CORRESPONDENCE

Neprilysin inhibitor–angiotensin II receptor blocker combination (sacubitril/valsartan): rationale for adoption in SARS-CoV-2 patients

On 11 March 2020, the World Health Organization (WHO) declared the coronavirus disease 2019 (COVID-19) outbreak as a 'pandemic'.[1] No valid therapy for COVID-19 is actually available. Severe acute respiratory syndrome-coronavirus-2 (SARS-CoV-2) is empirically treated with antivirals, antimalarics, tocilizumab, etc.[2] Production of a vaccine for COVID-19 has been attempted, although approval needs time.

We describe a possible, alternative approach for treating COVID-19. Lymphocyte count has been associated with increased disease severity risk.[3] Patients who died from COVID-19 showed a significantly lower lymphocyte count than survivors, therefore this should be closely monitored.[3] Repletion of lymphocytes could probably have beneficial effects on recovery.

A recent hypothesis suggests that the inhibition of the angiotensin 1 receptor (AT$_1$R) may provide benefits to COVID-19 patients.[4] This hypothesis is based on the observation that the SARS-CoV-2 virus uses angiotensin-converting enzyme 2 (ACE2) as a receptor to bind the virus to the bronchial cell membrane. The enzymes ACE and ACE2 belong to the same peptidase family but have two very different physiological functions. ACE cleaves angiotensin I to generate angiotensin II (Ang II), which binds to and activates AT$_1$R, and thus promoting vasoconstriction. ACE2 cleaves Ang II and generates angiotensin 1-7, a powerful vasodilator acting through Mas receptors. AT$_1$R antagonists are widely used in hypertensive patients but they increase the ACE2 cardiac expression in rats[5] and the urinary concentration of ACE2.[6]

