ACE2, Circumventricular Organs and the Hypothalamus, and COVID-19

Wei-Yi Ong1,2* · R. L. Satish1 · Deron R. Herr3

Received: 22 December 2021 / Accepted: 1 March 2022 / Published online: 22 April 2022
© The Author(s), under exclusive licence to Springer Science+Business Media, LLC, part of Springer Nature 2022

Abstract
The SARS-CoV-2 virus gains entry to cells by binding to angiotensin-converting enzyme 2 (ACE2). Since circumventricular organs and parts of the hypothalamus lack a blood–brain barrier, and immunohistochemical studies demonstrate that ACE2 is highly expressed in circumventricular organs which are intimately connected to the hypothalamus, and the hypothalamus itself, these might be easy entry points for SARS-CoV-2 into the brain via the circulation. High ACE2 protein expression is found in the subfornical organ, area postrema, and the paraventricular nucleus of the hypothalamus (PVH). The subfornical organ and PVH are parts of a circuit to regulate osmolarity in the blood, through the secretion of anti-diuretic hormone into the posterior pituitary. The PVH is also the stress response centre in the brain. It controls not only pre-ganglionic sympathetic neurons, but is also a source of corticotropin-releasing hormone, that induces the secretion of adrenocorticotropic hormone from the anterior pituitary. It is proposed that the function of ACE2 in the circumventricular organs and the PVH could be diminished by binding with SARS-CoV-2, thus leading to a reduction in the ACE2/Ang (1–7)/Mas receptor (MasR) signalling axis, that modulates ACE/Ang II/AT1R signalling. This could result in increased presynaptic activity/neuroendocrine secretion from the PVH, and effects on the hypothalamic–pituitary–adrenal axis activity. Besides the bloodstream, the hypothalamus might also be affected by SARS-CoV-2 via transneuronal spread along the olfactory/limbic pathways. Exploring potential therapeutic pathways to prevent or attenuate neurological symptoms of COVID-19, including drugs which modulate ACE signalling, remains an important area of unmet medical need.

Keywords COVID-19 · ACE2 · Circumventricular organs · Subfornical organ · Area postrema · Pineal gland · OVLT · Paraventricular nucleus of the hypothalamus

COVID-19 and Brain Blood Vessels, Glial Cells, and Neurons

Although the main clinical manifestations of COVID-19 are associated with respiratory or intestinal symptoms, reports of neurological signs and symptoms are increasing. The primary neurologic symptoms include ‘brain fog’ (81%), headache (68%), numbness/tingling (60%), dysgeusia (59%), anosmia (55%), and myalgia (55%). Most patients (85%) also report fatigue (Graham et al., 2021). The virus could potentially enter the brain through the circumventricular organs, disrupted blood–brain barrier, or retrograde transport via peripheral nerves (Kumar et al., 2021). Immune activation with astrocytosis, axonal damage, and blood–brain barrier leakage has been observed together with viral antigen and angiotensin-converting enzyme 2 (ACE2)-positive cells, at the blood–brain interface (Schwabenland et al., 2021). In addition, CSF levels of the chemokines CCL2 and CXCL8, and the blood vessel marker, vascular endothelium growth factor A are greater in severe COVID-19 cases than milder cases suggesting damage to the neurovascular unit (Bernard-Valnet et al., 2021). SARS-CoV-2 can stimulate extracellular neutrophils traps (NETs) in a process called NETosis. This normally functions as a defence against pathogens, but in excess may lead to increased reactive oxygen species (ROS) production in neutrophils and thrombus formation (Arcanjo et al., 2020; Pramitasuri et al., 2021). The SARS-CoV-2 spike protein reportedly binds to brain endothelial cells,
resulting in inflammatory changes and loss of blood–brain barrier integrity (Buzhdygan et al., 2020). Increased numbers of microglia and astrocytes, and elevated levels of pro-inflammatory markers are found in post-mortem specimens of the cerebral cortex in patients with COVID-19 (Boroujeni et al., 2021). Analysis of single-nucleus transcriptomes from the dorsolateral prefrontal cortex of patients with severe COVID-19 shows that transcriptional changes consistent with activated microglia are present, despite an absence of viral transcripts post-mortem (Fullard et al., 2021).

SARS-CoV-2 infection of cells occurs through the binding with ACE2. Virus binding to ACE2 induces conformational changes in the S1 subunit of its spike protein and exposes the S2’ cleavage site in the S2 subunit. The S2’ site is then cleaved by a protease (e.g. cathepsin or transmembrane protease serine 2 (TMPRSS2)) to expose a fusion peptide within the spike protein, that is able to attach to and induce fusion of the viral envelope with the host cell membrane, thus facilitating infection of the cell [for recent review, see (Jackson et al., 2022)]. Infection of neurons leads to activation of nicotinamide adenine dinucleotide phosphate oxidase 2 (NOX2), free radical formation, and release of ROS and inflammatory molecules (Sindona et al., 2021). Free radical attack alters the phospholipid composition of mitochondrial membranes (Clough et al., 2021). This could lead to a loss of mitochondrial membrane potential and activation of the NLRP3 inflammasome, resulting in increased expression of pro-inflammatory genes (Clough et al., 2021; Sita et al., 2021). Imaging mass spectrometry analysis indicates astrocytosis, axonal damage and blood–brain barrier leakage, together with the detection of viral antigen in ACE2-positive cells in vascular compartments in post-mortem cases of COVID-19 (Schwabenland et al., 2021). Oligodendrocytes could also be affected by the SARS-CoV-2 virus and/or activated microglia, and this might result in CNS demyelination (Pan et al., 2020).

