The Endoplasmic Reticulum–Plasma Membrane Junction: A Hub for Agonist Regulation of Ca^{2+} Entry

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Stimulation of cell-surface receptors induces cytosolic Ca^{2+} ($[\text{Ca}^{2+}]_i$) increases that are detected and transduced by effector proteins for regulation of cell function. Intracellular Ca^{2+} release, via endoplasmic reticulum (ER) proteins inositol 1,4,5-trisphosphate receptors (IP_3R) and ryanodine receptors (RyR), and Ca^{2+} influx, via store-operated Ca^{2+} entry (SOCE), contribute to the increase in $[\text{Ca}^{2+}]_i$. The amplitude, frequency, and spatial characteristics of the $[\text{Ca}^{2+}]_i$ increases are controlled by the compartmentalization of proteins into signaling complexes such as receptor-signaling complexes and SOCE complexes. Both complexes include protein and lipid components, located in the plasma membrane (PM) and ER. Receptor signaling initiates in the PM via phospholipase C (PLC)-mediated hydrolysis of phosphatidylinositol 4,5-bisphosphate (PIP_2), and culminates with the activation of IP_3R in the ER. Conversely, SOCE is initiated in the ER by Ca^{2+} -sensing stromal interaction molecule (STIM) proteins, which then interact with PM channels Orai1 and TRPC1 to activate Ca^{2+} entry. This review will address how ER–PM junctions serve a central role in agonist regulation of SOCE.

Calcium (Ca^{2+}) signaling mechanisms within a cell are tightly controlled to ensure generation of $[\text{Ca}^{2+}]_i$ signals with appropriate magnitude as well as spatial and temporal characteristics that are required for regulating specific physiological functions. $[\text{Ca}^{2+}]_i$ signals usually consist of an increase in cytosolic Ca^{2+} concentration, which is detected and transduced by effector proteins (Berridge 2006). In nonexcitable cells, such increases in $[\text{Ca}^{2+}]_i$ result from the stimulation of cell-surface receptors, primarily G-protein-coupled (GPCR) or tyrosine-kinase-coupled receptors, which lead to activa-

tion of phospholipase C (PLC) and hydrolysis of phosphatidylinositol 4,5-bisphosphate (PIP_2). This is a critical initial step in the Ca^{2+} -signaling cascade and results in the generation of two intracellular messengers: inositol 1,4,5-trisphosphate (IP_3) and diacyl glycerol (DAG). DAG is retained in the plasma membrane (PM) where it modulates a number of cellular functions such as regulation of ion channels and enzyme activation. IP_3 , a soluble metabolite, diffuses in the cytosol and binds to its receptor, the IP_3R , which is a Ca^{2+} channel localized in the endoplasmic reticulum (ER) membrane. The resulting release

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of Ca^{2+} from the ER lumen leads to an increase in $[\text{Ca}^{2+}]_i$, which is used to control many cellular functions, including ion channel regulation and gene expression (Mikoshiha 2015). Release of Ca^{2+} from the ER also causes a decrease in Ca^{2+} concentration within the ER lumen ($[\text{Ca}^{2+}]_{\text{ER}}$) and, depending on the magnitude, a substantial and sustained decrease in $[\text{Ca}^{2+}]_{\text{ER}}$ can be deleterious to cells, affecting processes such as protein synthesis and cell proliferation (Mekahli et al. 2011). The ER- Ca^{2+} pool is finite, and maintaining long-term signaling requires refilling of the ER- Ca^{2+} stores via Ca^{2+} influx from the extracellular medium. In the absence of Ca^{2+} influx, the release of Ca^{2+} from the ER would be transient in nature because of progressive run-down of $[\text{Ca}^{2+}]_{\text{ER}}$ and the lack of ER- Ca^{2+} store refilling by incoming extracellular Ca^{2+} . This review will discuss current concepts pertaining to IP_3 -induced Ca^{2+} signaling and the store-operated Ca^{2+} entry (SOCE) pathway (Putney 1990, 2017).

SOCE is mediated by PM Ca^{2+} channels activated in response to decreases in $[\text{Ca}^{2+}]_{\text{ER}}$ and inactivated by refilling of the ER- Ca^{2+} stores (Ambudkar 2006; Ambudkar et al. 2006; Putney 2017). The close coupling between SOCE and Ca^{2+} uptake into the ER suggests the formation of cellular microdomains, in which SOCE components in the PM are in close proximity to ER- Ca^{2+} -signaling components, such as sarco/ER Ca^{2+} ATPase (SERCA) and IP_3 Rs. Many studies have now established that SOCE is regulated within ER-PM junctions that are formed by interactions of ER-resident proteins with the PM lipid or protein components. The PM protein Orai1 was identified as a critical component of SOCE as it forms the pore of the highly Ca^{2+} -selective channel that is activated by ER- Ca^{2+} store depletion. This channel carries the Ca^{2+} release-activated Ca^{2+} current (I_{CRAC} ; Feske et al. 2006; Prakriya et al. 2006; Vig et al. 2006; Zhang et al. 2006; Hou et al. 2012; Yen et al. 2016). Orai1 has four transmembrane-spanning domains and assembles to form a hexameric channel that is activated by direct interaction with the ER proteins called STIMs, STIM1 and STIM2. Both STIM isoforms sense the decrease in $[\text{Ca}^{2+}]_{\text{ER}}$ via amino-terminal

