Clinicopathological characteristics of extremely young Korean multiple myeloma patients: therapeutic implications

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Received : August 3, 2016
Revised : October 18, 2016
Accepted : February 4, 2017

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Background/Aims: Although multiple myeloma (MM) is typically a disease of the elderly, a certain subset of extremely young patients exists. It is necessary to establish clinicopathological characteristics for this population.

Methods: We reviewed the medical records of MM patients whose age was 40 years or younger at diagnosis.

Results: A total of 32 patients were analyzed (male to female ratio 19:13, median age 37 years). According to International Staging System, 29%, 48%, and 16% were in stage I, II, and III, respectively. Light chain myeloma accounted for 30%. Clinically significant anemia, hypercalcemia, azotemia, and hypoalbuminemia were present in 29%, 28%, 13%, and 28%, respectively. Three or more lytic bone lesions were detected in 45% of the patients, whereas 13% had no lytic bone lesions. Regarding treatment, 79% of patients received autologous hematopoietic stem cell transplantation. After a median follow-up duration of 64 months, the 1-, 3-, and 5-year overall survival (OS) rates were 84%, 62%, and 54%, respectively. The median OS was 61 months for the entire cohort.

Conclusions: In our study, MM patients aged 40 years or younger at diagnosis showed no superior survival compared to those of the moderately elderly patients based on historical data.

Keywords: Multiple myeloma; Young adult; Hematopoietic stem cell transplantation; Treatment outcome

INTRODUCTION

Multiple myeloma (MM) is a disease of the elderly; it shows a rapidly increasing incidence with increased age. The peak age-specific incidence has been observed in people older than 80 years of age, where it reaches approximately 40 per 100,000 person-years [1]. Although MM is rare in young people, patients diagnosed before the age of 50 years account for 10% of the overall incidence, and approximately 2% are diagnosed before 40 years of age [2]. Various reports have examined the different clinical and laboratory features and treatment outcome for such young MM patients [2-8]. Although some of these data are debatable, those patients generally seem to be at lower International Staging System (ISS) and Durie-Salmon Staging System (DSS) stages at diagnosis, more frequently present with light-chain myeloma, have a lower proportion of measurable M-protein and bone marrow (BM) plasmacytosis, and have a lower frequency of related tissue injuries, including
hypercalcemia, azotemia, and anemia, compared with conventional elderly patients [3,4].

The overall survival (OS) of MM shows remarkable variation, ranging from a few months to more than 10 years. Because of this prognostic diversity, the necessity to stratify patients into divided risk groups has emerged. The prognosis of MM patients has been associated with several parameters of prognostic significance [2]. Those include the initial performance status, the ISS and DSS stage at diagnosis, indicators of related tissue injuries, plasma cell labeling index, and certain chromosomal aberrations [9,10]. However, the role of age as an independent prognostic factor, especially when the life expectancy of the corresponding general population is taken into account, has been part of debate. In a previous study comparing patients younger than 50 years at diagnosis with those aged 50 years or older, the younger group showed better survival in terms of observed survival rates; however, that group appeared to have a significantly worse outcome when the mortality was standardized with the expected death rate of the corresponding general population [2]. A more recently reported retrospective study also suggested that MM patients diagnosed before 40 years of age do not show significant differences in OS after autologous hematopoietic stem cell transplantation (aHSCT) compared to those aged 41 to 65 years [8]. In contrast, some large studies suggested that old age is associated with short survival [4,11]. Therefore, the importance of age as an independent prognostic factor in MM is still unclear.

Although evidence from several studies has implied that Asian MM patients could show clinical behavior and adverse drug reaction patterns that differ from those of Western patients [12-14], studies on the presenting features and long-term outcomes of young Asian MM patients have rarely been conducted. In this study, we analyzed the clinicopathological characteristics, the outcomes of several treatment approaches, and the prognosis of Korean MM patients diagnosed at age 40 years or younger.

