

## Inhibition of Rho-associated kinase blocks agonist-induced $\text{Ca}^{2+}$ sensitization of myosin phosphorylation and force in guinea-pig ileum

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1.  $\text{Ca}^{2+}$  sensitization of smooth muscle contraction involves the small GTPase RhoA, inhibition of myosin light chain phosphatase (MLCP) and enhanced myosin regulatory light chain ( $\text{LC}_{20}$ ) phosphorylation. A potential effector of RhoA is Rho-associated kinase (ROK).
2. The role of ROK in  $\text{Ca}^{2+}$  sensitization was investigated in guinea-pig ileum.
3. Contraction of permeabilized muscle strips induced by  $\text{GTP}\gamma\text{S}$  at pCa 6.5 was inhibited by the kinase inhibitors Y-27632, HA1077 and H-7 with  $\text{IC}_{50}$  values that correlated with the known  $K_i$  values for inhibition of ROK.  $\text{GTP}\gamma\text{S}$  also increased  $\text{LC}_{20}$  phosphorylation and this was prevented by HA1077. Contraction and  $\text{LC}_{20}$  phosphorylation elicited at pCa 5.75 were, however, unaffected by HA1077.
4. Pre-treatment of intact tissue strips with HA1077 abolished the tonic component of carbachol-induced contraction and the sustained elevation of  $\text{LC}_{20}$  phosphorylation, but had no effect on the transient or sustained increase in  $[\text{Ca}^{2+}]_i$  induced by carbachol.
5.  $\text{LC}_{20}$  phosphorylation and contraction dynamics suggest that the ROK-mediated increase in  $\text{LC}_{20}$  phosphorylation is due to MLCP inhibition, not myosin light chain kinase activation.
6. In the absence of  $\text{Ca}^{2+}$ ,  $\text{GTP}\gamma\text{S}$  stimulated  $^{35}\text{S}$  incorporation from  $^{35}\text{S}$ ATP $\gamma\text{S}$  into the myosin targeting subunit of MLCP (MYPT). The enhanced thiophosphorylation was inhibited by HA1077. No thiophosphorylation of  $\text{LC}_{20}$  was detected.
7. These results indicate that ROK mediates agonist-induced increases in myosin phosphorylation and force by inhibiting MLCP activity through phosphorylation of MYPT. Under  $\text{Ca}^{2+}$ -free conditions, ROK does not appear to phosphorylate  $\text{LC}_{20}$  *in situ*, in contrast to its ability to phosphorylate myosin *in vitro*. In particular, ROK activation is essential for the tonic phase of agonist-induced contraction.

Contractile stimulation of smooth muscle with physiological agonists leads to a rise in intracellular free  $\text{Ca}^{2+}$  concentration ( $[\text{Ca}^{2+}]_i$ ), calmodulin-dependent activation of myosin light chain kinase (MLCK), myosin regulatory light chain ( $\text{LC}_{20}$ ) phosphorylation, crossbridge cycling and force development. Potential for regulation of contraction by agonists is found at all levels. Recently, much attention has been focused on regulation of force that is independent of changes in  $[\text{Ca}^{2+}]_i$ , referred to as  $\text{Ca}^{2+}$  sensitization (Somlyo & Somlyo, 1994).  $\text{Ca}^{2+}$  sensitization occurs in response to stimulation of G protein-coupled receptors and can be mimicked by the non-hydrolysable GTP analogue  $\text{GTP}\gamma\text{S}$  or antagonized by  $\text{GDP}\beta\text{S}$  (Nishimura *et al.* 1988; Kitazawa *et al.* 1991a).

Cellular signalling pathways involved in  $\text{Ca}^{2+}$  sensitization converge on an increase in  $\text{LC}_{20}$  phosphorylation and analyses of kinase and phosphatase activities *in situ* have indicated that  $\text{LC}_{20}$  dephosphorylation is selectively reduced in this process, i.e.  $\text{Ca}^{2+}$  sensitization is mediated via inhibition of myosin light chain phosphatase (MLCP) (Kitazawa *et al.* 1991b; Kubota *et al.* 1992).

Several different mechanisms have been proposed to account for this inhibition of MLCP (for reviews see Somlyo & Somlyo, 1994; Hartshorne *et al.* 1998). These include dissociation of the subunits of the holoenzyme by arachidonic acid (Gong *et al.* 1992), phosphorylation of the myosin targeting subunit (MYPT) of MLCP by an unknown kinase (Ichikawa *et al.*

1996) or through the action of protein kinase C (PKC) (Masuo *et al.* 1994; Buus *et al.* 1998). The small GTPase RhoA (either the GTP $\gamma$ S-bound form or the constitutively active mutant, RhoA<sup>V14</sup>-GTP) induced Ca<sup>2+</sup> sensitization when added to permeabilized smooth muscle (Hirata *et al.* 1992; Noda *et al.* 1995; Gong *et al.* 1996; Otto *et al.* 1996). Conversely, inhibition of RhoA function (by ADP-ribosylation catalysed by exoenzyme C3 from *Clostridium botulinum* or by glucosylation catalysed by *Clostridium difficile* toxin B) reduced Ca<sup>2+</sup> sensitization (Otto *et al.* 1996; Gong *et al.* 1996; Akopov *et al.* 1998) and inhibited sustained contraction in intact smooth muscle (Fujihara *et al.* 1997; Lucius *et al.* 1998) indicating that RhoA plays an important role in this process. Ca<sup>2+</sup> sensitization requires translocation of RhoA to the plasma membrane where it interacts with an effector (Fujihara *et al.* 1997; Gong *et al.* 1997).

Several RhoA effectors, including Rho-associated kinase (ROK) (Leung *et al.* 1995; Ishizaki *et al.* 1996; Matsui *et al.* 1996), have been identified. Based on the observations that ROK phosphorylates myosin (Amano *et al.* 1996) and MYPT (causing MLCP inhibition) (Kimura *et al.* 1996), that recombinant constitutively-active ROK causes contraction in Triton X-100-skinned tissue (Kureishi *et al.* 1997) and that the selective ROK inhibitor Y-27632 inhibits phenylephrine-induced contraction of intact arterial smooth muscle (Uehata *et al.* 1997) and GTP $\gamma$ S-induced contraction of permeabilized arterial smooth muscle (Uehata *et al.* 1997; Fu *et al.* 1998), ROK has emerged as a candidate constituent of a signalling pathway leading to Ca<sup>2+</sup> sensitization.

It is interesting to note that longitudinal smooth muscle from the guinea-pig ileum, unlike other smooth muscles, demonstrates almost complete dependence on RhoA for Ca<sup>2+</sup> sensitization (Otto *et al.* 1996). It is, therefore, unlikely that Ca<sup>2+</sup> sensitization is a composite of various parallel pathways in this tissue. Guinea-pig ileum longitudinal smooth muscle therefore provides a suitable tissue for investigating the role of ROK in Ca<sup>2+</sup> sensitization of smooth muscle contraction.

## METHODS

### Animals

Guinea-pigs (250–400 g) were killed either by halothane inhalation or cervical dislocation as approved by the local Animal Care Committees at the University of Calgary and the University of Lund, respectively. A 20–30 cm long segment of the intestine just proximal to the ileocaecal valve was removed and washed in nominally Ca<sup>2+</sup>-free *N*-(2-hydroxyethyl)piperazine-*N'*-ethanesulphonic acid (Hepes)-buffered Krebs solution (for composition see 'Solutions'). Strips were prepared by tearing with fine forceps along natural lines of cleavage from the outer layer of ileal longitudinal smooth muscle. Taeniae coli were dissected from the colon and washed in Ca<sup>2+</sup>-free Hepes-buffered Krebs solution. Preparations with a width of 100–200  $\mu$ m, a thickness of 50–100  $\mu$ m and a length of 5–10 mm were thus obtained.

### Force determination and permeabilization

Muscle strips were attached with cellulose acetate glue to a force transducer (AME 801, SensoNor A/S, Horten, Norway) at one end

and a carbon fibre support at the other end. They were stretched shortly after mounting to  $\sim 0.2$  mN and thereafter the passive tension, determined in Ca<sup>2+</sup>-free solution, declined to a steady level of  $40 \pm 5$   $\mu$ N (2.4% of the maximum active plus passive force,  $n$  (no. of strips) = 47). Solution changes were rapidly effected by lowering a small platform supporting a Perspex cup and changing the cup for one containing the desired fresh solution. After equilibration in Hepes-buffered Krebs solution, the strips were contracted twice by immersion in high-K<sup>+</sup> solution (for composition see 'Solutions') at 22 °C then relaxed by immersion in Ca<sup>2+</sup>-free Hepes-buffered Krebs solution followed by intracellular substitution solution of pCa ( $-\log[\text{Ca}^{2+}]$ ) = 9. Strips were permeabilized with 0.05 mg ml<sup>-1</sup> (ileum) or 0.1 mg ml<sup>-1</sup> (taenia coli)  $\beta$ -escin in pCa 6.0 solution for 20–40 min at room temperature (20–22 °C), until force reached a plateau. The contractions elicited in pCa 4.5 solution immediately following permeabilization were used to normalize subsequent responses and these amounted to  $111.7 \pm 5.3\%$  ( $n = 47$ ) of the peak of the second contraction obtained in high-K<sup>+</sup> solution prior to permeabilization. The incubation volume was 200  $\mu$ l and agents or vehicle were added directly to the bath prior to change.

### Solutions

The Hepes-buffered Krebs solution used for intact strips had the following composition (mM): 135.5 NaCl, 5.9 KCl, 1.2 MgCl<sub>2</sub>, 11.6 glucose, 11.6 Hepes, 2.5 CaCl<sub>2</sub>, pH 7.35 at room temperature (20–22 °C). High-K<sup>+</sup> solution was obtained by replacing NaCl with equimolar KCl.

Solutions for permeabilized smooth muscle contained (standard solution): 3.2 mM MgATP; 2 mM free Mg<sup>2+</sup>, 12 mM phosphocreatine, 0.5 mM NaN<sub>3</sub>, 30 mM *N*-tris(hydroxymethyl)methyl-2-amino-ethanesulphonic acid, 0.5 mM dithioerythritol, 15 U ml<sup>-1</sup> creatine kinase, and 0.5  $\mu$ M calmodulin, pH 6.9 at room temperature. The desired free [Ca<sup>2+</sup>] for different solutions was obtained by mixing potassium ethyleneglycol-bis( $\beta$ -aminoethyl ether)-*N,N,N,N'*-tetra-acetic acid (EGTA) and K<sub>2</sub>CaEGTA, and the ionic strength was adjusted to 150 mM with potassium propionate. Rigor solution had the same composition as standard solution except it contained no ATP, phosphocreatine or creatine kinase; free [Mg<sup>2+</sup>] and ionic strength were maintained at 2 mM and 150 mM, respectively.

### Thiophosphorylation experiments

For mechanical experiments in which ATP $\gamma$ S was co-incubated with carbachol, GTP and ROK inhibitors, permeabilized muscle strips were first contracted in pCa 4.5 solution. After relaxation in pCa 9 solution, strips were washed (2  $\times$  5 min) in pCa 9 rigor solution before incubation with ATP $\gamma$ S (10  $\mu$ M) and additions as specified in the text. This was followed by three 5 min washes in pCa 9 solution containing ATP which ensured washout of carbachol and GTP, as determined in separate experiments. Strips were then contracted in pCa 6.5, 6.25 or 6.0 solution as specified. After a plateau response in pCa 4.5 solution was recorded, the chart recorder speed was increased to obtain relaxation times ( $t_{1/2}$ ) in pCa 9 solution. In some experiments, strips were finally exposed to 10  $\mu$ M microcystin-LR. The responses to microcystin-LR after 10 min incubation in the absence and presence of ATP $\gamma$ S were  $1.8 \pm 0.6$  and  $2.0 \pm 0.3$  mN ( $P > 0.4$ ,  $n = 8$ ), respectively. The corresponding values after 30 min incubation were  $1.5 \pm 0.5$  and  $1.6 \pm 0.5$  mN ( $P > 0.5$ ,  $n = 6$ ), respectively, suggesting that changes in maximal contractility were time dependent (i.e. a 20% reduction in 20 min) and not affected by the presence of the ATP $\gamma$ S during pre-incubations.