It is possible that the same mechanisms of COVID-19-induced damage in glial or endothelial cells described above could also occur in neurons. Increased levels of serum neurofilament light, indicating central and/or peripheral neuronal damage, are found in hospitalized patients with COVID-19, whereas no elevation is detected in milder cases (Paterson et al., 2021). Likewise, greater levels of CSF neurofilament light are present in critical cases of COVID-19 cases, compared to less severe cases (Garcia et al., 2021). RNA sequencing analyses of the amygdala of severe COVID-19 cases show increases in neuroinflammatory genes, but decreases in neuronal genes including those related to synaptic function (Piras et al., 2021). In addition, network analyses reveal a close relationship between COVID-19-induced neuroinflammation and pathways involved in Alzheimer’s disease (Zhou et al., 2021). Together, these findings indicate increased involvement of microglia, brain microvessels, and neurons with greater severity of COVID-19 (Stefano et al., 2021).

The SARS-CoV-2 virus enters cells by binding to ACE2 (Hoffmann et al., 2020). In this situation, ACE2 functions as a receptor for the spike protein of the SARS-CoV-2 and facilitates internalization of the virus and infection of the host cell (Jackson et al., 2022). It is also important to note that infection of cells by SARS viruses including SARS-CoV-2 results in a decrease of ACE2 expression (Kuba et al., 2005; Triana et al., 2021) and loss of ACE2 activity (Glowacka et al., 2010; Haga et al., 2010). This could prevent ACE2 from performing its normal function, to regulate or act as a ‘brake’ against ACE/Ang II/AT1R signalling (see below). Diminished effect on reducing angiotensin II (Ang II) signalling is suggested to contribute to injury in patients with COVID-19 (Sriram & Insel, 2020).

ACE2 and the Hypothalamus

ACE helps in the formation of Ang II from angiotensin I, and ACE2 cleaves angiotensin I and angiotensin II into angiotensin (1–9) and angiotensin (1–7) (Ang (1–7)), respectively. The metabolism of Ang II to the vasodilatory peptide Ang(1–7) by ACE2 is part of the ACE2/Ang (1–7)/Mas receptor (MasR) axis which helps to reduce blood pressure, as opposed to the ACE/Ang II/AT1R Receptor (AT1R) axis, which increases blood pressure (Fig. 1). Surface-Enhanced Laser Desorption Ionization–Time of Flight mass spectroscopic analyses to study Ang processing in normal mice show that not only ACE2 activity is found in the brain and kidney but also that the hypothalamus is the part of the brain that contains the highest ACE2 activity. This is in contrast to ACE, which is most active in the plasma (Elased et al., 2008). Overexpression of ACE2 induced by injection of adenovirus encoding ACE2 in the paraventricular hypothalamic nucleus results in attenuation of Ang II-induced hypertension in rats (Sriramula et al., 2011). On the other hand, selective knockdown of ACE2 in the subfornical organ and paraventricular hypothalamic nucleus in mice results in partial loss of the ability to regulate deoxycorticosterone acetate-salt induced neurogenic hypertension (Xia et al., 2015). Studies also show lower protein expression of ACE2 in spontaneously hypertensive rats, compared to normotensive rats (Wang et al., 2017). Moreover, downregulation of components of the ACE2/Ang (1–7)/MasR axis is present in spontaneously hypertensive rats (Han et al., 2020). These findings highlight an important role of the hypothalamic ACE2/Ang (1–7)/MasR axis in regulation of blood pressure.

Overexpression of ACE2 also significantly decreases anxiety-like behaviour in paradigms dependent on approach-avoidance conflict and novelty, but has no effect on basal and/or stress-induced
Proposed Function of ACE2 in Neurons

Fig. 1 Proposed general functions of ACE2 in neurons. ACE2 catalyses the breakdown of Ang II to Ang (1–7). The latter is an agonist of the Mas receptor (MasR), which has been colocalised with GABAergic neurons (in the amygdala) (Wang et al., 2016). Ang (1–7) causes inhibition of principal neurons by promoting GABAergic transmission in a MasR-dependent manner (Wang et al., 2016). This could occur via facilitation of GABA release through a nitric oxide-mediated pathway (Stragier et al., 2005) and possibly supplemented by an increase in GABA production via upregulation of glutamate decarboxylase 67 (GAD67) expression (studied in the pancreas) (Ma et al., 2020). GABA binds to GABA_A receptors to produce inhibition of the postsynaptic neuron. ACE2 is shed from the cell membrane, resulting in loss of activity through the action of another enzyme, ADAM17 (Xia et al., 2013). The latter is mostly expressed in glutamatergic projection neurons (Xu et al., 2019). In this manner, ACE2 and ADAM17 exert opposite effects on neuronal excitability (studied in presymptomatic projection neurons of the paraventricular hypothalamic nucleus) (Mukerjee et al., 2019). It is possible that the above schema could also apply to neurons in other parts of the brain including those in the cerebral cortex and hippocampus, which express lower levels of ACE2 (Doobay et al., 2007).

