KISS: Myeloma

Based on BJGP2018;e586 & NICE 2018, NG35 & NICE-CKS 2016

Just remind me...

- **Myeloma** is caused by a proliferation of monoclonal plasma cells in the bone marrow which secrete immunoglobulins, known as M proteins or paraproteins
- These paraproteins increase plasma viscosity and cause renal damage, and the proliferation of plasma cells leads to bone marrow suppression and can cause hypercalcaemia
- The median age of diagnosis is 70; only 15% of patients are aged under 60
- The typical features of myeloma give rise to symptoms of fatigue, bone and MSK pains, headache, nausea, recurrent infections etc
- Patients often present late with hypercalcaemia, pathological fractures or renal failure
- Prognosis is variable, with survival times varying from a few weeks to 20 years; most respond to initial treatment and have a period of stability before relapse after 2 to 5 years; younger fitter patients who have high dose therapy and stem cell transplantation can expect to survive for a median of 7 years
- MGUS = monoclonal gammopathy of uncertain significance
 - MGUS is a non-malignant condition with an abnormal paraprotein or M-protein
 - It is usually an asymptomatic, chance finding and once myeloma has been ruled out treatment is not necessary, however, patients have a chance of progression to myeloma or lymphoma at a rate of 1% per year with a latent period of up to 20 years

Diagnosis in primary care

- Consider possible myeloma in adults (especially aged over 60) presenting with non-specific symptoms e.g. fatigue, bone pain, recurrent infection, headaches etc.
- Check FBC, ESR, renal function and calcium studies in all patients
- A normal FBC and ESR is sufficient to rule out the disease in most patients presenting in primary care with nonspecific symptoms
- ESR is a better 'rule out' inflammatory marker for myeloma than CRP
- If FBC, ESR, creatinine, or calcium are abnormal or if the index of suspicion is higher arrange urgent serum electrophoresis and Bence-Jones protein urine assessment
- Serum electrophoresis and Bence-Jones protein are negative in 2% of people who have a non-secretory form of the disease, so refer if significant clinical suspicion persists
- Arrange XRays of symptomatic areas of bone pain to exclude pathological fracture
- Arrange urgent admission if significant hypercalcaemia or acute kidney injury
- If serum and/or urine protein electrophoresis suggest myeloma urgent haematology referral

Secondary care management

- Diagnostic confirmation is via further blood tests, bone marrow aspirate and biopsy and skeletal MRI
- Secondary care management depends on disease stage, prognosis and co-morbidities but may include bisphosphonates, corticosteroids, chemotherapy and immune-modulating drugs
- High dose chemotherapy and stem cell transplant may be offered in younger, fitter patients

Primary care management

- On-going support and holistic care, including lifestyle advice e.g. maintaining good hydration status
- Identification and prompt investigation of **possible new complications**, including: pathological fractures, spinal cord compression, impaired immunity, anaemia, bleeding disorders, acute kidney injury, cognitive impairment and stroke due to hyperviscosity
- Prompt treatment of infections with broad-spectrum antibiotics
- Vaccination, seasonal influenza and also pneumococcal
- Pain control (avoid NSAIDs due to renal risk)
- Management of depression and anxiety
- Supportive and palliative care