Prospective longitudinal study on quality of life in relapsed/refractory multiple myeloma patients receiving second- or third-line lenalidomide or bortezomib treatment

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Treatment advances for multiple myeloma (MM) that have prolonged survival emphasise the importance of measuring patients’ health-related quality of life (HRQoL) in clinical studies. HRQoL/functioning and symptoms of patients with relapsed/refractory MM (RRMM) receiving second- or third-line lenalidomide or bortezomib treatment were measured in a prospective European multicentre, observational study at different time points. At baseline, patients in the lenalidomide cohort were fraiier than in the bortezomib cohort with more rapid disease progression at study entry (more patients with Eastern Cooperative Oncology Group performance status >= 2, shorter time from diagnosis, more chronic heart failure, higher serum creatinine levels, more patients with dialysis required). About 40% of the patients receiving lenalidomide discontinued the study in < 6 months while 55% in the bortezomib cohort discontinued. No substantial HRQoL deterioration was observed for the first 6 months in patients with RRMM receiving one or the other treatment. For patients still on treatment at study completion (month 6), only the European Organization for Research and Treatment of Cancer Quality-of-Life Core domains of Diarrhoea and Global Health Status/QoL had worsened in the lenalidomide and bortezomib cohorts, respectively. A clinically meaningful deterioration in HRQoL was more often observed for patients who discontinued the study prior to 6 months in the bortezomib cohort than in the lenalidomide cohort.

ORIGINAL ARTICLE

INTRODUCTION

Multiple myeloma (MM) is an incurable haematological malignancy characterised by bone marrow infiltration of malignant plasma cells leading to impaired haematopoiesis, immuno-suppression and a high incidence of bone lesions that can cause pathologic fractures and severe bone pain.¹,² MM accounts for 10% of the malignant haematological diseases and approximately 1% of all cancer-related deaths in Western countries.³ It was estimated that, in 2015, 26 850 new cases of MM would be diagnosed and 11 240 patients would die from the disease in the United States of America.⁴ In Europe, MM is diagnosed in approximately 38 956 patients and claims about 24 296 lives each year.³

Despite the considerable improvements in the treatment, the majority of the MM patients will experience multiple subsequent relapses of their disease requiring subsequent treatment.⁵ There are increasingly more novel agent options in managing MM at diagnosis and relapse,⁶ including the proteasome inhibitors bortezomib, carfilzomib and recently US Food and Drug Administration-approved ixazomib,⁷ the immunomodulatory drugs pomalidomide, lenalidomide and thalidomide and recently Food and Drug Administration-approved daratumumab and elotuzumab.⁸,⁹ However, at some point MM can become refractory following multiple lines of treatment and resistant to the currently available therapies.¹⁰ While MM has no cure, successive lines of treatment can lead to greater risk of developing adverse reactions that could be in turn responsible for sequelae and create handicap impacting patients’ quality of life.

Health-related quality of life (HRQoL) is a multidomain concept that represents the patient’s perception of the effect of illness and treatment on physical, psychological and social aspects of life.¹¹ Assessing HRQoL is critical to better capture health aspects that matter to the patients themselves and that go beyond the prolongation of life.¹²

Among current treatments used, intravenously or subcutaneously administered bortezomib and orally administered lenalidomide showed statistically significant improvements in phase III trials for the treatment of relapsed/refractory MM (RRMM) in terms of overall response rate, complete response rate, time to progression and overall survival.¹³,¹⁴ However, even though a large body of evidence supports the clinical benefits of lenalidomide¹³,¹⁵,¹⁶ and bortezomib¹⁴,¹⁷ for RRMM patients, the literature is relatively scarce regarding the burden that treatment poses on patients’ HRQoL as reported directly by patients.¹⁸,¹⁹

In order to better understand how treatments for RRMM impact patients in terms of HRQoL in the real-life context, a European, prospective, multicentre, non-interventional, longitudinal study was conducted in RRMM patients beginning second- or third-line treatment with either lenalidomide or bortezomib, the two RRMM treatment options available at the time of study implementation. The objective of this study is to describe and better understand

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patients’ HRQoL when treated with bortezomib or lenalidomide for RRMM.

