

# BMJ Open Liver blood marker testing in UK primary care: a UK wide cohort study, 2004–2016

Polly Scutt,<sup>1</sup> Lu Ban,<sup>1,2</sup> Tim Card,<sup>1,3</sup> Colin John Crooks,<sup>3,4</sup> Neil Guha,<sup>3,4</sup> Joe West,<sup>1,3</sup> Joanne R Morling <sup>1,3</sup>

**To cite:** Scutt P, Ban L, Card T, *et al*. Liver blood marker testing in UK primary care: a UK wide cohort study, 2004–2016. *BMJ Open* 2022;**12**:e058967. doi:10.1136/bmjopen-2021-058967

► 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-2021-058967>).

Received 04 November 2021  
Accepted 17 March 2022



© Author(s) (or their employer(s)) 2022. Re-use permitted under CC BY. Published by BMJ.

<sup>1</sup>Lifespan and Population Health, University of Nottingham, Nottingham, UK

<sup>2</sup>European Office, Evidera, London, UK

<sup>3</sup>NIHR Nottingham Biomedical Research Centre, Nottingham University Hospitals NHS Trust and University of Nottingham, Nottingham, UK

<sup>4</sup>Nottingham Digestive Diseases Centre (NDDC), School of Medicine, University of Nottingham, Nottingham, UK

## Correspondence to

Joanne R Morling;  
[joanne.morling@nottingham.ac.uk](mailto:joanne.morling@nottingham.ac.uk)

## ABSTRACT

**Objective** We aimed to determine (1) the temporal trends of liver enzyme testing in UK general practice and (2) how these vary among different subgroups at risk of chronic liver disease (CLD).

**Design** Retrospective cohort study.

**Setting** UK primary care database (Clinical Practice Research Datalink (CPRD)), 2004–2016.

**Participants** Patients aged 18 years or over, registered in the CPRD from 1 January 2004 to 31 December 2016.

**Outcome measures** The frequency of testing recorded within the study period in general practice was calculated for: alanine aminotransferase (ALT); aspartate aminotransferase (AST); gamma glutamyl transferase (GGT); alkaline phosphatase (ALP); bilirubin and platelets. Analyses were conducted in subgroups of patients at high risk of developing liver disease.

**Results** The study cohort included 2 912 066 individuals with median follow-up of 3.2 years. The proportion of patients with at least one measurement for ALT, ALP, bilirubin or platelet test gradually increased over the course of the study period and fell for AST and GGT. By 2016, the proportion of the population receiving one of more tests in that year was: platelet count 28.0%, ALP 26.2%, bilirubin 25.6%, ALT 23.7%, GGT 5.1% and AST 2.2%. Those patients with risk factors for CLD had higher proportions receiving liver marker assessments than those without risk factors.

**Conclusions** The striking finding that AST is now only measured in a fraction of the population has significant implications for routine guidance which frequently expects it. A more nuanced approach where non-invasive markers are targeted towards individuals with risk factors for CLD may be a solution.

## INTRODUCTION

In the UK, liver disease is a significant and growing burden on the National Health Service (NHS) and is the UK's third most common cause of premature mortality<sup>1</sup>; between 2015 and 2017 it caused 26 265 premature deaths in England alone.<sup>2</sup> It is also a significant source of healthcare inequity, with the median age of death differing by 9 years between the most and least deprived quintiles.<sup>3</sup> There has been a 400% increase in liver disease mortality in the population as a

## Strengths and limitations of this study

- ⇒ Sampling frame: a significant strength is the use of a large national dataset (>15 mill people).
- ⇒ Data quality: the dataset used (Clinical Practice Research Datalink) known to be representative of the UK population in terms of age, gender and geographical location with robust quality controls.
- ⇒ Data validity: previously validated code lists were used for the identification of subgroups.
- ⇒ A key limitation is the lack of information on the indication for testing or the resultant actions which limits interpretation to some degree.
- ⇒ Since this study only includes people who attend general practice, and some of the individuals at highest risk of chronic liver disease will not be attending. Therefore, we underestimate the proportions potentially identified in systematic testing was employed.

whole since 1970 and nearly 500% increases in mortality observed in working age populations over in this period.<sup>4</sup>

