Optimising Cardiovascular Care of Patients with Multiple Myeloma

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Abstract

Multiple Myeloma (MM) is the third most common haematological malignancy, with increasing prevalence over recent years. Advances in therapy have improved survival, changing the clinical course of MM into a chronic condition and meaning that management of co-morbidities is fundamental to improve clinical outcomes. Cardiovascular (CV) events affect up to 7.5% of individuals with MM, due to a combination of patient, disease and treatment-related factors, and adversely impact survival. MM typically affects older people, many with pre-existing CV risk factors or established CV disease, and the disease itself can cause renal impairment, anaemia and hyperviscosity which exacerbate these further. Up to 15% of MM patients develop systemic amyloidosis - with prognosis determined by the extent of cardiac involvement. Management of MM generally involves administration of multiple treatment lines over several years as disease progresses, with many drug classes associated with adverse CV effects including high rates of venous and arterial thrombosis alongside heart failure. Recommendations for holistic management of MM patients now include routine baseline risk stratification including ECG and echocardiography, and administration of thromboprophylaxis drugs for patients treated with immunomodulatory drugs. Close surveillance of high risk patients with collaboration between haematology and cardiology is required, with prompt investigation in the event of CV symptoms, in order to identify and treat complications early. Decisions regarding discontinuation of cardiotoxic therapies should be made in a multidisciplinary setting, taking into account the severity of the complication, prognosis, expected benefits and the availability of effective alternatives.
1. Introduction

Multiple myeloma (MM) is a clonal malignant process arising from antibody-producing B-lymphocytes known as plasma cells. Owing to increased diagnosis and improved survival, its prevalence is markedly increasing, making it the third most common haematological malignancy in the United States (US), where 32,270 new cases were expected in 2020. (1) Over the past decade, ‘novel agents’ have significantly improved remission rates and survival, evolving MM into a chronic relapsing condition. Median survival is currently nearly 6 years, and additional ‘novel’ therapies are poised to further improve outcomes with a third of newly-diagnosed patients now expected to live 10 years or more from diagnosis. As a result, management of comorbidities and therapy-related toxicities are increasingly important to improve clinical outcomes and promote healthy survivorship in these patients.

Cardiovascular (CV) events are common in MM (up to 7.5% reported in phase-3 trials) (2) due to a combination of patient, disease and treatment-related factors, with impact on both CV and MM outcomes (Figure 1, Table 1). Observational studies have reported more frequent cardiac events for MM patients compared to noncancer age- and gender-matched controls, (3) and increased all-cause mortality and MM-specific mortality for newly-diagnosed MM patients with co-existing CV disease. (4) Together these data underpin the importance of assessment and treatment of CV risk factors and disease in this population to enable safe delivery of optimal MM-directed therapy over the course of many years.

In this review, we aimed to summarize the current knowledge and provide practical considerations on the management of CV complications in patients with MM.
2. Cardiovascular risk factors

2.1. Patient-related factors

MM typically affects older people, with a median age of 69 years at the time of diagnosis; (5) a population already at increased baseline CV risk. It is also more than twice as common in those of black ethnicity, males (1.3:1.0) and obese patients. (5) From retrospective data, almost two-thirds of newly-diagnosed MM patients had baseline CV disease, with reported prevalence of 14% for both arrhythmias and ischaemic heart disease. (6) Rates of baseline pre-existing CV risk factors are also high (7) and may be exacerbated by the metabolic effects of treatments, particularly steroids. In a large registry of almost 8000 MM patients, baseline prevalence of hypertension was 38%, with significantly greater risk of developing malignant hypertension over a 2-year follow-up period compared with matched non-MM patients. (7) Data from a US multicentre registry including over 2845 MM patients found baseline prevalence of diabetes at diagnosis to be 16% (8), rising to >30% after glucocorticoid therapy, with evidence for worse prognosis in this group. (9) Despite the high rates of obesity, most studies show reduced prevalence of dyslipidaemia in patients with newly-diagnosed stage-3 MM compared to controls. (3,10) This has been hypothesised to be secondary to increased clearance of cholesterol by cancer cells for growth and proliferation, and it has been demonstrated in preclinical studies to reverse after treatment. (10)

2. Disease-related factors

Renal dysfunction

Renal dysfunction is found in almost 50% of patients at the time of diagnosis, is often of multifactorial aetiology, and frequently requires dialysis in MM patients. (11) Alongside
increased risk for heart failure, arrhythmias and coronary events in patients with chronic impairment, acute renal failure (seen in up to 20% of MM patients, (11)) leads to fluid overload and electrolyte imbalances, increasing the potential for pulmonary oedema and arrhythmias.