MATERIALS AND METHODS

Patients

Patients were eligible for study enrolment if they had RRMM requiring second- or third-line treatment with either lenalidomide- or bortezomib-based regimens and with at least one measurable disease manifestation: any quantifiable serum monoclonal protein value (generally, but not exclusively, >1 g/dl immunoglobulin G M-protein or >0.5 g/dl immunoglobulin A) and, where applicable, urine light-chain excretion of ≥200 mg/24 h + presence of soft tissue (not bone) plasmacytomas as determined by clinical examination or applicable radiographs (that is, magnetic resonance imaging, computed tomographic scan) or a quantifiable plasma infiltration of the bone marrow as determined by bone marrow biopsy. Patients who were planned to receive a stem-cell transplant as part of the second-line treatment for MM and patients who were treated with a cytotoxic drug in combination with lenalidomide or bortezomib were excluded from the study.

This study was conducted in compliance with the Declaration of Helsinki and all current national regulations. In accordance to local requirements, the study protocol was reviewed and approved by the Independent Ethics Committee or to the inclusion of patients into the study. All patients gave written informed consent prior to their inclusion.

Study design

This was a European, prospective, multicentre, observational, longitudinal study conducted in six countries (Belgium, France, Germany, Ireland, Italy and United Kingdom). Patients were identified by their physician either through a prescreening and/or during the course of routine patient visits. Recruited patients were followed up for a maximum of 6 months. Physicians were asked to specify the reason for any patients not completing the study: disease progression, discontinuation of treatment, or any other reasons, including death, withdrawn consent or lost to follow-up.

Socio-demographic and clinical data were collected at baseline by the recruiting physicians. HRQoL was assessed using patient-completed questionnaires at baseline, 3 months and 6 months following treatment initiation and/or at study discontinuation.

Assessments

HRQoL was assessed using three questionnaires, the European Organization for Research and Treatment of Cancer (EORTC) Quality-of-Life Core (QLQ-C30),20,21 QLQ-Multiple Myeloma (QLQ-MM20)22,23 and QLQ-Chemotherapy-Induced Peripheral Neuropathy (QLQ-CIPN20)24 instruments. The QLQ-C30 includes 30 items distributed across six Functional domains (Cognitive, Emotional, Physical, Role and social functioning, Global health status/QoL) and nine Symptom domains (Appetite loss, Pain, insomnia). The QLQ-MM20 includes 20 items distributed across two Functional domains (Body image, Future perspective) and two Symptom domains (Disease symptoms, Side effects of treatment). The QLQ-CIPN20 includes 20 items distributed across three symptom domains (Autonomic scale, Motor scale, Sensory scale). For all questionnaires, all items were answered on a four-point Likert scale from ‘not at all’ to ‘very much’, except items 29 and 30 of QLQ-C30 that are answered on a seven-point Likert scale ranging from ‘very poor’ to ‘excellent’. Scores were converted to a range from 0 to 100; for Functional domains, higher scores indicate better functioning; for Symptom domains, higher scores indicate greater symptom.

Statistical analysis

Descriptive statistics were applied to describe the population, the individual components of the EORTC questionnaire and the change in EORTC questionnaire scores from baseline to month 3, month 6 or study discontinuation. Changes in EORTC scores were calculated only for patients who were able to complete the questionnaires at both baseline and the time point of interest.

Minimal important difference (MID), defined as the smallest change in a quality of life score considered important to patients, was estimated within the study to provide support for the interpretation of changes in scores.25 MID was calculated as 0.5 × SDmean with SDmean the s.d. of the score at baseline, for single-item domains, and as the s.e.m., defined as SDmean × (1 − n)1/2 with r the Cronbach’s alpha reliability coefficient, for multi-item domains.26

Patients in the bortezomib cohort were described by standard- or low-dose treatment: bortezomib treatment was considered as standard dose if ≥6 vials and as low dose if <6 vials within a 6-week period, a vial corresponding to an injection of dose of 1.3 mg/m2 body surface area as per the summary of product characteristics.

The statistical analysis was performed using the SAS software for Windows (Version 9.2, SAS Institute Inc., Cary, NC, USA).

