BMJ Open Wilson disease in the USA: epidemiology and real-world patient characteristics based on a retrospective observational health claims study

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ABSTRACT

Objectives To describe the epidemiology, patient characteristics and comorbidities in patients with Wilson disease (WD) in the USA.

Design Retrospective, population-based study.

Setting The study used the US Komodo claims database containing records regarding medical claims for over 120 million individuals.

Participants Patients with WD were identified via ICD-10 (10th revision of the International Classification of Diseases) code during the study period 2016–2019 and no age restriction was applied. A further stratification by disease subtype ('hepatic', 'neurologic' and 'psychiatric') was performed.

Main outcome measures WD prevalence was reported by age, sex and US census regions/divisions. Adjusted prevalence was calculated using age-specific prevalence standardised to the USA (2010 US census) and to the world (WHO 2000–2025) to enable comparisons across countries, using direct standardisation of prevalence estimates by age group.

Results Overall, 2115 patients with WD were identified during the study period. Among them, 56.8% had hepatic symptoms, 57.0% neurologic symptoms and 47.4% psychiatric symptoms. The most frequent manifestations in hepatic patients were liver signs and symptoms (90.8%), in neurologic patients cognitive defects (50.7%) and in psychiatric patients mood disorders (86.4%). The mean age in the overall cohort was 39.9 years. Prevalence estimation was based on 1481 patients with WD between 2017 and 2019. The 2017–2019 crude period prevalence was 21.2 patients per million (95% Cl: 20.1 to 22.3), with similar prevalence observed for both sexes.

Conclusions This study provides important real-world data on the diagnosed prevalence of WD in the USA and revealed the comorbidities associated with various disease subtypes, thereby providing a comprehensive basis for guiding physicians and policy makers in the management of this chronic disease.

INTRODUCTION

Wilson disease (WD) is an autosomal recessive condition which leads to an excessive copper deposition in body organs, particularly in the liver and the central nervous system.¹ This

STRENGTHS AND LIMITATIONS OF THIS STUDY

- ⇒ This study estimated the diagnosed prevalence of Wilson disease (WD) for the first time in the USA, by using a large, nationally representative claims database. The real-world nature of the data helps estimate the observed frequency of WD in the USA using ICD-10 (10th revision of the International Classification of Diseases) codes specific to WD.
- ⇒ The classification of patients into subtypes, and especially the novel approach to differentiate psychiatric symptoms from neurological symptoms, was a further strength of this study.
- ⇒ Besides general limitations inherent to claims data, the limitations specific to this study included a limited study period after the introduction of a new WDspecific ICD-10 code in 2015.
- ⇒ The used claims data did not capture information on over the counter medications and the treated patients represented the proportion of patients with prescription treatment only.

genetic disorder is caused by mutations of the P-type ATPase copper transporter *ATP7B* gene located in the human chromosome 13.² To date, more than 600 variants in the *ATP7B* gene have been described, and most patients with WD are compound heterozygous with two different *ATP7B* variants, complicating and prolonging genetic WD diagnosis.³ WD generally presents in childhood and young adulthood with the most common age of presentation between 10 and 20 years, though patients can occasionally present before the age of 5 years and after the age of 70 years.⁴

Clinical presentation of WD includes a combination of hepatic, neurologic, psychiatric and ophthalmologic symptoms. Psychiatric symptoms often precede the diagnosis of WD and include personality changes, depression, cognitive changes and anxiety.⁵ The neurologic symptoms of WD mostly refer to dysfunction in the extrapyramidal system

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Shona Fang; shona.fang@alexion.com including dysarthria, dystonia, gait abnormalities, tremor, parkinsonism, chorea and seizures.⁶ Psychiatric symptoms have been only recently recognised as independent manifestations of WD, as it was previously assumed that they occur together with neurological symptoms, and patients were often referred to as having neuropsychiatric disorders.⁷ The symptoms related to hepatic dysfunction in WD can range from an asymptomatic increase in liver enzyme levels to severe liver failure. Typically, early on in disease progression, there is a mild increase in transaminases, which then may progress to chronic active hepatitis, followed by fibrosis and cirrhosis.⁸

