities and the ethnic Danish population and whether there is an association between the frequency of consanguinity and the prevalence of IEM with an autosomal recessive mode of inheritance.

2. Materials and methods

2.1. Study design and patients

This study was retrospectively designed in Department of Clinical Genetics, Rigshospitalet. Based on results from the National Danish expanded neonatal screening for selected IEM that started in 2002 [5], the medical records of all neonates between 2002 to April 2017 with a true positive result for an IEM were reviewed for consanguinity. In
cases where the information stated in the medical records was not adequate, additional information was gathered through telephone contact with the parents of the affected children.

The selection of IEM was based on criteria such as national disease frequencies; availability of effective treatment, severity of disorder; benefit of early treatment; prevention of early death; no clinical signs at birth; detectability by an easy and precise screening method (for further details, see Lund et al.) [5].

Parental consanguinity in our study was defined according to the definition applied in clinical genetics as a union between two individuals related as second cousins or closer [6].

Fig. 1 shows a flow diagram of inclusion of cases. 196 neonates were reported with an IEM. Of these, 41 referred to the department from Greenland and The Faroe Islands were excluded. These are relatively small and homogenous populations, and we wanted to establish, in a population of significant size and ethnic diversity, if there was any significant difference in the frequency of consanguinity between ethnic Danes and ethnic minorities with an IEM with a recessive mode of inheritance. The remaining 155 patients comprised 117 ethnic Danes, including nine sets of sibs, and 38 immigrants (descendants), including three sets of sibs. The selection of IEM was based on criteria such as national disease frequencies; availability of effective treatment, severity of disorder; benefit of early treatment; prevention of early death; no clinical signs at birth; detectability by an easy and precise screening method (for further details, see Lund et al.) [5].

Parental consanguinity in our study was defined according to the definition applied in clinical genetics as a union between two individuals related as second cousins or closer [6].

Fig. 1 shows a flow diagram of steps in analysis. 838,675 newborns (2002-2017) were born to consanguineous parents or not and previous history of genetically affected sibs.

Demographic and statistical data on ethnic groups were gathered from the Danish Statistics Bureau, Statistics Denmark. We collected data on the number of inhabitants in Denmark (Danish citizenship) at April 1, 2017, number of inhabitants in Denmark defined as ethnic Danes, i.e. inhabitants with Denmark as country of origin at April 1, 2017, and number of Pakistani, Turkish, Afghan and Arab (Arab = Jordanians, Somalis, Palestinians) descendants in Denmark with Danish and foreign citizenship at April 1, 2017.

Inclusion criteria were a confirmed diagnosis of an autosomal recessive metabolic disorder, address in Denmark and knowledge of family history (± consanguinity).

2.2. The Danish Civil Registration System (CRS)

The Danish healthcare system is publicly funded. Data from hospitals are gathered in public registries linked with a unique personal identification number assigned to all inhabitants since 1968. The Danish Civil Registration System [7] is an administrative system containing individual information on the unique personal identification number, name, gender, date of birth, citizenship, identity of parents, place of residence, civil status, immigration and/or death. The CRS number is assigned to all persons with residential location in Denmark at birth or immigration.

2.3. Statistical concepts

Immigrants, descendants and country of origin are statistical concepts based on family relations, citizenship and country of birth. Since all the neonates in the ethnic minority groups were born in Denmark they are, according to the classification in Statistics Denmark, defined as descendants [8]. Statistics Denmark provides information on the number of immigrants and their descendants, their age, sex, citizenship, place of birth, country of origin and geographical distribution. The statistics are based on data obtained from the CRS.

2.4. Ethnicity

A number of different indicators can be applied when measuring ethnicity, e.g. country of birth, maternal/paternal country of birth, principal language at home, native language, citizenship of the person in question, and citizenship of the parents. We considered country of birth as an acceptable approximation for ethnicity. We defined ethnicity of the child by the mother's country of birth, similar to the definition used by Statistics Denmark. Thus, all births to women born outside of Denmark were referred to as children of the mother's country of birth, and children born to mothers who themselves were born in Denmark as ethnic Danish.

