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Current and emerging triplet combination therapies for relapsed and refractory multiple myeloma

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Summary

Despite significant improvement in outcomes have been observed for multiple myeloma (MM) patients over the past 10-15 years, mainly due to the introduction of novel agents targeting the tumor clone and the bone marrow microenvironment, treatment of refractory and/or relapsed (RR) disease remains a challenge, particularly for patients who have failed prior bortezomib- and lenalidomide-based therapies. More recently, new drugs with different mechanisms of action, including second generation proteasome inhibitors, third generation immunomodulatory drugs, histone deacetylase inhibitors and monoclonal antibodies, have been developed and are under investigation, further increasing treatment options for RRMM patients. Overall, novel agent-based triplet combinations demonstrated superior response rates and prolonged disease control when compared with two-drug regimens in several randomized clinical trials, without adding any relevant additional toxicity. Salvage triplet therapies are likely to play a key role in overcoming drug-resistance and hold promise to further improve long-term outcomes of RRMM patients.

Key Words: multiple myeloma , IMiDs , proteasome inhibitors , monoclonal antibodies , relapsed refractory

Introduction

Over the past decade, the introduction and widespread use of novel agents, such as the immunomodulatory drugs (IMiDs) thalidomide and lenalidomide, and the first-in-class proteasome inhibitor (PI) bortezomib, has dramatically improved the prognosis of multiple myeloma (MM) patients, resulting in a significant increase in the depth and duration of responses which have ultimately led to extended survival. [1] These novel drugs represent the backbone of many current standard-of-care therapies for young and elderly patients, both at diagnosis and in the relapse setting.

Despite these major advances, MM remains an incurable malignancy with the majority of patients experiencing relapse and requiring additional therapy. At every subsequent relapse the duration of response is progressively shorter and drug-resistance ultimately emerges. The outcomes of patients refractory to prior bortezomib- and lenalidomide-based therapies are dismal, with median progression-free survival (PFS) of only 5 months and overall survival (OS) of 9 months. [2] Therefore, newer drugs and novel combinations that show promise to overcome resistance to standard therapies are needed for relapsed/refractory (RR) MM.

New insights into the biology of the disease have prompted an intense research focused to the development of additional novel agents targeting specific pathways involved in tumour cell growth and survival. A plethora of new compounds have been extensively investigated in the last few years, both in the preclinical setting and in clinical trials. Due to their effectiveness some of these agents have received fast-track approval and have now moved to phase III and IV studies. In this regard, the third generation IMiD pomalidomide in combination with dexamethasone and the second generation PI carfilzomib as single agent have been approved by U.S. Food and Drug Administration (FDA) in 2013 for the treatment of RRMM patients who have received at least two prior treatment regimens, including both lenalidomide and bortezomib, and have demonstrated

disease progression on the last therapy. [3,4] Both molecules demonstrated a more potent *in vitro* activity than their respective first-in class compounds bortezomib and thalidomide. [4–8]

Pomalidomide, a derivate of thalidomide with a modified chemical structure, was demonstrated *in vitro* to have multiple mechanisms of action including direct antiproliferative, pro-apoptotic, and antiangiogenic effects, and inhibition of stromal cell support. Furthermore, pomalidomide showed modulatory effects on bone resorption and on the immune system. [9] Carfilzomib is an epoxyketone PI that irreversibly and selectively inhibits the chymotrypsin-like activity of the 20S proteasome, necessitating *de novo* protein synthesis to restore proteasome's activity. In preclinical studies carfilzomib demonstrates more potent proteasome inhibition with a minimal off-target activity compared to its predecessor bortezomib. In particular, the significantly less cross-reactivity with non-proteasomal proteases has been shown to correlate with a lack of carfilzomib-related neurotoxicity in preclinical studies. [5,10]

Although second generation PIs and third generation IMiDs have proven to be effective in the treatment of RRMM, [3,4] other agents with different mechanisms of action are being developed and hold promise to represent promising therapeutic options for MM treatment. Among emerging novel drugs with different mechanisms of action it is worth highlighting monoclonal antibodies (MoAb) and histone deacetylase inhibitors (HDACi).

