Key words

Petersen olefination

titanium-mediated radical cyclization

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Total Synthesis of Pyripyropene A *Tetrahedron* **2011**, *67*, 8195–8203.

Synthesis of Pyripyropene A

Significance: Pyripyropene A is a fungal metabolite isolated from *Aspergillus fumigatus* FO-1289–2501. It is a potent inhibitor of Acyl-CoA:cholesterol-acyltransferase. Several semi-synthetic derivatives show even greater inhibition.

Comment: Key steps in this synthesis include the titanium-mediated epoxide opening–radical cyclization ($\mathbf{A} \to \mathbf{D}$) and a Petersen-olefination/epoxide-opening sequence ($\mathbf{F} \to \mathbf{J}$). A more scalable sequence from \mathbf{D} to \mathbf{F} involving Me₄NHB(OAc)₃ as an alternative reducing agent was also reported.

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