Proprietary Renalase Platform

A first-in class Renalase agonist for Hyper-inflammation in:

Systemic Viral Infections
COVID-19
Acute Organ Injury

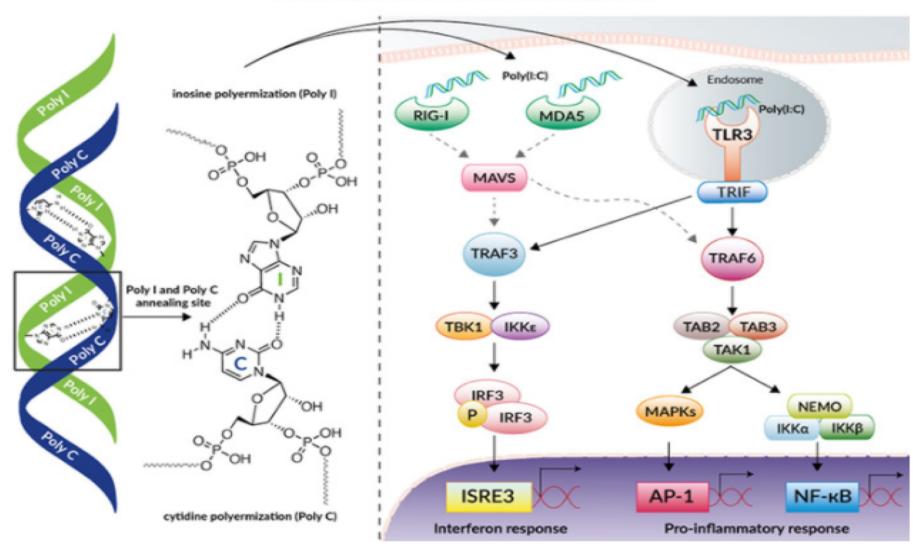
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Scientific Advisory Board member, Bessor Pharma
Co-founder and Chair of Scientific Advisory Board, Personal Therapeutics

The Renalase Agonist Platform

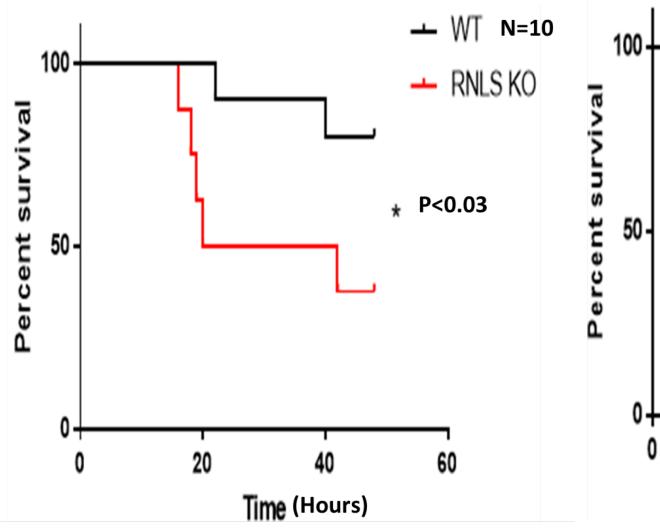
- Renalase (RNLS): a secreted protein that promotes cell survival and decreases inflammation through defined mechanisms
- Platform opportunities for broad indications
 - Hyper-inflammation in systemic viral infections, including COVID-19
 - Acute organ injury: Lung, Kidney, and Pancreas
- Proprietary RNLS mimetic peptide (BP-1002) with demonstrated preclinical proof of efficacy in acute diseases
 - IND targeted for Q1 2022
- Potential biomarker-linked therapeutic strategy with proprietary RNLS assay to optimize patient selection
- 3 NIH SBIR grants (>\$2.5 million)

