

Invited review

Regulation of TRP-like muscarinic cation current in gastrointestinal smooth muscle with special reference to PLC/InsP₃/Ca²⁺ systemAlexander V ZHOLOS¹*Department of Physiology, Queen's University Belfast, Belfast, UK***Key words**smooth muscle; muscarinic receptor; G protein; phospholipase C; InsP₃; calcium; TRP cation channel¹ Correspondence to Prof A V ZHOLOS.

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Abstract

Acetylcholine, the main enteric excitatory neuromuscular transmitter, evokes membrane depolarization and contraction of gastrointestinal smooth muscle cells by activating G protein-coupled muscarinic receptors. Although the cholinergic excitation is generally underlined by the multiplicity of ion channel effects, the primary event appears to be the opening of cation-selective channels; among them the 60 pS channel has been recently identified as the main target for the acetylcholine action in gastrointestinal myocytes. The evoked cation current, termed mI_{CAT} , causes either an oscillatory or a more sustained membrane depolarization response, which in turn leads to increases of the open probability of voltage-gated Ca²⁺ channels, thus providing Ca²⁺ entry in parallel with Ca²⁺ release for intracellular Ca²⁺ concentration rise and contraction. In recent years there have been several significant developments in our understanding of the signaling processes underlying mI_{CAT} generation. They have revealed important synergistic interactions between M₂ and M₃ receptor subtypes, single channel mechanisms, and the involvement of TRPC-encoded proteins as essential components of native muscarinic cation channels. This review summarizes these recent findings and in particular discusses the roles of the phospholipase C/InsP₃/intracellular Ca²⁺ release system in the mI_{CAT} physiological regulation.

Introduction

It has long been known that in visceral smooth muscle cells acetylcholine (ACh) acts as an excitatory neurotransmitter^[1]. Thus, as the primary excitatory transmitter released by enteric motor neurons, ACh plays a central role in the control of complex patterns of motility of the gastrointestinal (GI) tract^[2]. The excitatory input is received by G protein-coupled muscarinic ACh receptors (mAChR) expressed in postjunctional cells—smooth muscle myocytes with the involvement of interstitial cells of Cajal^[3–5]. ACh binding to mAChRs triggers a complex array of membrane and intracellular signals leading to membrane depolarization and smooth muscle contraction, respectively^[5,6].

GI muscles in various species and various regions of the digestive tract express predominantly, if not exclusively, the M₂ and M₃ subtypes of mAChRs which are present typically at a ratio between 5:1 and 4:1^[3,7–9]. The cholinergic contractile

response is usually accompanied by membrane depolarization and/or the occurrence of slow waves and accelerated action potential discharge with a concomitant increase of the membrane conductance^[10–16]. The excitatory effect of mAChR stimulation is mainly attributed to the activation of cation (Na⁺ and Ca²⁺) or chloride conductances, or sometimes to the inhibition of potassium conductance, with the secondary activation of voltage-dependent Ca²⁺ channels (VDCC). Therefore, the excitatory action of ACh can be seen as an integral and important mechanism of the contractile response because it initiates Ca²⁺ influx through VDCCs. This Ca²⁺ entry signal can summate with the well-documented ACh-induced intracellular Ca²⁺ release, and these additive Ca²⁺ sources are necessary to maintain smooth muscle contraction.

Indeed, pharmacological blockade of L-type Ca²⁺ channels does not prevent membrane depolarization in response to mAChR stimulation^[17] but strongly reduces the contrac-

tile response^[18–21], indicating that this voltage-dependent Ca^{2+} entry is important for the cholinergic contraction rather than membrane depolarization.

The cholinergic GI smooth muscle contraction is generally regarded as an M_3 response mediated by the main universal Ca^{2+} signalling pathway $G_{q/11}$ coupled to phospholipase C- β (PLC- β) activation, which in turn hydrolyzes phosphatidylinositol 4,5-bisphosphate (PIP_2) into 1,2-diacylsn-glycerol (DAG) and *D*-myo-inositol 1,4,5-trisphosphate (InsP_3)^[22]. By rapidly releasing Ca^{2+} from the sarcoplasmic reticulum InsP_3 evokes a transient intracellular Ca^{2+} concentration ($[\text{Ca}^{2+}]_i$) increase, thus initiating the myocyte contraction. Conversely, the major M_2 subtype apparently plays a central and direct role in many ion channel effects but its role in the cholinergic contraction is less clear. Unno *et al*^[21] recently addressed the specific roles of M_2 and M_3 receptors in agonist-evoked contraction of the guinea pig ileal longitudinal smooth muscle. Their results suggest that cholinergic contractions primarily originate from the integration of Ca^{2+} entry and Ca^{2+} sensitization of myofilaments. The authors thus conclude that M_3 -mediated Ca^{2+} store release might contribute to the contraction indirectly through potentiation of the electrical membrane responses.

