Virus interactions with the actin cytoskeleton—what we know and do not know about SARS-CoV-2

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Abstract

The actin cytoskeleton and actin-dependent molecular and cellular events are responsible for the organization of eukaryotic cells and their functions. Viruses, including severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), depend on host cell organelles and molecular components for cell entry and propagation. Thus, it is not surprising that they also interact at many levels with the actin cytoskeleton of the host. There have been many studies on how different viruses reconfigure and manipulate the actin cytoskeleton of the host during successive steps of their life cycle. However, we know relatively little about the interactions of SARS-CoV-2 with the actin cytoskeleton. Here, we describe how the actin cytoskeleton is involved in the strategies used by different viruses for entry, assembly, and egress from the host cell. We emphasize what is known and unknown about SARS-CoV-2 in this regard. This review should encourage further investigation of the interactions of SARS-CoV-2 with cellular components, which will eventually be helpful for developing novel antiviral therapies for mitigating the severity of COVID-19.

Introduction

In all eukaryotic cells, the actin cytoskeleton is indispensable for the anchoring and organization of the subcellular compartments and organelles. It regulates most cell functions, including cell–cell adhesion, cell movement, and cell division [1–3]. Cortical actin participates in all events related to the expression and presentation of membrane/cell surface molecules, the formation and movement of endocytic/phagocytic vesicles, virus entry, exocytosis, and virus egress. In the cell nucleus, actin regulates DNA repair, chromatin remodeling and condensation, and gene transcription [4, 5]. The two main forms of actin in eukaryotic cells are monomeric globular G-actin and polymerized filamentous F-actin. Actin polymerization and G/F actin dynamics are regulated by the small GTPase RhoA and actin-interacting proteins [2, 6–14]. Because of the pleiotropy of actin function, it is not surprising that the life cycle of viruses, from the point of cell entry to the egress of the progeny, is intimately connected with the actin cytoskeleton of the host cell and that viruses hijack various actin functions and modify or redirect them to enhance self-propagation. Viruses reconfigure and manipulate the actin cytoskeleton of the host to optimize the production of progeny. It is known that RNA viruses can promote actin polymerization and reorganization [15]. For example, the coronavirus transmissible gastroenteritis virus (TGEV) and porcine hemagglutinating encephalomyelitis virus (PHEV) use Rac1/Cdc42 small GTPases of the RhoA family to induce actin polymerization [15–17]. The virus also manipulates cell adhesion complexes and modifies intercellular contacts, allowing directional cell-to-cell...
spread. The different strategies used by different viruses to manipulate the actin cytoskeleton suggest that these strategies evolved independently [18, 19]. Here, we describe strategies involving the actin cytoskeleton that are used by various viruses during entry, assembly, and egress from the host cell and point out what is already known about severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) in this regard. This review should encourage further investigation of the mechanism used by SARS-CoV-2 and eventually help in the development of novel antiviral therapeutic or prevention strategies.