In contrast to Ang (1–7), Ang II has an excitatory effect on neurons (Fig. 2). Both in vivo and in vitro studies show that Ang II stimulates PVN neuronal activity (Bains et al., 1992; Cato & Toney, 2005; Li et al., 2003). This is likely through a direct effect on AT1 receptors (AT1R) which has been localized to neurons in the PVH and the circumventricular organs, including the organum vasculosum of the lamina terminalis, subfornical organ, area postrema, and median eminence (Sumners et al., 2020). An indirect effect involving inhibition of the astrocyte glutamate transporter has also been proposed (Stern et al., 2016). ACE2 is also known to have anti-inflammatory and anti-oxidative effects (Sriramula et al., 2011). This could occur via modulation of levels of ANG II which has both a pro-inflammatory effect via the production of tumour necrosis factor alpha (TNFα); and a pro-oxidative effect by increasing NADPH-oxidase expression and ROS production (Sriramula et al., 2013). Together, these findings indicate an important role of hypothalamic ACE2 in modulating the stress response.

ACE2 has an effect on regulating neuronal activity (Fig. 1). As noted above, ACE2 catalyses the breakdown of Ang II to Ang (1–7). The latter is an agonist of the Mas receptor (MasR), which has been colocalised with GABAergic neurons (in the amygdala) (Wang et al., 2016). Ang (1–7) causes inhibition of principal neurons by promoting GABAergic transmission in a MasR-dependent manner (Wang et al., 2016). This could occur via facilitation of GABA release through a nitric oxide-mediated pathway (Stragier et al., 2005) and possibly supplemented by an increase in GABA production via upregulation of glutamate decarboxylase 67 (GAD67) expression (studied in the pancreas) (Ma et al., 2020). GABA binds to GABA_A receptors to produce inhibition of the postsynaptic neuron. ACE2 is shed from the cell membrane, resulting in loss of activity through the action of another enzyme, ADAM17 (Xia et al., 2013). The latter is mostly expressed in glutamatergic projection neurons (Xu et al., 2019). In this manner, ACE2 and ADAM17 exert opposite effects on neuronal excitability (studied in presymptomatic projection neurons of the paraventricular hypothalamic nucleus) (Mukerjee et al., 2019). ADAM17 activity itself could be affected by oxidative stress. Mice that received an antioxidant, alpha-lipoic acid, showed modulation of oxidative stress-induced increased ADAM17 activity and decreased ACE2 activity in the hypothalamus (de Queiroz et al., 2015). It is possible that the above schema could also apply to neurons in other parts of the brain including those in the cerebral cortex and hippocampus, which express lower levels of ACE2 (Doobay et al., 2007).
The seven circumventricular organs are the organum vasculosum of the lamina terminalis (OVLT), subfornical organ, area postrema, pineal gland, subcommissural organ, median eminence, and neurohypophysis. Previous studies have shown high level of ACE2 expression in circumventricular organs (Doobay et al., 2007) and nuclei that are connected to the circumventricular organs. The latter includes the nucleus of the tractus solitarius, rostral ventrolateral medulla, and paraventricular nucleus of the hypothalamus (Doobay et al., 2007; Kar et al., 2010; Zucker et al., 2014). ACE2 is localised in neurons of the above nuclei (Doobay et al., 2007; Kar et al., 2010). The OVLT, subfornical organ, and area postrema are important in regulation of osmotic thirst, anti-diuretic hormone release, and blood pressure (Johnson et al., 1996; McKinley et al., 1992b). Their constituent neurons project not only to the paraventricular hypothalamic nucleus, but also the brainstem. Axons send collateral branches to the nucleus of the tractus solitarius, and rostral ventrolateral medulla, before terminating on pre-ganglionic sympathetic neurons in the intermediolateral horn of the spinal cord to increase sympathetic activity (Duan et al., 2020; Gu, 2021; Thomas, 2011).

