REVIEW

Non-alcoholic fatty liver disease: a practical approach to diagnosis and staging

Jessica K Dyson, Quentin M Anstee, Stuart McPherson

ABSTRACT
Non-alcoholic fatty liver disease (NAFLD) is now the commonest cause of abnormal liver function tests (LFTs) in the UK with approximately a third of the population being affected. The exact prevalence is not known, but population studies from the USA and China using magnetic resonance spectroscopy estimate that approximately 30% of the general population have steatosis. It is a spectrum of disease ranging from simple steatosis, to non-alcoholic steatohepatitis (NASH), through to advanced fibrosis and cirrhosis. The majority have simple steatosis, but approximately 10–30% develop NASH and the development of NASH cirrhosis is associated with a poor long-term prognosis. Patients with NASH have increased liver-related and cardiovascular mortality. Many patients with NAFLD remain undiagnosed, and recognising those at risk is the first step. Clinicians overly rely on abnormal liver enzymes to identify patients with NAFLD, so patients with significant liver disease can be overlooked, potentially missing opportunities for intervention. Although liver biopsy is the gold standard method for diagnosing and staging NAFLD, the majority of patients can be effectively diagnosed non-invasively with tests that are routinely available in the clinic today. This review discusses a pragmatic approach to diagnosis and staging of NAFLD so that patients at the highest risk of liver-related complications can be identified.

INTRODUCTION
As a result of increasing rates of obesity, non-alcoholic fatty liver disease (NAFLD) is now the most common cause of abnormal liver function tests (LFTs) in the UK. NAFLD is present when >5% of hepatocytes are steatotic in patients who do not consume excessive alcohol consumption (<20 g/day for women and <30 g/day for men) and ranges in severity from simple steatosis (fat without significant hepatic inflammation or hepatocellular injury), to steatohepatitis (fat with hepatocellular injury and hepatic inflammation), through to advanced fibrosis and cirrhosis. Although the exact prevalence of NAFLD in the UK is not known, population studies from the USA and China using the most accurate imaging modality for liver fat, magnetic resonance spectroscopy, estimate that approximately 28–31% of the general population have steatosis and 8% have a raised alanine transaminase (ALT) due to NAFLD. NAFLD frequently coexists with other liver diseases such as hepatitis C, haemochromatosis and alcoholic liver disease and has been shown to cause more rapid disease progression. Fatty infiltration of the liver can also be secondary to treatment with steatogenic drugs such as tamoxifen, amiodarone and steroids.

NATURAL HISTORY OF NAFLD
Up to 90% of patients with NAFLD have simple steatosis, which carries a relatively benign prognosis, with no overall increase in mortality. However, approximately 10–30% have the potentially progressive form of NAFLD, non-alcoholic steatohepatitis (NASH), which is associated with hepatocellular injury and inflammation. Approximately 25–40% of patients with NASH will develop progressive liver fibrosis, ultimately resulting in cirrhosis in 20–30%. The development of cirrhosis due to NASH is associated with a poor long-term prognosis. The 10-year mortality rate is 20% for subjects with Child-Pugh A disease and 45% will decompensate within 10 years of diagnosis. In addition, subjects with NASH cirrhosis are at significant risk of developing hepatocellular carcinoma.
patients have normal-range ALT levels (males <40 IU/L and females <31 IU/L), and even if elevated, the ALT typically falls (and AST may rise) as fibrosis progresses to cirrhosis. ALT values do not correlate with histological findings and are unhelpful in both the diagnosis of NAFLD and determining disease severity. Clinicians overly rely on abnormal liver enzymes to identify patients with NAFLD, so patients with significant liver disease can be overlooked, potentially missing opportunities for intervention. It has been repeatedly shown that 70–80% of subjects with central obesity and 50–80% of patients with type 2 diabetes have evidence of NAFLD on imaging. Therefore, a new approach is needed to use metabolic risk factors to identify subjects with NAFLD/NASH rather than relying on liver enzyme abnormalities.