Anaemia

Normocytic, normochromic anaemia is reported in 73% of newly-diagnosed MM patients, and in 97% over the course of their treatment.(12) Both bone marrow infiltration and chronic renal impairment contribute to anaemia, which in turn can lead to worsening heart failure symptoms and exacerbate pre-existing myocardial ischaemia.

Hyperviscosity and thrombosis

Hyperviscosity is commonly seen in plasma cell malignancies such as Waldenström’s macroglobulinemia and in the rare IgM-secreting-MM, in which hyperviscosity has been reported in 77% cases.(13) CV complications can occur due to coronary microthrombi, with resultant myocardial ischaemia and heart failure.

MM patients are also at higher risk of developing arterial thrombosis, manifest as stroke, myocardial infarction, or peripheral artery disease. Venous thromboembolism (VTE) is common, with multifactorial mechanisms including patient comorbidities (older age, immobility exacerbated by fractures and pain), inflammation and tumour-related pro-coagulant factors. This is further exacerbated by therapy, particularly immunomodulatory drugs that are key components of current optimal front-line regimens for MM.

Light chain cardiac amyloidosis

In patients with MM (or its precursors Smouldering Myeloma and Monoclonal Gammopathy of Uncertain Significance), proliferation of an abnormal clone of plasma cells overproduces circulating light-chains which can misfold and aggregate in tissues, leading to systemic light-
chain(AL) amyloidosis. In a cohort of newly diagnosed patients with MM, up to 15% develop overt systemic amyloidosis over the course of their disease, and nearly 30% have subclinical deposits in the subcutaneous fat, heart or liver on histology.(14) In this study, patients with incidental findings of asymptomatic amyloid did not differ in MM disease characteristics compared to those without amyloid, including whether the clone was kappa or lambda restricted. This finding emphasizes the importance of vigilance for the well-described clinical and biochemical features of amyloidosis.(14)

Of patients with systemic AL amyloidosis, 60% have cardiac involvement, with the potential for amyloid deposition within all structures of the heart, resulting in a wide spectrum of clinical manifestations.(15) Amyloid deposition within the extracellular space of the myocardium leads to restrictive pathophysiology and heart failure - initially with preserved left ventricular ejection fraction(LVEF). Circulating light-chains can also have direct toxic effects on the myocytes (proteotoxicity), leading to myocyte inflammation and apoptosis and subsequent systolic dysfunction.(16) Despite normal epicardial coronary arteries, perivascular infiltration can cause ischaemia and angina, and infiltration of the conducting system may result in conduction abnormalities requiring pacemaker implantation.(17) Atrial arrhythmias are found in half of patients,(18) with ventricular arrhythmias in 25%.(19)

In all patients with known MM, CV symptoms should prompt appropriate cardiac investigations, however diagnosis of amyloid can be challenging. ECG abnormalities (low voltages on limb leads and pseudo-infarct pattern) are highly specific (98%) for diagnosis of cardiac amyloid, however their sensitivity is low.(20) This contrasts with echocardiography, where reduced global longitudinal strain with apical sparing pattern has high sensitivity(93%) and specificity(82%).(21) Red flag markers for AL amyloidosis include echocardiographic features of biventricular hypertrophy with thickening of valves, discrepancy between left ventricular(LV) wall thickness and QRS voltage, atrioventricular block, repeated elevated
troponin levels and reduced LV global longitudinal strain with apical sparing (Figure 2). Patients with cardiac investigations suggestive of amyloidosis should undergo a myeloma screen including serum and urine electrophoresis with immunofixation and serum kappa/lambda free-light-chain quantification. Detection of a monoclonal protein suggestive of AL aetiology should prompt Haematology referral and tissue diagnosis for confirmation and typing.

Cardiac magnetic resonance incorporates tissue characterisation and has emerged as a useful diagnostic and prognostic tool in the evaluation of cardiac amyloidosis. Parametric mapping techniques using T1 measurement pre- and post-administration of gadolinium enable quantification of the extracellular volume (ECV) and provide a surrogate quantitative marker of amyloid burden. Gadolinium kinetics are abnormal, leading to difficulty nulling the myocardium, and there is generally diffuse (basal predominant) late gadolinium enhancement that may be subendocardial or transmural (Figure 2).