3. Results

During the 15 years succeeding the introduction of the National Danish expanded neonatal screening for selected IEM, 838,675 children were born in Denmark, Greenland and Faroe Islands. By April 2017, IEM with autosomal recessive mode of inheritance affecting the major biochemical areas of amino acid, organic acid, and carbohydrate metabolism had been diagnosed in 196 children born during this period. Table 1 shows the cases of IEM detected in ethnic Danes and ethnic minorities. All the patients are included, including cases among sibs.

In ethnic Danish population (Table 1) nine cases of disease were found among sibs. Additionally, one patient was a true screen positive for an autosomal recessive metabolic disorder (holocarboxylase synthetase deficiency), but further examination by the use of molecular genetic analysis showed that the patient was functionally a heterozygote, i.e. both mutations were on the same chromosome inherited
from his father. These ten cases were excluded. The remaining 107 patients represented 107 families with autosomal recessive disease.

Among ethnic minorities (Table 1), there were three cases of disease among sibs, which were excluded. The remaining 35 patients represented 35 families with autosomal recessive disease.

Table 1 shows that medium-chain acyl-CoA dehydrogenase deficiency (MCADD) was the most frequently found IEM among ethnic Danes as well as ethnic minorities. Among Danes 58% had MCADD; among ethnic minorities the frequency was 36.8%.

Table 2 shows the frequency of consanguinity in the different ethnic groups. Of the 33 sets of parents from ethnic minorities, about whom information about consanguinity was available, 20 were consanguineous, i.e. 60.6%. Among Danes, two cases of consanguinity were found among 93 families, i.e. 2.15%. The most popular form of consanguineous union was first cousin marriage, where the coefficient of inbreeding (F) equals 0.0625 for first cousin offspring. Among ethnic minorities, the 20 cases of consanguinity identified, one was a double first-cousin marriage (F = 0.25), 13 were first cousin unions, and six patients were progeny of second cousin unions (F = 0.0156). The two cases of consanguinity identified among ethnic Danes were second cousins (F = 0.0156). The prevalence of consanguinity was 28.2 times more frequent among ethnic minorities than among ethnic Danes. The frequency of consanguinity was conspicuously high among children of Pakistani, Afghan, Turkish and Arab origin (71.4%).

Table 3 shows the prevalence of IEM in the different ethnic groups. More than one case of disease among sibs were counted as a single case. In addition to a high frequency of consanguinity, the prevalence of disease was also significantly higher in ethnic minorities than in ethnic Danes. The frequency of IEM among Danes and ethnic minorities compared to the overall number of Danes and descendants of the above-mentioned ethnic groups were 0.2/10,000 and 5.35/10,000, respectively. As a result, the frequency of autosomal recessive metabolic disorders was 25.5 times higher among children of Pakistani, Turkish, Afghan and Arab origin than among ethnic Danes.

4. Discussion

On April 1st, 2017, immigrants and their descendants represented 12.9% of the total population in Denmark [9]. A large part of these, 8.3%, have origin from countries where consanguineous unions are common [2]. Additionally, figures from the latest population projection show that the proportion of descendants has significantly increased [9]. In the study, consanguineous unions were found to be more frequent among people from Pakistan, Turkey and Afghanistan. These are countries, where consanguineous unions are common, and they represent some of the largest ethnic minority groups in Denmark [2].

This is the largest report of IEM in the offspring of consanguineous unions in Denmark. Although the nationwide neonatal screening for IEM is limited to selected disorders, it is important to remember that IEM are a heterogeneous group of rare genetic disorders and that the total number of patients in the study diagnosed with an IEM is relatively high.

