

May 30, 2018



## ContraFect to Present New Data on CF-301 (exebacase) and Lysins Targeting Gram-Negative Pathogens at ASM Microbe 2018

YONKERS, N.Y., May 30, 2018 (GLOBE NEWSWIRE) -- [ContraFect Corporation](#) (Nasdaq:CFRX), a clinical-stage biotechnology company focused on the discovery and development of protein and antibody therapeutics for life-threatening, drug-resistant infectious diseases, today announced the presentation of new data on its lead drug candidate, CF-301 (exebacase), and its Gram-negative lysin discovery program at American Society for Microbiology (ASM) Microbe 2018, to be held from June 7-11, 2018, in Atlanta.

The presentations include results from new *in vivo*, *in vitro*, and surveillance studies of ContraFect's lead lysin CF-301 (exebacase), which is currently being studied in a Phase 2 clinical trial for the treatment of *Staphylococcus aureus* (*Staph aureus*) bacteremia, including endocarditis. Data will be presented which demonstrates CF-301's activity against a broad range of *Staphylococcus* and *Streptococcus* bacteria known to cause bacteremia and endocarditis, activity against contemporary clinical isolates of *Staph aureus* from U.S. hospitals, and ability to suppress emergence of vancomycin resistance. For the first time, data on CF-301's novel ability to synergize with and activate host factors in human serum will be presented. ContraFect will also present initial *in vitro* proof of principle data from the Gram-negative lysin discovery program demonstrating the bactericidal effect of its newly discovered lysins against multi-drug resistant (MDR) *Pseudomonas aeruginosa*.

"We are pleased to return to ASM this year and present important new data which expands our understanding of CF-301's spectrum of action of against a broad range of Staph and Strep species and provides new insight into the compound's novel ability to activate latent host factors in human blood which enhance bactericidal activity," said Cara Cassino, M.D., Chief Medical Officer and Executive Vice President of Research and Development at ContraFect. "We are also excited to unveil the first data on the bactericidal and anti-biofilm activity of Gram-negative lysins discovered in our laboratory against MDR *Pseudomonas aeruginosa*. We look forward to continuing to present new data from our active lysin research programs as we await top-line results from the ongoing Phase 2 trial of CF-301 later this year," continued Dr. Cassino.

### Presentation Details:

**Presentation Title:** Bacteriophage Lysins Can Be Engineered to Exert a Rapid and Potent

Bactericidal Effect against *Pseudomonas aeruginosa* in Serum

**Session Day & Time:** Sunday, June 10, 2018, 12:45 p.m. – 2:45 p.m. ET

**Poster Board Number:** SUNDAY - HMB LB14

**Session Title:** Session 384 - SUNDAY - HMB Late-breakers

**Presentation Title:** Lysin CF-301 Demonstrates Potent in Vitro Activity against a Range of *Staphylococcus* and *Streptococcus* Species Associated with Complicated Bacteremia and Infective Endocarditis in Humans

**Session Day & Time:** Sunday, June 10, 2018, 12:45 p.m. – 2:45 p.m. ET

**Poster Board Number:** SUNDAY - AAR LB5

**Session Title:** Session 403 - SUNDAY - AAR Late-breakers

**Presentation Title:** CF-301 Activity versus Contemporary *Staphylococcus aureus* Clinical Isolates from US Hospitals

**Session Day & Time:** Sunday, June 10, 2018, 12:45 p.m. – 2:45 p.m. ET

**Poster Board Number:** SUNDAY - 530

**Session Title:** Session 410 - AAR08 - New Antimicrobial Agents and New Research Technologies: Bacteriophage-Related Tools and Therapy

**Presentation Title:** Lysin CF-301 Administered in Addition to Vancomycin (VAN) Suppresses the Emergence of Reduced Susceptibilities to VAN Within Cardiac Vegetations in A Rabbit Model of MRSA Infective Endocarditis (IE)

**Session Day & Time:** Sunday, June 10, 2018, 12:45 p.m. – 2:45 p.m. ET

**Poster Board Number:** SUNDAY - 535

**Session Title:** Session 410 - AAR08 - New Antimicrobial Agents and New Research Technologies: Bacteriophage-Related Tools and Therapy

**Presentation Title:** Lysin CF-301 Activates Latent Host Factors in Human Blood to Potentiate Bacteriolysis

**Session Day & Time:** Sunday, June 10, 2018, 12:45 p.m. – 2:45 p.m. ET

**Poster Board Number:** SUNDAY - 536

**Session Title:** Session 410 - AAR08 - New Antimicrobial Agents and New Research Technologies: Bacteriophage-Related Tools and Therapy

The abstracts can be accessed through the [ASM Microbe website](#). Following the meeting, the presentation posters will be available on the [ContraFect website](#).

