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The Reaction of Phosphonyl Radicals with Hindered Carbonyl Compounds

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The Reaction of Phosphonyl Radicals with Hindered Carbonyl Compounds

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The reaction between phosphonyl radicals and substituted oand p-benzoquinone has been studied by ESR. The addition of
phosphonyl radicals to benzoquinone occurs at two sites, either
at carbonyl oxygen or at double bond of the cycle. The reaction
of phosphonyl radicals with p-benzoquinone and methylenquinones
proceeds in a similar fashion.

$$\begin{array}{c}
\downarrow 0 \\
0 \\
0 \\
0 \\
R^{1}R(0)P
\end{array}$$

The ESR analysis of the 3,6-ditert.buthyl-2-oxyphenoxy radicals, containing the phosphoryl group, indicates the covalent character of 0-P bond. In the case of 3,5-ditert.buthyl-o-benzo-quinone two cyclohexadienyl radicals are formed due to the attack of the phosphonyl radical at the different positions of the quinone cycle. While phosphonyl radicals are added only to the double bond of 4,4-dialkyl-2,6-di-tert.buthylcyclohexa-2,5-dion-1-one cycle, the reaction with tert.buthyl diphenoquinone occurs both at the double bond of the cycle (at the 1 (1')- and 2 (2')-positions) and at the oxygen atom.