**SARS-CoV-2 structure**

SARS-CoV-2 virions are spherical particles with a diameter of ~100 nm that contain a tightly packed positive-sense, single-stranded, 29,811-nucleotide-long genomic RNA that is “translation-ready”; i.e., it serves as an mRNA that can be translated directly (using the synthetic machinery of the host) to produce a set of viral proteins that are necessary for RNA replication and formation of new copies of the virus [20–24]. The RNA genome has 10–14 open reading frames (ORFs) and codes for 24–27 proteins [20, 25, 26]. Viral transcription and replication also rely on intricate long-range RNA–RNA interactions within the viral genome as well as interactions with host RNA molecules [27]. Viral RNA is surrounded by a protective coat, called a capsid, consisting of 1000 copies of the RNA-binding nucleocapsid (N) protein (Fig. 1). The N protein is highly phosphorylated, has a molecular weight of 46.5 kDa, and through the binding to a genomic packaging signal facilitates the packaging of the viral genome within the capsid [28, 29]. The capsid is surrounded by a membrane envelope, which is derived from the host cell but has viral proteins inserted into it: ~1000 copies of the membrane (M) protein, ~20 copies of the envelope (E) protein, and ~100 copies of the spike (S) protein (Fig. 1). The M protein has a molecular weight of 25–30 kDa, and it forms a dimer that shapes the viral membrane and promotes its binding to the capsid [29, 30]. The E protein has a molecular weight of 8–12 kDa and has ion channel activity. It is crucial for assembly of the virus and its release from the host cell [29]. The S proteins form the viral spikes. Each spike contains three copies of the S protein and has a length of ~10 nm [20, 29]. The S protein has a molecular weight of 150 kDa, is N-glycosylated, and belongs to the class I family of fusion proteins, which bind to receptors and mediate membrane fusion [29, 31]. The S protein consists of two subunits: S1 and S2. The S1 subunit contains the receptor-binding domain, and the S2 subunit mediates membrane fusion [32]. The S-protein-containing spikes of SARS-CoV-2 have high affinity for angiotensin-converting enzyme 2 (ACE2), which serves as a cell-surface receptor for the virus and facilitates its entry into the cell [33–35]. The binding of the virus to the host cells involves two steps: the first is docking of the virus to ACE2, and the second is activation of the spike protein by proteolytic cleavage at two sites by host-encoded proteases [36, 37]. Proteolytic cleavage itself is also a two-step process: the first cleavage occurs at the boundary of the S1 and S2 subunits in the spike protein precursor, and the second exposes the S2 subunit. The first cleavage step can occur either before or after binding to the receptor. The second cleavage can occur before endocytosis, through the activity of the serine protease TMPRSS2, or after endocytosis, mediated by endosomal proteases. The second cleavage primes the S2 subunit to induce fusion of the viral membrane with the cell membrane. In SARS-CoV-2 and Middle East respiratory syndrome coronavirus (MERS-CoV), but not SARS-CoV-1 (called SARS-CoV before the emergence of SARS-CoV-2), the boundary between S1 and S2 subunits contains a stretch of amino acids that can be recognized and cleaved by another serine protease called furin [32, 36, 37]. The S protein also has a sialic-acid-binding domain that interacts with sialic-acid-containing molecules on the surface of the host cell and facilitates virus entry [34]. This dual feature of the S protein might be one of the reasons for the high transmissibility of SARS-CoV-2 [33, 38]. Recent studies have suggested that the recruitment of SARS-CoV-2 to the cell surface is also facilitated by heparan sulfate (HS).
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proteoglycan surface molecules. HS interacts with the spike protein, increasing the virus concentration and promoting engagement with the ACE2 receptor [39]. Interestingly, the human coronaviruses HCoV-OC43 and HCoV-HKU1, which, like SARS-CoV-2, belong to the genus Betacoronavirus [40], have an additional gene encoding a hemagglutininesterase (HE) dimer protein [29]. In these viruses, the S-protein-containing spikes are interspersed among shorter ~8-nm stubby projections composed of the HE protein. The HE protein has a dual function [41]: it has a receptor-binding lectin domain that is specific for O-acetylated sialic acid (O-AcSia) and promotes attachment of the virus to sialic-acid-containing molecules on the surface of the host cell, and it has a receptor-destroying sialate-O-acetyl esterase that promotes virus release [33]. It seems that the S and HE proteins are functionally interdependent and co-evolved to optimize the balance between virus attachment and release [42]. Studies with SARS-CoV-1 have shown that the stage of binding to the receptor and virus entry requires ~10 min [20]. The process of making new virus particles within the cell lasts around 10 hours and ends with the release of ~1000 newly formed viruses [20]. Below, we summarize what happens inside the infected cell, from the moment of virus attachment to the receptor until the release of viral progeny, and how the actin cytoskeleton of the host is involved in these processes.