It has been demonstrated that the binding of virus to ACE2 leads to ACE2 down-regulation,
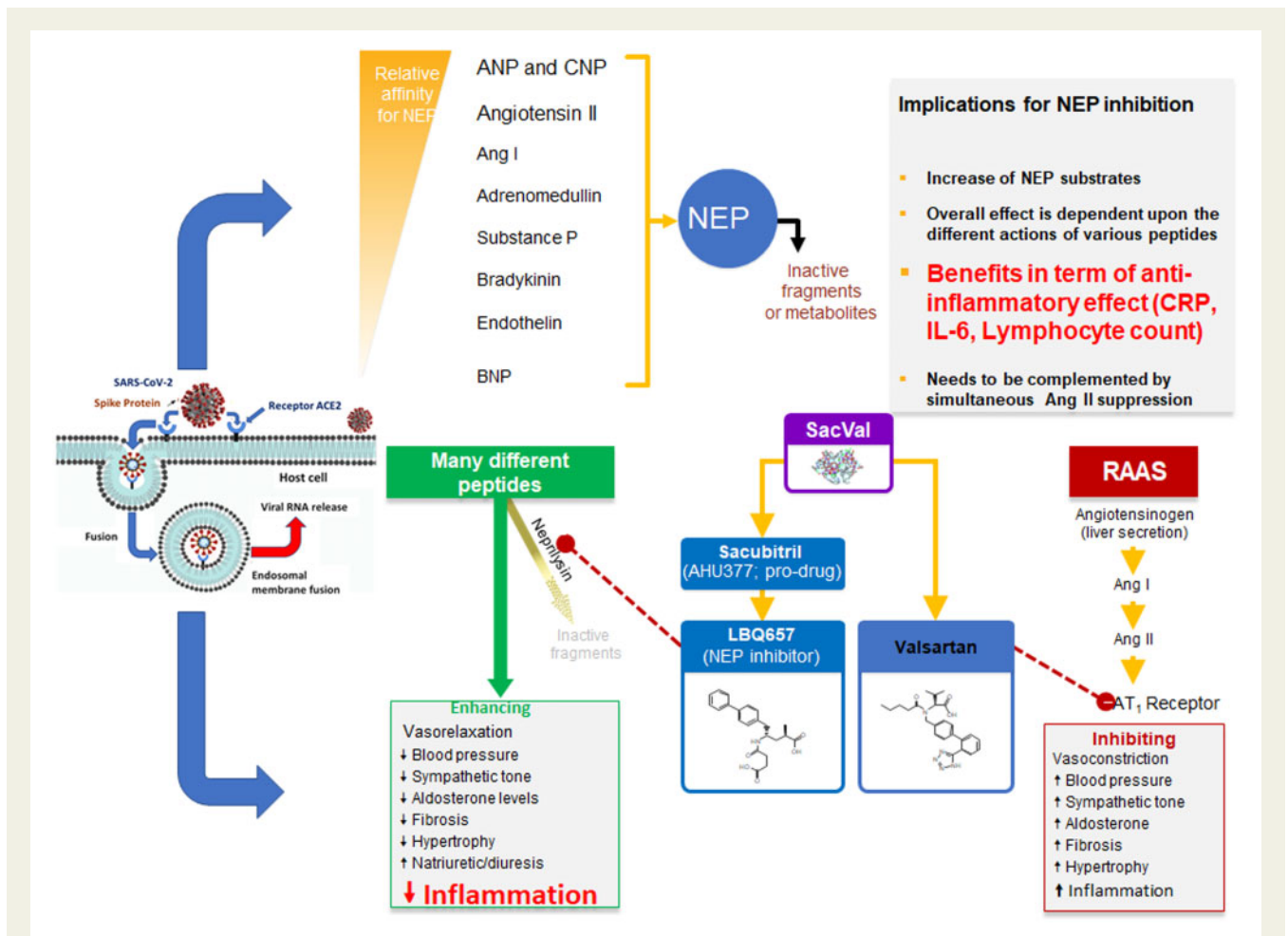
Figure 1 Double interaction between sacubitril and NEP and between valsartan and the AT1 receptor and their consequences on the development of the inflammatory response due to COVID-19 infection. RAAS, renin–renin–aldosterone system; SacVal, sacubitril/valsartan.

which increases the production of Ang II but reduces angiotensin 1-7. This contributes to increased AT_1-mediated pulmonary vascular permeability, thereby mediating increased lung pathology.[7] Therefore, higher ACE2 expression following chronic therapy with sartans may protect COVID-19 patients from acute lung injury rather than increasing the risk for SARS-CoV-2. Two complementary mechanisms may explain such a hypothesis: sartans will continue to block excessive angiotensin-mediated AT_1R activation due to the viral infection, and, in parallel, they will up-regulate ACE2, thus increasing angiotensin 1-7 production.

In such a setting, the role of neprilysin (NEP) and its inhibitor sacubitril should also be revised. Recently, Zhang et al.[8] demonstrated that sacubitril/valsartan reduced the concentration of pro-inflammatory cytokines and neutrophil count, while increasing lymphocyte count more than valsartan alone or placebo.[8] This finding might be related to the increase in plasma levels of atrial/brain/C-type natriuretic peptide, Ang I/II, substance P, bradykikin, and endothelin secondary to neprilisin inibition by sacubitril.[8] We have recently shown that early sacubitril/valsartan administration reduces high sensitivity C-reactive protein levels and increases lymphocyte count in patients with acute heart failure.[9]

These pieces of evidence support the biological plausibility of early administration of sacubitril/valsartan in COVID-19 patients, in order to maximize the anti-inflammatory effects of sacubitril and contain the effect of Ang I on the lungs (*Figure 1*).

Conflict of interest: none declared.

