1 Introduction

Myeloma is a malignancy of plasma cells. The normal role of the plasma cell is to secrete immunoglobulins (mainly immunoglobulin (Ig) A, IgG and IgM) to fight infection. Therefore immunoglobulins will be elevated as a result of infection or inflammation but usually in a polyclonal pattern (i.e. lots of different types of immunoglobulin are secreted). This is a normal physiological response to a reactive condition. When a clone of plasma cells turns malignant and expands it produces a single type of immunoglobulin – this is called a paraprotein, monoclonal protein or M protein. When a plasma cell neoplasm is suspected sending a sample for immunoglobulins will provide the clinician with the absolute levels of IgA, IgG and IgM as well as a result from serum protein electrophoresis and immunofixation which allows the identification of a monoclonal immunoglobulin versus polyclonal immunoglobulins.

2 Guideline scope

This guideline applies to adults who have had immunoglobulins and serum protein electrophoresis sent as part of a screen for myeloma. It will also help in the interpretation of immunoglobulins in general. It does not cover indications for immunoglobulin testing outside of the investigation of haematological malignancies.

3 Aim of the guideline

The aim of this document is to advise general practitioners and other hospital staff on the indications for testing immunoglobulins from a haematology perspective, how to monitor patients with a paraprotein and when to refer for haematology assessment.

4 When to send immunoglobulins

There are numerous indications for checking immunoglobulins which are not necessarily related to risk of plasma cell malignancy (for example immunoglobulins
can be part of a screen for liver disease). Therefore different specialties will have different guidelines on when to check immunoglobulins.

From a haematology perspective we usually suggest checking immunoglobulins in the investigation of myeloma or lymphoma. Myeloma is the most common malignancy that produces a monoclonal immunoglobulin. Patients with lymphoma and amyloidosis may also have a monoclonal immunoglobulin. From a haematology perspective we suggest sending immunoglobulins if there is clinical suspicion of a plasma cell disorder or:

- Unexplained hypercalcaemia
- Unexplained renal failure after standard investigations
- Unexplained anaemia (normal ferritin, B12, folate)
- Unexplained neuropathy
- Unexplained bone pains
- Unexplained elevated total protein or erythrocyte sedimentation rate (ESR)
- Bone lesions suspicious for myeloma on imaging

Screening otherwise healthy patients for a paraprotein is not advised.

5 What does the test include?

When immunoglobulins are requested the result will include the absolute level of IgA, IgG and IgM. The laboratory will also perform serum protein electrophoresis to look for a paraprotein. A high level of immunoglobulin does not necessarily mean there is a paraprotein as immunoglobulins are most commonly increased because of inflammation or infection. Similarly normal immunoglobulin levels do not exclude a small paraprotein.

Serum free light chains (SFLC) have replaced urine protein electrophoresis (also known as urine Bence Jones proteins) for the detection of light chain only myeloma. This is reflexed by the laboratory when there are relevant clinical details or there is a particular pattern on serum protein electrophoresis. If the suspicion of myeloma or plasma cell dyscrasia is high please request both serum protein electrophoresis and SFLC. SFLC are not required when the clinical suspicion is low or if the test is being done for high total protein or high ESR. It would be very unusual to have myeloma with normal immunoglobulin levels and normal serum protein electrophoresis.
6 What causes a paraprotein?

The most common causes of paraprotein are:

1) Monoclonal gammopathy of uncertain significance
   - Benign condition that is common in the over 70s (2-4% of people over the age of 70)
   - No features of myeloma or other plasma cell dyscrasia
   - Paraprotein (not total immunoglobulin) should be less than 30 g/L but mostly they will be less than 15 g/L
   - The SFLC ratio is often normal or only mildly abnormal
   - All cases of myeloma progress from MGUS but MGUS only rarely progresses to myeloma
   - MGUS prognosis can be calculated here: https://qxmd.com/calculate/calculator_148/mgus-prognosis
   - The risk of progression varies from 2% at 20 years to 58% at 20 years depending on level and type of immunoglobulin.

2) Myeloma
   - Malignant disorder of plasma cells resulting in production of monoclonal immunoglobulin
   - Patients are often symptomatic with anaemia, hypercalcaemia, renal failure, or bone disease
   - The paraprotein is usually IgG or IgA
   - Treated with chemotherapy and immunomodulatory agents

3) Plasmacytoma
   - A collection of malignant plasma cells in one area – usually a bone or soft tissue mass. Causes pain, swelling or fracture.
   - Treated with radiotherapy

4) Lymphoma – especially lymphoplasmacytic lymphoma (Waldenström’s macroglobulinaemia)
   - Causes anaemia, lymphadenopathy and B symptoms
   - The paraprotein is usually IgM

5) Amyloidosis
   - Systemic disorder which can result in neuropathy, cardiac failure, bleeding, gastro-intestinal problems, autonomic dysfunction, renal failure, nephrotic syndrome, arrhythmia, carpal tunnel syndrome, liver failure and macroglossia.
   - The paraprotein can be small
7 Flow chart if find a paraprotein

Paraprotein detected

Check FBC, U&E, bone profile, SFLC and urinalysis

Assess the patient clinically for bone pains, neuropathy, infections, B symptoms, hyperviscosity, heart failure and bleeding. Examine for lymphadenopathy and splenomegaly.