Ca^{2+} -binding domains, known as EF-hand domains, that are located in the ER lumen. Release of Ca^{2+} from the EF-hand domain triggers conformational changes to enable STIM1 or STIM2 protein clustering at the cell periphery where they recruit Orai1. The STIM carboxy-terminal tail mediates its binding to PM PIP_2 , whereas the cytosolic STIM-Orai1 activation region (SOAR; also known as CRAC activation domain [CAD]) interacts with and gates Orai1 (Liou et al. 2005; Roos et al. 2005; Brandman et al. 2007; Park et al. 2009; Yuan et al. 2009b; Bhardwaj et al. 2013). The STIM-Orai1 interaction, together with STIM1 scaffolding at the PM, culminates in the formation of ER-PM junctions that mark the sites of Ca^{2+} entry into the cell. Activation of clustered Orai1 channels generates higher $[\text{Ca}^{2+}]_i$ near the cytosolic mouth of the channels (referred to as local increase in $[\text{Ca}^{2+}]_i$) than when the channels are diffusely spread within the PM (Samanta et al. 2015). Both local and global $[\text{Ca}^{2+}]_i$ changes are used by cells to regulate distinct functions (Kar et al. 2011; Ong et al. 2012; Kar and Parekh 2015).

Transient receptor potential canonical (TRPC) channels, suggested as candidates for SOCE channels before the discovery of STIM and Orai, are also activated in response to receptor-stimulated PM PIP_2 hydrolysis, and contribute to agonist-induced $[\text{Ca}^{2+}]_i$ increases (Ambudkar et al. 2006; Abramowitz et al. 2007; Yuan et al. 2009a). TRPC channels regulate important physiological functions in several tissues such as exocrine glands, endothelium, cardiac muscle, neurons, and kidney cells (Ong et al. 2014). TRPC1, TRPC3, and TRPC4 have been shown to mediate Ca^{2+} entry following PM receptor stimulation and/or ER- Ca^{2+} store depletion. TRPC channel association with the Orai1/STIM1 complex is shown by the interaction of TRPC1 and TRPC3 with STIM1 (Yuan et al. 2007; Zeng et al. 2008). The localization of TRPC1 with Orai1/STIM1 in ER-PM junctions is a critical determinant of its activation (Pani et al. 2009; Cheng et al. 2011; de Souza et al. 2015). The polybasic domain of STIM1, specifically $^{684}\text{KK}^{685}$, which is distinct from the SOAR/CAD domain, activates TRPC1 by binding to $^{639}\text{DD}^{640}$ (Huang et al. 2006; Zeng et al.

2008). In cells expressing both TRPC1 and Orai1, both channels are activated by ER- Ca^{2+} depletion. Further, TRPC1-mediated Ca^{2+} entry enhances the $[\text{Ca}^{2+}]_i$ increase induced by Orai1 and contributes to sustained elevation of $[\text{Ca}^{2+}]_i$ (Ong et al. 2007). Importantly, Ca^{2+} signals generated by TRPC1 are used by cells for regulating functions distinct from those activated by Orai1; for example, NFAT is regulated by Ca^{2+} entry mediated by Orai1, whereas NF- κ B regulation involves TRPC1. Local $[\text{Ca}^{2+}]_i$ increase mediated by Orai1 is exclusively used for NFAT1 activation. The increase in $[\text{Ca}^{2+}]_i$ detected by calmodulin leads to activation of calcineurin, dephosphorylation of NFAT1, and translocation of the transcription factor to the nucleus. Global Ca^{2+} signals resulting from TRPC1 function are detected by $[\text{Ca}^{2+}]_i$ sensors, such as calmodulin kinase II, to cause NF- κ B activation. Additionally, in exocrine gland cells, TRPC1-mediated Ca^{2+} entry is required for regulation of Ca^{2+} -activated Cl^- and K^+ channel activities (Cheng et al. 2011; Hong et al. 2011; Ong et al. 2012; Sun et al. 2015).

The nature of SOCE-associated ER-PM junctions, their molecular components, and architecture, as well as their physiological relevance, are currently the subject of many studies in the field of Ca^{2+} signaling. Emerging data suggest that these junctions are not only involved in sensing agonist stimulation and regulating SOCE but also in compartmentalization of specific downstream targets and effector proteins that drive SOCE-dependent control of cell function. In the later sections of this article, we will discuss the role of ER-PM junctions as signaling hubs that coordinate agonist regulation of PIP_2 hydrolysis with depletion of ER- Ca^{2+} store and activation of SOCE. Precise spatial and temporal coordination of these processes generates specific Ca^{2+} signals that regulate different cell functions.