**Actin-driven “surfing” of the virus**

Some viruses, such as murine leukemia virus, human immunodeficiency virus, vesicular stomatitis virus, and human papillomavirus, slide (surf) on the surface of host cell protrusions called filopodia, which contain actin filaments, toward regions of the cell membrane with the highest endocytic activity [43] (Fig. 2A). Although viruses can also enter cells at other regions of the membrane, filopodium-dependent entry, due to the inherent ability of filopodia to bind and engulf various molecules, including viruses, is much more efficient [43]. Confocal time-lapse microscopy of murine leukemia virus (MLV) infecting human embryonic kidney HEK 293 cells showed that the virus, after attaching to the tip of a filopodium, moved downward toward the body of the cell [43]. The mechanism that initiates the surfing process is largely unknown, but it involves contact of the virus with...
its receptor(s) [43]. The surfing itself is powered by contraction (retrograde F-actin flow) of the filopodium’s actin filaments by myosin II motors located within the cortical actin at the base of the filopodium. This creates a dragging force that moves virus-receptor complexes to the base of the filopodium. Cytochalasin D, which blocks the barbed ends of actin filaments, inhibits surfing [18, 43]. Although there has been no description of SARS-CoV-2 surfing, high-resolution scanning microscopy has shown that SARS-CoV-2 adheres to the surface of filopodium-like projections of African green monkey kidney Vero cells, which would suggest that surfing might occur [44]. Recently, Seyran et al. [34] reported that, prior to binding to the ACE2 receptor, the SARS-CoV-2 S protein interacts (through the flat sialic-acid-binding domain in the N-terminal domain of the S1 subunit) with the sialic acid layer of the epithelium, allowing the virus to move along the epithelial surface until it encounters and interacts with the ACE2 receptor. It remains to be investigated whether this movement of the virus has anything to do with the above-described surfing mediated by retrograde F-actin flow.

**Virus entry into the cell**

Coronaviruses can enter cells using various routes, including phagocytosis, micropinocytosis, and membrane fusion [19]. However, recent studies have shown that the preferred mode of entry of SARS-CoV-2 is through binding to the ACE2 receptor and clathrin-mediated endocytosis [45], which depends on the proper arrangement of cortical actin (Fig. 3). Owczarek et al. [46] showed that both stabilization and depolymerization of actin interfere with endocytosis and entry of the common-cold-causing human coronavirus OC43. During virus entry, the virus-receptor complex recruits clathrin, which coats the invaginating membrane pit. The initiation of membrane invagination is controlled by a multiprotein complex containing the membrane-associated protein intersectin-1, which directs interactions between endocytic vesicles and the actin cytoskeleton, the epidermal growth factor receptor substrate 15 (Eps15), which is a scaffolding adaptor protein, and the membrane-sculpting F-BAR-domain-containing Fer/Cip4 homology-domain-only proteins 1 and 2 (FCHo1/2) [46]. FCHol/2 recruits intersectin-1 and Eps15 and induces membrane invagination [46, 47]. Eps15 [48] recruits adaptor protein 2 (AP2) [49], which recruits clathrin to form a lattice coating the membrane pit. The “neck” of the pit attracts the protein amphiphysin [50], which in turn recruits dynamin, which, via binding and hydrolysis of GTP, sever the “neck” of the pit from the cell membrane [51]. The scission results in the formation of a clathrin-coated endosomal vesicle containing the receptor and the virus (Fig. 3A) [46, 52].

**The endosomal phase of the virus**

The next step after the formation of the endosome containing the SARS-CoV-2-receptor complex is the acidification of the endosome through the proton-pump-dependent release of Ca$^{2+}$ from the endosome via the intracellular messenger nicotinic acid adenine dinucleotide phosphate (NAADP)-sensitive two-pore channel (TPC2) [53]. The acidification of the endosome occurs during its maturation from an early endosome to the late endosome. In the late endosome, the complex of the S protein and ACE2 receptor is cleaved by cathepsin L, which promotes the fusion of the viral and endosome membranes and the release of the viral RNA into the cytoplasm (Fig. 3A) [54]. Endosomal acidification inhibitors, such as chloroquine or bafilomycin A1, have been shown to drastically reduce RNA release and replication of SARS-CoV-2 [55]. The spatial arrangement of endosomes, their trafficking within the cell, and the recycling of endosomal membranes back to the cell surface are supported by actin filaments and microtubules and are powered by various motor proteins [56].

**Virus replication and translation**

The released viral RNA (genomic RNA, gRNA) undergoes replication, transcription, and translation to produce viral proteins and progeny RNAs [22, 57] (Fig. 3B). First, the two large open reading frames, ORF1a and ORF1b, of the viral RNA are translated into the polyproteins pp1a and pp1ab, which, after co-translational and post-translational modification, are converted into non-structural proteins (nsps), which participate in viral replication and transcription [22, 57, 58]. In the next step, viral gRNA, with the help of the nsps, generates negative-sense RNA intermediates, which are the templates for synthesis of gRNAs of the next generation and subgenomic RNAs (sgRNAs). The sgRNAs are translated to produce the viral structural proteins (N, E, M, and S), while the gRNAs are incorporated with the structural proteins into virus particles to produce new viral progeny (Fig. 3B). A recent analysis of the SARS-CoV-2 transcriptome showed that the viral RNA also has several ORFs of unknown function [22].