RESULTS

Study population

Patients’ disposition and characteristics. Out of the 274 patients enrolled from December 2010 to February 2014 by 33 sites, 258 (94.2%) patients met the selection criteria and were included in the study. More patients initiated a treatment with oral lenalidomide (n = 162) than with injectable bortezomib (n = 96) (Figure 1). Among the 96 patients receiving bortezomib, 41 (42.7%) had the drug administered intravenously and 11 (11.5%) subcutaneously; the data were missing for 44 (45.8%) patients. Most patients in both cohorts received concomitant dexamethasone. Some differences in baseline characteristics (Table 1) were observed between the lenalidomide and bortezomib cohorts: the oldest quartile of patients was somewhat older in the lenalidomide cohort (77–93 years) compared with the bortezomib cohort (74–85 years). Also, the proportion of Eastern Cooperative Oncology Group performance status >2 was slightly higher in the lenalidomide cohort (6.2%) vs the bortezomib cohort (3.1%). Time since MM diagnosis was longer by > 1 year in the bortezomib cohort (3.9 ± 3.0 years) compared with the lenalidomide cohort (2.8 ± 2.5 years); also the upper quartile of time from diagnosis in the bortezomib cohort was 4.9–20.5 years vs 3.4–12.4 years in the lenalidomide cohort (Table 1). The ad hoc statistical analysis showed that age and time since diagnosis were significantly different across the two cohorts (P < 0.05). This ad hoc analysis also showed that requirement for dialysis and treatment line were significantly different across the two cohorts (P < 0.05).

Although fewer patients in the lenalidomide cohort (13.0%) were dialytic compared with the bortezomib cohort (18.8%), slightly more had chronic heart failure (14.8% vs 8.3%, respectively). Median serum creatinine levels were equal between the lenalidomide and bortezomib cohorts, but the upper quartile in the lenalidomide cohort displayed far higher serum creatinine levels (2.46 vs 0.99 µmol/L).

Figure 1. Patients’ disposition and study completion by treatment group. Reasons for discontinuation are as reported by the physician. *Other reasons include death, loss to follow-up, withdrawn consent, other and missing (include two patients in the bortezomib cohort who never received treatment: one discontinued for consent withdrawal and the other for other reasons).
| Variable                                             | Bortezomib (N = 96) | Lenalidomide (N = 162) | P-value* |
|------------------------------------------------------|----------------------|-------------------------|----------|
| **Age, years**                                       |                      |                         |          |
| Mean (s.d.)                                          | 68.0 (9.1)           | 70.9 (9.8)              | 0.022    |
| Median (Q1–Q3)                                       | 69.0 (62.0–74.0)     | 72.0 (66.0–77.0)        |          |
| Min–max                                              | 38.0–85.0            | 29.0–93.0               |          |
| **Gender, %**                                         |                      |                         |          |
| Male                                                 | 57.3                 | 51.9                    | 0.397    |
| **Country, %**                                       |                      |                         |          |
| UK                                                   | 15.6                 | 6.2                     | 0.103    |
| Ireland                                              | 5.2                  | 7.4                     |          |
| Germany                                              | 13.5                 | 12.3                    |          |
| France                                               | 11.5                 | 19.8                    |          |
| Italy                                                | 34.4                 | 30.9                    |          |
| Belgium                                              | 19.8                 | 23.5                    |          |
| **Time since MM diagnosis, years**                    |                      |                         |          |
| Mean (s.d.)                                          | 3.9 (3.0)            | 2.8 (2.5)               | 0.001    |
| Median (Q1–Q3)                                       | 3.2 (1.9–4.9)        | 2.2 (1.2–3.4)           |          |
| Min–max                                              | 0.4–20.5             | 0.1–12.4                |          |
| **ECOG performance status, %**                       |                      |                         | 0.652    |
| 0                                                    | 35.4                 | 31.5                    |          |
| 1                                                    | 46.9                 | 45.1                    |          |
| 2                                                    | 11.5                 | 13.6                    |          |
| 3                                                    | 2.1                  | 5.6                     |          |
| 4                                                    | 1.0                  | 0.6                     |          |
| Missing                                              | 3.1                  | 3.7                     |          |
| **ISS, %**                                            |                      |                         | 0.831    |
| Stage I                                              | 16.7                 | 21.0                    |          |
| Stage II                                             | 18.8                 | 16.0                    |          |
| Stage III                                            | 17.7                 | 16.7                    |          |
| Missing                                              | 46.9                 | 46.3                    |          |
| **Treatment line, %**                                |                      |                         | 0.039    |
| Second line                                          | 84.4                 | 93.8                    |          |
| Third line                                           | 13.5                 | 6.2                     |          |
| Missing                                              | 2.1                  | 0.0                     |          |
| **Additional antimyeloma drugs, %**                   |                      |                         | 0.736    |
| Dexamethasone                                        | 92.7                 | 95.7                    |          |
| Prednisone                                           | 5.2                  | 3.1                     | 0.505    |
| **Dialysis required, %**                             |                      |                         | 0.020    |
| Yes                                                  | 0.0                  | 5.6                     |          |
| No                                                   | 97.9                 | 94.4                    |          |
| Missing                                              | 2.1                  | 0.0                     |          |
| **Serum creatinine, μmol/l**                         |                      |                         | 0.242    |
| Mean (s.d.)                                          | 104.8 (62.5)         | 145.0 (326.8)           |          |
| Median (Q1–Q3)                                       | 88.4 (71.0–106.1)    | 88.4 (70.7–114.0)       |          |
| Min–max                                              | 44.2–402.2           | 43.3–3933.8             |          |
| **Comorbidities, %**                                 |                      |                         |          |
| Neuropathy                                           | 26.0                 | 26.5                    | 0.718    |
| Diabetes                                             | 18.8                 | 13.0                    | 0.267    |
| Osteoporosis                                         | 14.6                 | 15.4                    | 0.330    |
| Chronic heart failure                                | 8.3                  | 14.8                    | 0.142    |
| Urogenital disorders                                 | 10.4                 | 13.6                    | 0.559    |
| Neuropathic pain                                     | 6.3                  | 14.8                    | 0.053    |
| Gastrointestinal/hepatobiliary disorders             | 14.6                 | 9.3                     | 0.170    |
| Depression                                           | 8.3                  | 4.9                     | 0.217    |
| Arthritis                                            | 3.1                  | 7.4                     | 0.412    |
| Chronic respiratory disorder                         | 7.3                  | 4.3                     | 0.240    |
| Visual impairment                                    | 6.3                  | 3.7                     |          |
| Stroke                                               | 3.1                  | 3.7                     | 0.030    |
| Hearing impairment                                   | 3.1                  | 3.7                     | 0.181    |