### Measurement of $[Ca^{2+}]_i$

Muscle strips were mounted in a bath staged on top of an inverted microscope for simultaneous recording of force and epifluorescence from 340 nm/380 nm excitation of the acetoxymethyl ester (AM) form of fura-2 with an emission wavelength of 510 nm using an IonOptix imaging system as described by Lindqvist *et al.* (1997). The tissue was loaded with 8  $\mu$ M fura-2 AM (Molecular Probes) in normal Hepes-buffered Krebs solution (loading solution) for 4 h at room temperature. Once every hour the strips were made to contract once by immersion in high- $K^+$  solution and then reimmersed in fresh loading solution. This loading protocol has previously been shown not to affect the mechanical performance of the tissue (Gomez & Swärd, 1997). After loading, each strip was allowed to equilibrate with perfusion for 30 min at 22 °C before the excitation light was turned on and data acquisition was started. Force and fluorescence signals were sampled at 10 Hz.

Strips were first contracted with carbachol (100  $\mu$ M) which was then washed out in normal Hepes-buffered Krebs solution ( $\geq 15$  min) during which time the lamp was turned off. After washout, strips were treated with 1-(5-isoquinolinesulphonyl)-homopiperazine (HA1077, 30  $\mu$ M) for 2–5 min before the lamp was turned on and the strips were contracted with carbachol in the presence of HA1077. Each experiment was concluded with an *in situ* calibration as described by Himpens *et al.* (1988). After background subtraction, fluorescence from excitation at 340 and 380 nm was used to calculate  $[Ca^{2+}]_i$  as described by Grynkiewicz *et al.* (1985). The dissociation constant of fura-2 for  $Ca^{2+}$  was taken to be 224 nM but is not known with certainty under the present intracellular conditions. The absolute values of  $[Ca^{2+}]_i$  reported should therefore be considered estimates.

### Quantification of $LC_{20}$ phosphorylation

For determination of  $LC_{20}$  phosphorylation levels, strips were tied as loops on stainless steel holders. The strips were then frozen at the desired times during contraction by rapidly (~1 s) submerging the entire holder in acetone mixed with 10% (w/v) trichloroacetic acid (TCA) and 20 mM dithiothreitol (DTT) that had been pre-cooled with dry ice. After lyophilization, protein extraction and urea-glycerol gel electrophoresis were performed as described by Weber *et al.* (1999). Separated proteins were blotted onto 0.2  $\mu$ m nitrocellulose (16 h, 30 V) and immunostained as described by Mita & Walsh (1997).  $LC_{20}$  phosphorylation was quantified by densitometric scanning using a Scanalytics Masterscan interactive densitometer using Camscan/RFLPscan 1.01 software (CSPI, Inc., Billerica, MA, USA). We observed a linear relationship between the chemiluminescence signal (optical density  $\times$  area of the band) and sample loading over the range 0–25  $\mu$ l. Loading levels of 10 or 15  $\mu$ l were therefore routinely used.

### Western blotting

Sheets of tissue were blotted on filter paper, weighed and homogenized using a Brinkmann Polytron in 15 volumes of homogenization buffer (composition: 50 mM tris(hydroxymethyl)aminomethane-HCl (Tris-HCl), pH 7.5, 400 mM NaCl, 2 mM EGTA, 1 mM ethylenediaminetetraacetic acid (EDTA), 1 mM DTT, 10  $\mu$ M phenylmethyl sulphonyl fluoride (PMSF), 10  $\mu$ g ml<sup>-1</sup> leupeptin, 1  $\mu$ g ml<sup>-1</sup> pepstatin and 1 mM benzamide). The homogenate was mixed with an equal volume of 2  $\times$  sodium dodecyl sulphate (SDS) sample buffer (Laemmli, 1970) and boiled. Proteins were separated by SDS-polyacrylamide gel electrophoresis (7.5–20% acrylamide gradient) and blotted onto nitrocellulose (0.2  $\mu$ m) in 25 mM Tris, 192 mM glycine, 20% (v/v) methanol, 0.1% (w/v) SDS, pH 8.3 at 40 V and 4 °C. Membranes were blocked for 1 h with 5% (w/v) dry milk in TBS-Tween (20 mM

Tris-HCl, pH 7.5, 0.5 M NaCl, 0.05% (v/v) Tween 20). After three 10 min washes in TBS-Tween, a primary antibody raised against the RhoA-binding domain of ROK $\beta$  was used at 1:200 dilution for detection of ROK. For RhoA detection, monoclonal antibody 26C4 (2  $\mu$ g ml<sup>-1</sup>; Santa Cruz Biotechnology Inc.) was used. MLCK was detected with a polyclonal antibody raised against chicken gizzard MLCK (0.25  $\mu$ g ml<sup>-1</sup>; Paul *et al.* 1995) and MYPT was detected with a monoclonal antibody raised against a 58 kDa fragment of chicken gizzard MYPT (1:2000 dilution; Okubo *et al.* 1994). Horseradish peroxidase-conjugated secondary antibodies, anti-rabbit and anti-mouse IgGs (both from Boehringer-Mannheim), were used at 1:5000 dilution. Membranes were incubated for 1 h each with primary and secondary antibody diluted in TBS-Tween with 1% dry milk. Incubations were followed by three 10 min washes in TBS-Tween. The blots were developed with the Supersignal CL-HRP substrate system (Pierce Chemical Co.) after a final 10 min wash in TBS.

### $[^{35}S]$ ATP $\gamma$ S experiments

Muscle strips were prepared as for  $LC_{20}$  phosphorylation experiments and permeabilized as described above. After washing (2  $\times$  5 min) in pCa 9 standard solution and then in pCa 9 rigor solution (2  $\times$  5 min) to remove residual ATP, strips were treated with 10  $\mu$ M  $[^{35}S]$ ATP $\gamma$ S (1 mCi ml<sup>-1</sup>) in pCa 9 rigor solution with additions as indicated in the figures. At the times indicated, groups of three muscle strips were snap-frozen in a dry ice-acetone slurry containing 10% (w/v) TCA and 20 mM DTT. Strips were removed from the holders, frozen briefly again and washed (8  $\times$  0.5 min) in acetone plus DTT. After lyophilization overnight, proteins were extracted in Eppendorf tubes with 200  $\mu$ l of 2  $\times$  SDS gel sample buffer for 2 h and then boiled. Proteins were separated by SDS-PAGE and stained with Coomassie Brilliant Blue. The wet gel enclosed in a sealed plastic bag was exposed in a Phosphorimager (Molecular Dynamics Storm 850) for 4–8 h for quantification of radioactivity. Radioactivity in the 130 kDa band was corrected for background and slight variations in protein loading levels, the latter obtained by scanning the stained gel (calponin or filamin bands) using a Pharmacia Biotech Image Master desktop scanning system (Uppsala, Sweden). Gels were also dried in a gel drier (model 583, BioRad) and exposed to X-ray film (Biomax MR, Kodak) for up to 14 days. The radiolabelled 130 kDa band was quantified by scanning the autoradiogram using the Image Master desktop scanning system.

### Protein purification

ROK was purified using a modification of the procedures used by Leung *et al.* (1995) and Matsui *et al.* (1996). Bovine brains were transported on ice from a local abattoir. The tissue was cleaned of adhering blood vessels and white matter in the brainstem and hemispheres with a scalpel. After weighing, the tissue was homogenized in a Waring blender with three volumes of homogenization buffer (HB, composition: 25 mM Tris-HCl, pH 7.5; 1 mM DTT, 5 mM EGTA, 10 mM MgCl<sub>2</sub>, 10  $\mu$ M PMSF, 1  $\mu$ g ml<sup>-1</sup> pepstatin, 1  $\mu$ g ml<sup>-1</sup> leupeptin and 10% (w/v) sucrose). The homogenate was centrifuged at 20 000 *g* for 30 min. The pellet was resuspended in two volumes of HB as before and an equal volume of HB with 4 M NaCl was added with continuous overhead stirring. After stirring for 1 h, the suspension was centrifuged 2  $\times$  1 h as before and the resulting pellets were discarded. The final supernatant was dialysed overnight against 3  $\times$  8 l of buffer A (composition: 20 mM Tris-HCl, pH 7.5, 1 mM EDTA, 5 mM MgCl<sub>2</sub>, 1 mM DTT).

Ammonium sulphate was slowly added to the dialysate (over 30 min) with overhead stirring to 40% saturation (w/v) followed by

centrifugation as before for 1–2 h. Supernatants were carefully discarded, the pellets were resuspended in adhering liquid plus ~10 ml of buffer A and dialysed against 3 × 5 l of buffer A. After centrifugation as above, the supernatant was loaded on a Q-Sepharose column (Pharmacia Biotech, Sweden) previously equilibrated with buffer A at a flow rate of 120 ml h<sup>-1</sup>. The column was washed for 1 h with buffer A and bound proteins were eluted with a 500 ml 0–0.8 M NaCl gradient in the same buffer, 3 ml fractions were collected. ROK was detected in selected fractions by Western blotting after separation of proteins on 7.5–20% SDS-PAGE gels. The primary antibody used was monoclonal antibody 1A1 raised against ROK $\alpha$  (1  $\mu$ g ml<sup>-1</sup>) and the secondary antibody goat anti-mouse IgG (Boehringer Mannheim, 1:5000 dilution). Fractions containing ROK $\alpha$  were pooled and dialysed against 3 × 2 l buffer B (composition: 25 mM 2-(N-morpholino)ethanesulphonic acid (Mes)-NaOH, pH 6.0, 0.5 mM MgCl<sub>2</sub>, 0.05% (w/v) Triton X-100, 0.1 M NaCl).

The ROK pool was subsequently loaded on a glutathione S-transferase (GST)–RhoA–GTP $\gamma$ S or a GST–RhoA<sup>V14</sup>–GTP $\gamma$ S affinity column previously equilibrated with buffer B plus 50  $\mu$ M GTP at 15 ml h<sup>-1</sup>. The column was washed for 1–2 h with buffer B plus 50  $\mu$ M GTP. Bound ROK $\alpha$  was eluted with 25 mM Tris, pH 8, 0.5 mM MgCl<sub>2</sub>, 0.05% (w/v) Triton X-100, 0.1 mM EDTA and 0.1 M NaCl. ROK-containing fractions, identified by enzymatic assays and Western blotting, were frozen.