Patients diagnosed with MM at the age of 40 years or younger from January 2000 to February 2015 were retrieved through a retrospective search of electronic medical records (EMRs). The Institutional Review Board of Seoul National University Hospital (SNUH) approved the study protocol (IRB approval number H-1305-629-491). All procedures were in accordance with the ethical standards of the institutional research committee and with the Declaration of Helsinki revised in the 2013 World Medical Association. The International Myeloma Working Group’s diagnostic and response criteria on the date of diagnosis and the response evaluation for each patient were used to enroll the patient and analyze the data regarding the treatment response. Monoclonal gammopathy of unknown significance, smoldering myeloma, amyloidosis, and POEMS (polyneuropathy, organomegaly, endocrinopathy, monoclonal gammopathy, and skin changes) syndrome were excluded.

The stage at diagnosis was classified according to both ISS, described by International Myeloma Working Group in 2005, and DSS, first described in 1975. The number and location of osteolytic lesions were analyzed by reviewing the skeletal survey images from X-rays and magnetic resonance imaging, if available. BM examination results were collected, including the proportion of plasma cell infiltration, conventional cytogenetics, and fluorescence in situ hybridization (FISH) for chromosomal aberrations with prognostic implications [15]. The starting date, regimen, maximal response, and the date of progression or death for first-line treatment, the bortezomib-containing regimen, and aHSCT were also reviewed. The causes of death were categorized into MM progression, infection, thrombosis, or others.

Statistical analysis
Progression-free survival (PFS) was defined as the interval from the beginning of each treatment to progression or death from any cause, whichever occurred first. OS was defined as the interval from diagnosis to death from any cause. Patients whose date of progression or death was uncertain or who were progression-free or alive on the date of the data collection were censored in the survival analysis using the last date they were known to be progression-free and alive. The PFS after first-line treatment, aHSCT and bortezomib-containing regimen and OS were calculated for each evaluable patient. A survival

METHODS

Eligible patients and data collection
Four institutes in South Korea participated in this study.
The curve was constructed using the Kaplan-Meier method for the entire cohort based on the observed mortality and used to estimate the median PFS and OS, 1-, 3-, and 5-year PFS and OS rates. To compare survival between two groups, a log-rank test was used. The results were considered statistically significant at two-sided \( p < 0.05 \). STATA statistical software version 12.0 (StataCorp LP, College Station, TX, USA) was used for computation.

**RESULTS**

**Baseline characteristics**

A total of 49 patients were retrieved via computer-based searching of the EMRs of four institutions. Among those patients, 13 and four were excluded from the analysis because of the erroneous input of the diagnosis and insufficient medical records, respectively. Of the 32 patients who were included in the analysis, 25 were treated at SNUH, four at Korea University Anam Hospital, two at Korea University Ansan Hospital, and one at Korea University Guro Hospital.

Patient characteristics are summarized in Table 1. The median age at diagnosis for all 32 patients was 37 years (range, 17 to 40). Of those, four patients (two males and two females) were younger than 30 years at diagnosis. The age distribution at diagnosis was similar irrespective of gender. Twenty patients were diagnosed before 2010 while the remaining patients were diagnosed in 2010 or thereafter. The most common subtype of M-protein was immunoglobulin G (IgG), followed by IgA and IgD. Light chain myeloma accounted for 30% of patients. Data regarding specific locations of bony lesions were available for 20 patients treated at SNUH. The most commonly involved bones were the vertebrae and skull, accounting for 60% and 55% of the patients with bone involvement, respectively; these were followed by the ribs, pelvis, femur, humerus, and scapulae, accounting for 40%, 30%, 30%, 20%, and 10%, respectively. Karyotypes of the BM cells determined by the conventional G-banding technique were available for 23 patients. The chromosomal abnormality was observed in 30% of the patients according to conventional karyotyping. When considering both of the conventional karyotyping and FISH, the proportion grew up to 59%. The prevalence of each adverse genetic lesion detected using FISH is