MoAb are directed against cell-surface receptors that are uniformly expressed by plasma cells and that impact fundamental cellular capabilities such as homing, adhesion and migration. Direct target-binding activates potent cytotoxic immune effector functions such as Ab-dependent cellular cytotoxicity (ADCC), complement-dependent cytotoxicity (CDC), activation of natural killer (NK) cells apoptosis, leading to MM cells lysis. Among the principal MoAb tested in RRMM, elotuzumab, targeting cell surface glycoprotein SLAMF7, failed to induce responses when used as a single agent, whereas significant activity was observed when it was combined either with

bortezomib or lenalidomide in phase I and II dose–escalation studies. [11,12] On the other hand, the anti-CD38 MoAb daratumumab exerted remarkable antitumor activity, which was further enhanced by the combination with lenalidomide. [13,14]

HDACi are involved in the unfolded protein response pathways and in the epigenetic regulation of gene expression, promoting cell proliferation and cell death. Despite demonstration of promising preclinical activity, monotherapy with the pan-HDACi vorinostat failed to provide any clinical benefit, while its use in combination with bortezomib resulted in a relevant increase in toxicity. [15] More promising results have been reported with the pan-HDACi panobinostat that has been granted fast track approval by the FDA in combination with bortezomib and dexamethasone for the treatment of RRMM, based on the increased PFS compared to bortezomib and dexamethasone demonstrated in a phase III study. [16]

Triplet *versus* double combinations

From what said it appears evident that various treatment options and drug combinations are currently available for RRMM patients. A growing body of evidence suggests that a three drug-based salvage therapy, often including a novel agent, could represent a key strategy to improve long-term treatment outcomes and overcome drug-resistance compared to standard treatments.

This review provides a summary of the main studies reported so far and aimed at exploring the role of three drug novel agent-based therapies for RRMM patients, with a particular focus on phase III trials prospectively comparing triplet versus doublet regimes. (Table.1)

First and second generation IMiDs: thalidomide- and lenalidomide-based regimens

Thalidomide was the first novel drug with well established activity in the RRMM setting, yielding an overall response rate (ORR) of 25% when used as a single agent, [17,18] and up to 50% when in combination with high dose dexamethasone. [19]

A higher efficacy was demonstrated with the subsequent addition of chemotherapeutic drugs, such as an alkylator [20,21] or an anthracycline [22]. Offidani et al. performed a pair-matched comparison of 94 RRMM patients who received a minimum of 3 to 6 cycles of either thalidomide-dexamethasone or the same doublet combined with liposomal pegylated doxorubicin (ThaDD), followed by maintenance therapy with thal-dex until relapse or toxicity. With the limits of a retrospective case-match analysis, ThaDD resulted in a significantly higher rate of at least minimal response (92% vs 63.5% in the control group, $p < 0.001$), including a superior rate of \geq partial response (PR, 75.5% vs 59.5%) and \geq near complete response (nCR, 30% vs 10.5%, $p=0.02$). This gain translated into a prolonged median PFS (21 vs 11.5 months for the control group) and OS (35.5 vs 20 months, respectively) for those patients receiving the triplet combination. [22] As expected the addition of an anthracycline was associated with a higher frequency of hematologic toxicity and infections which, however, were easily managed with adequate prophylaxis and supportive care and did not led to a higher treatment discontinuation. [22]

Results of a phase III randomized trial aimed at comparing the efficacy and safety of the triple combination bortezomib-thalidomide-dexamethasone (VTD) vs thalidomide-dexamethasone (TD) as first salvage therapy in MM patients relapsing after an autologous stem-cell transplantation were recently reported. [23] All patients received thalidomide at 200 mg daily for 1 year and dexamethasone at the total dose of 160 mg every 3 weeks. In addition, patients allocated to VTD treatment received intravenous bortezomib twice-weekly for the first eight 21-day cycles, and then once a week for additional 6 months. As already seen in preclinical studies

and in the upfront setting [24–26], the addition of bortezomib to TD resulted in superior efficacy in terms of high quality responses and prolonged disease control. On an intention-to-treat basis, VTD yielded a significantly higher rate of at least very good partial response (VGPR) as compared to TD (56% vs 35%, respectively, $p = 0.001$) and a two-fold higher probability to achieve CR+nCR (45% vs 21%, $p < 0.001$). Median time to progression was longer in patients receiving VTD than TD, (19.5 vs 13.8 months, $p = 0.001$), as well as median PFS (18.3 vs 13.6 months, $p = 0.001$), whereas the 2-year projected OS rates were not significantly different (71% vs 65%) between the two groups. Overall, grade 3-4 drug-related adverse events were more frequent in the VTD arm compared to TD (59% vs 36%, $p < 0.001$). In particular, the most frequent grade ≥ 3 adverse event was cumulative, dose-related, peripheral neuropathy, which occurred in 29% of patients in the VTD arm as compared to 12% in TD ($p = 0.001$). [23]

