Revive Therapeutics

A Specialty Life Sciences Company

RVV LISTED ON CSE™

Corporate Presentation

May 2020

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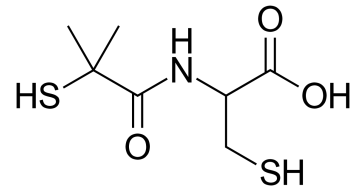
FORWARD LOOKING STATEMENTS

Certain statements contained in this presentation constitute forward-looking information within the meaning of securities laws. Forward-looking information may relate to our future outlook and anticipated events or results and may include statements regarding our future financial position, business strategy, budgets, litigation, projected costs, capital expenditures, financial results, taxes and plans and objectives. In some cases, forward-looking information can be identified by terms such as “may”, “will”, “should”, “expect”, “plan”, “anticipate”, “believe”, “intend”, “estimate”, “predict”, “potential”, “continue” or other similar expressions concerning matters that are not historical facts. These statements are based on certain factors and assumptions regarding, among other things, expected growth, results of operations, performance, and business prospects and opportunities. While we consider these assumptions to be reasonable based on information currently available to us, they may prove to be incorrect. Forward looking-information is also subject to certain factors, including risks and uncertainties that could cause actual results to differ materially from what we currently expect. These factors include, among other things, the availability of funds and resources to pursue development projects, the successful and timely completion of clinical studies, and the ability to take advantage of business opportunities, the granting of necessary approvals by regulatory authorities, and general economic, market and business conditions. For more exhaustive information on these risks and uncertainties you should refer to our most recently filed Annual Information Form which is available at www.sedar.com. Forward-looking information contained in this presentation is based on our current estimates, expectations and projections, which we believe are reasonable as of the current date. You should not place undue importance on forward-looking information and should not rely upon this information as of any other date. While we may elect to, we are under no obligation and do not undertake to update this information at any particular time.

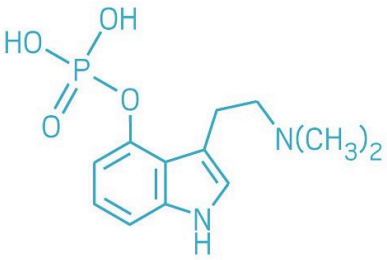
Revive Therapeutics

REVIVE THERAPEUTICS

- **Specialty life sciences company focused on repurposing drugs for rare disorders and infectious diseases**
- **Advancing clinical development of Bucillamine for infectious diseases, including COVID-19 (Phase 3)**
- **Developing novel Psilocybin and Cannabidiol therapeutics for various CNS and inflammatory disorders**
- **Robust patent portfolio covering methods and compositions of drugs**
- **Significantly undervalued compared to its peers**



Bucillamine



Psilocybin

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PHARMACEUTICAL STRATEGY

Clinical development of Psilocybin, Cannabidiol and Bucillamine

Targeting Rare Disorders and Infectious Diseases

Novel Uses, Formulations, and Delivery System

Targeting FDA Designations: Orphan, Fast track, Breakthrough therapy and Rare pediatric diseases

FDA Orphan Drug Designation

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PATENT PORTFOLIO

Title	USPTO No.	Status
Use of Bucillamine in the Treatment of Infectious Diseases, including COVID-19	62/991,996	Provisional patent filed
Use of Bucillamine in the Treatment of Gout	US9662305	Issued on May 30, 2017
Drug Delivery System	US 8642088 US 9545423 US 10104888	Issued on February 4, 2014 Issued on January 17, 2017 Issued on October 23, 2018
Psilocybin effervescent and psilocybin tablet - Solid Oral Pharmaceutical Compositions	62/985,052	Provisional patent filed
Psilocybin hard-shell capsules - Pharmaceutical Capsule Compositions	62/985,070	Provisional patent filed
Psilocybin gum drops - Pharmaceutical Gumdrop Compositions	62/985,084	Provisional patent filed
Psilocybin oral strips and transmucosal - Thin-Film Pharmaceutical Delivery System and Formulations	62/985,098	Provisional patent filed
Psilocybin - Pharmaceutical Formulations and Methods for Sublingual and Buccal Administration	62/984,590	Provisional patent filed
Methods for the Extraction and Crystallization of Psilocybin	62/985,360	Provisional patent filed
Use of Cannabidiol in the Treatment of Autoimmune Hepatitis	US 8242178	Issued on August 14, 2012