RhoA–GST and RhoA<sup>V14</sup>–GST were expressed in *E. coli* (strain pLysS, vector pGEX 2T) and purified from the bacterial cytosol according to Glover & Hames (1995) protocols 7, 8 and 10. RhoA–GST (7–10 mg) was incubated for 1 h at 30 °C in 20 mM Tris-HCl, pH 7.5, 10 mM EDTA, 1 mM DTT, 5 mM MgCl<sub>2</sub>, 0.03% (w/v) 3-(3-cholamido-propyl)dimethylammonio-1-propane-

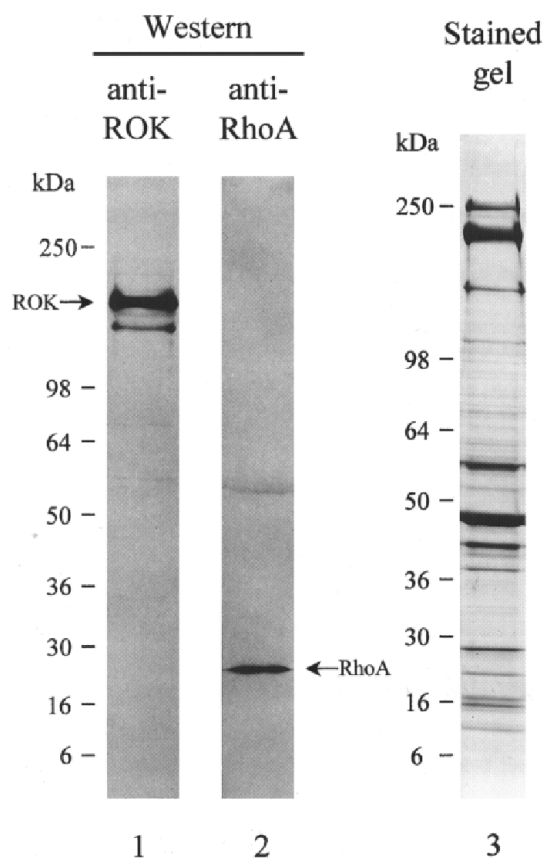
sulphonate (CHAPS), 1 mM 1,2-dimyristoyl-*sn*-glycero-3-phosphocholine (DMPC) and 50  $\mu$ M GTP $\gamma$ S to load RhoA with GTP $\gamma$ S. Glutathione–Sepharose (10 ml; Pharmacia Biotech), previously equilibrated with buffer A and dried, was then added. The mix was tumbled for 1 h at 4 °C before packing and equilibration of the affinity column.

Chicken gizzard MLCK (Ngai *et al.* 1984), rat brain PKC (a mixture of Ca<sup>2+</sup>-dependent  $\alpha$ ,  $\beta$  and  $\gamma$  isoforms) (Allen *et al.* 1994) and bovine brain calmodulin (Walsh *et al.* 1984) were purified as described previously. Chicken gizzard LC<sub>20</sub> was purified using a modification of the procedure of Hathaway & Haeberle (1983). Chicken gizzards were obtained from a local poultry packing plant and transported to the laboratory on ice. For PKC purification male Sprague-Dawley rats (300–500 g) were killed by inhalation of a lethal dose of halothane and decapitated, according to a research protocol consistent with the standards of the Canadian Council on Animal Care and approved by the local Animal Care Committee of the Medical Research Council of Canada.

#### Enzyme assays

ROK activity was assayed using a synthetic peptide (MYPT peptide, 0.2 mg ml<sup>-1</sup>) derived from the sequence surrounding the phosphorylatable threonine 654 (RQSRRTQGVTLTD) of the 130 kDa myosin targeting subunit of chicken gizzard MLCP (Ichikawa *et al.* 1996) as substrate under the following conditions: 50 mM Tris-HCl, pH 7.5, 2 mM EDTA, 7 mM MgCl<sub>2</sub>, 1 mM DTT, 0.15% (w/v) CHAPS and 0.35  $\mu$ M ROK.

MLCK activity was assayed under the following conditions: 25 mM Tris-HCl, pH 7.5, 60 mM KCl, 4 mM MgCl<sub>2</sub>, 0.1 mM CaCl<sub>2</sub>, 1 mM DTT, 0.1% (w/v) Tween 80, 10  $\mu$ M LC<sub>20</sub>, 0.5 nM MLCK and 0.6  $\mu$ M calmodulin.



**Figure 1. Identification of ROK and RhoA in guinea-pig ileum longitudinal smooth muscle**

Tissue homogenates were subjected to SDS-PAGE and either stained with Coomassie Brilliant Blue (lane 3) or immunoblotted with anti-ROK $\beta$  (lane 1) or anti-RhoA (lane 2). The equivalent of 0.625 mg wet weight of tissue was applied to lane 1, 2.5 mg wet weight of tissue to lane 2 and 2.5 mg wet weight of tissue to lane 3. Results are representative of three independent experiments.

PKC activity was assayed under the following conditions: 20 mM Hepes, pH 7.5, 10 mM MgCl<sub>2</sub>, 10 mM DTT, 0.15 mM CaCl<sub>2</sub>, 1 mg ml<sup>-1</sup> histone III-S, mixed micelles (composed of 0.3 mg ml<sup>-1</sup> phosphatidylserine and 62 µg ml<sup>-1</sup> 1,2-diolein in Triton X-100, 0.03% (w/v)), and 0.6 nM PKC. All assays were performed in an assay volume of 25 µl and reactions were started by addition of [<sup>γ</sup>-<sup>32</sup>P]ATP (200–500 c.p.m. pmol<sup>-1</sup>) to 0.25 mM at 30 °C. Reactions were stopped by spotting 22.5 µl of assay mix on Whatman P81 filter paper squares (1 cm × 1 cm) after 5 or 10 min (MLCK and PKC assays) or 30 min (ROK assay). Filter papers were then washed 3 times in phosphoric acid (0.5%, v/v) and once in acetone followed by drying and Čerenkov counting.

### Chemicals

HA1077, 1-(5-isoquinolinesulphonyl)-2-methylpiperazine (H-7), 1-(5-chloronaphthalene-1-sulphonyl)-1*H*-hexahydro-1,4-diazepine (ML-9) and microcystin-LR were purchased from Calbiochem. 3-(2-(4-(Bis(4-fluorophenyl)methylene)-1-piperidinyl)ethyl)-2,3-dihydro-2-thioxo-4[1*H*]-quinazolinone (R59022) was purchased from Sigma. Y-27632 (*R*-(+)-*trans*-*N*-(4-pyridyl)-4-(1-aminoethyl)-cyclohexanecarboxamide) was generously provided by Yoshitomi Pharmaceutical Industries, Ltd (Osaka, Japan). MYPT peptide was synthesized by Macromolecular Resources (Colorado State University). [<sup>γ</sup>-<sup>32</sup>P]ATP (>5000 Ci mmol<sup>-1</sup>) and [<sup>35</sup>S]ATPγS (>1000 Ci mmol<sup>-1</sup>) were purchased from Pharmacia Biotech. Triton X-100 (peroxide free) was purchased from Boehringer-Mannheim. L-α-Phosphatidylserine and 1,2-diolein were purchased from Serdary Research Laboratories (London, Canada). DMPC was from Avanti Polar Lipids, Inc. (Alabaster, AL, USA). Tween 20 (polyoxyethylene sorbitan mono-oleate) was purchased from

BioRad. Leupeptin, pepstatin A and DTT were from ICN Biomedicals, Inc. (Aurora, OH, USA). All other chemicals were of analytical grade or better and were purchased from Canlab (Edmonton, AB, Canada) or Sigma. Antibodies to ROKα and ROKβ were generously provided by Drs Thomas Leung and Louis Lim, The National University of Singapore. Anti-ROKα is a mouse monoclonal antibody (1A1) raised against the kinase domain (residues 112–392; Leung *et al.* 1995) and anti-ROKβ is a polyclonal antiserum raised in rabbits against a region (residues 670–1027) containing the Rho-binding domain (Leung *et al.* 1996). These antibodies recognize the same population of proteins in various rat tissues and therefore do not distinguish between ROKα and β (Leung *et al.* 1996).

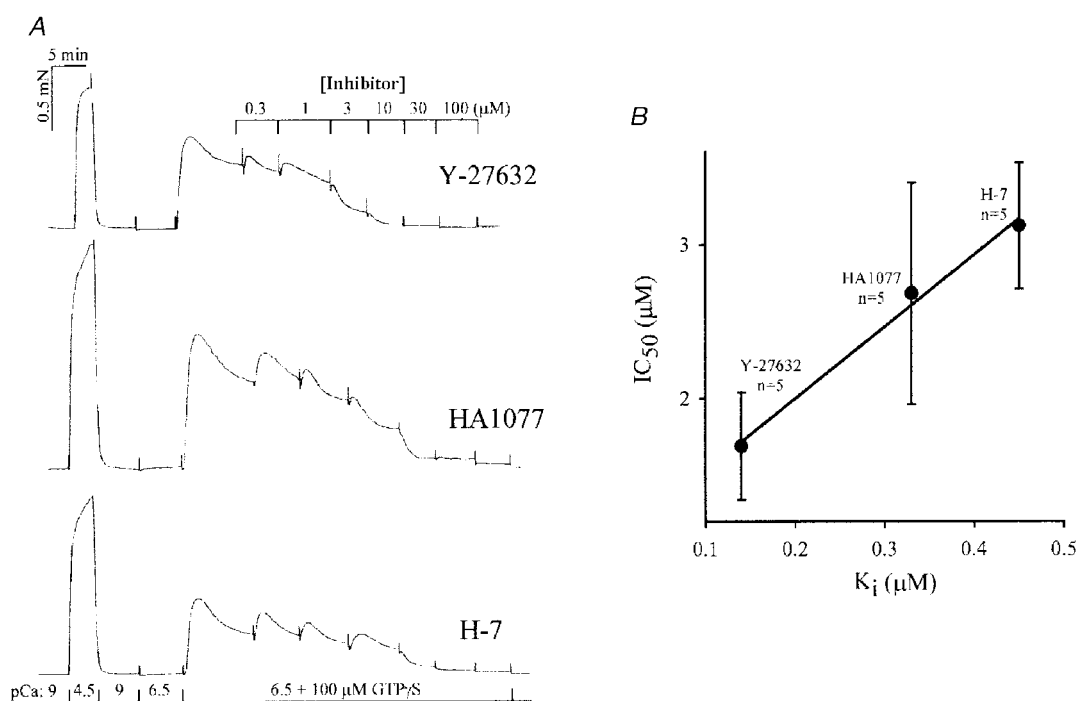
### Statistics

Student's *t* test, paired or unpaired as appropriate, was used for single statistical comparisons. For multiple comparisons, one-way analysis of variance followed by Dunn's test (Fig. 8*B*) was used. *P* < 0.05 was considered significant. Summarized data are presented as means ± s.e.m.

## RESULTS

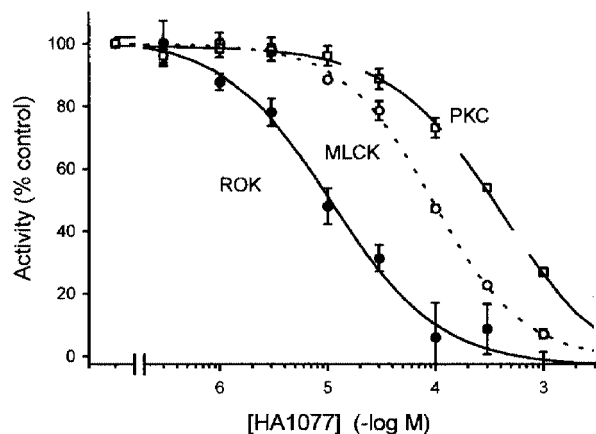
### Expression of RhoA and ROK in guinea-pig ileum

Figure 1 demonstrates by Western blotting with antibodies to ROKβ (lane 1) and RhoA (lane 2) that both ROK and RhoA are expressed in guinea-pig ileum longitudinal smooth muscle. The calculated molecular weights were as expected: ~160 kDa for ROK and 22 kDa for RhoA.



**Figure 2. Effects of kinase inhibitors on Ca<sup>2+</sup> sensitization of force induced in permeabilized guinea-pig ileum by GTPγS**

*A*, guinea-pig ileum longitudinal smooth muscle permeabilized with β-escin was contracted by increasing [Ca<sup>2+</sup>] from pCa 9 to 4.5 and relaxed by returning to pCa 9. Transfer to pCa 6.5 solution failed to elicit a significant contraction but subsequent transfer to pCa 6.5 + GTPγS (100 µM) evoked Ca<sup>2+</sup> sensitization of force. The effects of increasing concentrations of three kinase inhibitors (Y-27632, HA1077 and H-7) on Ca<sup>2+</sup> sensitization were recorded. *B*, correlation between IC<sub>50</sub> values for inhibition of Ca<sup>2+</sup> sensitization and the known K<sub>i</sub> values for inhibition of ROK (*n* = 5).