WD is a rare disease with recent worldwide clinical prevalence estimates ranging from approximately 16.7 to 25 patients per million,⁹ although the prevalence can vary across countries and may be higher in selected regions such as in some Asian communities.¹⁰ Interestingly, there is a discrepancy of WD prevalence estimates originating from epidemiological compared with genetic studies and recent genetic data with a considerably higher prevalence (139 per million) indicate that the prevalence of WD may be underestimated.¹¹ The possibility of underdiagnosis, misdiagnosis and the lack of accounting for incomplete variant penetrance in the genetic calculations were hypothesised as the main reasons for the observed inconsistency between genetic versus clinical data.^{11 12} For the USA, population-based epidemiological data are still scarce. A previous study reported a genetic birth prevalence of 18.2 per million births,¹³ and an abstract on cardiac manifestations in WD reported a prevalence of 26 patients per million, but no details of methodology.¹⁴

Thus, large population-based studies assessing the epidemiology of WD and its subtypes in the USA are needed, allowing for a greater characterisation around the spectrum of disease severity and the diagnosed prevalence of WD. Here, we present data from a recently completed real-world evidence study that used US claims data to investigate various aspects of WD in a real-life setting including epidemiology and patient demographic and clinical characteristics.

METHODS

Study design

This retrospective, population-based observational study used health insurance claims data to assess WD epidemiology and patient characteristics using the US Komodo Health claims database available from 2012 to 2020.¹⁵ Komodo's private payor-complete database contains records regarding prescription and/or medical claims for over 120 million individuals, collected from more than 150 private insurers in the USA, including Medicaid managed-care and Medicare Advantage plans. This study used the closed claims available in the database. Closed claims come directly from the payer and provide the complete patient journey, such as full medical and/or prescription benefit information including insurance

Population

Patients were selected based on their first observed WD diagnosis (index event) in the claims database between 2016 and 2019 (study period). The diagnosis was based on code (ICD-10 code E83.01 or ICD-9-CM code 275.1 prior to 1 October 2015). The standard ICD-9 diagnostic code for WD was not specific to WD and could also Found and the use another rare condition called Menkes disease. The WD-specific ICD-10 code (E83.01) was introduced in the USA on 1 October 2015. A WD diagnosis was defined the USA on 1 October 2015. A WD diagnosis was defined as at least one inpatient or two outpatient visits with a WD ICD-10 code, separated by at least 30 days during of the study period. For the selected patients with WD, a follow-up period of at least one calendar year (defined as postindex period) was examined, to observe their treatment journey. The follow-up period was defined from index until the last available claim. No age restriction was imposed. Patients were excluded from the study if they **G** had at least one prescription for a copper replacement **o** drug, a drug used to treat Menkes disease, during the uses related to text study period.

Cohorts and subgroups

The cohort presented here includes patients that fulfilled the criteria for at least one WD diagnosis at any time during the study period (2016–2019) and a continuous enrolment for at least one calendar year before the index event. Separately, prevalence was assessed in the period between 2017 and 2019 for patients that had a history of at least one WD diagnosis at any time during that period and a continuous enrolment for at least two calendar years. This separate assessment period for prevalence d estimation was based on the WD-specific diagnosis code \triangleright becoming effective in the USA only in 2015. Due to the possibility of miscoding, the most accurate prevalence estimation of WD can be made between 2017 and 2019, **a** , and as this period includes at least one full calendar year since the introduction of the new code.

A further stratification by WD subtypes was performed, according to ICD-10 codes (see online supplemental tables 1-3) based on the presence of signs and symptoms for each category. The subtypes 'hepatic', 'neurologic' and 'psychiatric' were not mutually exclusive or strictly delineated but were based on the presence of signs and **g** symptoms. Patients were assigned to the hepatic subtype if they were diagnosed with liver signs and symptoms, acute hepatitis (not viral), cirrhosis (decompensated or compensated), liver failure, portal hypertension or hepatocellular carcinoma. Patients assigned to the neurologic subtype had a diagnosis of tremor, parkinsonism or akinetic rigid syndrome, gait abnormalities/ataxia, dysarthria, dystonia, chorea, dysphagia, myopathy, seizures, migraine, somatoform autonomic dysfunction or cognitive disorder. Patients diagnosed with mood disorders,

