

Ferring and Rebiotix, a Ferring Company, have the largest and most robust clinical trial program ever conducted in microbiome-based therapeutics. The pivotal Phase 3 results for investigational RBX2660 are the first of its kind to demonstrate efficacy as early as first recurrence of *C. difficile* infection (CDI).

Details Of RBX2660 Clinical Trials:

World's most robust clinical program in microbiomebased therapeutics

Phase 2 trials, including a randomized double-blind placebo controlled Phase 2B trial

Phase 3 trials, including the pivotal PUNCH™ CD3 trial and PUNCH™ CD3 Open-Label study with expanded patient inclusion criteria (presented at DDW® 2021)

Retrospective assured active treatment (AAT) trial

The Decade-long Development Program Has:

> 1,000 patients enrolled



24 months of patient evaluation from two studies

 longest follow-up period of any microbiome-based therapeutic

The Program Is:

Designed to be representative of people physicians treat everyday

The only one:

- utilizing the same manufacturing process for product through the entire clinical program
- → showing:
 - repeated efficacy over placebo + standard of care
 - consistent safety across the clinical trial program

CDI Is A Major Health Threat

CDI is an urgent public health threat requiring immediate action.

people in the U.S. each year

Kills ~30K

500,000 cases a year in the U.S.

Approx.

35% of people within 8 weeks after first diagnosis

Antibiotics – which are the standard of care for treatment

of CDI – treat the disease but also are the primary risk

factor for a vicious cycle of recurrence, causing a

CDI recurs

in up to





Microbiome Therapeutics Development