Central to this hypothesis is the notion that muscarinic cation channel activation is strongly potentiated by $[\text{Ca}^{2+}]_i$ ^[23,24]. Although mI_{CAT} in smooth muscle cells has been extensively characterized as an M_2 -mediated pertussis toxin-sensitive response primarily resulting from G_o activation^[25–30], accumulating evidence suggests that mI_{CAT} is in fact a mixed M_2/M_3 response^[31–35]. Therefore, the hypothesis that an interaction between M_2 and M_3 receptors plays a crucial role in the contractile response^[21] is consistent with the synergistic activation of the muscarinic cation channels by both mAChR subtypes. This review discusses mI_{CAT} regulation with the involvement of various M_2 -, M_3 -, and Ca^{2+} -dependent signaling pathways.

Muscarinic effects on ion channels in GI smooth muscles

It has long been known that the depolarizing effect of ACh and other muscarinic agonists on GI smooth muscles is mediated mostly by an increase in the Na^+ permeability^[12]. However, the reversal potential of carbachol-evoked currents in the intestinal smooth muscle is approximately -10 mV and sensitive to changes in both $[\text{Na}^+]_o$ and $[\text{K}^+]_o$, but relatively insensitive to changes in $[\text{Cl}^-]_o$ ^[15], which indicates a non-selective increase in membrane permeability.

More recently, multiple muscarinic ion channel effects have been described in voltage-clamped cells, which include

generation of inward cationic or chloride current, inhibition or potentiation of voltage-dependent Ca^{2+} current, and modulation of several types of K^+ currents (reviewed in detail elsewhere^[3,5,6,36,37]).

Consistent with the direct role of the M_2 receptor, some ion channel effects of muscarinic agonists are reduced or even abolished by pertussis toxin treatment (eg, cation channel activation, BK_{Ca} channel inhibition), which selectively uncouples M_2 receptors from G_i/G_o proteins. Conversely, M_3 receptors typically modulate ion channels indirectly through $G_{q/11}$ coupling to PLC- β activation and accumulation of InsP_3 and DAG. InsP_3 evokes a transient $[\text{Ca}^{2+}]_i$ increase by Ca^{2+} release, whereas DAG initiates the translocation and activation of protein kinase C (PKC) in a Ca^{2+} -dependent manner. These second messenger pathways are often interposed between M_3 receptors and target channels, notably those sensitive to $[\text{Ca}^{2+}]_i$ (ie, potentiation of cationic, Cl^- and BK_{Ca} channels and inhibition of VDCC).

From these studies, clear implications arise concerning the functional importance of the numerous muscarinic effects on ion channels. Opening of cationic and Cl^- channels will produce membrane depolarization, thus promoting Ca^{2+} influx for the contractile response. As both types of channels are $[\text{Ca}^{2+}]_i$ -sensitive they were proposed to be involved in a positive feedback loop whereby Ca^{2+} entry and Ca^{2+} release promote membrane depolarization and further Ca^{2+} influx^[38,39], termed Ca^{2+} -induced Ca^{2+} entry^[40]. This action might be further facilitated by the muscarinic inhibition of several types of K^+ channels, such as BK_{Ca} , delayed rectifier and K_{ATP} channels^[37]. There are also mechanisms that provide negative feedback control by either limiting (VDCC channel inhibition, mI_{CAT} desensitization) or terminating (BK_{Ca} activation) cholinergic membrane depolarization.

General properties of mAChR-operated cation channels

Muscarinic effects on many different channels, which occur in parallel, raise the challenging task of understanding their relative importance, especially in the context of the interactions between M_2 and M_3 receptors. However, activation of mI_{CAT} is undoubtedly one of the major mechanisms of GI smooth muscle excitation. Cation channels carrying mI_{CAT} in many smooth muscles have relatively low Ca^{2+} permeability^[23,41,42] (eg, with 110 mmol/L extracellular CaCl_2 and 145 mmol/L intracellular CsCl, a $P_{\text{Ca}}/P_{\text{Cs}}$ ratio of 2.8 was estimated^[43]); therefore the major pathway for Ca^{2+} entry is through VDCC. However, in some smooth muscles, Ca^{2+} permeability of this channel is more substantial providing an additional Ca^{2+} influx^[44–46]. For example, in tracheal myocytes

the fraction of the cationic current carried by Ca^{2+} at -60 mV under physiological ion gradients was estimated at 14%^[45], but in guinea pig gastric myocytes it amounts to only 1%^[43]. The relative channel permeability to various monovalent cations was evaluated in gastric myocytes. The ratio was $\text{Rb}^+:\text{K}^+:\text{Cs}^+:\text{Na}^+:\text{Li}^+=1.1:1.1:1.0:0.98:0.8$ with a negligible anionic component^[47]. However, the amplitude of the inward current was largest with Cs^+ as the main permeant cation, and smallest with Na^+ (ie, the rank order was $\text{Cs}^+>\text{K}^+>\text{Li}^+>\text{Na}^+$), and unitary conductance was in the order Cs^+ (34 pS) $>$ Na^+ (25 pS) $>$ Li^+ (21 pS)^[47].