**Figure 3. Sensitivity of ROK, MLCK and PKC to the kinase inhibitor HA1077**

The effects of increasing concentrations of HA1077 on ROK (●), MLCK (○) and PKC activities (□) were determined as described in Methods. Activity in the presence of vehicle was expressed as 100%.

Figure 1 (lane 3) shows a Coomassie Blue-stained gel of the tissue homogenate used for Western blotting.

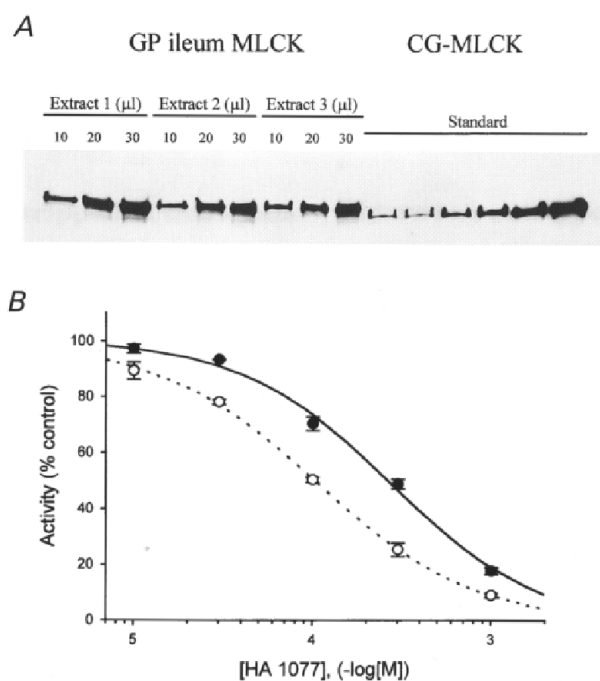
#### Effects of kinase inhibitors on $\text{Ca}^{2+}$ sensitization of force in guinea-pig ileum

Addition of GTP $\gamma$ S (100  $\mu\text{M}$ ) to solution bathing  $\beta$ -escin-permeabilized guinea-pig ileum longitudinal smooth muscle strips at a sub-threshold  $[\text{Ca}^{2+}]$  (pCa 6.5) induced a contraction that peaked and then declined to a steady level corresponding to  $25.9 \pm 2.0\%$  ( $n = 20$ ) of the preceding reference contraction in pCa 4.5 solution (Fig. 2A), i.e. GTP $\gamma$ S induced  $\text{Ca}^{2+}$  sensitization. Subsequent addition of the kinase inhibitors Y-27632, HA1077 and H-7 led to concentration-dependent relaxation to  $< 5\%$  residual force. Force was well maintained in the presence of vehicle alone over the same number of solution changes for the same duration ( $-0.7 \pm 13.0\%$ ,  $n = 5$ ). The calculated  $\text{IC}_{50}$  values for inhibition of  $\text{Ca}^{2+}$  sensitization ( $1.7 \pm 0.4 \mu\text{M}$  for Y-27632,  $2.7 \pm 0.7 \mu\text{M}$  for HA1077 and  $3.1 \pm 0.4 \mu\text{M}$  for H-7)

correlated well ( $r^2 = 0.99$ ; Fig. 2B) with the reported  $K_i$  values for inhibition of ROK *in vitro* (Uehata *et al.* 1997).

HA1077 was used to inhibit ROK in most subsequent experiments due to its ready commercial availability. Figure 3 confirms its selectivity for inhibition of ROK over MLCK and PKC with  $\text{IC}_{50}$  values of  $10.7 \pm 2.0 \mu\text{M}$  (ROK),  $95.2 \pm 4.7 \mu\text{M}$  (MLCK) and  $425 \pm 41 \mu\text{M}$  (PKC). The PKC inhibitor chelerythrine ( $\leq 100 \mu\text{M}$ ) and the tyrosine kinase inhibitor genistein ( $\leq 100 \mu\text{M}$ ) had no significant effect on ROK activity (results not shown).

Many of the subsequent experiments designed to achieve specific inhibition of ROK activity in intact and permeabilized smooth muscle were carried out at 30  $\mu\text{M}$  HA1077. From the data in Fig. 3 it is apparent that, under the *in vitro* conditions used in this experiment, MLCK activity was inhibited by  $\sim 20\%$  at this concentration of HA1077. It should be noted, however, that the concentration of MLCK in this experiment was low,



**Figure 4. Quantification of MLCK levels in guinea-pig ileum and the effect of [ATP] on inhibition of MLCK by HA1077**

A, guinea-pig ileum longitudinal smooth muscle proteins were extracted as described in Methods, subjected to SDS-PAGE and immunoblotted with anti-MLCK. Three loading levels of three independent extracts and different known amounts of purified chicken gizzard MLCK (0.05, 0.1, 0.15, 0.2, 0.4 and 0.8  $\mu\text{g}$ ) were applied to the gel. MLCK bands were quantified by densitometric scanning. A standard curve relating the intensity of the chicken gizzard MLCK bands (optical density  $\times$  area of the band) to micrograms of MLCK loaded was linear with an  $r^2$  value of 0.98. B, the effect of increasing concentrations of HA1077 on the activity of purified MLCK was assayed as described in Methods at 0.25 (○) and 1 mM ATP (●).

0.05  $\mu\text{g ml}^{-1}$  or 0.5 nM, which was necessary to ensure linear phosphorylation time courses. The tissue concentration of MLCK in guinea-pig ileum longitudinal smooth muscle was calculated to be  $1.8 \pm 0.1 \mu\text{M}$  ( $n = 3$ ) assuming 50% cell water (Fig. 4A) so that significantly higher concentrations of HA1077 would be required to inhibit MLCK *in situ*. This estimate of the tissue concentration of MLCK is similar to values previously published for other smooth muscle tissues (2–4  $\mu\text{M}$ , Walsh *et al.* 1984; Tansey *et al.* 1994) and is significantly higher than the tissue concentration of ROK (estimated to be 0.37  $\mu\text{M}$  in chicken gizzard from the yield of purified enzyme reported by Feng *et al.* 1999). The ROK concentration used in Fig. 3 (0.35  $\mu\text{M}$ ) was close to this estimated tissue concentration. HA1077 inhibits ROK by competing with ATP. Figure 4B shows that increasing the ATP concentration in the *in vitro* MLCK assay towards the physiological concentration shifts the concentration–response curve to the right so that at 1 mM ATP 30  $\mu\text{M}$  HA1077 had no significant inhibitory effect. The  $\text{IC}_{50}$  values were calculated to be  $105.8 \pm 4.7 \mu\text{M}$  HA1077 at 0.25 mM ATP and  $273.5 \pm 31.9 \mu\text{M}$  HA1077 at 1 mM ATP. Taken together, these results indicate that 30  $\mu\text{M}$  HA1077 is unlikely to have any inhibitory effect on MLCK *in situ*. Confirmatory evidence will be presented later (Table 3).

Using a protocol similar to that in Fig. 2, GTP (50  $\mu\text{M}$ ) and carbachol (100  $\mu\text{M}$ ) at pCa 6.5 increased force to a steady level of  $25.7 \pm 3.1\%$  ( $n = 6$ ) of the reference contraction at pCa 4.5 (Fig. 5A). HA1077 caused a concentration-dependent relaxation to near-basal force, with maximal effect at  $\sim 30 \mu\text{M}$  ( $n = 6$ ). Contraction induced at pCa 5.75 amounted to  $96.0 \pm 4.0\%$  ( $n = 8$ ) of the reference contraction at pCa 4.5 and was not affected by HA1077 at concentrations as high

**Table 1. Phorbol ester induces  $\text{Ca}^{2+}$  sensitization of force in permeabilized guinea-pig taenia coli but not ileum**

[PDBu] ( $\mu\text{M}$ )	Steady-state force	
	Taenia coli (% maximum)	Ileum (% maximum)
0	$31.6 \pm 4.4$ ( $n = 10$ )	$16.0 \pm 2.5$ ( $n = 12$ )
1	$78.0 \pm 5.5^*$ ( $n = 10$ )	$13.1 \pm 4.4$ ( $n = 4$ )
10	—	$16.8 \pm 3.0$ ( $n = 8$ )

Steady-state force was measured in  $\beta$ -escin-permeabilized guinea-pig taenia coli or ileum at pCa 6.25 in the absence or presence of phorbol 12,13-dibutyrate (PDBu, 1 or 10  $\mu\text{M}$ ). Values were normalized to the reference contraction at pCa 4.5 in each experiment and are given as means  $\pm$  S.E.M. \*Significantly different from the corresponding response in the absence of PDBu ( $P < 0.001$ ; Student's *t* test).

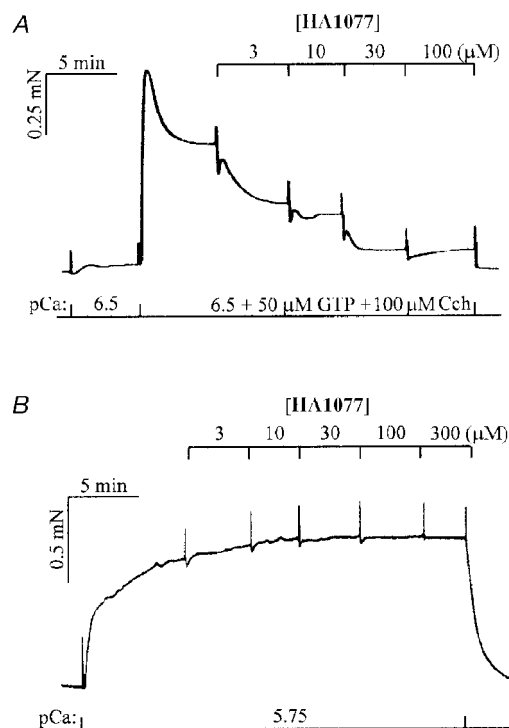
as 300  $\mu\text{M}$  ( $n = 8$ ; Fig. 5B). These results implicate ROK activation in agonist-induced  $\text{Ca}^{2+}$  sensitization.

#### Lack of $\text{Ca}^{2+}$ sensitization in guinea-pig ileum in response to PKC activation

PKC activation has been implicated in  $\text{Ca}^{2+}$  sensitization in a variety of smooth muscle tissues (Masuo *et al.* 1994; Sato *et al.* 1994; Gailly *et al.* 1997; Buus *et al.* 1998; Li *et al.* 1998). As shown in Table 1, however, phorbol 12,13-dibutyrate (1 or 10  $\mu\text{M}$ ) failed to induce  $\text{Ca}^{2+}$  sensitization in  $\beta$ -escin-permeabilized guinea-pig ileum longitudinal smooth muscle. On the other hand, the same stock solution of phorbol ester significantly increased force at pCa 6.25 in  $\beta$ -escin-permeabilized guinea-pig taenia coli at a final

**Figure 5. Effects of HA1077 on  $\text{Ca}^{2+}$  sensitization of force evoked by agonist plus GTP and  $\text{Ca}^{2+}$ -induced contraction**

**A**, force in  $\beta$ -escin-permeabilized guinea-pig ileum longitudinal smooth muscle was activated at a sub-threshold [ $\text{Ca}^{2+}$ ] (pCa 6.5) by addition of 100  $\mu\text{M}$  carbachol (Cch) and 50  $\mu\text{M}$  GTP. Cumulative addition of HA1077 inhibited this  $\text{Ca}^{2+}$  sensitization. In control strips ( $n = 6$ ), Cch + GTP-induced steady-state force at pCa 6.5 was maintained for at least 40 min.  $\text{Ca}^{2+}$  sensitization induced by Cch + GTP was inhibited by 100  $\mu\text{M}$  atropine, verifying that the Cch effect was mediated by muscarinic receptors. **B**, force was maximally activated at pCa 5.75 and was unaffected by cumulative addition of HA1077 up to 300  $\mu\text{M}$ .