### About ContraFect:

ContraFect is a biotechnology company focused on discovering and developing therapeutic protein and antibody products for life-threatening, drug-resistant infectious diseases, particularly those treated in hospital settings. An estimated 700,000 deaths worldwide each year are attributed to antimicrobial-resistant infections. We intend to address life threatening infections using our therapeutic product candidates from our lysin and monoclonal antibody platforms to target conserved regions of either bacteria or viruses (regions that are not prone to mutation). ContraFect's initial product candidates include new agents to treat antibiotic-resistant infections such as MRSA (Methicillin-resistant *Staph aureus*) and influenza. ContraFect's lead product candidate, CF-301, is currently in a Phase 2 clinical trial for the treatment of *Staph aureus* bacteremia, including endocarditis and is the first lysin to enter clinical studies in the U.S. ContraFect is also conducting research focused on the discovery

of lysins to target Gram-negative bacteria.

### **About CF-301 (exebacase):**

CF-301 (exebacase) is a recombinant bacteriophage-derived lysin with potent bactericidal activity against *Staph aureus*, a major cause of blood stream infections, or bacteremia. CF-301 has the potential to be a first-in-class treatment for *Staph aureus* bacteremia. It has a novel, rapid, and specific mechanism of bactericidal action against *Staph aureus* and does not impact the body's natural bacterial flora. By targeting a conserved region of the cell wall that is vital to bacteria, resistance is less likely to develop to CF-301. Combinations of CF-301 with standard of care antibiotics significantly increased bacterial killing and survival in animal models of disease when compared to treatment with antibiotics or CF-301 alone. In addition, in vitro and in vivo experiments have shown that CF-301 is highly active against biofilm infections. CF-301 was licensed from The Rockefeller University and is being developed at ContraFect. It is the first lysin to enter clinical studies in the U.S.

### **Forward-Looking Statements:**

This press release contains, and our officers and representatives may make from time to time, "forward-looking statements" within the meaning of the U.S. federal securities laws. Forward-looking statements can be identified by words such as "projects," "may," "will," "could," "would," "should," "believes," "expects," "anticipates," "estimates," "intends," "plans," "potential," "promise" or similar references to future periods. Examples of forward-looking statements in this release include, without limitation, statements regarding new data and results on CF-301 and ContraFect's Gram-negative lysin discovery program, our ability to discover and develop protein and antibody therapeutics for life-threatening, drug-resistant infectious diseases, our ability to address life threatening infections using our therapeutic product candidates from our lysin and monoclonal antibody platforms to target conserved regions of either bacteria or viruses, whether our initial product candidates can treat antibiotic-resistant infections such as MRSA and influenza, our ability to discover new lysins targeting Gram-negative bacteria, the potential for CF-301 to be a treatment for *Staph aureus* bacteremia, including endocarditis, our ability to obtain top-line results from the ongoing Phase 2 trial later this year, *in vivo*, *in vitro* and surveillance studies, data on bactericidal and anti-biofilm activity of Gram-negative lysins against MDR *Pseudomonas aeruginosa* and statements made regarding presentations. Forward-looking statements are statements that are not historical facts, nor assurances of future performance. Instead, they are based on ContraFect's current beliefs, expectations and assumptions regarding the future of its business, future plans, strategies, projections, anticipated events and trends, the economy and other future conditions. Because forward-looking statements relate to the future, they are subject to inherent risks, uncertainties and changes in circumstances that are difficult to predict and many of which are beyond ContraFect's control, including those detailed in ContraFect's filings with the Securities and Exchange Commission. Actual results may differ from those set forth in the forward-looking statements. Important factors that could cause actual results to differ include, among others, our ability to develop treatments for drug-resistant infectious diseases. Any forward-looking statement made by ContraFect in this press release is based only on information currently available and speaks only as of the date on which it is made. Except as required by applicable law, ContraFect expressly disclaims any obligations to publicly update any forward-looking statements, whether written or oral, that may be made from time to time, whether as a result of new

information, future developments or otherwise.

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