In patients with abnormal LFTs, alternative causes of liver disease (or cofactors) should be excluded, including alcohol excess, drug-induced liver injury, viral hepatitis, autoimmune liver disease, haemochromatosis, coeliac disease and Wilson’s disease (in patients <45 years old). Autoantibodies are also frequently detected at a low titre in subjects with NAFLD (antinuclear antibody (ANA) ≥1:160 and/or antismooth muscle antibody (ASMA) ≥1:40) and are usually associated with normal IgG levels and do not generally indicate autoimmune hepatitis. Raised ferritin levels are common in NAFLD and usually reflect underlying inflammatory activity or insulin resistance. A transferrin saturation <45% rules out haemochromatosis. If there is uncertainty about the diagnosis of NAFLD, then a liver biopsy should be considered.

The NAFLD Liver Fat Score is calculated using the presence of the metabolic syndrome, type 2 diabetes, fasting serum insulin, fasting serum AST and the AST/ALT ratio (AAR). In a cohort of 470 patients, a score greater than −0.640 predicted NAFLD with a sensitivity of 86% and specificity of 71%. Using cut-off scores of −1.413 and ≥1.257 gave 95% sensitivity for the prediction of NAFLD (with 52% and 51% specificity, respectively). However, this score does not distinguish between the different stages of NAFLD.

Imaging assessment of steatosis
Once suspected clinically, fatty infiltration of the liver can be confirmed with imaging. Ultrasonography is widely used as a first-line investigation for hepatic steatosis that provides a qualitative assessment of fatty infiltration of the liver. Ultrasound is very effective in diagnosing steatosis where >33% of hepatocytes are steatotic but can be unreliable with lesser degrees of steatosis. Therefore, the finding of a normal liver on ultrasound does not rule out mild fatty infiltration of the liver. Other imaging modalities such as CT or MRI can also detect hepatic steatosis, but they are not routinely used in the assessment of steatosis. MRI and proton magnetic resonance spectroscopy (MRS) are the most accurate non-invasive measures of steatosis.

### Table 1 Risk factors for NAFLD

<table>
<thead>
<tr>
<th>Risk Factor</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>Higher risk with increasing age</td>
</tr>
<tr>
<td>Metabolic syndrome (table 2)</td>
<td>70–90% of patients have NAFLD</td>
</tr>
<tr>
<td>Gender</td>
<td>Commoner in men</td>
</tr>
<tr>
<td>Certain ethnic groups</td>
<td>High risk in Hispanics</td>
</tr>
<tr>
<td>Dietary factors</td>
<td>High cholesterol and saturated fats</td>
</tr>
<tr>
<td></td>
<td>High fructose intake</td>
</tr>
<tr>
<td></td>
<td>Low carbohydrate</td>
</tr>
<tr>
<td></td>
<td>Caffeine may be protective</td>
</tr>
<tr>
<td>Obstructive sleep</td>
<td>Increased risk of hepatic fibrosis</td>
</tr>
<tr>
<td>apnoea</td>
<td>Patatin-like phospholipase domain-containing 3 (PNPLA3) gene</td>
</tr>
</tbody>
</table>

NAFLD, non-alcoholic fatty liver disease.
NAFLD.

Thresholds have not yet been clearly defined in patients with NAFLD, and the diagnostic features of the metabolic syndrome have been studied, and the diagnostic uncertainty or if non-invasive staging is adequate using non-invasive strategies. Liver biopsy provides an assessment of hepatic steatosis, hepatocellular injury, inflammation and fibrosis. The presence of hepatocyte ballooning degeneration in association with steatosis is the key histological feature that distinguishes NASH from simple steatosis.39 During apoptosis, caspases are activated and cleave various substrates, including cytokeratin-18 (CK-18), a major intermediate filament protein in hepatocytes. Hepatocyte apoptosis releases cleaved CK-18 fragments to the bloodstream that can be detected with an ELISA.40 Studies have demonstrated that the M30 antibody can identify patients with NASH with reasonable accuracy.41–43 Feldstein et al.42 showed that the level of plasma-cleaved CK-18 fragments was an

