MYELODYSPLASTIC SYNDROME

Randomized phase 2 trial of pevonedistat plus azacitidine versus azacitidine for higher-risk MDS/CMML or low-blast AML

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To the Editor

There is a critical unmet need for novel treatments for higher-risk myelodysplastic syndromes (MDS), higher-risk chronic myelomonocytic leukemia (CMML), and low-blast (LB) acute myeloid leukemia (AML). For patients ineligible for stem cell transplant (SCT), standard therapy with hypomethylating agents, such as azacitidine and decitabine, is not curative, with most patients relapsing within 2 years [1–3].

Pevonedistat is the first small-molecule inhibitor of the neural precursor cell expressed, developmentally down-regulated 8 (NEDD8)-activating enzyme (NAE); NAE facilitates conjugation of the small ubiquitin-like protein, NEDD8, which activates cullin-RING E3 ubiquitin ligases (CRLs) [4–6]. Inhibition of NAE by pevonedistat prevents degradation of CRL substrates integral to tumor cell growth, proliferation, and survival, thereby leading to cancer cell death [4–6]. Pevonedistat + azacitidine demonstrated preclinical synergistic antitumor activity in AML xenografts and was well tolerated in patients with untreated AML, with promising clinical activity [7]. Based on these results, this phase 2, multicenter, global, randomized, controlled, open-label trial (NCT02610777) compared pevonedistat + azacitidine versus single-agent azacitidine in patients with higher-risk MDS/CMML and LB-AML who had not previously received a hypomethylating agent.

The study enrolled adults with morphologically confirmed higher-risk MDS, non-proliferative CMML, or LB-AML (20–30% myeloblasts in bone marrow); these patients were eligible for enrollment because the diseases are part of the higher-risk MDS spectrum, and were included in the pivotal randomized study that demonstrated significant improvement in overall survival (OS) with azacitidine versus conventional care regimens [3, 8, 9]. Patients with MDS/CMML were required to have very-high, high, or intermediate risk according to the revised international prognostic scoring system (IPSS-R);
patients with intermediate-risk IPSS-R (>3–4.5 points) had ≥5% bone marrow myeloblasts (see Supplementary Appendix for detailed eligibility criteria).

Patients were randomized 1:1 to receive either pevonodistat 20 mg/m² (intravenous) on days 1/3/5, plus azacitidine 75 mg/m² (intravenous or subcutaneous) on days 1–5/8/9, or azacitidine alone on the same schedule, in 28-day cycles, and stratified into four categories: LB-AML, and MDS/CMML with IPSS-R risk of very-high/high/intermediate. Treatment continued until unacceptable toxicity, relapse, transformation to AML (defined according to World Health Organization classification as ≥20% blasts in blood or marrow and 50% increase in blast count from baseline [8]), progressive disease (PD), or the initiation of subsequent anticancer therapy or hematopoietic SCT. Patients with PD could continue treatment if deriving clinical benefit if their disease had not transformed to AML.

The study was initially powered for a primary endpoint of event-free survival (EFS; defined as the time from randomization to death or transformation to AML in higher-risk MDS/CMML, or death in LB-AML). In consultation with regulatory agencies following completion of enrollment, the primary endpoint was changed to OS, with EFS as a secondary endpoint. Other secondary and exploratory endpoints are listed in the Supplementary Appendix. Response assessment was based on modified international working group (IWG) criteria for MDS for patients with higher-risk MDS/CMML [10] and revised recommendations of the IWG for AML for patients with LB-AML [11]. Disease assessments were based on local bone marrow aspirate blast counts and transfusions, and central laboratory data. Toxicity was evaluated according to National Cancer Institute Common Terminology Criteria for Adverse Events, version 4.03. Further details of assessments and statistical analysis are provided in the Supplementary Appendix.

Overall, 120 patients from 45 sites in 12 countries were enrolled (pevonodistat + azacitidine: 58 patients; azacitidine: 62 patients) (Supplementary Fig. 1). Baseline demographics and disease characteristics were generally well-balanced between arms (Supplementary Table 1).