The high frequency of MCADD is, especially for the Danish group, significantly higher in ethnic minorities than in ethnic Danes, which is the largest report of IEM in the offspring of consanguineous unions in Denmark. Although the nationwide neonatal screening for IEM is limited to selected disorders, it is important to remember that IEM are a heterogeneous group of rare genetic disorders and that the total number of patients in the study diagnosed with an IEM is relatively high.
(58.0%), not a striking observation given that MCADD is the most prevalent non-PKU congenital metabolic disorder in Western countries \[10\]. Moreover, increased frequencies of mutations have been found following the implementation of neonatal screening \[5,11\].

Compared to the ethnic Danish population, the prevalence of autosomal recessive IEM is conspicuously high among ethnic minorities in the study (Table 3). The findings show a highly significant association between consanguinity and IEM. This association with consanguinity is predictable since all cases of IEM in the study have an autosomal recessive mode of inheritance, and migrants from countries where consanguinity is common tend to preserve traditional patterns of marriage \[12,13\].

The high proportion of Pakistani, Turkish, Afghan and Arab patients might be the result of the high rate of parental consanguinity. Although control for medical and sociodemographic variables, e.g. maternal age, body mass index, occupation, education, smoking, alcohol consumption, nutrition, parity, medical problems during pregnancy could have been instructive, it is highly unlikely that most of these variables would increase the frequency of a disease with an autosomal recessive mode of inheritance \[6,14\].

Some well-investigated studies on risk factors for congenital anomalies that also controlled for non-genetic variables have demonstrated a highly significant association between consanguinity and congenital anomalies. These studies had contemporary control groups and were of sufficient size to achieve significance in comparisons between the offspring of consanguineous and those of non-consanguineous unions. In a Norwegian study \[15\], the proportion of congenital anomalies in children of Pakistani origin born to first cousins that could be attributed to consanguinity was 28%, similar to what was reported from a UK study (31%) \[16\]. The Norwegian and the UK study reported an adjusted relative risk for congenital anomaly of 2.15 and 2.19, respectively, for children of Pakistani origin who were the product of a first cousin union. The Danish Pakistani community merits particular attention. Many studies have identified consanguinity as a causal factor for the elevated rates of congenital anomalies among some ethnic minorities, particularly the Pakistani community. Specifically for IEM, another prospective study on births in Birmingham during the 1980s concluded that if the tradition of consanguineous marriage was abandoned by the Pakistani community, a 60% reduction in deaths and severe morbidity would be achieved \[17\]. An associated study further reported that the overall prevalence of IEM in UK Pakistani children was ten-fold higher than in children of European heritage, among whom parental consanguinity was estimated to be 0.2% \[1\]. Comparing countries separately, our findings show that children of Pakistani origin had the second highest prevalence of IEM, and the highest proportion of consanguinity. Consequently, consanguinity was 46.5 times more frequent among Pakistani descendants than among ethnic Danes (2.15%). The prevalence of IEM among Pakistani descendants was 6.5/10,000 (Table 3). As a result, the frequency of autosomal recessive metabolic disorders was 31 times higher among children of Pakistani origin than among ethnic Danes (0.21/10,000).

A direct comparison between the children of Pakistani and Turkish origin show that the prevalence of IEM is significantly higher among children of Pakistani origin. As in the Pakistani group, almost all the children of Turkish origin had consanguineous parents. However, Turks and their descendants comprise the largest minority group in Denmark and are almost three times as big as the Pakistani community. Furthermore, a nationally representative study indicated that 56.4% of marriages in Pakistan were between first or second cousins, with little or no evidence of any change in prevalence during the preceding 40 years \[18\]. Studies conducted in Turkey indicated that about 20.1% of marriages were between first or second cousins \[19,20\]. Direct comparison of the studies is difficult because of the different reporting methods used. However, the data suggest that frequency of consanguineous unions is generally higher among Pakistanis.

Newborns of Afghanistan origin had the highest prevalence of IEM (Table 3). This is most likely because consanguineous unions are common among Afghans, and that immigration from Afghanistan has increased significantly since 2001. A study conducted in 2011/2012 indicated that over 50% of marriages in Afghanistan were between consanguineous couples \[21\].

