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Mechanisms underlying cannabinoid inhibition of presynaptic Ca^{2+} influx at parallel fibre synapses of the rat cerebellum

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Activation of CB1 cannabinoid receptors in the cerebellum acutely depresses excitatory synaptic transmission at parallel fibre–Purkinje cell synapses by decreasing the probability of glutamate release. This depression involves the activation of presynaptic 4-aminopyridine-sensitive K^+ channels by CB1 receptors, which in turn inhibits presynaptic Ca^{2+} influx controlling glutamate release at these synapses. Using rat cerebellar frontal slices and fluorometric measures of presynaptic Ca^{2+} influx evoked by stimulation of parallel fibres with the fluorescent dye fluo-4FF, we tested whether the CB1 receptor-mediated inhibition of this influx also involves a direct inhibition of presynaptic voltage-gated calcium channels. Since various physiological effects of CB1 receptors appear to be mediated through the activation of PTX-sensitive proteins, including inhibition of adenylate cyclases, activation of mitogen-activated protein kinases (MAPK) and activation of G protein-gated inwardly rectifying K^+ channels, we also studied the potential involvement of these intracellular signal transduction pathways in the cannabinoid-mediated depression of presynaptic Ca^{2+} influx. The present study demonstrates that the molecular mechanisms underlying the CB1 inhibitory effect involve the activation of the PTX-sensitive G_i/G_o subclass of G proteins, independently of any direct effect on presynaptic Ca^{2+} channels (N, P/Q and R (SNX-482-sensitive) types) or on adenylate cyclase or MAPK activity, but do require the activation of G protein-gated inwardly rectifying (Ba^{2+} - and tertiapin Q-sensitive) K^+ channels, in addition to 4-aminopyridine-sensitive K^+ channels.

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In the cerebellar cortex, activation of presynaptic type 1 cannabinoid receptors (CB1) (Herkenham *et al.* 1991) inhibits excitatory synaptic transmission at parallel fibre (PF)–Purkinje cell (PC) synapses by decreasing the probability of glutamate release (Levenes *et al.* 1998). This results from the activation of presynaptic 4-aminopyridine-sensitive K^+ channels which in turn inhibit presynaptic Ca^{2+} influx controlling glutamate release at these synapses (Daniel & Crepel, 2001). However, in the latter study, we did not determine whether this CB1 receptor-mediated inhibition of presynaptic Ca^{2+} influx also implies a direct inhibition of presynaptic voltage-gated calcium channels (VGCCs). At PF–PC synapses there are at least three pharmacologically distinguishable types of VGCCs that synergistically control glutamate release (Mintz *et al.* 1995). In these fibres, presynaptic Ca^{2+} influx

occurs through (i) ω -agatoxin IVA (ω -Aga-IVA)-sensitive P/Q-type VGCCs, (ii) ω -conotoxin GVIA-sensitive N-type VGCCs and (iii) other VGCCs resistant to ω -Aga IVA, ω -conotoxin GVIA and dihydropyridines (Mintz *et al.* 1995). These other VGCCs might correspond to R-type channels, or to more than one type of as-yet-unidentified VGCC (see Wu & Sagau, 1997). In heterologous expression systems and cell cultures, cannabinoid receptor activation has been shown to inhibit P/Q- and N-type VGCCs (Caulfield & Brown, 1992; Mackie & Hille, 1992; Mackie *et al.* 1995; Pan *et al.* 1996; Twitchell *et al.* 1997; Sullivan, 1999; Nogueron *et al.* 2001). On the other hand, the inhibitory effect of cannabinoids in rat striatal slices involves N- but not L-, P- or Q-type VGCCs (Huang *et al.* 2001), while in slices from the mouse nucleus accumbens or lateral amygdala the same effect does not involve any

of these channels (Robbe *et al.* 2001; Azad *et al.* 2003). Finally, in hippocampal brain slice preparations, endocannabinoids, acting as retrograde messengers, mediate a selective presynaptic inhibition of N-type Ca^{2+} channels (Wilson *et al.* 2001). Such direct inhibition of presynaptic VGCCs might also be operational at cerebellar PFs following activation of presynaptic CB1 receptors, and thus might participate in the reduction of glutamate release at PF–PC synapses.

Cannabinoid CB1 receptors have an amino acid sequence typical of G protein-coupled receptors (Matsuda *et al.* 1990), and many physiological effects of cannabinoids appear to be mediated through activation of PTX-sensitive G_i/G_o proteins (Caulfield & Brown, 1992; Mackie & Hille, 1992; Holwett, 1995; Mackie *et al.* 1995; Pan *et al.* 1996; Twitchell *et al.* 1997; Prather *et al.* 2000; Nogueron *et al.* 2001) and negative coupling to adenylate cyclase (for a review see Childers & Deadwyler, 1996; also Holwett, 1985; Pacheco *et al.* 1993; Jung *et al.* 1997). However, other studies have reported that CB1 receptor activation can also stimulate adenylate cyclase activity, probably through the G_s subclass of G proteins (Glass & Felder, 1997; Maneuf & Brotchie, 1997; Felder *et al.* 1998). Finally, agonist stimulation of CB1 receptors may activate other signal transduction pathways via the PTX-sensitive G_i/G_o proteins. In particular, evidence that the CB1 receptor is functionally coupled through these proteins to the mitogen-activated protein kinases (MAPKs) (Bouaboula *et al.* 1995, 1999; Derkinderen *et al.* 2001, 2003) or to the inwardly rectifying K^+ channels (Mackie *et al.* 1995; McAllister *et al.* 1999) has been reported.

Here we show that CB1 inhibition of presynaptic Ca^{2+} transients evoked by PF stimulation is mediated through the activation of the PTX-sensitive G_i/G_o subclass of G proteins. This effect is independent of any direct inhibition of presynaptic Ca^{2+} channels (N, P/Q and R (SNX-482-sensitive) types) and of any modulation of adenylate cyclase or MAPK activity, but does involve activation of G protein-gated inwardly rectifying (Ba^{2+} and tertiapin-sensitive) K^+ channels, in addition to 4-aminopyridin-sensitive K^+ channels. These presynaptic molecular events were explored in rat cerebellar slices using fluorometric methods and the fluorescent low affinity Ca^{2+} -sensitive dye fluo-4FF.

Methods

Preparation of cerebellar slices

Animal care and all experimental procedures used during experiments were in accordance with CNRS national

guidelines. Coronal or saggital slices were prepared from the vermis of the cerebellum of male Sprague-Dawley rats aged 17–36 days. In brief, the rats were stunned and decapitated, and cerebellar slices (200–250 μm thick) were cut with a vibroslicer (Campden Instruments Ltd). The slices were kept at room temperature for at least 1 h before recording in a storage chamber of saline solution saturated with 95% O_2 –5% CO_2 . This solution contained (mM): NaCl, 124; KCl, 3; NaHCO_3 , 24; KH_2PO_4 , 1.15; MgSO_4 , 1.15; CaCl_2 , 2; glucose, 10 (330 mosmol l^{-1} , pH 7.35 at 25°C). For experiments with the K^+ channel blocker Ba^{2+} , slices were perfused with an external HEPES-based Ringer solution saturated with O_2 that contained (mM): NaCl, 138; KCl, 3; MgCl_2 , 2; CaCl_2 , 2; glucose, 10; HEPES 10 (330 mosmol l^{-1} , adjusted to pH 7.35). The recording chamber was perfused at a rate of 1.8 ml min^{-1} with the saline solutions and the GABA_A receptor antagonist bicuculline methiodide (10 μM , Sigma Aldrich, St Quentin Fallavier, France) added.

Drugs and solution

All pharmacological agents were applied to cerebellar slices by direct addition to the saline solution, with the exception of pertussis toxin (PTX). In the experiments involving PTX treatment, the cerebellar slices were incubated in the saline solution containing the toxin (2 $\mu\text{g ml}^{-1}$) from 1 h after their preparation and for 14–15 h before the recording session. Stock solutions of ω -agatoxin TK (Sigma Aldrich; Alomone Labs, Israel) ω -conotoxin GVIA (Sigma Aldrich; Alomone Labs), SNX-482 (Alomone Labs) and tertiapin-Q (Sigma Aldrich) were prepared in distilled water. Stock solutions of nifedipine (ICN Pharmaceuticals, Orsay, France) were dissolved in ethanol (final concentration of ethanol 0.1%). All these stock solutions were kept at -20°C until the day of experiment, and individual drugs were diluted in saline solution to their final concentration just before application. PD98059 and WIN55,212-2 were purchased, respectively, from Calbiochem and Tocris (Illkirch, France), SR141716-A was a kind gift from Sanofi-Recherche (Montpellier, France). Stocks of PD98059, WIN55,212-2, SR141716-A and forskolin were prepared in dimethylsulfoxide (DMSO), kept at -20°C until the day of experiment, then added to the saline solution at the desired concentration just before application (final concentration of DMSO 0.1%). After each experiment the perfusion system and the recording chamber were carefully washed with dilute ethanol and distilled water, to avoid any carry-over of cannabinoids. All other compounds were obtained from Sigma (Sigma Aldrich), unless stated otherwise.