**Table 2 Features of the metabolic syndrome**

<table>
<thead>
<tr>
<th>Feature*</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Central obesity</td>
<td>Waist circumference: ≥94 cm for men and ≥80 cm for women (ethnicity specific measurements)</td>
</tr>
<tr>
<td>Impaired fasting glucose</td>
<td>&gt;5.6 mmol/L or on treatment</td>
</tr>
<tr>
<td>Hypertriglyceridemia</td>
<td>&gt;1.7 mmol/L or on treatment</td>
</tr>
<tr>
<td>Low HDL cholesterol</td>
<td>&lt;1.0 mmol/L for men or on treatment</td>
</tr>
<tr>
<td>Hypertension</td>
<td>&gt;135/85 mmHg or on treatment</td>
</tr>
</tbody>
</table>

**Table 3 NAFLD activity score (NAS)**

<table>
<thead>
<tr>
<th>Histological feature</th>
<th>Score</th>
<th>Category definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Steatosis</td>
<td>0</td>
<td>&lt;5%</td>
</tr>
<tr>
<td></td>
<td>1</td>
<td>5–33%</td>
</tr>
<tr>
<td></td>
<td>2</td>
<td>34–66%</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>&gt;66%</td>
</tr>
<tr>
<td>Plus</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hepatocyte ballooning</td>
<td>0</td>
<td>None</td>
</tr>
<tr>
<td></td>
<td>1</td>
<td>Few</td>
</tr>
<tr>
<td></td>
<td>2</td>
<td>Many</td>
</tr>
<tr>
<td>Inflammation</td>
<td>0</td>
<td>None</td>
</tr>
<tr>
<td></td>
<td>1</td>
<td>1–2 foci per ×20 field</td>
</tr>
<tr>
<td></td>
<td>2</td>
<td>2–4 foci per ×20 field</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>&gt;4 foci per ×20 field</td>
</tr>
<tr>
<td>NAS total 0–8</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fibrosis</td>
<td>0</td>
<td>No fibrosis</td>
</tr>
<tr>
<td></td>
<td>1a</td>
<td>Zone 3 mild perisinusoidal fibrosis</td>
</tr>
<tr>
<td></td>
<td>1b</td>
<td>Zone 3 moderate perisinusoidal fibrosis</td>
</tr>
<tr>
<td></td>
<td>1c</td>
<td>Periportal/portal fibrosis only</td>
</tr>
<tr>
<td></td>
<td>2</td>
<td>Zone 3 + periportal/portal fibrosis</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>Bridging fibrosis</td>
</tr>
<tr>
<td></td>
<td>4</td>
<td>Cirrhosis</td>
</tr>
</tbody>
</table>

**Fibrosis score 0–4**

A score of ≥5 with steatosis and hepatocyte ballooning is generally considered diagnostic of non-alcoholic steatohepatitis (NASH), but patients can still have NASH with lower NAS scores if steatosis and hepatocyte ballooning are present. NAFLD, non-alcoholic fatty liver disease.

### Liver biopsy for NAFLD

Although frequently not required for diagnosis, a liver biopsy is the definitive investigation for NAFLD and provides an assessment of hepatic steatosis, hepatocellular injury, inflammation and fibrosis. The presence of hepatocyte ballooning degeneration in association with steatosis is the key histological feature that distinguishes NASH from simple steatosis. The ‘NAFLD activity score’ (NAS) is the most widely used histological grading and staging system for NAFLD (table 3).35 The SAF score (encompassing an assessment of steatosis (S), activity (A) and fibrosis (F)) has been introduced more recently, which may be more accurate in identifying NASH.36 However, the majority of patients with NAFLD can be diagnosed and staged adequately using non-invasive strategies. Liver biopsy should be used for subjects where there is diagnostic uncertainty or if non-invasive staging is indeterminate.37

### Differentiating steatosis from steatohepatitis without liver biopsy

Knowledge of whether a patient has simple steatosis or NASH is very important prognostically. Subjects with simple steatosis have a good long-term prognosis with low rates of liver-related morbidity and mortality, and therefore do not require specific liver-related treatment. However, patients with NASH can progress to cirrhosis and therefore should be more actively managed to try and prevent disease progression. Unfortunately there is no widely available simple blood test or imaging modality that can differentiate simple steatosis from NASH. Clinically, the risk of steatohepatitis increases with the number of metabolic risk factors. Seventy per cent of centrally obese patients with hypertension and diabetes have steatohepatitis on liver biopsy.17 Therefore, until effective blood tests are available, metabolic risk factor profiling could be used to identify patients for further investigations. The search for an accurate biomarker of NASH is an active area of clinical research, and there have been some recent advances (see ref. 38 for a comprehensive review).