We acknowledge that a combination of biradier endogamy, especially in relation to the Pakistani community, arrangement within ethno-linguistic groups (Hazara, Pashtun etc.) and consanguinity are likely responsible for the high disease frequencies. However, we do not have any nation-wide data available on biradier membership among the different ethnic groups to comment on it in the context of our study. Further, the relatively high number of diagnosed cases \[41\] referred from Greenland and The Faroe Islands is most likely explained by a founder effect \[22\]. The latter is a geographically isolated group of islands that originates largely from colonization by a small number of Norwegians about 1000 years ago. In a previous study by the authors (A.L., F.S.), it was observed that carnitine transporter deficiency (CTD) and holocarboxylase synthetase deficiency (HLCSD) were relatively frequent among patients referred from The Faroe Islands to our department for diagnosis and treatment \[23\]. CTD and HLCSD have an estimated carrier frequency of about 1:20 for both disorders \[23\]. The birth rates of HLCSD and CTD among Faroese are about 1:1700 (DK 1:100,000) and 1:300 (DK 1:100,000), respectively. This provides an example of how most likely a founder effect and genetic drift have resulted in a unique national genetic disease profile.

Although there was a conspicuously high association between consanguinity and IEM, analysis of some non-genetic variables (age, parity, taxable income, household size), would have been instructive. Danish \[24,25\] and many other international studies \[26–31\] have demonstrated that the high rate of infant death with inherited congenital disorders as a primary cause among some ethnic groups cannot be explained by the aforementioned variables. Furthermore, some of the studies that controlled for these risk factors still found consanguinity to be a major risk factor for congenital anomalies \[15,16\]. Moreover, by investigating IEM with a recessive mode of inheritance, it is highly unlikely that the association reported would be strongly affected. This leads to the question of perhaps consanguinity being a major risk factor for the higher death rates from congenital abnormalities among some ethnic minorities. With ongoing demographic changes in Western countries, differences in health among offspring of some ethnic minorities have been observed in a number of studies where an increased risk of death in infancy was seen for offspring of immigrants \[24–31\]. A Danish population-wide study showed an increased stillbirth and infant mortality rate among children of mothers of ethnic Turkish and Pakistani origin compared with ethnic Danish children \[24\]. A majority of the deaths were due to congenital anomalies, which is consistent with other international studies that found that inherited congenital conditions were one of the main causes of the excess risk of mortality. A Dutch study on early child mortality (0–2 years) found that deaths from hereditary diseases in the first 2 years of life were four to five times higher among Turkish immigrants \[32\]. Another study found that the leading cause of infant death among infants of mothers born in Pakistan and Bangladesh was congenital anomalies \[33\].

The study is the largest report of data for IEM in the offspring of consanguineous unions in Denmark. Our findings confirm that the offspring of consanguineous unions have an increased risk for IEM. Couples contemplating such unions should be advised of these risks; however, advice should be given with sociological awareness. Ethnic groups can have different perceptions of genetic disease and risk factors. This can have an impact on their preferences toward genetic counseling and treatment \[34\]. Linguistic barriers may also limit comprehension of genetic issues. A study found increased attendance of Pakistanis at a medical genetic clinic if a Pakistani doctor would also be present perhaps due to a better understanding of language and culture \[35\].

Demographic changes have significantly impacted on the patient
profile of clinical genetic services in many countries [3,4]. As a result, knowledge about the health outcomes of consanguinity is pertinent for health care professionals providing clinical genetic services and to primary care takers to assist consanguineous couples to make informed decisions about family planning. Perhaps even education programmes at secondary school level, and leaflets at relevant languages on recessive diseases and consanguinity might prove beneficial. The results of this study will hopefully increase knowledge and inform health personnel, who work with communities at increased risk.

Conflict of interest

The authors declare no conflict of interest.

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