Nonstationary noise analysis of mI_{CAT} induced by intracellular perfusion of GTP γ S in ileal myocytes suggested a mean channel open probability (P_o) of 0.48 (at -40 mV during maximal conductance activation) and a total number of channels (n) of approximately 750^[48]. Similar values ($P_o=0.43$ and $N=830$) were derived from direct single channel measurements in outside-out patches exposed to 50 $\mu\text{mol/L}$ carbachol, suggesting channel density on average of one channel per 6.45 μm^2 ^[49]. The channels appear to be clustered as two-thirds of patches were "blank" and approximately 20% of patches had 2–7 active channels. This uneven distribution of active channels in membrane fragments might reflect not only clustering of the channels themselves, but also a tight co-localization of mAChR and associated signaling molecules.

In ileal myocytes, the agonist concentration dependence of the channel activation is characterized by a mean EC_{50} value of 7.6 $\mu\text{mol/L}$ ^[31]. The range of carbachol concentration over which cationic conductance increases in these single isolated cells is thus identical to that measured in the intact tissue^[15]. Although in different cells the EC_{50} values varied from approximately 1 to 30 $\mu\text{mol/L}$, their frequency distribution was normal. Thus, within the cell population there seem to be no subgroups of cells, which would differ in their sensitivity to carbachol. Notably, the depolarizing effect of muscarinic agonists saturates at a lower agonist concentration, which can be explained by a large functional reserve in this system owing to the high input resistance of GI myocytes.

Pharmacological properties of mI_{CAT}

Selective blockers of the muscarinic cation current are lacking, although a number of other ion channel blockers can efficiently inhibit mI_{CAT} . These include the following Ca^{2+} channel blockers (IC_{50} values in parentheses): Zn^{2+} (38 $\mu\text{mol/L}$); Cd^{2+} (98 $\mu\text{mol/L}$); Ni^{2+} (131 $\mu\text{mol/L}$); Co^{2+} (700 $\mu\text{mol/L}$); and Mn^{2+} (1 mmol/L)^[40,45,50,51]. Their blocking action is practically voltage-independent, with little change in the reversal potential or sensitivity to the agonist. K^+ channel or-

ganic blockers inhibit cationic current in a voltage-dependent manner, with quinidine being the most potent ($\text{IC}_{50}=0.25$ $\mu\text{mol/L}$), followed by quinine (1.0 $\mu\text{mol/L}$), 4-aminopyridine (3.3 mmol/L), TEA^+ (4.1 mmol/L), and procaine (1–5 mmol/L)^[40,52–54]. Caffeine, a Ca^{2+} -releasing agent, also blocks cationic current (~ 10 mmol/L)^[53]. Diphenylamine-2-carboxylate derivatives (DCDPC and flufenamic acid) strongly inhibit Ca^{2+} -activated cationic channels and Cl^- channels, and block mI_{CAT} with IC_{50} values of approximately 30 $\mu\text{mol/L}$ ^[53].

SK&F 96365, a common blocker of receptor-stimulated Ca^{2+} entry, showed a most peculiar blocking action on mI_{CAT} , which was time- and voltage-dependent producing concomitant alteration of the steady-state current-voltage (I - V) relationship^[55]. At a constant SK&F 96365 concentration the degree of the mI_{CAT} inhibition was a sigmoidal function of the membrane voltage with a slope factor of approximately 13 mV and half-maximal inhibition at approximately 30 mV. This unique time- and voltage-dependent mode of SK&F 96365 action on the native channel might be very useful for its identification in future comparative expression studies. Extracellular spermine and putrescine also blocked mI_{CAT} in a voltage-dependent manner with IC_{50} values (at -40 mV) of approximately 1 and 5 mmol/L , respectively^[56]. This inhibitory action was relieved by membrane depolarization. Intracellular polyamines also inhibited mI_{CAT} , but its complex N-shaped activation curve did not seem to result from this effect.

Voltage dependence of mI_{CAT} : synergy with G protein activation

mI_{CAT} that is evoked in GI myocytes either by mAChR activation or by intracellular application of GTP γ S in the absence of an agonist has interesting voltage-dependent properties. Its steady-state I - V curve shows double rectification around the reversal potential, and with hyperpolarization the current first increases in keeping with an increase in the driving force, reaches a maximum at approximately -40 to -50 mV, then declines. In some cells mI_{CAT} can even be completely lost when the membrane potential reaches approximately -100 to -120 mV. By analogy with the classical voltage-gated channels, a Boltzmann relationship was often used to describe this voltage-dependence (although its usefulness is limited to negative potentials because the activation curve is overall N- rather than S-shaped), and a two-state (one open and one closed) model was introduced as an underlying mechanism^[57]. The voltage dependence was characterized by the potential of half-maximal activation of approximately -50 mV and the slope factor in the range 15–27 mV^[57–59]. A receptor-operated cationic current

displaying a similar U-shaped I - V relationship at negative potentials was described in many different smooth muscle tissues^[38,40,45,51,52,54,60,61], suggesting that this current is ubiquitous.