**Table 2. Inhibition of diacylglycerol kinase fails to increase carbachol-induced force in permeabilized guinea-pig ileum**

[Carbachol] ( $\mu\text{M}$ )	Steady-state force		<i>P</i>
	Control (% maximum)	R59022 (% maximum)	
1	17.7 $\pm$ 1.7	7.7 $\pm$ 2.2*	0.005
3	27.1 $\pm$ 2.6	12.3 $\pm$ 2.7*	0.004
10	36.3 $\pm$ 4.3	21.7 $\pm$ 3.9*	0.03

Steady-state force was measured in  $\beta$ -escin-permeabilized guinea-pig ileum longitudinal smooth muscle in response to the indicated concentrations of carbachol at pCa 6.25 in the absence and presence of the diacylglycerol kinase inhibitor R59022 (10  $\mu\text{M}$ ). Values were normalized to the reference contraction at pCa 4.5 in each experiment and are given as means  $\pm$  s.e.m. ( $n = 6$ ). \* Significantly different from the corresponding response in the absence of R59022 with the indicated *P* values.

concentration of 1  $\mu\text{M}$ . Furthermore, treatment of permeabilized guinea-pig ileum longitudinal smooth muscle at pCa 6.25 with the diacylglycerol kinase inhibitor R59022 did not increase carbachol-induced force, as would be expected if PKC activation evoked  $\text{Ca}^{2+}$  sensitization, but actually significantly reduced carbachol-induced force (Table 2). The lack of detectable PKC-mediated  $\text{Ca}^{2+}$  sensitization in guinea-pig ileum longitudinal smooth muscle makes this tissue particularly well suited for investigation of the functional role of ROK.

### Effects of HA1077 on $\text{LC}_{20}$ phosphorylation in permeabilized guinea-pig ileum

To determine if ROK inhibition by HA1077 was associated with changes in the level of  $\text{LC}_{20}$  phosphorylation, permeabilized muscle strips were frozen 5 min after exposure to the conditions shown in Fig. 6.  $\text{LC}_{20}$  phosphorylation levels were very low in relaxed muscle strips at pCa 9 and

did not increase significantly when the  $[\text{Ca}^{2+}]_i$  was increased to pCa 6.5, consistent with the lack of a contractile response at this  $[\text{Ca}^{2+}]_i$  (Figs 2 and 5).  $\text{LC}_{20}$  phosphorylation was significantly increased by  $\text{GTP}\gamma\text{S}$  at pCa 6.5 and 5.75 and this was prevented by pretreatment with HA1077 (30  $\mu\text{M}$ ). On the other hand, the increase in  $\text{LC}_{20}$  phosphorylation induced by increasing  $[\text{Ca}^{2+}]_i$  to pCa 5.75 without  $\text{GTP}\gamma\text{S}$  was unaffected by 30  $\mu\text{M}$  HA1077.

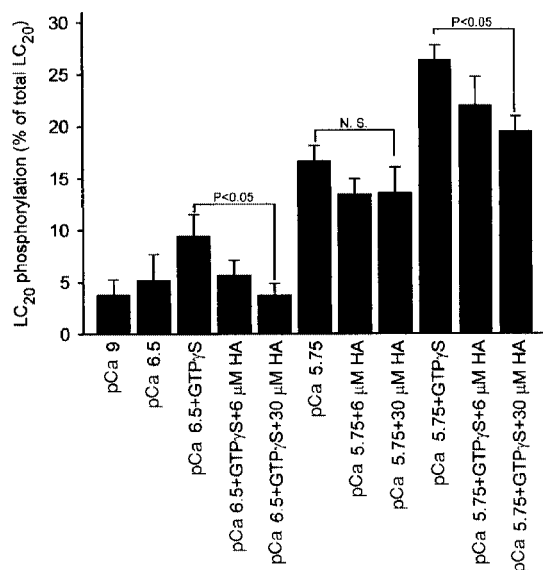
### Effects of HA1077 on force and $[\text{Ca}^{2+}]_i$ in intact guinea-pig ileum muscle strips stimulated with carbachol

Stimulation of intact fura-2 AM-loaded guinea-pig ileum longitudinal smooth muscle strips with carbachol (100  $\mu\text{M}$ ) led to a transient increase in  $[\text{Ca}^{2+}]_i$  which settled at a sustained level significantly above resting  $[\text{Ca}^{2+}]_i$  (Fig. 7A). Force peaked rapidly, relaxed partially, then increased again before relaxing very slowly (Fig. 7B). Pre-treatment with HA1077 (30  $\mu\text{M}$ ) had no effect on the  $[\text{Ca}^{2+}]_i$  response to carbachol (Fig. 7C) but completely abolished the tonic phase of the contractile response (Fig. 7D). The phasic component of the contractile response to carbachol was slightly reduced by HA1077 (compare Fig. 7B and D).

Data from seven  $[\text{Ca}^{2+}]_i$ -force experiments are summarized in Fig. 7E and F. Whereas integrated  $[\text{Ca}^{2+}]_i$  was not affected by 30  $\mu\text{M}$  HA1077, integrated force was markedly reduced.

### Effect of HA1077 on $\text{LC}_{20}$ phosphorylation in intact guinea-pig ileum

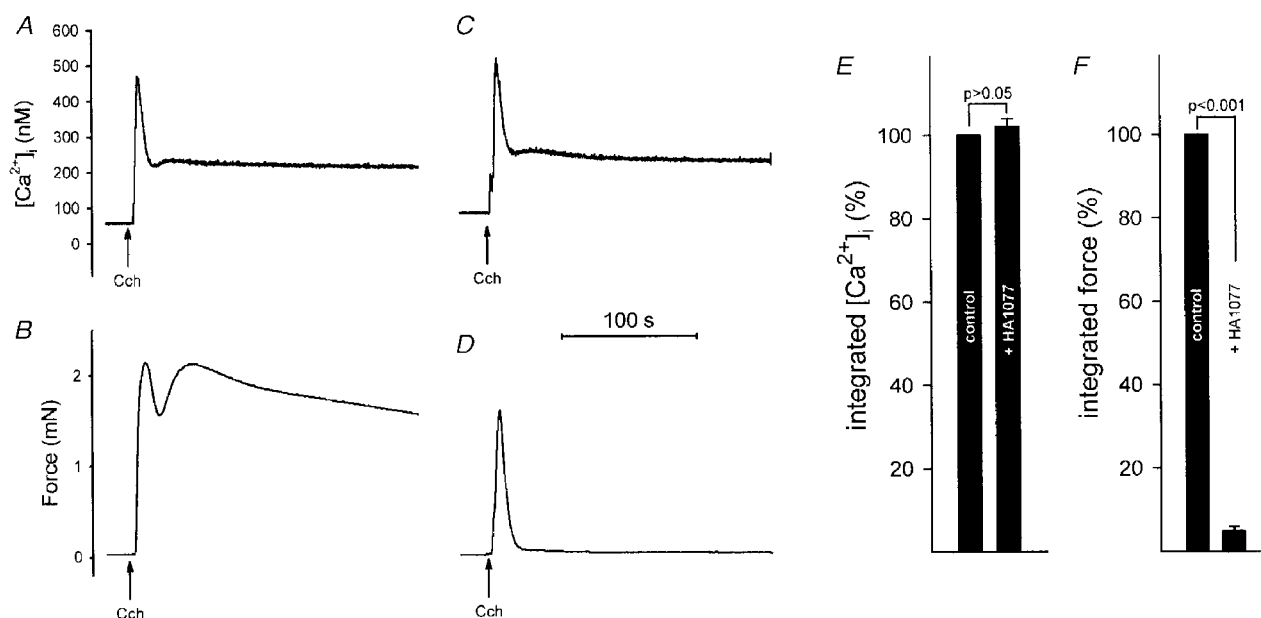
Intact guinea-pig ileum longitudinal smooth muscle strips were frozen prior to stimulation (0 s) and at 3, 6, 15, 30, 60 and 200 s after stimulation with carbachol (100  $\mu\text{M}$ ) in the absence or presence of HA1077 (6 or 30  $\mu\text{M}$ ) and  $\text{LC}_{20}$  phosphorylation levels were quantified. A typical experimental result is shown in Fig. 8A. The level of phosphorylation was close to zero prior to stimulation either in the absence or presence of HA1077. In the absence of



**Figure 6. Effect of HA1077 (HA) on the  $\text{GTP}\gamma\text{S}$ -induced increase in  $\text{LC}_{20}$  phosphorylation in permeabilized guinea-pig ileum**

$\beta$ -Escin-permeabilized guinea-pig ileum smooth muscle strips were exposed to the indicated conditions for 5 min, following which the strips were snap-frozen and  $\text{LC}_{20}$  phosphorylation levels quantified as described in Methods. Columns and bars represent the means and s.e.m.s. Columns 1–5,  $n = 5$ ; column 6,  $n = 11$ ; columns 7–11,  $n = 6$ . N.S., not significant.



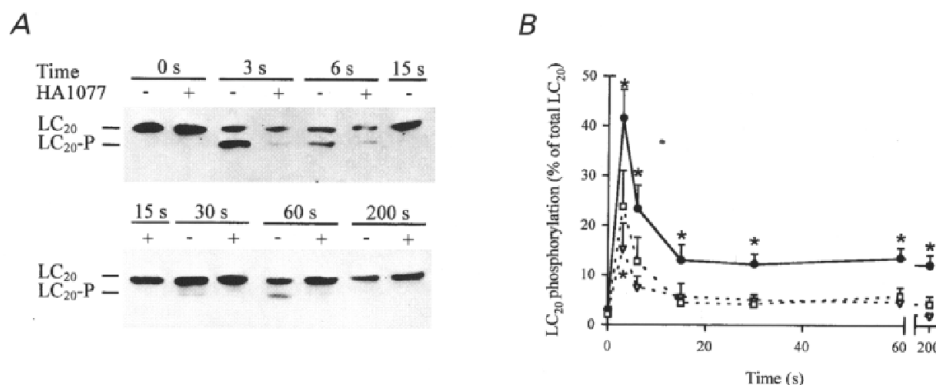


**Figure 7. Effects of HA1077 on carbachol-induced changes in  $[Ca^{2+}]_i$  and force in intact guinea-pig ileum**

Intact guinea-pig ileum longitudinal smooth muscle strips were loaded with fura-2 AM.  $[Ca^{2+}]_i$  (A and C) and force (B and D) were recorded following stimulation with carbachol ( $100 \mu M$ ) in the absence (A and B) and presence (C and D) of HA1077 ( $30 \mu M$ ). HA1077 was present 4 min prior to, and during, the carbachol-induced contraction. Integrated and normalized  $[Ca^{2+}]_i$  (E) and integrated and normalized force (F) were recorded in controls and after exposure to HA1077 ( $n = 7$ ).

