BMJ Open Cohort study of diagnostic delay in the clinical pathway of patients with chronic wounds in the primary care setting

Kirsti Ahmajärvi ^(b), ¹ Kirsi Isoherranen, ² Maarit Venermo³

To cite: Ahmajärvi K, Isoherranen K, Venermo M. Cohort study of diagnostic delay in the clinical pathway of patients with chronic wounds in the primary care setting. BMJ Open 2022;12:e062673. doi:10.1136/ bmjopen-2022-062673

 Prepublication history and additional supplemental material for this paper are available online. To view these files. please visit the journal online (http://dx.doi.org/10.1136/ bmjopen-2022-062673).

Received 16 March 2022 Accepted 05 October 2022

Check for updates

C Author(s) (or their employer(s)) 2022. Re-use permitted under CC BY-NC. No commercial re-use. See rights and permissions. Published by BMJ.

¹Doctoral Programme in Populational Health and **Department of General Practice** and Primary Health Care, Helsinki University Hospital, Helsinki, Finland ²Department of Dermatology and allergology. University of Helsinki and Inflammation Center, Helsinki University Hospital, Helsinki, Finland ³Vascular Surgery Department, University of Helsinki and Helsinki University Hospital, Helsinki, Finland

Correspondence to Dr Kirsti Ahmajärvi;

kirsti.ahmajarvi@helsinki.fi

ABSTRACT

Objectives Exact wound diagnosis is essential for successful wound management and a holistic care of the patient suffering from a wound. Wound management has been traditionally seen as a nursing area, but this can lead to considerable delays in wound diagnostics. A diagnostic delay has been recognised as an element of diagnostic error, which, in turn, affects patient safety. The aim of this cohort study was to examine diagnostic delays of chronic wound within primary care.

Setting A specialised diagnostic unit, a wound care team, was established in the primary healthcare with the objective of reducing diagnostic and treatment delays in primary care.

Participants The data consists of 197 consecutive patients attending their first appointment with the wound care team in 2016. The collected data included basic demographics, information about the clinical pathway. including doctor's appointments in primary and specialised care, as well as the International Classification of Diseases 10th Revision (ICD-10) diagnostic codes.

Primary and secondary outcome measures The diagnostic delays were calculated in days and divided into three groups: (1) patient-related delay, (2) diagnostic delay and (3) organisational delay.

Results The median duration of a patient-related delay was 2 days (IQR 0-14), whereas a physician's first evaluation was performed at a median of 8 (1-32) days from wound appearance and the correct diagnosis by the wound care team was established in a median of 57 (33-100) days. The organisational delay from first contact to diagnosis was a median of 41 (22-80) days. Only one in three patients had a diagnostic delay of less than 4 weeks. **Conclusions** According to this study, the diagnostic delay occurs within primary care, as an organisational delay from first contact to correct diagnosis. It is possible to arrange an optimal pathway of care in which a holistic wound care process starts within primary care.

INTRODUCTION

Chronic wounds pose a significant burden to healthcare, constituting roughly 2%-6% of all healthcare costs.¹⁻³ According to a recent study in the UK, the annual prevalence of wounds increased by 71% between 2012/2013 and 2017/2018 and the patient management costs increased by 48% in real terms.⁴ In addition to costs, chronic wounds cause

STRENGTHS AND LIMITATIONS OF THIS STUDY

- \Rightarrow Strengths of the study include a unique data which contain primary care patients suffering from chronic wounds.
- \Rightarrow Strengths include also systematic and detailed data review and collection.
- \Rightarrow Limitations include the possibility of interpretation bias.
- \Rightarrow Limitations include also the possibility of error in defining the moment of 'the right diagnosis'.
- \Rightarrow There is no input for patient and public involvement in the study design, which could be seen as a limitation.

substantial suffering at the individual level, leading to an impaired quality of life, social isolation and mental health problems.⁵⁻⁷ Wound management can be successful only when the wound is correctly diagnosed and treated accordingly.^{8 9} Wound care has been traditionally viewed as measures related to **E** the assessment of the wound bed, which can obscure the importance of the holistic care of 🦉 the patient.¹⁰

In many European countries, wound a general practitioners (GPs) or nurses.¹¹ This **g** wound patients should receive a timely evaluation by a qualified healthcare professional who can make the correct diagnosis, plan holistic treatment and make the necessary referrals.¹² This process should aim to avoid diagnostic error, which has been recognised by the WHO as a global challenge to patient & safety.¹³ Diagnostic error includes an incorrect or delayed diagnosis, which leads to patient harm or to inappropriate or delayed treatment. The diagnostic errors mainly occur within primary care or at the emergency department, where physicians lack the appropriate tools and sufficient time to make accurate decisions.¹⁴¹⁵

In 2013, a special wound care team was established in the primary healthcare system

text

and

Protected by copyright, including for uses related to

of the Helsinki area. The wound care team consists of a wound care nurse and a GP specialised in wound care. This team has the possibility to consult a podiatrist and/ or vascular surgeon. Patients are referred to a wound care team consultation from all primary care units: health centres, home care units and nursing homes. The instructions for the primary care personnel were to react early and refer patients suffering from a non-healing wound within (2-)4 weeks of wound appearance in order to have the wound appropriately diagnosed. The main focus of the wound care team was to discover the correct diagnosis as early as possible and, thereafter, to initiate proper treatment accordingly.

The purpose of this study was to evaluate the delay in the diagnostic process in the clinical pathway of wound patients who were referred for a consultation by the wound care team during 2016. The delays were divided into system-related and patient-related delays.

MATERIAL AND METHODS

This prospective cohort study analysed the characteristics and medical history of 197 consecutive patients who visited the wound care team in primary healthcare during 2016. The information was collected at the first wound care team appointment.