ACE2 and Circumventricular Organs that are Intimately Connected to the Hypothalamus

The seven circumventricular organs are the organum vasculosum of the lamina terminalis (OVLT), subfornical organ, area postrema, pineal gland, subcommissural organ, median eminence, and neurohypophysis. Previous studies have shown high level of ACE2 expression in circumventricular organs (Doobay et al., 2007) and nuclei that are connected to the circumventricular organs. The latter includes the nucleus of the tractus solitarius, rostral ventrolateral medulla, and paraventricular nucleus of the hypothalamus (Doobay et al., 2007; Kar et al., 2010; Zucker et al., 2014). ACE2 is localised in neurons of the above nuclei (Doobay et al., 2007; Kar et al., 2010). The OVLT, subfornical organ, and area postrema are important in regulation of osmotic thirst, anti-diuretic hormone release, and blood pressure (Johnson et al., 1996; McKinley et al., 1992b). Their constituent neurons project not only to the paraventricular hypothalamic nucleus, but also the brainstem. Axons send collateral branches to the nucleus of the tractus solitarius, and rostral ventrolateral medulla, before terminating on pre-ganglionic sympathetic neurons in the intermediolateral horn of the spinal cord to increase sympathetic activity (Duan et al., 2020; Gu, 2021; Thomas, 2011).

Organum Vasculosum of the Lamina Terminalis (OVLT)

The OVLT is located in the anterior wall of the third ventricle, approximately midway between the optic chiasm and the anterior commissure (McKinley et al., 2004). Unlike other species, fenestrations in the capillary endothelial cells have not been observed in the human OVLT. Nevertheless, reports of imbibed silver suggest altered blood–brain barrier characteristics of the human OVLT (Landas et al., 1985). The major afferent inputs to the OVLE appear to come from several hypothalamic nuclei (median preoptic, lateral preoptic, anterior, lateral, dorsomedial, and ventromedial nuclei) and extrahypothalamic regions (subfornical organ, locus coeruleus, central grey) (Camacho & Phillips, 1981). Studies in the rat indicate that the OVLE project to the median preoptic nucleus immediately dorsal to the OVLT, the supraoptic nucleus (Camacho & Phillips, 1981), and the paraventricular hypothalamic nucleus (Sunn et al., 2001). The OVLT has a role as an osmoreceptor, a mediator of the febrile response, and
subfornical organ to the paraventricular hypothalamic nucleus and supraoptic nucleus change the excitability of anti-diuretic hormone and oxytocin neurons projecting to the posterior pituitary (Ferguson et al., 1984a); as well as that of anti-diuretic hormone containing neurons projecting to the dorsolateral medulla (Ferguson et al., 1984b). Studies using a combination of Fos staining and tract tracing show that neurons in the subfornical organ which project to the supraoptic and paraventricular hypothalamic nuclei, can be activated by hypertonicity or by circulating levels of Ang II or relaxin (Larsen & Mikkelsen, 1995; Oldfield et al., 1994; Sunn et al., 2001). Reduction in blood flow to the renal artery or increased sympathetic activity via renal nerves results in secretion of renin from juxtaglomerular cells in the tunica media of the renal arteries as they enter the glomeruli. Renin converts angiotensinogen to angiotensin I. Most of the angiotensin-converting enzyme is located in endothelial cells [Reviewed in (Barrett et al., 2016)], but some of the blood-borne angiotensin I reaches the subfornical organ where it is converted to Ang II. This leads to activation of neurons in the subfornical organ, induction of water drinking, anti-diuretic hormone secretion, and a central pressor response (McKinley et al., 1992a, 1997; Simpson, 1981).

**Pineal Gland**

The pineal gland in humans is a solid organ located in the midline roof of the third ventricle. It contains astrocytes and pinealocytes, which secrete melatonin. The central part of the gland highly vascularized by large sinusoid capillaries and its peripheral part poorly vascularized by small and fine blood vessels (Duvernoy et al., 2000). The major neural to the pineal gland is from the peripheral nervous system, i.e. sympathetic innervation from the superior cervical ganglion (Kappers, 1965). Signals from the retina are transmitted via the retinohypothalamic tract to the suprachiasmatic nucleus of the hypothalamus; from here, information is relayed via the paraventricular hypothalamic nucleus to the intermedialateral cell column of the spinal cord. The latter sends axons to synapse in the superior cervical ganglion; thereafter, postganglionic axons travel along the great cerebral vein of Galen to enter the pineal gland and innervate pinealocytes (Larsen et al., 1998; Teclemariam-Mesbah et al., 1999). It is through this pathway that information regarding changes in the day-night cycle and seasonal alterations in illumination is transmitted to the pineal gland to regulate melatonin secretion (Tamarkin et al., 1985). mRNA for angiotensinogen, angiotensin receptor type 1A (AT1aR) and 1B (AT1bR), and ACE have been detected in the pineal glands of rats. The presence of angiotensin receptors points to a role for renin-angiotensin system in the physiology of the pineal gland (Baltatu et al., 1998).