At data cutoff for the final analysis of this randomized proof-of-concept study, median follow-up was 21.4 and 19.0 months in the pevonodistat + azacitidine and azacitidine arms, respectively. In the intent-to-treat (ITT) population, pevonodistat + azacitidine demonstrated clinically meaningful increases in OS (median 21.8 months versus 19.1 months; HR: 0.539; P = 0.045; Fig. 1d). Patients with higher-risk MDS were more likely to achieve a response with pevonodistat + azacitidine versus azacitidine (ORR 79.3% versus 56.7%); the CR rate was nearly doubled (51.7% versus 26.7%) and duration of response was also improved (median 34.6 versus 13.1 months) (Fig. 1e).

Data on EFS and OS in prespecified subgroups, time to treatment failure (TTF), transformation to AML, transfusion independence, and subsequent SCT are available in Supplementary Appendix/Supplementary Figs. 2–6. In patients with higher-risk MDS, TTF was longer (median 19.7 versus 13.6 months; HR: 0.521; P = 0.025) and the rate of transfusion independence in patients with higher-risk MDS who were transfusion-dependent at baseline was higher with pevonodistat + azacitidine versus azacitidine alone (69.2% versus 47.4%; P = 0.228).

In LB-AML, median OS (equivalent to EFS) trended longer with pevonodistat + azacitidine versus azacitidine (23.6 months versus 16.0 months, P = 0.081; Fig. 1f) although there was no ORR increase (52.9% versus 60.0%; CR/CRi 41.2% versus 60.0%). However, the LB-AML population was small (n = 32), and differences in baseline rate of AML with myelodysplasia-related changes (71% versus 42% with pevonodistat + azacitidine versus azacitidine) and differing proportions of patients with adverse risk according to European LeukemiaNet 2017 guidelines (59% versus 26%) may have affected response rates.

In higher-risk CMML, median OS was 21.7 months versus not evaluable (NE) (Fig. 1g), and median EFS was 21.0 months versus NE (Fig. 1h) with pevonodistat + azacitidine versus azacitidine, respectively. ORR was 77.8% with pevonodistat + azacitidine versus 75.0% with azacitidine (Supplementary Table 2). Although there was no observed benefit, the small number of patients with higher-risk CMML (17 total) precludes meaningful conclusions.

At data cutoff, patients in the pevonodistat + azacitidine arm had received a median of 13.0 cycles (range: 1–37) of pevonodistat and 13.0 cycles (range: 1–39) of azacitidine; patients in the azacitidine arm received a median of 8.5 cycles (range: 1–41) of azacitidine. The higher number of treatment cycles with pevonodistat + azacitidine compared
with azacitidine alone was consistent with the observed longer duration of response in the combination arm. To determine if the slightly higher number of patients with very-high-risk (VHR)-MDS in the azacitidine arm (n = 16/35) versus the combination arm (n = 10/32) may have contributed to the potential benefit observed with pevonedistat + azacitidine, sensitivity analyzes for OS and EFS (statistical analysis details provided in the Supplementary Appendix) demonstrated that the treatment effect was maintained after stratification adjustment for IPSS-R risk category. Median pevonedistat dose intensity was 98.7%; median azacitidine dose intensity was similar between treatment arms (pevonedistat + azacitidine: 96.9%, azacitidine: 98.2%).

Overall, the safety profile of pevonedistat + azacitidine was comparable to that of azacitidine alone (Table 1).
Table 1 Overall safety profile, and most common any-grade and grade ≥ 3 TEAEs occurring in ≥10% of patients.