During a negative voltage step the current increased instantaneously and proportionally to the increase in the driving force, but then it exponentially relaxed to a smaller steady-state level with a time constant in the order of 50–150 ms which decreased with hyperpolarization^[52,57]. After repolarization to the holding potential, monoexponentially decaying tail currents were observed.

This behavior is characteristic for voltage-gated ion channels and thus consistent with the idea that some charged groups (“voltage sensor”) influence the channel P_o . However, as this channel is primarily gated by mAChR and associated G protein activation (ie, depolarization by itself hardly activates it), the gating mechanisms could be undoubtedly far more complex. Indeed, it turned out that the voltage range of the channel activation was one of the primary targets of G protein activation^[59]. In some cells even a completely linear I - V curve could be seen over the range of 0 to -120 mV at high agonist concentration (300 μ mol/L carbachol) or following sufficiently long 200 μ mol/L GTP γ S infusion. With desensitization, however, the I - V relationship again acquired its usual U-shape^[62]. Associated with these changes in the I - V shape, there was also a pronounced change in the rate of relaxation of the current during a voltage step. The physiological significance of this G protein modulation of the cationic channel voltage dependence is obviously in creating a system where there is cross-talk between mAChR/G protein activation, channel opening and membrane depolarization.

This interesting “interchangeability” of the effects of membrane potential and G protein activation on channel gating suggests that a strong synergy exists in this system, to an extent where it is difficult to distinguish between the primary and secondary gating stimuli. Indeed, if, instead of using the Boltzmann relation, mI_{CAT} amplitude measured at different potentials is plotted against the agonist concentration and data is fitted by the Hill relation, this reveals the main effect of voltage on the agonist sensitivity, characterized by approximately 80% EC_{50} reduction between -120 and 80 mV^[63,64]. There is also a significant reduction of the latency of the response, acceleration of the agonist on-rate and reduction of the off-rate associated with membrane depolarization. All these observations imply that some ligands produced by mAChR stimulation might interact with the channel in a voltage-dependent manner, rather than there being a classical “voltage sensor” in its structure. In con-

nection with this, it is notable that the channel voltage range of activation is also affected by PLC- β inhibition and by $[Ca^{2+}]_i$ ^[65,66]. This is relevant to the recently proposed linkage between mI_{CAT} and TRPC4/5 channels^[17,67–69] as TRP channels are generally lacking the full complement of charged amino acids (ie, classical “voltage sensors”) in the S4 segment^[70].

It should be noted, however, that recent single channel studies began to reveal intrinsic voltage-dependent channel gating, which was not evident in previous whole-cell measurements. Although membrane depolarization alone could hardly induce any significant whole-cell current, in single channel measurements spontaneous background channel activity was revealed. It was characterized by long closings and only brief openings consistent with the C1–O1 gating (see below) and it was channel gating in this pair of states that showed the most prominent voltage dependence^[49,71]. Membrane depolarization strongly accelerated these spontaneous openings (the main effect of voltage is to shorten the long closed state of the channel) although they remain too brief to generate any measurable whole-cell current.

Signal transduction: synergy between M_2 and M_3 receptors

Early studies have shown that mI_{CAT} in GI myocytes can be almost completely blocked by treating the cells with pertussis toxin^[25,27,28,35] which raised an immediate possibility that it was the M_2 receptor subtype that triggered mI_{CAT} . Consistent with this, experiments with the use of antibodies against the α -subunits of a range of G-proteins pinpointed $G_{\alpha o}$ involvement and excluded the role of the $\beta\gamma$ -subunits of G-proteins^[26,29].

Subsequent pharmacological studies confirmed the primary role of M_2 receptor activation in mI_{CAT} generation, but, quite unexpectedly, mI_{CAT} was also found to be strongly dependent on the activation of the M_3 receptor subtype^[30–35,72]. This was evident from the effects of M_3 -selective muscarinic antagonists, which, particularly at low concentrations, did not displace the agonist curve but produced a substantial reduction of the maximum cationic conductance^[31,32].