Calcium-sensitive fluorometric measurements

In coronal slices, presynaptic parallel fibre (PF) tracts were labelled by local application of a saline solution containing the membrane-permeant form of the fluorescent low affinity calcium-sensitive dye fluo-4FF AM (100 μM , Molecular Probes), as previously described (Daniel & Crepel, 2001). Cerebellar slices were positioned on an upright microscope (Nikon or Zeiss) and calcium-sensitive fluorometric changes were measured through the $\times 40$ or $\times 60$ water-immersion objective. The experiments were conducted at 28°C and started at least 45 min after loading. Optical signals were recorded through a 20 $\mu\text{m} \times 50 \mu\text{m}$ window, placed on the visible narrow band of labelled PFs, approximately 400–700 μm away from the loading site. Parallel fibres located in the recording window were stimulated every minute, with a single 100 Hz train of five to seven electrical stimuli, through a saline-filled glass electrode positioned in the molecular layer between the loading site and the recording site. A confined region of labelled PFs was illuminated with a xenon or mercury lamp. The fluo-4FF filter sets used were 485DF22 (485 \pm 22 nm) for excitation, and DM505 (505 nm) dichroic and 530DF30 (530 \pm 30 nm) for emission. Fluorometric signals were collected with a photometer and analysed on- and off-line using the Acquis1 computer program. The fluorescence data were expressed as $\Delta F/F$, where F is the baseline fluorescence intensity, and ΔF is the change induced by PF stimulation. When an unlabelled region of the slice background fluorescence was greater than 5% of the fluorescence of the indicator, the fluorescence data were corrected for background fluorescence.

Electrophysiology

Whole-cell patch-clamp recordings of PCs from sagittal slices were performed at a somatic level with an Axopatch-1D amplifier (Axon instruments). Patch pipettes (2–3.5 M Ω) were filled with an internal solution containing (mM): KCl, 140; NaCl, 8; Hepes, 10; EGTA, 5, CaCl₂, 0.5, ATP-Mg, 2 (pH 7.3 with KOH; 300 mosmol l⁻¹). As reported previously (Goossens *et al.* 2001), PCs were clamped at -70 mV and PFs were stimulated at 0.33 Hz through an extracellular glass saline-filled monopolar electrode, placed at the surface of the slice to evoke PF-mediated excitatory postsynaptic currents (EPSCs).

Results

Consistent with our previous report (Daniel & Crepel, 2001), a train of five to seven stimulations of PFs

induced reproducible transient increases in presynaptic fluorescence, which returned to resting levels within a few hundred milliseconds (a in Fig. 1A–B). The peak amplitude of the fluorescence transients was $7.8 \pm 0.6\%$ ($\Delta F/F$) (mean \pm s.e.m., $n = 65$).

To test the hypothesis that activation of presynaptic CB1 receptors at PF–PC synapses reduces glutamate release, not only by activation of presynaptic 4-aminopyridine-sensitive K⁺ channels (Daniel & Crepel, 2001), but also by directly inhibiting presynaptic VGCCs, we used specific toxins that target the different Ca²⁺ channels known to participate in the neurotransmitter release process at PF–PC synapses (Mintz *et al.* 1995).

Firstly, using ω -agatoxin TK, which blocks presynaptic P/Q-type Ca²⁺ channels, we investigated the potential involvement of these channels in the cannabinoid-mediated inhibition of presynaptic Ca²⁺ transients evoked by PF stimulation. In accordance with previous data (Mintz *et al.* 1995), a 10 min bath application of ω -agatoxin TK (250 nM) irreversibly inhibited the Ca²⁺-sensitive fluorescence transients elicited by PF, with a resulting mean decrease in amplitude of the fluorescence transients (plateau) of $38.2 \pm 3.7\%$ (mean \pm s.e.m., $n = 4$, Fig. 1A). From this plateau, subsequent application of the CB1 receptor agonist WIN55,212-2 (1 μM) further inhibited presynaptic Ca²⁺ transients with a resulting final mean decrease in amplitude of $63.7 \pm 7.0\%$ ($n = 4$, Fig. 1A). Thus, the WIN55,212-2-mediated reduction of the fluorescence transients, expressed as a percentage of the amplitude of these signals recorded at the plateau level following the ω -agatoxin TK effect, was $\cong 38\%$. This value was not significantly different (Student's t test; $P > 0.6$) of our previously published results in which application of this CB1 agonist alone resulted in a $\cong 33\%$ reduction of the presynaptic fluorescence transients (Daniel & Crepel, 2001). Moreover, the depressant effect of WIN55,212-2 on these transients was reversed by subsequent application of the CB1 receptor antagonist SR141716-A (1 μM) ($n = 4$, Fig. 1A), demonstrating the specific involvement of CB1 receptors.

To study the potential involvement of N-type Ca²⁺ channels in the cannabinoid-mediated inhibition of presynaptic Ca²⁺ transients, we employed ω -conotoxin GVIA, which specifically blocks these Ca²⁺ channels. Again in agreement with the previous report by Mintz *et al.* (1995), a 10 min bath application of ω -conotoxin GVIA (250 nM) irreversibly inhibited presynaptic fluorescence transients by $27.9 \pm 4.7\%$ ($n = 6$, Fig. 1B). Subsequent application WIN55,212-2 (1 μM) further reduced presynaptic Ca²⁺ transients, resulting in a final mean amplitude reduction of $56.7 \pm 2.3\%$ ($n = 6$, Fig. 1B). Here again, the WIN55,

212-2-induced inhibitory effect, expressed as a percentage of the amplitude of the fluorescence signals at the plateau level following ω -conotoxin GVIA effect, was $\cong 37\%$, i.e. a value not significantly different (Student's *t* test; $P > 0.3$) to that observed with WIN55,212-2 application alone (Daniel & Crepel, 2001).

Parallel fibres also possess other VGCCs that control glutamate release and are distinguishable from N and P/Q types, since part of presynaptic Ca^{2+} influx remains unaffected by combined application of ω -Aga-IVA and ω -conotoxin GVIA (Mintz *et al.* 1995). Some of these channels resistant to the aforementioned blockers could