| Characteristic                  | Frequency/Totala (%) |
|--------------------------------|----------------------|
| **Sex**                        |                      |
| Male                           | 19/32 (59)           |
| Female                         | 13/32 (41)           |
| **ECOG**                       |                      |
| 0                              | 4/31 (13)            |
| 1                              | 13/31 (42)           |
| 2                              | 8/31 (26)            |
| 3                              | 5/31 (16)            |
| 4                              | 1/31 (3)             |
| **ISS**                        |                      |
| I                              | 10/31 (32)           |
| II                             | 15/31 (48)           |
| III                            | 6/31 (19)            |
| **DSS**                        |                      |
| I                              | 5/31 (16)            |
| II                             | 8/31 (26)            |
| III                            | 18/31 (58)           |
| **Subtype**                    |                      |
| A                              | 27/31 (87)           |
| B                              | 4/31 (13)            |
| **Subtype Light chain**        |                      |
| κ                               | 4/30 (13)            |
| λ                               | 5/30 (17)            |
| **Whole immunoglobulin**       |                      |
| Immunoglobulin G               | 14/30 (47)           |
| Immunoglobulin A               | 5/30 (17)            |
| Immunoglobulin D               | 2/30 (7)             |
| κ                               | 13/30 (43)           |
| λ                               | 8/30 (27)            |
| **Hemoglobin < 10 g/dL**       | 9/31 (29)            |
| **Platelet < 130 × 10^3/μL**    | 7/31 (23)            |
| **Calcium** ≥ 10 mg/dL         | 9/32 (28)            |
| **Creatinine ≥ 2 mg/dL**        | 4/32 (13)            |
| **Albumin < 3.5 g/dL**          | 9/32 (28)            |
| **B2MG ≥ 3.5 mg/dL**            | 14/29 (48)           |
| **LDH above normal**           | 10/27 (37)           |
| **Lytic bone lesions**         |                      |
| 0                              | 4/31 (13)            |
| 1–2                            | 13/31 (42)           |
| ≥ 3                            | 14/31 (45)           |
| **BM plasma cells, %**         |                      |
| 10–30                          | 15/27 (56)           |
| ≥ 30                           | 7/27 (26)            |
| **Biopsy-proven plasmacytoma** |                      |
| 20/31 (65)                     |                      |

ECOG, Eastern Cooperative Oncology Group; ISS, International Staging System; DSS, Durie-Salmon Staging System; B2MG, β-2 microglobulin; LDH, lactate dehydrogenase; BM, bone marrow.

aTotal indicates the number of patients with sufficient data for categorization.
bCorrected calcium level, calculated from measured calcium + 0.8 × (4 – albumin).
summarized in Table 2. The chromosomal aberrations included in the table were selected based on previous reports showing their significant association with shorter PFS and OS [10,16]. Regarding initial manifestation, 20 (65%) out of the 31 patients evaluated for detectable tumors using an imaging study were found to have plasmacytomas which were pathologically confirmed. Those patients showed a tendency toward a low probability of BM plasmacytosis (BM plasma cell fraction ≥ 10%; \( p = 0.137 \) with a Fisher exact test).

**Table 2. Frequency of adverse chromosomal alterations**

| Chromosomal alteration | Genetic involvement | Frequency/Total\(^a\) (%) |
|------------------------|---------------------|-------------------------|
| del(17p)               | p53 deletion        | 1/9 (11)                |
| t(4;14)                | IGH/FGFR3 rearrangement | 1/10 (10)          |
| t(14;16)               | IGH/MAF rearrangement | 0/11 (0)               |
| t(14;20)               |                     | 0/6 (0)                 |
| 1q gain               |                     | 4/15 (26)               |
| del(13q)              | rbi deletion        | 4/17 (24)               |
| del(9)                | p16 (CDKN2A) deletion | 1/16 (6)               |

FISH, fluorescence in situ hybridization.