Abbreviations: ECOG, Eastern Cooperative Oncology Group; ISS, International Staging System; MM, multiple myeloma. *Non-parametric P-value using T-test for continuous variables, Chi² or Fisher’s exact for categorical variables; in bold P-value < 0.05.
levels (114–2934 μmol/l) as compared with the bortezomib cohort (106–402 μmol/l). This was also reflected by the fact that 9 patients (5.6%) in the lenalidomide cohort required dialysis vs no patients in the bortezomib cohort. Slightly more patients in the lenalidomide cohort had baseline neuropathic pain (14.8 vs 6.3% in the bortezomib group) (Table 1). Altogether, these characteristics suggest frailer patients in the lenalidomide cohort with a more rapid disease progression at baseline than in the bortezomib cohort.

Out of the 162 patients receiving lenalidomide, 64 (39.5%) discontinued the study before 6 months; out of the 96 patients receiving bortezomib, 53 (55.2%) discontinued the study before 6 months. Reasons for discontinuing the study are shown in Figure 1. Twenty patients (64.5%) in the standard-dose bortezomib cohort discontinued the study and 25 patients (43.9%) in the low-dose bortezomib cohort (data not shown). Eight patients could not be classified as standard- or low-dose bortezomib because of too few bortezomib dosing administrations during the study. Detailed information of the dosage of treatment received during the study is presented in Table 2.

Mean study duration was about 5 months in the lenalidomide cohort and 4 months in the bortezomib standard- and low-dose cohorts.

Longitudinal HRQOL results for lenalidomide and bortezomib cohorts

EORTC questionnaires were completed at baseline by 93 (96.9%) patients in the bortezomib cohort and 158 (97.5%) patients in the lenalidomide cohort, at month 3 by 65 (100%) and 122 (100%) patients, at month 6 by 43 (100%) and 94 (95.9%) patients and at study discontinuation by 29 (54.7%) and 27 (42.2%) patients in the bortezomib and lenalidomide cohorts, respectively. Baseline HRQoL scores are presented in Table 3.

At study completion (month 6), HRQoL reductions from baseline reaching MID were observed for 1 of the 22 domains in each

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### Table 2. Description of study treatment received by patients along the study

| Variable                                      | Bortezomib (N = 162) | Lenalidomide (N = 162) |
|-----------------------------------------------|-----------------------|------------------------|
| **Standard dose (N = 31)**                    |                       |                        |
| Treatment duration, months Mean (s.d.)         | 3.8 (1.4)             | 4.1 (2.1)              |
| Median (Q1–Q3)                                | 3.7 (2.4–5.1)         | 4.7 (2.4–5.6)          |
| Min–max                                       | 1.5–6.9               | 0.2–10.9               |
| **Cumulative dosage, mg/m² or mg** Mean (s.d.)| 25.8 (8.7)            | 10.8 (8.8)             |
| Median (Q1–Q3)                                | 24.9 (19.5–31.2)      | 7.8 (5.2–10.4)         |
| Min–max                                       | 10.5–44.1             | 2.3–41.6               |
| **Average dosage, mg/m² or mg per cycle** Mean (s.d.) | 4.7 (0.7)             | 3.6 (1.7)              |
| Median (Q1–Q3)                                | 5.1 (4.2–5.2)         | 2.7 (2.5–4.8)          |
| Min–max                                       | 2.9–5.8               | 1.6–7.5                |