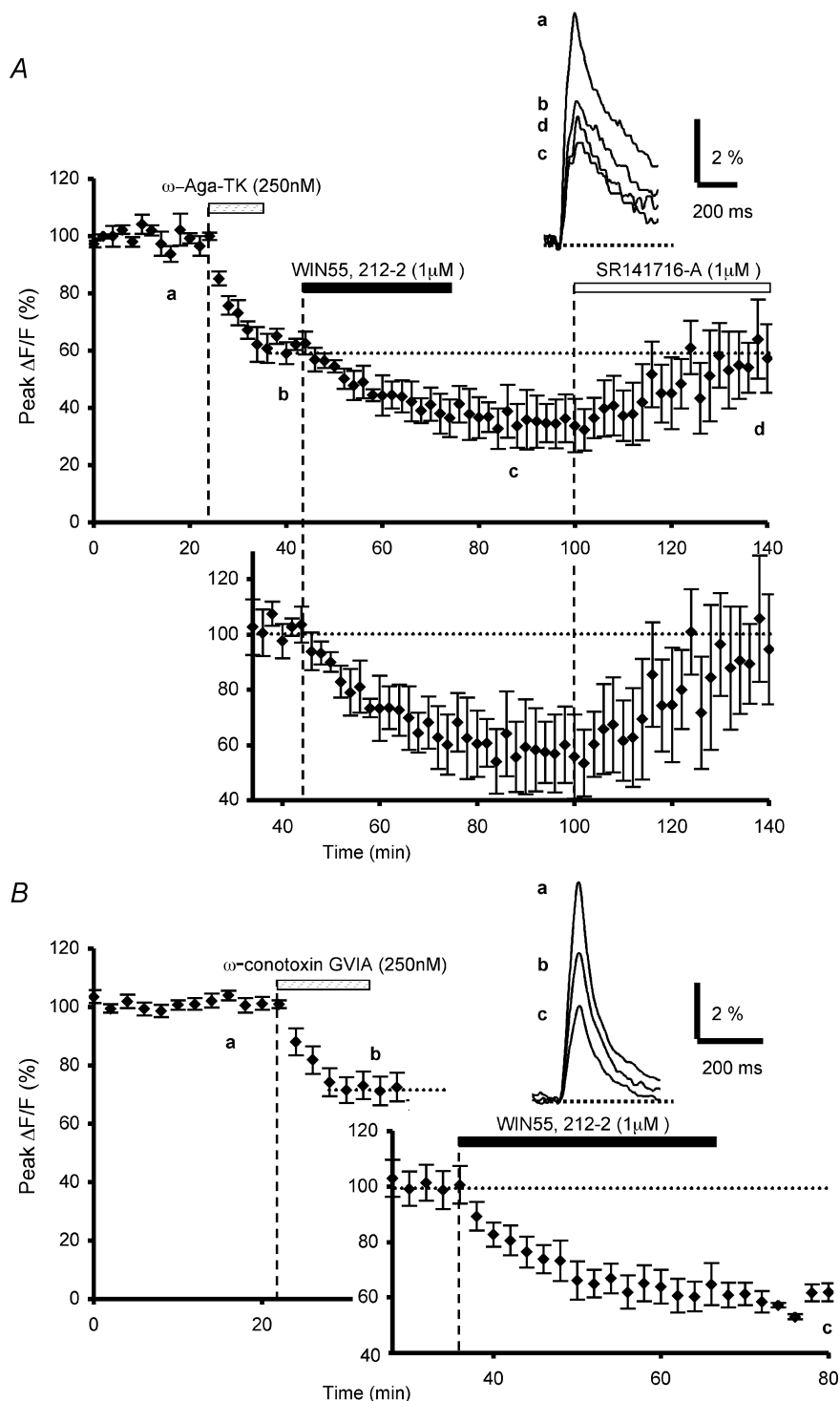


Figure 1. Effects of ω -agatoxin TK or ω -conotoxin GVIA preapplication on WIN55,212-2-mediated inhibition of presynaptic calcium influx

A, the plot represents the normalized amplitudes of peak fluo-4FF fluorescence transients ($\Delta F/F$) evoked by 5–7 stimulations (delivered at 100 Hz) of the parallel fibres recorded as a function of time, before, during and after sequential bath application of ω -agatoxin TK (250 nM) and WIN55,212-2 (1 μ M). Each point is the mean \pm s.e.m. of 4 separate experiments. Note that the inhibitory effect of WIN55,212-2 (1 μ M) on the normalized amplitude of peak fluorescence transients was reversed by bath application of the selective CB1 antagonist SR141716-A (1 μ M, $n = 4$). The top inset displays superimposed averaged fluorescence changes in one of these experiments, recorded at the indicated times. Each trace is an average of 5–10 consecutive trials. The bottom inset (graph) shows the WIN55,212-2-mediated reduction of the fluorescence transients normalized to the plateau level obtained after application of ω -agatoxin TK. *B*, the same as in *A* with sequential application of ω -conotoxin GVIA (250 nM) and WIN55,212-2 (1 μ M) ($n = 6$). Insets as in *A*. Note that preapplication of ω -agatoxin TK or ω -conotoxin GVIA does not prevent the WIN55,212-2-mediated depressant effect on presynaptic Ca^{2+} influx.

be R-type Ca²⁺ channels (Ellinor *et al.* 1993; Zhang *et al.* 1993). While these channels have not been formally identified on PFs, their existence has been hypothesized by Randall & Tsien (1995) and demonstrated by Tottene *et al.* (2000) in rat cerebellar granule neurones in primary culture. In these neurones, the R-type Ca²⁺ channels are all encoded by the pore-forming α_{1-E} subunit gene (Soong *et al.* 1993), probably with different isoforms, which could explain their different pharmacological properties (Tottene *et al.* 2000). Using the recently developed polypeptide toxin SNX-482, a selective antagonist of the α_{1-E} subunit (Newcomb *et al.* 1998), Tottene *et al.* (2000) have shown that some of the native R-type Ca²⁺ channels recorded in the soma of cerebellar granule cells are sensitive to this toxin, and some are resistant. Ten minute bath applications of SNX-482 (100 nM) significantly reduced

the presynaptic fluorescence transients by $17.5 \pm 4.4\%$ ($n = 3$, Fig. 2A), suggesting that R-type (SNX-482-sensitive) VGCCs are present on PFs. However, this toxin did not abolish the subsequent decrease of fluorescent transients induced by WIN55,212-2. Indeed, the resulting mean decrease in amplitude of these transients was $55.3 \pm 9.1\%$ after sequential application of SNX-482 and WIN55,212-2 ($n = 3$, Fig. 2A). Thus, the WIN55,212-2-induced inhibitory effect, expressed as a percentage of the amplitude of the fluorescence signals at the plateau level following SNX-482 application, was $\cong 34\%$, a value not significantly different to that observed in control conditions (Student's *t* test; $P > 0.8$) (see Daniel & Crepel, 2001) and after application of ω -agatoxin TK (Student's *t* test; $P > 0.7$) or ω -conotoxin GVIA (Student's *t* test; $P > 0.5$) (see above).

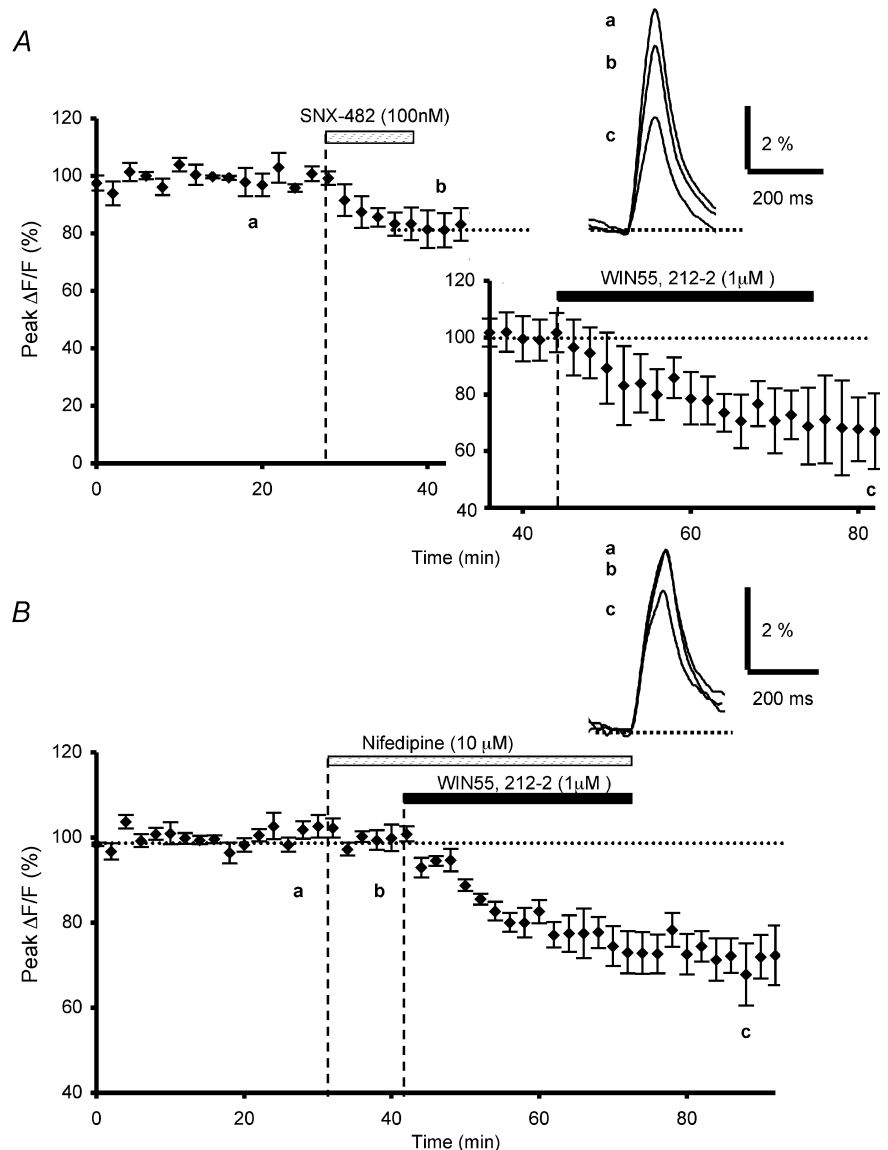


Figure 2. Effects of SNX-482 or nifedipine preapplication on WIN55,212-2-mediated inhibition of presynaptic calcium influx

A, time course of fluo-4FF fluorescence transients before, during and after sequential bath application of SNX-482 (100 nM) and WIN55,212-2 (1 μ M) ($n = 3$). The insets show fluorescence transients in one of these experiments and the WIN55,212-2-mediated reduction of the fluorescence transients normalized to the plateau level obtained after application of SNX-482. Note that preapplication of SNX-482 does not prevent the WIN55,212-2-mediated depressant effect on presynaptic Ca²⁺ influx. **B**, the same as in **A** with application of nifedipine (10 μ M) and WIN55,212-2 (1 μ M) ($n = 5$). Insets as in **A**. Note that application of nifedipine does not alter the amplitude of presynaptic Ca²⁺ influx and does not prevent the WIN55,212-2-mediated depressant effect on Ca²⁺ influx.