HA1077,  $LC_{20}$  phosphorylation increased rapidly to a peak at 3 s and then declined to almost zero at 15 s followed by a second increase between 30 and 60 s. By 200 s, phosphorylation had fallen again to almost zero. This biphasic behaviour was observed in 10 of 13 experiments, but the times at which the first reduction and the second peak of  $LC_{20}$  phosphorylation took place were variable. The

average data presented in Fig. 8B ( $n = 6-16$ ), therefore, show only a rapid increase in  $LC_{20}$  phosphorylation that peaked at 3 s and then declined to a sustained level that was significantly greater than the resting level ( $P < 0.05$ ). In the presence of 6 or  $30 \mu M$  HA1077,  $LC_{20}$  phosphorylation also increased rapidly in response to carbachol but returned quickly to resting levels. HA1077 reduced the transient



**Figure 8. Effect of HA1077 on carbachol-induced  $LC_{20}$  phosphorylation in intact guinea-pig ileum**

Intact guinea-pig ileum longitudinal smooth muscle strips were treated with carbachol ( $100 \mu M$ ) in the absence and presence of HA1077 (6 or  $30 \mu M$ ) for the indicated times at which the tissue was snap-frozen and  $LC_{20}$  phosphorylation quantified as described in Methods. A, representative Western blots of  $LC_{20}$  after separation of the phosphorylated from the unphosphorylated species by urea-glycerol gel electrophoresis. B, quantitative data pooled from several experiments ( $n = 6-16$ ). Control (●),  $6 \mu M$  HA1077 (▽),  $30 \mu M$  HA1077 (□). \* Significant differences from  $LC_{20}$  phosphorylation levels at time zero.

**Table 3. Effects of  $\text{Ca}^{2+}$ , GTP $\gamma$ S (100  $\mu\text{M}$ ) and HA1077 on the rate of microcystin-LR-induced contraction of permeabilized guinea-pig ileum**

Conditions	$t_{1/2}$ (s)	Steady-state force (mN)	$n$
pCa 9	$326.7 \pm 55.7$	$1.31 \pm 0.20$	8
pCa 9 + GTP $\gamma$ S	$305.2 \pm 40.5$	$1.71 \pm 0.21$	8
pCa 6.5	$46.0 \pm 2.6$	$1.69 \pm 0.25$	7
pCa 6.5 + GTP $\gamma$ S	$47.9 \pm 2.4$	$1.51 \pm 0.19$	7
pCa 6.5 + GTP $\gamma$ S + 6 $\mu\text{M}$ HA1077	$47.0 \pm 4.2$	$1.72 \pm 0.11$	7
pCa 6.5 + GTP $\gamma$ S + 30 $\mu\text{M}$ HA1077	$46.1 \pm 4.2$	$1.68 \pm 0.13$	6

$\beta$ -Escin-permeabilized guinea-pig ileum longitudinal smooth muscle strips were induced to contract by addition of microcystin-LR (10  $\mu\text{M}$ ) under the conditions indicated. The time to half-maximal contraction ( $t_{1/2}$ ) and amplitude of steady-state force were determined. Values are given as means  $\pm$  S.E.M.

peak of LC<sub>20</sub> phosphorylation and abolished the second peak (sustained component) of LC<sub>20</sub> phosphorylation (Fig. 8A and B).

### ROK induces LC<sub>20</sub> phosphorylation via inhibition of MLCP

Previous analyses have indicated that the mechanism of increased LC<sub>20</sub> phosphorylation induced by GTP $\gamma$ S or agonists plus GTP involves inhibition of MLCP activity rather than increased MLCK activity. To determine if the ROK-mediated increase in LC<sub>20</sub> phosphorylation in guinea-pig ileum is due to MLCP inhibition, MLCK activation or direct phosphorylation of myosin by ROK, we carried out the following experiments. Firstly, to determine if ROK affects MLCK activity, we examined the effects of GTP $\gamma$ S in the absence and presence of HA1077 on the rates of contraction induced by the phosphatase inhibitor microcystin-LR (Table 3). Increasing  $[\text{Ca}^{2+}]$  from pCa 9 to pCa 6.5 markedly reduced the half-time of microcystin-LR-induced contraction ( $t_{1/2}$ ) from  $>300$  s to  $<50$  s ( $P < 0.001$ ),

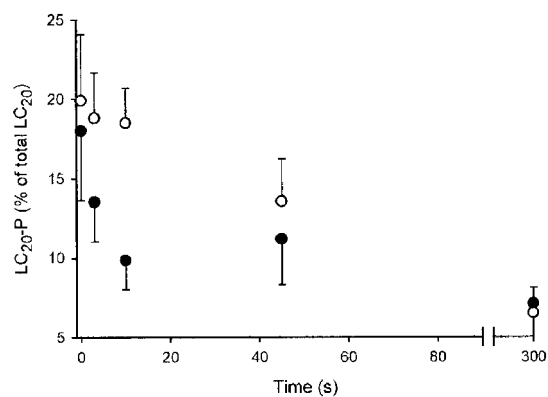
**Table 4. Effect of HA1077 on the rate of relaxation of permeabilized guinea-pig ileum**

Solution	$t_{1/2}$		$P$
	Control (s)	HA1077 (s)	
pCa 9	$34.6 \pm 1.7$	$22.3 \pm 1.4^*$	0.0016
pCa 9 rigor	$32.7 \pm 0.45$	$21.0 \pm 1.9^*$	0.0008
pCa 9 rigor + 300 $\mu\text{M}$ ML-9	$50.4 \pm 3.9$	$36.6 \pm 2.2^*$	0.013

$\beta$ -Escin-permeabilized guinea-pig ileum longitudinal smooth muscle strips were maximally contracted at pCa 4.5 in the presence of GTP $\gamma$ S (100  $\mu\text{M}$ ). HA1077 (30  $\mu\text{M}$ ) or vehicle (control) was added 5 min prior to relaxation in three different solutions: pCa 9 containing ATP and an ATP regenerating system, pCa 9 without ATP or an ATP regenerating system (rigor conditions) or pCa 9 under rigor conditions in the presence of ML-9 (300  $\mu\text{M}$ ). All relaxing solutions contained GTP $\gamma$ S (100  $\mu\text{M}$ ). Values are given as means  $\pm$  S.E.M. ( $n = 5$ ). \*Significantly different from the corresponding response in the absence of HA1077 with the indicated  $P$  values.

but the amplitude of steady-state force was the same in both conditions. GTP $\gamma$ S (100  $\mu\text{M}$ ) had no significant effect on  $t_{1/2}$  or the amplitude of steady-state force at pCa 9 or 6.5. Furthermore, HA1077 (6 or 30  $\mu\text{M}$ ) had no effect on  $t_{1/2}$  or steady-state force at pCa 6.5 in the presence of GTP $\gamma$ S. We conclude, therefore, that ROK has no effect on MLCK. These results also suggest that ROK does not phosphorylate myosin directly *in situ*.

To determine whether ROK activation leads to inhibition of MLCP, we examined the effect of HA1077 on the rate of relaxation under conditions of zero MLCK activity. Contraction of permeabilized guinea-pig ileum longitudinal smooth muscle was elicited at pCa 4.5 in the presence of 100  $\mu\text{M}$  GTP $\gamma$ S. We then used three sets of conditions to inhibit MLCK activity: pCa 9, pCa 9 without ATP or an ATP regenerating system (rigor conditions) and pCa 9 without ATP in the presence of the MLCK inhibitor ML-9 (Table 4). Under all three conditions, the rate of relaxation was significantly enhanced by HA1077 (Table 4), suggesting that ROK activation leads to inhibition of MLCP. To test



**Figure 9. Effect of HA1077 on the rate of LC<sub>20</sub> dephosphorylation in permeabilized guinea-pig ileum**

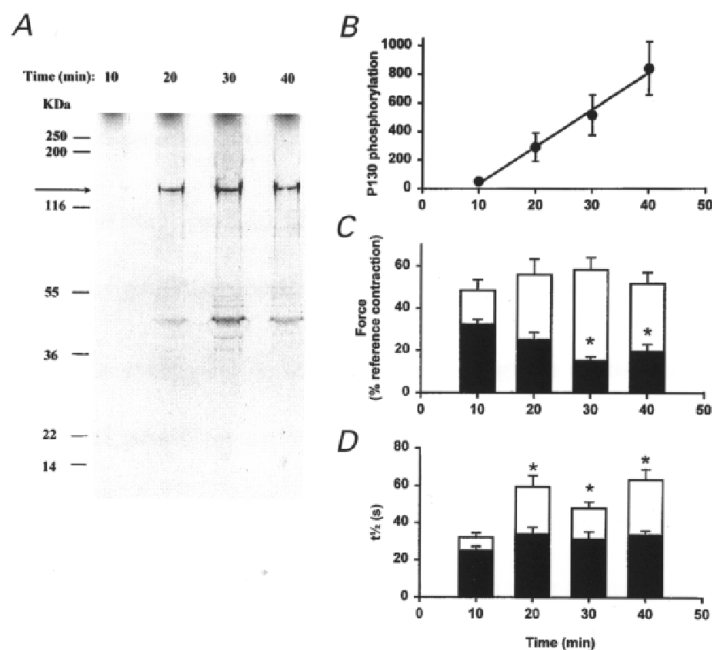
$\beta$ -Escin-permeabilized guinea-pig ileum longitudinal smooth muscle strips were maximally contracted at pCa 4.5 in the presence of GTP $\gamma$ S (100  $\mu\text{M}$ ). HA1077 (30  $\mu\text{M}$ , ●) or vehicle (control, ○) was added 5–10 min prior to relaxation which was initiated at time zero by transfer to rigor solution at pCa 9 containing GTP $\gamma$ S (100  $\mu\text{M}$ ). Tissues were snap-frozen at the indicated times for quantification of LC<sub>20</sub> phosphorylation levels (LC<sub>20</sub>-P) as described in Methods.

this hypothesis more directly, we measured LC<sub>20</sub> dephosphorylation at various times during relaxation (Fig. 9). Strips were again contracted in pCa 4 solution with GTP $\gamma$ S for 5 min and then either maintained in the same solution for 5–10 min or transferred to solution containing HA1077 for 5–10 min. Strips were frozen immediately (0 s) or transferred to pCa 9 rigor solution with GTP $\gamma$ S in the absence or presence of HA1077 and frozen at the indicated times for quantification of LC<sub>20</sub> phosphorylation. Inhibition of ROK by HA1077 clearly increases the rate of myosin dephosphorylation (the half-time of dephosphorylation was  $50 \pm 11$  s ( $n=6$ ) in the absence of HA1077 and  $15 \pm 4$  s ( $n=6$ ) in the presence of HA1077;  $P=0.02$ ), indicating that ROK activation leads to inhibition of MLCP.

To test the hypothesis that the mechanism of inhibition of MLCP activity by ROK involves phosphorylation of MYPT, muscle strips were incubated with [<sup>35</sup>S]ATP $\gamma$ S in pCa 9 rigor solution containing 100  $\mu$ M ML-9 and thiophosphorylation of MYPT visualized by autoradiography as described by

Trinkle-Mulcahy *et al.* (1995). We observed time-dependent incorporation of <sup>35</sup>S predominantly into two proteins of 42 and 130 kDa (Fig. 10A and B). In Western blotting experiments, the 130 kDa protein comigrated with MYPT (results not shown). Thiophosphorylation of the 130 kDa protein was associated with enhanced Ca<sup>2+</sup> sensitivity and slowing of relaxation (increased  $t_{1/2}$ ) as shown in Fig. 10C and D, respectively. Collectively, these data support the hypothesis that phosphorylation of MYPT correlates with MLCP inhibition and Ca<sup>2+</sup> sensitization of contraction.