One well established mechanism of the possible M_3 receptor convergence on the M_2 -activated mI_{CAT} is a $[Ca^{2+}]_i$ rise due to $InsP_3$ -induced Ca^{2+} release. This is operational because cation channel opening is strongly potentiated by intracellular Ca^{2+} . A better documented example of such a role of M_3 receptors in mI_{CAT} modulation is concurrent mI_{CAT} and $[Ca^{2+}]_i$ oscillations^[23,66,73–76]. However, even when this Ca^{2+} -mediated link is disabled, for example, by strongly buffering $[Ca^{2+}]_i$, M_3 -dependent regulation of mI_{CAT} is

still present^[31,32] and a significant correlation of muscarinic agonist potencies between the M_2/mI_{CAT} and the $M_3/PLC-\beta/InsP_3$ systems still exists^[34].

The interaction between M_2 and M_3 receptors in producing mI_{CAT} shows up somewhat differently in guinea pig gastric myocytes under “unclamped” $[Ca^{2+}]_i$ conditions, in which case the M_3 selective blockade does not reduce the maximal response, but instead reduces the agonist sensitivity of the M_2 -induced current^[35].

Intracellular Ca^{2+} dependence of mI_{CAT} : synergy with the PLC- $\beta/InsP_3$ system

The $[Ca^{2+}]_i$ -dependency aspect of mI_{CAT} regulation has received perhaps the most attention. An elevation of $[Ca^{2+}]_i$ by itself is somewhat insufficient to induce mI_{CAT} , but when the channel is primed by an activated G protein its opening is greatly potentiated by a rise in $[Ca^{2+}]_i$ ^[23,24,30,38,40,44,60,73,74,76]. However, when $[Ca^{2+}]_i$ is abnormally low (eg, at a high EGTA or BAPTA concentration inside the cell) no or very little channel activation is possible even at maximal mAChR stimulation^[24,44,61,74]. Thus, it follows that intracellular Ca^{2+} has both a permissive and a facilitating effect on channel gating.

With low $[Ca^{2+}]_i$ buffering, mAChR stimulation or direct G protein activation by intracellular GTP γ S produces $[Ca^{2+}]_i$ oscillations, which result from periodic release of Ca^{2+} from the intracellular stores. These cause temporally closely related oscillations of the cationic current^[23,27,74]; the frequency of these oscillations is strongly modulated by Ca^{2+} influx^[77]. Interestingly, $[Ca^{2+}]_i$ rise results in a considerable negative shift of the activation curve by approximately 30 mV^[64], that is, linearization of the $I-V$ relationship occurs similarly to that caused by increasing receptor stimulation. Both $[Ca^{2+}]_i$ and mI_{CAT} oscillations are similarly sensitive to Ca^{2+} store depletion produced by $InsP_3$, thapsigargin or cyclopiazonic acid, as well as to the $InsP_3$ receptor blocker heparin^[23,30,74,77,78].

The relationship between $[Ca^{2+}]_i$ and mI_{CAT} amplitude was quantified by buffering $[Ca^{2+}]_i$ at different levels using either EGTA/ Ca^{2+} or BAPTA/ Ca^{2+} mixtures or, in the same cell, by varying the amount of Ca^{2+} entering the cell through VDCC^[24,64,65]. These experiments revealed a calcium EC_{50} value of approximately 200–400 nmol/L. However, prolonged $[Ca^{2+}]_i$ elevation (eg, $[Ca^{2+}]_i$ buffering at 500 nmol/L) caused strong mI_{CAT} desensitization even if the current was induced by GTP γ S without receptor stimulation^[64,79], an effect presumably related to PKC activation^[80]. At the single channel level, this biphasic $[Ca^{2+}]_i$ dependence is seen as a peak of channel P_O with 100 nmol/L Ca^{2+} at the internal side of the membrane compared with either 30 or 500 nmol/L $[Ca^{2+}]_i$; no

effect of $[Ca^{2+}]_i$ on the unitary conductance was observed^[81].

Interestingly, mI_{CAT} was found to be most responsive to rapid changes in $[Ca^{2+}]_i$ caused by abrupt flash photolysis release of $InsP_3$, but was rather insensitive to $[Ca^{2+}]_i$ elevation produced in small steps by NP-EGTA photorelease^[76]. Moreover, a striking difference was found between the effects of flash-released $InsP_3$ and caffeine applications, as Ca^{2+} release through ryanodine receptors (RyR) failed to potentiate mI_{CAT} . This can be attributed, at least in part, to the differential distribution of $InsP_3$ and RyRs within a smooth muscle myocyte, as type I $InsP_3$ receptors were predominantly localized in a close juxtaposition to the plasma membrane but RyRs were mostly found in the central region of the cell^[76]. Even more intriguingly, Ca^{2+} sparks induced STOCs but failed to potentiate mI_{CAT} . This dependence of mI_{CAT} on a global rather than local $[Ca^{2+}]_i$ rise is similar to differential activation of STOCs and $I_{Cl(Ca)}$ by Ca^{2+} sparks and waves, respectively, in rat portal vein myocytes^[82].

