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Satisfaction and Experiences with Talquetamab: Results from Qualitative Patient and Physician Research

BACKGROUND

This study aimed to explore **the experiences of patients receiving talquetamab**, a bispecific antibody therapy approved for relapsed or refractory multiple myeloma (MM), which targets GPRC5D and CD3 receptors. Despite the therapy's demonstrated efficacy, there is limited understanding of patient experiences and physician perspectives regarding this first-in-class treatment.

METHODOLOGY

The methodology involved a qualitative study of U.S. adults diagnosed with multiple myeloma who took talquetamab for at least 4 months, and physicians who had experience with talquetamab. The overall study design was approved by both Johnson & Johnson's Methods Review Board and Sterling IRB.

PHASE 1



Patients were recruited through HealthTree Cure Hub Registry, a platform sponsored by HealthTree that enables patients to voluntarily share their health information.



They participated in 2-hour virtual interviews to discuss their experiences **1-2 months before and 4-5 months after** starting talquetamab.



Topics included **GPRC5D-related symptoms, overall treatment impact, and satisfaction.** The interviews occurred from February to April 2024.

PHASE 2



Patients and physicians met in a 2.5-hour facilitated roundtable in Chicago, IL, in May 2024, to review Phase 1 findings and generate ideas for symptom management.

RESULTS

PHASE 1

Total Participants: 12 participants

Gender: Male (67%), Female (33%)

Education: Some college (17%), College (42%),

Graduate degree (42%)

Age: 50-59 (8%), 60-69 (8%), 70-79 (84%)

Race: White (84%), Hispanic (8%),

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Black/African American (8%)

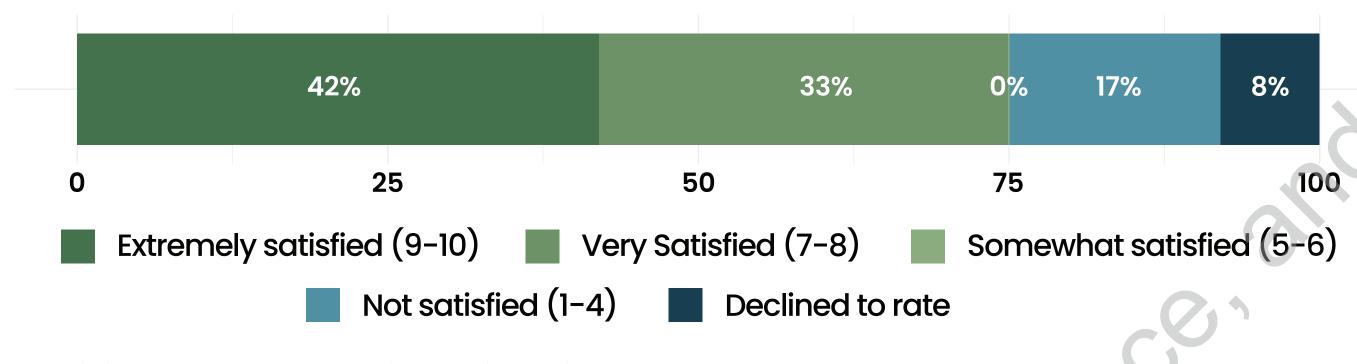
PHASE 2

Total Participants: 3 patients, 3 physicians Gender: Male (67%), Female (33%) Education: College (67%), Graduate degree (33%)

Age: 60-69 (33%), 70-79 (67%)
Race: White (100%)

OVERALL SATISFACTION WITH TALQUETAMAB

Patients reported **high overall satisfaction**, with most (75%) rating their satisfaction an 8 or higher on a 10-point scale:



Physicians also reported high satisfaction. Physicians said responses have been "very durable," and patients are doing "very well." One noted, "I'm very happy. As a physician, it's very nice to see the responses that we're seeing."

PATIENT QUOTES ON SATISFACTION

"This is the best response we've had to any of the therapies to date."

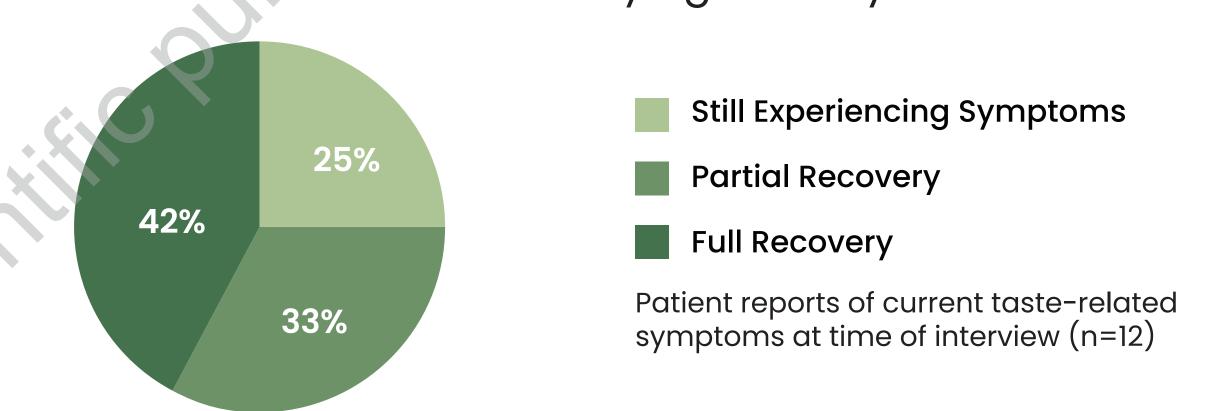
"If it wasn't for talquetamab, I wouldn't have gotten into MRD negativity when I went into CAR T-cell therapy. So despite all those toxicities, I was very grateful, because God knows what would've happened. I'd probably be dead by now."