**Area Postrema**

The area postrema is located in the dorsomedial medulla oblongata as two prominences bulging into the most caudal aspect of the fourth ventricle (McKinley et al., 2004). This nucleus and the specialized region of the adjacent nucleus of the tractus solitarius have permeable capillaries (Gross et al., 1990). Ablation of the area postrema in animals interferes with taste aversion, due to nausea-inducing stimuli (Berger et al., 1973). The area postrema is long thought to be the ‘chemoreceptor trigger zone’ for vomiting. This nucleus receives afferents from the lateral parabrachial nucleus, the adjacent nucleus of the tractus solitarius (Shapiro & Miselis, 1985; van der Kooy & Koda, 1983), and the paraventricular hypothalamic nucleus (Shapiro & Miselis, 1985). In turn, it projects to the nucleus of the tractus solitarius, nucleus ambiguus, and noradrenergic neurons of the caudal ventrolateral medulla (McKinley et al., 2004). The unique position of the area postrema at the point of entry of visceral sensory information, as well as lack of a blood–brain barrier that exposes it to the circulation, enables it to integrate and modulate homeostatic responses in the body (Gross et al., 1990; Miselis et al., 1987; Shaver et al., 1991).
**Median Eminence**

The median eminence and arcuate nucleus of the hypothalamus have a deficient blood–brain barrier [Reviewed in (Haddad-Tovolli et al., 2017)]. Moreover, dense binding of radiiodinated analogs of Ang II have been observed in the human median eminence and arcuate nucleus (McKinley et al., 1987).

**Subcommissural Organ**

Unlike other mammals, the human subcommissural organ is only clearly evident in the foetus and new-born. In the adult it has almost completely disappeared (McKinley et al., 2004).

**The Hypothalamus and COVID-19 (Fig. 2)**

There has been increasing attention to the role of the hypothalamus and brainstem autonomic centres in the pathophysiology of COVID-19 (Chigr et al., 2020). The fact that circumventricular organs lack a blood–brain barrier (Haddad-Tovolli et al., 2017), together with the high levels of ACE2 expression in circumventricular organs and hypothalamus (Doobay et al., 2007), suggests that they could be easy entry points for SARS-CoV-2 into the brain via the circulation. On the corollary, the observation that some of the ACE2-positive regions of the brain lack a blood–brain barrier, could imply that they are amenable to treatment with antiviral agents/other therapeutic agents, that otherwise have difficulty crossing the barrier. Hypothalamic pathology is found in a case of COVID-19 (Pascual-Goni et al., 2020), and involvement of the hypothalamic/pituitary gland as well as other endocrine glands has been suggested to result in an ‘endocrine phenotype’ of the disease (Frara et al., 2021; Mussa et al., 2021; Puig-Domingo et al., 2021).

The paraventricular nucleus of the hypothalamus receives abundant connections from the circumventricular organs which contain high level of ACE2 expression, and itself expresses ACE2 protein (Doobay et al., 2007), as mentioned above. These are part of a circuit to regulate osmolarity in the blood through secretion of anti-diuretic hormone (ADH) from the paraventricular hypothalamic nucleus. It is interesting that hyponatremia has been found in a relatively high proportion of COVID-19 patients. The Health Outcome Predictive Evaluation for COVID-19 (HOPE) study found that 20.5% of COVID-19 patients had hyponatremia at admission (Ruiz-Sanchez et al., 2020), and greater morbidity and mortality are reported in hospitalized patients with hyponatremia (Tzoulis et al., 2021). Many of the cases of hyponatremia likely occur in the setting of Syndrome of Inappropriate Secretion of Antidiuretic Hormone (SIADH) (Frara et al., 2021; Habib et al., 2020; Yousaf et al., 2020). The latter is a disorder of impaired water excretion caused by the inability to suppress the secretion of anti-diuretic hormone. If water intake exceeds the reduced urine output, the ensuing water retention leads to the development of hyponatremia. This could be due to loss of function of ACE2 as a result of binding to SARS-CoV-2, and consequent increase in ACE signalling. Changes in the subfornical organ neurons likely affect its output to the paraventricular hypothalamic nucleus. Interference with ADH release into the posterior pituitary via an effect on the subfornical organ and paraventricular hypothalamic nucleus has been postulated to lead to hydromineral electrolytic imbalance in some patients with COVID-19 (de Melo et al., 2021).

The paraventricular hypothalamic nucleus is also the stress response centre in the brain. Neurons in this nucleus coordinate this response through their neuronal connections, and via their neuroendocrine secretion of CRH (Kim et al., 2019). The latter has several actions, but its main role is the central driver of the HPA axis. It is proposed that inflammatory mediators released at the site of COVID-19 infection are transmitted as stress signals to cause dysfunction to the complex neurological circuit of the paraventricular hypothalamic nucleus, and result in interference with the modulation of stress (Mackay, 2021). Higher levels of cortisol have been detected in patients with severe COVID-19 compared to less severe cases (Guven & Gultekin, 2021; Tan et al., 2020). These findings are consistent with the findings of preclinical studies, that mice with overexpression of ACE2 in the hypothalamus show reduced plasma corticosterone level in response to restraint stress. These animals also exhibit decreased anxiety-like behaviour in the “elevated-plus maze” and open field test (Wang et al., 2018). High level of stress might induce downstream effects on hippocampal neurogenesis in the dentate gyrus, leading to deficits in the formation of new memories (McEwen, 1999). It is proposed that the function of ACE2-positive neurons in the paraventricular hypothalamic nucleus is affected by binding with the SARS-CoV-2 virus, resulting in interference with modulation of Ang II function and reduced modulation of stress/anxiety, in patients with COVID-19. Some of the other psychiatric consequences of COVID-19 patients such as post-viral fatigue states, aberrant daytime oscillation in alertness, disturbed sleep cycles, and significant fluctuating anxiety have also been postulated to be due to altered function of the paraventricular hypothalamic nucleus (Rosenweig et al., 2020).