Outcome in systemic AL amyloidosis is primarily determined by cardiac involvement, and staging (Revised Mayo Classification) involves measurement of troponin and NT-pro-BNP, alongside serum free light-chains levels, with cardiac biomarkers also used to assess response to therapies. (22) Prognosis ranges from 6 months for stage IV patients without treatment to more than 5 years for stage I patients who undergo stem-cell-transplantation. (22)

2.3. Treatment-related factors

The aim of treatment in MM is to achieve and maintain a deep remission for the maximum duration, and patients are likely to be exposed to multiple treatment lines as their disease progresses (Figure 3). The selection and sequence of drugs used depend on an individual patient’s performance status, comorbidities, response to and toxicities from previous therapies, patient preference and availability of increasingly costly ‘novel’ agents, which may be via
recruitment to clinical trials. Many of these drugs have associated CV adverse effects resulting in a range of toxicities(Table 2)(23).

**Proteasome inhibitors**

Proteasome inhibitors (PIs) are the mainstay of MM treatment. These drugs inhibit the ubiquitin-proteasome system, leading to intracellular accumulation of aggregated proteins that is disproportionally toxic to MM cells. Given the high metabolic rate of cardiomyocytes, these cells are sensitive to the cytotoxic effects of PIs. The risk of cardiotoxicity in MM patients is about 4% with the first-in-class drug bortezomib, with higher rates in the elderly and with concomitant steroid use.(24)

Carfilzomib is a newer more potent and irreversible proteasome inhibitor which, despite reduced dose-limiting peripheral neuropathy than with bortezomib, is associated with greater risk of CV complications.(2) The phase-3 ENDEAVOUR trial reported higher rates of adverse events in the carfilzomib compared with bortezomib arms, including grade 3 hypertension and heart failure.(2) Recent prospective data from n=95 MM patients reported CV events in 51% of patients treated with carfilzomib compared with 17% with bortezomib (any grade of severity, mainly heart failure, coronary artery events, hypertension, arrhythmia and pulmonary hypertension). Despite generally not requiring discontinuation of therapy, these events were associated with worse MM progression-free and overall survival. Of note, natriuretic peptides were highly predictive of CV complications, supporting a role for biomarker surveillance in patients receiving these treatments.(25)

Ixazomib, the first orally available proteosome inhibitor, has demonstrated a favourable cardiotoxic profile, with no differences in complications when compared to placebo,(26) suggesting that cardiotoxicity is not a drug class effect of PIs.

**Immunomodulatory imide drugs (IMiDs)**
IMiDs such as Thalidomide, Lenalidomide and Pomalidomide are also widely-used for patients with both newly diagnosed, relapsed and refractory MM, often in combination with PIs and corticosteroids. They act in several ways modifying the immune-microenvironment but also directly targeting the malignant cells.

IMiDs are associated with increased risk of both venous and arterial thrombosis,(27) due to their pro-coagulant and pro-inflammatory effects. Thrombosis rates rise further when IMiDs are used in combination with other MM treatments: up to 25% with dexamethasone(28) and 58% in patients receiving dexamethasone and anthracyclines(29).

**Alkylation agents**

Traditional cytotoxic chemotherapy still retains a role in the treatment of MM, with cyclophosphamide commonly used in combination with novel agents. Cardiotoxicity is rare with cyclophosphamide, but includes severe congestive heart failure(30) which usually occurs within days of administration, and is increased by other factors including high dose, combination with other cardiotoxic drugs or radiotherapy, and older age.(31) High dose melphalan is frequently given to ablate the bone marrow prior to autologous stem cell transplantation, with up to 11% of patients developing atrial fibrillation.(32) Simultaneous treatment with bendamustine, bortezomib and dexamethasone was associated with 6.5% of CV adverse events in a phase-2-trial including 75 patients, including heart failure and sudden cardiac death in one patient.(33)

**Corticosteroids**

Alongside the widely-known adverse CV effects of long-term corticosteroid use, concomitant administration with other cardiotoxic regimens may potentiate their risk. In a phase-3-trial of lenalidomide with different doses of dexamethasone, patients receiving high-dose
dexamethasone experienced higher rates of VTE and pulmonary embolism than those receiving low-dose dexamethasone. (34)

