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# Delphi Xe3 Slip File Generator

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## Delphi Xe3 Slip File Generator

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=====*/
#ifndef BOOST_SPIRIT_INCLUDE_QI_ACTION #define BOOST_SPIRIT_INCLUDE_QI_
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page.Peroxisome Proliferator-Activated Receptors (PPARs) form a subfamily of the nuclear  
hormone receptor superfamily. These receptors control the expression of genes involved in  
lipid metabolism, cell differentiation, and inflammation (H. Tan et al. (1995) J. Biol. Chem.  
270:12953-12956; S. Kliewe (1995) Endocr. Rev. 16:1-22). PPARs contain three distinct  
domains: an N-terminal ligand binding domain, a DNA binding domain, and a C-terminal  
ligand binding domain. The ligand binding domain of PPARs may also be referred to as a  
ligand binding pocket, or LBP. PPARs are activated by fatty acids, fatty acid metabolites and  
eicosanoids. PPARs and their target genes have been implicated in the regulation of lipid  
metabolism and inflammation in the human cardiovascular system. PPARs are activated by  
fatty acids, fatty acid metabolites and eicosanoids. PPARs interact with other nuclear  
receptors, including members of the steroid hormone receptor superfamily, via the C-

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terminal ligand binding domain (H. Tan et al. (1995) J. Biol. Chem. 270:12953-12956; S. Kliewe (1995) Endocr. Rev. 16:1-22). PPARs are activated by fatty acids, fatty acid metabolites and eicosanoids. The ligand binding domain of PPARs may also be referred to as a ligand binding pocket, or LBP. The PPARs ( $\alpha$ ,  $\beta/\delta$ , and  $\gamma$ ) are distinct receptors that show some level of selectivity for arachidonic acid. These PPARs are activated by a number of compounds found in food products, including omega-3 and omega-6 fatty acids and their derivatives. Based on their pharmacological profiles, PPAR $\alpha$  is reported to be activated by a number of hypolipidemic drugs including clofibrate and Wy-14643