Finally, in keeping with previous results by Mintz *et al.* (1995), presynaptic fluorescence transients evoked by PF stimulation were unaffected ($n = 5$, Fig. 2B) by a 10 min bath application of the dihydropyridine derivative nifedipine ($10 \mu\text{M}$), a blocker of L-type Ca^{2+} channels (Rampe & Triggle, 1990). This confirms that these channels are not involved in the glutamate release process at PF–PC synapses, even if they are present on cerebellar granule neurones in culture (Randall & Tsien, 1995; Pearson *et al.* 1995). In addition, in the presence of this toxin, the WIN55,212-2-mediated inhibitory effect on presynaptic fluorescence signals was $\cong 29\%$ ($n = 5$, Fig. 2B), a value not significantly different (Student's *t* test; $P > 0.5$) to that observed in control conditions (see Daniel & Crepel, 2001), demonstrating that treatment with an L-type Ca^{2+} channel blocker did not alter the cannabinoid-mediated depressant effect.

We also investigated possible cross-interactions between CB1 receptors and the different presynaptic Ca^{2+} channels by combined toxin treatment (ω -agatoxin TK (250 nM), ω -conotoxin GVIA (250 nM) and SNX-482 (100 nM)), followed by addition of WIN55,212-2. Co-application of these blockers decreased fluorescence transients by $70.1 \pm 1.2\%$ ($n = 4$, Fig. 3). Subsequent application WIN55,212-2 ($1 \mu\text{M}$) further reduced presynaptic Ca^{2+} transients, with a resulting final mean

decrease in amplitude of $79.5 \pm 1.4\%$ ($n = 4$, Fig. 3). Thus, the WIN55,212-2-induced inhibitory effect expressed as a percentage of the amplitude of the fluorescence signals at the plateau level reached with the combined blockers, was $\cong 32\%$, a value not significantly different to that observed in control conditions (Student's *t* test; $P > 0.8$) (see Daniel & Crepel, 2001) and after application of ω -agatoxin TK (Student's *t* test; $P > 0.5$), or ω -conotoxin GVIA (Student's *t* test; $P > 0.1$), or SNX-482 (Student's *t* test; $P > 0.7$) alone (see above).

Taken together, these data demonstrate that the CB1 receptor-mediated depression of presynaptic Ca^{2+} influx is not due to a direct inhibition of presynaptic P/Q-, N- and R (SNX-482-sensitive)-type VGCCs.

In an attempt to determine whether the intracellular signals mediating the WIN55,212-2-induced inhibition of glutamate release required the activation of the G_i/G_o subclass of G proteins, slices were treated with PTX ($2 \mu\text{g ml}^{-1}$) for 14–15 h before the recording session (see Methods). In these conditions, WIN55,212-2 application had no discernable effect on fluorescence transients, since, on average, 20 min after application of this cannabinoid agonist, the decrease in amplitude of the fluorescence transients was $4.1 \pm 5.1\%$ ($n = 5$, Fig. 4A). In contrast, control slices incubated in the same experimental conditions for an equivalent period

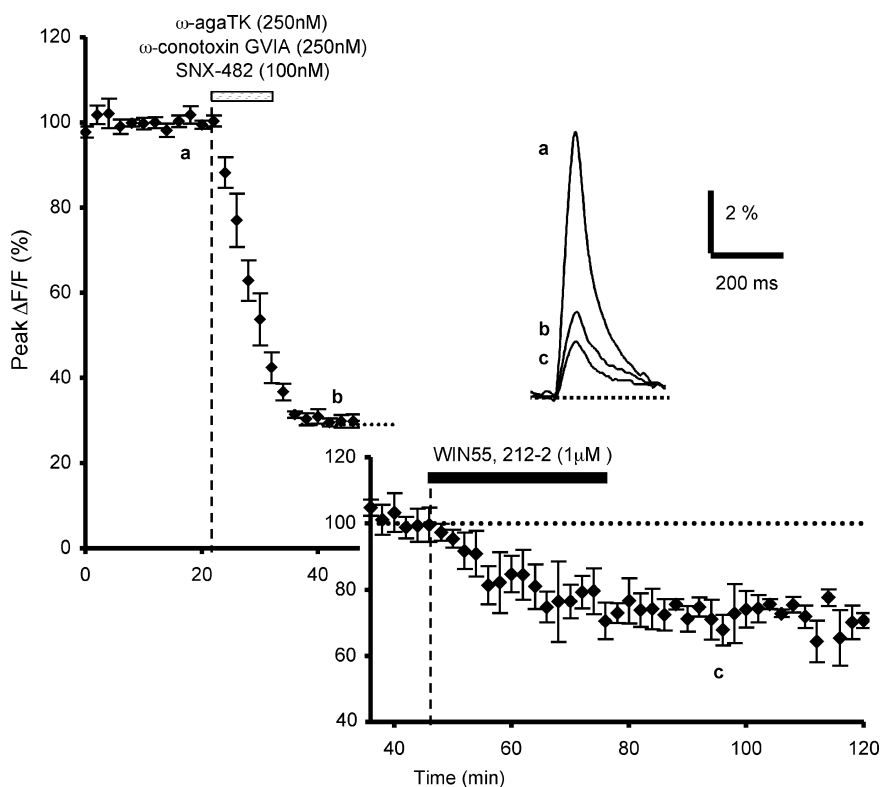


Figure 3. Effects of preapplication of combined toxin treatment on WIN55,212-2-mediated inhibition of presynaptic calcium influx

Time course of fluo-4FF fluorescence transients before, during and after bath application of combined blockers (ω -agatoxin TK (250 nM), ω -conotoxin GVIA (250 nM) and SNX-482 (100 nM)) followed by application of WIN55,212-2 ($1 \mu\text{M}$) ($n = 4$). The insets show the fluorescence transients in one of these experiments (top) and the WIN55,212-2-mediated reduction of the fluorescence transients normalized to the plateau level obtained after combined toxin treatment (bottom).

of time, but without PTX, showed a clear WIN55,212-2-mediated decrease in the amplitude of fluorescence transients, averaging $29.7 \pm 5.9\%$ ($n = 3$, Fig. 4B). Thus, under these conditions, the magnitude of the WIN55,212-2-mediated inhibitory effect ($\cong 30\%$), was not significantly different (Student's t test; $P > 0.4$) to that observed when the interval between slice preparation and the recording session was around 3–6 h (see Daniel & Crepel, 2001). Moreover, in the present experiments, the

peak amplitude of the fluorescence transients recorded in control slices incubated for long periods of time ($8.0 \pm 1.9\%$ ($\Delta F/F$), $n = 3$), was not significantly different from those recorded in control slices without incubation ($8.2 \pm 0.7\%$ ($\Delta F/F$), $n = 46$) (Student's t test; $P < 0.01$). Therefore, these data demonstrate that the inhibitory action of WIN55,212-2 on presynaptic Ca^{2+} transients is mediated exclusively through the G_i/G_o subclass of G proteins.