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PRODUCT PIPELINE

Focus on Infectious Diseases, including COVID-19, and Rare Disorders

Product	Indication	Stage of Development	Regulatory Status	Market
Bucillamine	Infectious Diseases (COVID-19)	Filing IND	IND for Phase 3	TBD
Psilocybin	Undisclosed	Pre-clinical	Target FDA Orphan Status	+ $500M
Psilocybin	Undisclosed Multiple Indications	Pre-clinical	Target FDA Orphan Status	TBD
CBD	Liver Diseases (Autoimmune Hepatitis)	Filing IND	Received FDA Orphan Status IND for Phase 2	+ $100M 50k US Pop

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INFECTIOUS DISEASE OPPORUNTIY

- **Focus on Bucillamine in the treatment of infectious diseases**
  - Well-known safety profile and prescribed for arthritis in Japan and South Korea for over 30 years
- **Revive's clinical history with Bucillamine**
  - Obtained 2 FDA INDs with Bucillamine and FDA orphan drug status
  - FDA Phase 2 clinical study for acute gout flares and cystinuria
- **Bucillamine (BUC) scientific rationale as an intervention for COVID-19** *(see Appendix)*
  - BUC is 16x more potent than particularly N-acetylcysteine (NAC); NAC has shown to prevent acute lung injury caused by influenza virus
  - BUC shown superior function in restoring glutathione and therefore greater potential to prevent acute lung injury during influenza infection
  - BUC also shown to prevent oxidative and reperfusion injury in heart and liver tissues
  - BUC proven safety and MOA similar to NAC, but with much higher potency
- **Advancing towards FDA Phase 3 clinical study for COVID-19**

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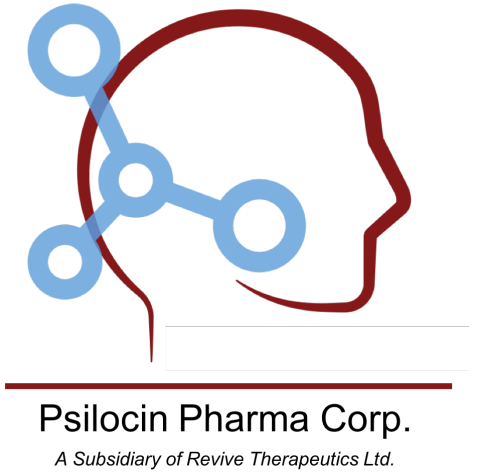
PSYCHEDELIC OPPORTUNITY

- **Acquired Psilocin Pharma Corp.**
  - Derrick Welch, Founder with 14 years of HC experience; 5 years in Cannabis
  - Worked with Xanthic Bio Pharma and Green Growth Brands
  - Developed water Soluble THC and CBD products (Beverages, effervescent tablets)



- **Novel Psilocybin formulations, extraction and purification methods**
  - Suited for pharmaceutical development and recreational markets where legal

- **Patent pending Psilocybin formulations (natural synthetic derived)**
  - Capsules, Sublingual Spray, Gel Cap, Effervescent Tablets, and Oral Strips

- **Targeting rare diseases, mental health and addiction**

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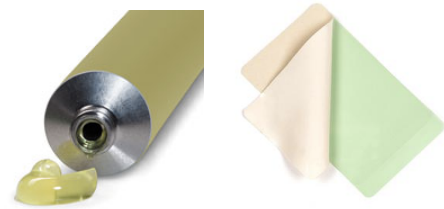
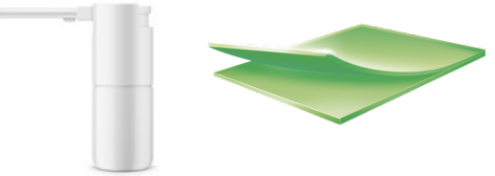
FORMULATION & DELIVERY TECHNOLOGY