### Table 3. Baseline EORTC scores in the lenalidomide cohort and bortezomib dosage cohorts, presented as mean (s.d.)

| EORTC questionnaire | Domain                          | Bortezomib (N = 96) | Lenalidomide (N = 162) |
|---------------------|---------------------------------|---------------------|------------------------|
| QLQ-C30             | Global health status/QoL        | 54.6 (25.7)         | 54.8 (23.8)            |
|                     | Physical functioning            | 69.1 (27.1)         | 63.7 (27.0)            |
|                     | Role functioning                | 58.9 (34.5)         | 56.7 (36.9)            |
|                     | Emotional functioning           | 69.6 (25.7)         | 70.3 (23.8)            |
|                     | Cognitive functioning           | 79.5 (21.1)         | 77.1 (25.6)            |
|                     | Social functioning              | 69.8 (31.7)         | 68.6 (32.6)            |
|                     | Fatigue                         | 42.6 (28.1)         | 43.0 (28.7)            |
|                     | Nausea and vomiting             | 6.7 (15.0)          | 8.5 (18.9)             |
|                     | Pain                            | 36.1 (34.1)         | 39.7 (33.8)            |
|                     | Dyspnoea                        | 21.5 (26.6)         | 26.9 (28.6)            |
|                     | Insomnia                        | 32.6 (33.3)         | 27.2 (32.2)            |
|                     | Appetite loss                   | 19.9 (28.4)         | 21.4 (31.2)            |
|                     | Constipation                    | 19.1 (26.4)         | 26.5 (33.5)            |
|                     | Diarrhoea                       | 7.3 (16.3)          | 9.7 (21.1)             |
|                     | Financial difficulties          | 15.6 (26.8)         | 13.6 (24.2)            |
|                     | Body image                      | 79.1 (30.9)         | 79.5 (30.1)            |
|                     | Future perspective              | 54.3 (28.0)         | 52.7 (30.0)            |
|                     | Disease symptoms                | 26.0 (22.5)         | 27.9 (22.8)            |
|                     | Side effects of treatment       | 18.1 (13.5)         | 20.8 (15.3)            |
| QLQ-CIPN20          | Autonomic scale                 | 11.1 (15.5)         | 14.0 (20.0)            |
|                     | Motor scale                     | 11.9 (13.5)         | 17.9 (17.6)            |
|                     | Sensory scale                   | 12.3 (15.1)         | 16.7 (19.2)            |

**Abbreviations:** EORTC; European Organization for Research and Treatment of Cancer; QLQ-C30, Quality-of-Life Core; QLQ-CIPN20, QLQ-Chemotherapy-Induced Peripheral Neuropathy; QLQ-MY20, QLQ-Multiple Myeloma; QoL, quality of life.
cohort: Diarrhoea domain in the lenalidomide cohort (mean change (s.d.) of 10.9 (27.1), indicating a worsening of the symptom) and Global health status/QoL domain in the bortezombib cohort (mean change (s.d.) of −8.5 (22.7), indicating a worsening of HRQoL). For all other domains, changes over time did not reach the MID. A slight deterioration in HRQoL was consistently observed over time in both lenalidomide and bortezombib cohorts for all other domains, except Financial difficulties, Pain, Disease symptoms and Future perspective domains, where a slight improvement was observed (Figure 2).

For patients who discontinued the study prior to 6 months owing to disease progression or discontinuation of treatment,

Figure 2. Description of changes in domains scores for the three EORTC questionnaires over time in the lenalidomide and bortezombib cohorts. Bort: bortezombib, with N = 59–62 at month 3, 40–42 at month 6 and 27–29 at discontinuation; Len: lenalidomide, with N = 113–120 at month 3, 90–93 at month 6 and 23–27 at discontinuation; MID corresponding to a meaningful worsening (straight line), and a meaningful improvement (dashed line), MID defined as standard error of measurement for multi-item domains and as 0.5 × SD at baseline for single-item domains.
clinically meaningful declines in HRQoL exceeding the MID were observed in 8 of the 22 domains for the bortezomib cohort (Global health status/QoL, Role functioning, Social functioning, Fatigue, Dyspnoea, Diarrhoea, Motor scale and Sensory scale domains) and in 1 of the 22 domains in the lenalidomide cohort (Motor scale domain) (Figure 2).