Thus, one possibility is that the spatiotemporal pattern of the calcium signal is an important determinant of the $[Ca^{2+}]_i$ -dependent modulation of mI_{CAT} . However, the recently proposed link between mI_{CAT} and TRPC4/5 proteins^[67–69] raises other intriguing possibilities and questions. In different studies, TRPC4 and TRPC5 have been shown to form either store- or receptor-operated cation channels that can be activated with or without the involvement of $InsP_3$ ^[83,84]. Overall, many properties of mI_{CAT} are highly similar to receptor-operated cation channels formed by TRPC4 and TRPC5, which includes the complex effects of intracellular Ca^{2+} (ie, permissive, potentiating and desensitizing action)^[85].

Further similarities of mI_{CAT} and TRPC4/5 activation include regulation through the PLC pathway. Several recent studies revealed the importance of PLC in mI_{CAT} activation^[66,68,86]. In murine gastric myocytes the current was inhibited by the anti- $G_{q/11}$ antibody, the PLC blocker U-73122 and the IICR inhibitor 2-APB, but was insensitive to the anti- G_o antibody^[68]. By contrast, in guinea pig ileal myocytes PLC inhibition abolished mI_{CAT} without the involvement of DAG, $InsP_3$ or Ca^{2+} store depletion^[66,86]. Strikingly, the anti- $G_{q/11}$ antibody was ineffective in ileal myocytes^[29,86], raising questions about the PLC isoforms involved in mI_{CAT} activation in these cells. One possibility is that M_2 receptors can also couple to PLC activation through $\beta\gamma$ dimers released from $G_{i/o}$ proteins^[87,88] and this link can obscure the effects of the anti- $G_{q/11}$ antibody. However, such a role of $\beta\gamma$ -subunits would be inconsistent with the lack of the effects of the $G\beta$ -antibody as well as $\beta\gamma$ dimer infusion in the same cells^[29]. Even more importantly, recent studies have shown that mI_{CAT} is lacking in cells isolated from M_2 or M_3 knockout mouse^[89],

but if the M_2 /PLC linkage were sufficient to support its activation the current would be present in the M_3 subtype knock-out mouse. Tyrosine kinase-dependent pathways also play a role in mI_{CAT} regulation^[90], therefore the role of receptor tyrosine kinases that couple to PLC- γ deserves further investigation. Finally, there is a possibility that some soluble PLC isoforms are involved in mI_{CAT} activation, which are not inhibited by G protein antibodies^[91].

The role of DAG as a possible intermediary between PLC activation and mI_{CAT} was also addressed in these studies, however both in gastric and ileal myocytes OAG, an analogue of DAG, failed to induce any significant current. Thus, the signalling events downstream of PLC activation remain unknown. It is also unclear how the PLC pathway activates TRPC4/5 channels; one possibility that needs to be explored is that the channels are activated by the depletion of the PLC substrate, like PIP₂, rather than by the PLC products. However, PLC generates other potential signalling molecules, such as poly-unsaturated fatty acids (PUFAs), and the role of the numerous lipid messengers in mI_{CAT} activation remains to be explored.

It also remains unknown whether intracellular Ca²⁺ directly binds to the channel protein or exerts its modulatory effect by an intermediate enzymatic step. Following agonist application there is a considerable latency of approximately 230 ms and a time lag of approximately 1.2 s between peaks of [Ca²⁺]_i and mI_{CAT} during the first Ca²⁺ wave, but subsequent [Ca²⁺]_i oscillations are mirrored by mI_{CAT} very closely^[76]. These kinetics data thus suggest that channel Ca²⁺ “priming”, or permissive effect, might be indirect, but once established the channel might be regulated by Ca²⁺ in a more direct manner. Possible intermediates include calmodulin and myosin light chain kinase^[61,92]. It should be also noted that the cAMP/PKA pathway is not involved in mI_{CAT} regulation^[66].

Channel gating mechanism: cyclical transitions between 4 connected open and closed states

Summarizing the signal transduction pathways, the muscarinic cation channel is a voltage- and Ca²⁺-sensitive channel gated by G α -GTP in a PLC-dependent manner, as shown in Figure 1. Various ligands could produce kinetically distinct channel conformations, therefore it was reasonable to expect a similarly complex channel gating mechanism. Single channel activity was studied in membrane patches isolated from guinea pig and murine ileal and gastric myocytes. Cation channels with voltage-dependent properties consistent with the whole-cell current behavior had unitary conduc-

tances of 35 pS (guinea pig gastric myocytes^[47]), 57 pS (guinea pig ileal myocytes^[49]) and 70 pS (murine ileal myocytes^[81]). These differences to some extent are related to different ion conditions, for example, for recordings in guinea pig gastric myocytes 2 mmol/L external Ca²⁺ was used, and in murine ileal myocytes an addition of 2.5 mmol/L Ca²⁺ to the external solution was shown to reduce unitary conductance from 70 to 46 pS.

