Fauci, Anthony (NIH/NIAID) [E] From: Sent: Mon. 9 Mar 2020 09:57:48 +0000 To: Cassetti, Cristina (NIH/NIAID) [E]

(b) (6) Cc:

Subject: FW: Connecting with Tony Fauci

Attachments: Baricitinib as potential treatment for 2019-nCoV acute respiratory disease.pdf

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Please take a look and respond
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----Original Message----
From: Collins, Francis (NIH/OD) [E] <
Sent: Monday, March 9, 2020 5:29 AM
To: Fauci, Anthony (NIII/NIAID) [E]
                                                     (b) (6) >; Erbelding, Emily (NIH/NIAID) [E]
                    (b) (6) >; Austin, Christopher (NIH/NCATS) [E]
                                                                                   (b) (6)>
                                                        (b) (6)
Cc: Tabak, Lawrence (NIH/OD) [E]
Subject: FW: Connecting with Tony Fauci
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Passing this on in case Bill Chin's idea might be of interest.

Francis

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----Original Message----
From: William Chin
                                        (b) (6)>
Sent: Sunday, March 8, 2020 10:48 PM
To: Collins, Francis (NIH/OD) [E]
                                                   (b) (6)
Cc: Baker, Rebecca (NIH/OD) [E]
                                                     (b) (6)>; Austin, Christopher (NIH/NCATS) [E]
                 (b) (6) >; Roger Glass
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Subject: Connecting with Tony Fauci

Hi Francis et al, I write to get a message to Tony, who clearly is on the COVID-19 front lines of these days and "everywhere." At this early stage, you may agree that it is possible that containment and/or mitigation will not completely solve the current pandemic. Instead, effective treatment or amelioration of the worst complications is necessary. In this spirit, I have an idea that was spurred by a letter in Lancet last month authored by AI Benevolent (attached). In this communication they suggest that baricitinib (Oluminant; a JAK 1/2 inhibitor registered by Lilly for the treatment of rheumatoid arthritis), using in silico techniques, might be useful in the treatment advanced COVID-19 pneumonia/ARDS identified. Baricitinib could possibly blunt the cytokine storm seen in the most severely affected patients via inhibition of JAK1/2, but also decrease viral entry in AT2 pulmonary cells and myocardial cells via inhibition of GAK and AAKI. It is has a relatively short plasma half-life and hence could be more useful than Roche's tocilizumab/Actemra). Baricitinib has a good safety profile although as a drug to treat autoimmunity it is formally contraindicated in patients with infections, this could be offset by treatment patients with anti-virals such as Gilead's remdesivir, etc. Perhaps you folks have already thought about/discussed its use but if not I'd like a chance to chat about my additional thoughts. Thanks. Bill