



Little is formally documented on the lived experience of Lyme disease - the state of diagnosis, treatment & care and the impact on the lives of those infected. Lyme Resource Centre (LRC) conducted an online survey to gather information on the impact of Lyme disease in the United Kingdom & Republic of Ireland.

111

responses from residents of Scotland who reported being diagnosed with Lyme disease by a qualified health professional; these responses were used for this analysis 86%

were bitten in Scotland of which

>80% bitten in Highlands & Islands

60%

were bitten in local fields, woodlands farmlands

20%

bitten in the garden

33%

were not aware of need to protect against tick bites prior to illness **42%** 

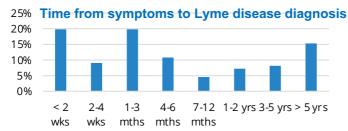
were not aware of Lyme disease prior to illness

## Scotland must prioritise raising awareness of tick bites and Lyme disease

In a letter of June 2019, Scotland's CMO directs practitioners to NICE guideline 95 on management of Lyme disease, which "aims to raise awareness of when Lyme disease should be suspected and ensure that people have prompt and consistent diagnosis and treatment".

From onset of symptoms

64%
respondents
had an
Erythema
Migrans
(EM) rash



71% not diagnosed within 4

weeks

**36%** 

not diagnosed for > 1 year 15%

not diagnosed for > **5 years** 

## Timely diagnosis of Lyme disease in Scotland must improve

NICE guideline 95 states that 'Prompt antibiotic treatment reduces the risk of further symptoms developing and increases the chance of complete recovery".

68%

had difficulty accessing treatment

93%

had antibiotics

Time from symptoms to first antibiotic

**57%** 

delay of > one month

44%

delay of > 3 months

19%

experienced delay of 2 years or more

**78%** 

did not fully recover after first antibiotic Of 68% who had difficult accessing treatment.

86%

say lack of disease awareness & expertise amongst health professionals was top barrier to treatment

## Prompt, effective treatment for Lyme disease must be available across Scotland

NICE guideline 95 also states that 'Symptoms can be more severe and long-lasting if treatment is delayed'

**72%** have ongoing symptoms

55% have been unwell for

> 2 years

Most commonly reported ongoing symptoms

32% have been unwell for > 6 years

Fatigue

Joint pains

Muscle aches

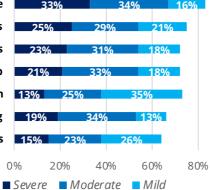
Impaired sleep

Anxiety, depression

Impaired thinking

199

ty, depression 13% 19% 19% 15% 15% 15%



"I was fiercely independent and strong before, now I'm vulnerable and rely on others good will and love. I lost career, family, friends, income, independence. It's hard to do anything now.....I wouldn't wish this on anyone."

Scotland must meet the needs of those with ongoing consequences of Lyme disease





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## **AWARENESS**

## Scotland must prioritise raising awareness of tick bites and Lyme disease

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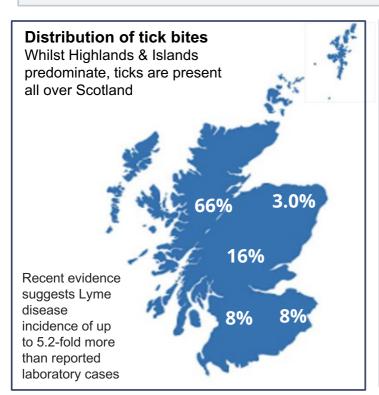
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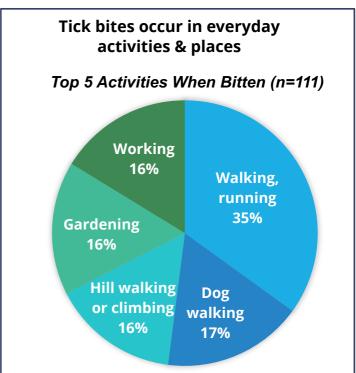
bitten in the garden

33%

were not aware of need to protect against tick bites prior to illness 42%

were not aware of Lyme disease prior to illness





Location when tick bite occurr	ed (n=111)
Local fields, woodlands	29%
Farmland, countryside	31%
In the garden	20%
Mountains, valleys, forests, wilds	41%
Indoors	2%

**30%** didn't recall a tick bite

Of those who found & removed a tick, only

40%

used a recognised tick removal tool

In a letter of June 2019, Scotland's CMO states - "Frontline healthcare practitioners have a key role not only in the early diagnosis and management of Lyme disease cases, but also in promoting awareness of ticks and tick borne infections amongst their patients. NHS Boards should promote awareness raising by frontline Healthcare Professionals, including those in Primary Care and GP practices."





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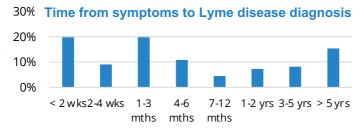
### **DIAGNOSIS**

## Timely diagnosis of Lyme disease in Scotland must improve

In a letter of June 2019, Scotland's CMO directs practitioners to NICE guidance on management of Lyme disease. NICE guidance "aims to raise awareness of when Lyme disease should be suspected and ensure that people have prompt and consistent diagnosis and treatment".