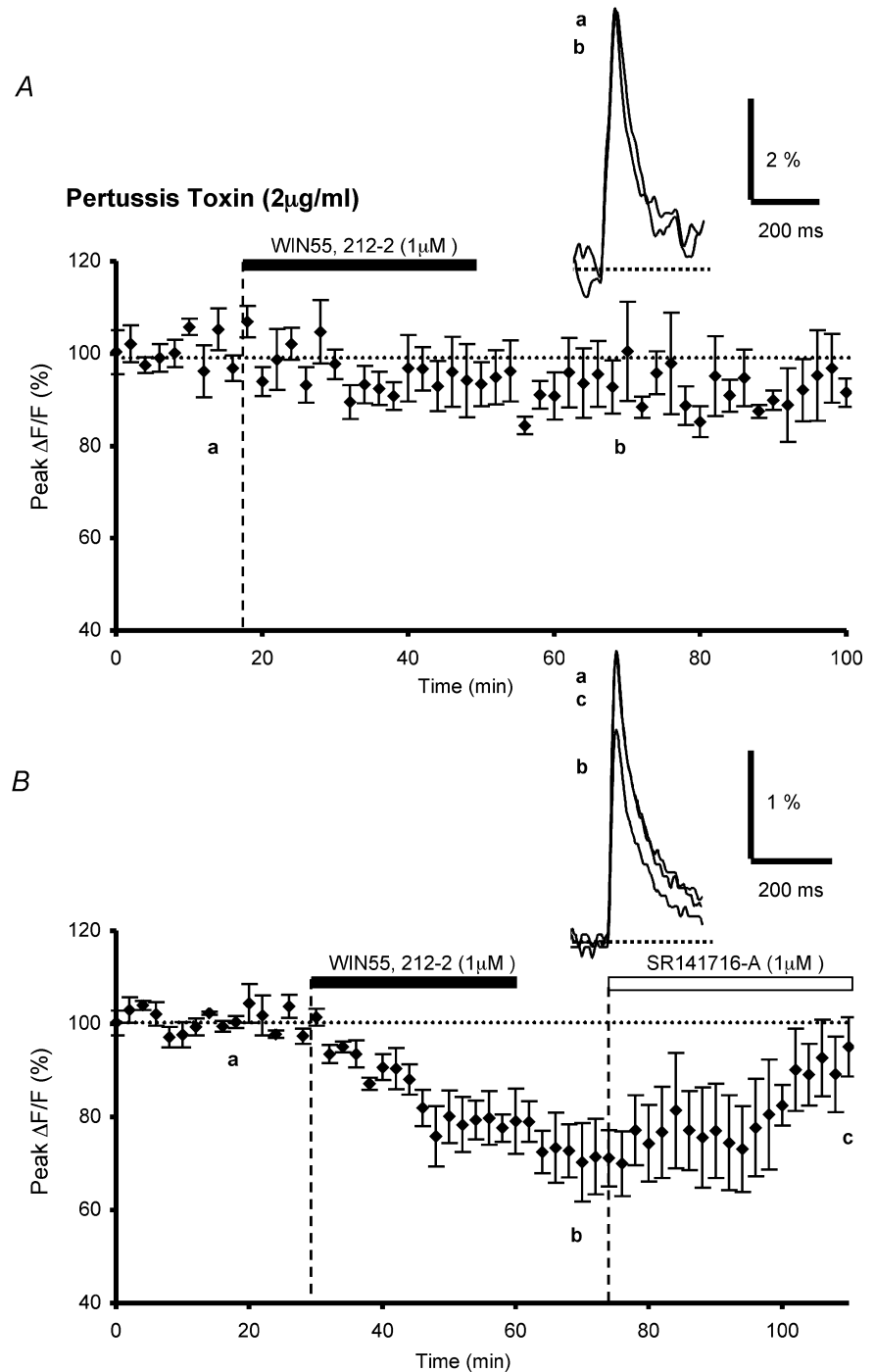


Figure 4. Effects of pertussis toxin treatment on WIN55,212-2-mediated inhibition of presynaptic calcium influx

A, plot of normalized amplitudes of peak fluorescence transients before, during and after bath application of WIN55,212-2 (1 μM) ($n = 5$). Experiments were performed with slices preincubated with pertussis toxin (2 $\mu\text{g ml}^{-1}$) during 14–15 h. Note that PTX preincubation prevents the WIN55,212-2-induced inhibition of peak fluorescence transients. The insets represent superimposed averaged fluorescence changes in one of these experiments. **B**, the same as in **A** with slices incubated in the same experimental conditions for an equivalent period of time, but without PTX ($n = 3$). Note the clear inhibitory effect of WIN55,212-2 on the fluorescence transients and reversal of the effect by bath application of the selective CB1 antagonist SR141716-A (1 μM , $n = 4$). Inset as in **A**.

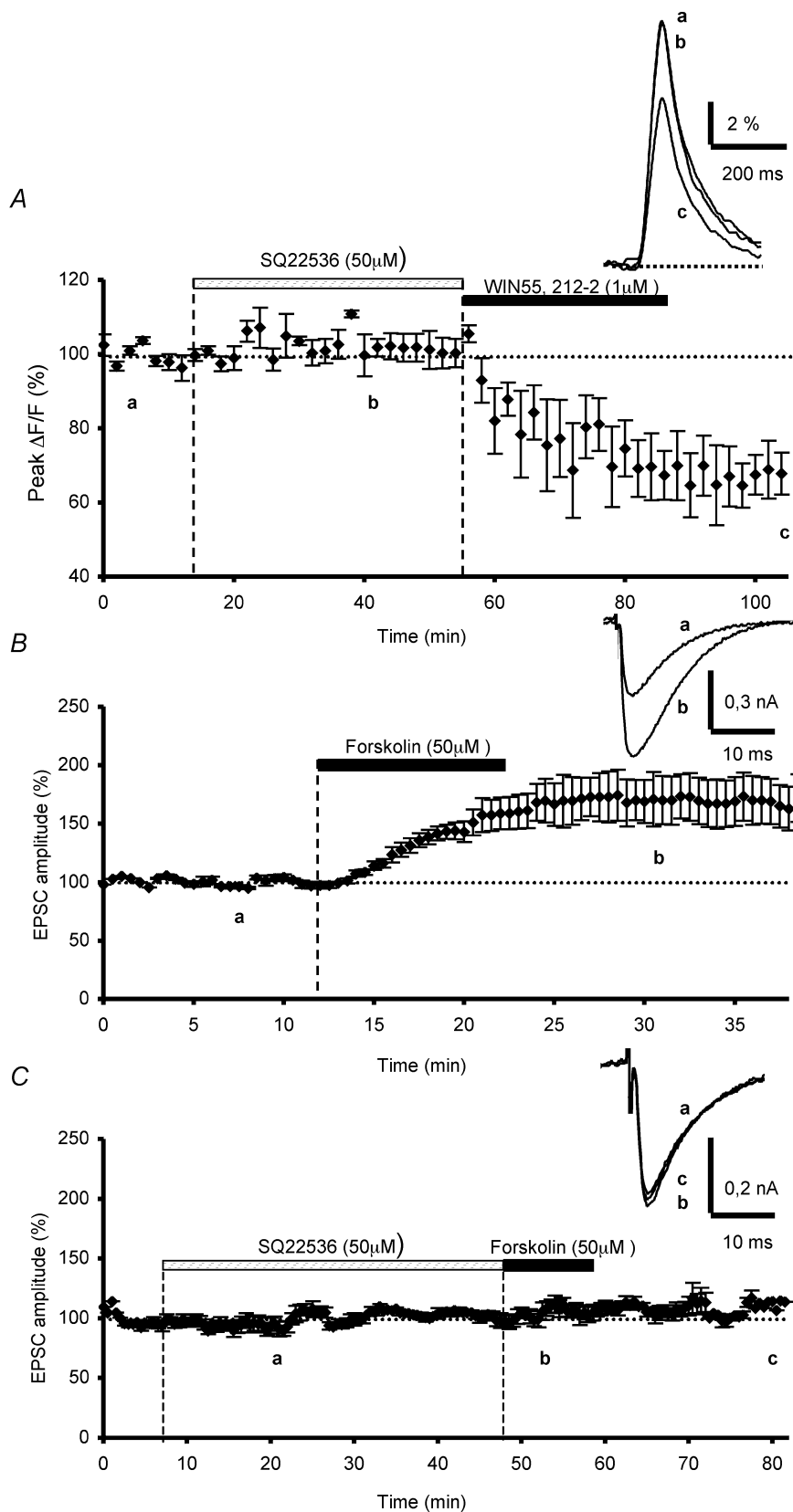


Figure 5. Effects of adenylate cyclase blockade on WIN55,212-2-mediated inhibition of presynaptic calcium influx and on forskolin-induced enhancement of parallel fibre synaptic responses

A, plot of normalized amplitudes of peak fluorescence transients before, during and after bath sequential application of SQ 22,536 (50 μ M) and WIN55,212-2 (1 μ M) ($n = 4$). Note that in these experiments, the adenylate cyclase inhibitor did not prevent the WIN55,212-2-induced inhibition of peak fluorescence transients. The insets represent superimposed averaged fluorescence changes in one of these experiments. *B*, plot of normalized amplitudes of PF-mediated EPSCs as a function of time before, during and after bath application of forskolin (50 μ M) ($n = 4$). Note the large enduring increase of PF-mediated EPSCs amplitude. Inset, superimposed averaged PF-mediated EPSCs recorded in one of these experiments. Each trace is an average of 10 consecutive trials. *C*, prevention of the forskolin-induced long-term enhancement of the normalized amplitudes of PF-mediated EPSCs by prior bath application SQ 22,536 (50 μ M) ($n = 5$).

To address the question of a possible participation of adenylate cyclase in the CB1 receptor-mediated depression of presynaptic Ca^{2+} influx, we carried out experiments with SQ 22,536, an analogue of adenosine that appears to be a broad-spectrum blocker of adenylate cyclase (Harris *et al.* 1979). A 40 min bath application of SQ 22,536 prior to WIN55,212-2 application, had no effect on control fluorescence transients evoked by PF stimulation ($n = 4$, Fig. 5A). Moreover, this inhibitor did not prevent the subsequent inhibitory effect of WIN55,212-2 on these transients, since the cannabinoid agonist inhibition of fluorescence transients was $35.2 \pm 6.6\%$ on average ($n = 4$, Fig. 5A), which was not significantly different (Student's *t* test; $P > 0.7$) to the WIN55,212-2-mediated inhibitory effect in control conditions (see Daniel & Crepel, 2001).

