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Introduction

- Non-alcoholic fatty liver disease (NAFLD) and non-alcoholic steatohepatitis (NASH) have an estimated UK prevalence of 20-30% and 2-12% respectively.^{1,2}
- Challenges around diagnosis, such as non-specific symptoms and biopsy diagnosis, mean that NAFLD & NASH are often diagnosed in later stages of the disease.
- Identifying pathways to diagnosis may improve earlier diagnoses and identify cohorts suitable for inclusion in clinical trials.

Objectives

- To identify the pathways by which patients are diagnosed with NAFLD/NASH.
- To describe the patient characteristic, survival, laboratory testing and diagnostic testing of patients in the identified pathways.

Methods

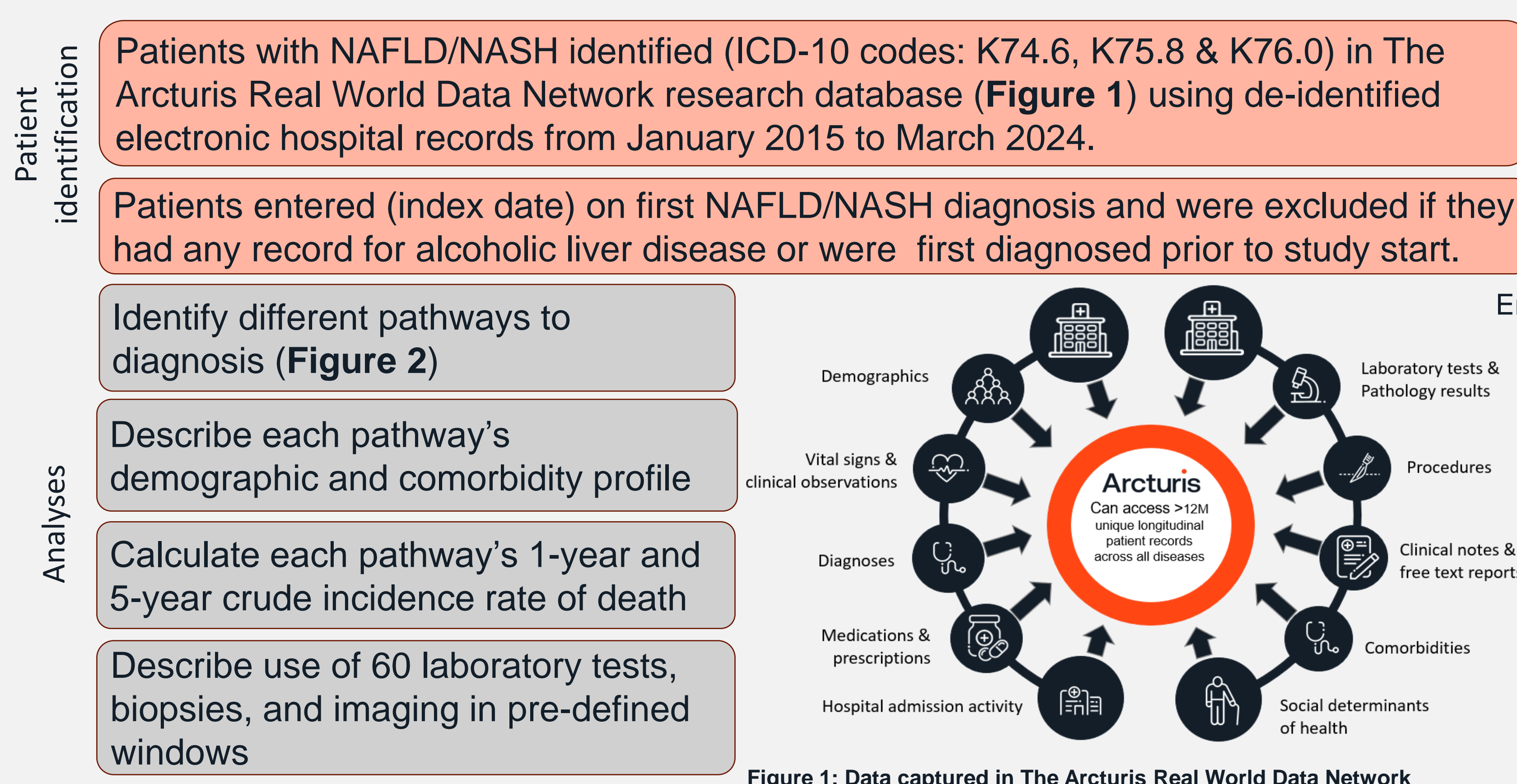


Figure 1: Data captured in The Arcturis Real World Data Network

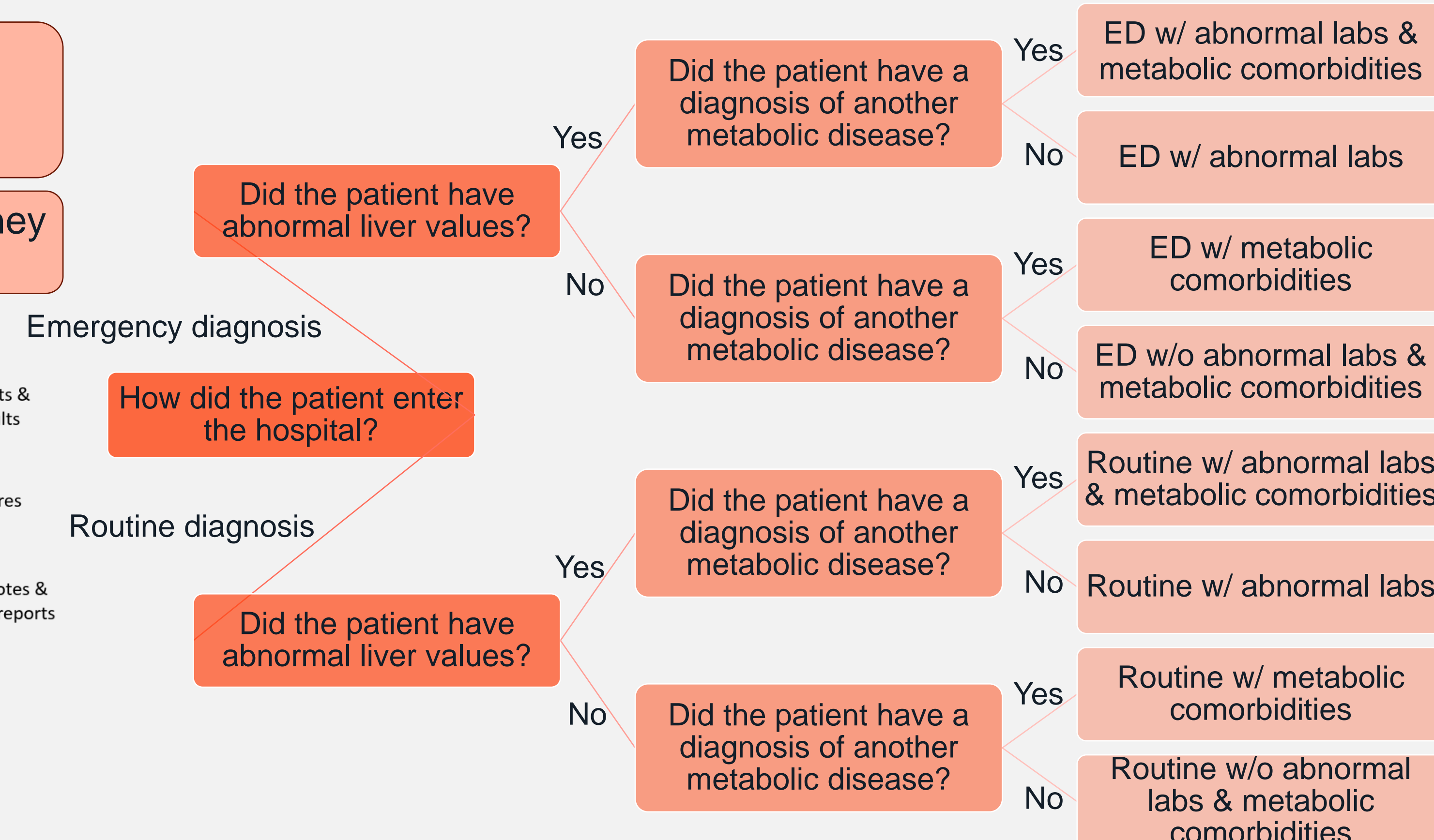


Figure 2: Decision tree depicting the eight pathways to diagnosis

Results - Characteristics

- Of 22,444 patients diagnosed with NAFLD/NASH, 8,855 had abnormal labs and metabolic comorbidities (59% emergency, 41% routine), 6,151 had just abnormal labs (58% emergency, 42% routine), 4,278 had just metabolic comorbidities (40% emergency, 60% routine), 3,160 had neither abnormal labs nor metabolic comorbidities (41% emergency, 59% routine).
- Patients without comorbidities were on average ~10 years younger than those with metabolic comorbidities.
- Other demographic characteristics between the pathways were broadly similar.
- The 1-year incidence rate varied between 284.78 (95% CI 191.75, 377.81) and 2,839.77 (95% CI 2,662.60, 3,016.94) per 10,000 person years, Figure 3, with emergency department admissions and patients with abnormal labs having higher rates.