Three independent reports since 2014 have highlighted the need for the early detection of chronic liver disease (CLD) including the Chief Medical Officer report (2012),<sup>5</sup> the All-Party Parliamentary Hepatology Group Inquiry<sup>6</sup> and the Lancet commission,<sup>4</sup> in order to allow intervention and change the course of the disease. A number of organisations have now developed guidance advocating the use of non-invasive fibrosis markers in risk stratification.<sup>7–9</sup> Despite this, many existing community diagnostic pathways for detection and onward referral of suspected CLD are based on traditional liver enzyme tests which lack accuracy and result in delays to diagnosis.<sup>10</sup>

The optimal non-invasive fibrosis marker is yet to be determined, however, there are simple algorithms involving easily accessible measures such as aspartate aminotransferase (AST) and platelets that can be conducted in primary care, for example, aspartate to

platelet ratio index,<sup>11</sup> Fibrosis-4 score (FIB-4)<sup>12</sup> and CIRRhosis Using Standard tests (CIRRUS).<sup>13</sup> However, there is little understanding about how liver blood tests are currently used in UK general practice in order to support the implementation of changing practice and policy.

Given the rising prevalence of lifestyle related CLD and growing knowledge of non-invasive fibrosis measures, one could hypothesise that there should have been a shift away from traditional liver blood testing over time (shifting to non-invasive assessment). The aim of this study was to determine (1) the temporal trends of liver blood testing in UK general practice and (2) how these vary among different subgroups at risk of CLD.

## MATERIALS AND METHODS

### Data source

A population-based cohort study was conducted using the Clinical Practice Research Datalink (CPRD). The CPRD contains primary care data on 15.5 million people from 734 practices in the UK and is considered representative of the UK population.<sup>14</sup> Data are anonymised at patient and practice level and contain information on patient demographics, consultations, diagnoses, referrals and prescriptions. Clinical information is entered using READ codes which was a standard clinical terminology system used in the UK. For a subset of English practices (58% of UK CPRD practices), primary care data can be linked with the Hospital Episode Statistics (HES) dataset containing information for all hospital admissions.<sup>15 16</sup> The population for this study consists only of patients from these practices eligible for linkage with the HES dataset. This was a fully anonymised databased study not requiring ethical approval. This use of the data for this study was approved by the Independent Scientific Advisory Committee for CPRD and the Medicines and Healthcare products Regulatory Agency and assigned reference Protocol 19\_256.

### Study population

Patients aged 18 years or over, registered in the CPRD from 1 January 2004 to 31 December 2016, and having at least 1 day of registration with a practice eligible for linkage with the HES dataset were eligible for inclusion in the study. Patients with a diagnosis of CLD before the start of their follow-up period were excluded from the population. Patients were followed up starting at the latest of either the day after the date of current registration with their general practitioner (GP) practice, the start of the study period or the date the GP practice was labelled 'up to standard'. Follow-up ended at the earliest of either the date of death, date the patient transferred out of the GP practice, last date of data collection for the GP practice the patient is registered with, the end of the study period or the date of diagnosis with CLD in primary care (see online supplemental table S1).

## Outcomes

The frequency of testing recorded within the study period in general practice for the following liver blood tests was calculated: alanine aminotransferase (ALT); AST; gamma glutamyl transferase (GGT); alkaline phosphatase (ALP); bilirubin and platelet count. These markers were selected as being routinely used in UK primary care for the assessment of liver function. Abnormal results for each test were defined as: ALT result >50 (IU/L); AST result >40 (IU/L); ALP result >130 (IU/L); GGT result >50 (IU/L); bilirubin result >21 (IU/L); platelet result <150 (platelets/mcl).

## Subgroups

Analyses were conducted in the following subgroups of patients at high risk of developing CLD: presence of type 2 diabetes mellitus (T2DM) defined using READ codes (see online supplemental table S1); obesity defined as a body mass index >30 calculated using height and weight measures; use of alcohol defined as excessive use of alcohol using READ codes (see online supplemental table 1) or recorded >14 units per week alcohol consumption. For all subgroups, follow-up for an individual patient started at the date of diagnosis in primary care. Patients who were diagnosed with CLD within their follow-up period had their follow-up shortened to end 3 months before their date of diagnosis with CLD. An analysis of the subgroup of patients not included in any of these high-risk subgroups was also performed.