AGONIST-DEPENDENT GENERATION OF $[\text{Ca}^{2+}]_i$ SIGNALS

Signaling Complexes and ER-PM Junctions

Two major types of stimuli that trigger $[\text{Ca}^{2+}]_i$ increases in nonexcitable cells are those acting on

GPCRs (e.g., muscarinic and α 1-adrenergic receptors) and those acting on tyrosine kinase-coupled receptors (e.g., growth factor receptors and T-cell receptor [TCR]; Berridge 2017). Activation of either class of PM receptors leads to the assembly of a receptor-associated signaling complex that includes G proteins, PLCs, and downstream targets such as adenylyl cyclases, RyR, and IP_3R . A family of proteins that function as regulators of G protein signaling (RGS) has been suggested to serve as scaffolds in the assembly of such signaling complexes. Additionally, other scaffolding proteins (e.g., HOMER and A-kinase anchoring proteins [AKAPs]) and PM lipid domains have also been implicated in this process. Such association between proteins of the ER and PM results in the formation of ER-PM junctions. An important function of these junctions, which contain PM receptor-signaling proteins and the IP_3R , ensures that IP_3 is effectively delivered from the site of its synthesis in the PM to its receptor in the ER membrane with minimal diffusion or hydrolysis. Targeted delivery of IP_3 increases the response speed and organizes the Ca^{2+} signals in spatially segregated areas within the cell (Delmas et al. 2002). Interaction of IP_3Rs with other proteins, such as PLCs, receptor of activation protein C kinase 1 (RACK1), AKAPs, or HOMER, may bring them close to the PM receptor-signaling complex to facilitate IP_3 delivery in a selective and rapid manner. Notably, some of these proteins also interact with TRPC channels and Orai1 (Ong et al. 2014; Prole and Taylor 2016). Thus, Ca^{2+} signaling is organized to be rapid and specific through compartmentalization of signaling proteins within privileged ER-PM junctions that are assembled to initiate the PLC cascade and IP_3 -induced Ca^{2+} release from the ER (Thillaiappan et al. 2019).

TRPC channels reside in PM lipid raft domains (LRDs), which are also populated by a number of receptor-coupled signaling proteins (Ong et al. 2014). These channels assemble and form signaling complexes with PLC, IP_3R , G proteins, and scaffolding proteins such as RACK1 (TRPC3) and HOMER (TRPC1; Yuan et al. 2003; Bandyopadhyay et al. 2008; Ong et al. 2014). Taken together, these findings showed

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that PM Ca^{2+} influx channels, activated in response to PIP_2 hydrolysis, are recruited into the same junctions as the signaling complex that causes hydrolysis of PIP_2 . Importantly, biochemical and total internal reflection microscopy (TIRFM) data show interactions between TRPC1 and TRPC3 with STIM1 and Orai1 (Ong et al. 2007; Yuan et al. 2007; Cheng et al. 2008, 2011; Zeng et al. 2008). These findings place Orai1/STIM1 together with TRPC channel within close proximity to the receptor-signaling complexes and IP_3 Rs. Further confirmation is provided by a recent study showing that native IP_3 Rs are preclustered in cellular areas in which STIM1 accumulates after store depletion (Thilalaiappan et al. 2017). It has been suggested that this allows STIM1 to sense decreases in $[\text{Ca}^{2+}]_{\text{ER}}$ near the IP_3 Rs, and rapidly respond to activate

Orai1 within the same microdomain. This would also lead to regulation of TRPC1 function (further discussed below), which is directed to regions where Orai1/STIM1 cluster via a fast endocytic recycling pathway (de Souza et al. 2015) or interaction with PM caveolin-1 ([Cav-1]; Brazer et al. 2003; Pani et al. 2009). It is not clear whether the receptor and SOCE-signaling complex involve the formation of smaller “sub” or “nano” domains within a larger ER–PM junction or whether they are both segregated within a single large domain (summarized in the model shown in Fig. 1). Research in the field of lipid rafts has suggested that smaller rafts containing different components can undergo fusion to allow proteins segregated in two different domains to be incorporated into one domain, enabling protein–protein interactions to

Figure 1. Organization of endoplasmic reticulum–plasma membrane (ER–PM) junctions involved in agonist-regulated store-operated Ca^{2+} entry (SOCE). Binding of an agonist/ligand to its G-protein-coupled receptor in the PM triggers a signaling cascade that results in phospholipase C (PLC)-mediated hydrolysis of phosphatidylinositol 4,5-bisphosphate (PIP_2) in the PM and generation of the second messenger inositol 1,4,5-trisphosphate (IP_3). When IP_3 binds to its receptor (IP_3R) in the ER membrane, it causes release of Ca^{2+} from the ER- Ca^{2+} stores. Alternatively, the second messenger NAADP can activate the ryanodine receptors (RyRs) to cause Ca^{2+} release from the ER. The resulting decrease in Ca^{2+} concentration within the ER lumen ($[\text{Ca}^{2+}]_{\text{ER}}$) is sensed by stromal interaction molecules STIM1 and STIM2, which aggregate and translocate to the cell periphery to activate Orai1. Ca^{2+} entry via Orai1 raises the local $[\text{Ca}^{2+}]$ (denoted by boxed $[\text{Ca}^{2+}]_{\text{i}}$) near the channel pore, which is required for the trafficking of TRPC1 to and its insertion into the PM, as well as activation of NFAT signaling (not shown). Following channel insertion into the PM, TRPC1 is activated by STIM1. At the moment, it has not been shown that STIM2 also activates transient receptor potential canonical 1 (TRPC1). The Ca^{2+} entry via TRPC1 contributes to a global increase in $[\text{Ca}^{2+}]_{\text{i}}$, which is required for other downstream Ca^{2+} -dependent processes such as NF- κ B activation and fluid secretion in salivary glands. Additionally, the sarco/ER Ca^{2+} -ATPase (SERCA) and PM Ca^{2+} -ATPase (PMCA) pumps also modulate the $[\text{Ca}^{2+}]_{\text{i}}$ by pumping Ca^{2+} into the ER or outside of the cell, respectively. Assembly of the proteins within these junctions coordinate PM–ER signaling, which then induces ER–PM signaling. Precise attunement of the functional and physical interactions between ER and PM proteins is critical for the cell.