64% respondents had an Erythema Migrans

(EM) rash



From onset of symptoms

71%

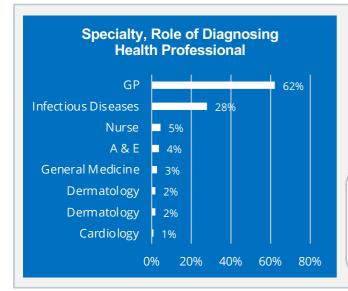
not
diagnosed not
within 4 diagnosed
weeks for > 1 year

15%

not
diagnosed
for > 5
years

The CMO's letter also stated that "the vast majority of Lyme disease cases can be diagnosed and treated within primary care without the need for specialist input".

Survey evidence suggest this isn't consistently happening



for **82%** 

respondents, self, family member, friend or colleague first suspected Lyme disease

"Multiple negative tests and misdiagnosis resulted in years of (me) being undiagnosed and treated." At least 16

# different medical specialties

involved with respondents Lyme related symptoms

"An earlier diagnosis; earlier treatment would have limited damage done to my body by Lyme related infections...."

50+

different
medical
conditions
suggested or
diagnosed before
Lyme disease
diagnosis

Myalgic-Encephalomyelitis

Hamoprous Mental-Health Anxiety Vitamin-Deficiency

Post-Viral-Fatigue-Syndrome

Mental-Hauth Anxiety Vitamin-Deficiency

Post-Viral-Fatigue-Syndrome

Morana Osteoarthritis

Proventi Depression

Migran Osteoarthritis

Rheumaniod-Arthritis

Rheumaniod-Arthritis

Rheumaniod-Vindranio

Migran Osteoarthritis

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National Migran Osteoarthritis

Rh





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### **TREATMENT**

## Prompt, effective treatment for Lyme disease must be available across Scotland

experienced

delay of 2 years

or more

NICE guideline 95 states that 'Prompt antibiotic treatment reduces the risk of further symptoms developing and increases the chance of complete recovery".

Survey evidence highlights missed opportunities for effective treatment.

**68%** 

had difficulty accessing treatment

93%

had antibiotics

Time from symptoms to first antibiotic

**57%** 

delay of > one month

44%

delay of > 3 months

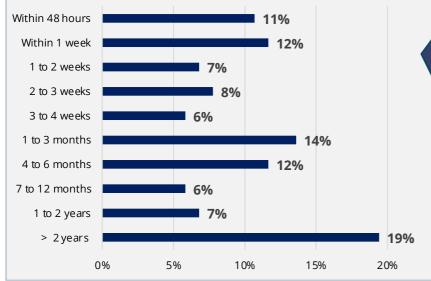
**78%** 

did not fully recover after first antibiotic Of 68% who had difficult accessing treatment.

86%

**say lack of disease awareness & expertise** amongst health professionals was **top barrier** to treatment





"Had to fight - got treatment from GP despite having the tick, rash was not bulls eye so not accepted as EM despite having symptoms.

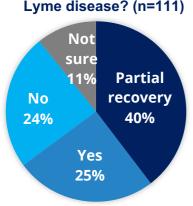
When still had symptoms following doxycycline was refused further treatment-- sent tick for testing it was positive. GP contacted ID who stated no further treatment.

MSP contacted and intervened on my behalf resulting in full

on my behalf resulting in full course of amoxicillin in line with NICE guidelines which did resolve symptoms."

Just **25%** have recovered from Lyme disease

# Have you now recovered from Lyme disease? (n=111)



Of 68% who had difficulty accessing treatment Key Reasons for Difficulty Accessing Treatment		
Lack of disease awareness / expertise of health professionals		
Misdiagnosis resulling in delayed diagnosis and Irealment		
Diagnosis excluded due to negative / inconclusive blood test*		
Restrictive treatment guidelines (*despite Lyme symptom	s) <b>49%</b>	
Cost of private treatment		
Distance to travel for treatment		
Stigma associated with Lyme disease	27%	

68%

have been unwell with Lyme disease for a year or more

32%

have been unwell for > 6 years

**72%** 

have ongoing symptoms



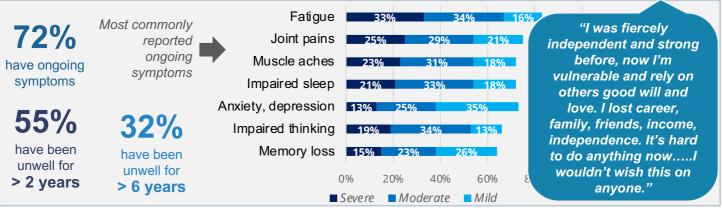


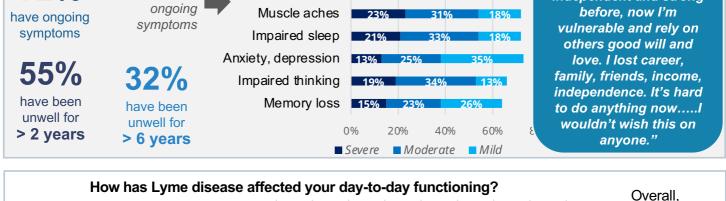
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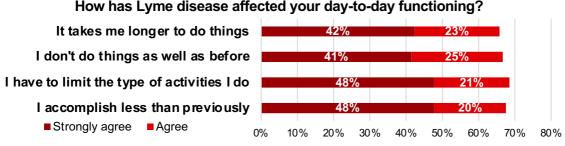
#### ONGOING NEEDS

