Analysis of Time to Next Treatment in Melflufen and Dexamethasone-Treated Patients With Relapsed/Refractory Multiple Myeloma

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BACKGROUND

• In a patient with progressing relapsed/refractory multiple myeloma (RRMM), an important clinical objective is to achieve durable disease stabilization (stable disease [SD] or better), especially in patients with moderate to low disease burden. With each relapse, a patient's prognosis worsens, and time to the next relapse decreases^{1,2}

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- Time to next treatment (TTNT) reflects time of disease stabilization in patients with RRMM and may be a relevant parameter of clinical benefit for patients, clinicians, and
- TTNT is an important parameter to help assess cost of care and is used in real-world evidence (RWE) to assist treatment decisions and support economic reimbursement modeling^{1,3,4}
- Longer TTNT has previously been associated with lower costs in
- Melflufen is a lipophilic peptide-conjugated alkylator that rapidly delivers a highly cytotoxic payload into myeloma cells through peptidase activity (**Figure 1**)⁵⁻⁹
- O-12-M1 is a phase 1/2 study with melflufen plus dexamethasone in 62 patients with RRMM who had ≥2 prior lines of therapy, prior exposure to at least an IMiD and a proteasome inhibitor, and disease progression on last line of therapy. Final study results were presented
- In O-12-M1, cycle lengths of 21 and 28 days were assessed, and a cycle length of 28 days was determined to be optimal with regard to dose intensity and safety profile¹⁰
- The phase 2 part of O-12-M1 showed encouraging activity¹⁰
- Median 4-5 prior lines of therapy, 44% high-risk cytogenetics, 67% double refractory, 44% pomalidomide refractory, 60% ISS stage 2-3
- Overall response rate (ORR; ≥partial response), 31%; median duration of response (DOR), 8.4 months
- Clinical benefit rate (CBR; ≥minimal response), 49%
- Median progression-free survival (PFS), 5.7 months - Median overall survival (OS), 20.7 months

OBJECTIVES

METHODS

N=45

Patients with RRMM and

≥2 prior lines of therapy including

lenalidomide and bortezomib and

PD on or within 60 days of

completion of last therapy

- The most common grade 3 and 4 treatment-emergent adverse events (TEAEs) were hematologic. Grade 3 and 4 nonhematologic toxicity was infrequent with an infection rate of 9% and no bleeding
- Patients on the 21-day cycle had substantially more grade 3 and 4 TEAEs than patients on the 28-day cycle

exploratory, post hoc analysis of the phase 2 part of the O-12-M1 study

Figure 2. Phase 2 O-12-M1 Study Design (NCT01897714)

PFS, progression-free survival; PD, progressive disease; ORR, overall response rate; OS, overall survival; RRMM, relapsed/refractory multiple myeloma.

• To assess TTNT with melflufen plus dexamethasone in patients with RRMM in an

 Including patients with SD, a large proportion of patients treated with melflufen had clinical benefit in O-12-M1¹⁰ Disease stabilization (≥SD), 76%

Peptidase-enhanced activity in multiple myeloma cells

The hydrophilic alkylator

payloads are entrapped⁷⁻⁹

Melflufen is 50-fold more potent than melphalan in myeloma cells in vitro

due to increased intracellular alkylator activity^{5,7}

• To describe subsequent treatments in an RRMM patient population with a median of 4 to

Melflufen 40 mg was administered intravenously on day

1 of each 21- or 28-day cycle with oral dexamethasone

40 mg given weekly for up to 8 cycles or longer at the

TTNT was retrospectively reviewed and was defined in

dexamethasone to first subsequent therapy or death,

whichever occurred first. An analysis of TTNT, in which

deaths were censored, was performed to allow comparison

with historical data (where both methods have been used)

by International Myeloma Working Group criteria

Response was assessed by the investigator at each cycle

line with guidelines as time from start of melflufen plus

discretion of the investigator and sponsor

Figure 1. Melflufen Mechanism of Action

Melflufen is rapidly taken

up by myeloma cells due

to its high lipophilicity^{5,7}

pFPhe, para-fluoro-L-phenylalanine.