Total paraprotein (not total immunoglobulin level) is
• ≥15 g/L for IgG paraproteins
• ≥10 g/L for IgA or IgM paraproteins
Or the SFLC ratio is <0.125 or >8 or an individual light chain is above 500 mg/L.
Or the patient has a IgD or IgE paraprotein (any value)

Yes

Haematology referral (via 2WW if fulfils criteria). If unclear discuss via Advice and Guidance. If super urgent (cannot wait 24-48 hours) discuss with SpR on call*

No

Repeat immunoglobulins, SFLC, FBC, U&E, bone profile in four months.

Calculate the risk of progression to myeloma using https://qmrd.com/calculator/calculator_148/mgu-prognosis

Patients should be given written information – suitable organisations include Myeloma UK, Bloodwise or Macmillan. Patients should be encouraged to report any signs of myeloma or plasma cell dyscrasia.

Yes

Symptoms?***
• Unexplained hypercalcaemia
• Unexplained renal failure after standard investigations
• Unexplained anaemia (normal ferritin, B12, folate)
• Unexplained neuropathy
• Unexplained bone pains or osteoporotic crush fractures or lytic lesions on imaging
• Features of amyloidosis (e.g. proteinuria, cardiac failure, neuropathy)
• Neutropenia <1.2 x 10⁹/L
• Thrombocytopenia <100 x 10⁹/L
• If the patient is under 40
• Symptomatic hyperviscosity, e.g. visual disturbance, neurological symptoms, bleeding
• B symptoms
• Lymphadenopathy/splenomegaly

No

Paraprotein is stable. Repeat every six months for two years. If stable for two years then can reduce frequency of monitoring to annually. If there are concerns or uncertainties please discuss via Advice and Guidance. Some patients may have numerous comorbidities or frailty and therefore continued monitoring may not be appropriate; especially if risk of progression is low.

Yes

No

Total paraprotein (not total immunoglobulin level) is
• ≥15 g/L for IgG paraproteins
• ≥10 g/L for IgA or IgM paraproteins
Or the SFLC ratio is <0.125 or >8 or an individual light chain is above 500 mg/L.
Or if paraprotein has increased by 5 g/L or by 25%
Or if concerning symptoms** (see boxes above)

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8 No paraprotein found but results abnormal

**Hypogammaglobulinaemia**
One or more immunoglobulin class reduced below the reference range. In adults, please check SFLC for light chain only myeloma. If SFLC ratio is normal then look for clinical signs of lymphoma and check for lymphocytosis. If none, evaluate for acquired or congenital immunodeficiency. This may require discussion with an immunologist. Please note that mild decreases in IgM are common in elderly patients and that selective IgA deficiency is seen in 1 in 600 people and is generally an asymptomatic condition. In children, this test would only usually be requested by a specialist haematologist or immunologist, but please seek advice if there are concerns.

**Hypergammaglobulinaemia**
One or more immunoglobulin class increased above the reference range. The report may say ‘increased gamma zone’ or the serum protein electrophoresis will be normal. This means there is a polyclonal rise in immunoglobulins which can be seen in reactive cases e.g. infection, inflammation, malignancy or liver disease. Isolated rises in IgA are common in the elderly. Investigation of the underlying cause may be warranted depending on symptoms and clinical suspicion. This sort of result is not indicative of myeloma and does not warrant a haematology referral.

**Multiple banding**
This is often seen in inflammatory conditions. Repeating the result in four to six months to ensure no larger band appears is recommended. If multiple banding is still present investigate for inflammatory condition.

**Possible band**
Repeating the result in four to six months to ensure no larger band appears is recommended. If a possible band is persistent then follow MGUS flow chart.

9 Interpretation of SFLC when there is no paraprotein

It is normal to have more serum kappa light chains than lambda. Light chains increase in the serum in renal failure due to under excretion and in inflammation due to overproduction. Although the individual values are important the ratio between the kappa and lambda light chains is more important because if the individual light chains are proportionally increased then the ratio will be normal.

The normal ratio is 0.26 -1.65 if the eGFR is ≥60 mL/min and 0.37-3.1 if the eGFR <60 mL/min. If it is outside of these ranges and the:

- Ratio is <0.125 or >8 or an individual light chain is above 500 mg/L – refer to haematology as per paraprotein flow chart
- Ratio 0.125-0.2 or 5-8 – probable light chain MGUS – follow paraprotein flow chart for monitoring
- Abnormal but between 0.2 and 5 – minor abnormality of SFLC. In presence of normal immunoglobulin levels myeloma very unlikely. Likely causes include inflammation or abnormal renal function. Suggest discuss via Advice and Guidance.
10 Further advice

If further advice is needed then please contact a haematologist via the Advice and Guidance system. If the patient is known to a haematologist please write to the clinician directly. In an emergency please discuss with the on call specialist registrar. Please don’t phone the on call specialist registrar for day-to-day routine queries as they are very unlikely to know the patient and may not have access to all the available test results.

11 Training, Implementation, Resource Implications

Education about management of paraproteins and myeloma is provided for specialty trainees in haematology as part of their regional training programme.

12 Evidence Review and Evaluation

The evidence was reviewed and evaluated by Dr Andrew McGregor, Consultant Haematologist. The guideline was written by Dr Andrew McGregor and was reviewed and amended by Dr Brigit Greystoke, Consultant Haematologist, Professor Graham Jackson, Consultant Haematologist, Dr Suzanne Elcombe, Consultant Immunologist and Ashleigh Rainey, Healthcare Scientist.