The replication of the viral RNA of positive-strand (+) viruses and its transcription occurs within virus-induced replication organelles situated in the vicinity of the host cell nucleus (Fig. 3B). This replication compartment, which was identified recently in cells infected with SARS-CoV-1 [59], creates a protective niche for viral replication and transcription. It is derived from the ER or other membranous structures of the host cell and consists
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of double-membrane vesicles (DMVs), double-membrane cups, called double-membrane spherules (DMSs), convoluted membranes (CMs), or/and tubules [57, 60–62]. We do not know if actin participates in the establishment and maintenance of the replication organelle. However, the involvement of the actin cytoskeleton in the formation and structure of other membranous organelles in eukaryotic cells [2] would imply that it does. It is known that actin regulates various transcription factors in eukaryotic cells and that binding to actin is necessary for the activity of RNA polymerases [5]. This implies that the transcription of viral RNA should also be actin-dependent. This assumption is supported by older studies showing that the actin cytoskeleton is involved in the replication and transcription of human parainfluenza virus type 3 [63]. Moreover, mutational analysis of respiratory syncytial virus (RSV) has shown that actin and profilin are necessary for viral transcription and that the divalent-cation-binding domain of actin is necessary for activation of viral RNA synthesis [64, 65]. Another study has shown that viruses that replicate in the nucleus interact with nuclear actin for replication, assembly, and export of mRNA or
capsids [66]. The fact that interactions with actin are necessary for many viral functions suggests that transcription of SARS-CoV-2 RNA might also be actin-dependent. However, despite that, to our knowledge, there is no direct information available on this subject.

**Virus assembly and egress**

The newly synthesized viral proteins and RNAs are transported to the ER/Golgi intermediate compartment (ERGIC, also referred to as the vesicular-tubular cluster [VTC]) [67], where the SARS-CoV-2 RNA coated with N proteins is enveloped by a host-cell membrane containing viral M, E, and S proteins (Fig. 3C). From the ERGIC, the virus enters the trans-Golgi network (TGN)/Golgi complex, where the proteins undergo posttranslational modification [68]. Some time ago, a peculiar structure called “the juxtanuclear actin ring” was described in coronavirus-infected cells (Fig. 3B). The ring, composed of actin filament layers surrounding the cell nucleus and the adjacent ER and Golgi compartments, facilitates viral genome replication and protein synthesis [19, 69]. The sequestration of the molecules needed for virus replication, transcription, and translation by the actin ring may concentrate them in the vicinity of ER and Golgi for further processing. Another function of the perinuclear actin ring was described in human adenovirus infection. In this case, the adenovirus early region 4 ORF4 protein (E4orf4) induces, via the myosin II motor/Src kinase/Cdc42/N-Wasp/RhoA kinase and Rac1 pathways, actin polymerization around the nucleus. Subsequently, the actin ring, by recruiting endosomes and disrupting vesicular trafficking and membrane dynamics, activates an endo-lysosomal-based cell death pathway [70].

The egress of newly assembled virus particles from the cell occurs via exocytosis [54, 71–73] (Fig. 3C). Recent studies have shown that SARS-CoV-2 and other betacoronaviruses use a unique exit strategy. Instead of using the conventional secretory pathway used by other viruses, they use deacidified lysosomes to egress from the cell [68]. The exact mechanism of deacidification of the lysosomes, which also deactivates their lytic enzymes, is still unknown. Ghosh et al. [68] have suggested that the deacidification of lysosomes is either a direct consequence of too much cargo (viruses) and/or a dysfunctional proton pump or that some viral proteins have a viroporin function, which, by forming hydrophilic pores, modifies membrane permeability and Ca$^{2+}$ homeostasis, leading to lysosome deacidification. It is also still unknown how the viruses are translocated from the TGN/Golgi to lysosomes. Ghosh et al. [68] propose two possible routes: One route is a transfer from the Golgi to multivesicular bodies (MVBs), which mature into lysosomes via the activity of the Rab7 GTPase pathway (Fig. 3C). The fact that the inhibition of Rab7 causes a reduction in both the formation of lysosomes and the egress of the virus supports this hypothesis [68]. Another, more complicated option is the retrograde transport of the virus from the Golgi back to the ER/ERGIC and then to lysosomes. A modification of the second option is the possible involvement of microphagy, with fragments of the ER containing the virus being engulfed directly by lysosomes [68]. Studies on the coronavirus avian infectious bronchitis virus have shown that budding of the virus from the cell membrane and its egress are actin-filament-dependent and involve interaction between actin and amino acids A159 and K160 of the viral membrane protein M [19, 74]. Using scanning electron microscopy and atomic force microscopy, it was shown that, just before its egress, SARS-CoV-1 induces the formation of pseudopodia and thickening of the cortical actin, which, by bending the membrane, expels the virus progeny from the host cell [75].