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Table 1 Selected baseline characteristics of patients with Wilson disease (WD), overall and by subtype						
	Hepatic*, n=1202	Neurologic*, n=1206	Psychiatric*, n=1003	Overall, n=2115		
Age at index date (years), mean (SD)	40.4 (19.08)	42.3 (20.70)	39.2 (20.19)	39.9 (20.06)		
Male, N (%)	607 (50.5%)	659 (54.6%)	558 (55.6%)	1096 (51.8%)		
US region of residence at index, N (%)						
Northeast	254 (21.1%)	246 (20.4%)	182 (18.2%)	462 (21.8%)		
South	376 (31.3%)	398 (33.0%)	332 (33.1%)	647 (30.6%)		
Midwest	275 (22.9%)	292 (24.2%)	249 (24.8%)	502 (23.7%)		
West	297 (24.7%)	270 (22.4%)	240 (23.9%)	504 (23.8%)		
Physician specialty of primary WD diagnosis, N (%)†						
Gastroenterology	197 (16.4%)	117 (9.7%)	90 (9.0%)	279 (13.2%)		
General/family practice	136 (11.3%)	133 (11.0%)	124 (12.4%)	266 (12.6%)		
Ophthalmology/optometry	61 (5.1%)	59 (4.9%)	50 (5.0%)	101 (4.8%)		
Neurology	41 (3.4%)	75 (6.228%)	48 (4.8%)	88 (4.2%)		
Follow-up period (years), mean (SD)	2.2 (1.28)	2.1 (1.25)	2.1 (1.26)	2.2 (1.27)		
Never treated, N (%)	962 (80.0%)	1044 (86.6%)	881 (87.8%)	1755 (83.0%)		

*These groups are not mutually exclusive.

†This is the primary physician specialty for the first claim with WD as the primary diagnosis. Due to limitations in the analysis of the prevalent WD population, it cannot be inferred with certainty that it represents the first diagnosis.

paranoia/schizophrenia, psychosis or personality disorders were assigned to the psychiatric subtype.

Variables

Data on patient characteristics (age, sex, region of residence) and physician specialty at baseline (ie, at first observed WD diagnosis during the study period) were ascertained. In addition, the proportion of patients never treated and distribution of subtypes and manifestations were ascertained during the study period. Comorbidities/concurrent diagnoses were assessed during the study period (2016-2019). Overall, age-specific, sex-specific and region-specific prevalence (crude and age-adjusted

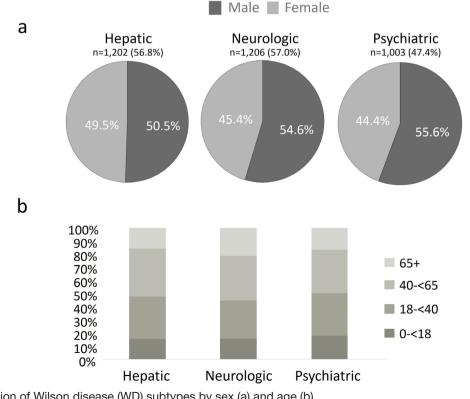


Figure 1 Distribution of Wilson disease (WD) subtypes by sex (a) and age (b).

 Table 2
 Most frequent (top 3) Wilson disease (WD)

 manifestations by subtype

	·		% overall cohort	
Manifestations	Ν	% subtype	(n=2115)	
Hepatic subtype* (n=1202)				
Liver signs and symptoms	1092	90.8%	51.6%	
Cirrhosis	456	37.9%	21.6%	
Hepatitis	347	28.9%	16.4%	
Neurologic subtype* (n=1206)				
Cognitive defects	611	50.7%	28.9%	
Ataxia and gait abnormalities	436	36.2%	20.6%	
Dysphagia	328	27.2%	15.5%	
Psychiatric subtype* (n=1003)				
Mood disorders	867	86.4%	41.0%	
Mix of symptoms, children	224	22.3%	10.6%	
Paranoia and schizophrenia	199	19.8%	9.4%	
*These groups are not mutually exclusive.				

period prevalence) were assessed in the period between 2017 and 2019. A rationale for considering a period prevalence rather than an annual prevalence was that the identification of patients with WD was based on ICD-10 codes, without considering WD treatment or laboratory records, and the assumption that patients with WD who are well managed using over the counter (OTC) or prescription medication may not necessarily visit a health-care provider (HCP) each year. A further reason for using

a period prevalence was to account for possible misdiagnosis in the first year after the introduction of the ICD-10 code for WD.