**DISCUSSION**

The objective of this prospective, observational European 6-month study was to investigate the HRQoL of patients diagnosed with RRMM receiving second- or third-line lenalidomide or bortezomib treatment. In this real-world setting, the results showed that HRQoL was not substantially impaired under continued treatment with either lenalidomide- or bortezomib-based regimens. Only a change indicating a worsening in the Diarrhoea domain from baseline to month 6 was observed in the lenalidomide cohort, and a change indicating a worsening in the Global health status/QoL domain was observed in the bortezomib cohort.

Patients who discontinued therapy early showed worsening of 8 of the 22 EORCT domains in the bortezomib group but only in 1 domain in the lenalidomide group. One can hypothesise that this could be linked to the fact that patients on bortezomib who discontinued early more frequently have symptoms on relapse compared with patients on lenalidomide.

In our study, we did observe some differences in patient populations, but as this is a real-world setting, patient populations should not be directly compared between treatment cohorts. Other considerations are that several factors may impact a physician’s decision to administer one treatment, such as: patient’s performance status, prior line of therapy, disease characteristics and risk factors, and local national limitations for drug funding approval guidelines.

As an observational study, physician choice could have contributed to the imbalance in the number of patients in each treatment cohort and in dosing schedule for bortezomib with approximately 60% of patients receiving low-dose bortezomib, with a mean (s.d.) average dosage of 3.6 (1.7) mg/m² body surface area per 6-week period.

According to the summary of product characteristics, the recommended treatment duration of bortezomib for RRMM is four cycles followed by four cycles in the case of response or stable disease. In our study, the median duration was approximately 4 months, which is short compared with the label. However, in an Italian retrospective study of 85 patients with second-line therapy, the median treatment duration was 4.1 months, so it may be the case that the actual duration of treatment is lower than the summary of product characteristics in normal clinical use.

Despite being the second most common haematological cancer, MM remains rare. In the context of this study conducted with patients with RRMM starting second- or third-line treatment, small sample sizes could be expected in particular when focussing on subgroups of patients at discontinuation. Further research is required to draw more robust conclusions. Differentiation of HRQoL at discontinuation between patients discontinuing the study due to disease progression and due to discontinuation of treatment would also have been interesting, but it could not be made here owing to the small sample size in the subgroups.

In addition to sample size, another limitation of this study is the low number of patients who received subcutaneous bortezomib, now a common form of treatment in RRMM. This prevented us from drawing meaningful conclusions with regard to QoL between subcutaneously and intravenously administered bortezomib.

Finally, while it is generally recommended using both distribution- and anchor-based approaches and a range of MIDs rather than a unique value, MIDs defined in our study were similar to the ones previously reported by Kvaan et al. in 2010 which were defined using values representing minimal changes that patients regard as a definite improvement or deterioration.

This study focusses on RRMM; however, with the approval of lenalidomide in combination with dexamethasone in frontline MM in February 2015 by the European Medicines Agency (EMA), the question on bortezomib vs lenalidomide has also become relevant for newly diagnosed MM (NDMM) patients. HRQoL at discontinuation due to disease progression was analysed separately from individual HRQoL measurement time points in the FIRST trial comparing lenalidomide in combination with dexamethasone to the combination of melphalan, prednisone and thalidomide in NDMM. Improvements in HRQoL were generally maintained for patients on continued treatment with lenalidomide–dexamethasone. The VISTA study also investigated HRQoL outcomes in NDMM patients randomised to melphalan–prednisone in combination or not with bortezomib, but there was no distinction made between patients completing the study and patients discontinuing the study early. Findings on lenalidomide and bortezomib in the context of HRQoL in RRMM, especially at discontinuation, may well be similar in NDMM given the treatments’ general mode of action. However, it requires further studies on HRQoL in frontline MM in order to confirm this hypothesis. Also, one cannot exclude that the decreased HRQoL observed upon treatment discontinuation, may be partly a result of disease progression and not only a result of a direct drug effect.