**Actin comets**

Many viruses, including vaccinia virus, Ebola virus, Marburg virus, and insect baculoviruses, use actin “tails” or so-called “actin comets” to be propelled within the cytoplasm to the cell periphery and/or to the outside of the infected cell (Fig. 2B). Viral proteins induce actin nucleation via the activity of the Arp2/3 complex. This is followed by interactions with the myosin and kinesin motors and actin polymerization/depolymerization cycles, which propel the virus. Because the comet-associated viruses were observed not only inside the cytoplasm but also in the vicinity of cell–cell junctions, it has been proposed that the propelling force of the actin comet allows the virus to pass through the actin-reinforced cellular junction and be delivered to an adjacent cell [76–78]. So far, there is no information about whether any coronaviruses, including SARS-CoV-2, use actin comets for movement inside and outside of the cell.

**Actin and direct cell-to-cell transmission of the virus**

Direct cell-to-cell transmission, which is faster and much more efficient than cell-free spread and allows the immune response, especially antibody-mediated immunity, to be circumvented, is used efficiently by many viruses, including SARS-CoV-2 [70, 78–80]. Below, we describe several possible routes of direct transmission of the virus between cells. All of these routes of transmission involve the actin cytoskeleton of the host cell.

a. **Tunneling nanotubes (TNTs)**

Tunneling nanotubes (TNTs) are long, thin (50–1500 nm in diameter), F-actin-based open-ended channels used
by neighboring cells to exchange large cargo, such as protein complexes, lipids, nucleic acids, vesicles, pathogens, and organelles (Golgi, ER, lysosomes, endosomes, and mitochondria) at the distance of hundreds of micrometers [78, 79, 81–85] (Fig. 4). Thicker TNTs, which are used for the transport of very large cargo, in addition to F-actin, also contain microtubules. The traveling speed of the cargo through the TNTs is very fast, reaching ~7 nm/s [85]. Viruses can also “surf”, docking to heparan sulfate (HS) receptors on the surface of TNTs for long distances (Fig. 4). The same receptors can also induce the fusion of the virus with the TNT membrane and induce endocytic internalization of the virus into the TNT lumen [85] (Fig. 4). Actin polymerization required for the formation of TNTs is regulated by Rho GTPase signaling, including the Rac1/Cdc42/Pak1 pathways [86]. Many viruses, such as HIV-1, influenza virus, and coronaviruses, induce the formation of TNTs to spread between cells [85, 86]. Interestingly, TNTs formed by Mycobacterium tuberculosis (which is known to enhance HIV-1 pathology) are used by the coinfected HIV for cell-to-cell transfer [86]. We do not know if SARS-CoV-2 uses TNTs to spread between cells. However, the observations that the coronaviruses murine hepatitis virus (MHV) and SARS-CoV-1 promote the formation of filopodia and membrane ruffling using the Rac1/Cdc42/Pak1 pathways and that SARS-CoV-2 is able to bind to the surface HS suggest that it can also induce the formation of TNTs. This should certainly be studied further in the context of SARS-CoV-2, as it may offer novel therapeutic targets for preventing or treating COVID-19.

b. Virological synapses

Many viruses, including the retroviruses human T cell leukemia virus type 1 (HTLV-1), human immunodeficiency virus type 1 (HIV-1), and simian immunodeficiency virus (SIV), spread from cell to cell through a virus-induced structure called the virological synapse (VS) or the infectious synapse, which shares similarities with neurological, immunological, and plant synapses [87–90]. During synapse formation, pre-synaptic and post-synaptic cells adhere to each other, but their membranes do not fuse. The virological synapse (VS) transmits viruses directly from the infected (pre-synaptic) to the uninfected (post-synaptic) cell [87]. Also, HIV-1, after being captured and internalized by dendritic cells, is transferred through the immunological synapse to interacting T cells. However, studies have indicated that not only internalized virus is transmitted but also virus particles attached to the surface of dendritic cells. This type of viral transmission is called transinfection [91].