\(^a\)Total indicates the number of patients tested for each chromosomal alteration.

**Response to treatment**

Regarding treatment, the first-line chemotherapeutic regimens administered are summarized in Table 3. Two patients were transferred to other institutions right after diagnosis, leaving no records regarding treatment. Either vincristine/doxorubicin/dexamethasone (VAD) alone or two cycles of VAD subsequently followed by bortezomib/thalidomide/dexamethasone (VTD) was the most frequently used regimen, followed by thalidomide/dexamethasone (TD). Among the patients whose response to the first-line treatment could be evaluated, either a complete response (CR) or a very good partial response (VGPR) was achieved in 14 patients (48%) after first-line chemotherapy with or without subsequent aHSCT. Of these patients, five were not eligible for aHSCT because of early progression before stem cell collection (\( n = 2 \)), the patient’s refusal (\( n = 2 \)), and suicide immediately after the initial response evaluation (\( n = 1 \)). Regarding the 26 patients with relevant data, 18 (69%) eventually experienced progressive disease (PD). The median PFS after first-line treatment was 16 months (95% confidence interval [CI], 9 to 30) with 1-, 3-, and 5-year PFS rates of 65% (95% CI, 42% to 81%), 27% (95% CI, 16% to 48%), and 14% (95% CI, 3% to 34%), respectively (Fig. 1A). The patients who received aHSCT as a first-line treatment showed a tendency toward improved PFS compared with the patients who did not (median 17 months vs. 3 months, \( p = 0.101 \)) (Fig. 1B).

**Table 3. First-line therapeutic regimens**

| Regimen          | Frequency/Total\(^a\) (%) | CR or VGPR, n (% rate) | OR, n (% rate) |
|------------------|--------------------------|------------------------|----------------|
| VAD ± VTD        | 12/30 (40)               | 5/12 (42)              | 9/12 (75)      |
| TD               | 8/30 (27)                | 5/8 (62.5)             | 6/8 (75)       |
| CTD              | 2/30 (7)                 | 1/2 (50)               | 1/2 (50)       |
| MP               | 2/30 (7)                 | 0/2 (0)                | 1/2 (50)       |
| VD               | 2/30 (7)                 | 1/2 (50)               | 1/2 (50)       |
| VCD              | 1/30 (3)                 | 0/1 (0)                | 0/1 (0)        |
| MPT              | 1/30 (3)                 | 1/1 (100)              | 1/1 (100)      |
| CD               | 1/30 (3)                 | 1/1 (100)              | 1/1 (100)      |
| Dexamethasone only | 1/30 (3)               | NA                     | NA             |

CR, complete response; VGPR, very good partial response; OR, overall response; VAD, vincristine/doxorubicin/dexamethasone; VTD, bortezomib/thalidomide/dexamethasone; TD, thalidomide/dexamethasone; CTD; cyclophosphamide/thalidomide/dexamethasone; MP, melphalan/prednisolone; VD, bortezomib/dexamethasone; VCD, bortezomib/cyclophosphamide/dexamethasone; MPT, melphalan/prednisolone/thalidomide; CD, cyclophosphamide/dexamethasone; NA, not available.

\(^a\)Total indicates the number of patients with sufficient data for categorization.
Bortezomib/dexamethasone (VD) with or without other agents (such as thalidomide, lenalidomide, doxorubicin, or cyclophosphamide) was administered in 16 patients, of whom 31% achieved CR or VGPR, 25% achieved partial response (PR), 25% achieved stable disease, and 19% achieved PD. Only three of these patients received a bortezomib-containing regimen as a first-line treatment. Radiation therapy was performed in seven patients for the palliation of symptoms.