- **Delivering naturally extracted and synthetic psilocybin and cannabinoids**



- **PSILOCYBIN - Precise dosed formulations i.e. capsules, sublingual spray, gel caps, effervescent tablets and oral/transmucosal strips**
  - Releases (rapid, controlled, sustained), improved bioavailability, no first-pass metabolism

- **CANNABIDIOL - Novel combination of composites allowing for multiple delivery formats, potential synergistic and therapeutic effects**
  - **Tannin** (plant-based) - antibacterial, antifungal, antioxidant and wound healing
  - **Chitosan** (shrimp-based) - blood-clotting and antimicrobial properties



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LIVER DISEASE OPPORTUNITY

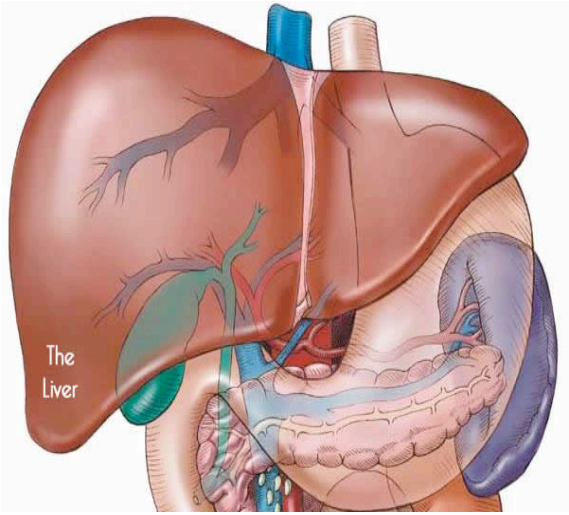
- **Focus on Autoimmune Hepatitis (AIH)**

- **AIH - rare disease (~ 76k patients in US) causing liver inflammation**
  - Drawbacks of current therapies (steroids): Severe side effects in 13%, relapse after drug withdrawal in 50%-86%*

- **Obtained FDA orphan drug status for CBD in the treatment of AIH**

- **Seeking to file FDA IND to conduct Phase 2 clinical study in patient affected by AIH**

- **Big Pharma interest in liver diseases**
  - Allergan acquisition of Tobira for $1.7 billion
  - Novartis license of Conatus drug for $650 million
  - Gilead acquisition of Nimbus for $1.2 billion



Source: World J Gastroenterol. 2010 Feb 28; 16(8): 934–947

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STRATEGIC PARTNERS

Pharm-Olam
Helping Create A Healthier World
Celebrating 25 Years of Excellence

Infectious Diseases CRO

Psilocin Corp.

Acquired March 2020 Psychedelic pharma

WISCONSIN
UNIVERSITY OF WISCONSIN-MADISON

R&D partnership Psychedelic pharma

RED LIGHT HOLLAND
Underground to Mainstream

Supply and collaboration of truffles

UNIVERSITY OF SOUTH CAROLINA

License of cannabidiol for treatment of Autoimmune Hepatitis

WARF
Wisconsin Alumni Research Foundation

License of cannabinoid delivery technology

Sanyal BIOTECHNOLOGY

Research collaboration for liver diseases

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VALUE DRIVING MILESTONES IN 2020