Lenalidomide in combination with high-dose dexamethasone (RD) was approved for use in RRMM in 2006 on the basis of two parallel randomised phase III trials demonstrating its superiority versus high-dose dexamethasone alone in terms of ORR and median PFS. [27,28] Based on the results of a subsequent study for newly diagnosed MM patients providing demonstration of the inferior short-term OS and higher toxicity of RD compared to lenalidomide and low-dose dexamethasone (Rd), [29] this latter combination has formed the backbone of many triplet therapies for RRMM patients incorporating either a PI or a MoAb, which will be discussed more in details in the following subsections.

Third generation IMiDs: pomalidomide-based regimens

Pomalidomide combined with low-dose dexamethasone (pom-dex) proved to be a feasible and effective salvage therapy for heavily pre-treated RRMM patients who had exhausted prior responsiveness to bortezomib and lenalidomide. [30,31] In particular, in the MM-03 trial

comparing pom-dex vs high-dose dexamethasone in patients who had received a median of 5 lines of prior therapy, the doublet pom-dex regimen yielded a 32% ORR, with median PFS and OS values of 4 and 13 months, respectively. [3]

More recently, the addition of a third drug to pom-dex has been explored in several phase I/II studies, in order to further improving clinical outcomes. Larocca et al. evaluated dosing, safety profile and efficacy of pomalidomide-cyclophosphamide-prednisone (PCP) in patients who had failed 1 to 3 prior lines of therapy, including lenalidomide. Despite using a monthly dose of pomalidomide of 70 mg (inferior to the standard dose of 84 mg/month) and a low dose of cyclophosphamide (namely 50 mg every other day), the PCP regimen induced an ORR of 51% and a median PFS of 10.4 months, with 69% of patients alive at 1 year. Most common grade 3-4 hematologic toxicities were consistent with what observed in previous studies. [32]

Similarly, preliminary data from a phase II randomised trial showed that pom-dex combined with oral cyclophosphamide yielded higher ORR and a two-fold longer median PFS (9.2 vs 4.4 months) compared to pom-dex alone. [33]

Even more promising results seemed to emerge for the combination of pom-dex with a first or second generation PI. Impressive rates of at least PR, averaging approximately from 75 to 80%, were reported for both pom-dex-bortezomib and pom-dex-carfilzomib triplets, with median PFS of 12 and 17 months, respectively. [34,35] The combination of pomalidomide-carfilzomib-dexamethasone (CPD) is currently being further tested in three ongoing phase I/II clinical trials for RRMM patients previously exposed and refractory or intolerant to lenalidomide. Results from a phase I study including 32 patients previously treated with a median of 6 lines of therapy have recently been published. [36] By study design, all patients were lenalidomide refractory and 91% were also bortezomib refractory. In this heavily pretreated population, at least a minimal response (MR) was observed in 66% of the patients, including 16% of VGPR. The efficacy of the CPD regimen

was retained across all cytogenetic subgroups, including those high-risk. Side effects were manageable, with the main toxicity being hematological. In particular, 44% of patients experienced grade III-IV neutropenia, a value superimposable to that seen in this setting using pom-dex alone. Grade III non hematological adverse events occurred in 1 patients each and included congestive heart failure, transient ischemic attack and pulmonary embolus; two treatment related deaths were reported. [36]

Proteasome inhibitors-based regimens

Based on the results of the phase III APEX trial, the first-in-class PI bortezomib received FDA and EMEA approval in 2005 as single agent therapy for relapsed MM patients who had received at least one prior line of therapy. [37,38] In subsequent phase II studies, the combination of bortezomib and dexamethasone (VD) demonstrated added clinical benefit, without adversely affecting the safety profile, and is actually considered as a reference regimen in the relapse setting. [39]