Data analysis

Descriptive statistics were calculated for demographic and baseline characteristics (overall and by subtype). Means and SDs were provided for continuous or discrete data. Frequencies and percentages were provided for categorical data. For WD manifestations, the frequencies of occurrence of specific ICD-10 codes within subtypes and within the overall patient cohort were calculated and the three most frequent manifestations were listed by subtype. For comorbidities, all recorded diagnoses (ICD-10 codes level 3) at any time during the study period were ranked according to frequency and a list of the 10 most frequent comorbidities in the overall cohort were reported. In addition, the Charlson Comorbidity Index (CCI; version of 2011), an assessment tool designed to predict mortality in patients with multiple comorbidities,¹⁶ was assessed for each patient and a mean score including SD (higher scores indicating greater mortality risk and more severe comorbid conditions) was calculated for the overall cohort. Crude period prevalence was calculated from the number of WD cases identified and was expressed as patients per million including 95% CIs. The denominator was the number of patients with **ö** continuous enrolment spanning the period of interest (2017–2019) and the numerator was any patient from the denominator with at least one WD diagnosis claim during the period of interest. Prevalence was reported by age, sex and US census regions/divisions. Adjusted prevalence was calculated using age-specific prevalence standardised to the USA (2010 US census) and to the world (WHO 2000-2025) to enable comparisons across countries, using direct standardisation of prevalence estimates by age group.

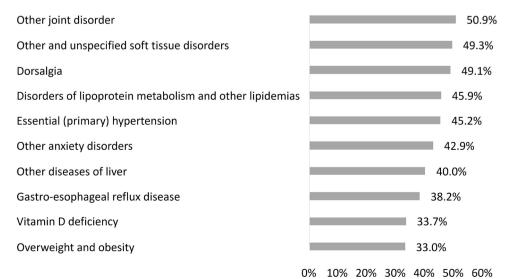


Figure 2 Most frequent (top 10) comorbidities/concurrent diagnoses of patients with Wilson disease (WD).

Table 3 Crude and age-standardised Wilson disease (WD)

	N	Prevalence per million (95% CI)
Crude prevalence (overall)	1481	21.2 (20.1 to 22.3)
Age (years)		
<18	256	12.4 (10.9 to 14.0)
18 to <40	490	27.1 (24.7 to 29.5)
40 to <65	581	26.1 (24.0 to 28.2)
65+	154	17.1 (14.4 to 19.8)
Sex		
Male	697	21.2 (19.7 to 22.8)
Female	784	21.1 (19.7 to 22.6)
Region		
Northeast	344	24.1 (21.6 to 26.7)
South	439	18.4 (16.7 to 20.1)
Midwest	355	22.2 (19.9 to 24.5)
West	319	21.6 (19.2 to 24.0)
Missing	24	
US-adjusted prevalence (overall)*	1481	22.0 (20.9 to 23.1)
WHO-adjusted prevalence (overall)†	1481	21.4 (20.4 to 22.5)

*Standardised to the age distribution of the total US population in year 2010, according to Census Bureau.

†Standardised to the age distribution of the total world population in year 2000-2025, according to WHO.

Patient and public involvement

Patients or the public were not actively involved in this observational study.

RESULTS

Baseline characteristics

The observed cohort included 2115 patients with WD identified between 2016 and 2019. Selected baseline characteristics of the overall cohort and by subtype are displayed in table 1. The mean age in the overall cohort was 39.9 years (SD=20.1 years). The lowest mean age was reported for the psychiatric subtype (39.2 years, SD=20.2 years) and the highest for the neurologic subtype (42.3 years, SD=20.7 years). Overall, 51.8% were male patients. A larger portion of patients came from the South (30.6%)and fewer from the Northeast (21.8%) with little difference between subtypes. While for a majority (54.2%) of overall WD cases no information was available regarding the specialists managing the primary WD diagnosis, the available results showed that patients were mostly in the care of gastroenterologists (13.2%) and general/ family practitioners (12.6%). Hepatic patients were more frequently seen by gastroenterologists (16.4%), as compared with the other subtypes (9.7% of neurologic

patients and 9.0% of psychiatric patients). On average, the follow-up time in the study was 2.2 years (SD=1.3 vears) with little difference between subtypes. Among all patients, the majority (83.0%) were never treated with a reimbursable WD prescription medication during the study period.