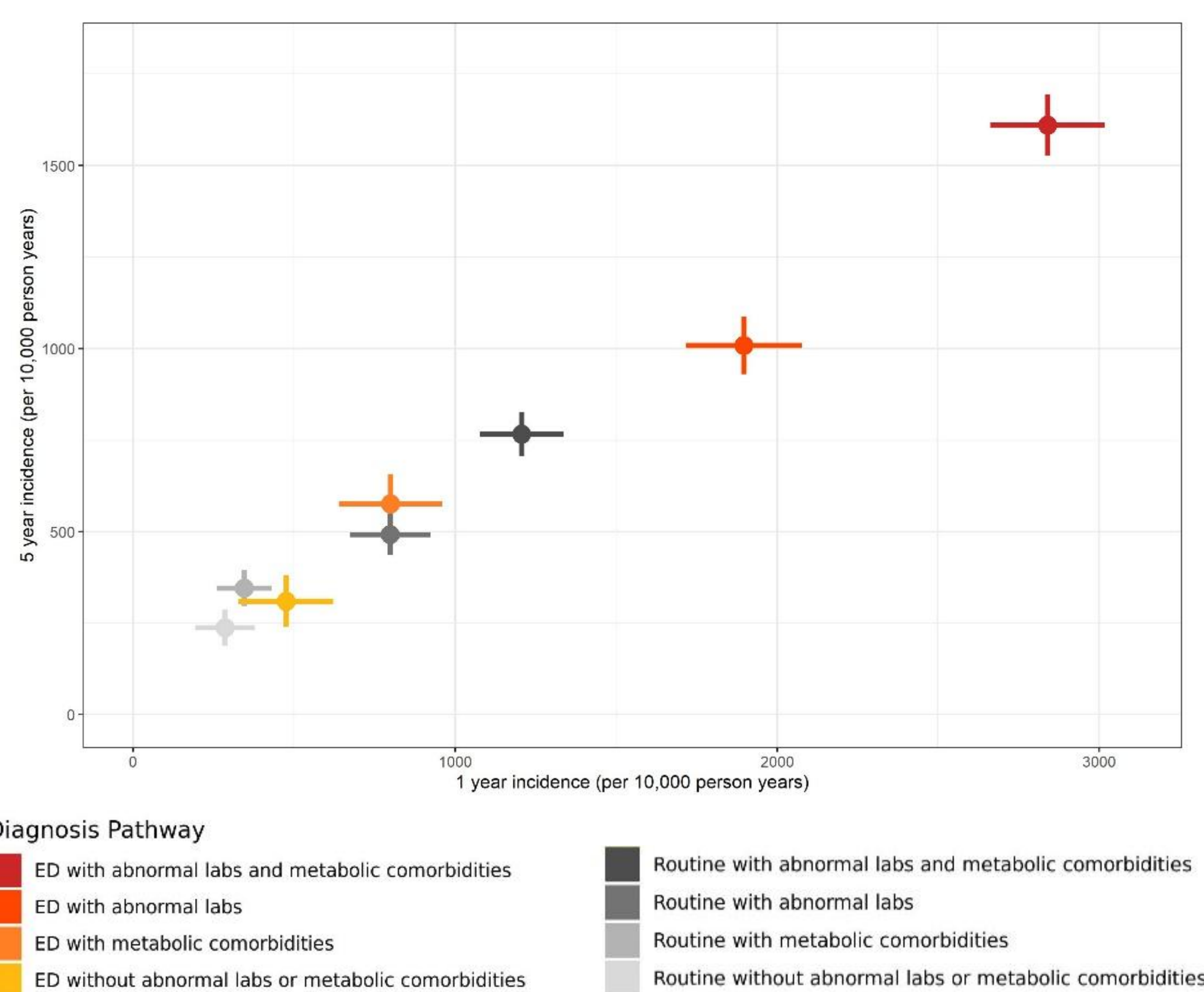


Figure 3: Overall survival incidence rate at 1 (x-axis) and 5 (y-axis) years after index date of all eight diagnosis pathways

Results - Testing

- 64.9% of liver biopsies were undertaken in the month before index date, range: 15.4% in the ED with metabolic comorbidities pathway to 81.3% in the routine pathway with no abnormal liver labs or metabolic comorbidities, Figure 4a.
- Abdominal ultrasounds occurred frequently prior to diagnosis, Figure 4b.
- Around diagnosis abdominal ultrasound were more common in the emergency department.
- Liver laboratory tests, such as albumin and ALP, were frequently tested throughout the windows, with peak testing around the index date, Figure 4c.
- Testing for GGT, secondary liver test, was higher in routine care diagnosed patients, Figure 4d.