Data were collected from electronic patient records. The collected data consisted of patients' background factors (sex, age, comorbidities, medication, previous wounds, state of mobility and living standards, need of home care, smoking, blood sugar levels and lipids), as well as the date of wound appearance, the date of the patient's first contact with a primary care unit and physician's appointment, the date of consulting the wound care team and the date of admission to a specialist care unit if needed. In order to analyse the diagnostic process, additional information was collected on signs of infection and bacterial swab results, on whether the Ankle Brachial Index (ABI) was measured, or pulse palpation occurred, or whether neuropathy was detected with a monofilament test. Observations of oedema and any blood test analyses regarding the wound were also recorded. Furthermore, information was gathered from radiological examinations, if performed, as well as any further investigations within specialist care, such as toe pressure and angiography results. The treatment plan was evaluated and compared with the diagnostic methods and the International Classification of Diseases 10th Revision (ICD-10) code. (online supplemental table 1).

Delays were calculated at different points of the care pathway, starting from wound appearance. The different types of delays included: (1) patient-related delay (time from wound appearance to the patient's first contact with healthcare providers), (2) diagnostic delay (time from the onset of the wound to the first physician's appointment where the initial diagnosis was made) and (3) organisational delays within primary care in arriving at the correct diagnosis and treatment (from the first contact with the

Table 1 Categorisation and diagnostic variation in the clinical pathway of a patient with a wound

Diagnostic categories	Primary care physician (n=155)	Wound care team physician (n=197)	Specialist care physician (n=111)
No diagnosis*	26	2	1
Arterial wound	4	16	26
Venous or oedematous ulcer	15	57	17
Diabetic foot ulcer	4	24	15
Pressure ulcer	12	29	9
Post-traumatic wound	16	23	3
Atypical wound	0	8	7
Mixed-aetiology ulcer	0	7	3
Infectious wound	42	11	10
Foot malformation or pressure ulcer	0	7	1
Wound of unspecified aetiology	36	13	19

'The category was defined as 'no diagnosis' when a patient had been seen by a physician but there was no ICD-10-coded diagnosis in the patient records ICD-10, International Classification of Diseases 10th Revision

primary healthcare unit to the wound care team consultation). Some patients needed a referral to a specialist consultation, his delay was also considered.

Diagnostic codes were collected as ICD-10 codes and to text were compared with the diagnoses made by the primary care physician, by the wound care team physician and by the specialist. As the number of different diagnostic codes was high, we categorised the diagnoses into 10 groups (table 1, figure 1A,B). In the grouping process, we also included, in addition to the diagnostic code, information on how the wound had appeared and which diagnostic tools had been used to arrive at the specific conclusion and treatment plan.

In order to avoid bias caused by outliers, 16 patients whose wound had persisted for over 365 days prior to the wound team consultation were excluded from the delay

 analysis.

 Patient and public involvement

 No patient or public involvement has occurred in planning, managing, designing or carrying out this research.

 RESULTS

 A total of 197 patients were included in this study. The mean are use 71 years and 106 (52,5%) of the patients

mean age was 71 years, and 106 (53.5%) of the patients were women. The basic demographics and risk factors of the patients are reported in tables 2–3.

The majority of the patients were living at home (n=172; 86.9%) either without any support (n=130) or with home care (n=42). The patients' living status is presented in table 2. Almost half of the patients had had wounds earlier (48.7%). As regards comorbidities, 39.1% had diabetes, 11.7% had ischaemic heart disease, 17.3% had memory disorders and 9.1% had a mental health condition. Only

Protected by copyright, including for uses rel

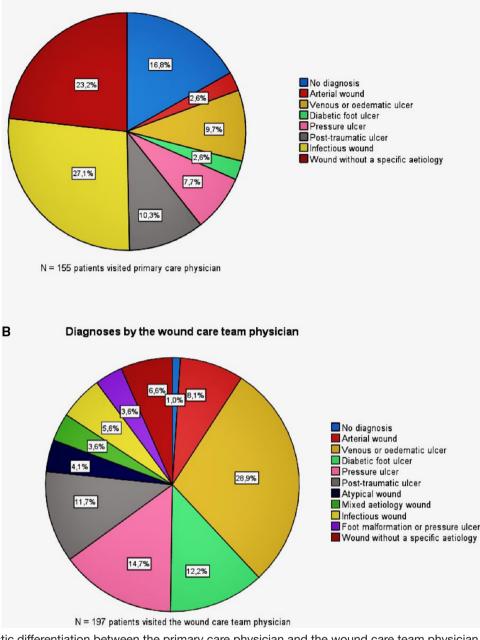
ated

ല

ā



A Diagnoses by the primary care physician





3.6% of the patients had no comorbidities. Overweight (61.4%), obesity (27.9%) and neuropathy (43.7%) were relatively common. Venous insufficiency or a previous deep venous thrombosis had been diagnosed in only 13.2%. As can be seen in tables 3 and 4, the patients had several comorbidities and heterogenous medications. Almost half of the patients used analgesics (44.7% used non-steroidal anti-inflammatory drugs or paracetamol and 15.2% used opioids), whereas different psychopharmaceuticals were used by 11.7%–17.8% of the patients.

Diagnostics

Forty-two (21.3%) patients were not seen by a primary care physician before they visited the wound care team,

meaning that the diagnostic process was not even started before this visit. Of the 155 patients who had been seen by a physician prior to the wound care team, 129 (83.2%) had a recorded diagnosis code (ICD-10), while 26 (16.8%) patients remained undiagnosed. Thus, 34.5% of the patients (n=68) received their first diagnosis by the wound care team.

Out of the patients who were seen by a primary care physician, 85 (58.8%) had no delay (median 0 days), meaning that the patients visited an emergency room and were seen by a physician immediately. The diagnoses for these patients mainly comprised infectious wounds (n=30, 35.3%) and wounds with no specific cause (n=21,

	Female (n)	%	Male (n)	%	Total (n)	%
Sex	106	53.8	91	46.2	197	100
Age (years)						
Mean	78	39.6	69	97.2	71	36.0
Median	80	40.6	73	37.1	41	20.8
Mobility (n)						
Walking	38	19.3	56	28.4	94	47.7
Walking with assistance device	33	16.8	20	10.2	53	26.9
Walking with device only indoors	16	8.1	5	2.5	21	10.7
Wheelchair	11	5.6	8	4.1	19	9.6
Bedridden	8	4.1	2	1.0	10	5.1
Residence (n)						
Home	57	28.9	73	37.1	130	66.0
Home with home care	30	15.2	12	6.1	42	21.3
Assisted living facility	14	7.1	5	2.5	19	9.6
24/7 care nursing home	5	2.5	1	0.5	6	3.0

Of those patients who saw a primary care physician (n=155), 36.2% (n=56) had clinical signs of infection according to the patient records. However, as many as 94 (60.6%) patients were treated with antibiotics, and 82 (52.9%) had a bacterial swab taken.