Q2	Submit FDA IND for Phase 3 clinical study of Bucillamine in the treatment of COVID-19
Q2	Initiate Phase 3 clinical study of Bucillamine in the treatment of COVID-19 in the U.S.
Q2	Submit IND for Phase 2 clinical study of CBD in the treatment of Autoimmune Hepatitis
Q2/3	Pre-IND meeting with FDA for Bucillamine in various infectious diseases (undisclosed)
Q3	Initiate Phase 2 clinical study of CBD in the treatment of Autoimmune Hepatitis
Q3	Pre-IND meeting with FDA for Psilocybin (undisclosed indications)
Q3/4	Results from Phase 3 clinical study of Bucillamine in the treatment of COVID-19
Q4	Results from Phase 2 clinical study of CBD in the treatment of Autoimmune Hepatitis
Q4	Submit FDA IND for Phase 2 clinical study of Bucillamine (undisclosed indication)

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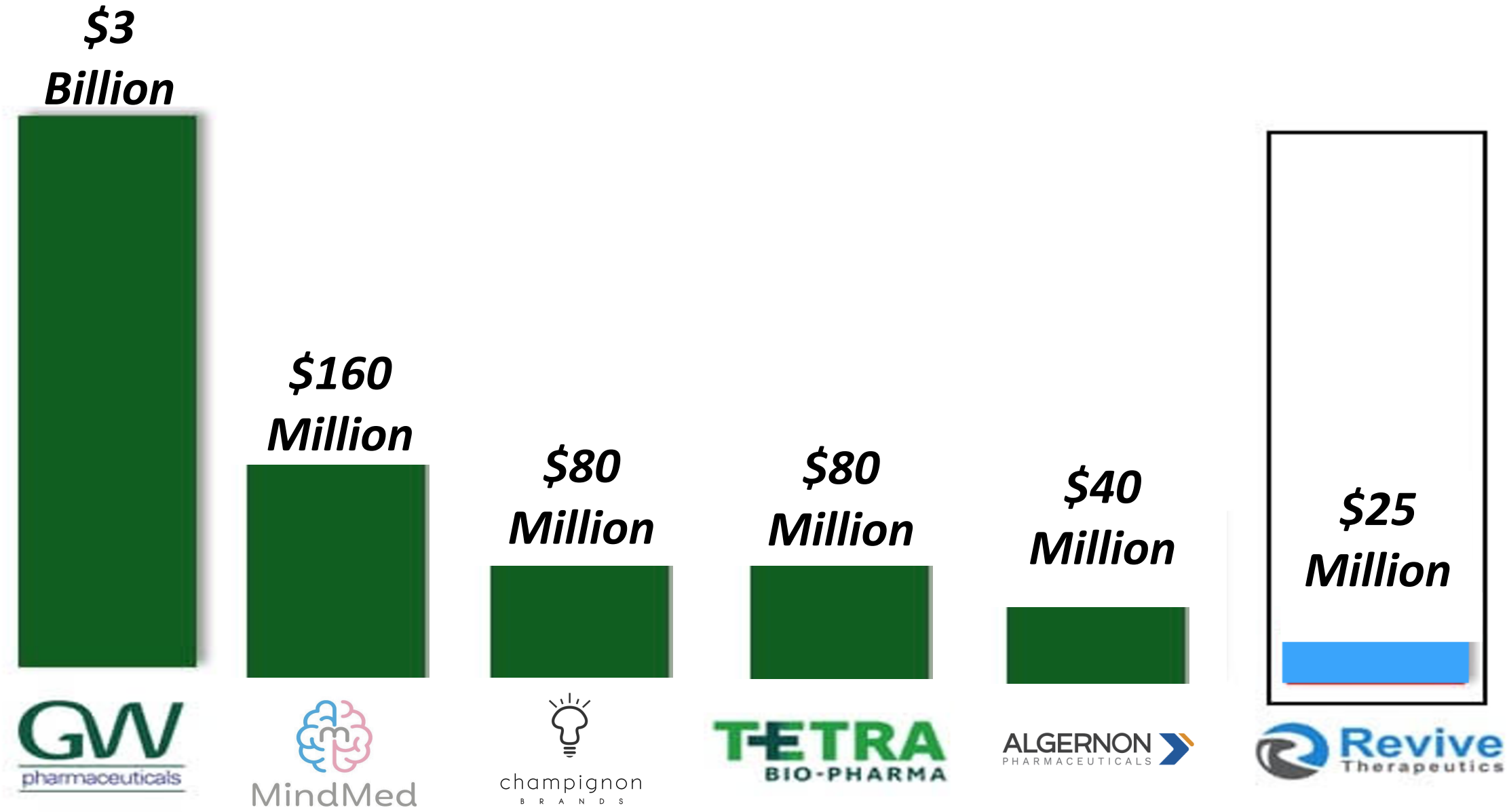
Peer Comparison *(Market Caps)*