Among the 29 patients with relevant data, 23 (79%) received aHSCT at any stage of treatment. Of these, 18 (62%) received it as a part of a first-line treatment. Four patients received aHSCT more than once, either as a tandem transplantation (n = 3) or at separate stages (n = 1). Before aHSCT, 36% of the patients were in CR or VGPR, and 55% were in PR. After aHSCT, 64% achieved CR or VGPR, and 23% achieved PR, yielding an overall response rate of 87%. Throughout the entire course of treatment, six patients did not receive aHSCT: two refused aHSCT; two were preparing for allogeneic hematopoietic stem cell transplantation (alloHSCT) because of secondary aplastic anemia and MDS; one died without having responded to chemotherapy; and one committed suicide during the induction of chemotherapy.

Overall survival

With a median follow-up duration of 64 months (interquartile range, 35 to 105) for the entire cohort, 15 out of 32 patients died. The 1-, 3-, and 5-year OS rates were 84% (95% CI, 66% to 93%), 62% (95% CI, 42% to 77%), and 54% (95% CI, 34% to 71%), respectively. The median OS was 61 months (95% CI, 33 to not reached months) for the entire cohort (Fig. 1C). No significant difference in OS was observed between subgroups divided by whether treated

Figure 1. (A) Progression-free survival (PFS) after first-line treatment. (B) PFS after first-line treatment stratified by whether the patients received autologous hematopoietic stem cell transplantation (aHSCT) as part of the first-line treatment. (C) Overall survival (OS). (D) OS stratified by whether the patients received aHSCT as part of the first-line treatment.
with aHSCT as the first-line treatment or not (median 50 months vs. 75 months, \( p = 0.596 \)) (Fig. 1D). There was a tendency toward improved OS for patients with a lower ISS stage (median 75 months [I] vs. 50 months [II] vs. 33 months [III], \( p = 0.363 \)), a \( \kappa \) over a lambda subtype (median 75 months vs. 42 months, \( p = 0.161 \)), whole immunoglobulin subtype over light chain myeloma (median 75 months vs. 29 months, \( p = 0.219 \)), and those with biopsy-proven plasmacytoma at diagnosis (median 42 months vs. 61 months, \( p = 0.340 \)); though, no statistical significance was observed (Fig. 2).

The most common cause of death was myeloma progression (67%), followed by infection (20%). Therapy-related mortality after first-line induction chemotherapy or aHSCT, defined as death within 4 weeks after each treatment, was not observed in our cohort.

**DISCUSSION**

To the best of our knowledge, our study is the first case series report focusing on MM patients aged 40 years or younger at diagnosis in South Korea. In this study, we analyzed clinicopathological characteristics of extremely young MM patients to detect and accentuate any small differences in clinical features compared with the relatively well-studied population of elderly patients. Although several studies have been conducted to address the distinct clinical features of these patients in other ethnic groups, many large-scale studies used a cutoff age of 50 years [2,4]. Important clinical characteristics of cohorts in various previous studies being compared with our cohort are summarized in Table 4. Several presenting features in our cohort were in accordance with previous reports [2-4,8,17]. To be specific, the patients...
were in relatively low ISS stages and had a good perfor-
mance status at diagnosis compared with the historical
controls of elderly MM patients (32%, 48%, and 19% vs.
26%, 39%, and 34% for stage I, II, and III, respectively,
in patients aged 50 years or older) [4]. The proportion
of lytic bone lesions involving three or more parts (45%)
was comparable to that in another large MM cohort of
patients younger than 50 (48%) [4].

However, some notable findings were revealed in
our study data. First of all, a remarkably large propor-
tion (30%) was confirmed to have light chain myeloma,
compared to 12% and 13% among patients younger than
50 [2,4] and 12% and 21% among those aged 40 years
or younger in other studies [5,8]. Furthermore, the BM
plasma cell fraction below 30% was observed in 74%
compared to 28% and 42% among similar age groups in
previous studies [4,5]. Organ injuries indicated by ane-
mia, hypercalcemia, azotemia, and hypoalbuminemia
were observed at a low frequency compared to both of
the young and old age groups in a previous large MM
cohort study [4]. Most importantly, despite many favor-
able prognostic factors mentioned above such as low
stages at diagnosis, good performance status, and a low
frequency of tissue injuries, the median OS was only 61
months. This was much shorter than those observed in
other two cohorts, one including patients aged 41 to 65
(longer than 80.7 months) and the other including pa-
tients aged 60 to 65 (89 months) [8,11]. Therefore, we
could not find any tendency toward improved survival
for young patients compared to moderately elderly pa-
tients despite previous reports suggesting that young
age is a good prognostic factor [4,11].