Of the 129 patients who had a diagnosis before the first appointment with the wound care team, the same diagnosis was made by the team and the primary care physician in 59 (45.7%) of the cases. The concordant diagnoses most often entailed pressure ulcers, infectious ulcers, as well as venous and post-traumatic ulcers. Specialist care was received by 111 patients. The same diagnosis (ICD-10) was made by all three points in the clinical pathway in only 24 (12.2%) cases, and the majority of these comprised infectious ulcers, followed by a venous aetiology and wounds without a specific diagnosis. (Table 5)

Of the patients who visited a specialist, the same diagnosis was made by the wound care team physician and the specialist in 75.5% of the cases (tables 5 and 6). The concordant diagnoses most often comprised diabetic foot ulcers (20.5%), arterial ulcers (19.3%) and venous or oedematous ulcers (15.7%). In the remaining 24.6%(n=27) of the patients, the diagnosis made by the wound care team was revised in specialist care, mostly comprising arterial ulcers (40.7%) that were usually referred for further investigations with a suspicion of an arterial wound. The ulcers that turned out to be arterial wounds were diagnosed by the wound care team as diabetic foot ulcers (14.8%), wounds of mixed aetiology (18.5%), foot malformations (14.8%) and oedematous ulcers (37.0%).

Post-traumatic wounds were categorised into one category. In the primary care setting, there were 16

Delays

The median time from wound appearance to the first đ healthcare contact was 2 days (IQR 0-14 days, range text 0-351 days) and from wound appearance to the first evaluation by a physician 8 days (IOR 1-32 days, range 0-314 days). The majority of the patients had their first healthcare contact at the emergency department where a physician also examined the patient, or at the district nurse's office at a health centre with the possibility of an immediate physician's consultation. The median time from the ≥ onset of the wound to the first wound care team appointment was 57 days (IQR 33-100 days, range 2-358 days). The median time between the first healthcare contact ğ and the wound care team appointment was 41 days (IQR 22-80 days: range 1-484 days). Only one in three patients (n=61) had an organisational delay of less than 4 weeks between the first contact with health services and the appointment with the wound care team.

Half of the patients (n=113, 57.4%) were referred to a secondary healthcare unit to be seen by a specialist, most often by a vascular surgeon (n=67), followed by a plastic surgeon (n=43) and a dermatologist (n=13). Twenty-one (18.6%) patients were referred to two or more specialists. The median delay from the first appointment with the wound care team to the appointment with the secondary healthcare specialist was 21 days (IQR 8-55, min-max -58 to 235; range 293 days).

The median time from the appearance of the wound to the final diagnosis was 57 days (IQR 33-101; min-max 2 to 358; range 356 days).

The delays in different subgroups are presented in table 7.

Table 3 Description of the sample	e (n=197, %	of total) in i	ndividuals; com	orbidities a	nd risk factors		
	Female (n)	%	Male (n)	%	Total (n)	%	P value*
Comorbidities							
Hypertension	60	30.5	50	25.4	110	55.8	
Heart failure	25	12.7	11	5.6	36	18.3	p=0.044
lschaemic heart disease	13	6.6	10	5.1	23	11.7	
Atrial fibrillation	37	18.8	24	12.2	61	31.0	
Respiratory condition	27	13.7	11	5.6	38	19.3	
Cancer	14	7.1	14	7.1	28	14.2	
Mental health condition	9	4.6	9	4.6	18	9.1	
Dementia/memory disorder	25	12.7	9	4.6	34	17.3	p=0.013
Diabetes	30	15.2	47	23.9	77	39.1	p<0.001
Peripheral arterial disease	16	8.1	24	12.2	40	20.3	p=0.042
Kidney malfunction	12	6.1	15	7.6	27	13.7	
Rheumatoid arthritis	15	7.6	6	3.0	21	10.7	p=0.091
Liver malfunction	0	0.0	6	3.0	6	3.0	p=0.007
Spinal stenosis	5	2.5	3	1.5	8	4.1	
Gout	3	1.5	8	4.1	11	5.6	p=0.064
Haematological condition	5	2.5	6	3.0	11	5.6	
Chronic pain disorder	2	1.0	0	0.0	2	1.0	
Urinary condition	4	2.0	10	5.1	14	7.1	p=0.045
Cerebrovascular disorder	15	7.6	13	6.6	28	14.2	
Dermatological disease	3	1.5	2	1.0	5	2.5	
Musculoskeletal disorder	21	10.7	7	3.6	28	14.2	p=0.018
No comorbidities	3	1.5	4	2.0	7	3.6	· ·
Risk factors							
Previous wounds	46	23.4	50	25.4	96	48.7	
Previous deep venous thrombosis	9	4.6	1	0.5	10	5.1	p=0.020
Venous insufficiency	9	4.6	7	3.6	16	8.1	•
Chronic oedema	3	1.5	2	1.0	5	2.5	
Chronic cellulitis	10	5.1	5	2.5	15	7.6	
Previous amputation	4	2.0	1	0.5	5	2.5	
Smoking (n)	13	6.6	28	14.2	41	20.8	p=0.001
Drug misuse	2	1.0	5	2.5	7	3.6	p=0.001
Alcohol abuse	3	1.5	11	5.6	14	7.1	P-0.010
Overweight (BMI 24–30)	59	29.9	62	31.5	121	61.4	
Obesity (BMI over 30)	26	13.2	29	14.7	55	27.9	
High cholesterol (diagnosis)	22	11.2	23	11.7	45	22.8	
LDL over 3.0	23	11.7	19	9.6	43	21.3	
Joint malformation	8	4.1	3	1.5	11	5.6	p=0.010
Neuropathy (diagnostic coded)	8	4.1	18	9.1	26	13.2	p=0.010 p<0.001
Neuropathy (monofilament test	o 32	4.1	54	27.4	86	43.7	h<0.001
posit.)							
MRSA	2	1.0	6	3.0	8	4.1	p=0.048
Hemiplegia	2	1.0	2	1.0	4	2.0	
HbA1c (n=153)							
Mean (SD)	43 (12.9)		49 (16.8)		46 (15.1)		p=0.018†
							Continued