**Anthracyclines**

The role of anthracyclines in treatment of MM is now predominantly reserved for aggressive relapsed extramedullary disease in younger, fitter patients. The risks of systolic dysfunction and heart failure with anthracyclines are well documented (35) and arise from interaction with topoisomerase-IIβ within the cardiac myocytes leading to apoptosis, mitochondrial dysfunction and generation of reactive oxygen species. Overall incidence of anthracycline-related cardiotoxicity is widely variable, but is usually dose dependent (risk is substantially higher above cumulative dose of 240 mg/m² doxorubicin or equivalent) with higher prevalence in patients with pre-existing CV risk factors. (35)

**Novel agents**

Over recent years, more effective and safer therapies are emerging for MM treatment. CV complications appear rare with these drugs in clinical trials (36–38), however most excluded patients with significant pre-existing CV disease. Daratumumab, an anti-CD38 monoclonal antibody whose use is rapidly expanding, is commonly associated with new-onset hypertension, (36) however panobinostat, a nonselective histone deacetylase inhibitor, has minimal cardiotoxicity apart from QT prolongation (generally <500ms) in nearly 4% of patients. (39) VTE has been observed in up to 6% of patients treated with the anti-SLAMF7 monoclonal antibody elotuzumab. (37) The recently published phase-1 DREAMM-1 trial of belantamab-mafodtin, an antibody-drug targeting B-cell maturation antigen to deliver the cytotoxic agent auristatin-F directly inside MM cells, reported no significant CV complications in heavily pre-treated patients with relapsed or refractory MM. (38)
These novel agents do however still require long-term surveillance in real-world clinical settings, particularly if given in combination with other cardiotoxic treatments, or in patients at higher CV risk.

**Stem cell transplantation**

Following diagnosis and risk stratification, all patients are evaluated to determine eligibility for autologous hematopoietic stem cell transplantation (HSCT), which is proven to significantly prolong progression-free and overall survival. HSCT strategy is highly centre dependent but is not commonly offered to patients with significant comorbidities (including CV disease) due to high risk of severe complications.

HSCT survivors have a 4-fold increased risk of CV late effects compared to the general population. This elevated CV risk depends on many factors including prior MM treatment exposure, the conditioning regimen used, and patient-related risk factors including impaired LVEF. Data on the long-term risk of LV dysfunction following HSCT are lacking. Bleeker et al. reported a clinically-significant LVEF decline in 1.6% of patients following autologous-HSCT, however no association was observed with melphalan dosing and risk of cardiotoxicity. A minority of patients undergo allogeneic-HSCT, which may be potentially curative, but is associated with high rates of mortality and graft-versus-host disease.

**Radiotherapy**

Data from patients with other cancers suggest that those receiving radiotherapy experience up to 2-fold more fatal CV events than non-radiated patients, and baseline CV risk factors, cumulative dose and age at exposure are established risk factors for radiotherapy related cardiac
damage. (42) Up to one-third of MM patients receive radiotherapy at some point during the course of the disease. Generally, however this is locally directed at plasmacytomas and painful bony lesions, and rarely includes the heart within the high-dose field, therefore CV risk is likely lower.

3. Practical considerations in the management of cardiac complications in patients with Multiple Myeloma

Given the combination of improved survival, high rates of pre-existing risk factors and CV disease, and the potential for cardiotoxicity from therapies, CV considerations should be incorporated into decision-making for MM patients.

Baseline risk stratification

All patients should undergo a baseline CV risk assessment prior to initiating or changing treatments, including a thorough history and physical examination, and optimisation of CV risk factors. An ECG and echocardiogram alongside cardiac biomarkers (troponin and NT-proBNP) should be performed at baseline to identify patients with subclinical CV disease (allowing for its management and optimization) and serve as future reference if cardiac complications develop. (43,44) Formal risk stratification proformas are available for patients receiving anthracyclines, PIs and IMiDs for MM treatment. (43)

Surveillance and treatment of CV complications during MM therapy

During treatment, CV symptoms should prompt immediate investigation. Evidence-based protocols for routine systematic surveillance for CV complications with particular treatment regimens are lacking. These should however be individualised - based on patient and treatment-related risk factors, and should include echocardiography, ECG, blood biomarkers for those at
highest risk (Figure 4). Recently published European guidelines recommend surveillance of high-risk patients receiving anthracyclines with echocardiogram every 2-cycles of treatment and 6-months after therapy. (44)

Cancer treatment-related systolic dysfunction is considered as a decline of LVEF to below 50% or >10% reduction from baseline falling below the lower limit of normal, and should prompt initiation of heart failure therapy with β-blockers and angiotensin-converting enzyme inhibitors. Other parameters such global longitudinal strain (GLS) should also be considered, and a decrease falling below −18% or a >15% relative decrease from baseline to below the limit of normal may be considered an early sign of cardiotoxicity (44).