References

1. Rothan HA, Byrareddy SN. The epidemiology and pathogenesis of coronavirus disease (COVID-19) outbreak. *J Autoimmun* 2020;**109**:102433.
2. Cunningham AC, Goh HP, Koh D. Treatment of COVID-19: old tricks for new challenges. *Crit Care* 2020;**24**:91.
3. Yang X, Yu Y, Xu J, Shu H, Xia J, Liu H, Wu Y, Zhang L, Yu Z, Fang M, Yu T, Wang Y, Pan S, Zou X, Yuan S, Shang Y. Clinical course and outcomes of critically ill patients with SARS-CoV-2 pneumonia in Wuhan, China: a single-centered, retrospective, observational study. *Lancet Respir Med* 2020;doi: 10.1016/S2213-2600(20)30079-5.
4. Gurwitz D. Angiotensin receptor blockers as tentative SARS-CoV-2 therapeutics. *Drug Dev Res* 2020; doi: 10.1002/ddr.21656.
5. Wang X, Ye Y, Gong H, Wu J, Yuan J, Wang S, Yin P, Ding Z, Kang L, Jiang Q, Zhang W, Li Y, Ge J, Zou Y. The effects of different angiotensin II type 1 receptor blockers on the regulation of the ACE–AngII–AT1 and ACE2-Ang(1-7)–Mas axes in pressure overload-induced cardiac remodeling in male mice. *J Mol Cell Cardiol* 2016;**97**:180–190.
6. Furuhashi M, Moniwa N, Mita T, Fuseya T, Ishimura S, Ohno K, Shibata S, Tanaka M, Watanabe Y, Akasaka H, Ohnishi H, Yoshida H, Takizawa H, Saitoh S, Ura N, Shimamoto K, Miura T. Urinary angiotensin-converting enzyme 2 in hypertensive patients may be increased by olmesartan, an angiotensin II receptor blocker. *Am J Hypertens* 2015;**28**: 15–21.
7. Kuba K, Imai Y, Rao S, Gao H, Guo F, Guan B, Huan Y, Yang P, Zhang Y, Deng W, Bao L, Zhang B, Liu G, Wang Z, Chappell M, Liu Y, Zheng D, Leibbrandt A, Wada T, Slutsky AS, Liu D, Qin C, Jiang C, Penninger JM. A crucial role of angiotensin converting enzyme 2 (ACE2) in SARS coronavirus-induced lung injury. *Nat Med* 2005;**11**:875–879.
8. Zhang H, Liu G, Zhou W, Zhang W, Wang K, Zhang J. Neprilysin inhibitor–angiotensin II receptor blocker combination therapy (sacubitril/valsartan) suppresses atherosclerotic plaque formation and inhibits inflammation in apolipoprotein E-deficient Mice. *Sci Rep* 2019;**9**:6509.
9. Acanfora D, Scicchitano P, Acanfora C, Maestri R, Goglia F, Incalzi RA, Bortone AS, Ciccone MM, Uguccioni M, Casucci G. Early initiation of sacubitril/valsartan in patients with chronic heart failure after acute decompensation: a case series analysis. *Clin Drug Investig* 2020;doi: 10.1007/s40261-020-00908-4.

Domenico Acanfora[1],
Marco Matteo Ciccone[2],
Pietro Scicchitano[2,3]*,
Chiara Acanfora[1,4], and
Gerardo Casucci[1]

[1]Unit of Internal Medicine, San Francesco Hospital, Viale Europa 21, 82037 Telese Terme (BN), Italy; [2]Section of Cardiovascular Diseases, Department of Emergency and Organ Transplantation, University of Bari, School of Medicine, Bari, Italy; [3]Cardiology Unit, Hospital 'F. Perinei' ASL BA, Altamura, Bari, Italy; and [4]Department of Biotechnological and Applied Clinical Sciences, University of L'Aquila, L'Aquila, Italy

*Corresponding author. Cardiology Section, Hospital 'F. Perinei' ASL BA, Altamura, Bari, Italy. Tel.: +39 080 3108286, Email: piero.sc@hotmail.it; pietrosc.83@libero.it Correspondence