	Female (n) %	Male (n) %	Total (n) %	P value*
Median (IQR)	40	43	41(37–52)	
BMI (n=178)				
Mean (SD)	27.4 (8.2)	29.0 (6.2)	28 (7.4)	
Median	26	28	26 (23–32)	
fP-Kol-LDL (n=169)				
Mean (SD)	2.5 (0.83)	2.4 (0.86)	2.5 (0.84)	
Median	2.4	2.3	2.4 (1.8–3.0)	
fP-Gluk (n=173)				
Mean (SD)	6.0 (1.8)	7.3 (3.4)	6.6 (2.7)	p=0.002†
Median	5.8	6.7	5.9 (5.3–6.9)	

*Pearson's x2, difference between female and male patients.

†One-way analysis of variance test.

BMI, body mass index; fP-Gluk, fasting plasma glucose; fP-Kol-LDL, fasting plasma low-density lipoprotein Cholesterol; HbA1c, glycated hemoglobin; LDL, low-density lipoprotein Cholesterol; MRSA, Methicillin-resistant Staphylococcus aureus.

DISCUSSION

It is well-known that an early diagnosis of the underlying cause of a chronic wound is essential for wound healing and for avoiding amputations.^{8 16 17} However, there are only a few studies describing the importance of wound diagnosis and the deleterious effects of diagnostic delays.¹⁸⁻²⁰ In the current study, we investigated the diagnostic processes and delays in wound care in the Helsinki area. The patients' first contact with healthcare services after wound appearance was prompt, the median delay being only 2 days. In stark contrast, it took 57 days from the appearance of the wound before the patient was seen by the wound care team for the first time. In our material, only 31.0% of the patients visited the wound care team within 4 weeks of the first healthcare appointment. This caused a significant delay in reaching the correct diagnosis of the wound. We also discovered that only 65.5% of all patients had a recorded wound diagnosis before the first wound care team appointment and that this diagnosis matched the final diagnosis in approximately 50% of the cases. Accordingly, in European countries, the delay in diagnosing diabetic foot ulcers (DFU) was over 3 weeks in 21%–34% of the patients. The shortest average time from event to diagnosis was 10 days in the UK, 14 days in Spain and France and 20 days in Germany.¹²

In Finland, wound patients are evaluated and treated mainly during primary care appointments, including home care and nursing homes.²¹ In the mentioned European countries, the healthcare professionals who participate in the diagnostic process vary considerably. In the UK, only 6% of GPs completely agreed that the care and management of DFUs is the GP's responsibility, and 22% did not diagnose DFUs. Instead, in 27% of the cases, district nurses made the diagnosis. In the UK, the approach seems more often to be multidisciplinary, as 49% of the GPs referred patients to a podiatrist if needed.

Protected by copyright, including for In all four countries, GPs were able to refer patients with DFU to specialised multidisciplinary clinics.²

In the UK, DFUs were diagnosed by a GP in only 45% rel of the cases, and most of the wounds were diagnosed by a district nurse or practice nurse.¹²

ated The optimal treatment pathways for wound patients to include patient surveillance and an early detection of wound healing problems. Errors in the pathway may lead to delays and, consequently, even fatal errors, such as a amputations, in some patients.¹⁷

We took a closer look at the ICD-10 codes assessed by the primary care physicians, wound care team physicians and specialists and found that they differed from each other significantly. This highlights the complexity of wound diagnostics. Surprisingly, only 12.2% of the patients ⊳ had the same diagnosis throughout the whole pathway of care, and these mostly comprised infectious wounds. Based on our data, we assume that there was a significant overdiagnosis of infections in primary care, since an infectious wound was diagnosed in 32.6% of the cases and antibiotics were prescribed for 63.2% of the patients in the primary care setting. Similar results have been obtained in Sweden, where the use of a national wound register diminished the use of antibiotics.²³ Evidence of difficulties in the diagnostics of wound infections is also found in a study of GPs recognising and treating wound **b** infections—according to the results, GPs mostly desired further knowledge about when to start or stop treatment (81%-82%), about topical antimicrobials (80%-68%)and about when to prescribe antibiotics (82%-95%).²⁴

Another diagnostic challenge was the diagnosis of an ischaemic wound and DFU in primary care. Only four diabetic and arterial ulcers were diagnosed in primary care. In contrast, the wound care team diagnosed a DFU in 54 patients, and 26 patients were referred to a specialist when an ischaemic wound was suspected. This problem

. uses

	Female		Male		Total		2
Medication	(n)	%	(n)	%	(n)	%	X ²
Cardiac and vessel medicine							
Anticoagulant	41	20.8	26	13.2	67	34.0	p=0.164
Antithrombotic	29	14,7	28	14.2	57	28.9	
Beta blocker	57	28.9	41	20.8	98	49.7	
Diuretic	45	22.8	32	16.2	77	39.1	
Calcium channel blocker	27	13.7	31	15.7	58	29.4	
Angiotensin-converting enzyme-blocker	38	19.3	47	23.9	85	43.1	p=0.018
Statin	40	20.3	35	17.8	75	38.1	
Diabetes medicine							
Oral diabetes medicine	16	8.1	21	10.7	37	18.8	p=134
Insulin	12	6.1	30	15.2	42	21.3	p<0.001
Psychopharmaceuticals							
Dementia medicine	16	8.1	6	3.0	22	11.2	p=0.066
Antidepressant	14	7.1	11	5.6	25	12.7	
Benzodiazepine	13	6.6	10	5.1	23	11.7	
Sleeping pills	23	11.7	12	6.1	35	17.8	p=0.135
Analgesia (mild)	57	28.9	31	15.7	88	44.7	p=0.008
Opiates	18	9.1	12	6.1	30	15.2	
Immune system medicine							
Cancer medicine	2	1.0	3	1.5	5	2.5	
Immunosuppressive	11	5.6	4	2.0	15	7.6	p=0.124
Cortisone per oral	12	6.1	4	2.0	16	8.1	p=0.083
Cortisone cream	4	2.0	0	0.0	4	2.0	p=0.064
Supplements							
Thyroxin	16	8.1	3	1.5	19	9.6	p=0.006
Calcium supplement	46	23.4	16	8.1	62	31.5	p<0.001
Folic acid	7	3.6	3	1.5	10	5.1	
B ₁₂ supplement	11	5.6	12	6.1	23	11.7	
Vitamin D supplement	44	22.3	22	11.2	66	33.5	p=0.013
Nutrition supplement	6	3.0	2	1.0	8	4.1	
Magnesium supplement	7	3.6	4	2.0	11	5.6	
Vitamin K supplement	11	5.6	9	4.6	20	10.2	
Other							
Inhaler/nebuliser	26	13.2	16	8.1	42	21.3	
Proton pump inhibitor	39	19.8	20	10.2	59	29.9	p=0.030
Urine medicine	8	4.1	12	6.1	20	10.2	
Skin cream	19	9.6	10	5.1	20	10.2	p=0.190