Based on preclinical demonstration that bortezomib enhanced the sensitivity of myeloma cells to conventional chemotherapy [40] and of its synergistic clinical activity when combined with other agents, [41] several phase I/II trials have explored the combination of bortezomib with anthracyclines, [42–44], IMiDs [45] and bendamustine [46–48]. All these studies showed that the combination of VD with a third agent in patients exposed to a median of 1 to 5 prior therapies provided a rate of at least PR in the 61-72% range, with a manageable toxicity. Offidani and colleagues investigated the role of the triplet therapy bendamustine-bortezomib-dexamethasone (BBD) in 75 RRMM patients that had received less than 4 prior lines of therapy. In the 70 evaluable patients treatment with BBD provided an ORR (at least a PR) of 71.5%, with 16% and 18.5% of patients achieving a CR and a VGPR, respectively. With a follow-up of 1 year the median time-to-

progression (TTP) was 16.5 months and the 1-year overall survival was 78%. [46] Similar results were obtained by the French group, which investigated the BBD regimen in 73 RRMM patients. After 4 cycles of BBD at least a PR was achieved in 57.6% of patients, with 10.9% and 16.5% achieving a CR and a VGPR respectively. With a median follow up of 15.7 months the median PFS was 10.8 months and the median OS was 23 months. [48]

More recently, in the phase III randomized PANORAMA-1 study the combination of VD with the HDCAi panobinostat was compared with VD and placebo in a population of more than 760 RRMM patients. The simultaneous targeting of different pathways involved in the unfolded protein response, e.g. the ubiquitin-proteasome pathway by bortezomib and the aggresome protein degradation pathway by HDACi, provided the rationale for this triplet regimen. The addition of panobinostat to VD resulted in a significantly higher frequency of CR/nCR (27.6%) compared to VD-placebo (15.7%) ($p < 0.0001$), a gain which translated into a significantly longer PFS (12 vs 8.1 months, $p < 0.0001$). Importantly, PFS benefit was observed across pre-specified subgroups, including patients aged 65 years or older, those with ISS 2-3 and patients previously exposed to bortezomib. At the time of the analysis, no significant difference was reported in median OS (33.6 months for panobinostat and 30.4 for placebo group). The triplet regimen was associated with a superior frequency of grade 3-4 adverse events, including diarrhoea (26% vs 8% for VD alone), fatigue (24% vs 12%), and haematological toxicities, mainly thrombocytopenia (67% vs 31%) and lymphopenia (53% vs 40%). [16]

As stated above, the combination of lenalidomide with emerging new molecules has been extensively investigated in the last few years. Recently, the ASPIRE trial prospectively compared Rd plus carfilzomib versus Rd in a large population of relapsed myeloma patients who had received 1 to 3 prior treatments. [49] The addition of carfilzomib to Rd for eighteen 28-day cycles followed by Rd until progression or undue toxicity resulted in a remarkable improvement of PFS (26.3 months

vs 17.6 in the control arm, $p < 0.001$) and a 31% decrease in the relative risk of progression or death compared to standard Rd. The advantage in PFS offered by the triplet combination was retained across all predefined subgroups, including patients previously exposed to bortezomib and IMiDs, and those with a high-risk cytogenetic profile at study entry. Two-year estimates of OS were 73.3% and 65%, respectively ($p=0.04$). The 3-drug regimen was associated with a higher rate of CR or better (31.8% vs 9.3%, $p < 0.001$) and a superior ORR (87% vs 66.7%, $p < 0.001$). Overall, no relevant differences were observed between the 2 groups in the frequency of reported grade ≥ 3 adverse events or treatment discontinuation due to toxicity, although patients in the carfilzomib arm experienced more common side effects than those treated with Rd, including diarrhea, cough and hypertension. Of note, the incidence of any-grade peripheral neuropathy was 17% in both groups of patients. [49] Based on these results, in July 2015 the FDA approved Carfilzomib in combination with lenalidomide and dexamethasone for the treatment of patients with relapsed MM who have received 1 to 3 prior lines of therapy.

Based on its activity as single agent in RRMM patients previously exposed to a median of 4 prior lines of treatment, [50] the oral PI ixazomib is also being evaluated in combination with Rd versus the doublet Rd in an ongoing phase III clinical trial.

Monoclonal antibodies-based regimens

Among the new agents explored for the treatment of MM, MoAbs have emerged as an attractive targeted strategy based on the wide range of antigens expressed on the surface of plasma cells, in analogy with what previously seen in non Hodgkin Lymphoma, where the advent of the antiCD20 MoAbs dramatically changed the treatment scenario. [51,52] Encouraging results have been recently provided by the ELOQUENT-2 trial, a phase III randomised study comparing the efficacy and safety of Rd with or without elotuzumab in RRMM patients. [53] The rationale for