occur in a regulated manner (Ong and Ambudkar 2015). Such a mechanism would bring agonist-signaling components within close proximity to SOCE components to facilitate interactions between them. There are no data to directly show this. However, it has been reported that the association of Orai1/STIM1 complexes with PM lipid domains may be altered following stimulation of Ca^{2+} entry (Maleth et al. 2014). Additionally, remodeling of ER-PM junctions has also been reported (further discussed below), which could reorganize the signaling complex and modulate protein function.

Stimulus Intensities and Activation of SOCE

Agonist-induced, IP_3 -dependent release of Ca^{2+} from the ER causes $[\text{Ca}^{2+}]_i$ elevation, the pattern and amplitude of which varies depending on the cell type and stimulus intensity. Low agonist concentrations typically trigger an oscillatory pattern of $[\text{Ca}^{2+}]_i$ increase. Intermediary agonist concentrations induce either more frequent Ca^{2+} oscillations or Ca^{2+} oscillations arising from an elevated baseline. High agonist concentrations can cause a sustained elevation of $[\text{Ca}^{2+}]_i$ (Bird and Putney 2005; Putney and Bird 2008; Ong et al. 2015). Both intracellular Ca^{2+} release via IP_3 Rs and extracellular Ca^{2+} influx via SOCE or other Ca^{2+} channels contribute to the elevations in $[\text{Ca}^{2+}]_i$. Several Ca^{2+} flux mechanisms in the cell can contribute to Ca^{2+} oscillations. In some cases, positive feedback regulation of PLC by Ca^{2+} causes IP_3 levels to oscillate, which can drive periodic release of Ca^{2+} from the ER (Kawabata et al. 1996; Nash et al. 2001). However, Ca^{2+} oscillations have also been detected in cells with stable IP_3 levels (Dupont et al. 2011). Importantly, $[\text{Ca}^{2+}]_i$ oscillations in many cell types can be supported for a short time period in the absence of external Ca^{2+} because of repetitive Ca^{2+} release and reuptake into ER. Such $[\text{Ca}^{2+}]_i$ oscillations are primarily controlled by regulation of IP_3 R function by IP_3 and $[\text{Ca}^{2+}]_i$. The Ca^{2+} -dependent activation of IP_3 Rs, followed by their Ca^{2+} -dependent inhibition, can fully explain Ca^{2+} oscillations in most cells (Iino 1990; Bezprozvanny et al. 1991). In the absence of Ca^{2+} influx, there is an incremental loss of Ca^{2+} con-

tent in the ER during the oscillations as a result of the efflux of Ca^{2+} from the cell by the PM Ca^{2+} pump PMCA. Sustained Ca^{2+} oscillations or elevation in $[\text{Ca}^{2+}]_i$, which are critical for regulating most cellular functions, are dependent on Ca^{2+} entry, primarily via the SOCE mechanism (Parekh 2011; Thillaiappan et al. 2019).

SOCE modifies the pattern of Ca^{2+} oscillations by increasing the frequency and/or amplitude of the Ca^{2+} signal with a more sustained elevation of $[\text{Ca}^{2+}]_i$ being generated at higher stimulus intensities. It is important to consider the status of ER- Ca^{2+} stores at different stimulus intensities. At low agonist concentrations, as a result of less IP_3 generation and oscillatory $[\text{Ca}^{2+}]_i$, there is less depletion of ER- Ca^{2+} stores. Conversely, a higher agonist concentration produces a greater amount of IP_3 and substantial depletion of $[\text{Ca}^{2+}]_{\text{ER}}$. Thus, SOCE modulation requires differential Ca^{2+} sensing of ER- Ca^{2+} stores across the range of $[\text{Ca}^{2+}]_{\text{ER}}$ that result from agonist stimulation of cells. Although STIM1 and STIM2 share many properties, they have key differences that are critical in determining agonist regulation of SOCE. The different Ca^{2+} affinities of their EF-hand domains allow the proteins to differentially respond at high and low $[\text{Ca}^{2+}]_{\text{ER}}$, and regulate the response of SOCE to different stimulus intensities (Liou et al. 2005; Brandman et al. 2007; Carrasco and Meyer 2011; Ong et al. 2015; Subedi et al. 2018). STIM1 has a relatively high affinity for Ca^{2+} (about 200 μM) and thus can respond only when $[\text{Ca}^{2+}]_{\text{ER}}$ is substantially decreased. Such STIM1-activating $[\text{Ca}^{2+}]_{\text{ER}}$ decreases are typically achieved only with high agonist concentrations, or when SERCA function is compromised in cells. On the other hand, STIM2 has lower affinity for Ca^{2+} (around 400 μM) and can sense small changes in $[\text{Ca}^{2+}]_{\text{ER}}$ (note that ambient $[\text{Ca}^{2+}]_{\text{ER}}$ is around 600–800 μM). Thus, STIM2 is well placed to respond to low agonist concentrations that lead to small decreases in $[\text{Ca}^{2+}]_{\text{ER}}$, such as those observed in cells with $[\text{Ca}^{2+}]_i$ oscillations. Together, STIM2 and STIM1 can regulate activation of SOCE across a wide range of agonist concentrations and consequent $[\text{Ca}^{2+}]_{\text{ER}}$ depletion levels. Indeed, previous findings have shown that STIM2