is also detected in a multicentre study performed in four European countries. The researchers found that, even though GPs described neuropathy and peripheral arterial disease as cofactors in the DFU development, they investigated DFUs with additional tests in only half of the cases; this entailed monofilament tests in 21%-43% and, more often, pulse palpation or the measurement of the

ABI in 78%–90% of the cases, but diabetic foot infection was tested in only 7%–20% of the investigated cases.¹¹

Our main finding was that there was an organisational delay in reaching a timely diagnosis. The time between wound appearance and the first contact with a physician was adequate, the median being 8 days and two-thirds (58.8%, n=85) of the patients had their first evaluation

Differentiation of the diagnoses when they remained unchanged or were revised over the clinical pathway Table 5

	Diagnoses that r	emained unchanged		Diagnoses that were r	evised
Categorised diagnostic groups	Within primary care	Throughout the entire clinical pathway	By wound care team and specialist care	Primary care physician's diagnosis	Wound care team physician's diagnosis
Arterial wound	0	0	16	4	7
Venous or oedematous ulcer	13	5	17	2	24
Diabetic foot ulcer	2	1	13	2	11
Pressure ulcer	11	3	9	1	8
Post-traumatic wound	15	3	3	0	2
Atypical wound	0	0	7	0	4
Mixed-aetiology wound	0	0	2	0	3
Infectious wound	10	8	9	32	1
Foot malformation or pressure ulcer	0	0	1	0	5
Wound of unspecified aetiology	8	4	6	27	3

at the first contact with health services. This means that a physician's examination was mostly available at the ER or as a rapid consultation during a nurse's appointment at the health centre. However, our study shows that, among these patients, the initial physician's evaluation very seldom leads to a correct diagnosis and treatment.

Regarding wound patients, ER assessment should include the three most important and acute aetiologies, such as infection, ischaemia and diabetes,²⁵ but otherwise the ER might not be the optimal setting for diagnosing wounds.

Post-traumatic ulcers could be a subgroup of oedematous leg ulcers due to the same management approach, namely compression therapy, which should be assessed immediately after vascular/arterial causes have been ruled out. Hence, according to the present data, oedematous and post-traumatic ulcers accounted for 40.6% of all the ulcers and were treated with compression.

We found that the wound diagnostic process was good enough in the wound care team, as the diagnosis did not change for 75.5% of the patients who were referred by the team to specialist care. In the remaining 24.5%, in whom the diagnosis made by a specialist differed from the diagnosis made by the wound care team, the final

Protected by copyright, includ diagnosis was confirmed using diagnostic tools that were not available in the health centre. In these cases, however, the wound care team often had a default diagnosis to base the referral on, and specialist care then responded to this idea, most often leading to the correct treatment. Indeed, the referral was very useful for these patients. uses rela

On the other hand, there were ordinary delays in receiving a specialist evaluation, as the median delay was 21 days, where the largest differences were between the subgroups of patients living in institutions and those living at home. This might be explained by the advance care planning among nursing home residents. However, previous studies propose delays of less than 2 weeks in diagnosing arterial ulcers and DFUs to avoid leg losses.¹⁷ As a response to the challenge of timely referral to the ata . correct department for treatment there are globally several multidisciplinary wound clinics to have only one place to send a patient for a consultation.²⁶⁻²⁸

Previous studies suggest that diagnostic errors are often preceded by common symptoms, followed by common diagnoses.^{29 30} Studies have shown that the most frequent frequent is 'premature closure', meaning 'the tendency to stop considering other possibilities after reaching g and similar technologies a diagnosis'.^{31 32} In the UK, diagnosis-related and

Table 6 Diagnostic different	iation through the treatment pathway		
	The same diagnosis* throughout entire treatment pathway	The same diagnosis within primary care†	The same diagnosis by wound care team and specialist
1=the same	24	60	83
0=different	47	67	27
Total	71	127 [‡]	110
% of diagnosed§ ¶ patients	21.8	47.2	75.5
% of 197 (whole sample)	12.2	30.5	42.1

*Treatment pathway: primary care physician, wound care team (physician) and specialist care.

†Primary care physician and wound care team (physician).

‡Two patients were undiagnosed in the wound care team. They were of those of the 129 diagnosed in the primary care.

§One hundred and fifty-five patients visited a doctor in primary care before the appointment with the wound care team, but only 129 patients were diagnosed

¶One hundred and ten patients were diagnosed in specialist care.