combining a MoAb with an IMiD is based on the synergistic effect of direct activation and engagement of the host's natural killer cells by elotuzumab, and the enhancement of antibody-dependent cellular toxicity exerted by lenalidomide. Six hundred forty-six patients, with 1 to 3 prior therapies and a documented disease progression after their most recent treatment, were randomly assigned to receive Rd alone until disease progression or unacceptable toxicity, or the same combination plus elotuzumab. The combination Elotuzumab-Rd resulted in an extended PFS as compared with the control arm (19.4 months vs 14.9 months, respectively, $p < 0.001$), a benefit maintained regardless of patients age, number of prior therapies, previous exposure to lenalidomide or the presence of high-risk cytogenetic abnormalities, such as del 17(p) or t(4;14). The ORR was 79% in the elotuzumab group vs 66% in the control group ($p < 0.001$). Concerning safety profile, the most common grade 3-4 adverse events were lymphocytopenia, which was reported in 77% of patients receiving elotuzumab and 49% of patients in the Rd arm, neutropenia and fatigue. Infusion reactions to elotuzumab occurred in 10% of patients and were of mild grade. [53]

In another recent phase II study, 152 patients were randomized to receive either VD alone or VD in combination with Elotuzumab until disease progression or unacceptable toxicity. Despite the achievement of a similar ORR, the triplet regimen improved median PFS (9.7 vs 6.9 months for VD, $p=0.08$). More common \geq grade 3 adverse events were thrombocytopenia and infections, with mild grade 1-2 infusion reactions occurring in 7% of patients in the elotuzumab group. [54]

Based on the favourable toxicity profile and efficacy of the anti-CD38 MoAb Daratumumab as single agent in heavily pretreated and refractory MM patients, [55] subsequent phase I/II studies explored Daratumumab combined with Rd in RRMM, with a further enhanced ORR, averaging 75%. [14,56,57] These results prompted the initiation of phase III clinical trials of Daratumumab in combination with either Rd or VD, whose results are eagerly awaited. The combination of

daratumumab with pomalidomide and dexamethasone is also being evaluated in ongoing phase I trials. [58]

Expert commentary

The plethora of emerging novel drugs currently under investigation and in various stages of clinical development has dramatically expanded the therapeutic armamentarium for RRMM. However, in the absence of prospective randomized trials directly comparing the different newer combinations in the same setting of patients, a standard of care for the management of RRMM cannot be defined. More conventional regimens, such as those including alkylators and conventionally used in newly diagnosed patients should also not be forgotten, as they can still have a role in the RR setting. In this respect regimens such as Bortezomib-Melphalan-Prednisone (VMP) or Bortezomib-Cyclophosphamide-Dexamethasone (VCD), or even the more old combinations MP and CP, could be considered for the treatment of RRMM patients, proven that patients are not refractory to these drugs.

With so many available treatment options, and in the lack of a defined standard of care for this category of patients, the choice of optimal salvage therapy and treatment sequencing still remains a special challenge and depends on several disease-related and patient-related factors, such as risk stratification, age, comorbidities, previous therapies actually received and related quality and duration of response, not to mention drug availability and costs. In addition, a careful balance between efficacy and potential side effects is required, in particular considering that patients with advanced or end-stage MM are more fragile and at risk of developing severe complications compared with those at earlier relapse.

Five years view

Despite the caveat mentioned above (possible higher toxicities for 3-drugs compared to 2-drugs regimen), a three-drug based therapy for RRMM is likely to be superior to a double regimen. This considerations are based on the results of several large, phase III clinical trials, demonstrating that the addition of a third drug to the backbone of Rd and VD increases the depth and duration of response and prolongs PFS up to unprecedented values, sometimes rivalling those seen in newly diagnosed disease, without meaningfully affecting patient safety and quality of life. Thanks to these trials most of this newer drugs have received, or are currently receiving, FDA and EMA approval, making it possible also for patients outside clinical trials to benefit from them. In light of these observations, triplet combination regimens actually represent a key option in overcoming drug resistance and improving long-term treatment outcomes in RRMM patients. The availability of multiple drugs with different mechanism of actions makes it reasonable to postulate that in the future a plethora of therapeutic regimens, differently combining PI, IMiDs, HDACi and MoAb will be available for RRMM patients, ultimately increasing the survival expectancy of this particularly difficult group of patients. Whether a cure for MM patients will eventually be achieved is a burning and interesting question, that needs to be carefully addressed. Future and ongoing studies, also looking into the biology of MM and on the mechanism of resistance development, will help us answer this question and to balance risk and benefit in the treatment of myeloma.