The decision on whether to hold or discontinue cardiotoxic therapy should be made on individual basis, taking into account the severity of the complication, the overall prognosis, expected benefits of treatment, and the availability of effective alternatives.

Statins have been associated with improved MM-related survival and overall survival in large retrospective studies (45) but prospective studies are currently lacking. Thrombocytopenia occurs frequently during MM treatment meaning that coronary intervention in patients with coronary disease requires careful risk-benefit assessment and should balance evidence for efficacy against bleeding risks with dual-antiplatelet therapy. Similarly, for patients with atrial arrhythmias, anticoagulant choice must balance bleeding risk, platelet levels, compliance and potential drug interactions. Decision-making should be collaborative in a multidisciplinary setting involving both haematologists and cardio-oncologists, ideally with patient involvement.

**Thromboprophylaxis for patients treated with IMiDs**

A thromboprophylaxis strategy has been proposed by the International Myeloma Working Group and the American Society of Clinical Oncology (Figure 5), (46,47) for patients receiving IMiDs-based regimens. This should include risk stratification, with lower risk patients offered
either aspirin or prophylactic low-molecular-weight heparin (LMWH), and LMWH or warfarin prescribed for higher-risk patients. Direct oral anticoagulants (DOACs) may be considered dependent on patient wishes, although unlicensed for this indication. (48)

**AL amyloidosis**

Management of AL amyloidosis involves suppressing light-chain production via treatment of the underlying plasma cell dyscrasia, and supportive treatment for CV complications. Specific evidence for benefit of standard heart failure therapies in this context is lacking, and autonomic and renal involvement commonly prevent administration. (16) Management is therefore predominantly directed towards symptomatic management of congestive heart failure with diuretics. High rates of ventricular arrhythmias and conduction abnormalities are seen early during chemotherapy, therefore pre-treatment ambulatory ECG monitoring should be considered for risk stratification, with cardiac monitoring during chemotherapy initiation for those with abnormalities. Digitalis should usually be avoided due to potential binding to the amyloid fibrils, predisposing these patients to toxicity. Implantable cardioverter defibrillators for primary prevention have been shown to be associated with high rate of appropriate shocks in a retrospective study, but no survival benefit. (49)

Newer treatments that directly target amyloid fibrils are emerging. Doxycycline has demonstrated to interfere with amyloid fibril formation and to abrogate light-chain toxicity, leading to improved survival in patients with cardiac AL amyloidosis, (50) but placebo-controlled randomised clinical trials are awaited.

**Conclusion**

Life expectancy is improving for patients with MM due to the emergence of new treatments and patients can live now more than a decade. High baseline CV risk in addition to MM-related disease factors and treatment cardiotoxicities lead to CV events in up to 7.5% of this
population. Multidisciplinary collaboration between cardiology and haematology is recommended to minimise the risk of CV complications and to further improve outcomes.

**Patients and public involvement**

Patients or the public were not involved in the design, or conduct, or reporting, or dissemination plans of our research.

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References


Figure 1. Cardiovascular considerations in patients with multiple myeloma – interplay between pre-existing risk factors and disease and treatment related complications.

AL, light chain amyloidosis; LV, left ventricular

Myeloma related factors
- Renal impairment
- Anaemia
- Hyperviscosity
- Cardiac AL amyloidosis

Patient related factors
- Age
- Hypertension
- Diabetes Mellitus
  (ethnicity, male gender)

Therapy related factors
- Cardiotoxic chemotherapy
- Corticosteroids
- Stem-cell transplant
- Radiotherapy

Increased risk of cardiovascular complications
- Hypertension
- Heart failure
  - LV systolic impairment
  - Amyloid restrictive cardiomyopathy
- Coronary artery disease
- Arterial thrombotic events
- Accelerated atherosclerosis
- Supraventricular and ventricular arrhythmias
- Venous thromboembolism
- Pulmonary hypertension
Table 1. Cardiovascular risk factors in patients with multiple myeloma