## Statistical analysis

Characteristics of the population were compared using  $\chi^2$  or Student's t-test as appropriate to the data distribution. The frequency of liver blood testing was presented as the proportion of patients with one or more tests out of the total eligible population over the study period. The frequency of abnormal test results was calculated and presented as the proportion of non-missing test results with an abnormal value. The number of tests performed per year on an individual was calculated by dividing the number of tests performed in the individual's follow-up period divided by the total length of their follow-up period. The proportion of patients with an AST test within 6 weeks following an abnormal ALT test result was calculated.

All analyses were conducted overall and stratified by sex, age group (18–29, 30–39, 40–49, 50–59, 60–69, 70–79, 80+ years) and calendar year. Analyses were performed on the whole study population and in the risk subgroups.

Analyses were performed using SAS V.9.4.

## Patient and public involvement

This study involved members of the Nottingham Digestive Diseases Biomedical Research Unit Patient Advisory Group at the following stages: research design and funding application, lay dissemination and discussion of results.

## RESULTS

### Characteristics

The study cohort included 2 912 066 individuals with follow-up during the years 2004–2016 (median follow-up

3.2 years, IQR 1.3–6.9). Of these, the predefined risk factor subgroups contained: 550 185 (19%) with obesity, 384 011 (13%) with excess alcohol use, 120 305 (4%) with T2DM and 2 235 938 (77%) with none of the three risk factors. One thousand four hundred and eighty individuals had all three risk factors.

The most frequently measured blood marker was platelet count, with 49% of patients having at least one platelet count measured during their follow-up period. The least commonly measured was the AST level with only 12% of patients having at least Exeter@123

in their follow-up. For all tests, the prevalence of testing increased with increasing age, with the highest proportion of patients being tested in the 70–80 year age category. Markers were more frequently measured in women and this difference was statistically significant for all markers ( $p < 0.0001$ ). Full details are given in [table 1](#).

Of those participants having tests the median number of tests undertaken each year was 1, however, some individuals had in excess of 100 of the same test per year. Platelet count was most likely to be tested more than once in an individual with the other liver markers being similar (for additional detail see online supplemental table S2).

### Prevalence of marker measurement over time

The proportion of patients in the study population with at least one measurement for ALT, ALP, bilirubin or platelet test gradually increased over the course of the study period (2004–2016) but conversely fell for AST and GGT markers ([figure 1](#) and [table 2](#)). By 2016, the proportion of the population receiving one or more of each test in that year was: platelet count 28.0%, ALP 26.2%, bilirubin 25.6%, ALT 23.7%, GGT 5.1% and AST 2.2%.

### Prevalence of abnormal measures

The proportion of all tests being measured as abnormal remained generally static over the study period ([figure 2](#)). Of the 3 922 529 (total number) of ALT test, 343 474 (8.8%) had an abnormal value. The first abnormal ALT test for each patient ( $n=1\ 60\ 191$ ) was paired with an AST test measurement within 6 weeks for 13 997 (8.7%). The proportion of measurements with abnormal values for all other markers was also low: AST (7.5%), ALP (7.9%), GGT (24.6%), bilirubin (4.7%), platelets (16.0%) and these proportions remained stable over the study period.

### Risk factor subgroup analyses

The prevalence of liver marker testing over time by the subgroups (no liver risk factors, excess alcohol consumption and/or obesity) showed similar trends to those for the whole population and are shown in online supplemental figure S1 and online supplemental table S3.

People with T2DM had a notably higher prevalence of testing for all markers (eg, in 2016 ALT measured in 68.8% of those with T2DM vs 15.3% and 21.9% of those with alcohol excess and obesity, respectively). However, the rates of decline in measurement of AST and GGT were also faster in those with diabetes than the other

groups; for AST falling from 24.3% in 2004 to 6.5% in 2016 versus 6.5% and 9.8% to 3.0% and 3.4% of those with alcohol excess and obesity, respectively; and for GGT falling from 28.6% in 2004 to 13.1% in 2016 versus 9.7% and 11.8% to 7.3% and 7.7% of those with alcohol excess and obesity, respectively.