The formation of any synapse, including the VS, involves the polarization of F-actin filaments and Rho GTPase signaling [92–95]. The actin network at the synapse not only plays a structural role but is a scaffold for signaling molecules and the clustering of virus particles [92, 93]. It seems that, at least in the case of HIV-1, the synapse contains a local, F-actin-depleted zone that facilitates virus entry by

![Fig. 4](image-url) Hypothetical TNT-dependent transfer of SARS-CoV-2. Binding of the S protein of SARS-CoV-2 to a heparan sulfate (HS) proteoglycan molecule (which acts as a co-receptor for ACE2 binding) present on the cell surface, induces the formation of a tunneling nanotube (TNT) between the infected cell and a neighboring uninfected cell. The virus is transferred within the lumen of the TNT, which contains actin filaments. Rho GTPase signaling including the Rac1/Cdc42/Pak1 pathways regulates actin polymerization, which is required for the formation of the TNT. The virus can also, after binding to HS, surf on the surface of the TNT. The virus on the surface of the TNT can also be internalized into the TNT interior. Although there are no data showing that SARS-CoV-2 spreads through TNTs, the fact that its S protein binds to HS and that the closely related SARS-CoV-1 can induce various types of cytoplasmic extensions, suggests that it does.
removing the physical cortical actin barrier [96]. Like retroviruses, coronaviruses also use the VS for direct cell-to-cell transmission. Yang et al. [97] have described VS transmis-
sion of SARS-CoV-1 between dendritic cells (DC) and target cells. Recently, additional studies have suggested that the ability of the SARS-CoV-2 to invade the nervous system involves the transmission of the virus through neurological synapses [98–100].

c. Virus interaction with the cellular junctions

A direct transfer of viruses between cells can also occur through intercellular junctions. Many viruses have evolved various strategies to alter different types of cell junctions to allow viral penetration. There are three main categories of cell junctions: Tight (occluding) junctions support the exchange of small hydrophilic molecules. Anchoring junctions (adherens junctions, desmosomes, focal adhesions, and hemidesmosomes) attach cells to each other or the extracellular matrix. Adherens junctions and focal adhesions are connected to bundles of actin filaments, while desmosomes and hemidesmosomes are connected to intermediate filaments. Gap (communicating) junctions transfer small (below 1000 Da) proteins, amino acids, nucleotides, small metabolites, ions, regulatory molecules, and secondary messengers [82, 101, 102]. Studies have shown that hepatitis C virus and retroviruses may enter through tight junctions, human papilloma virus (HPV) may enter through the adherens junctions, and HIV may modify gap junctions for entry [103]. Retroviruses modify the distribution of the tight-junction proteins occludin and claudin-1 and destroy tight junctions via the RhoA/ROCK/MLC signaling pathways [103–105]. Hepatitis C virus also affects occludin and claudin-1, and these proteins may also promote endocytosis of the virus [103, 106, 107]. HPV enters through adherens junctions. It modifies the distribution of adherens junction proteins such as β-catenin, disrupting their association with E-cadherin and the actin cytoskeleton [103, 108]. HIV hijacks gap junctions to spread apoptosis-inducing factors to uninfected cells and destroys, via the mitogen-activated protein kinase (MAPK) pathway, adherens and tight junctions [103, 109]. Studies have shown that the intermediate filament protein vimentin, connected to adherence junctions, is essential for several phases of the viral life cycle, including entry, transport, replication, assembly, and egress [110]. Intermediate filaments, actin, and microtubules are also crucial for the entry and propagation of dengue virus, Zika virus, and West Nile virus [111]. Recent studies have shown that alterations in cell junctions induced by SARS-CoV-2 are highly damag-
ing for patients [112, 113]. Humans have over 100 proteins containing PDZ (post-synaptic density protein) domains, some of which are components of cell junctions. The enve-
lope (E) proteins of SARS-CoV-2 and SARS-CoV-1 have a PDZ-binding motif (PBM). The binding of the viral PBM to PALS1 (protein associated with Lin seven 1) present in human cell junctions relocates PALS1 from the junctions to the ERGIC compartment, where the virus is assembled. The envelope protein also binds and dislocates other PDZ-containing junctional proteins, including the adherens junction protein syntenin and the tight junction ZO-113 (zonula occludens-113) protein, which regulates the actin-relevant RhoA/ROCK pathway [114]. Disruption of the junctional proteins in the inter-epithelial junctions of the lungs causes vascular leakage, damages the alveoli, and induces acute respiratory distress syndrome (ARDS) in COVID-19 patients [112, 115].