The paraventricular hypothalamic nucleus and several other hypothalamic nuclei are the source of releasing and inhibiting hormones, which control the secretion of hormones from the anterior pituitary. Gonadotropin-releasing hormone neurons are located primarily in the medial preoptic area; growth-hormone-inhibiting
hormone (= somatostatin) secreting neurons are found in the periventricular nucleus; thyrotropin-releasing hormone (TRH) and corticotropin-releasing hormone (CRH) secreting neurons are located in the paraventricular nucleus; while growth hormone releasing (GRH) neurons are located in the arcuate nucleus [Reviewed in (Barrett et al., 2016)]. These hormones are released from nerve terminals in the median eminence of the hypothalamus and carried via the bloodstream to the anterior pituitary, where they cause secretion of anterior pituitary hormones. The median eminence and arcuate nucleus of the hypothalamus have a deficient blood–brain barrier [Reviewed in (Haddad-Tovolli et al., 2017)] and could be readily affected by circulating SARS-CoV-2. Damage to the hypothalamus and/or pituitary gland as a result of COVID-19 has been suggested to result in anomalies of the hypothalamus–pituitary–thyroid axis (Caron, 2020; Malik et al., 2021). This could result in a central hypothroidism (Sandru et al., 2021), besides potential damage from the virus to the thyroid gland itself (Croce et al., 2021). COVID-19 injury of the hypothalamus is also proposed to lead to malfunction of the hypothalamic–pituitary–testicular axis (Ardestani Zadeh & Arab, 2021; Selvaraj et al., 2021) with possible effects on testicular function (Selvaraj et al., 2021).

Besides potential infection from the bloodstream, the hypothalamus might also be affected by transneuronal spread of SARS-CoV-2, through its neuronal connections with the olfactory and limbic systems. These pathways have been postulated to be a potential route of spread of SARS-CoV-2 from the olfactory epithelium (Baig & Sanders, 2020; Bougakov et al., 2021; Jiao et al., 2021; Mussa et al., 2021). After receiving inputs from the olfactory epithelium in the nasal cavity, the olfactory bulb projects via the medial olfactory stria to the septal nuclei, which in turn projects to the hypothalamus [Reviewed in (Getz, 2007)]. The olfactory bulb also projects via the lateral olfactory stria to the amygdala, which projects to the hypothalamus via the stria terminalis. Other parts of the lateral olfactory stria terminate in the entorhinal cortex, which projects to the hippocampus, and the latter projects to the mammillary bodies of the hypothalamus via the fornix [Reviewed in (Getz, 2007)]. Previous studies have shown the ability of SARS-CoV to induce neuronal death in mice by invading the brain via the olfactory epithelium (Netland et al., 2008). Direct infection of neurons in organoids by SARS-CoV-2 has also been demonstrated (Song et al., 2021). Decreased 18F-FDG PET metabolism in olfactory/limbic regions and the hypothalamus has been found in a pilot study of COVID-19 patients (Guedj et al., 2021), suggesting that olfactory and limbic regions are affected by the virus.

Conclusion

A better understanding of the role of ACE2 in circumventricular organs and the hypothalamus may help in appreciating the effects of neurological symptoms of COVID and its underlying mechanisms. Given the socio-economic impact and drain on healthcare resources from COVID and the scale and persistence of the pandemic, exploring potential therapeutic pathways to prevent or attenuate these long-lasting neurological symptoms, including drugs which modulate ACE signalling, remains an important area of unmet medical need.

Acknowledgements This work was supported by grants from the Ministry of Education (NUHSRO/2019/051/T1/Seed-Mar/04) and the National Medical Research Council of Singapore (HLCA21Jan-0019).

Declarations

Conflict of interest The authors have no conflicts of interest to declare.

References

Arcanjo, A., Logullo, J., Menezes, C. C. B., de Souza Carvalho, T. C., Dos Reis, M. C., de Castro, G. M. M, et al. (2020). The emerging role of neutrophil extracellular traps in severe acute respiratory syndrome coronavirus 2 (COVID-19). *Science and Reports*, 10(1), 19630. https://doi.org/10.1038/s41598-020-76781-0

Ardestani Zadeh, A., & Arab, D. (2021). COVID-19 and male reproductive system: Pathogenic features and possible mechanisms. *Journal of Molecular Histology*, 52(5), 869–878. https://doi.org/10.1007/s10735-021-10003-3