In guinea pig ileal myocytes, the 57 pS channel has at least 8 kinetically distinct states, 4 open and 4 closed. Analysis of adjacent dwell times revealed strong correlations, which suggested connections between them, as shown in Figure 2. These are features present in many other ligand-gated ion channels, such as nAChR or BK_{Ca} channels. One unusual property of the mAChR-gated channel is the presence of prominent regular cycles of P_O that occurred due to a variable number of long openings between consecutive long shuttings, or, in other words, due to periodical shifts of gating between the two main modes, the low- P_O mode in the C1–O1 states (long closings and brief openings) and the high- P_O mode in the C4–O4 states (conversely, long openings and brief closings).

Presently the origin of the channel voltage dependence in this channel mechanism is clear but the nature of ligands stabilizing various open states needs further exploration. Thus, membrane potential affects vertical transitions but does not cause any net horizontal redistribution between the states. Interestingly, as the channel activity progresses from low- to high- P_O mode (ie, gating shifts from left to right in Figure 2) channel gating becomes less voltage-dependent. Thus, the C1 dwell time shows the strongest voltage dependence, but the C4 mean dwell time remains unaltered at potentials between -120 and -10 mV.

Considering how channel ligands could produce these various kinetically distinct channel conformations, it is important to note that channel P_O according to this scheme increases in 3 stages, and that the O1 and O2 states could hardly generate any significant integral current as their joint contribution to the overall P_O is only 3%. Thus, any signals that can induce only these 2 open states would produce a tiny whole-cell current; nevertheless, such signals might be crucially important for further progression of the channel activation towards the C3–O3 and C4–O4 gating. The O3 and O4 states were estimated to generate, respectively, 24% and 73% of the whole-cell current at maximal receptor activation.

Because channel gating similar to the C1–O1 gating is seen in the absence of mAChR activation it seems to be an intrinsic voltage-dependent channel property. Based on the

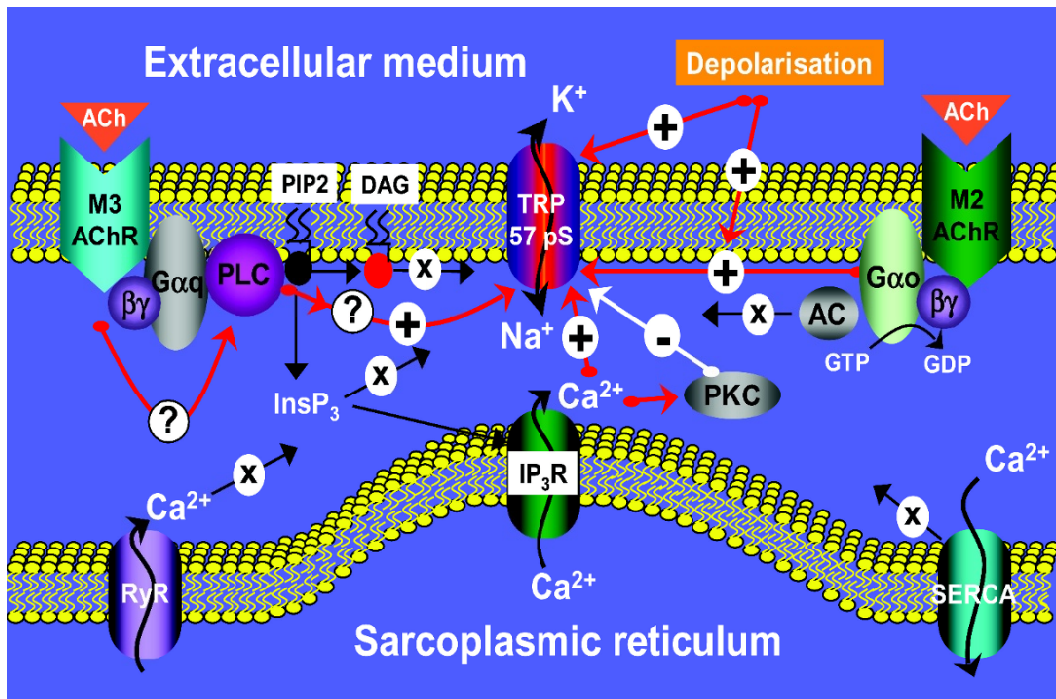


Figure 1. Multiple signal transduction pathways linking M₂ and M₃ receptor activation to TRP-like cation channel opening in GI myocytes. Many activation pathways (+) act in synergy but the precise nature of their interaction remains unclear, particularly regarding the possibility of direct G_{αo}-GTP and Ca²⁺ binding to the channel, the enigmatic M₃ or M₂ link to the PLC activation bypassing the G_{q/11} protein, and the events downstream of PLC activation. Ca²⁺-dependent activation of PKC appears to be the main mechanism of mI_{CAT} desensitization (-). However, several pathways do not seem to play a role in mI_{CAT} regulation (x); these include DAG, βγ-dimers of G proteins, Ca²⁺ store depletion (eg, SERCA inhibition), Ca²⁺ release through RyRs, direct action of InsP₃ and adenylyl cyclase (AC) inhibition.