The effect of GTP $\gamma$ S on MYPT thiophosphorylation was then examined by coincubation of muscle strips with [<sup>35</sup>S]ATP $\gamma$ S and GTP $\gamma$ S in pCa 9 rigor solution. Results of a typical experiment are shown in Fig. 11A, which demonstrates increased thiophosphorylation of MYPT in the presence of GTP $\gamma$ S. The Coomassie Blue-stained filamin band was used to correct for variations in protein loading levels (Fig. 11B). Cumulative data (Fig. 11C) indicate significant increases in MYPT thiophosphorylation levels in



**Figure 10.** Thiophosphorylation of a 130 kDa protein correlates with Ca<sup>2+</sup> sensitization of contraction

$\beta$ -Escin-permeabilized guinea-pig ileum longitudinal smooth muscle strips were incubated with 100  $\mu$ M ML-9 in pCa 9 rigor solution and 10  $\mu$ M [<sup>35</sup>S]ATP $\gamma$ S was added at time zero. Muscle strips were snap-frozen at the indicated times and proteins extracted and separated by SDS-PAGE as described in Methods. The gels were stained with Coomassie Blue and thiophosphorylated proteins detected by autoradiography. A, a representative autoradiogram. The 130 kDa thiophosphorylated protein (identified as MYPT by Western blotting) is indicated by the arrow. The positions of molecular weight markers are indicated to the left. Note the absence of LC<sub>20</sub> thiophosphorylation. B, cumulative data ( $n=3$ ) showing the time course of MYPT thiophosphorylation. Values were normalized to the Coomassie Blue-stained calponin band to correct for variations in protein loading levels. C, to evaluate the mechanical effect of MYPT thiophosphorylation, permeabilized muscle strips were incubated in the absence (■) or presence of ATP $\gamma$ S (□) as described above. Tissues were then washed in pCa 9 solution ( $3 \times 5$  min) and contracted in pCa 6.0 solution. \* Significant difference ( $P < 0.05$ ) from 10 min controls ( $n \geq 10$ ). D, after contraction in response to immersion in pCa 4.5 solution, relaxation was recorded at high chart speed to obtain  $t_{1/2}$  values for relaxation. \* Significant difference from the 10 min plus ATP $\gamma$ S value. In C and D, values recorded in the presence of ATP $\gamma$ S differ significantly from controls at all times.

the presence of GTP $\gamma$ S at 5 and 10 min. It is also noteworthy that no thiophosphorylation of LC<sub>20</sub> was detected (Fig. 11A). Thiophosphorylation for 40 min (under conditions defined in Fig. 10) did not lead to attenuation of the subsequent sensitization of force by GTP $\gamma$ S at pCa 6.25 (Fig. 11D), suggesting that MYPT phosphorylation had not reached saturation by this time.

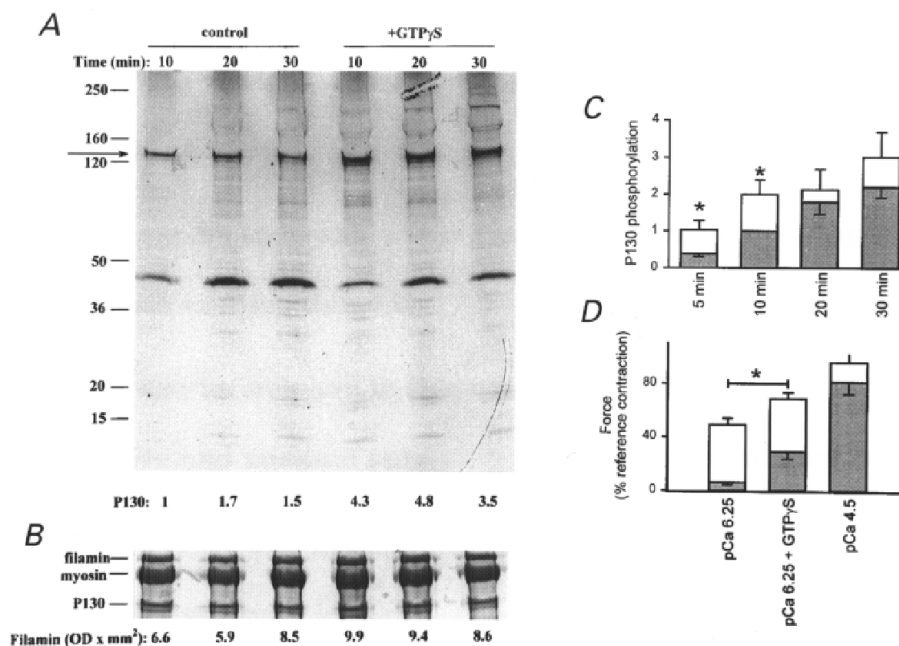
To determine the mechanical effect of enhanced MYPT phosphorylation it was necessary to determine the reversibility of Ca<sup>2+</sup> sensitization of force by GTP $\gamma$ S. A 10 min incubation with 100  $\mu$ M GTP $\gamma$ S in pCa 9 rigor solution without ATP $\gamma$ S caused subsequent Ca<sup>2+</sup> sensitization after more than 15 washes (5 min each) in fresh solution with no significant attenuation of the effect (results not shown). The effect of GTP $\gamma$ S is therefore irreversible within the time frame of the experiment, due perhaps to very slow dissociation from RhoA. Ca<sup>2+</sup> sensitization by carbachol plus GTP, on the other hand, was rapidly reversible, requiring fewer than three 5 min washes (results not shown; see also Otto *et al.* 1996), and co-incubation of ATP $\gamma$ S (10  $\mu$ M) with carbachol (100  $\mu$ M) and GTP (10  $\mu$ M) for 10 min, while not leading to increased

basal force in the subsequent washes in pCa 9 solution (with MgATP), approximately doubled the subsequent response to pCa 6.25, as shown in Fig. 12. This effect was completely reversed by co-incubation with Y-27632 (30  $\mu$ M) and attenuated by HA1077 (30  $\mu$ M). The elevated response to pCa 6.25 after ATP $\gamma$ S pre-treatment (Figs 10C and 11D) was not affected by either Y-27632 or HA1077, suggesting that the kinase responsible for basal MYPT phosphorylation is probably distinct from ROK.

Finally, the effect of ROK inhibition on GTP $\gamma$ S-induced MYPT thiophosphorylation was examined. As shown by the representative autoradiogram (Fig. 13A) and the cumulative data (Fig. 13B), the GTP $\gamma$ S-induced increase in MYPT thiophosphorylation was inhibited by HA1077.

## DISCUSSION

The results presented in this paper indicate that ROK activation is the predominant, if not the only, mechanism causing Ca<sup>2+</sup> sensitization in guinea-pig ileum longitudinal smooth muscle. The high degree of selectivity of Y-27632 for ROK (Uehata *et al.* 1997) and the observed correlation



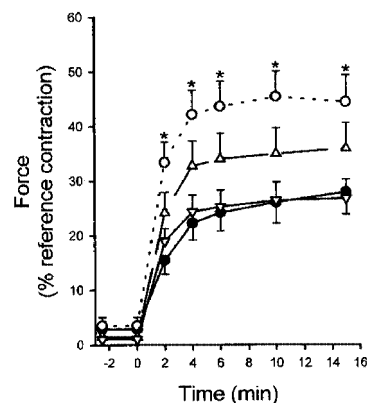
**Figure 11. GTP $\gamma$ S enhances MYPT thiophosphorylation**

$\beta$ -Escin-permeabilized guinea-pig ileum longitudinal smooth muscle strips were incubated in pCa 9 rigor solution containing 10  $\mu$ M [<sup>35</sup>S]ATP $\gamma$ S in the absence or presence of 1  $\mu$ M GTP $\gamma$ S. Muscle strips were processed as described in the legend to Fig. 10. *A*, a representative autoradiogram. Numbers below each lane indicate the radioactivity in the 130 kDa band normalized to the 10 min control value. *B*, a portion of the Coomassie Blue-stained gel to show the filamin band that was used to correct for variations in protein loading levels. Numbers below the gel lanes denote the optical density  $\times$  area of the filamin band in each lane. *C*, cumulative data ( $n \geq 5$ ) showing the time course of MYPT thiophosphorylation in the absence (shaded bars) and presence (open bars) of GTP $\gamma$ S. Values are corrected for background and protein loading and are normalized to the control (no GTP $\gamma$ S) value at 10 min. *D*, permeabilized muscle strips were incubated in pCa 9 rigor solution containing 100  $\mu$ M ML-9 in the absence (shaded bars;  $n = 4$ ) or presence (open bars;  $n = 4$ ) of 10  $\mu$ M ATP $\gamma$ S. After washing, muscle strips were contracted in pCa 6.25 solution. At the ensuing plateau, 100  $\mu$ M GTP $\gamma$ S was added. Finally, muscle strips were contracted in pCa 4.5 solution.

**Figure 12. Addition of carbachol and GTP during ATP $\gamma$ S pre-incubation enhances the subsequent response to Ca $^{2+}$  which is blocked by ROK inhibition**

$\beta$ -Escin-permeabilized guinea-pig ileum longitudinal smooth muscle strips were pre-incubated in pCa 9 rigor solution containing 10  $\mu$ M ATP $\gamma$ S with the following additions: none (●); 100  $\mu$ M carbachol and 10  $\mu$ M GTP (○); 100  $\mu$ M carbachol, 10  $\mu$ M GTP and 30  $\mu$ M Y-27632 (▽); or 100  $\mu$ M carbachol, 10  $\mu$ M GTP and 30  $\mu$ M HA1077 (Δ). Muscle strips were then washed in pCa 9 solution containing MgATP to ensure washout of acute sensitization by carbachol plus GTP. Force development upon re-addition of MgATP was not significant in either condition but differed during the subsequent pCa 6.25-induced contraction (initiated at time zero).

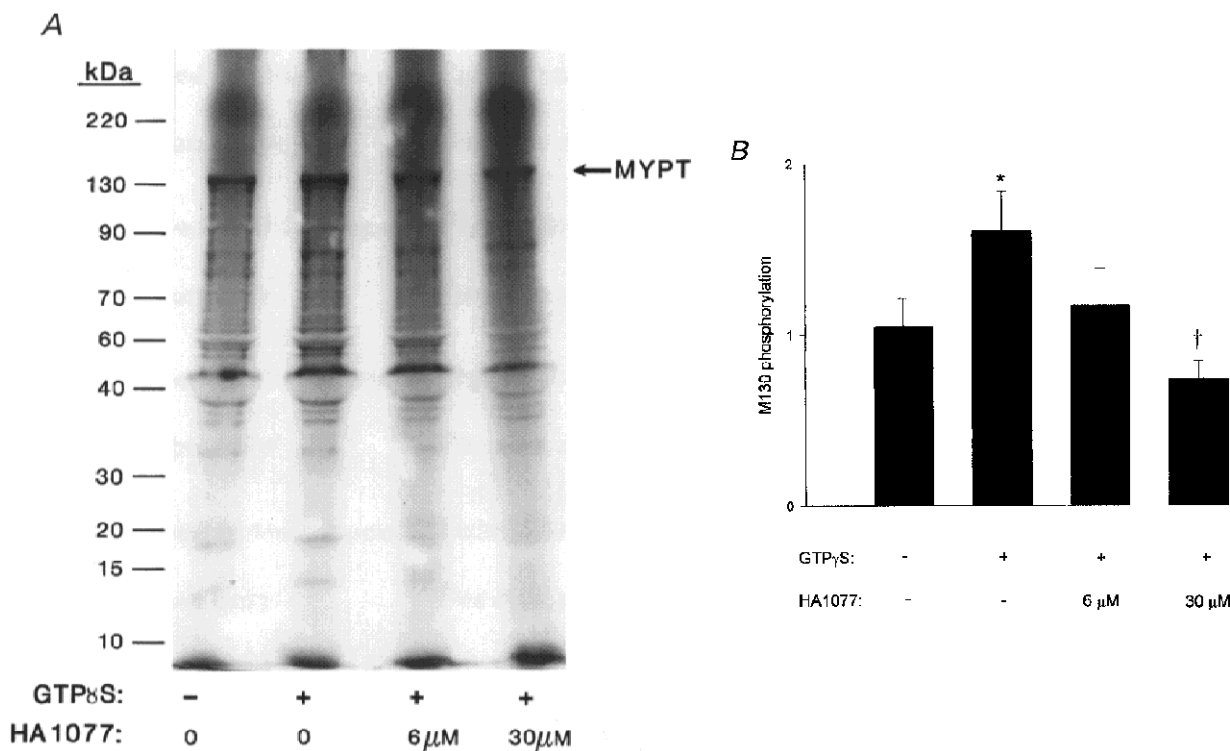
\*Significant difference compared with control (ATP $\gamma$ S alone).



between inhibition of Ca $^{2+}$  sensitization and inhibition of isolated ROK by three inhibitors (Y-27632, HA1077 and H-7), suggests that the relevant inhibited kinase is indeed ROK.