Poly (I:C): A General Model of Hyper-inflammation

General Poly(I:C) structure and signaling pathway



Renalase agonist BP-1002 improves survival in Poly (I:C) viral mimic model

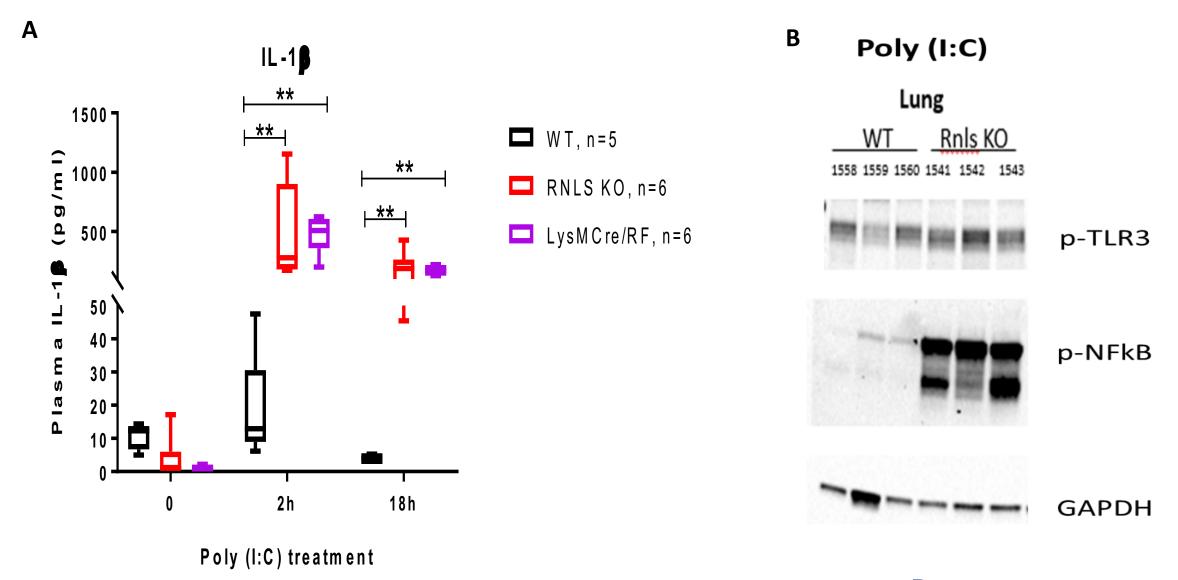


N=7 RNLS agonist: 100% P<0.0001 Control: 40% N=8 20 Time (hours)

Increased mortality in RNLS KO Exposed to Poly (I:C)

BP-1002 rescues RNLS KO phenotype

RNLS deficiency dramatically increases Poly (I:C) mediated inflammatory cytokines production through NF-κB, ISRE3 activation in mouse model



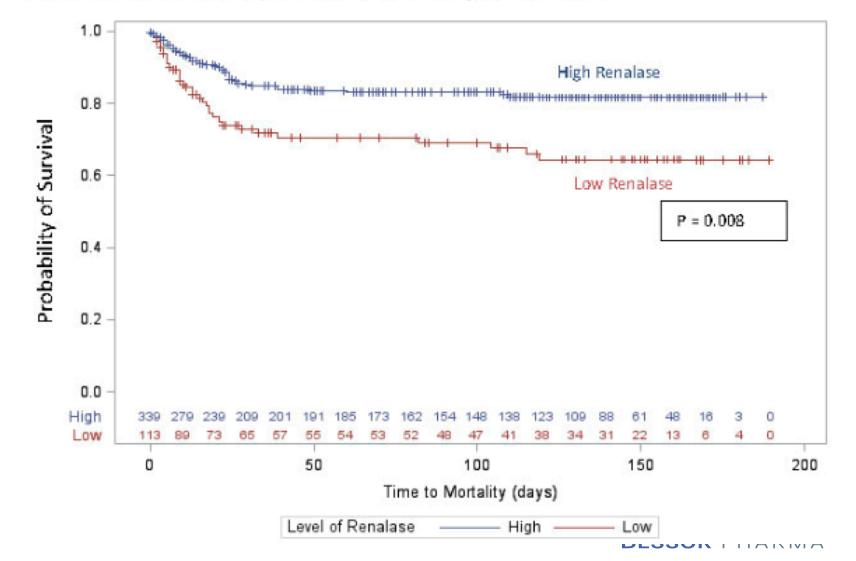
Renalase Levels Correspond with Mortality in COVID-19 patients