Key issues

- Despite these major advances, MM remains an incurable malignancy, with patients ultimately succumbing to their disease
- Outcomes of patients who have failed prior bortezomib- and lenalidomide-based therapies are particularly dismal
- Second generation proteasome inhibitors, third generation immunomodulatory drugs, histone deacetylase inhibitors and monoclonal antibodies have been developed and are under investigation, further increasing treatment options for RRMM patients
- Novel agent-based combinations are likely to increase treatment options for RRMM patients
- Triplet combinations demonstrated superior response rates and prolonged disease control when compared with two-drug regimens
- The addition of a third agent does not add any relevant additional toxicity.
- Salvage triplet therapies are likely to play a key role in overcoming drug-resistance and it is believed that they will further improve long-term outcomes of RRMM patients.
- A standard of care for the management of RRMM still cannot be defined.
- Optimal salvage therapy and treatment sequencing are still the matter of debate

Conflict of interests:

Michele Cavo has received honoraria and has been a member of the advisory board for Celgene, Janssen and Millennium. The authors have no other relevant affiliations or financial involvement with any organization or entity with a financial interest in or financial conflict with the subject matter or materials discussed in the manuscript apart from those disclosed.

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Table 1: Principal trials comparing two- versus three-drug combinations in RRMM

Author	Study	Pts	Drugs	ORR (≥ PR)	P value	nCR/CR	P value	PFS (mo)	P value	OS	P value
Offidani (2008)[46]	Retrospective,	47	Thal-dex	59.5%	0.07	10.5%	0.002	11.5	0.009	20 mo	0.01
	case match	47	ThaDD	75.5%		30%		21		35.5 mo	
Garderet (2012)[23]	Randomized	134	TD	72%	0.003	21%	0.001	13.6	0.001	65% @ 2-y	0.09
	Phase III	135	VTD	87%		45%		18.3		71% @ 2-y	
Stewart (2015)[49]	Randomized	396	Rd	66.7%	< 0.001	9.3% *	< 0.001	17.6	< 0.001	65.0% @ 2-y	/
	Phase III	396	Rd-Carfilzomib	87.1%		31.8 %		26.3		73.3% @ 2-y	
Lonial (2015)[53]	Randomized	325	Rd	66%	< 0.001	11% *	/	14.9	< 0.001	NA	/
	Phase III	321	Rd-Elotuzumab	79%		11%		19.4			
San Miguel (2014)[16]	Randomized	381	VD-placebo	54.6%	0.09	15.7%	< 0.001	8.1	< 0.001	30.4	0.26
	Phase III	387	VD-Panobinostat	60.7%		27.6%		12		33.6	
Jakubowiak (2015) [12]	Randomized	75	VD	63%	NA	NA	/	6.9	0.08	NA	/
	Phase II	77	VD-Elotuzumab	66%		9.7					

Abbreviations: ORR: overall response rate; PR: partial response; nCR: near complete response; CR: complete response; PFS: progression free survival; OS overall survival; ThaDD: thalidomide-dexamethasone-doxorubicin; TD: thalidomide-dexamethasone; VTD: bortezomib-thalidomide-dexamethasone; Rd: lenalidomide- low dose dexamethasone; VD: bortezomib-dexamethasone; NA: not available.

* only ≥CR

Table 2: Principal grade III-IV toxicities of three-drug combination regimen in RRMM

Author	Treatment	Pts	Anemia	Neutropenia	Thrombocytopenia	Infections	PNP	Cardiac events	TE	GI events
Offidani (2008)[46]	ThaDD	47	NR	25%	3 pts	23%	2%	0	6 pts	NR
Garderet (2012)[23]	VTD	13	8%	11%	17%	14%	31%	2%	6%	8%
Stewart (2015)[49]	Rd-Carfilzomib	39	17.9%	29.6%	16.6%	1.8%	17%	7.1%	NR	5.6%
Lonial (2015)[53]	Rd-Elotuzumab	32	19%	34%	19%	NR	NR	NR	NR	6%
San Miguel (2014)[16]	VD-Panobinostat	38	18%	35%	68%	13%	17%	NR	NR	46%
Jakubowski (2015) [12]	VD-Elotuzumab	77	NR	NR	9%	19%	NR	NR	NR	NR

Abbreviations: PNP: peripheral neuropathy; TE: thromboembolic events; GI: gastrointestinal events, including diarrhoea, constipation, vomiting, nausea; NR: not reported