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tunes the sensitivity of SOCE activation at low agonist concentration (Ong et al. 2015). STIM2 has been shown to be important in maintaining both basal Ca^{2+} entry and Ca^{2+} influx in response to weak levels of stimulation (Brandman et al. 2007). Moreover, STIM1 is much more effective than STIM2 at gating Orai1, resulting in more Ca^{2+} entry via the channel, which is critical for cell function. STIM2 causes relatively weak Orai1 channel activation, which leads to less $[\text{Ca}^{2+}]_i$ increase. $[\text{Ca}^{2+}]_i$ signals generated by Orai1/STIM1 or Orai1/STIM2 complexes have been reported to cause activation of distinct cellular functions in mast cells (Kar et al. 2012; Thiel et al. 2013). Despite the contribution of STIM2 in Orai1 function, STIM1 has been established as the critical regulatory component for Orai1 activation. SOCE and SOCE-dependent cell function, at both low and high agonist concentration, are almost completely eliminated in cells lacking STIM1 (Luik et al. 2008; Oh-Hora et al. 2013; Ong et al. 2015; Subedi et al. 2018). A major question arising from these findings is how does STIM1 activate Orai1 at $[\text{Ca}^{2+}]_{\text{ER}}$ that is not low enough to trigger its activation. Indeed, TIRFM shows that STIM1 does not respond to low [agonist] (Ong et al. 2015).

Recent findings have revealed a novel function for STIM2 in regulating SOCE. STIM2 mediates recruitment of STIM1 to ER-PM junctions, facilitating assembly of STIM1/Orai1 at low agonist concentrations (Ong et al. 2015; Subedi et al. 2018). Further, interaction of STIM1 with STIM2 causes a conformational change in the former that leads to STIM1-activating Orai1 even when $[\text{Ca}^{2+}]_{\text{ER}}$ is not low enough to elicit a STIM1 response (Subedi et al. 2018). This scaffolding role for STIM2 appears to be critical for SOCE-dependent regulation of cell functions, such as cytokine synthesis in stimulated T lymphocytes, which is dependent on the assembly of Orai1/STIM1 complexes and strong Orai1 activation (Oh-Hora et al. 2008; Diercks et al. 2018). The relatively stronger PIP_2 -binding domain in STIM2 is a key determinant of its ability to cluster and generate ER-PM junctions in response to low level stimuli (Bhardwaj et al. 2013). Importantly, the carbox-

yl terminus of STIM2 has been shown to be in a more open conformation that allows the protein to be in an activated state to recruit and activate Orai1 in cells with minimal depletion of ER- Ca^{2+} stores (Subedi et al. 2018; Zheng et al. 2018). This ability of STIM2 to respond to small depletions of ER- Ca^{2+} stores and recruit Orai1/STIM1 under conditions in which STIM1 cannot respond on its own plays a critical role in the regulation of cell function.

REGULATION OF TRPC CHANNELS IN ER-PM JUNCTIONS

All TRPC channels are activated in response to stimulation of PIP_2 hydrolysis by agonists. Some of them are activated by agonist-induced depletion of ER- Ca^{2+} stores (Ambudkar et al. 2006; Yuan et al. 2009a; Ong et al. 2014), whereas others are dependent on metabolites arising from receptor-stimulated PIP_2 hydrolysis (Hurst et al. 1998; Liu et al. 2000; Strübing et al. 2001). The available data are the most consistent and strongest in support of a role for TRPC1 in SOCE. Nonetheless, a few studies have shown TRPC3 and TRPC4 to be activated by store depletion. Importantly, TRPC1 and TRPC3 are dependent on Orai1 and STIM1 for their function, as loss of either of these latter components eliminates TRPC-mediated Ca^{2+} entry. In contrast, knockdown of TRPC channels only causes partial reduction of Ca^{2+} entry. However, loss of TRPC function is reflected by a change in the phenotype of the current induced by store depletion. In cells expressing both TRPC and Orai channels, the current is typically nonselective, referred to as store-operated calcium current (I_{SOC}). Loss of TRPC channels results in attenuation of the I_{SOC} current. Instead, I_{CRAC} carried by the residual Orai1 channel can be detected (Cheng et al. 2011). In cells containing both Orai1 and TRPC1, both channels contribute to $[\text{Ca}^{2+}]_i$ increase. Nevertheless, Ca^{2+} entry mediated by each channel is used by cells to regulate distinct functions.