Once inside the myeloma cell,

melflufen is immediately

cleaved by peptidases⁷⁻⁹

5 prior lines of therapy

After PD or start of

patients were

followed for survival

every 3 months for

up to 24 months

- Given the large proportion of patients who achieved disease stabilization, melflufen has
- potential to delay progressive disease (PD) and extend TTNT in patients with RRMM

Peptidases are expressed in several

cancers, including multiple myeloma¹¹⁻¹³

Melflufen rapidly induces

rreversible DNA damage

leading to apoptosis of

Melflufen

Peptidase

pFPhe (carrier)

Alkylator payload

• In total, 27 out of 41 patients (66%) received subsequent therapy (4 patients were still progression free at the time of data cutoff), which is in line with previously reported data in advanced RRMM in which 39% to 72% of patients received subsequent therapy¹⁴⁻¹⁶

Reasons for discontinuation were significantly different for patients on the

- 9 Patients who discontinued because of AEs had grade 4 thrombocytopenia after the last cycle of melflufen (all in the 21-day cycle length group), with a median PFS of 9.3 months (95% CI, 6.5-not calculable [NC]), median DOR of 9.0 months (95% CI, 3.0-NC), and a median time between progression and start of subsequent therapy of 6 days
- 8 out of the 9 patients (89%) received subsequent therapy

As of 9 November 2017, 45 patients were treated in O-12-M1

At data cutoff, 44 patients (98%) had discontinued melflufen plus

RESULTS

dexamethasone (Table 1)

21-day cycle than on the 28-day cycle

Table 1. Patient Disposition (and Supplemental Safety Information)

Disposition, n (%)	ITT (N=45)	21-Day Cycle (n=28)	28-Day Cycle (n=17)
Discontinued treatment	44 (98)	28 (100)	16 (94)
Reason for discontinuation AE PD Lack of response Death Completed 8 cycles	18 (40) 14 (31) 1 (2) 2 (4) 9 (20)	16 (57) 7 (25) 0 0 5 (18)	2 (12) 7 (41) 1 (6) 2 (12) 4 (24)
Ongoing treatment	1(2)	0	1(6)
Grade 3/4 thrombocytopeniab	28 (62)	24 (86)	4 (24)

AE, adverse event; ITT, intention-to-treat; PD, progressive disease Discontinuation rate due to AE was high in the 21-day cycle group due to thrombocytopenia. Please see supplemental safety information.

Table 2. Baseline Patient Characteristics for the ITT Population and by Dosing Cycle Length

Characteristic	ITT (N=45)	21-Day Cycle (n=28)	28-Day Cycle (n=17)	
Median age (range), years	66 (47-78)	64 (48-74)	70 (47-78)	
Gender (male/female), %	67/33	64/36	71/29	
ISS stage at study entry (I/II/III), %	33/40/20	32/43/21	35/35/18	
Median no. prior lines (range)	4 (2-14)	4 (2-14)	4 (2-14)	
Albumin level at study entry ≥3.5 g/dL, n (%) <3.5 g/dL, n (%)	31 (69) 14 (31)	18 (64) 10 (36)	13 (76) 4 (24)	
High-risk cytogenetics, ^a n (%)	20 (44)	12 (43)	8 (47)	
High LDH (1.5×ULN), n (%)	23 (51)	15 (54)	8 (47)	
Single refractory, ^b n (%)	41 (91)	25 (89)	16 (94)	
Double refractory, n (%)	30 (67)	18 (64)	12 (71)	
Triple refractory,d n (%)	3 (7)	2 (7)	1(6)	
Alkylator refractory, e n (%)	24 (53)	16 (57)	8 (47)	

ISS, International Staging System; ITT, intention-to-treat; LDH, lactate dehydrogenase; PI, proteasome inhibitor; ULN, upper limit of normal.