HA1077, used in the majority of the present experiments, shows good selectivity for ROK over PKC and MLCK (Uehata *et al.* 1997; this study). However, it also inhibits cAMP- and cGMP-dependent protein kinases with  $K_i$

values ~5-fold greater than for that ROK (Asano *et al.* 1989). This possible complication is irrelevant in the context of this work since inhibition of the cyclic nucleotide-dependent kinases would cause contraction rather than relaxation (cf. Lee *et al.* 1997, and references therein). Furthermore, we failed to observe relaxation of  $\beta$ -escin-permeabilized ileum by a combination of the adenylyl cyclase activator forskolin (10  $\mu$ M) and the phosphodiesterase



**Figure 13. Inhibition of the GTP $\gamma$ S-induced increase in MYPT thiophosphorylation by ROK inhibition**

$\beta$ -Escin-permeabilized guinea-pig ileum longitudinal smooth muscle strips were incubated in pCa 9 rigor solution. At time zero, 10  $\mu$ M [ $^{35}$ S]ATP $\gamma$ S was added without or with 1  $\mu$ M GTP $\gamma$ S in the absence or presence of HA1077 (6 or 30  $\mu$ M) as indicated. Muscle strips were processed after 10 min as described in the legend to Fig. 10. *A*, a representative autoradiogram. *B*, cumulative data ( $n = 8$ ) showing the GTP $\gamma$ S-induced increase in MYPT thiophosphorylation and its prevention by HA1077. \*Significant difference from control. †Significant difference from value measured in the presence of GTP $\gamma$ S alone. More thiophosphorylated bands are visible in *A* than in Fig. 11*A* because the gel was exposed to the X-ray film for a longer period of time.

inhibitors rollipram (30  $\mu\text{M}$ ) and OPC3911 (3  $\mu\text{M}$ ) (results not shown).

The lack of effect of PDBu in guinea-pig ileum, which may relate to a low level of expression of a particular PKC isoform or the endogenous cytosolic protein inhibitor of myosin light chain phosphatase, CPI-17 (Li *et al.* 1998), suggests that conventional and/or novel PKC isoforms play a minor role (if any) in  $\text{Ca}^{2+}$ -independent modulation of contractility in the ileum. This is in contrast to rat mesenteric small arteries in which PKC inhibition by calphostin C caused complete attenuation of  $\text{Ca}^{2+}$  sensitization (Buus *et al.* 1998) and rabbit femoral artery and portal vein in which a pseudosubstrate peptide inhibitor based on PKC $\alpha$  reduced  $\text{Ca}^{2+}$  sensitization and myosin phosphorylation (Gailly *et al.* 1997).

In permeabilized guinea-pig ileum, addition of  $\text{GTP}\gamma\text{S}$  increased  $\text{LC}_{20}$  phosphorylation and this was reversed by ROK inhibition. At the lower  $[\text{Ca}^{2+}]_i$  (pCa 6.5), changes in  $\text{LC}_{20}$  phosphorylation by  $\text{GTP}\gamma\text{S}$  and HA1077 were relatively small (significantly different only after normalization to pCa 6.5 in each experiment) compared with the higher  $[\text{Ca}^{2+}]_i$  (pCa 5.75) but may nevertheless be important if the  $\text{LC}_{20}$  phosphorylation–force relationship is very steep. This has indeed been demonstrated to be the case in several smooth muscles. For instance, Pfitzer & Arner (1998) showed that force changed from <10 to >90% for an ~8% change in  $\text{LC}_{20}$  phosphorylation in the guinea-pig ileum. The present data thus support the hypothesis that the change in force by ROK inhibition is mediated by changes in  $\text{LC}_{20}$  phosphorylation in the permeabilized tissue.

To assess the functional effects of ROK inhibition under more physiological circumstances, intact strips were used. In strips loaded with fura-2 AM, contraction in response to carbachol was reduced by ROK inhibition to a transient peak coinciding with peak  $[\text{Ca}^{2+}]_i$ .  $[\text{Ca}^{2+}]_i$  on the other hand was unchanged. The effect of ROK inhibition by HA1077 in guinea-pig ileum is almost identical to the effect of treatment with toxin B from *Clostridium difficile* which glucosylates the Ras-related low molecular mass GTPases of the Rho subfamily thereby inactivating them. This treatment caused complete reduction of sustained contraction without associated changes in  $[\text{Ca}^{2+}]_i$  (Lucius *et al.* 1998). ROK may thus be the sole RhoA effector for  $\text{Ca}^{2+}$  sensitization in this tissue.

Even though ROK inhibition abolished the tonic phase of contraction this does not imply that the rise in  $[\text{Ca}^{2+}]_i$  and the consequent increase in MLCK activity is not important during this phase. It has been demonstrated frequently that extracellular  $\text{Ca}^{2+}$  is necessary for maintained contraction and that inhibition of MLCK reduces or abolishes maintained contraction. An interesting comparison can be made between our results in intact and permeabilized muscles. After a transient peak following stimulation of intact muscle with carbachol,  $[\text{Ca}^{2+}]_i$  settled at a steady level of  $202 \pm 33$  nM which corresponds to a pCa of 6.69 and this was associated

with a sizeable contraction that was, however, prevented by ROK inhibition. In the permeabilized tissue, stimulation with pCa 6.5 in the absence of  $\text{GTP}\gamma\text{S}$ , i.e. a slightly higher  $[\text{Ca}^{2+}]_i$ , while drastically increasing the rate of contraction in response to the phosphatase inhibitor microcystin-LR, hardly elevated steady-state force relative to that achieved at pCa 9. Although a time-dependent change in the dependence of contraction on  $\text{LC}_{20}$  phosphorylation (latch) is likely to be important for this difference, our results also imply that physiological elevations in  $[\text{Ca}^{2+}]_i$  and consequent MLCK activation are not sufficient to cause a full force response unless GTPase-dependent pathways leading to MLCP inhibition are simultaneously activated. It therefore seems likely that maintained contraction in response to agonist stimulation depends on both MLCK and ROK activities.

$\text{LC}_{20}$  phosphorylation in response to carbachol in the intact tissue was elevated only at 3 s in the presence of HA1077, whereas carbachol-induced  $\text{LC}_{20}$  phosphorylation in the absence of HA1077 was elevated for at least 200 s. The design of these experiments, with controls run *versus* each HA1077 concentration (6 and 30  $\mu\text{M}$ ) individually, precludes direct comparison of the effects of the two concentrations but allows us to conclude that selective ROK inhibition in the intact tissue is associated with an altered phosphorylation–time profile. Interestingly, Lucius *et al.* (1998) did not detect an increase in  $\text{LC}_{20}$  phosphorylation following toxin B treatment, even though the phasic component of the contractile response to carbachol plus GTP was retained. They concluded therefore that the carbachol-induced force transient occurred independently of myosin phosphorylation. Our results with HA1077 show that ROK inhibition reduces, but does not eliminate, the phasic increase in  $\text{LC}_{20}$  phosphorylation in response to carbachol treatment. Whether or not this lower level of  $\text{LC}_{20}$  phosphorylation accounts for the phasic contractile response is uncertain.

Our analysis of the dynamics of contraction and myosin phosphorylation indicates that inhibition of ROK affects the rate of dephosphorylation. The effects of agonist or  $\text{GTP}\gamma\text{S}$  were similarly found to be associated with reduced MLCP activity (Kitazawa *et al.* 1991*b*) with no effect on MLCK activity. The use of microcystin and/or  $\text{ATP}\gamma\text{S}$  in determination of the effect of  $\text{GTP}\gamma\text{S}$  on phosphorylation rates *in situ* might cause activation of ROK-catalysed myosin phosphorylation by hyperphosphorylation of upstream signalling components. This would be detected by direct ROK inhibition and does not seem to be the case since HA1077 failed to affect rates of contraction and thus presumably the rate of phosphorylation in the present study. Although rates of phosphorylation were not measured directly, these contractions were much too slow to be limited by crossbridge turnover (~1/s) and, if rates of phosphorylation were increased by  $\text{GTP}\gamma\text{S}$  (contrary to several reports in the literature), an explanation would need to be invoked for the increased lag phase between

phosphorylation and force since  $t_{1/2}$  values for contraction were unchanged.

There are additional observations which suggest that ROK acts primarily through inhibition of MLCP activity and not through direct phosphorylation of myosin. Firstly, activation of ROK by GTP $\gamma$ S did not induce Ca<sup>2+</sup> sensitization of force at pCa 9 (results not shown) and did not thiophosphorylate LC<sub>20</sub> whereas it did thiophosphorylate MYPT (Figs 11 and 13). Secondly, co-incubation of ATP $\gamma$ S with carbachol plus GTP at pCa 9 did not lead to significant force development upon re-addition of MgATP (as would have been expected if myosin had been phosphorylated) but only enhanced the subsequent response to pCa 6.25 (Fig. 12). Studies with purified ROK, on the other hand, have demonstrated that intact myosin and isolated LC<sub>20</sub> are both good ROK substrates *in vitro* (Amano *et al.* 1996) and addition of a constitutively active fragment of ROK caused contraction via direct phosphorylation of myosin at serine-19, the MLCK site (Kureishi *et al.* 1997).

It is interesting to note that the effect of ROK inhibition did not appear to be attenuated by the lack of ATP in the solutions (Table 3). This may be due either to a slow turnover of phosphate on MYPT or to the presence of residual ATP in the tissue. The latter possibility is more likely since no qualified effort, such as apyrase treatment, was made to remove high energy phosphates and human ROK has been shown to have an unusually low  $K_m$  for ATP (0.1  $\mu$ M; Uehata *et al.* 1997), although Feng *et al.* (1999) determined a higher value (30.8  $\mu$ M) for the chicken gizzard enzyme.

It is also clear from our results that there is no absolute Ca<sup>2+</sup> requirement for GTP $\gamma$ S-induced MYPT phosphorylation. The Ca<sup>2+</sup> dependence of sensitization of force and myosin phosphorylation thus appears to be downstream of RhoA/ROK activation and MYPT phosphorylation and is probably due to MLCK as discussed above.

Results from physiological studies such as this, and the studies of Kitazawa *et al.* (1991*b*), Kubota *et al.* (1992) and Noda *et al.* (1995), are thus not reconcilable with biochemical investigations with respect to direct phosphorylation of myosin. The cumulated physiological evidence argues that direct ROK-catalysed phosphorylation of myosin in the contractile apparatus is not of sufficient magnitude to cause contraction, at least in the absence of Ca<sup>2+</sup>. It could be speculated that ROK is spatially restricted, with limited access to myosin, but this would require a mobile messenger such as RhoA to translocate to the contractile apparatus following activation at the membrane. ROK isoform variation could perhaps also explain the lack of evidence for direct myosin phosphorylation and could contribute to variability of smooth muscle responses to neurotransmitters, autacoids and drugs *in vivo*. Whatever the case, the present study indicates that ROK plays an important role in myosin phosphorylation and contraction in intestinal smooth muscle and that this occurs through modulation of MLCP activity.

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