Table 4. Comparison of patient characteristics between several multiple myeloma cohort studies

| Characteristic                  | This cohort (n = 32) | Ludwig et al. (2008) [4] | Corso et al. (1998) [2] | Cheema et al. (2009) [8] | Chretien et al. (2014) [11] |
|--------------------------------|---------------------|--------------------------|------------------------|--------------------------|----------------------------|
|                                | Male sex, %          | Younger (n = 1,689) | Older (n = 8,860)     | Younger (n = 61) | Older (n = 295) | Younger (n = 38) | Older (n = 608) | Older (n = 813) |
| Male sex, %                    | 59                  | 61                       | 57                     | 52                       | 57                       | 61                       | 60                       | NA                       |
| Age, yr, median (range)        | 37 (17–40)          | 36 (20–49)               | 62 (50–93)             | 45 (33–49)               | 61 (50–87)               | 37.2 (29–40)             | 56.4 (41–65)             | NA (60–65)               |
| ISS, %                         |                     |                          |                        |                          |                          |                          |                          |                           |
| I                              | 32                  | 39                       | 26                     | 30                       | 34                       | 48                       | NA                       | 30                       |
| II                             | 48                  | 35                       | 39                     | 13                       | 20                       | NA                       | NA                       | 40.40                    |
| III                            | 19                  | 27                       | 34                     | 57                       | 46                       | NA                       | NA                       | 29.70                    |
| Subtype, %                     | Light chain only    | 30                       | 13                     | 10                       | NA                       | NA                       | 21                       | NA                       |
| Immunoglobulin G               | 47                  | 60                       | 60                     | 65                       | 67                       | 53                       | NA                       | NA                       |
| Immunoglobulin A               | 17                  | 21                       | 25                     | 18                       | 20                       | 18                       | NA                       | NA                       |
| Immunoglobulin D               | 7                   | 3                        | 3                      | 1                        | 1                        | 8                        | NA                       | NA                       |
| Hemoglobin < 10 g/dL, %        | 29                  | 37                       | 41                     | NA                       | NA                       | NA                       | NA                       | NA                       |
| Ca²⁺ 10 mg/dL, %               | 28                  | 33                       | 34                     | NA                       | NA                       | NA                       | NA                       | NA                       |
| Cr ≥ 2 mg/dL, %                | 13                  | 15                       | 17                     | 8                        | 14                       | 25                       | 16                       | NA                       |
| BMPC 30%, %                    | 26                  | NA                       | NA                     | NA                       | NA                       | 72                       | 74                       | NA                       |
| Bone disease, %                | 87                  | 79                       | 78                     | 43                       | 34                       | 76                       | 74                       | NA                       |
| Upfront aHSCT, %               | 62                  | 41                       | 24                     | NA                       | NA                       | 100b                     | 31.60                    | 100c                     |
| Median OS, mon                 | 61                  | NA                       | NA                     | NA                       | NA                       | 81.4                     | > 80.7d                  | 89                       |

NA, not available; ISS, International Staging System; BMPC, bone marrow plasma cell; aHSCT, autologous hematopoietic stem cell transplantation; OS, overall survival.

aCorrected calcium level, calculated from measured calcium + 0.8 × (4 – albumin).
bUpfront aHSCT was included in the inclusion criteria.
cAll patients were treated with 3–4 cycles of induction chemotherapy followed by 1–2 courses of high-dose melphalan.
dMedian OS was recorded only from the date of aHSCT and not from the date of diagnosis.
might be partially due to the high proportion of light chain myeloma in our cohort which is associated with poor survival and relatively low proportion of patients who received upfront aHSCT (Table 4). However, considering that most previous studies included patients who had been diagnosed at least 5 to 10 years earlier than our cohort when recently developed highly effective therapeutic agents such as bortezomib, thalidomide, and lenalidomide were not available, the lack of a superior outcome in our cohort compared with these series of moderately elderly patients is notable.