People with no risk factors for liver disease had the lowest prevalence of liver marker testing for all markers, however, did still follow the same trends over time—increasing for ALT, ALP, bilirubin and platelets, and falling for AST and GGT.

## DISCUSSION

We found that while the majority of liver blood markers have shown increased rates of use in general practice over the past 10 years there was wide variation by both marker and subgroups of the population. Most notably, the use of AST has fallen to only 2% per annum among all general practice users.

The striking finding that AST is now only measured in a fraction of the population has significant implications for policy and practice. Major international guidelines, including American, European and British<sup>7 17 18</sup> all use non-invasive markers for investigating liver disease at a community level. AST is a critical component of FIB-4 which has been suggested as a first line test; to rule out significant disease. The absence of AST as a routinely collected marker presents a major barrier to the current implementation of pathways that attend to the aforementioned guidelines. This finding is consistent with other publications, where for example, in the assessment of liver fibrosis in individuals with a diagnosis of non-alcoholic fatty liver disease only 11% had the necessary measures to allow the assessment of FIB-4 in the UK (rising to 54% in Catalonia, Spain).<sup>19</sup> Furthermore, we found that <9% of abnormal ALT measurements also had an AST measured within a 6-week window.

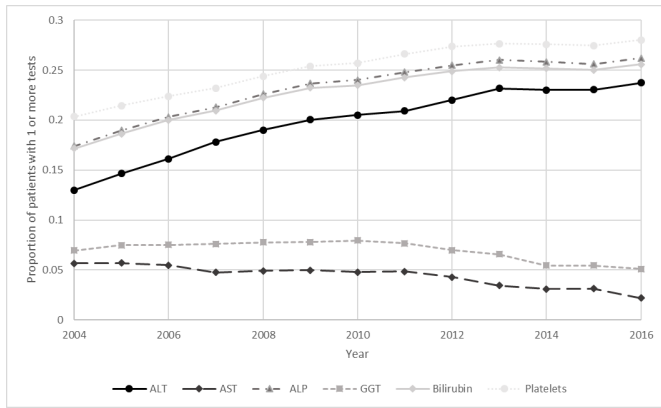
The decision to prioritise ALT measurement over AST may have been driven by a push for efficiency savings<sup>20</sup> with ALT being considered more valuable as it is more liver specific. However, AST may be a more sensitive indicator of chronic liver injury<sup>21–23</sup> especially when used as a ratio with ALT. In some regions an AST is automatically added if the ALT measure is abnormal to facilitate the AST/ALT ratio.<sup>24</sup> Over the 12-year period examined nearly 40% of the population had at least one ALT measurement. This far exceeds the proportion of the known population dying prematurely of liver disease (estimated at 26 265 premature deaths in England in 2015–2015), or the prevalence of recognised hepatic cirrhosis (estimated at 76.3 per 100 000 in 2001).<sup>26</sup> Though the level of CLD in the UK is not known it is unlikely therefore that these tests are all done in those who have it or even are at high risk, and we therefore have to question why they are being performed and the opportunity cost it represents. Existing evidence suggests they are more often measured as part of routine monitoring than for