As a control, we performed additional electrophysiological experiments to test the efficiency of this pharmacological inhibitor of adenylate cyclase in our conditions. At PF-PC synapses, elevation of cAMP levels by forskolin, an activator of adenylate cyclase, has been shown to durably enhance neurotransmitter release (Salin *et al.* 1996; Chen & Regher, 1997), through a presynaptic mechanism that does not alter presynaptic Ca^{2+} influx or resting Ca^{2+} levels, but directly increases the probability of vesicular release (Chen & Regher, 1997). Thus, we reasoned that this long-term enhancement of synaptic strength involving adenylate cyclase, downstream from Ca^{2+} influx, might provide a sensitive means to test the efficiency of the pharmacological inhibitor SQ 22,536. According to previous reports (Salin *et al.* 1996; Chen & Regher, 1997), a clear enhancement in the amplitude of PF-mediated

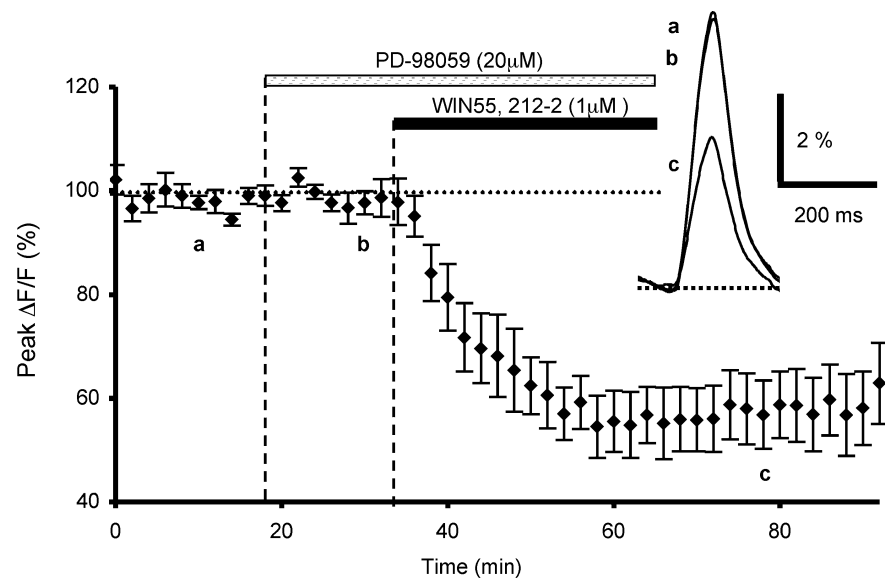
EPSCs was seen following 10 min bath applications of $50 \mu\text{M}$ forskolin, averaging $164.6 \pm 16.6\%$ ($n = 4$, Fig. 5B). In contrast, preapplication of SQ 22,536 ($50 \mu\text{M}$) for 40 min, rendered forskolin ineffective in enhancing PF-mediated responses ($n = 5$, Fig. 5C), demonstrating that in our experimental conditions, adenylate cyclase was totally blocked. Taken together, these data establish that an adenylate cyclase-coupled signalling pathway does not mediate the depressant effect of WIN55,212-2 on presynaptic Ca^{2+} influx.

In another set of experiments, we examined the possible involvement of MAPKs in the CB1 receptor-mediated depression of presynaptic Ca^{2+} influx. We used PD98059, a cell-permeable specific inhibitor of MAPKs (Alessi *et al.* 1995). PD98059 ($20 \mu\text{M}$) was bath applied 15 min prior to and during WIN55,212-2 application. This inhibitor affected neither control fluorescence transients evoked by PF stimulation ($n = 6$, Fig. 6), nor the subsequent depressant effect of WIN55,212-2 on these transients, since the cannabinoid agonist inhibition of fluorescence transients was $40.9 \pm 5.8\%$ on average ($n = 6$, Fig. 6), which was not significantly different (Student's *t* test; $P > 0.2$) to the WIN55,212-2-mediated inhibitory effect in control conditions (see Daniel & Crepel, 2001).

Finally, the actions of K^+ channels blockers on the CB1 receptor-mediated inhibition of presynaptic fluorescence transients elicited by PF stimulations, were assessed. We previously found that WIN55,212-2 no longer had any effect on these transients, when presynaptic K^+ channels were blocked by application of 1 mM 4-aminopyridine (4-AP) (Daniel & Crepel, 2001). Since, this concentration of 4-AP is fairly unselective (Coetzee *et al.* 1999),

Figure 6. Effects of MAPK on WIN55,212-2-mediated inhibition of presynaptic calcium influx

Plot of normalized amplitudes of peak fluorescence transients before, during bath application of PD-98059 ($20 \mu\text{M}$) and coapplication with WIN55,212-2 ($1 \mu\text{M}$), and after this combined treatment ($n = 6$). Note that in these experiments, the MAPK inhibitor does not prevent the WIN55,212-2-induced inhibition of peak fluorescence transients. The insets represent superimposed averaged fluorescence changes in one of these experiments



we performed new experiments with this blocker at non-saturating doses ($200 \mu\text{M}$). As previously described (Daniel & Crepel, 2001), bath application of 4-AP alone increased the amplitude of presynaptic fluorescence transients evoked by PF stimulation compared to those recorded in control conditions, since the resulting mean increase on the amplitude of these transients with $200 \mu\text{M}$ of this blocker was $625.8 \pm 37.1\%$ ($n = 8$, not illustrated; see Daniel & Crepel, 2001). Thus, in the presence of $200 \mu\text{M}$ 4-AP, to reduce the amplitude of these transients to values similar to those recorded in control medium, the extracellular Ca^{2+} concentration was lowered from 2 to 0.2 mM , with corresponding changes in the Mg^{2+} concentration to maintain the osmolarity ($n = 8$, not illustrated). In such conditions, when the amplitude of the fluorescence transients was brought back to control levels, the depressant effect of WIN55,212-2 on these transients was $20.4 \pm 0.6\%$ on average ($n = 4$, Fig. 7A), i.e. a value significantly (Student's *t* test; $P < 0,05$) smaller than that observed with WIN55,212-2 in control experiments. Thus, the present experiments show that the WIN55,212-2-mediated inhibitory effect on presynaptic Ca^{2+} influx is partially blocked by $200 \mu\text{M}$ 4-AP.

We then tested the hypothesis that CB1 receptor-mediated inhibition of presynaptic Ca^{2+} influx elicited by PF stimulation involves the activation of G protein-gated inwardly rectifying K^{+} channels (GIRKs) by determining whether the inhibitory effect of WIN55,212-2 was modulated by Ba^{2+} or tertiapin-Q, two blockers of this channel family, even though it is known that Ba^{2+} also affects other K^{+} channels and that tertiapin only affects some subtypes of GIRKs (Slesinger *et al.* 1997; McAllister *et al.* 1999; Jin *et al.* 1999; Jin & Lu, 1999; Takigawa & Alzheimer, 2002). In these experiments, bath application of $300 \mu\text{M}$ Ba^{2+} , a non-selective blocker of GIRKs, did not appreciably alter the absolute fluorescence, suggesting that it does not seem to sizeably affect the basal Ca^{2+} level concentration in PFs ($n = 5$). Moreover, the amplitude and the duration of presynaptic fluorescence transients evoked by PF stimulations were unaffected by this blocker, compared to those recorded in control conditions ($n = 5$, Fig. 7B). In contrast; in the presence of $300 \mu\text{M}$ Ba^{2+} , the depressant effect of WIN55,212-2 on fluorescence transients was only $21.8 \pm 2.1\%$ on average ($n = 5$, Fig. 7B), i.e. a value significantly (Student's *t* test; $P < 0,05$) smaller than that observed with WIN55,212-2 in control conditions (see Daniel & Crepel, 2001). These experiments demonstrate the blocking action of Ba^{2+} on the depressant effect of WIN55,212-2 on presynaptic Ca^{2+} influx. Since even low concentrations of Ba^{2+} affect various types of K^{+} channels in addition to GIRKs

(see Discussion), we subsequently employed tertiapin-Q, which selectively inhibits some subtypes of this K^{+} channel family with nanomolar affinity. A 20 min bath application of tertiapin-Q (100 nM), prior to WIN55,212-2 application, had no effect on control fluorescence transients evoked by PF stimulation ($n = 6$, Fig. 8A). In contrast, in the presence of this blocker, the depressant effect of WIN55,212-2 on fluorescence transients was only $19.9 \pm 1.9\%$ on average ($n = 6$, Fig. 8A), i.e. a value significantly (Student's *t* test; $P < 0,02$) smaller than that observed with WIN55,212-2 in control experiments (see Daniel & Crepel, 2001).

Finally, we tested whether combined application of tertiapin-Q and 4AP would produce a larger reduction of the WIN55,212-2-induced inhibition on fluorescence transients, than the reduction observed with either blocker applied alone. When 4-AP (used at non-saturating doses of $200 \mu\text{M}$ with reduced extracellular Ca^{2+} concentration) and tertiapin-Q (100 nM) were applied together, the depressant effect of WIN55,212-2 on the presynaptic fluorescence transients was only $12.9 \pm 3.7\%$ on average ($n = 4$, Fig. 8B). Together, these data strongly suggest that the inhibitory action of WIN55,212-2 on presynaptic Ca^{2+} transients can be attributed to the activation of 4-AP-sensitive and tertiapine-Q-sensitive K^{+} channels.