Several studies have shown that TRPC1 and Orai1 cocluster and interact with STIM1 within ER-PM junctions. STIM1 is required for clustering, as well as activation, of the channels.

LRDs are enriched in cholesterol and sphingolipids, and have been proposed to provide a stable platform for the assembly of Ca^{2+} -signaling complexes (Pani and Singh 2009; Lingwood and Simons 2010; Ong and Ambudkar 2012). The structural integrity of LRD appears to be vital for SOCE as disruption, using methyl- β -cyclodextrin to sequester cholesterol, has been shown to attenuate SOCE in many cell types. Moreover, LRDs in the PM are involved in localizing TRPC1 channels to the ER–PM junctions (Lockwich et al. 2000; Brownlow et al. 2004; Kannan et al. 2007; Jardin et al. 2008; Pani et al. 2008; Bomben et al. 2011). Disruption of LRD causes impairment of TRPC1 assembly with Orai1, STIM1, and the type 2 isoform of the inositol 1,4,5-triphosphate receptor ($[\text{IP}_3\text{R}2]$; Brownlow and Sage 2003; Brownlow et al. 2004; Pani et al. 2008; Galan et al. 2010). TRPC1 and STIM1 partition into, and interact within, LRDs following store depletion (Lockwich et al. 2000; Berthier et al. 2004; Kannan et al. 2007; Pani et al. 2008; Formigli et al. 2009). Recruitment of TRPC1 into LRDs is mediated by Cav-1, whereas STIM1 is likely to interact directly with the PM phospholipids via its carboxy-terminal polybasic domain. Subsequent studies have shown that the Orai1/STIM1 complexes are also recruited into a microdomain that is PIP_2 -rich and contains Cav-1 (Jha et al. 2013; Maleth et al. 2014). Cav-1 has been also shown to regulate Orai1 function and trafficking (Yu et al. 2010; Yeh and Parekh 2015). These studies further highlight the importance of specialized PM domains in the assembly and regulation of SOCE channels.

Several studies have addressed the critical functional interaction between Orai1 and TRPC1. Strong evidence supports the suggestion that Ca^{2+} entry via Orai1 triggers recruitment of TRPC1 to the PM (Cheng et al. 2011). Thus, coordinated regulation of TRPC1 surface expression by Orai1 and gating by STIM1 provides a mechanism for rapidly modulating and amplifying Orai1-generated Ca^{2+} signals (Ong et al. 2012). By recruiting ion channels (e.g., TRPC channels) and other signaling components (e.g., adenylyl cyclase and NFAT-signaling complexes), Orai1 and STIM1 concertedly

impact a variety of critical cell functions. The specific effector proteins that are involved in sensing the respective Ca^{2+} signals for regulating cell function are not yet known for TRPC1. Although AKAP79 provides a scaffold to coordinate Orai1-mediated NFAT activation (Kar et al. 2014), there are no data to show a role for AKAP79 in TRPC1-mediated regulation of cell function. Whereas other scaffolds, such as RACK1, Homer, and NHERF (Na^+/H^+ exchange regulatory factor), have been associated with TRPC proteins, it remains to be seen whether these represent signaling complexes in different ER–PM junctions (Ong et al. 2014).

ER- Ca^{2+} STORE RELEASE—A CRITICAL LINK BETWEEN RECEPTOR SIGNALING AND SOCE

IP_3Rs

IP_3 links receptor-stimulated, PLC-mediated, hydrolysis of PIP_2 in the PM with ER- Ca^{2+} release via the IP_3R . This critical step, leading to a decrease in $[\text{Ca}^{2+}]_{\text{ER}}$, is also the trigger for STIM/Orai1 clustering, TRPC recruitment, and activation of SOCE. Thus, signals are transmitted from the PM to the ER and back to the PM (denoted in the model in Fig. 1). The generation and concentration of IP_3 , as well as the location and activity of IP_3R , are critical factors in the coordinated regulation of SOCE. IP_3R function is exquisitely regulated by a number of different cellular factors. Although gating is primarily induced by IP_3 , Ca^{2+} exerts an important biphasic effect on IP_3R , which enables the channel to be activated when $[\text{Ca}^{2+}]_i$ is low but inhibited when $[\text{Ca}^{2+}]_i$ is elevated above a certain threshold (Iino 1990; Bezprozvanny et al. 1991; Thillaiappan et al. 2019). IP_3Rs are also regulated by cAMP-dependent protein kinase A (PKA). IP_3R phosphorylation enhances IP_3 binding affinity (Taylor 2017). Because Ca^{2+} entering via SOCE regulates Ca^{2+} -dependent adenylyl cyclases within ER–PM junctions, IP_3R in the vicinity of ER–PM junctions could be sensitized by cAMP to respond to low IP_3 concentrations. Thus, there is a feedforward regulation of IP_3R by SOCE via generation of cAMP

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and activation of PKA (Thillaiappan et al. 2019). Conversely, adenylyl cyclases that are inhibited by Ca^{2+} will be suppressed by SOCE, causing a decrease in ambient cAMP concentration and dampening of IP_3R function (Willoughby et al. 2014).