**Table 1** Characteristics of participants (ever measurements)

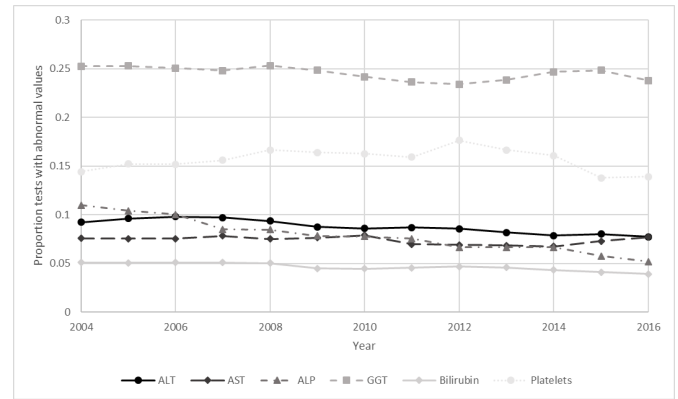
	Whole population (n)	ALT		AST		ALP		GGT		Bilirubin		Platelet count	
		1+tests	1+abnormal	1+tests	1+abnormal	1+tests	1+abnormal	1+tests	1+abnormal	1+tests	1+abnormal	1+tests	1+abnormal
All	n	1 112 879	160 191	284 274	33 743	1 261 596	140 433	459 754	124 475	1 246 003	99 633	1 414 798	301 127
	%	38.2	14.4	9.8	11.9	43.3	11.1	15.8	27.1	42.8	8.0	48.6	21.3
Sex													
Male	n	484 471	102 962	125 241	19 852	547 936	57 766	213 668	76 266	542 549	62 838	555 508	124 432
	%	35.1	21.3	9.1	15.9	39.7	10.5	15.5	35.7	39.3	11.6	40.3	22.4
Female	n	628 408	57 229	159 033	13 891	713 660	82 667	246 086	48 209	703 454	36 795	859 290	176 695
	%	41.0	9.1	10.4	8.7	46.5	11.6	16.1	19.6	45.9	5.2	56.0	20.6
Age group, years													
18–29	n	196 244	19 026	46 304	3311	230 084	15 495	70 498	7038	225 208	15 343	328 696	65 280
	%	17.6	9.7	4.1	7.2	20.6	6.7	6.3	10.0	20.1	6.8	29.4	19.9
30–39	n	246 015	35 067	61 081	6127	287 277	18 522	95 123	19 394	281 707	18 460	370 007	74 137
	%	24.6	14.3	6.1	10.0	28.7	6.4	9.5	20.4	28.2	6.6	37.0	20.0
40–49	n	266 922	42 817	66 307	7793	307 249	19 276	108 280	30 790	303 132	20 614	324 775	62 005
	%	37.1	16.0	9.2	11.8	42.8	6.3	15.1	28.4	42.2	6.8	45.2	19.1
50–59	n	221 016	36 113	54 770	7251	251 719	24 052	91 799	30 896	248 874	16 947	247 761	47 522
	%	46.4	16.3	11.5	13.2	52.9	9.6	19.3	33.7	52.3	6.8	52.0	19.2
60–69	n	187 210	24 880	46 118	5646	210 752	25 495	77 627	25 207	208 949	16 616	200 591	39 762
	%	57.9	13.3	14.3	12.2	65.2	12.1	24.0	32.5	64.7	8.0	62.1	19.8
70–79	n	133 592	11 776	33 557	3637	150 577	23 758	54 137	16 123	149 271	13 196	145 700	31 741
	%	63.0	8.8	15.8	10.8	71.0	15.8	25.5	29.8	70.3	8.8	68.7	21.8
80+	n	106 411	6586	25 860	2436	120 156	28 719	41 925	11 671	118 880	9989	123 721	28 849
	%	58.3	6.2	14.2	9.4	65.8	23.9	22.6	28.2	65.1	8.4	67.7	23.3

ALP, alkaline phosphatase; ALT, alanine aminotransferase; AST, aspartate aminotransferase; GGT, gamma glutamyl transferase.





**Figure 1** Prevalence of liver enzyme testing among adults over time. ALP, alkaline phosphatase; ALT, alanine aminotransferase; AST, aspartate aminotransferase; GGT, gamma glutamyl transferase.



**Figure 2** Prevalence of abnormal values of liver blood tests in adults over time. ALP alkaline phosphatase; ALT alanine aminotransferase; AST aspartate aminotransferase; GGT gamma glutamyl transferase.

CLD identification,<sup>27 28</sup> and that discontinuation of such drugs rarely results.<sup>29</sup> If all these abnormalities were to be followed up (in accordance with existing guidance) there would be significant implications for downstream services. This includes the cost of a full liver screen, liver ultrasound and onward consultation and investigation in secondary care (eg, national tariff for ultrasound scan £75.50, new patient consultant led hepatology outpatient appointment is £208.56.<sup>30</sup> Furthermore, there is growing evidence that in advanced liver disease many individuals have a normal ALT,<sup>10 31</sup> so the growth in use of this marker as a trigger for further assessment may still not identify liver disease.