^aDefined as del(17p), t(4;14), t(14;16), t(14;20), or gain(1q). ^bAt least 1 PI or IMiD. ^cAt least 1 PI and IMiD. ^dAt least 1 PI and IMiD and daratumumab

Median TTNT when censoring for deaths was 10.6 months (95% CI,

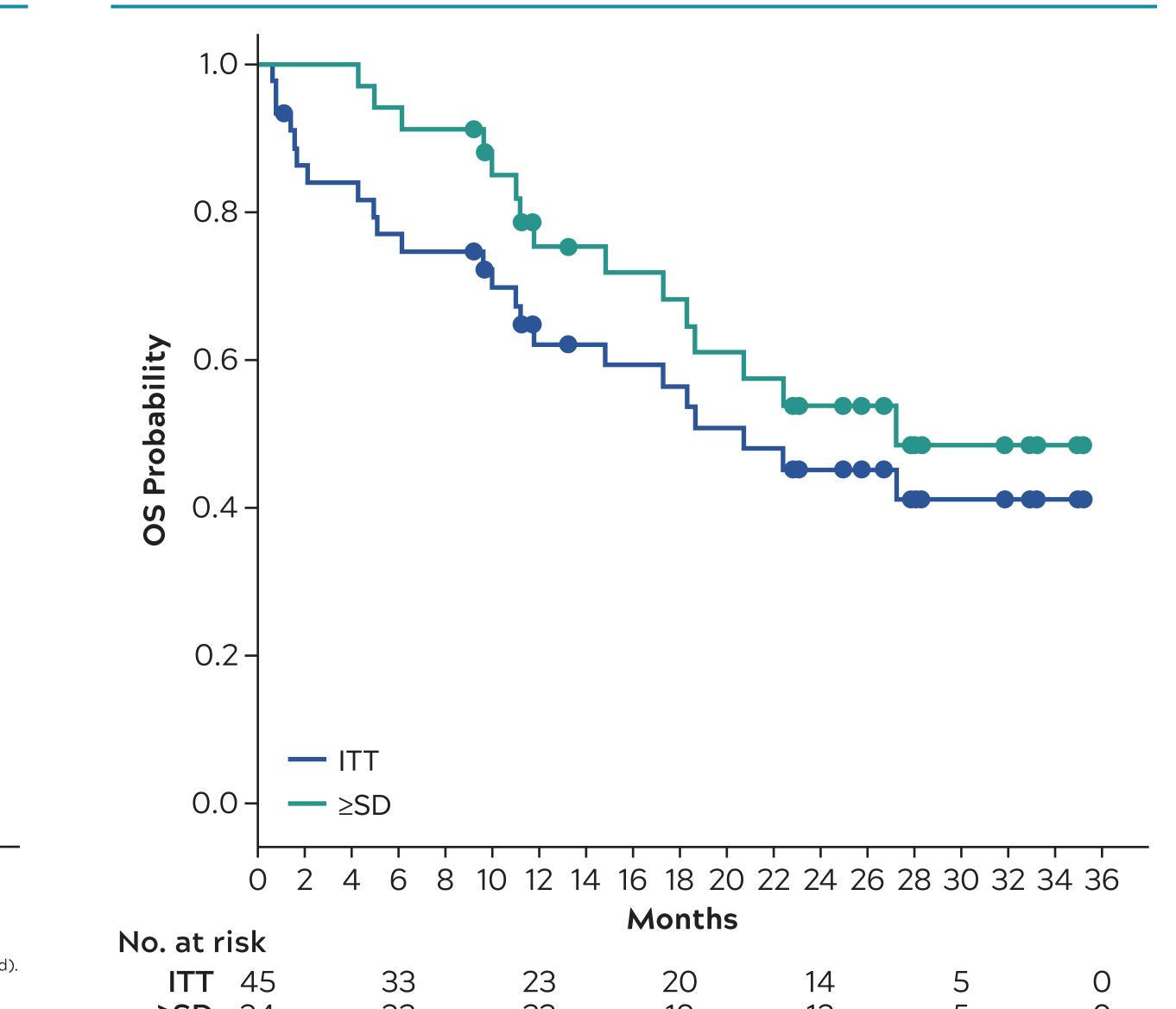
8.0-12.3; **Figure 3**)

Median TTNT was 7.9 months (95% CI, 5.68-11.01), with 40 events in

 Median PFS was 5.7 months (95% CI, 3.7-9.3), with 41 events in 45 patients (**Figure 3**)

TT, intention-to-treat; PD, progressive disease; PFS, progression-free survival; TTNT, time to next treatment.