NASH is associated with increased apoptosis, so serum markers of apoptosis may be valuable in distinguishing NASH from simple steatosis.39 During apoptosis, caspases are activated and cleave various substrates, including cytokeratin-18 (CK-18), a major intermediate filament protein in hepatocytes. Hepatocyte apoptosis releases cleaved CK-18 fragments to the bloodstream that can be detected with an ELISA.40 Studies have demonstrated that the M30 antibody can identify patients with NASH with reasonable accuracy.41–43 Feldstein et al.42 showed that the level of plasma-cleaved CK-18 fragments was an

---


---

Downloaded from http://fg.bmj.com/ on September 7, 2016 - Published by group.bmj.com
independent predictor of NASH with an AUROC for a diagnosis of NASH compared with ‘nor’ or ‘border-
line’ NASH of 0.83. In that study, cleaved CK-18 frag-
ments were 71% sensitive and 85% specific for NASH at a cut-off of 279 U/L. Another CK-18 assay (M65 ELISA) is available that detects both cleaved and intact plasma CK-18 fragments and as a result is a marker of cell death by apoptosis and necrosis.\(^{44}\) Further validation is required to establish whether this assay is more accurate in differentiating NASH from simple steatosis.\(^{44}\) Although CK-18 is still being evaluated, some units are starting to incorporate this test into investigation algorithms for patients with NAFLD.\(^{45}\)

Another potentially interesting biomarker of NASH is the serum marker of matrix turnover, terminal peptide of procollagen III (PIIINP). In a study of 172 patients, PIIINP differentiated between simple stea-
tosis and NASH with reasonable accuracy in patients with both mild and advanced fibrosis (AUROC 0.77–
0.82 in patients with F0–2 fibrosis and 0.82–0.84 in patients with F0–3 fibrosis). Moreover, PIIINP was accurate in identifying patients with either NASH or advanced fibrosis (AUROC 0.85–0.87). While this needs to be validated, PIIINP might offer a useful test to identify the highest risk patients for further investigation.\(^{46}\)

### Non-invasive staging of liver fibrosis

Staging fibrosis is essential in all patients with NAFLD to identify subjects with advanced fibrosis who are at risk of liver-related complications.

#### Simple non-invasive markers of fibrosis

Hepatocellular dysfunction and portal hypertension result from advancing hepatic fibrosis. This may be reflected in ‘routine’ blood tests such as liver function tests (low albumin), full blood count (thrombocyto-
penia) and coagulation profile (prolonged prothrom-in time). These tests provide an indirect measure of fibrosis and are potentially appealing non-invasive

<table>
<thead>
<tr>
<th>Score</th>
<th>Indices</th>
<th>Calculation</th>
<th>Interpretation</th>
</tr>
</thead>
<tbody>
<tr>
<td>BARD score</td>
<td>BMI, AST/ALT ratio, T2DM</td>
<td>Weighted sum:</td>
<td>Validated in 827 patients with biopsy proven NAFLD fibrosis(^{47}) Score ≥2: Se 0.91, Sp 0.66, NPV 0.96 AUROC 0.81 for stage 3–4 fibrosis</td>
</tr>
<tr>
<td>NAFLD fibrosis score</td>
<td>Age, Hyperglycaemia, BMI, Platelet count, Albumin, AST/ALT ratio</td>
<td>Age×AST (IU/L)/platelet count (×109/L)×ALT (IU/L)</td>
<td>Validated in 733 patients with NAFLD(^{48}) AUROC 0.88 for stage 3–4 fibrosis</td>
</tr>
<tr>
<td>FIB-4 score</td>
<td>Age, AST, ALT</td>
<td>Age×AST (IU/L)/platelet count (×109/L)×(\sqrt{ALT}) (IU/L)</td>
<td>Validated in 541 patients with biopsy-proven NAFLD AUROC 0.80 for stage 3–4 fibrosis(^{49})</td>
</tr>
</tbody>
</table>