Ultimately, there remains no cure for MM, and while we have made significant advances in the length of time that patients diagnosed with MM are living with the disease, it is important to ensure that with this longevity there is an acceptable quality of life, as reflected in their HRQoL functions. This study has enabled us to observe, in a real-world setting, the impact that continuous treatment over a 6-month period had on the patients’ HRQoL scores and showed that they did not substantially deteriorate, despite receiving treatment with associated diverse events that could have potentially impacted patient’s well-being. Importantly, some differences in HRQoL deterioration were observed between bortezomib and lenalidomide at the time of discontinuation of treatment.

**CONFLICT OF INTEREST**

PM, XL, CK, IVB and MTP were Consultant/Advisor for Celgene and received honoraria from Celgene. XL also received honoraria and travel support from Janssen, Takeda, BMS, Novartis, Amgen, Leopharma and Sanofi. PB and PL declare employment and equity ownership in Celgene. During the course of study and manuscript development, PL was an employee of Celgene. HG, former employee of Mapi, and BA, employee of Mapi, were Consultant/Advisor for Celgene and received research funding from Celgene.

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**REFERENCES**

1. Palumbo A, Anderson K. Multiple myeloma. *N Engl J Med* 2011; 364: 1046–1060.
2. Kristinsson SY, Minter AR, Korde N, Tan E, Landgren O. Bone disease in multiple myeloma and precursor disease: novel diagnostic approaches and implications on clinical management. *Expert Rev Mol Diagn* 2011; 11: 593–603.
3. Ferlay J, Soerjomataram I, Ervik M, Dikshit R, Eser S, Mathers C et al. GLOBOCAN 2012 v1.0, Cancer Incidence and Mortality Worldwide: IARC CancerBase No. 11 [Internet]. International Agency for Research on Cancer: Lyon, France, 2013. [cited 11 September 2015]; available from http://globocan.iarc.fr.
4. NCI (National Cancer Institute). Surveillance, Epidemiology, and End Results (SEER) Program. SEER Stat Fact Sheets: Myeloma. [cited 11 September 2015]; available from http://seer.cancer.gov/statfacts/html/myel.html.
5. Lonial S. Relapsed multiple myeloma. *Hematology Am Soc Hematol Educ Program* 2010; 2010: 303–309.
6 Spicka I. Advances in multiple myeloma therapy during two past decades. Comput Struct Biotechnol J 2014; 10: 38–40.

7 Merin NM, Kelly KR. Clinical use of proteasome inhibitors in the treatment of multiple myeloma. Pharmaceuticals (Basel) 2014; 8: 1–20.

8 Poh A. Multiple myeloma gets three new drugs. Cancer Discov 2016; 8: 4.

9 Ghosh N, Bunwai RF, Massa A, Bhutani M. Expanding role of lenalidomide in hematologic malignancies. Cancer Manag Res 2015; 7: 105–119.

10 Anderson KC, Kyle RA, Rajkumar SV, Stewart AK, Weber D, Richardson P et al. Clinically relevant end points and new drug approvals for myeloma. Leukemia 2008; 22: 231–239.

11 US Department of Health and Human Services FDA Center for Drug Evaluation and Research, US Department of Health and Human Services FDA Center for Biologics Evaluation and Research, US Department of Health and Human Services FDA Center for Devices and Radiological HealthGuidance for industry. Guidance for Industry Patient-Reported Outcome Measures: Use in Medical Product Development to Support Labeling Claims 2009. Available at https://www.fda.gov/downloads/drugs/guidances/ucm193282.pdf.

12 Basch E, Abernethy AP, Mullins CD, Reeve BB, Smith ML, Coons SJ et al. Recommendations for incorporating patient-reported outcomes into clinical comparative effectiveness research in adult oncology. J Clin Oncol 2012; 30: 4249–4255.

13 Dimopoulos M, Spencer A, Attal M, Prince HM, Harousseau JL, Dmoszynska A et al. Lenalidomide plus dexamethasone for relapsed or refractory multiple myeloma. N Engl J Med 2007; 357: 2123–2132.

14 Richardson PG, Sonneveld P, Schuster MW, Irwin D, Stadtmueller EA, Facon T et al. Bortezomib or high-dose dexamethasone for relapsed multiple myeloma. N Engl J Med 2005; 352: 2487–2498.

15 Palumbo A, Hajek R, Delforge M, Kroppf M, Petrucci MT, Catalano J et al. Continuous lenalidomide treatment for newly diagnosed multiple myeloma. N Engl J Med 2012; 366: 1759–1769.