A more nuanced approach where non-invasive markers are targeted towards individuals with risk factors for CLD

may be one solution. From a diagnostic perspective it increases the pre-test probability of having disease and indeed this approach has been shown to be cost effective regardless of choice of biomarker<sup>32 33</sup> and region studied.<sup>34</sup> Within CPRD those patients with risk factors for CLD, as expected, had higher proportions receiving liver markers assessment than those without risk factors. However, this was still very varied by 2016, with 70% of individuals with T2DM having an ALT measure that year, more than double those with obesity and nearly three times those with alcohol excess—with all three groups having similar proportions of abnormal results. While AST testing was more frequent among those with risk factors than in those without it was still very low (<8% in all groups). Therefore, from an implementation perspective

**Table 2** Annual frequency of testing per patient in those with at least one test

		ALT	AST	GGT	ALP	Bilirubin	Platelet count
Median (IQR) maximum number of tests		1 (1–2) 108	1 (1–1) 47	1 (1–1) 45	1 (1–2) 131	1 (1–2) 108	1 (1–2) 92
1	n	1 914 577	436 400	691 189	2 129 817	2 200 429	1 972 278
	%	74.2	75.5	75.6	70.2	74.1	60.1
2	n	457 080	99 123	155 168	610 628	528 607	849 020
	%	17.7	17.1	17.0	20.1	17.8	25.9
3	n	121 616	24 862	40 197	164 364	139 762	185 136
	%	4.7	4.3	4.4	5.4	4.7	5.6
4	n	39 173	8279	13 221	61 498	45 583	153 538
	%	1.5	1.4	1.4	2.0	1.5	4.7
5	n	15 569	3336	4960	23 572	18 166	29 952
	%	0.6	0.6	0.5	0.8	0.6	0.9
6–10	n	22 702	4909	7084	32 002	26 262	73 577
	%	0.9	0.8	0.8	1.1	0.9	2.2
11+	n	9349	1400	2610	11 234	10 248	18 286
	%	0.4	0.2	0.3	0.4	0.3	0.6

ALP, alkaline phosphatase; ALT, alanine aminotransferase; AST, aspartate aminotransferase; GGT, gamma glutamyl transferase.



it would make sense to focus efforts of obtaining AST and ALT in these groups, appreciating as step change in management is needed.

The strengths of this population approach are driven by the use of a dataset known to be broadly representative of the UK population in terms of age, gender and geographical location with robust quality controls<sup>14</sup> and also the use of validated code lists for subgroup identification.<sup>35</sup> It is therefore reasonable to assume that our findings regarding the level of testing overall and in subgroups are representative of what is happening in the UK. A key limitation is the lack of information on the indication for testing or the resultant actions which clearly limits interpretation to some degree. Additionally, since this study only includes people who attend the GP, some of the individuals at highest risk of CLD will not be attending, the estimates of the proportion of tests which would be abnormal with more systematic testing may be less accurate. A further issue is the lack of information to allow assessment of different liver blood testing systems, for example, which areas 'package' different blood tests together or where abnormal results automatically trigger additional tests.

In conclusion, large numbers of liver blood markers are being measured annually in UK primary care. At present, they are not suitable for risk stratifying high risk populations for CLD as the key element (AST) required to calculate non-invasive fibrosis markers is missing. However, the highest risk groups are receiving regular blood testing (69% of those with diabetes and 22% of those with obesity) so routine or opportunistic risk stratification could be feasible with limited additional expense to the NHS.

**Contributors** The authors confirm contribution to the paper as follows. JRM, JW, TC contributed to study conception and design. JRM contributed to data collection. JRM, PS, LB, JW, TC, CJC contributed to analysis and interpretation of results. JRM, PS, NG contributed to draft manuscript preparation. All authors reviewed the results and approved the final version of the manuscript. JRM is the guarantor with overall responsibility for the manuscript.

**Funding** This study was funded as part of a Medical Research Council Clinician Scientist award held by JRM (grant number MR/P008348/1).

**Competing interests** None declared.

**Patient and public involvement** Patients and/or the public were involved in the design, or conduct, or reporting, or dissemination plans of this research. Refer to the Methods section for further details.

**Patient consent for publication** Not applicable.

**Ethics approval** Not applicable.

**Provenance and peer review** Not commissioned; externally peer reviewed.

**Data availability statement** Data may be obtained from a third party and are not publicly available. This data was extracted following approval from the Clinical Practice Research Datalink (CPRD), <https://www.cprd.com/>. Applications for access should be directed directly to CPRD.