<table>
<thead>
<tr>
<th>Patient-related factors</th>
<th>Evidence</th>
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<tbody>
<tr>
<td>Older age</td>
<td>Median age 69 years at diagnosis;(5) CV complications rise with age.</td>
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<tr>
<td>Male predominance</td>
<td>MM 1.3 times more common in males(5)</td>
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<td>Obesity</td>
<td>MM more common in overweight people(5)</td>
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<tr>
<td>Black ethnicity</td>
<td>MM 2-3x more common with black ethnicity(5)</td>
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<tr>
<td>Hypertension</td>
<td>Present in 38% at baseline, malignant hypertension in up to 4/1000 person-years in MM patients with history of hypertension (7)</td>
</tr>
<tr>
<td>Diabetes</td>
<td>Present in 16% at baseline and in &gt;30% after glucocorticoid therapy; associated with worse prognosis(8,9)</td>
</tr>
<tr>
<td>Dyslipidaemia</td>
<td>Lower prevalence of dyslipidaemia in MM patients compared to controls (increased clearance of cholesterol?)(3,10,51)</td>
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<table>
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<tr>
<th>Disease-related factors</th>
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<tr>
<td>Renal dysfunction</td>
<td>50% prevalence at diagnosis; multifactorial aetiology; associated with increased risk of acute and chronic CV complications(11)</td>
</tr>
<tr>
<td>Anaemia</td>
<td>73% prevalence at diagnosis, 97% during over the disease course (12); worsens heart failure and pre-existing myocardial ischaemia</td>
</tr>
<tr>
<td>Hyperviscosity</td>
<td>Common in Waldenström’s Macroglobulinemia and in IgM-secreting-MM;(13) can lead to coronary microthrombi</td>
</tr>
<tr>
<td>Thrombosis</td>
<td>High risk of venous and arterial thrombosis due to comorbidities, cancer-related procoagulability, and treatments;</td>
</tr>
<tr>
<td>Light chain amyloidosis</td>
<td>Up to 15% of MM patients develop systemic amyloidosis; cardiac involvement leads to heart failure and arrhythmias and is associated with poor prognosis(15)</td>
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<tr>
<th>Treatment-related factors</th>
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<tbody>
<tr>
<td>Proteasome inhibitors (PIs)</td>
<td>Associated with heart failure, coronary artery events, hypertension, arrhythmia and pulmonary hypertension;(25) carfilzomib &gt; bortezomib</td>
</tr>
<tr>
<td>Category</td>
<td>Description</td>
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<tr>
<td><strong>Immunomodulatory drugs (IMiDs)</strong></td>
<td>Associated with increased risk of both venous and arterial thrombosis (up to 25% in patients treated with IMiDs and dexamethasone); indication for thromboprophylaxis (27, 46, 47)</td>
</tr>
<tr>
<td><strong>Alkylating agents</strong></td>
<td>Cyclophosphamide is associated with rare but severe heart failure; melphalan associated with AF and supraventricular tachycardia (32)</td>
</tr>
<tr>
<td><strong>Corticosteroids</strong></td>
<td>Associated with exacerbated hypertension and diabetes, and increased CV risk; increased other drug's risk of CV complications (34)</td>
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<tr>
<td><strong>Anthracyclines</strong></td>
<td>Associated with LVEF decline and heart failure (35)</td>
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<td><strong>New-novel agents</strong></td>
<td>Daratumumab associated with hypertension; Panobinostat increases QT interval; Elotuzumab associated with VTE (37)</td>
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<td><strong>Stem cell transplantation</strong></td>
<td>HSCT survivors have a 4-fold increased risk of CV late effects compared to the general population (40)</td>
</tr>
<tr>
<td><strong>Radiotherapy</strong></td>
<td>Increased risk of coronary artery disease and valvular disease when the heart is exposed (uncommon for MM patients) (42)</td>
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</table>
Table 2. Cardiovascular adverse drug reactions associated with agents used to treat MM in Europe, according to the European summary of product characteristics. AMI, acute myocardial infarction; LVEF, left ventricular ejection fraction; MM, multiple myeloma; SV, supraventricular; VTE, venous thromboembolism

