

レプトスピラのホスホリパーゼ  
 I. *Leptospira biflexa* におけるホスホリパーゼ A<sub>1</sub> とリゾホスホリパーゼの存在

柳原 保武\* 谷山 忠義\* 美崎 英生\* 鈴木 康夫\*  
 松本 亮\* 三瀬 一二

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Phospholipases of *Leptospira*

I. Presence of Phospholipase A<sub>1</sub> and Lysophospholipase in *Leptospira biflexa*

**ABSTRACT** The hydrolysis of phosphatidylethanolamine, phosphatidylcholine, lysophosphatidylcholine, and trioleoylglycerol by *Leptospira biflexa* strain Urawa was studied *in vitro*.

Phospholipase A<sub>1</sub> was identified by the formation of <sup>32</sup>P- and <sup>14</sup>C-labeled lysoderivatives from <sup>32</sup>P-phosphatidylcholine, <sup>32</sup>P-phosphatidylethanolamine, or 1-acyl-2-[1-<sup>14</sup>C]-oleoyl-*sn*-glycero-3-phosphorylcholine. Phospholipase A<sub>1</sub> activity was independent of lipase in the microorganism since <sup>14</sup>C-labeled trioleoylglycerol was scarcely attacked under the same conditions in which the phospholipids were hydrolyzed.

Lysophospholipase activity was also demonstrated using <sup>32</sup>P- and non-labeled lysophosphatidylcholine.

The activity of phospholipase A<sub>1</sub> was found in a broad range of pH but no optimal pH was determined. The pH optimum of lysophospholipase was 8.0. Both enzymes were labile to heat.

Phospholipase C activity, however, could not be detected because no radio-active di- and mono-acylglycerol was found in the experiment with 1-acyl-2-[1-<sup>14</sup>C]-oleoyl-*sn*-glycero-3-phosphorylcholine as the substrate.

It was inferred that phosphatidylethanolamine, which was the major component of phospholipids in leptospirae, was hydrolyzed serially by phospholipase A (A<sub>1</sub> and/or A<sub>2</sub>?) and lysophospholipase to glycerophosphorylethanolamine *via* 2-acyl-type-lyso-derivative as one metabolic pathway of the substrate.

抄録 *Leptospira biflexa* 浦和株のホスファチジルエタノールアミン, ホスファチジルコリン, リゾホスファチジルコリン及びトリオレイルグリセロールの加水分解を調べた。ホスホリパーゼ A<sub>1</sub> は <sup>32</sup>P-ホスファチジルコリン, <sup>32</sup>P-ホスファチジルエタノールアミンや 1-アシル-2-[1-<sup>14</sup>C]オレイル-*sn*-グリセロ-3-ホスホリルコリンから <sup>32</sup>P-と <sup>14</sup>C-標識のリゾ誘導体の作られることで確認した。ホスホリパーゼ A<sub>1</sub> 活性は, <sup>14</sup>C-標識トリオレイルグリセロールにほとんど作

用しないことから、レプトスピラ中のリパーゼとは別のものであった。リゾホスホリパーゼ活性も  $^{32}\text{P}$ -と非標識リゾホスファチジルコリンを用いて確認した。ホスホリパーゼ  $A_1$  には至適 pH は認められず、広い pH 範囲で活性が認められた。リゾホスホリパーゼの至適 pH は 8.0 であり、両酵素とも熱に不安定であった。ホスホリパーゼ C 活性は認められなかった。

\* Shizuoka College of Pharmacy 静岡薬科大学