ORGANIZATION

Management

- **Michael Frank**
  Chairman and CEO
- **Carmelo Marrelli**
  Chief Financial Officer
- **Derrick Welsh**
  Founder, Psilocin Pharma Corp.

Clinical

- **Dr. Kelly McKee, Jr., MD, MPH**
  Chief Scientific Officer, Consultant
- **Dr. David Boulware, MD, MPH, CTropMed, FIDSA**
  Clinical and Scientific Advisor
- **Dr. Onesmo Mpanju, PhD**
  FDA Regulatory Affairs, Consultant

Board of Directors

- **Michael Frank**
  Chairman and CEO
- **William Jackson**
  Director
- **Joshua Herman**
  Director
- **Christian Scovenna**
  Director
- **Andrew Lindzon**
  Director

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STOCK INFORMATION

Ticker	RVV (CSE)
Share Price	CAD $0.145 (May 7, 2020)
52 week High/Low	CAD $0.23 / $0.025
Capital Structure	189,408,777 common shares (267,075,486 fully-diluted)
Market Cap	CAD ~ $27,500,000

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Appendix – Bucillamine Scientific Rationale for COVID-19

Current antiviral interventions for influenza have exhibited modest efficacy, especially in improving mortality in at-risk populations, such as the elderly.[1,2] Novel antivirals have been plagued by poor oral bioavailability and lack of efficacy when not delivered early.1 This is because these drugs mostly act to prevent the early processes of virus binding to cells or viral replication.[2] Thiols, particularly N-acetylcysteine (NAC), with antioxidant and reducing activity have been investigated as effective therapies that abrogate the potential for influenza to cause severe disease.[3,4,5] Restoration of glutathione, the major intracellular thiol antioxidant, is a critical functional activity of NAC.[6] Reactive oxygen species (ROS) generation during influenza virus infection aggravate destructive inflammation and programmed death of epithelial cells.[7] Studies in human cells and animal models have shown that NAC works to prevent acute lung injury caused by influenza virus infection through inhibition of these ROS-mediated mechanisms.[4,7] NAC has been investigated clinically and found to significantly attenuate clinical symptoms associated with influenza infection, especially in elderly at-risk patients.[5] While NAC is easily taken up by cells and has low toxicity, clinical efficacy has required long-term and high-dose administration because of modest relative potency, limiting its clinical applicability.

Bucillamine (N-(mercapto-2-methylpropionyl)-l-cysteine), which has a well-known safety profile and is prescribed in the treatment of rheumatoid arthritis in Japan and South Korea for over 30 years, is a cysteine derivative with 2 thiol groups that is 16-fold more potent than NAC as a thiol donor in vivo, giving it vastly superior function in restoring glutathione and therefore greater potential to prevent acute lung injury during influenza infection.[8] Bucillamine has also been shown to prevent oxidative and reperfusion injury in heart and liver tissues[8] and is highly cell permeable for efficient delivery into cells.[8,9] Bucillamine has unrealized potential for the treatment of influenza with both proven safety and proven mechanism of action similar to that of NAC, but with much higher potency, mitigating the previous obstacles to using thiols therapeutically. It is also reasonable to hypothesize that similar processes related to ROS are involved in acute lung injury during nCov-19 infection, possibly justifying the investigation of Bucillamine as an intervention for COVID-19.

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Appendix – Bucillamine Scientific Rationale for COVID-19

References

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