**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 4.0 Unported (CC BY 4.0) license, which permits others to copy, redistribute, remix, transform and build upon this work for any purpose, provided the original work is properly cited, a link to the licence is given, and indication of whether changes were made. See: <https://creativecommons.org/licenses/by/4.0/>.

#### ORCID iD

Joanne R Morling <http://orcid.org/0000-0003-0772-2893>

#### REFERENCES

- Public Health England. Liver disease: applying all our health [Internet]. Available: <https://www.gov.uk/government/publications/liver-disease-applying-all-our-health/liver-disease-applying-all-ourhealth>
- Public Health England. Liver Disease Profiles [Internet]. Available: <http://fingertips.phe.org.uk/profile/liver-disease> [Accessed February 23, 2016].
- Public Health England. The 2nd Atlas of variation in risk factors and Healthcare for liver disease in England. Reducing unwarranted variation to improve Health Outcomes and Value [Internet]. Available: <https://www.gov.uk/government/publications/liver-disease-applying-all-our-health/liver-disease-applying-all-ourhealth> [Accessed October 3, 2021].
- Williams R, Aspinall R, Bellis M, *et al*. Addressing liver disease in the UK: a blueprint for attaining excellence in health care and reducing premature mortality from lifestyle issues of excess consumption of alcohol, obesity, and viral hepatitis. *Lancet* 2014;384:1953–97 <http://linkinghub.elsevier.com/retrieve/pii/S0140673614618389>
- Davies S. Chief medical officer annual report 2011: volume 1, chapter 2C, P 163. Dep health Novemb 2012. Available: [https://www.gov.uk/government/uploads/system/uploads/attachment\\_data/file/141773/CMOAnnualReport2011Chapter2cpdf](https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/141773/CMOAnnualReport2011Chapter2cpdf)
- Liver Disease: Today's complacency, tomorrow's catastrophe. All-Party parliamentary hepatology group (APPHG) inquiry into improving outcomes in liver disease. Available: <https://www.iasorg.uk/uploads/APPHG20report20March20201420FINAL.pdf>
- Newsome PN, Cramb R, Davison SM, *et al*. Guidelines on the management of abnormal liver blood tests. *Gut* 2018;67:6–19.
- National Institute for Health and Care Excellence. NICE guidelines [NG49] Non-alcoholic fatty liver disease (NAFLD): assessment and management. Published online July 2016. Available: <https://www.nice.org.uk/guidance/ng49> [Accessed September 8, 2016].
- National Institute for Health and Care Excellence. NICE guidelines [NG50] Cirrhosis in over 16s: assessment and management. Published online July 2016. Available: <https://www.nice.org.uk/guidance/ng50> [Accessed September 8, 2016].
- Chalmers J, Wilkes E, Harris R. Development and implementation of a commissioned pathway for the identification and stratification of liver disease in the community. *Frontline Gastroenterol* 2019:Published online.
- Wai C-T, Greenon JK, Fontana RJ, *et al*. A simple noninvasive index can predict both significant fibrosis and cirrhosis in patients with chronic hepatitis C. *Hepatology* 2003;38:518–26.
- Vallet-Pichard A, Mallet V, Nalpas B, *et al*. FIB-4: an inexpensive and accurate marker of fibrosis in HCV infection. Comparison with liver biopsy and fibrotest. *Hepatology* 2007;46:32–6.
- Hydes T, Moore M, Stuart B, *et al*. Can routine blood tests be modelled to detect advanced liver disease in the community: model derivation and validation using UK primary and secondary care data. *BMJ Open* 2021;11:e044952.
- Herrett E, Gallagher AM, Bhaskaran K, *et al*. Data resource profile: clinical practice research Datalink (CPRD). *Int J Epidemiol* 2015;44:827–36.
- Herbert A, Wijlaars L, Zylbersztejn A, *et al*. Data resource profile: hospital episode statistics admitted patient care (HeS APC). *Int J Epidemiol* 2017;46:1093–1093i.
- Padmanabhan S, Carty L, Cameron E, *et al*. Approach to record linkage of primary care data from clinical practice research Datalink to other health-related patient data: overview and implications. *Eur J Epidemiol* 2019;34:91–9.
- Berzigotti A, Tsochatzis E, Boursier J. EASL clinical practice guidelines on non-invasive tests for evaluation of liver disease severity and prognosis – 2021 update. *J Hepatol. Published online* 2021.