AAR, AST/ALT ratio; AUROC, area under receiver operating characteristic; BMI, body mass index; IFG, impaired fasting glucose; NAFLD, non-alcoholic fatty liver disease; NPV, negative predictive value; Se, sensitivity; Sp, specificity; T2DM, type 2 diabetes mellitus.

72% of patients scoring below 1.3 or above 2.67. Other studies have confirmed that the FIB-4 score is slightly better than other non-invasive tests in diagnosing advanced fibrosis in NAFLD, including in subjects with normal range ALT levels.20 32 33

All these simple non-invasive tests for fibrosis have good NPVs and can therefore exclude advanced fibrosis in patients with NAFLD who have low scores. As they can be calculated in all patients with routine blood tests, they offer an excellent method of identifying patients with mild disease who can be managed in primary care. However, the PPVs for these tests are modest (ranging from 27 to 79%), meaning that clinicians should consider further investigation to look for advanced fibrosis in patients with an intermediate or high score for their chosen test.20

Fibroscan

Fibrotic livers have reduced elasticity due to the deposition of fibrous tissue in the hepatic parenchyma. TE (Fibroscan) gives a ‘liver stiffness measurement’ (LSM) using pulsed-echo ultrasound as a surrogate marker of fibrosis.54 The LSM correlates well with the degree of hepatic fibrosis in a range of liver diseases, including NAFLD.54 55 In a study of 246 patients with biopsy-proven NAFLD, TE achieved high AUROCs for the detection of ≥stage 2 fibrosis, ≥stage 3 fibrosis and cirrhosis (0.84, 0.93 and 0.95, respectively) and performed better than a number of simple non-invasive scores in the staging of fibrosis.56 In that study, TE had a high NPV of 96% for ≥stage 3 fibrosis at a cut-off of 7.9 kPa but only modest PPV (52% at 7.9 kPa and 72% at 9.6 kPa). A low LSM reliably excludes advanced fibrosis, but the optimum cut-offs for clinical use are yet to be determined.

However, there are significant limitations to using TE in NAFLD. Results may be invalid in older patients (>52 years) and those with central obesity (BMI >35 kg/m²) or type 2 diabetes.57 For obese patients, the Fibroscan XL probe has been developed that is associated with fewer LSM failures (1.1% vs 16%) than the M probe and was accurate for the diagnosis of ≥F2 fibrosis and cirrhosis (AUROC 0.83 and 0.94, respectively).58 However, even with the XL probe, 10% of patients with a BMI >28 kg/m² have a difference of ≥2 fibrosis stages between TE and liver biopsy.59

Acoustic radiation force impulse

Another imaging technique that has the potential for the non-invasive assessment of fibrosis is acoustic radiation force impulse (ARFI). This technique uses conventional B-mode ultrasonography to generate an ultrasonic pulse and measure the response of the liver tissue as shear wave velocity.60 The median velocity measured by ARFI increases with the degree of fibrosis.61 In one study of 54 patients with NAFLD, the AUROC for the diagnosis of stage 3 or 4 fibrosis was 0.97.62 Although further validation is necessary, this technique is becoming increasingly available on ultrasound machines and has the potential to stage liver fibrosis at the time of liver ultrasound.

Commercial non-invasive fibrosis tests

The Enhanced Liver Fibrosis (ELF) test is a commercial panel of markers of matrix turnover: tissue
inhibitor of matrix metalloproteinase 1 (TIMP1), hyaluronic acid and PIIINP. This test performs slightly better than the NAFLD fibrosis score for diagnosing moderate fibrosis (AUROC 0.90 vs 0.86) and severe fibrosis (AUROC 0.93 vs 0.89), but combining the two tests gives an AUROC of 0.93 for moderate fibrosis and 0.98 for severe fibrosis.