## Scotland must meet the needs of those with ongoing consequences of Lyme disease

NICE guideline 95 states that 'Symptoms can be more severe and long-lasting if treatment is delayed'. Survey evidence suggest significant ongoing needs for many with Lyme disease.







**Strongly Agree** or Agree that Lyme disease has affected functioning

"I am not able to work, care for my young family, (even) function for a whole day."

Has having Lyme disease affected you in other ways?	
Affected social life and hobbies	68%
Financial loss	50%
Affected relationships with family and friends	49%
Career consequences	43%
Stigma associated with Lyme disease	32%
Loss of independence	32%
No consequences	23%
Educational consequences	19%

"It has ruined my life I am not the same person I used to be."

""It's hard to condense how much this has impacted me. I can no longer make plans that cannot be cancelled at short notice, I can't travel abroad...... have retained a handful of friends who understand what it is like to not have day to day control of my health. I can no longer undertake tasks such as shopping and have a husband who now acts as my carer."

#### What could have been better?

"More awareness of Lyme disease in Scotland and knowledge on how it should be treated and the fact that sometimes treatment doesn't work. Communication between GP and other specialists to be made aware of Lyme diagnosis."

#### About Lyme Resource Centre (LRC)







Lyme Resource Centre is a Scottish registered charity (SCIO: SC049151) focused on educating the public and healthcare professionals about Lyme disease & other tick-borne illnesses <a href="https://www.lymeresourcecentre.com">https://www.lymeresourcecentre.com</a>

#### **About the Survey**

From 1st February to 30th April 2024, LRC ran a **Lyme Disease Experience Survey** in UK and Republic of Ireland, to explore awareness of ticks and Lyme disease and *capture the lived experiences of people affected by Lyme disease*, including diagnosis, treatment and health outcomes.

The survey aimed to drive meaningful change for people who contract and live with Lyme disease and its consequences. The survey sought to capture responses from people who had fully recovered from Lyme disease as well as those with ongoing symptoms.

In researching the approach to the survey LRC consulted with the CEO of MyLymeData, a USA based, patient-powered research project with significant experience in surveys and patient data collection for Lyme disease.

Survey Monkey was chosen as the platform for an online survey due to its ease of use, functionality, and cost effectiveness.

#### **Survey Method**

To avoid concerns over data privacy and to encourage open, honest responses, the survey was conducted anonymously and participation was voluntary.

Participants were recruited via

- · open invitation on the LRC website
- social media channels (Facebook, Instagram, LinkedIn, X [Twitter])
- other organisations including Lyme Disease UK, TickTalk Ireland, Lyme Disease Alba, and others who shared and promoted the survey
- organisations whose members or followers use the outdoors professionally, educationally or socially (e.g. National Farmers Union, Forest Schools Association, Ramblers Association)
- individuals, who raised awareness of the survey with their own networks.

#### Survey Respondents

Out of 491 completed responses from UK and Republic of Ireland between 1st February 2024 and 30<sup>th</sup> April 2024, 475 reported a diagnosis of Lyme disease confirmed by a qualified health professional; 111 of these responses were from Ireland and formed the basis of this analysis. Respondents who could not report a diagnosis of Lyme disease by a qualified health professional (including self-diagnosis) were excluded.

Respondents were asked to confirm their answers were accurate to the best of their knowledge prior to submitting their survey response. The large number of completed responses (almost 500) should minimise the impact of any response inaccuracies.

There was no direct access to patients themselves or to any medical records.

We are mindful that some people who fully recovered from Lyme disease may have been unaware of the survey - we aimed to address this by promoting the survey via outdoor organisations.

People with undiagnosed Lyme disease are not represented in this survey.

#### References

- Scottish Government, Chief Medical Officer Letter re: Identification and Management of Lyme Disease Educational Resources, July 2019 https://www.publications.scot.nhs.uk/files/cmo-2019-07.pdf
- NICE Guideline [NG95] Lyme Disease https://www.nice.org.uk/guidance/ng95
- Incidence and management of Lyme disease: a Scottish general practice retrospective study, Mavin et al, BJGP Open, April 2024 <a href="https://pubmed.ncbi.nlm.nih.gov/38670577/">https://pubmed.ncbi.nlm.nih.gov/38670577/</a>