"Within a few months ... the disease was undetectable, all my extramedullary disease, like my big lump, that all went away, all my broken bones were healing up."

"I couldn't be happier with the positive effects it's had on my disease burden. I'm down to zero. But getting to this point, it's been a bit of a tough go."

EXPERIENCE WITH GPRC5D-RELATED SYMPTOMS

Oral symptoms such as distorted taste/loss of taste and dry mouth were common with varying severity:



- Ten lost weight, but most (92%) mantained adequate nutrition.
- Skin- and nail-related symptoms were common but less impactful on quality of life (83% considered these symptoms mild).
- Overall, Experience with GPRC5D-related Symptoms were **not treatment-limiting**, and no patients had to miss a dose due to them. Patients perceived these symptoms as bearable and noted the treatment was effective at controlling the disease.

CONCLUSION

- This study described patient experiences and reflected how patients perceive the benefit/risk profile of talquetamab.
- Symptoms were perceived as bearable and high satisfaction scores were driven by strong clinical effectiveness.
- Patients and physicians agreed that real-world feedback from patients, practical tips, and management strategies are vitally important for patients and physicians for this first-in-class novel therapy.

IMPROVEMENTS TO QUALITY OF LIFE

Ratings largely stayed consistent from 1-2 months before starting to 4-5 months after starting, but **they noted improvements more often than declines in several main areas.** Physicians affirmed these findings as consistent with their observations.

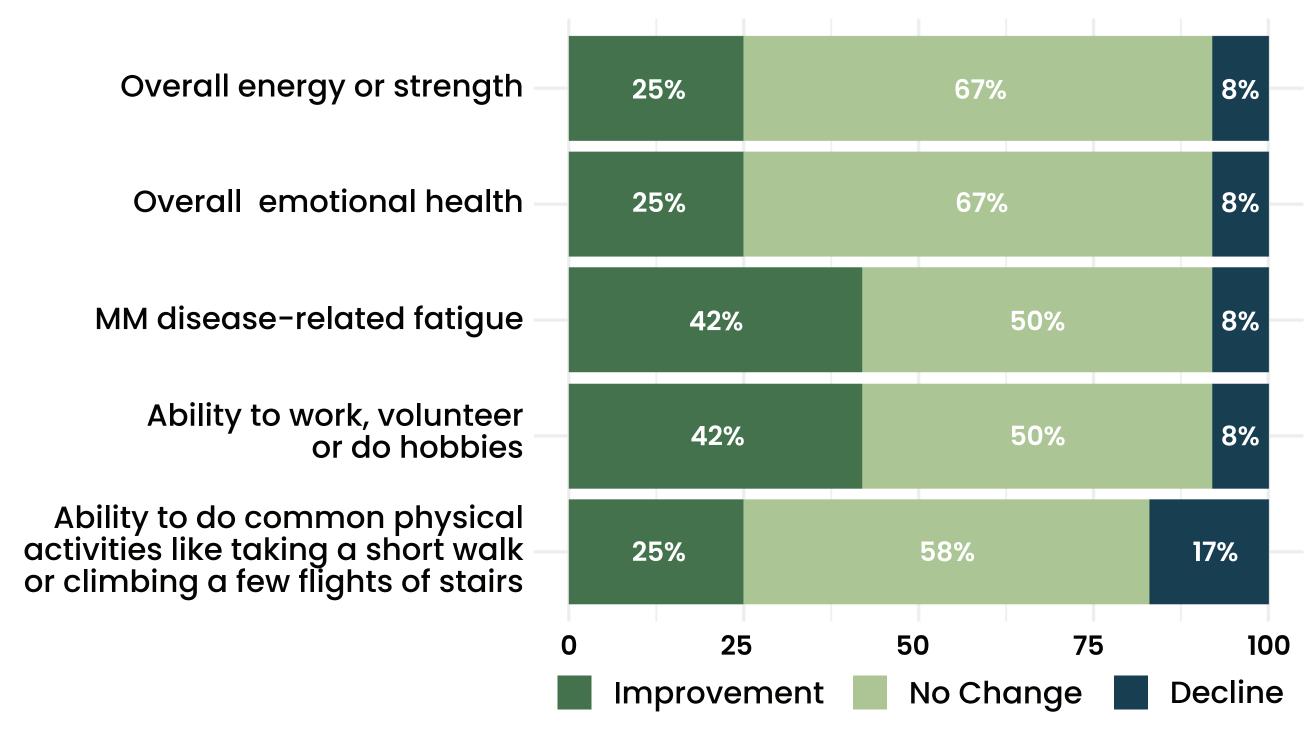
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Data reflects patient ratings which changed by 2 or more points on a 10-point scale from baseline (1-2 months prior to starting talquetamab) to post-test (4-5 months after starting). (n=12)

PHYSICIAN AND PATIENT SUGGESTIONS FOR SYMPTOM MANAGEMENT STRATEGIES