đ

e

and

Protected by copyright, including for uses related to text and data mining. Al training, and similar technologies.	Enseignement Superieur (ABES) .	BMJ Open: first published as 10.1136/bmjopen-2022-062673 on 21 November 2022. Downloaded from http://bmjopen.bmj.com/ on June 10, 2025 at Agence Bibliographique de I
--	---------------------------------	---

Delays are calculated and analysed with the inclusion criterion 'wound appearance within 365 days prior to wound care team appointment'	ed with	the inclusion criterion w	vound appearance within 36	5 days prior to wound care te	am appointment'.		
	c	Wound appearance - first contact to healthcare	Wound appearance to first physician evaluation	Wound appearance to wound care team	Delay from first contact to wound care team (organisational delay within primary care)	Delay wound care team to specialist care	Mann-Whitney U
All patients	182	2 (0-14; 0 to 351; 351)	8 (1–32; 0 to 314; 314)	57 (33-101; 2 to 358; 356)	42 (22–80; 1 to 484; 483)	21 (7–52; –58 to 252; 414)	
Male	81	3 (0-24; 0 to 351; 351)	9 (1-37; 0 to 314; 314)	69 (37-111; 2 to 358; 356)*	44 (23-85; 2 to 484; 482)	23 (3-48; -58 to 235; 293)	*p=0.058
Female	101	1 (0–8; 0 to 295; 295)	8 (1–24; 0 to 295; 295)	54 (30–96; 2 to 306; 304)*	41 (22–76; 1 to 264; 263)	20 (8–58; 0 to 176; 176)	
Age under 65 years ¹	46	6 (1-27; 0 to 298; 298)*	9 (3–30; 0 to 142; 142)	62 (36-100; 11 to 320; 309)	37 (22–76; 6 to 382; 376)	28 (14–48; 1 to 182; 181)	*1 vs 2; p=0.005
Age 65–80 years ²	62	0 (0-14; 0 to 258; 258)*	10 (0–38; 0 to 314; 314)	61(41-106; 10 to 337; 327)	49 (25–88; 4 to 264; 260)	16 (2–50; 0 to 235; 235)	*1 vs 3; p=0.003
Age over 80 y ³	74	1 (0-8; 0 to 351; 351)*	7 (1–32; 0 to 295; 295)	53.5 (30–98; 2 to 358; 356)	40 (22-78; 1 to 484; 483)	16 (6–56; –58 to 167; 225)	
DM+	72	1 (0–15; 0 to 142; 142)	10 (1–37; 0 to 142; 142)	59 (36-102; 2 to 324; 322)	45 (26–75; 2 to 245; 243)	14 (3–48; 0 to 235; 235)	No statistical
DM-	110	2 (0-13; 0 to 351; 351)	7 (1–32; 0 to 314; 314)	56 (31-101; 4 to 358; 354)	40 (22-84; 1 to 482; 483)	26 (8-56; -58 to 167; 225)	Difference
Living at home	160	2 (0-15; 0 to 351; 351)	9 (1–33; 0 to 314; 314)	57 (33-102; 2 to 358; 356)	42 (22-83; 1 to 382; 381)	20 (5-48; 0 to 235; 235)*	*p=0.010
Living in institution	22	1 (0–8; 0 to 295; 295)	3 (0-14; 0 to 295; 295)	55 (34–86; 7 to 306; 299)	43 (24–72; 7 to 484; 477)	56 (16–143; –58 to 176; 234)*	
Walking; outdoors	134	3 (0-15; 0 to 351; 351)*	8 (1–27; 0 to 246; 246)	56 (33–97; 4 to 358; 354)	40 (22–76; 1 to 382; 381)	22 (6–56; –58 to 235; 293)	*p=0.047
Not-walking; staying indoors	48	0 (0–5; 0 to 298; 298)	9 (1–55; 0 to 314; 314)	71 (33-108; 2 to 337; 335)	54 (24–94; 2 to 484; 482)	16 (7–46; 0 to 182; 182)	
Delay before wound care team under 28 days	61	6 (0–20; 0 to 351; 351)*	7 (3–24; 0 to 295; 295)	26 (19–41; 2 to 358; 356)*	18 (12–22; 1 to 28; 27)	22 (4–42; 0 to 167; 167)	*p<0.001
Delay before wound care team over 28 days	121	0 (0–8; 0 to 258; 258)	9 (0–34; 0 to 314; 314)	73 (51–112; 29 to 337; 308)	70 (42–101; 29 to 484; 455)	20 (7–55; –58 to 235; 293)	
Unchanged diagnosis (PC-WT- spec.)	20	3 (2–12; 0 to 246; 246)	7 (3–19; 0 to 246; 246)	51 (32–80; 5 to 267; 262)	35 (20-74; 2 to 186; 184)	23 (8–45; 0 to 89; 89)	
Different diagnosis	43	1 (0-17; 0 to 258; 258)	8 (0-42; 0 to 314; 314)	71 (37–111; 4 to 337; 333)	57 (30–89;1 to 245; 244)	18 (7–98; 0 to 235; 235)	
The same diagnosis within primary care	, 55	2 (0-7; 0 to 246; 246)	3 (1–18; 0 to 246; 246)	52 (31–78; 5 to 267; 262)*	40 (20-71; 2 to 186; 184)	33 (9–56; 0 to 176; 176)	*p=0.042
Different diagnosis	61	1 (0-24; 0 to 258; 258)	17 (1–56; 0 to 314; 314)	73 (37–126; 4 to 337; 333)	53 (31–98; 1 to 245; 244)	15 (7–64; 0 to 235; 235)	
Age categories are 1 under 65 years, 2 DM, Diabetes mellitus; PC, Primary car	between e; spec, s	65-80 years, 3 over 80 years. pecialist care; WT, Wound car	There is a significancy in patient- e Team.	related delay in seeking health serv	Age categories are 1 under 65 years, 2 between 65-80 years, 3 over 80 years. There is a significancy in patient-related delay in seeking health services after onset of the wound between age group under 65 year older than 65 years. DM, Diabetes mellitus; PC, Primary care; specialist care; WT, Wound care Team.	group under 65 year older than 65	years.

assessment-related incidents were the highest causes of patient harm.²⁹ As a conclusion, the ability to use differential diagnostic methods is key when diagnosing wound aetiologies. There is always a danger of diagnosing a wound incorrectly, if possibilities to perform differential diagnostics are lacking.^{19 30}