Fibrotest (FT) is a commercial panel of biochemical markers of fibrosis that is widely used in France. In NAFLD, FT can diagnose advanced fibrosis with modest accuracy (AUROCs 0.75–0.86 for stage 2–4 fibrosis and 0.81–0.92 for stage 3–4 fibrosis). Using a FT cut-off of 0.30 gives a 90% NPV for advanced fibrosis (sensitivity 77%), and a FT cut-off of 0.70 had a 73% PPV for advanced fibrosis (specificity 98%). However, this test is not widely available in the UK.

A PRAGMATIC APPROACH TO DIAGNOSIS AND STAGING OF NAFLD IN CLINICAL PRACTICE

NAFLD is very common and the majority of patients have mild disease, but patients with advanced NASH need to be identified to offer treatment and surveillance for liver-related complications. With the current lack of a simple, widely available biomarker for NASH, a pragmatic diagnostic and staging approach is needed. One such approach for the investigation and assessment of disease severity in patients with NAFLD is shown in figure 1.

In brief, the first stage involves the identification of patients with NAFLD either with metabolic risk factor profiling, LFTs or imaging. If steatosis is confirmed and other causes of liver disease are excluded, a clinical diagnosis of NAFLD can be made. The second stage involves risk stratification to determine a patient’s stage of disease. This should be initially undertaken non-invasively with a locally available test (eg, FIB-4 score, NAFLD fibrosis score, TE, ARFI, CK-18). Patients who are identified as ‘low’ risk of NASH or advanced fibrosis can be managed in primary care with modification of their metabolic risk factors. Patients who are ‘indeterminate’ or ‘high’ risk should undergo further assessment (often requiring a liver biopsy) to determine the stage of disease. Risk stratification means patients can then be managed appropriately as will be discussed in ‘Non-alcoholic fatty liver disease: a practical approach to management’ by Dyson et al.

Machado et al. have recently proposed a similar algorithm for patients with NAFLD to guide when liver biopsy is needed. They used the NAFLD Fibrosis Score and TE to evaluate fibrosis and CK-18 fragments to evaluate NASH. The management pathway for patients would be very similar as with the algorithm we propose, but CK-18, and even TE, are not available in many centres, which is reflected in our algorithm.

CONCLUSIONS

NAFLD is a very common condition affecting approximately 30% of the population and can cause significant liver disease in a proportion of patients. Accurate diagnosis and staging is important in determining the appropriate long-term management for patients with NAFLD.

Key points

- Alanine transaminase (ALT) levels are a poor predictor of non-alcoholic fatty liver disease (NAFLD).
- Ultrasound is the first-line imaging test for patients with suspected steatosis (good accuracy if >30% of hepatocytes are steatotic).
- Liver fat decreases as fibrosis increases.
- Risk of NAFLD/NASH directly related to presence and severity of the metabolic syndrome.
- Simple steatosis carries benign prognosis.
- NASH carries poor prognosis with increased liver-related and cardiovascular mortality.

Aims:
- to identify individuals at risk of NAFLD
- to risk stratify patients with NAFLD
- to focus care on patients with NASH.

Contributors All authors contributed equally to this review.

Competing interests None.

Provenance and peer review Not commissioned; externally peer reviewed.

Open Access This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 3.0) 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 and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/3.0/

REFERENCES


Non-alcoholic fatty liver disease: a practical approach to diagnosis and staging

Jessica K Dyson, Quentin M Anstee and Stuart McPherson

*Frontline Gastroenterol* 2014 5: 211-218 originally published online December 24, 2013
doi: 10.1136/flgastro-2013-100403

Updated information and services can be found at:
http://fg.bmj.com/content/5/3/211

These include:

**References**
This article cites 75 articles, 12 of which you can access for free at:
http://fg.bmj.com/content/5/3/211#BIBL

**Open Access**
This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 3.0) 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 and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/3.0/

**Email alerting service**
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

**Topic Collections**
Articles on similar topics can be found in the following collections

Open access (44)

**Notes**

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/