Virus-induced syncytia

Pathological examination of the lungs of the patients with severe cases of COVID-19 infection shows the presence of SARS-CoV-2-infected, multinucleated syncytia of pneumocytes [116, 117]. These patients are also often severely lymphopenic; i.e., they have a deficit of lymphocytes, which decreases their ability to fight the infection [118, 119]. It is known that many viruses, including poxviruses, paramyxoviruses, herpesviruses, and retroviruses, induce the fusion of infected cells into multinuclear syncytia, which eventually die [120]. Recent studies have shown that SARS-CoV-2 infection causes the formation of giant syncytia and the destruction of tight junctions in the lung airway epithelium [113]. There are two main types of syncytia: (1) homotypic syncytia consisting of cells of the same type and (2) heterotypic syncytia, which result from the fusion of cells of different types. Recently, Zhang et al. [119] described another type of syncytia in the lungs of COVID-19 patients. These syncytia consist of lung cells containing internalized lymphocytes (Fig. 5). The cell fusion leading to the formation of syncytia requires the rupture of the membranes of the fusing partners, followed by membrane rejoining. The whole process is facilitated by fusogenic proteins and involves the restructuring of the supportive platform of the actin cytoskeleton underlying the cell membranes. The spike protein of SARS-CoV-2 is a fusogen with homology to mammalian syncytin-1, which is involved in the formation of the syncytial layers of the placent [31]. The model and the details of the fusion process are described in references [31, 119]. Virus-induced syncytia not only enhance virus spread but may also serve as a tool for the elimination of immune cells that fight the virus. Studies of COVID-19 patients showed that SARS-CoV-2 induces both homotypic syncytia of pneumocytes, and heterotypic cell structures containing fused lung cells with the engulfed lymphocytes inside [36, 119]. These studies suggest that
SARS-CoV-2-infected cells express the viral S (spike) protein on their surface. The interaction of the spike protein with the ACE2 receptor of the neighboring cell activates the calcium-activated lipid scramblase TMEM16F (anoctamin 6), which exposes phosphatidylserine on the cell surface and triggers the formation of a syncytium, which subsequently engulfs the immune cells [116, 122]. These studies suggest that syncytium-mediated lymphocyte internalization and elimination may be one of the causes of lymphopenia in COVID-19 patients [119–121]. Recent studies indicate the antihelminthic drug niclosamide suppresses the activity of TMEM16F and inhibits spike-induced syncytium formation by SARS-CoV-2 [123].

**Conclusions**

The information presented here indicates that interactions of viruses, including SARS-CoV-2, with the actin cytoskeleton and actin-related cell functions and signaling pathways, are of crucial importance for viral functions, infectivity, and pathogenicity. It is also clear that our knowledge about the interactions of SARS-CoV-2 with actin and other components of the cytoskeleton is still very limited and requires further studies. A better understanding of these interactions and processes and the participating viral and cellular components involved can help in the design of novel antiviral therapies and/or strategies to mitigate the severity of COVID-19 and its damaging, long-term effects. The conventional approaches to fighting viral infections include vaccinations, antiviral drugs, and preventing overactivation of the immune response, the last of which is especially important in COVID-19, in which the overproduction of inflammatory factors can cause a cytokine storm in the patient’s lungs [36]. A recent analysis of gene expression profiles in the SARS-CoV-1/SARS-CoV-2 infections using Gene Ontology (GO) and MetaCore showed that the upregulation of actin/cytoskeleton remodeling via the RhoA pathway is crucial for infection [124]. Thus, we propose that interference with actin-dependent functions of the virus and/or host immune cells using RhoA pathway inhibitors, such as the multiple sclerosis drugs fingolimod and siponimod, might be a novel therapeutic approach to fighting viral diseases [125].

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Declarations

Conflict of interest None of the authors has a conflict of interest.

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