Omega-3 and omega-6 DPA inhibits Ca²⁺-sensitization of vascular smooth muscle contraction induced by sphingosylphosphorylcholine via inhibiting Rho-kinase activation and translocation Ying Zhang, Min Zhang, Bochao Lyu, Hiroko Kishi, and Sei Kobayashi^{*}

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Supplementary Video S1 Time-lapse video of SPC-induced contraction of vascular smooth muscle cells. Images were acquired at 30 second intervals for 30 min.

Supplementary Video S2 Time-lapse video of SPC-induced contraction of vascular smooth muscle cells pretreated with n-3 DPA (60 μM) for 30 min. Images were acquired at 30 second intervals for 30 min.

Supplementary Video S3 Time-lapse video of SPC-induced contraction of vascular smooth muscle cells pretreated with n-6 DPA (60 μM) for 30 min. Images were acquired at 30 second intervals for 30 min.

The chemical structures of n-3 DPA, n-6 DPA and EPA.



- a SPC-induced contraction in the presence of vehicle.
- b 40 mM K⁺-induced contraction in the presence of vehicle.
- c The inhibitory extent of contraction induced by n-3 DPA or n-6 DPA was calculated by a percentage of the response to the contraction induced by 30 μM SPC.



a: The contraction of VSM induced by SPC.
b: The contraction of VSM in the presence of DPA.
c: The contraction of VSM in the presence of vehicle.
Inhibitory ratio of DPA (%) = (a-b-c)/a × 100%.

Induction of contractile type of human CASMCs after

serum-free medium treatment for 2 days. Scale bar =200 μ m.

b

a Phase-contrast micrograph of human CASMCs

in normal medium.

b Phase-contrast micrograph of human CASMCs in serum-free medium for 2 days.





- a Control image of immunofluorescent staining in the presence of second antibody. Scale bar = 10 μ m.
- b Translocation of Fyn induced by SPC in the presence of EPA and DPA. Scale bar = 100 μ m.



b



As a reference, the picture (EPA+ SPC+) obtained from our previous publication (Ref. 27) is shown.