WD subtypes and manifestations

Among the cohort, more than half had hepatic and/or neurologic symptoms (56.8% and 57.0%, respectively) σ and around half had psychiatric symptoms (47.4%). Most patients had overlapping subtypes, presenting concomitantly with hepatic, neurologic and/or psychiatric signs and symptoms (see online supplemental figure 1). For instance, 734 (34.7%) presented with at least two concomitant conditions, whereas 422 (20.0%) patients with WD presented with the three of them together. Figure 1 illustrates the distribution of sex (a) and age (b) within subtypes. Slightly more male than female patients had neurologic (54.6%) and psychiatric (55.6%) symptoms, whereas the sex distribution was equal in the hepatic subtype. With regard to age distribution, most patients in all subtypes were in the age groups between 18-39 uses reia years and 40-64 years. The distribution of young (0-17 years) and older (above 65 years) patients was also similar between subtypes, though there were slightly more older patients in the neurologic subtype (21% vs 16% (hepatic)) and 17% (psychiatric)).

The most frequent manifestations associated with the e different subtypes and their proportion in the overall cohort are highlighted in table 2. Hepatic patients most frequently experienced liver signs and symptoms (90.8%), cirrhosis (37.9%) and hepatitis (28.9%). Neurologic a patients most frequently experienced cognitive defects (50.7%), ataxia and gait abnormalities (36.2%) and dysphagia (27.2%). Psychiatric patients most frequently ≥ experienced mood disorders (86.4%), a mix of symptraining, and toms in children (22.3%) and paranoia/schizophrenia (19.8%).

Comorbidities/concurrent diagnoses

Patients with WD were diagnosed with a broad spectrum of comorbidities. The most common clinical conditions diagnosed during the study period included other joint disorders (50.9%), unspecified soft tissue disorders (49.3%), dorsalgia (49.1%), disorders of lipoprotein metabolism and other lipidemias (45.9%), essential hypertension (45.2%), anxiety disorders (42.9%) and $\textbf{\texttt{G}}$ other diseases of liver (40.0%) (figure 2). The extent of concurrent diagnoses and comorbidities was also reflected by a mean CCI Score of 1.92 (SD=2.68).

Prevalence

Prevalence estimation was based on 1481 patients with WD between 2017 and 2019. The 2017–2019 crude period prevalence was 21.2 patients per million (95% CI: 20.1 to 22.3), with similar prevalence observed for both sexes (male: 21.2 per million, 95% CI: 19.7 to 22.8 and female: 21.1 per million, 95% CI: 19.7 to 22.6) (table 3). The crude period prevalence peaked among young adults in the 18 to 39 years age group (27.1 per million, 95% CI: 24.7 to 29.5), followed closely by adults 40 to 64 years old (26.1 per million, 95% CI: 24.0 to 28.2). The highest crude period prevalence was recorded in the Northeast (24.1 per million, 95% CI: 21.6 to 26.7), followed by the Midwestern (22.2 per million, 95% CI: 19.9 to 24.5), Western (21.6 per million, 95% CI: 19.2 to 24.0) and Southern (18.4 per million, 95% CI: 16.7 to 20.1) regions of the USA.

Adjusted period prevalence was calculated using agespecific prevalence standardised to the USA (2010 US census) and to the world (WHO 2000–2025). Between 2017 and 2019, both the US-adjusted and WHO-adjusted period prevalences were similar (22.0 per million, 95% CI: 20.9 to 23.1 and 21.4 per million, 95% CI: 20.4 to 22.5, respectively) to the crude period prevalence observed in this study.

DISCUSSION

Principal findings and implications

There is a gap of knowledge regarding the epidemiology and clinical characteristics of patients with WD from a national and international perspective. In this large population-based study, we assessed WD patient characteristics and epidemiology in the USA.