Discussion

The primary finding of this study, which extends earlier work on the mechanisms of cannabinoid receptor-mediated inhibition of excitatory synaptic transmission at PF-PC synapses (Levenes *et al.* 1998; Daniel & Crepel, 2001), is that cannabinoids do not directly inhibit the two major presynaptic N- and P/Q-type VGCCs required for evoked neurotransmitter release at these synapses (Mintz *et al.* 1995). Cannabinoids have been found to inhibit N-type and/or P/Q-type VGCCs in non-neuronal cell lines (Mackie & Hille, 1992), transfected cells (Mackie *et al.* 1995), cultured hippocampal neurones (Twitchell *et al.* 1997; Sullivan, 1999) and primary cultures of cerebellar granule neurones (Nogueron *et al.* 2001). Since, in brain slice preparations, our findings show that the activation of CB1 receptors by a cannabinoid agonist does not interfere with presynaptic VGCCs, it cannot be excluded that the signal transduction events linked to cannabinoid receptor activation in cell lines or primary cultures may be different. Nevertheless, the molecular mechanisms underlying the cannabinoid-mediated effects studied in brain slice preparations depend on the structure or neuronal cell type studied. For instance, presynaptic CB1 receptor-mediated depression of glutamatergic synaptic inputs involves the

inhibition of N-type but not P/Q type Ca²⁺ channels in the striatum (Huang *et al.* 2001), whereas such direct inhibition of VGCCs is absent in the nucleus accumbens (Robbe *et al.* 2001) and lateral amygdala (Azad *et al.* 2003), as here in the cerebellar cortex. In contrast, blockade of N-type Ca²⁺ channels (Wilson *et al.* 2001) or blockade

of all subtypes of VGCCs by cadmium a non-selective blocker (Hoffman & Lupica, 2000) abolishes the CB1-mediated inhibition of GABAergic neurotransmission in the hippocampus.

In addition, the present experiments show that application of SNX-482, a blocker of a part

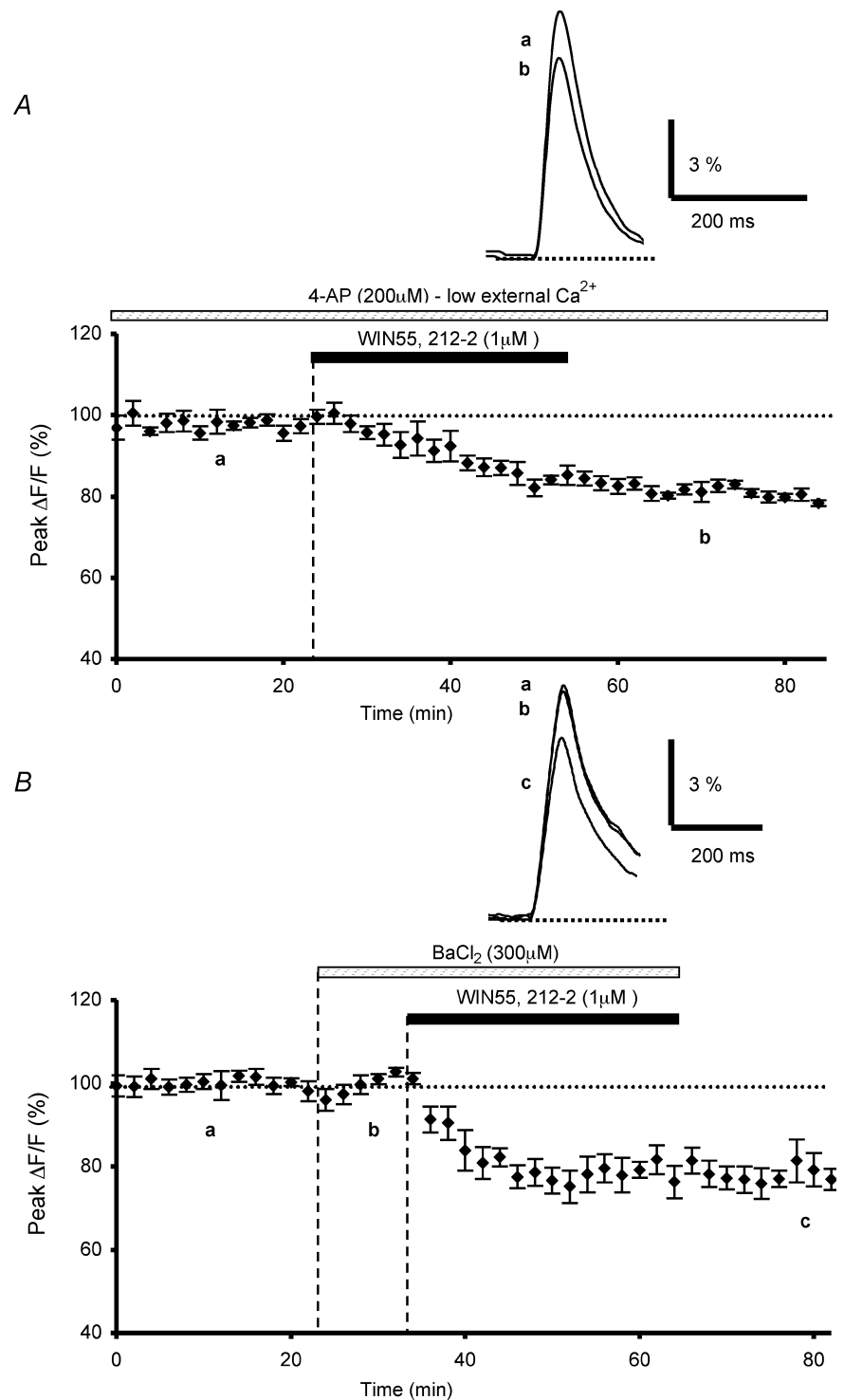


Figure 7. Effects of K⁺ channel blockers 4-AP and Ba²⁺ on WIN55,212-2-mediated inhibition of presynaptic calcium influx

A, plot of normalized amplitudes of peak fluorescence transients before, during and after application of WIN55,212-2 (1 μM), in the presence of 4-AP (200 μM) with low extracellular calcium concentration (0.2 mM) (*n* = 4). Note that in these experiments, bath-applied 4-AP partially prevents the WIN55,212-2-induced inhibition of peak fluorescence transients. The insets represent superimposed averaged fluorescence changes in one of these experiments. **B**, the same as in **A** with applications of Ba²⁺ (300 μM) and WIN55,212-2 (1 μM) (*n* = 5). Inset as in **A**. Note that this K⁺ channel blocker also partially prevents the WIN55,212-2-induced inhibition of peak fluorescence transients.

of R-type Ca^{2+} channels, is able to reduce the amplitude of presynaptic Ca^{2+} transients evoked by PF stimulation. This observation strongly supports the fact that PFs bear not only N- and P/Q-type Ca^{2+} channels, but also R-type (SNX-482-sensitive) Ca^{2+} channels, which could contribute to fast excitatory synaptic transmission in rat cerebellar cortex, as previously demonstrated in rat

hippocampus (Gasparini *et al.* 2001). Interestingly, our observation demonstrates that these channels are not involved in the cannabinoid receptor-mediated inhibition of presynaptic Ca^{2+} transients.

Taken together, our data demonstrate that the direct inhibition of presynaptic P/Q-, N- and R (SNX-482 sensitive)-type VGCCs is not involved in cannabinoid