Baig, A. M., & Sanders, E. C. (2020). Potential neuroinvasive pathways of SARS-CoV-2: Deciphering the spectrum of neurological deficit seen in coronavirus disease-2019 (COVID-19). *Journal of Medical Virology*, 92(10), 1845–1857. https://doi.org/10.1002/jmv.26105

Bains, J. S., Potyok, A., & Ferguson, A. V. (1992). Angiotensin II actions in paraventricular nucleus: Functional evidence for neurotransmitter role in efferents originating in subfornical organ. *Brain Research*, 599(2), 223–229. https://doi.org/10.1016/0006-8993(92)90395-p

Baltatu, O., Lippoldt, A., Hansson, A., Ganten, D., & Bader, M. (1998). Local renin-angiotensin system in the pineal gland. *Molecular Brain Research*, 54(2), 237–242. https://doi.org/10.1016/s0169-328x(97)00339-2

Barrett, K. E., Barman, S. M., Boitano, S., & Brooks, H. L. (2016). Ganong’s review of medical physiology. Mc Graw Hill.

Berger, B. D., Wise, C. D., & Stein, L. (1973). Area postrema damage and bait shyness. *Journal of Comparative and Physiological Psychology*, 82(3), 475–479. https://doi.org/10.1037/h0034112

Bernard-Valnet, R., Perriot, S., Canales, M., Pizzarotti, B., Caranzano, L., Castro-Jimenez, M., et al. (2021). Encephalopathies associated with severe COVID-19 present neurovascular unit alterations without evidence for strong neuroinflammation. *Neurology - Neuroimmunology Neuroinflammation*. https://doi.org/10.1212/NXI.0000000000001029
Guven, M., & Gultekin, H. (2021). Could serum total cortisol level at admission predict mortality due to coronavirus disease 2019 in the intensive care unit? A prospective study. Sapo Paulo Medical Journal, 139(4), 398–404. https://doi.org/10.1590/1516-3180.2020.0722.R1.230202

Habib, M. B., Sardar, S., & Sajid, J. (2020). Acute symptomatic hyponatremia in setting of SIADH as an isolated presentation of COVID-19. Idcesases, 21, e00859. https://doi.org/10.1016/j.idcr.2020.e00859

Haddad-Tovolli, R., Dragan, N. R. V., Ramalho, A. F. S., & Velloso, L. A. (2017). Development and function of the blood–brain barrier in the context of metabolic control. Frontiers in Neuroscience, 11, 224. https://doi.org/10.3389/fnins.2017.00224

Haga, S., Nagata, N., Okamura, T., Yamamoto, N., Sata, T., Yamamoto, N., et al. (2010). TACE antagonists blocking ACE2 shedding caused by the spike protein of SARS-CoV are candidate antiviral compounds. Antiviral Research, 85(3), 551–555. https://doi.org/10.1016/j.antiviral.2009.12.001

Han, W., Wang, M., Zhai, X., Gan, Q., Guan, S., & Qu, X. (2020). Chemical renal denervation-induced upregulation of the ACE2/Ang (1–7)/Mas axis attenuates blood pressure elevation in spontaneously hypertensive rats. Clinical and Experimental Hypertension, 42(7), 661–668. https://doi.org/10.1080/10641963.2020.1772812

Han, Y., Sun, H. J., Li, P., Gao, Q., Zhou, Y. B., Zhang, F., et al. (2012). Angiotensin-(1–7) in paraventricular nucleus modulates sympathetic activity and cardiac sympathetic afferent reflex in renovascular hypertensive rats. PloS ONE, 7(11), e49866. https://doi.org/10.1371/journal.pone.0049866

Hoffmann, M., Kleine-Weber, H., Schroeder, S., Kruger, N., Herrler, T., Erichsen, S., et al. (2020). SARS-CoV-2 cell entry depends on ACE2 and TMPRSS2 and is blocked by a clinically proven protease inhibitor. Cell, 181(2), 271–280.e278, https://doi.org/10.1016/j.cell.2020.02.052

Jackson, C. B., Farzan, M., Chen, B., & Choe, H. (2022). Mechanisms of SARS-CoV-2 entry into cells. Nature Reviews Molecular Cell Biology, 23(1), 3–20. https://doi.org/10.1038/s41580-021-00418-x

Jiao, L., Yang, Y., Yu, W., Zhao, Y., Long, H., Gao, J., et al. (2021). A crucial role of angiotensin converting enzyme 2/angiotensin-(1–7)/Mas axis attenuates the cardinal activity and cardiac sympathetic afferent reflex in renovascular hypertensive rats. Frontiers in Neuroscience, 15(4), 2609–2627. https://doi.org/10.1523/jneurosci.05-04-02609.1995

Li, D. P., Chen, S. R., & Pan, H. L. (2003). Angiotensin II stimulates spinally projecting paraventricular neurons through presynaptic disinhibition. Journal of Neuroscience, 23(12), 5041–5049. https://doi.org/10.1523/jneurosci.23-12-05041.2003

