**Synthesis of Tetrahydropyrans through Formal [4+2] Cycloaddition**

**Aromatic aldehydes:**

\[
\begin{align*}
\text{R}^1 & = \text{Ph, 4-MeOC}_6\text{H}_4, 4-\text{BrC}_6\text{H}_4, (E)-\text{CH}=\text{CHMe} \\
\text{R}^2 & = \text{Ph, 4-MeC}_6\text{H}_4, 3-\text{BrC}_6\text{H}_4, 2-\text{FC}_6\text{H}_4, 4-\text{F}_2\text{CC}_6\text{H}_4, 4-\text{MeOC}_6\text{H}_4, 3-\text{O}_2\text{NC}_6\text{H}_4, 2-\text{ClC}_6\text{H}_4, 4-\text{ClC}_6\text{H}_4, (E)-\text{CH}=\text{CHPh}
\end{align*}
\]

**Aliphatic aldehydes:**

\[
\begin{align*}
\text{R}^1 & = \text{Ph, 4-MeOC}_6\text{H}_4, 4-\text{BrC}_6\text{H}_4, (E)-\text{CH}=\text{CHMe} \\
\text{R}^2 & = \text{n-Pent, i-Pr, Cy}
\end{align*}
\]

**One-pot procedure:**

\[
\begin{align*}
\text{Ar} & = 4-\text{MeOC}_6\text{H}_4 \\
R & = \text{Ph, 3-BrC}_6\text{H}_4, 4-\text{MeC}_6\text{H}_4
\end{align*}
\]

**Significance:** Although insertion reactions with cyclopropanes are well known in organic chemistry, analogous reactivity with cyclobutanes has remained largely overlooked. The authors describe a Lewis acid catalyzed reaction between cyclobutanes and aldehydes to yield substituted tetrahydropyrans in high diastereomeric purity. The reaction is further modified into a formally [(2+2)+2] one-pot process, which gives the products in a higher overall yield. In the future of this chemistry, chiral ligands can potentially be applied to yield enantiopure products.

**Comment:** Both aliphatic and aromatic aldehydes were successfully applied in the transformation, provided that the correct catalyst was used. The mechanism of the reaction is not well understood. In competition experiments products with more electron-rich aldehydes are favored, yet electron-poor aldehydes react more rapidly. The latter observation is proposed to result from catalyst poisoning and not from enhanced reactivity. Interestingly, enantiopure starting materials give rise to nearly racemic products, suggesting that racemization of the cyclobutane stereocenter occurs during the reaction. Notably, only cyclobutanes bearing aromatic substituents were utilized ($R^1 = \text{Ar}$), which may be a requirement for facile C–C bond cleavage.