Low plasma levels are associated with worse outcomes in COVID-19. 2020. Wang, et al MedRxiv (Initial cohort of 51 COVID-19 Subjects)

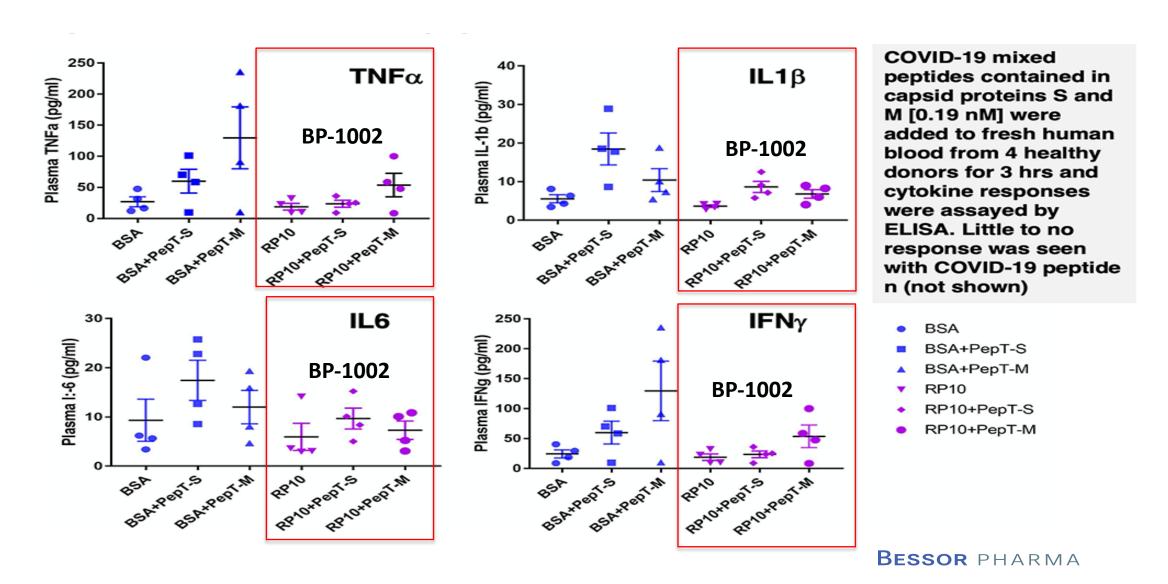
Updated cohort: N=458 COVID-19 Subjects

Paper under review

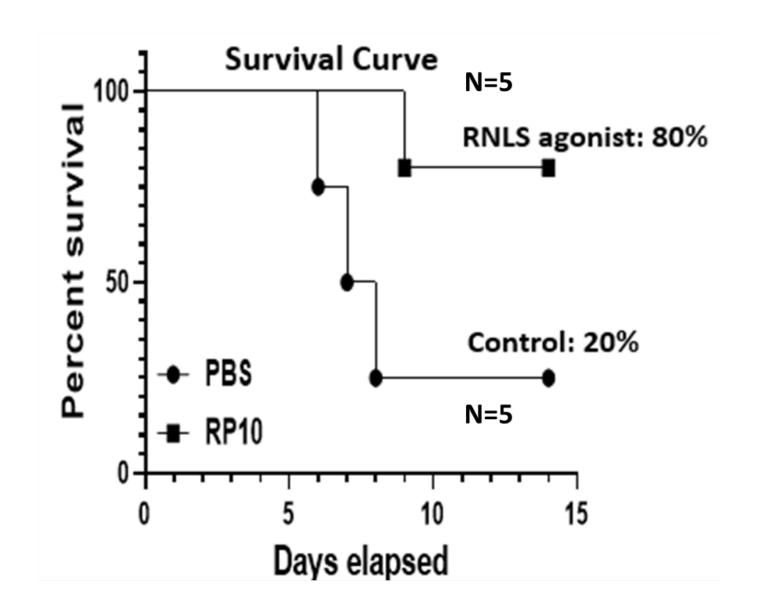
Renalase levels correspond with mortality in COVID-19