Figure 4. OS in the ITT Population (N=45) igure 3. TTNT and PFS in the ITT Population (N=45)



An event was defined as death for the OS analysis ITT, intention-to-treat; OS, overall survival; SD, stable disease.

 Median OS in the total population was 20.7 months (95% CI, 11.8-NC), with 23 events in 45 patients, and 27.2 months (95%CI, 18.3-NC) in patients with SD or better (Figure 4)

Table 3. TTNT With Melflufen in O-12-M1 and Other Agents in RRMM

Agent or Regimen	No. of Patients	Prior Lines of Therapy	Death as an Event or Censored	Median TTNT, mo
Melflufen	45	4	Censored	10.6
Pomalidomide or carfilzomib ¹⁷	Pomalidomide: 264 Carfilzomib: 190	2+	Censored	Pomalidomide: 11.9 Carfilzomib: 9.4
Melflufen	45	4	Event	7.9
VRd or KRd ¹⁸	VRd: 343 KRd: 139	1-3	Event	VRd: 12.9 KRd: 8.7
Daratumumab ¹⁹	126	4	Event	5.9

- KRd, carfilzomib + lenalidomide + dexamethasone; RRMM, relapsed/refractory multiple myeloma; TTNT, time to next treatment; VRd, bortezomib + lenalidomide + dexamethasone.
- The data presented suggest a TTNT with melflufen plus dexamethasone that is as good as or better than that of other RWE studies of agents in RRMM (**Table 3**)

SUBSEQUENT THERAPY

medical need in this patient population

0.6

— PFS

45 patients (**Figure 3**)

TTNT or death

An event was defined as PD or death, whichever occurred first for the PFS analysis.

 In total, 66% of the patients went on to receive subsequent therapy (27/41 patients, with 4 patients still progression free at the time of the data cutoff)

Variability in subsequent therapy was high, indicating a high unmet

 The majority of patients (52%) received single agent with or without steroid therapy, and approximately half of the patients receiving subsequent therapy (44%) received ≥2 subsequent lines of therapy

CONCLUSIONS AND FUTURE DIRECTIONS

- Melflufen plus dexamethasone treatment results in disease stabilization (≥SD) in 76% of patients, which translates to a median TTNT of 7.9 months (10.6 months when censoring at time of death) in heavily pretreated patients with RRMM, which compares favorably with other relevant trials
- The median OS of 20.7 months in this advanced RRMM population suggests that melflufen therapy is associated with a long-term benefit, allowing patients to receive further treatment to control disease
- Longer TTNT for patients with RRMM is associated with clinical benefit as well as health economic value for payors. The reported ranges of median PFS/TTNT values in real-world are generally shorter than those in phase 3 clinical studies.²⁰ Real world data will be gathered for melflufen in
- Results support those of previous reports showing the promising efficacy profile of melflufen for the treatment of RRMM
- Furthermore, data from O-12-M1 suggest that, in addition to the established clinical benefit from ≥PR and ≥MR responses, SD is also a clinically meaningful response in patients with RRMM treated with melflufen plus dexamethasone and that such patients should continue to stay on treatment. This will be further investigated in ongoing and future studies
- The high variability in subsequent therapies after melflufen plus dexamethasone indicates a lack of good treatment options and a significant unmet medical need in patients with advanced RRMM
- OP-103 OCEAN is an ongoing phase 3, randomized, global study that is further evaluating the efficacy and safety of melflufen plus dexamethasone versus pomalidomide plus dexamethasone in patients with RRMM refractory to lenalidomide (NCT03151811)

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DISCLOSURES

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28-Day cycle

Melflufen + dexamethasone

21-Day cycle

(n=28)

Primary endpoint: ORR

Secondary endpoints: PFS, OS, safety

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