- 18 Chalasani N, Younossi Z, Lavine JE, *et al.* The diagnosis and management of nonalcoholic fatty liver disease: practice guidance from the American association for the study of liver diseases. *Hepatology* 2018;67:328–57.
- 19 Alexander M, Loomis AK, Fairburn-Beech J, *et al.* Real-World data reveal a diagnostic gap in non-alcoholic fatty liver disease. *BMC Med* 2018;16:1–11.
- 20 Xu Q, Higgins T, Cembrowski GS. Limiting the testing of AST: a diagnostically nonspecific enzyme. *Am J Clin Pathol* 2015;144:423–6.
- 21 Daniel S, Ben-Menachem T, Vasudevan G, *et al.* Prospective evaluation of unexplained chronic liver transaminase abnormalities in asymptomatic and symptomatic patients. *Am J Gastroenterol* 1999;94:3010–4.
- 22 Whitehead MW, Hawkes ND, Hainsworth I, *et al.* A prospective study of the causes of notably raised aspartate aminotransferase of liver origin. *Gut* 1999;45:129–33.
- 23 Lilford RJ, Bentham L, Girling A, *et al.* Birmingham and Lambeth liver evaluation testing strategies (BALLETS): a prospective cohort study. *Health Technol Assess* 2013;17:i-xiv, 1–307.
- 24 Dillon JF, Miller MH, Robinson EM, *et al.* Intelligent liver function testing (iLFT): a trial of automated diagnosis and staging of liver disease in primary care. *J Hepatol* 2019;71:699–706.
- 25 Public Health Profiles. Accessed June 25, 2020. <https://fingertips.phe.org.uk/profile/liver-disease>
- 26 Fleming KM, Aithal GP, Soleymani-Dodaran M, *et al.* Incidence and prevalence of cirrhosis in the United Kingdom, 1992–2001: a general population-based study. *J Hepatol* 2008;49:732–8.
- 27 Standing HC, Jarvis H, Orr J, *et al.* GPs' experiences and perceptions of early detection of liver disease: a qualitative study in primary care. *Br J Gen Pract* 2018;68:e743–9.
- 28 Patel PJ, Banh X, Horsfall LU, *et al.* Underappreciation of non-alcoholic fatty liver disease by primary care clinicians: limited awareness of surrogate markers of fibrosis. *Intern Med J* 2018;48:144–51.
- 29 Nakafero G, Grainge MJ, Card T. What is the incidence of methotrexate or leflunomide discontinuation related to cytopenia. *liver enzyme elevation or kidney function decline? Rheumatology* 2021;(keab254).
- 30 UK Department of Health. National Cost Collection: National Schedule of NHS costs - Year 2019–20 - NHS trust and NHS foundation trusts. Available: <https://www.england.nhs.uk/national-cost-collection/>
- 31 Fracanzani AL, Valenti L, Bugianesi E, *et al.* Risk of severe liver disease in nonalcoholic fatty liver disease with normal aminotransferase levels: a role for insulin resistance and diabetes. *Hepatology* 2008;48:792–8.
- 32 Srivastava A, Jong S, Gola A, *et al.* Cost-comparison analysis of FIB-4, ELF and fibroscan in community pathways for non-alcoholic fatty liver disease. *BMC Gastroenterol* 2019;19:122.
- 33 Tanajewski L, Harris R, Harman DJ, *et al.* Economic evaluation of a community-based diagnostic pathway to stratify adults for non-alcoholic fatty liver disease: a Markov model informed by a feasibility study. *BMJ Open* 2017;7:e015659.
- 34 Serra-Burriel M, Graupera I, Torán P, *et al.* Transient elastography for screening of liver fibrosis: cost-effectiveness analysis from six prospective cohorts in Europe and Asia. *J Hepatol* 2019;71:1141–51.
- 35 Ratib S, West J, Crooks CJ, *et al.* Diagnosis of liver cirrhosis in England, a cohort study, 1998–2009: a comparison with cancer. *Am J Gastroenterol* 2014;109:190–8.