BP-1002 blunts multiple COVID-19 induced inflammatory cytokines in human blood ex vivo



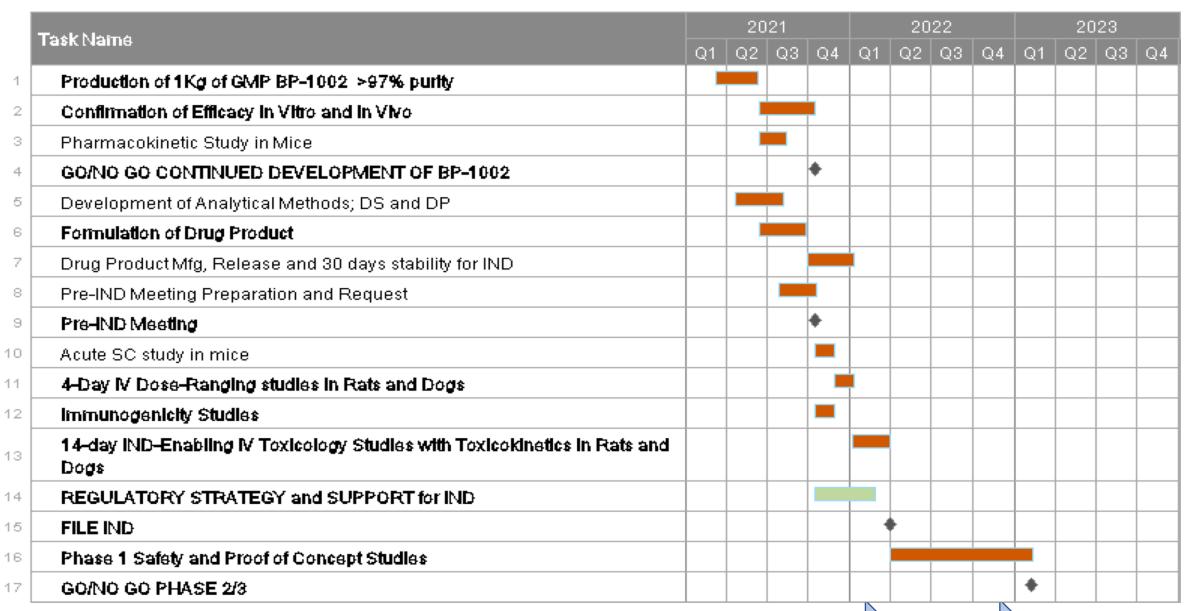
RNLS agonist BP-1002 (RP10) improves survival in COVID-19 mouse model



BP-1002 vs Standard Therapy in COVID-19

- Advantageous Side Effect Profile Over Dexamethasone
 - BP-1002 does not suppress adrenals and thymus; Dexamethasone does
 - Dexamethasone increases risk of opportunistic infections
- Potential therapeutic advantage over Tocilizumab,
 - BP-1002 targets multiple cytokines and is also a survival factor that prevents cell death
 - Tocilizumab only targets IL-6 Receptor
- Potential for predictive RNLS Assay to optimize patient selection and timing of administration of BP-1002

Development plan to clinical trials: IND in Q1 2022



\$5.5M \$10M

BP-1002 in Hyper-Inflammation

- Broad utility in systemic viral infections and acute organ injury
- Low Renalase associated with worse outcome in COVID-19 patients
 - 1st indication: COVID-19
- Chemically synthesized novel small RNLS mimetic peptide, designed for improved stability and efficacy
 - Treats severe disease and reduces mortality in animal models
 - Scalable for chemical manufacture
 - Composition of matter patent filed
 - Licensed and developed in collaboration with Bessor Pharma
 - IND in Q1 2022
- Platform potential for biomarker-linked therapeutics for acute diseases.

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