| AE, n (%)                              | Pevonedistat + azacitidine n = 58 | Azacitidine alone n = 62 | Total N = 120 |
|----------------------------------------|-----------------------------------|-------------------------|---------------|
| Any AE, n (%)                          | 57 (98)                           | 62 (100)                | 119 (99)      |
| Any drug-related AE, n (%)             | 44 (76)                           | 50 (83)                 | 94 (78)       |
| Any grade ≥ 3 AE, n (%)                | 52 (90)                           | 54 (87)                 | 106 (88)      |
| Any drug-related grade ≥ 3 AE, n (%)   | 26 (45)                           | 29 (47)                 | 55 (46)       |
| Any serious AE, n (%)                  | 40 (69)                           | 39 (63)                 | 79 (66)       |
| Any drug-related serious AE, n (%)     | 9 (16)                            | 10 (16)                 | 19 (16)       |
| AE leading to discontinuation, n (%)   | 10 (17)                           | 13 (21)                 | 23 (19)       |
| On-study deaths, n (%)                 | 5 (9)                             | 10 (16)                 | 15 (13)       |
| Most common any-grade AE (≥10% of patients), n (%) |                                    |                         |               |
| Constipation                           | 21 (36)                           | 29 (47)                 | 50 (42)       |
| Nausea                                 | 20 (34)                           | 28 (45)                 | 48 (40)       |
| Pyrexia                                | 22 (38)                           | 25 (40)                 | 47 (39)       |
| Anemia                                 | 18 (31)                           | 28 (45)                 | 46 (38)       |
| Cough                                  | 22 (38)                           | 21 (34)                 | 43 (36)       |
| Neutropenia                            | 20 (34)                           | 18 (29)                 | 38 (32)       |
| Fatigue                                | 12 (21)                           | 25 (40)                 | 37 (31)       |
| Diarrhea                               | 19 (33)                           | 17 (27)                 | 36 (30)       |
| Febrile neutropenia                    | 15 (26)                           | 18 (29)                 | 33 (28)       |
| Anemia                                 | 17 (29)                           | 12 (19)                 | 29 (24)       |
| Dyspnea                                | 13 (22)                           | 15 (24)                 | 28 (23)       |
| Thrombocytopenia                       | 14 (24)                           | 14 (23)                 | 28 (23)       |
| Vomiting                               | 14 (24)                           | 13 (21)                 | 27 (23)       |
| Decreased appetite                     | 11 (19)                           | 12 (19)                 | 23 (19)       |
| Edema peripheral                       | 12 (21)                           | 8 (13)                  | 20 (17)       |
| Epistaxis                              | 13 (22)                           | 6 (10)                  | 19 (16)       |
| Pneumonia                              | 9 (16)                            | 10 (16)                 | 19 (16)       |
| Back pain                              | 10 (17)                           | 8 (13)                  | 18 (15)       |
| Neutrophil count decreased             | 12 (21)                           | 6 (10)                  | 18 (15)       |
| Arthralgia                             | 5 (9)                             | 12 (19)                 | 17 (14)       |
| Dizziness                              | 8 (14)                            | 8 (13)                  | 16 (13)       |
| Hypokalemia                            | 4 (7)                             | 11 (18)                 | 15 (13)       |
| Abdominal pain                         | 4 (7)                             | 10 (16)                 | 14 (12)       |
| Fall                                   | 7 (12)                            | 7 (11)                  | 14 (12)       |
| Pain in extremity                      | 10 (17)                           | 4 (6)                   | 14 (12)       |
| Platelet count decreased               | 7 (12)                            | 7 (11)                  | 14 (12)       |
| Insomnia                               | 6 (10)                            | 7 (11)                  | 13 (11)       |
| Headache                               | 3 (5)                             | 9 (15)                  | 12 (10)       |
| Most common grade ≥ 3 AE (≥10% of patients), n (%) | 19 (33)                           | 17 (27)                 | 36 (30)       |
| Neutropenia                            | 15 (26)                           | 18 (29)                 | 33 (28)       |
| Febrile neutropenia                    | 11 (19)                           | 17 (27)                 | 28 (23)       |
| Anemia                                 | 11 (19)                           | 14 (23)                 | 25 (21)       |
| Neutrophil count decreased             | 12 (21)                           | 6 (10)                  | 18 (15)       |
| Pneumonia                              | 7 (12)                            | 6 (10)                  | 13 (11)       |

AE adverse event, TEAE treatment-emergent adverse event.

Grade ≥ 3 TEAEs were reported in 90% of patients with pevonedistat + azacitidine versus 87% with azacitidine. The most frequent grade ≥ 3 TEAEs were neutropenia (33% versus 27%), febrile neutropenia (26% versus 29%), anemia (19% versus 27%), and thrombocytopenia (19% versus 23%). The addition of pevonedistat to azacitidine did not result in additional myelosuppression, which is important for patients with disease- and age-related comorbidities and azacitidine dosing was not compromised. Consequently, patients could remain on treatment for longer with pevonedistat + azacitidine versus azacitidine alone. This contrasts with prior studies, in which the addition of a second agent to azacitidine led to increased toxicity, resulting in azacitidine dose reductions or shorter dosing schedules [12, 13]. On-study deaths occurred in 9% of pevonedistat + azacitidine-treated patients versus 16% with azacitidine. The 60-day mortality rate was 3.4% versus 12.9%; causes of death within 60 days included acute cardiac failure and multi-organ failure (both n = 1) with pevonedistat + azacitidine, and gastric necrosis, hypoxia, multiorgan failure, pneumonia, the progression of MDS, sepsis, subdural hematoma, and unknown factors (all n = 1) with azacitidine alone.