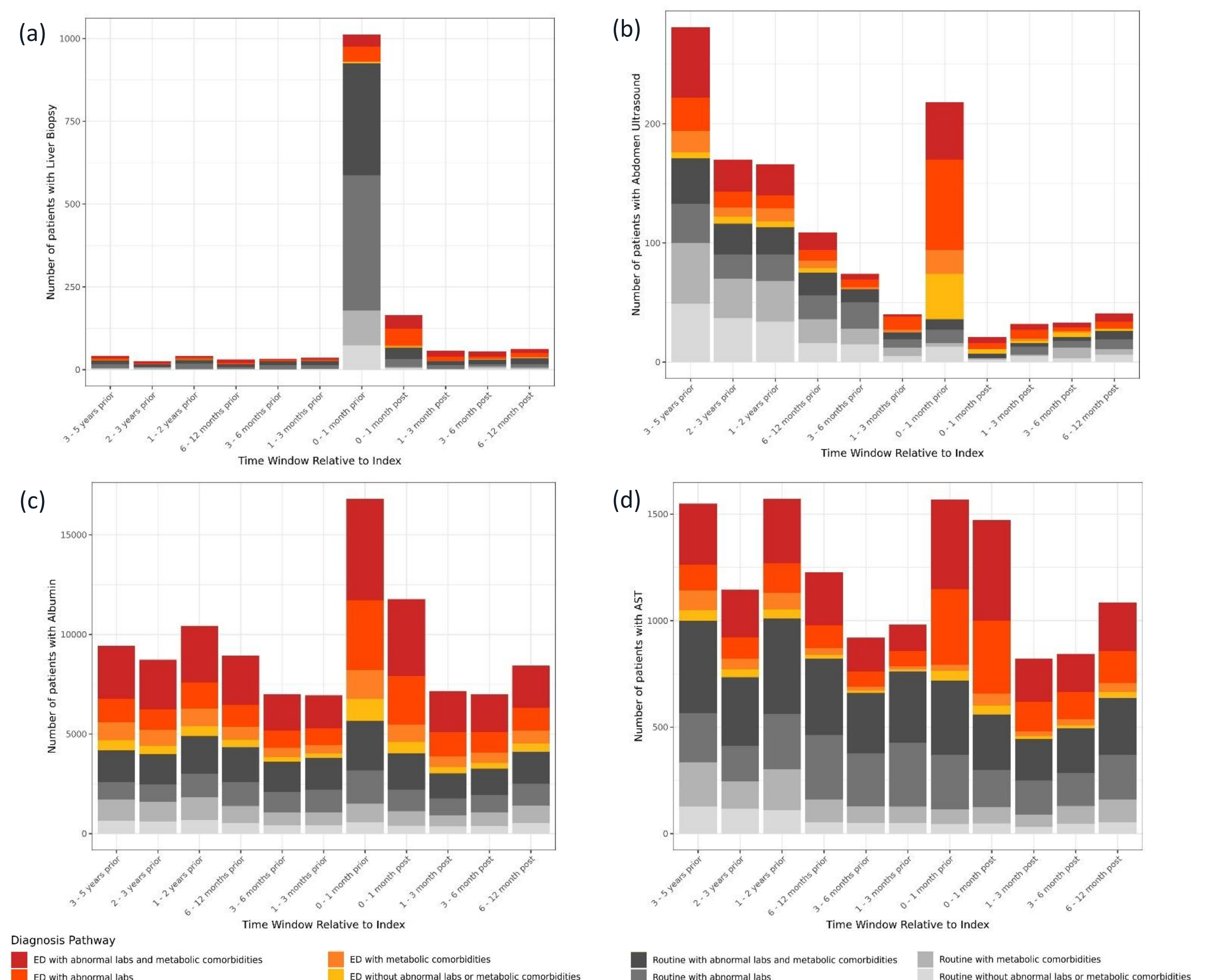


Figure 4: Stacked bar charts depicting number of patients with (a) liver biopsy, (b) abdominal ultrasound, (c) Albumin lab test and (d) GGT lab test

Conclusions

- We have identified and characterised 8 potential pathways to NAFLD/NASH diagnosis.
- Patient demographics were generally similar between pathways to diagnosis.
- Pathway to diagnosis impacted survival, likelihood of liver biopsy, imaging and liver laboratory tests.
- These pathways can help inform recruitment for clinical trials assessing novel therapies for NAFLD/NASH patients.

Acknowledgements

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References

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- Harrison, S. *et al.* Prevalence and stratification of NAFLD/NASH in a UK and US cohort using non-invasive multiparametric MRI, 2018.