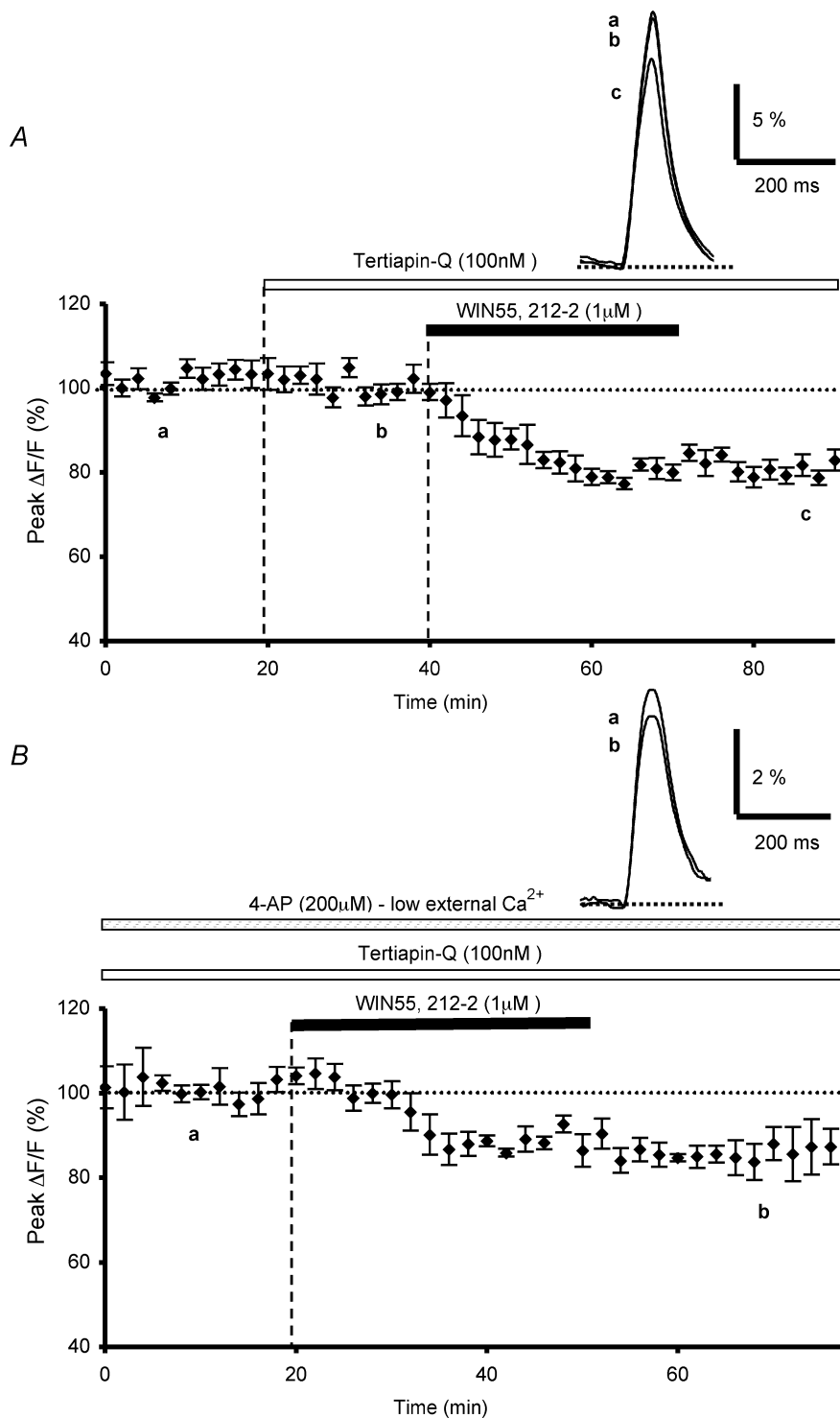


Figure 8. Effects of K^{+} channel blocker tertiapin-Q alone or combined with 4-AP on WIN55,212-2-mediated inhibition of presynaptic calcium influx

A, plot of normalized amplitudes of peak fluorescence transients before, during bath application of tertiapin Q (100 nM) and coapplication with WIN55,212-2 (1 μM), and after washout of WIN 55, 212-2 ($n = 6$). Note that this K^{+} channel blocker partially prevents the WIN55,212-2-induced inhibition of peak fluorescence transients. The inset represents superimposed averaged fluorescence changes in one of these experiments. *B*, plot of normalized amplitudes of peak fluorescence transients before, during and after application of WIN55,212-2 (1 μM), in the continuous presence of bath applied tertiapin-Q (100 nM) and 4-AP (200 μM) with a low extracellular calcium concentration (0.2 mM) ($n = 4$). Inset as in *A*. Note that in these conditions, tertiapin-Q and 4-AP drastically reduce the WIN55,212-2-induced inhibition of peak fluorescence transients.

receptor-mediated inhibition of excitatory synaptic transmission at PF–PC synapses.

Our finding that PTX pretreatment abolishes the WIN55,212-2-mediated inhibitory effect on presynaptic Ca²⁺ influx evoked by PF stimulation indicates that CB1 receptor activation inhibits glutamatergic synaptic transmission at PF–PC synapses through a PTX-sensitive G_i/G_o subclass of G proteins. A well-documented action of cannabinoids mediated through such proteins, is the inhibition of adenylate cyclase activity. Such inhibition has been described in many different cell types (Holwett, 1995; Childers & Deadwyler, 1996; Shen *et al.* 1996), including cultured cerebellar granule cells (Pacheco *et al.* 1993; Jung *et al.* 1997). Using a pharmacological tool inhibiting this cyclase in cerebellar slice preparations, we have presented evidence that CB1 activation is unlikely to decrease synaptic transmission at PF–PC synapses through modulation of adenylate cyclase activity. This finding agrees with previous reports showing that in nucleus accumbens (Robbe *et al.* 2001) or in hippocampal slices (Wilson *et al.* 2001) activation of presynaptic CB1 receptors inhibits, respectively, glutamate or GABA release, independently of an adenylate cyclase-coupled signalling pathway. Since the inhibition of adenylate cyclase activity is one of the predominant effects of cannabinoid receptor activation in primary cultures of neurones, and particularly in cultured cerebellar granule cells (Pacheco *et al.* 1993; Jung *et al.* 1997), it cannot be excluded that these cultured cells may develop their own intracellular signal transduction pathways linked to cannabinoid receptors. However, another possibility is that the activation of cannabinoid receptors in cerebellar slices effectively leads to inhibition of adenylate cyclase activity in PFs, but as proposed, this molecular mechanism is independent of the signalling pathway underlying the control of synaptic transmission at PF–PC synapses.

Activation of MAPK, mediated through a PTX-sensitive G protein, is another well-established mechanism induced by CB1 activation in heterologous expression systems (Bouaboula *et al.* 1995, 1999) and in hippocampal slices (Derkinderen *et al.* 2001, 2003). However, in the present study, interfering with MAPK activity did not affect the depressant effect of WIN55,212-2 on presynaptic Ca²⁺ influx, indicating that the MAPK pathway is not involved in the CB1 receptor-mediated decrease of excitatory synaptic transmission at PF–PC synapses.

Finally, although the molecular mechanisms downstream of the PTX-sensitive G_i/G_o proteins remain to be elucidated, the present study demonstrates that the CB1-mediated inhibition of synaptic transmission at PF–PC synapses requires activation of GIRKs. However,

although such a functional coupling between this subtype of G protein-activated K⁺ channels (sensitive to PTX and inhibited by Ba²⁺ ions) and the GABA_B receptor has previously been demonstrated in cerebellar granule cells (Slesinger *et al.* 1997), the exact identity of the K⁺ channels underlying the CB1-mediated inhibition of presynaptic Ca²⁺ transients remained to be determined. Indeed, the K⁺ channel blocker Ba²⁺, even at low concentrations, not only inhibits GIRKs, but is also known to affect Ca²⁺-activated K⁺ channels (Katz *et al.* 1995), delayed rectifier K⁺ channels, K⁺ channels closed by cytosolic ATP (K_{ATP} channels) (Hille, 1992) and two-pore-domain K⁺ channels (Lesage & Lazdunski, 2000; Han *et al.* 2002). Even though Ba²⁺ is not a highly selective blocker of GIRKs, our data obtained with tertiapin-Q, a selective inhibitor of some subtypes of this K⁺ channel family, demonstrates that the activation of these channels participates in the reduction of glutamate release by cannabinoids. Moreover, immunohistological and *in situ* hybridization studies have demonstrated that GIRK subunits are abundant in the cerebellar cortex, where they are concentrated primarily in the granule cell layer but are also found in the molecular layer (Liao *et al.* 1996; Ponce *et al.* 1996; Karschin *et al.* 1996). However, their precise location on the granule cell axons remains to be established. In addition, the present data together with our previous study (Daniel & Crepel, 2001), demonstrate the participation of 4-AP-sensitive K⁺ channels in cannabinoid receptor-mediated inhibition of excitatory synaptic transmission at PF–PC synapses. Nevertheless, because the selectivity of 4-AP is not high, even when it is used at low concentrations (Mathie *et al.* 1998), one cannot unambiguously attribute the inhibition observed with this blocker to a particular K⁺ channel subtype. Further studies to determine which subtypes of 4-AP-sensitive K⁺ channels are involved in this inhibition would be worthwhile. Taken together, our results show that the activation of presynaptic CB1 receptors inhibits glutamatergic transmission via voltage-dependent K⁺ channels (4-AP-sensitive) and G protein-gated inwardly rectifying K⁺ channels (Ba²⁺- and tertiapin-Q-sensitive); an effect that is also found in other brain areas (Robbe *et al.* 2001; Azad *et al.* 2003).

In conclusion, using the exogenous cannabinoid agonist WIN55,212-2, the present study identifies presynaptic targets which are involved in CB1 receptor-mediated acute inhibition of neurotransmitter release at PF–PC synapses. Further studies will be necessary to show whether the retrograde regulation of excitatory PF–PC synaptic transmission by endogenous cannabinoids (Kreitzer & Regehr, 2001) shares similar molecular mechanisms. It also remains to be determined to what extent the

effects of cannabinoids on cerebellar function, leading to impairment of motor performances, are attributable to such modulation of synaptic transmission.

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