Treatment with pevonedistat + azacitidine or azacitidine alone was associated with similar patient-reported symptoms, functioning, and health-related quality of life (HRQoL) (Supplementary Appendix/Supplementary Fig. 7). Baseline mutational profiling data suggest that the numerically higher ORR observed with pevonedistat + azacitidine occurred across prognostic subgroups, including in patients harboring poor prognostic mutations (Supplementary Appendix/Supplementary Figs. 8–10).

In summary, this randomized, proof-of-concept phase 2 study demonstrated clinical efficacy with pevonedistat + azacitidine in patients with higher-risk MDS and LB-AML. The OS, EFS, and ORR benefits were particularly promising among patients with higher-risk MDS, as was the OS benefit in LB-AML. The addition of pevonedistat to azacitidine resulted in a comparable safety profile to azacitidine alone, no increased myelosuppression, and azacitidine dose intensity was maintained. The combination of azacitidine and pevonedistat appears less myelosuppressive than azacitidine and venetoclax and more applicable to outpatient treatment [14]. Given the encouraging clinical activity in combination with azacitidine, its novel mechanism of action, and its nonmyelosuppressive safety profile, pevonedistat may be an ideal combination partner with other agents, such as venetoclax, as the treatment landscape evolves.

Data availability

The datasets, including the redacted study protocol, redacted statistical analysis plan, and individual participants data supporting the results reported in this article, will be made available within 3 months from initial request to researchers.
who provide a methodologically sound proposal. The data will be provided after its de-identification, in compliance with applicable privacy laws, data protection, and requirements for consent and anonymization.

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Author contributions MAS, LA, RJF, and DVF wrote the first draft of the manuscript. All authors conceived and/or designed the work that led to the submission, acquired data, and/or played an important role in interpreting the results, revised the manuscript, approved the final version, and agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Compliance with ethical standards

Conflict of interest MAS reports a membership or advisory role for Takeda/Millennium and Bristol-Myers Squibb. JW reports consulting for Jazz Pharmaceuticals, Takeda, Genentech, Rafael Pharma, Pfizer, and Celgene. AR reports consultancy within the past two years for Janssen, Pfizer, Novartis, Servier, research funding from Amgen, Bayer, Takeda, and honoraria directly received from an entity from Janssen, Pfizer, Novartis, and Servier. MAS reports consulting and research funding from Novartis and Celgene and research funding from Janssen. PFL reports expert testimony for Celgene and Novartis, and an advisory board role for Celgene, Pfizer, and Menarini. JFZ reports consultancy, researching funding, and honoraria from Celgene and Takeda, consultancy from AsystBio Laboratories, and research funding from Arog Pharmaceuticals Inc., Forty Seven, Inc., Merck, and Tolero Pharmaceuticals, Inc. JFZ also reports honoraria from Abbvie Inc., Agios, Daiichi Sankyo, Genentech, and Pfizer. MDC reports a membership or advisory role and consulting for Celgene, Novartis, and Takeda. CG reports an advisory board role for Celgene, Pfizer, Novartis, Janssen, Incyte, and Amgen. DS reports consultancy and honoraria from Takeda, Celgene, Novartis, Janssen Cilag, and AbbVie, expert testimony from Celgene and Janssen Cilag, and membership or advisory role for Belgian College for reimbursement of orphan drugs. RJF reports employment for Millennium Pharmaceuticals, Inc., a wholly-owned subsidiary of Takeda Pharmaceutical Company Limited, consultancy for BeyondSpring Pharmaceuticals, and an ownership interest in Takeda, Bristol Myers Squibb, Teva, Baxter, Gilead Sciences, and Pfizer. DZ, SF, and DVF report employment for Millennium Pharmaceuticals, Inc., a wholly-owned subsidiary of Takeda Pharmaceutical Company Limited. IB reports employment and an ownership interest in Millennium Pharmaceuticals, Inc., a wholly-owned subsidiary of Takeda Pharmaceutical Company Limited. LA reports honoraria directly received from an entity from Celgene, Takeda, Abbvie, and Jazz Pharmaceuticals. The remaining authors declare that they have no conflict of interest.