Approximately two-thirds of the cohort showed hepatic and/or neurologic symptoms, which is consistent with the natural disease course, as WD mostly affects the liver and brain.¹⁷ Typically, hepatic symptoms precede the onset of neurologic symptoms and may thus be diagnosed at a younger age.² In line with this, though not as evident in the age distribution by subtypes, the mean age at index in the hepatic subtype was slightly lower compared with the neurologic subtype. In agreement with our data on sex distribution in the neurologic and psychiatric subtypes, a registry-based study observed that the neuropsychiatric WD form occurred more frequently in men (67%) versus women (49%).¹⁸

Our results also revealed the extent of comorbidities associated with various disease subtypes, as some of the most frequently reported manifestations were quite severe (eg, around 40% of patients in the hepatic subtype presented with cirrhosis). The frequency of hepatic, neurologic, and psychiatric presentations, as well as the manifestations within these subtypes, varies considerably in other published studies.⁴ Considering the great variability in WD symptoms, it is likely that there is referral bias. A review of several independent case series suggested that dysarthria, gait abnormality/ataxia and dystonia are the most frequent manifestations in the neurologic subtype, which partially reflects our findings.⁴ The higher rates of dystonia and dysarthria reported in the literature may be explained by the fact that only symptomatic conditions that require treatment get captured in the claims under the ICD-10 diagnosis code. Furthermore, a study

from China reported distributions of hepatic manifestations (eg, 22% cirrhosis and 11% cirrhotic complications) which were similar to our findings.¹⁹

Patients with WD in our study were in the typical age range for first symptom onset and disease progression (between the second and sixth decades of life). We found a slightly lower mean age in the psychiatric subtype compared with the other subtypes, which might reflect that psychiatric manifestations are often the first symptoms.' The earlier diagnosis of psychiatric WD may highlight a critical gap in care for these patients, as neurologic and psychiatric involvements have been previously assessed in tandem, with neurologic symptoms 9 becoming more apparent later in life. Given the recessive nature of WD, the results confirmed the expectation of **8** a relatively even distribution of cases across sexes, which also aligns with previous studies conducted in France,²⁰ and Hong Kong.¹⁹ The geographical distribution within the USA, with a larger portion of patients coming from the South and fewer from the Northeast, might correspond to the distribution of major university centres in the USA where patients with WD are typically diagnosed. Given the nature of the disease, it is not surprising that uses rela patients were primarily in the care of gastroenterologists (most pronounced in hepatic patients), though a considerable fraction of patients was also diagnosed or treated by general or family practitioners who might be the first contact points. The high rate of patients never treated đ with a reimbursable WD medication during the study te period was likely due to the use of OTC zinc preparations which are not covered by insurers in the USA, the refusal to take medications and rely solely on a low-copper diet, or the hypothesis that some physicians used the WD diaga nosis code to initiate further testing for the patients but did not prescribe a WD medication at that initial stage. Further investigation on this aspect is warranted given the chronic nature of this disease.

Furthermore, our results demonstrated that patients with WD were diagnosed with a broad spectrum of comorbidities, most frequently affecting joints, soft tissues and the cardiovascular system, as well as pain symptoms and metabolic disorders. Comorbidities of WD are rarely described in the literature and vary considerably; however, our findings are at least partially consistent with comorbidities reported in the literature. Kruger *et al* and Dziezyc *et al* described that the most common complications in patients with WD were pain, renal, neurologic, cardiac, skin, osteoarticular or endocrinologic complications and included other organ disturbances.^{21 22}

Only a few population-based studies assessing WD epidemiology have been performed to date worldwide. It is challenging to compare the results from this study with other reports, because of the heterogeneity of the disease and of the heterogeneity in population, diagnosis, and methodologies employed in other studies assessing WD epidemiology. However, the observed prevalence estimates (US-adjusted prevalence: 22.0 patients per million) align well within the WD prevalence range reported in other claims-based studies in France (15 per million),²³ and China (17.9 per million),¹⁹ while estimates from a claims-based study in South Korea were slightly higher (38.7 per million).²⁴ Two recent systematic literature reviews assessed WD prevalence data originating from population-based epidemiological as well as genetic studies.^{11 25} Sandahl et al reported a crude populationbased prevalence between 25 and 34.5 per million,²⁵ and underlined that specific populations in Croatia, Sardinia, Israel, Costa Rica, Middle Eastern countries, Pakistan and India had considerably higher prevalence estimates, related to either consanguinity or higher mutation frequencies. Gao et al performed a meta-analysis resulting in a pooled population-based prevalence of 14 patients per million.¹¹ Both reviews mentioned that at least in some studies a higher genetic versus clinically based prevalence could be observed. Gao et al reported a pooled prevalence at birth of 127 per million.¹¹ Overall, it seems evident that many factors such as the still uncertain mutational spectrum and penetrance of WD variants, the unclear effect of combined mutations and epigenetic factors, methodological differences in studies, underdiagnosis, changes in diagnostic awareness and treatment options over time, as well as geographic factors may influence WD prevalence estimates. Thus, it is important to assess epidemiological data from large, population-based studies, as the one presented here.

