Outcomes of Candida Bone and Joint Infections in Eight Patients from a Phase 3 Open-label Study (FURI)

JW Sanders¹, CG Morse¹, OA Cornely², P Koehler², R Krause³, J. Prattes³, P. Munoz⁴, G. Weiss⁵, NE Azie⁶, DA Angulo⁶ ¹Wake Forest University, ²University of Cologne, ³Medical University of Graz, ⁴Hospital General Universitario Gregorio Maranon, ⁵Medical University of Innsbruck, ⁶SCYNEXIS, Inc.

BACKGROUND

- Candida osteoarticular infections are often preceded by candidemia with the intervertebral discs and knee joints the most common area for candidemic seeding.
- Candida osteomyelitis has significant morbidity and diagnosis is often delayed & difficult to treat. Treatment courses are usually long and there are limited oral options available for patients who have an azole-resistant infection.
- Oral ibrexafungerp is an investigational broadspectrum glucan synthase inhibitor antifungal with activity against Candida and Aspergillus species, including azole- and echinocandin-resistant strains.
- A Phase 3 open-label, single-arm study of ibrexafungerp (FURI; NCT03059992) is ongoing for the treatment of patients with fungal disease refractory or who are intolerant of to standard of care antifungal therapy.
- Patients enrolled in the FURI study were from 22 centers in US, UK and EU who were treated with ibrexafungerp for severe mucocutaneous or invasive fungal infections from 2016-2020.

of gastrointestinal origin. No deaths due to progressive fungal disease FURI PATIENT FLOW were reported Treatment: Survival: Survival: (Days 3-5, 7-10, and EoT: (up to 90 days of Screening Baseline: Week 6 after EoT Day 84 after Day 42 after every 14 days treatment with (Day -1) Treatment Day 1 Visit through end of ibrexafungerp) Treatment Day 1 Treatment Day 1

METHODS

FURI subjects were eligible for enrollment if they had:

- Proven or probable severe mucocutaneous candidiasis,
- Invasive candidiasis or aspergillosis, other fungal diseases,
- Evidence of treatment failure, intolerance, or toxicity related to a currently approved standard-of-care antifungal treatment, or
- Unable to receive an approved oral antifungal option (e.g., susceptibility of the organism) and a continued IV antifungal therapy was clinically undesirable or unfeasible.
- An independent Data Review Committee (DRC) provided an assessment of treatment response for patients who completed therapy by October 2020.

SAFETY RESULTS

Ibrexafungerp was well-tolerated with the most common treatment-related adverse events being

EFFICACY RESULTS					
Site	AGE	SEX	Case type	Isolated organism	
Bone-spondylodiscitis	50	Male	Intolerant	Candida albicans	
Bone-spondylodiscitis	75	Female	Refractory	Candida tropicalis	
Bone-spondylodiscitis	58	Male	Refractory	Candida albicans	
Bone-spondylodiscitis	86	Male	IV step-down	Candida glabrata	
Bone-tibia	50	Male	IV step-down	Candida albicans and glabrata	
Bone-zygomatic arch	50	Male	Refractory	Candida glabrata	
Articular knee (prosthetic joint infection)	75	Female	IV step-down	Candida glabrata	
Bone-spondylodiscitis	86	Male	Intolerant	Candida albicans	

- zygomatic arch.
- All patients with bone or joint infections were white.
- ٠ vears of age.
- The median days of therapy for this group was 210.5 days.

Anatomic Location (n)	Comple Partial Respons
Bone/spondylodiscitis (5)	2 (40%)
Knee /prosthetic joint (1)	1 (100%
Tibia (1)	1 (100%
Zygomatic arch (1)	1 (100%
Totals (N=8)	5 (63%)



treatment (EoT)



There were 8 subjects (6 male, 2 female) who were diagnosed with various bone and joint infections, 5 with spinal infections, 1 with a knee/prosthetic joint infection and 2 subjects with osteomyelitis, one in the tibia and one in the

The mean age for this subpopulation at time of enrollment was 55.6 years; the median age at enrollment was 58

te/ **Progression of** Stable Indeterminate Disease Disease 1 (20%) 1 (20%) 1 (20%) 0 0 0 0 0 0 %) 0 0 0 1 (13%) 1 (13%) 1 (13%)



CONCLUSIONS

- Five of the 8 patients who were refractory or intolerant to standard of care showed a good therapeutic response.
- These data indicate that oral ibrexafungerp provides a promising therapeutic response option for patients with fungal infections of the bones or joints.