As a solution, a broader range of differential diagnostic possibilities as regards the origin of the wound would probably help in the first evaluation and in avoiding diagnostic error and delay in the treatment.^{33 34} Our study shows that the problem for wound management lies in the primary care; that is, wounds that should be referred to multidisciplinary care are not recognised. The solution is continuous education of primary care physicians and nurses focusing on basic differential diagnostics of chronic wounds instead of wound management per se. Education is needed also to bring up the awareness of the triageremembering that also an acute wound may turn into a chronic wound which needs quick response and treatment. One practical tool to tackle these diagnostic challenges could be the use of checklists^{35–37} and digitalised wound diagnostic tools (³⁸, submitted for publication). It has been determined in other contexts that there are tools for avoiding fatal errors in differential diagnostics, such as existing guidelines and, to be regarded with a grain of salt, electronic aids in decision-making.³⁹⁻⁴¹

Limitations

The limitations of this study are related to the variety of aetiologies behind chronic wounds. There are no generally agreed-upon diagnostic codes to be used for chronic wounds, and the differentiation potential is enormous. Most often wounds are coded as merely a wound of unspecified aetiology (L97, L98), or they are S-coded, which refers to a traumatic wound. Therefore, it is difficult to define when a diagnosis is correct or not.

Outliers also constituted a limitation of the study, as we could not include them in the data analysis. Some patients in the material had suffered from a wound for several years. Despite this, they were referred to the wound care team when it was established in 2013. In our delay analyses, we tried to avoid this bias by selecting patients whose wounds had appeared less than 1 year prior to the appointment with the wound care team.

CONCLUSION

It seems that the diagnostic delay of wound patients occurs within primary care. It is an organisational delay and causes patient harm, as the patients are not receiving a timely and correct diagnosis and treatment. Infectious wounds seem to be easy to detect, but there is a risk of overdiagnosis, leading to an overuse of antibiotics. However, primary care physicians seem to pay little attention to distinguishing arterial insufficiency or DFU. BMJ Open: first published as 10.1136/bmjopen-2022-062673 on 21 November 2022. Downloaded from http://bmjopen.bmj.com/ on June 10, 2025 at Agence Bibliographique de I Enseignement Superieur (ABES) . Protected by copyright, including for uses related to text and data mining, Al training, and similar technologies.

The delay before seeing a primary care physician was not substantial, but the physicians' differential diagnostic approaches did not cover peripheral arterial disease or DFU. Consequently, the delay before being seen by the wound care team was over 1 month, which is a long time when treating DFU, especially those of vascular origin.

Based on our results, we propose that it is possible to arrange an optimal treatment pathway within a primary care setting, where a holistic wound care process is initiated, provided that there is organisational support, knowledge, skills and a multidisciplinary team available. It has been demonstrated that such an approach does not even require any additional resources, but rather a rearrangement of the patient care.^{16 42} We also suggest that the specialist care clinics could play a supportive role in the treatment of complex wounds, while the primary care system could take responsibility for the holistic wound care.

Ethics

Not applicable/no human participants included. This is a retrospective registry-based study. Data were anonymised before the authors assessed them for the purpose of the study. Due to the nature of the study ethical approval was not required for the study. Study was approved in the Institutional Review Board (IRB) of the Abdominal Centre, Helsinki University Hospital and in the IRB of the city of Helsinki.

Twitter Kirsi Isoherranen @Kirsilsoherran1

Contributors KA is the responsible guarantor of the content and has contributed to the designing of the study, conducting, collecting and analysing the data and reporting of the work described in the article. KI has contributed to the following parts of the study: designing of the study and reporting, reviewing and editing of the final manuscript. MV has contributed to the following parts in this study: designing of the study and reporting, reviewing and editing of the manuscript, reviewing and editing of the final manuscript. The corresponding author attests that all listed authors meet authorship criteria and that no others meeting the criteria have been omitted.

Funding KA has received a personal working grant from the University of Helsinki, The Finnish Wound Association and The Finnish Association for General Practice. This research received no specific grant from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests All authors have completed the ICMJE uniform disclosure form at Helsinki and declare: KI and MV have no support from any organisation for the submitted work, KA has received a working grant from the University of Helsinki, The Finnish Wound Association and The Finnish Association for General Practice; no financial relationships with any organisations that might have an interest in the submitted work in the previous 3 years; no other relationships or activities that could appear to have influenced the submitted work.

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 Not applicable.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available upon reasonable request.

Supplemental material This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content

includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation and adaptation or otherwise.

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

ORCID iD

Kirsti Ahmajärvi http://orcid.org/0000-0001-9106-5915

REFERENCES

- Lindholm C, Searle R. Wound management for the 21st century: combining effectiveness and efficiency. *Int Wound J* 2016;13 Suppl 2:5–15.
- 2 Phillips CJ, Humphreys I, Fletcher J, *et al.* Estimating the costs associated with the management of patients with chronic wounds using linked routine data. *Int Wound J* 2016;13:1193–7.
- 3 Guest JF, Ayoub N, McIlwraith T, et al. Health economic burden that different wound types impose on the UK's National health service. Int Wound J 2017;14:322–30.
- 4 Guest JF, Fuller GW, Vowden P. Cohort study evaluating the burden of wounds to the UK's National Health Service in 2017/2018: update from 2012/2013. *BMJ Open* 2020;10:e045253.
- 5 Sen CK. Human wound and its burden: updated 2020 compendium of estimates. Adv Wound Care 2021;10:281–92.
- 6 Herber OR, Schnepp W, Rieger MA. A systematic review on the impact of leg ulceration on patients' quality of life. *Health Qual Life Outcomes* 2007;5:44.
- 7 Green J, Jester R, McKinley R, *et al.* The impact of chronic venous leg ulcers: a systematic review. *J Wound Care* 2014;23:601–12.
- 8 Mooij MC, Huisman LC. Chronic leg ulcer: does a patient always get a correct diagnosis and adequate treatment? *Phlebology* 2016;31:68–73.
- 9 Dean S. Leg ulcers causes and management. *Aust Fam Physician* 2006;35:480–4.
- 10 Atkin L, Bućko Z, Conde Montero E, et al. Implementing timers: the race against hard-to-heal wounds. J Wound Care 2019;23:S1–50.
- 11 Garcia-Klepzig JL, Sánchez-Ríos JP, Manu C, et al. Perception of diabetic foot ulcers among general practitioners in four European countries: knowledge, skills and urgency. J Wound Care 2018;27:310–9.
- 12 Sánchez-Ríos JP, García-Klepzig JL, Manu C, et al. Referral of patients with diabetic foot ulcers in four European countries: patient follow-up after first GP visit. J Wound Care 2019;28:S4–14.
- 13 Sheikh A, Donaldson L, Dhingra-Kumar N. Diagnostic errors: technical series on safer primary care. Licence: CC BY-NC-SA 3.0 IGO2016. Geneva World Health Organization; 2016.
- 14 Kostopoulou O, Delaney BC, Munro CW. Diagnostic difficulty and error in primary care--a systematic review. *Fam Pract* 2008;25:400–13.
- 15 Irving G, Neves AL, Dambha-Miller H, et al. International variations in primary care physician consultation time: a systematic review of 67 countries. BMJ Open 2017;7:e017902.
- 16 Laakso M, Honkasalo M, Kiiski J, et al. Re-organizing inpatient care saves legs in patients with diabetic foot infections. *Diabetes Res Clin Pract* 2017;125:39–46.
- 17 Noronen K, Saarinen E, Albäck A, et al. Analysis of the elective treatment process for critical limb ischaemia with tissue loss: diabetic patients require rapid revascularisation. Eur J Vasc Endovasc Surg 2017;53:206–13.
- 18 Walker CM, Bunch FT, Cavros NG, et al. Multidisciplinary approach to the diagnosis and management of patients with peripheral arterial disease. *Clin Interv Aging* 2015;10:1147–53.