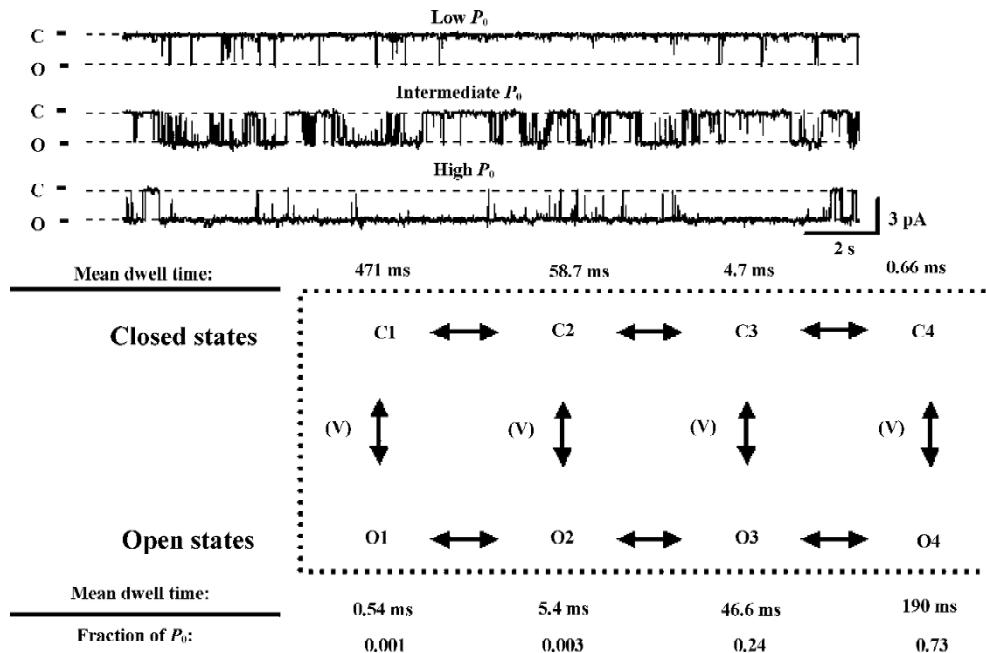


Figure 2. Channel gating mechanism with mean dwell times as measured in outside-out patches held at -40 mV and exposed to external 50 μmol/L carbachol or internal 200 μmol/L GTPγS^[49]. Typical examples of channel gating in the low (C1-O1), intermediate (C2-O2 and C3-O3) and high (C4-O4) P₀ modes are shown. C, closed; O, open.

above considerations, one can also suggest that any permissive effects (eg, $[Ca^{2+}]_i$, the PLC/InsP₃ system) are fulfilled through the occurrence of the intermediate states, such as O2, whereas full channel activation in the longer O3 and O4 open states requires cooperative interactions of all channel ligands, including G α -GTP. This hypothesis can mechanistically account for many permissive and major synergistic links in mI_{CAT} activation, which were discussed in this review, and might offer a useful model for future experimental tests.

Conclusion

The past decade has led to significant progress in our understanding of the roles of M₂ and M₃ receptors in GI smooth muscle excitation through the detailed studies of mI_{CAT} , a primary depolarizing current. Many important synergies have been revealed, including cross-talk between receptor subtypes, Ca²⁺ and InsP₃, and the crucial roles of both G_o and PLC- β activation. However, exactly how G_o protein is involved in mI_{CAT} generation and the activator in the PLC pathway remain unknown. PLCs are complexly regulated by various receptors, and it appears that they are one of the most important merging points in the pathways linking M₂ and M₃ receptors to cation channels. Synergistic mechanisms also exist in receptor-mediated PLC activation, thus more research is needed on the PLC-dependent modulation of mI_{CAT} . The mouse model of mI_{CAT} has been recently validated^[93], and it shows all the essential features of the complex regulation of mI_{CAT} discussed in this review. Thus, future significant progress can be expected through the studies of mI_{CAT} in genetically modified mice lacking certain receptors or other putative elements of the complexly intervened signal transduction pathways. The sequence of ligand interactions with the channel also remains unknown, but if established it will provide the most conclusive evidence for the specific roles of M₂ and M₃ receptors in smooth muscle excitation. The recently proposed TRPC4/5 connection raises further challenging questions, for example, the reasons why heterologously expressed TRPC4/5 channels can be activated by G_{q/11}-coupled receptors alone, but the native channel requires simultaneous activation of M₂ receptor and G_o.

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