A few case reports or case series have implied the existence of some unusual features of manifestation of young MM patients. For instance, a specific subgroup has been shown to manifest initially with multiple osteolytic lesions or extramedullary involvement without BM infiltration of myeloma cells. These patients seemed to have a lower tumor burden and better survival than conventional MM patients of similar ages [3,5]. Another subgroup has been characterized with multifocal liver sinusoidal infiltration of myeloma cells, presenting with fatigue, weight loss, and abdominal distension as initial symptoms [6]. In response to these findings, we separately analyzed the patients who initially had biopsy-proven plasmacytoma. The low proportion of BM plasmacytosis in this subset of patients suggests that a part of this subgroup has presented with only extramedullary plasmacytoma without BM infiltration of neoplastic cells. However, these patients seemed to have short survival compared with those without plasmacytoma at diagnosis, which is not accordant with the previous reports indicating a favorable outcome of macrofocal MM [3,5].

High-dose therapy (HDT) followed by aHSCT has long been considered a standard treatment for MM patients younger than 65 years of age and shown to be superior to conventional combination chemotherapy in terms of response rate, PFS, and OS [18]. Therefore, virtually all patients enrolled in this study were candidates for HDT plus aHSCT. Nevertheless, 10% of the patients were initially treated with a regimen containing melphalan, a myelotoxic agent that can compromise the BM stem cell reservoir and thus hinder stem cell harvest. One of the patients was diagnosed in 2008 and eventually received aHSCT after second-line VD, but the remaining two patients, diagnosed in 2002 and 2007, respectively, did not receive aHSCT through their entire course of treatment. Considering that the superiority of HDT plus aHSCT over conventional therapy was demonstrated and reported in 2003 in a multicenter randomized controlled trial [18], it is unclear why these patients were initially treated with myelotoxic agents. Although a full medical history was not available for these patients, a reasonable explanation might be that their performance or comorbidity was too poor to deliver HDT at the initiation of treatment.

Our study has several limitations. First, this study was based on a small number of patients; therefore, the results should be treated with caution. Second, the source of data is a retrospective medical chart review, which made many clinical and laboratory parameters for individual patients unavailable. Third, because the national death registry data were available only for the 25 patients treated at SNUH, some patients who might have already died have been censored in the survival analysis.

Despite the limitations, a strength of our study is that a vast range of clinicopathological characteristics were addressed comprehensively for the entire cohort. The proportions of missing data were small for each analyzed variable. Although the number of studied patients is small, this study is still valuable considering the rarity of reports regarding young MM patients in Asian population. Although many laboratory parameters suggested a good prognosis in our cohort, those were not actually linked to an OS benefit compared to the moderately elderly patients in historical data [8,11]. Our study implies that an extremely young age at diagnosis does not always predict a good response to treatment. Rather, heterogeneous groups of individual patients may exist within those young MM patients. Thus, for those who are expected to show a poor response to current treatments despite their young age, the efficacy of an alternative therapeutic strategy, such as frontline alloHSCT, should be further investigated. The reason for this prognostic heterogeneity remains unclear.

**KEY MESSAGE**

1. In Korean multiple myeloma patients aged 40 years or younger, light chain myeloma accounted for a larger proportion while bone marrow plasmacytosis and end organ injuries were less
Conflict of interest
No potential conflict of interest relevant to this article was reported.

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