It is important to note that although IP_3Rs are localized throughout the ER, the distribution is not uniform. The location of IP_3R in the cell determines the spatial characteristics of IP_3 -generated Ca^{2+} signals, and also likely defines the location of other signaling proteins that are regulated by the Ca^{2+} released from the ER. In polarized cells like exocrine gland acinar cells, IP_3Rs are enriched in the apical region where the Ca^{2+} signal initiates following stimulation of cells (Yule et al. 1997; Lee et al. 2006; Petersen and Tepikin 2008). Importantly, STIM1, Orai1, and TRPC channels also overlap, at least partially, with IP_3R in this region, suggesting close association between IP_3Rs and the SOCE-signaling components in the PM (Hong et al. 2011). It has been reported that there are two populations of IP_3R : mobile (~75%) and immobile receptors. The latter group are found in discrete clusters at the ER membrane localized immediately beneath the PM. Interestingly, the small population of immobile IP_3R clusters represent the sites of Ca^{2+} release induced by agonists, despite the population of mobile receptors. Further, following ER- Ca^{2+} release, STIM1 accumulates at ER-PM junctions that are located adjacent to immobile IP_3Rs . These findings proposed that the immobile IP_3R clusters are “licensed” receptors positioned to sense IP_3 in the vicinity of the receptor-signaling complex in which it is generated. STIM1 localized near these IP_3R can sense this decrease in local $[\text{Ca}^{2+}]_{\text{ER}}$ and rapidly move to the ER-PM junctions to recruit and activate Orai1 (Thillaiappan et al. 2017). Together, these interesting findings place IP_3R as a central Ca^{2+} -signaling protein that links the receptor-mediated signaling events with those that result in SOCE activation. A key point to be considered is that both GPCR- and SOCE-associated signaling components are compartmentalized within ER-PM junctions. Localization of STIM1 in the close vicinity of IP_3R suggests cross talk between these two func-

tionally linked signaling complexes, in which the function of one protein may influence that of another. Alternatively, remodeling of SOCE-associated ER-PM junctions could bring the SOCE-associated signaling components close to the IP_3R /GPCR-signaling complex. This would allow IP_3R to rapidly respond to IP_3 generation and STIM proteins to efficiently detect a decrease in the ER locally near the site of Ca^{2+} release, and activate Orai1 and TRPC channels. Currently, available data do not resolve between these two possible scenarios, which may not be mutually exclusive.

RyRs and T-Cell Receptor

TCR engagement leads to depletion of Ca^{2+} in the ER- Ca^{2+} stores followed by activation of SOCE, primarily via Orai1 (Lewis and Cahalan 1989; Zweifach and Lewis 1996). The cellular location where T cells make contact with antigen-presenting cell, the immunological synapse (IS), represents the signaling center in the cell. It has been shown that Orai1 and STIM1 are rapidly recruited to the IS, resulting in enhanced localized calcium influx at this site. Further, Orai1 and STIM1 colocalized with IS proteins, CD3, CD28, and LFA-1. Two types of T-lymphocyte potassium channel, Kv1.3 and KCa3.1, that are indirectly involved in functional Ca^{2+} signaling, are also recruited to the IS. Although SOCE is not required for IS formation, blocking Ca^{2+} entry compromises its long-term stability (Panyi et al. 2004; Beeton et al. 2006; Nicolaou et al. 2007). For example, SOCE is involved in the reorganization of actin, which helps regulate the intensity and duration of TCR signaling (Hartzell et al. 2016).

The earliest intracellular signals that occur after T-cell activation are local Ca^{2+} microdomains, which occur in the sub-PM region, close to where the TCR is activated. Ryanodine receptor type 1 (RyR1), NAADP, and Ca^{2+} entry contribute to these Ca^{2+} microdomains (Wolf et al. 2015; Wolf and Guse 2017). Recently, Ca^{2+} microdomain formation has also been described in unstimulated T cells, which require Orai1, STIM1, and STIM2, and which are preclustered within ER-PM junctions (Diercks et al. 2018).

Following cell stimulation, the number of Ca^{2+} microdomains increased locally near the site of stimulation. This increase requires Orai1, STIM2, RyR1, and NAADP. The latter complex, involving ER- Ca^{2+} release, rapidly increases the number of Ca^{2+} microdomains, causing global spread of Ca^{2+} signals to promote full T-cell activation. Dependent on TCR stimulation, NAADP, IP_3 , and cADPR are formed sequentially to bind and activate their respective target Ca^{2+} channels in the ER such as RyRs and IP_3 Rs (Streb et al. 1983; Guse et al. 1999; Gasser et al. 2006). The preformed Orai1/STIM1/STIM2 complexes may enable the SOCE machinery in T cells to respond very quickly to ER- Ca^{2+} store depletion.