- 19 Eliassen A, Vandy F, McHugh J, et al. Marjolin's ulcer in a patient with chronic venous stasis. Ann Vasc Surg 2013;27:1182.e5–1182.e8.
- 20 Janowska A, Oranges T, Chiricozzi A, et al. Synergism of therapies after postoperative autograft failure in a patient with melanoma of the foot misdiagnosed as a pressure ulcer. Wounds 2018;30:E41–3.
- 21 Ahmajärvi KM, Isoherranen KM, Mäkelä A, et al. A change in the prevalence and the etiological factors of chronic wounds in Helsinki metropolitan area during 2008-2016. Int Wound J 2019;16:522–6.
- 22 Manu C, Lacopi E, Bouillet B, et al. Delayed referral of patients with diabetic foot ulcers across Europe: patterns between primary care and specialised units. J Wound Care 2018;27:186–92.
- 23 Öien RF, Forssell HW. Ulcer healing time and antibiotic treatment before and after the introduction of the registry of ulcer treatment: an improvement project in a national quality registry in Sweden. *BMJ Open* 2013;3:e003091.
- 24 Woo KY. Physicians' knowledge and attitudes in the management of wound infection. *Int Wound J* 2016;13:600–4.
- 25 Salava A, Isoherranen K. Rule of three I's: a mnemonic aid in undergraduate teaching of lower limb wounds. *J Wound Care* 2021;30:1030.
- 26 Gottrup F. A specialized wound-healing center concept: importance of a multidisciplinary department structure and surgical treatment facilities in the treatment of chronic wounds. *Am J Surg* 2004;187:S38–43.
- 27 de Leon J, Bohn GA, DiDomenico L, *et al.* Wound care centers: critical thinking and treatment strategies for wounds. *Wounds* 2016;28:S1–23.
- 28 Kim PJ, Attinger CE, Steinberg JS, et al. Building a multidisciplinary hospital-based wound care center: nuts and bolts. *Plast Reconstr Surg* 2016;138:241S–7.
- 29 Carson-Stevens A, Hibbert P, Williams H, et al. Characterising the nature of primary care patient safety incident reports in the England and wales national reporting and learning system: a mixed-methods agenda-setting study for general practice. *Health Serv Deliv Res* 2016;4:1–76.
- 30 Ely JW, Kaldjian LC, D'Alessandro DM. Diagnostic errors in primary care: lessons learned. J Am Board Fam Med 2012;25:87–97.
- 31 Norman GR, Eva KW. Diagnostic error and clinical Reasoning. Med Educ 2010;44:94–100.
- 32 Graber ML, Franklin N, Gordon R. Diagnostic error in internal medicine. *Arch Intern Med* 2005;165:1493–9.
- 33 Schiff GD, Hasan O, Kim S, et al. Diagnostic error in medicine: analysis of 583 physician-reported errors. Arch Intern Med 2009;169:1881–7.
- 34 Maude J. Differential diagnosis: the key to reducing diagnosis error, measuring diagnosis and a mechanism to reduce healthcare costs. *Diagnosis* 2014;1:107–9.
- 35 Snyder RJ, Jensen J, Applewhite AJ, *et al.* A standardized approach to evaluating lower extremity chronic wounds using a checklist. *Wounds* 2019;31:S29–44.
- 36 Kapp S, Santamaria N. The "self-treatment of wounds for venous leg ulcers checklist" (STOW-V Checklist V1.0): part 1-development, pilot and refinement of the checklist. *Int Wound J* 2022;19:705–13.
- 37 Wiseman JT, Fernandes-Taylor S, Gunter R, *et al.* Inter-rater agreement and checklist validation for postoperative wound assessment using smartphone images in vascular surgery. *J Vasc Surg Venous Lymphat Disord* 2016;4:320–8.
- 38 Kaari S, Vähätalo M, Ahmajärvi K. A digital wound management checklist to support clinical decision-making: a qualitative validation study
- 39 Bond WF, Schwartz LM, Weaver KR, et al. Differential diagnosis generators: an evaluation of currently available computer programs. J Gen Intern Med 2012;27:213–9.
- 40 Riches N, Panagioti M, Alam R, et al. The effectiveness of electronic differential diagnoses (DDX) generators: a systematic review and meta-analysis. PLoS One 2016;11:e0148991.
- 41 El-Kareh Ř, Hasan O, Schiff GD. Use of health information technology to reduce diagnostic errors. *BMJ Qual Saf* 2013;22 Suppl 2:ii40–51.
- 42 Sanders AP, Stoeldraaijers LGMC, Pero MWM, et al. Patient and professional delay in the referral trajectory of patients with diabetic foot ulcers. *Diabetes Res Clin Pract* 2013;102:105–11.