Strengths and limitations

This study benefitted from several strengths. Data elements extracted from claims, some of which are not generally available in literature or patient registries, are collected routinely in clinical practice and represent realworld activities and outcomes. The real-world nature of the data helps estimate the observed frequency of diagnosed diseases using ICD-10 codes. The ICD-10 codes for WD are highly specific, despite any limitations in coding or misdiagnosis which may occur. The Komodo claims data are large and nationally representative based on comparisons made to the US census estimates (by age, sex and region). The closed claims provide a longitudinal history that facilitates the analysis of the natural history of the disease, healthcare utilisation and treatment dynamics. Of note, the crude prevalence observed in this study between 2017 and 2019 was comparable to the US age-standardised adjusted results, reaffirming the representativeness of the Komodo data for rare disease research in the USA. The classification of patients into subtypes, and especially the novel approach to differentiate psychiatric symptoms from neurological symptoms, was a further strength of this study.

There are some general limitations inherent to claims data, such as the potential for errors in diagnosis coding or record keeping at the point of the HCP. Since claims data are used for billing purposes, they only include records for the insured population; therefore, projections of the US population assume similarities between the insured and uninsured patients. Leveraging secondary data requires an algorithm for case identification and validated algorithms specific to WD do not currently exist. Claims data contain only reimbursed medical services and thus for example, results of lab tests or patient-reported outcomes are not captured. In addition, these claims data in general have limited follow-up, for example, due to patients switching insurance plan or losing coverage or privacy restrictions limiting data linkage.

In addition, there were some limitations specific to this study. Epidemiological trends over time were hard to assess given the limited study period and the introduction of the new WD-specific ICD-10 code. Only symptomatic conditions were captured under the ICD-10 code; therefore, no inferences can be made regarding pre-symptomatic patients with WD. The claims data did not capture information on laboratory testing well, and results are not available in claims data. Further the claims data did not capture OTC medications and thus the treated patients represented the proportion of patients with prescription treatment only. The proportion of patients without a claim for a WD-specific treatment was quite high and unexpected however and whether this truly reflects a large proportion of patients on OTC zinc is unclear. Further investigation is needed as to the proportion of patients with WD receiving treatment in the USA.

CONCLUSIONS

This claims study provides important real-world data on the prevalence of WD in the USA and revealed the extent of comorbidities associated with various disease subtypes. The results of this study extend existing research findings and provide a comprehensive epidemiological basis for guiding physicians and policy makers in the management of this chronic disease.

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Contributors SF, CS, KHW and PH were responsible for the design, writing and editing of the final version of the manuscript. SF, CS and HC were responsible for the execution of the study and HC was responsible for the analysis of the data. SF is the guarantor. All authors had critically reviewed the manuscript and approved its final version.

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Competing interests CS and HC declare being employees of Oracle America, which received funding from Alexion Pharmaceuticals to conduct this study. PH has no financial relationships to disclose. KHW declares no conflicts of interest that pertains to this work. SF is an employee of Alexion and AstraZeneca Rare Disease and owns stock in AstraZeneca.

Patient and public involvement Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.

Patient consent for publication Not applicable.

Ethics approval This was study was approved by the IRB Pearl Pathways. The study was conducted in accordance with legal and regulatory requirements, as well as with scientific purpose, value and rigour and followed generally accepted research practices described in Good Pharmacoepidemiology Practices guidelines issued by the International Society for Pharmacoepidemiology.

Provenance and peer review Not commissioned; externally peer reviewed.

Open access

Data availability statement Data may be obtained from a third party and are not publicly available. The data that support the findings of this study are available from Komodo Health. Restrictions apply to the availability of these data, which were used under licence for this study. Access to data must be requested through Oracle.

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