ACCESSORY PROTEINS WITHIN ER-PM JUNCTIONS—TETHERING AND LIPID TRANSFER PROTEINS

SOCE-associated ER-PM junctions are suggested to contribute to maintenance and/or replenishment of PM lipids, including sterols and phosphoinositides. Once in the junctions, STIM1 and Orai1 trigger the recruitment of proteins that induce remodeling of PM lipids as well as the underlying cytoskeleton (Jha et al. 2013; Sharma et al. 2013; Jing et al. 2015). Extended synaptotagmins (E-Syts1, -2, and -3) are ER proteins that tether to the PM and promote the formation and stabilization of ER-PM junctions. These proteins have a lipid-binding domain (SMP domain), which has been implicated in lipid transfer between membranes independently of vesicular transport (Herdman and Moss 2016). In E-Syts, the SMP domain binds phosphatidylcholine, phosphatidylethanolamine, phosphatidylinositol (PI), and phosphoserine (Schauder et al. 2014). The conserved domain 2 (C2 domain) has been shown to mediate the Ca^{2+} -dependent binding of E-Syts to PM PIP_2 , which is required for tethering the proteins to the PM. The ER-PM gap and characteristics of the junctions are determined by the carboxy-terminal lengths of the proteins. E-Syts2 and -3 contribute to formation of constitutive junctions, which display further narrowing following stimulation. In contrast, E-Syt1 is primar-

ily intracellular in unstimulated cells and recruited to ER-PM junctions after activation of SOCE, as it is anchored to PM PIP_2 in a Ca^{2+} -dependent manner (Min et al. 2007; Herdman and Moss 2016). Agonist stimulation of the cell induces formation of new junctions at which STIM1 co-localizes with Orai1 and E-Syt1. Although knockout of all three E-Syts is required to significantly reduce the number of ER-PM junctions, silencing E-Syt1 attenuates Orai1/STIM1 clustering (Giordano et al. 2013; Idevall-Hagren et al. 2015). However, the effect of E-Syt knockdown on SOCE has not been clearly established. Whereas one study shows that there is no effect on Orai1 function (Giordano et al. 2013), another shows diminished histamine-induced Ca^{2+} pulses (Chang et al. 2013). It can be suggested that ER-PM junctions containing E-Syts2 and -3 represent preformed sites where SOCE components can assemble immediately following ER- Ca^{2+} store depletion, whereas E-Syt1-dependent junctions represent newly formed junctions. Whether the function of E-Syts is determined by the stimulus intensity and how they contribute to STIM2 function are not yet known.

RASSF4, a member of the RASSF family of proteins that are proposed to function as adaptor or scaffolding proteins for large signaling complexes, modulates STIM1 translocation into the ER-PM junctions by regulating the ER-PM tethering functions of E-Syts2 and -3. Further, RASSF4 regulates the PM targeting of Arf6, an upstream regulator of phosphatidylinositol phosphate kinase (PIP5K ; Chen et al. 2017). Loss of endogenous RASSF4 significantly reduces the steady-state levels of PIP_2 , leading to adverse effects on junction formation and stability and SOCE activation. Conversely, overexpression of RASSF4 enhanced the formation of ER-PM junctions and clustering of STIM1 and SOCE (Chen et al. 2017). Arf6 can activate phospholipase D (PLD) and, thus, act synergistically to increase PM- PIP_2 levels. PLD also hydrolyzes phosphatidylcholine to generate phosphatidic acid (PA), which may act as a cofactor for activating PIP5K (Dickson 2017). PA also promotes recruitment of Nir2 to ER-PM junctions, which boosts the replenishment of PM PIP_2 levels following PLC-mediated hydrolysis

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of PIP₂. Both STIM1 and Nir2 colocalized with MAPPER, a marker for SOCE-associated ER–PM junctions (Chang et al. 2013). The synergistic actions of E-Syt1, Nir2, RASSF4, and Arf6 provide a positive feedback mechanism to replenish PM PI, PI-4-phosphate, and PIP₂ at contact sites where SOCE occurs, enabling cells to respond to further agonist stimulation and prolonging the duration of Ca²⁺ signaling (Chang et al. 2013; Chen et al. 2017; Dickson 2017). It is interesting that Ca²⁺-induced feed-forward regulation of E-Syts and other regulatory proteins involved in lipid transfer are localized in/or in close proximity to the SOCE domain. This allows PM PIP₂ to be maintained at levels optimal for assembly of the STIM/Orai1 complex and PLC-stimulated hydrolysis following cell stimulation. Whether lipid microdomain(s) involved in agonist-stimulated PIP₂ hydrolysis and SOCE vary and are differentially regulated needs further study.

CONCLUDING REMARKS

The nature of SOCE-associated ER–PM junctions, their molecular components and architecture, as well as their physiological relevance, are currently the subject of many studies in the field of Ca²⁺ signaling. Emerging data suggest that these junctions are not only hubs for sensing agonist stimulation and regulating SOCE but also for compartmentalizing specific downstream targets and effector proteins that drive SOCE-dependent regulation of cell function. Future studies should focus on the identification of the components involved in assembly and remodeling of these junctions. Furthermore, most of the currently available data have been obtained from studies with cell lines, using over-expressed proteins. Thus, information regarding the native status of these proteins and junctions is lacking. Further studies will be required to fill in this knowledge gap and increase our understanding of how receptor signaling, culminating in intracellular Ca²⁺ release, functionally and physically interact with, and regulate, SOCE. It is possible that tissue function will determine localization of the junctions as well as diversity in the components. Nevertheless, signaling

from PM to the ER and, conversely, from the ER to the PM is central to agonist regulation of SOCE.

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