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References

1. Kadia TM, Jabbour E, Kantarjian H. Failure of hypomethylating agent-based therapy in myelodysplastic syndromes. Semin Oncol. 2011;38(Oct):682–92.
2. Prebet T, Gore SD, Esterni B, Gardin C, Itzykson R, Thepot S, et al. Outcome of high-risk myelodysplastic syndrome after azacitidine treatment failure. J Clin Oncol. 2011;29(Aug):3322–7. 20
3. Fenaux P, Multi OJ, Hellstrom-Lindberg E, Santini V, Finelli C, Giagounidis A, et al. Efficacy of azacitidine compared with that of conventional care regimens in the treatment of higher-risk myelodysplastic syndromes: a randomised, open-label, phase III study. Lancet Oncol. 2009;10(Mar):223–32.
4. Brownell JE, Sintchak MD, Gavin JM, Liao H, Bruzese FJ, Bump N, et al. Substrate-assisted inhibition of ubiquitin-like protein-activating enzymes: the NEDD8 E1 inhibitor MLN4924 forms a NEDD8-AMP mimetic in situ. Mol Cell. 2010;37(Jan):102–11. 15
5. Soucy TA, Smith PG, Milhollen MA, Berger AJ, Gavin JM, Adhikari S, et al. An inhibitor of NEDD8-activating enzyme as a new approach to treat cancer. Nature. 2009;458(Apr):732–6. 9
6. Soucy TA, Smith PG, Rolfe M. Targeting NEDD8-activated cullin-RING ligases for the treatment of cancer. Clin Cancer Res. 2009;15(Jun):3912–6. 15
7. Swords RT, Coutre S, Mari S, Zeidner JF, Foran JM, Cruz J, et al. Pevonedistat, a NEDD8 E1 inhibitor, is an effective therapy for chronic myelomonocytic leukemia compared to azacitidine in patients with AML. Blood. 2018;131(Mar):1415–24. 29
8. Vardiman JW, Harris NL, Brunning RD. The World Health Organization (WHO) classification of the myeloid neoplasms. Blood. 2002;100(Oct):2292–302. 1
9. Zandberg DP, Huang TY, Ke X, Baer MR, Gore SD, Smith SW, et al. Treatment and outcomes for chronic myelomonocytic leukemia compared to myelodysplastic syndromes in older adults. Haematologica. 2013;98(Apr):584–90.
10. Chevenet BD, Greenberg PL, Bennett JM, Lowenberg B, Wijermans PW, Nimer SD, et al. Clinical application and proposal for modification of the International Working Group (IWG) response criteria in myelodysplasia. Blood. 2006;108(Jul):419–25. 15
11. Chevenet BD, Bennett JM, Kopecky KJ, Buchner T, Willman CL, Estey EH, et al. Revised recommendations of the International Working Group for diagnosis, standardization of response criteria, treatment outcomes, and reporting standards for therapeutic trials in acute myeloid leukemia. J Clin Oncol. 2003;21(Dec):4642–9. 15
12. Garcia-Manero G, Montalban-Bravo G, Berdeja JG, Abaza Y, Jabbour E, Eshell J, et al. Phase 2, randomized, double-blind study of pracinostat in combination with azacitidine in patients with
untreated, higher-risk myelodysplastic syndromes. Cancer. 2017;123(May):994–1002. 15

13. Sekeres MA, Othus M, List AF, Odenike O, Stone RM, Gore SD, et al. Randomized phase II study of azacitidine alone or in combination with lenalidomide or with vorinostat in higher-risk myelodysplastic syndromes and chronic myelomonocytic leukemia: North American intergroup study SWOG S1117. J Clin Oncol. 2017;35(Aug):2745–53. 20

14. DiNardo CD, Jonas BA, Pullarkat V, Thirman MJ, Garcia JS, Wei AH, et al. Azacitidine and venetoclax in previously untreated acute myeloid leukemia. N Engl J Med. 2020;383(Aug):617–29. 13