16 Weber DM, Chen C, Niezvisky R, Wang M, Belch A, Stadtmueller EA et al. Lenalidomide plus dexamethasone for relapsed multiple myeloma in North America. N Engl J Med 2007; 357: 2133–2142.

17 San Miguel JF, Schlag R, Khuaigea NK, Dimopoulos MA, Shiplberg O, Kroppf M et al. Bortezomib plus melphalan and prednisone for initial treatment of multiple myeloma. N Engl J Med 2008; 359: 906–917.

18 Kvam AK, Waage A. Health-related quality of life in patients with multiple myeloma–does it matter? Haematologica 2015; 100: 704–705.

19 Sonneveld P, Verelst SG, Lewis P, Gray-Schoepfer V, Hutchings A, Nixon A et al. Review of health-related quality of life data in multiple myeloma patients treated with novel agents. Leukemia 2013; 27: 1959–1969.

20 Aaronson NK, Ahmedzai S, Bergman B, Bullinger M, Cull A, Duez NJ et al. The European Organization for Research and Treatment of Cancer QLQ-C30: a quality-of-life instrument for use in international clinical trials in oncology. J Natl Cancer Inst 1993; 85: 365–376.

21 Osoba D, Zee B, Pater J, Warr D, Kaizer L, Latrelle J. Psychometric properties and responsiveness of the EORTC quality of Life Questionnaire (QLQ-C30) in patients with breast, ovarian and lung cancer. Qual Life Res 1994; 3: 353–364.

22 Cocks K, Cohen D, Wisloff F, Sezer O, Lee S, Hillebrandt JG, Delattre JY et al. An international field study of the reliability and validity of a disease-specific questionnaire module (the QLQ-MY20) in assessing the quality of life of patients with multiple myeloma. Eur J Cancer 2007; 43: 1670–1678.

23 Stead ML, Brown JM, Velikova G, Kaasa S, Wisloff F, Child JA et al. Development of an EORTC questionnaire module to be used in health-related quality-of-life assessment for patients with multiple myeloma. European Organization for Research and Treatment of Cancer Study Group on Quality of Life. Br J Haematol 1999; 104: 605–611.

24 Postrma TJ, Aaronson NK, Heimans JJ, Muller MJ, Hildebrandt JG, Delattre JY et al. The development of an EORTC quality of life questionnaire to assess chemotherapy-induced peripheral neuropathy: the QLQ-CIPN20. Eur J Cancer 2005; 41: 1135–1139.

25 Jaeschke R, Singer J, Guyatt GH. Measurement of health status. Ascertain the minimal clinically important difference. Control Clin Trials 1989; 10: 407–415.

26 Norman GR, Sloan JA, Wyrciwn KW. Interpretation of changes in health-related quality of life: the remarkable universality of half a standard deviation. Med Care 2003; 41: 582–592.

27 Pantani L, Zamagni E, Zannetti BA, Pezzi A, Tacchetti P, Brioli A et al. Bortezomib and dexamethasone as salvage therapy in patients with relapsed/refractory multiple myeloma: analysis of long-term clinical outcomes. Ann Hematol 2014; 93: 123–128.

28 Petrucci MT, Finsinger P, Chisini M, Gentilini F. Subcutaneous bortezomib for multiple myeloma treatment: patients’ benefits. Patient Prefer Adherence 2014; 8: 939–946.

29 Kvam AK, Wisloff F, Fayers PM, Gentilini F. Minimal important differences and response shift in health-related quality of life; a longitudinal study in patients with multiple myeloma. Health Qual Life Outcomes 2010; 8: 79.

30 European Medicines Agency - Committee for Medicinal Products for Human Use (CHMP). Revlimid: Summary of opinion EMA/CHMP/91664/2015 Rev 1 issued on 18 February 2015. [cited 24 June 2016]; Available from http://www.ema.europa.eu/docs/en_GB/document_library/Summary_of_opinion/human/000717/WC500179310.pdf.

31 Benboubker L, Dimopoulos MA, Dispensieri A, Catalano J, Belch AR, Cavo M et al. Lenalidomide and dexamethasone in transplant-ineligible patients with myeloma. N Engl J Med 2014; 371: 906–917.

32 Delforge M, Dhawan R, Robinson Jr D, Meunier J, Regnault A, Essertine DL et al. Health-related quality of life in elderly, newly diagnosed multiple myeloma patients treated with VMP vs. MP; results from the VISTA trial. Eur J Haematol 2012; 89: 16–27.

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