Ma, X., Gao, F., Chen, Q., Xuan, X., Wang, Y., Deng, H., et al. (2020). ACE2 modulates glucose homeostasis through GABA signaling during metabolic stress. Journal of Endocrinology, 246(3), 223–236. https://doi.org/10.1530/joe-19-0471

Mackay, A. (2021). A paradigm for post-covid-19 fatigue syndrome analogous to ME/CFS. Frontiers in Neuroscience, 12, 701419. https://doi.org/10.3389/fnneu.2021.701419

Malik, J., Zaidi, S. M. J., Waqar, A. U., Khawaja, H., Malik, A., Ishaq, U., et al. (2021). Association of hypothyroidism with acute COVID-19: A systematic review. Expert Review of Endocrinology and Metabolism, 16(5), 251–257. https://doi.org/10.1080/17446651.2021.1968830

Martins Lima, A., Xavier, C. H., Ferreira, A. J., Raizada, M. K., Walukat, G., Velloso, E. P., et al. (2013). Activation of angiotensin-converting enzyme 2/angiotensin-(1–7)/Mas axis attenuates the cardinal activity to acute emotional stress. American Journal of Physiology-Heart and Circulatory Physiology, 305(7), H1057–1067. https://doi.org/10.1152/ajpheart.00433.2013

McEwen, B. S. (1999). Stress and hippocampal plasticity. Annual Review of Neuroscience, 22, 105–122. https://doi.org/10.1146/annurev.neuro.22.1.105

McKinley, M. J., Allen, A., Clevers, J., Denton, D. A., & Mendelsohn, F. A. (1986). Autoradiographic localization of angiotensin receptors in the sheep brain. Brain Research, 375(2), 373–376. https://doi.org/10.1016/0006-8993(86)90761-4

McKinley, M. J., Allen, A. M., Clevers, J., Paxinos, G., & Mendelsohn, F. A. (1987). Angiotensin receptor binding in human hypothalamus: Autoradiographic localization. Brain Research, 420(2), 375–379. https://doi.org/10.1016/0006-8993(87)91260-1

McKinley, M. J., Badoer, E., & Oldfield, B. J. (1992a). Intravenous angiotensin II induces Fos-immunoreactivity in circumventricular organs of the lamina terminalis. Brain Research, 594(2), 295–300. https://doi.org/10.1016/0006-8993(92)91138-5

McKinley, M. J., Bicknell, R. J., Hards, D., McAllen, R. M., Vivas, L., Weisgerber, R. S., et al. (1992b). Efferent neural pathways of the lamina terminalis. In G. Paxinos, & J. K. Mai (Eds.), Progress in brain research (Vol. 91, pp. 395–402). Elsevier.

McKinley, M. J., Clarke, I. J., & Oldfield, B. J. (2004). Circumventricular organs. In G. Paxinos, & J. K. Mai (Eds.), The human nervous system. (2nd ed.). Elsevier.

McKinley, M. J., Colvill, L. M., Giles, M. E., & Oldfield, B. J. (1997). Distribution of Fos-immunoreactivity in rat brain following a dipsogenic dose of captopril and effects of angiotensin receptor

Kopers, J. A. (1965). Survey of the innervation of the epiphysis cerevisi.
Sunn, N., McKinley, M. J., & Oldfield, B. J. (2001). Identification of
Tamarkin, L., Baird, C. J., & Almeida, O. F. (1985). Melatonin: A
Sumners, C., Alleyne, A., Rodríguez, V., Pioquinto, D. J., Ludin, J.
Stragier, B., Hristova, I., Sarre, S., Ebinger, G., & Michotte, Y. (2005).
Stern, J. E., Son, S., Biancardi, V. C., Zheng, H., Sharma, N., & Patel,
Tzoulis, P., Waung, J. A., Bagkeris, E., Hussein, Z., Biddanda, A.,
Thomas, G. D. (2011). Neural control of the circulation. 
Triana, S., Metz-Zumaran, C., Ramírez, C., Kee, C., Doldan, P., Shah-
Wang, K., Xu, Y., Yang, W., & Zhang, Y. (2017). Insufficient hypotha-
van der Kooy, D., & Koda, L. Y. (1983). Organization of the projec-
Wang, L., de Kloet, A. D., Pati, D., Hiller, H., Smith, J. A., Pioquinto,
Wang, L. A., de Kloet, A. D., Smeltzer, M. D., Cahill, K. M., Hiller,
Xia, H., de Queiroz, T. M., Sriramula, S., Feng, Y., Johnson, T., Mun-
Xia, H., Sriramula, S., Chhabra, K. H., & Lazartigues, E. (2013).
Xia, J., Molinas, A. J. R., Mukerjee, S., Morgan, D. A., Rahmouni, K.,
Yousaf, Z., Al-Shokri, S. D., Al-Soub, H., & Mohamed, M. F. H. (2021).
Zhou, Y., Xu, J., Hou, Y., Leverenz, J. B., Kallianpur, A., Mehra, R.,
Zucker, I. H., Xiao, L., & Haack, K. K. (2014). The central renin-angio-
Publisher’s Note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.