<table>
<thead>
<tr>
<th>Proteasome inhibitors</th>
<th>Immunomodulators</th>
<th>Alkylating agents</th>
<th>Anthracyclines</th>
<th>Newer therapies</th>
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<tbody>
<tr>
<td>Bortezomib</td>
<td>Carfilzomib</td>
<td>Thalidomide</td>
<td>Lenalidomide</td>
<td>Pomalidomide</td>
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<tr>
<td>Impaired LVEF / Heart failure</td>
<td>Chest pain / ischaemia / AMI</td>
<td>Hypertension</td>
<td>Hypotension</td>
<td>VTE / Pulmonary embolism</td>
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<td>Uncommon</td>
<td>Common</td>
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<td>Rare</td>
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<td>Very Common</td>
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- Very Common (≥ 1/10)
- Common (≥ 1/100 to < 1/10)
- Uncommon (≥ 1/1,000 to < 1/100)
- Rare (<1/1,000) / Not known
- Not reported
**Figure 2. AL cardiac amyloidosis imaging.**

Top panel: Echocardiographic findings in patients with advanced AL cardiac amyloidosis. (A) Parasternal long axis view and showing concentric left ventricular hypertrophy (particularly noted in the interventricular septum (IVS) and posterior wall (PW)). (B) Pulse wave Doppler showing restrictive left ventricular inflow pattern (E/A ratio > 2.0, the deceleration time is low and the e/e’ ratio is elevated (the latter not shown in this picture)). (C) Strain pattern characteristic of an infiltrative process, with global reduction of the longitudinal strain sparing the apical segments. Mid and bottom panels: Cardiac magnetic resonance findings. (D) Four chamber and (G) short axis steady state free precession cines demonstrating left ventricular hypertrophy. (H) and (E) Abnormal gadolinium kinetics with diffuse and subendocardial late gadolinium enhancement (white arrowheads). (F) Native T1 map showing a high value of T1 (reference value 970 – 1050 ms). (I) Extracellular volume (ECV) map showing elevated ECV (normal <30%). IVS: interventricular septum, LA: left atria, LV: left ventricle, PW: posterior wall, RA: right atria, RV: right ventricle
Figure 3. Agents commonly used for the treatment of Multiple Myeloma in the United Kingdom.

HSCT, hematopoietic stem cell transplantation; UK, United Kingdom

Newly diagnosed Multiple Myeloma mandating treatment

Assessment of fitness for high dose Melphalan with Autologous Haematopoietic Stem Cell Transplant (HSCT)

**HSCT ELIGIBLE**

**FIRST LINE**

PI + IMID + STEROID ‘TRIPLET’ TO BEST RESPONSE

Commonly:
Bortezomib + Thalidomide + Dexamethasone
Bortezomib + Lenalidomide + Dexamethasone

\[ \downarrow \]

HSCT

\[ \downarrow \]

Expectant management until relapse

or:
Lenalidomide maintenance (not yet funded in UK)

**SECOND LINE**

Daratumumab + Bortezomib + Dexamethasone
Lenalidomide + Dexamethasone if no prior exposure
Consider 2nd HSCT if prior long response and remains fit

**THIRD LINE**

Ixazomib + Lenalidomide + Dexamethasone
Bortezomib + Panobinostat + Dexamethasone

**BEYOND THIRD LINE**

Strongly consider trials, drugs not previously exposed to, retreatment with previously effective drugs, or:
Pomalidomide
Daratumumab
Isatuximab + Pomalidomide + Dexamethasone
Figure 4. Cardiovascular risk assessment in MM patients and practical recommendations.

*Recommended for all patients but essential for high risk patients. Based on the recently published position statement on baseline risk assessment in cancer patients from the European Society of Cardiology in collaboration with the International Cardio-Oncology Society (43)

** Please see Figure 4 for specific recommendations (47)

*** Please note that specific evidence for benefit of standard heart failure therapies in patients with cardiac amyloidosis are lacking.

CV, cardiovascular; ECG, electrocardiogram; IMiD, immunomodulatory drugs; LMWH, low-molecular-weight heparin; LVEF, left ventricular ejection fraction; MM, multiple myeloma
Figure 5. Venous thromboembolic events (VTE) risk assessment and recommendations

IMiD, immunomodulatory drugs; LMWH, low-molecular-weight heparin; MM, multiple myeloma; VTE, Venous thromboembolism