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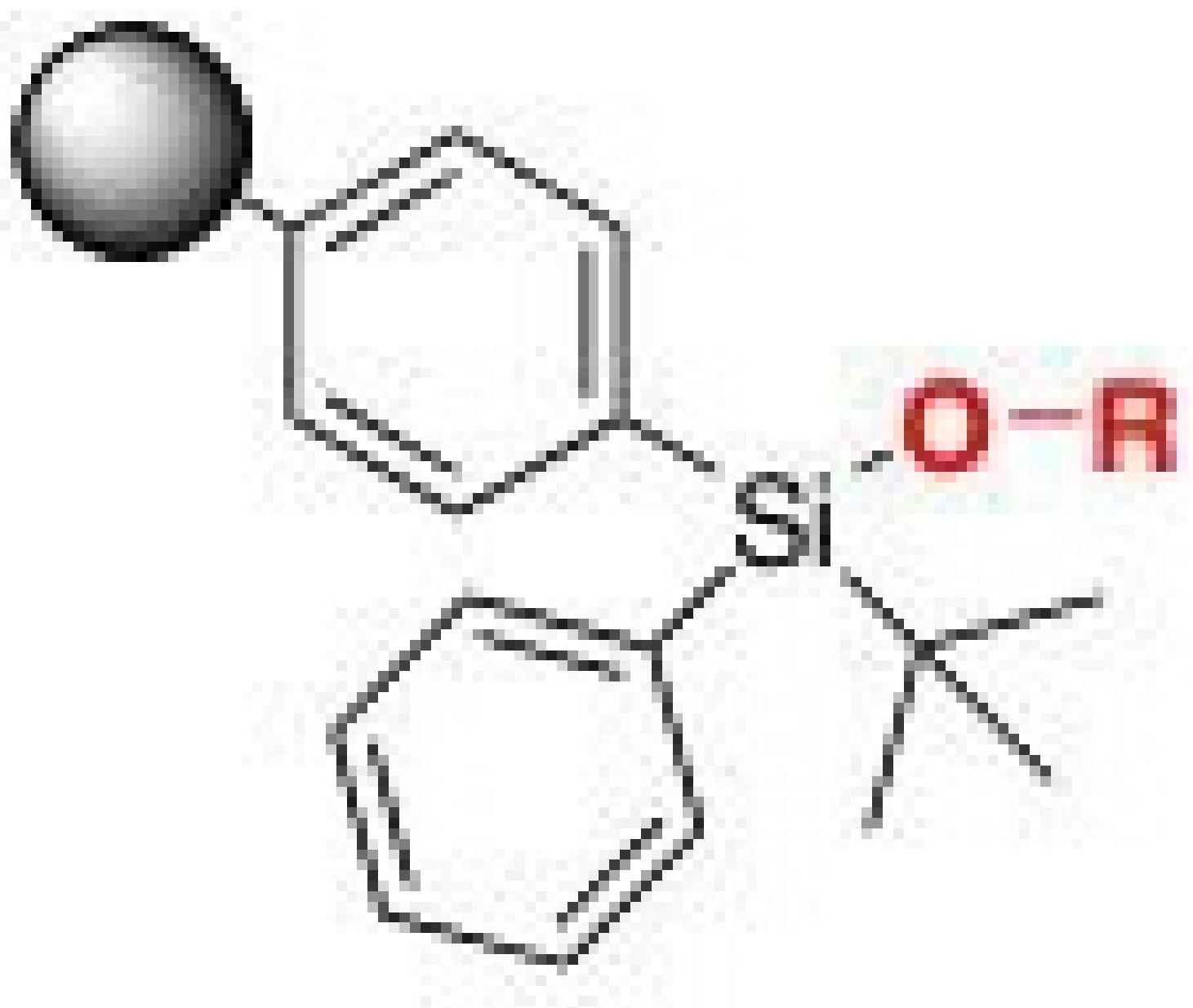
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## TBDAS Linker

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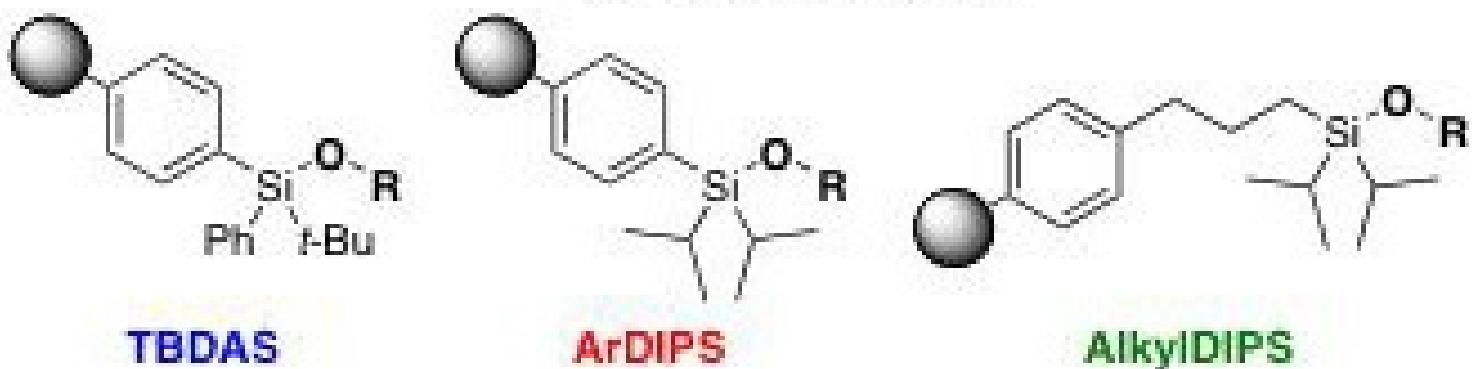
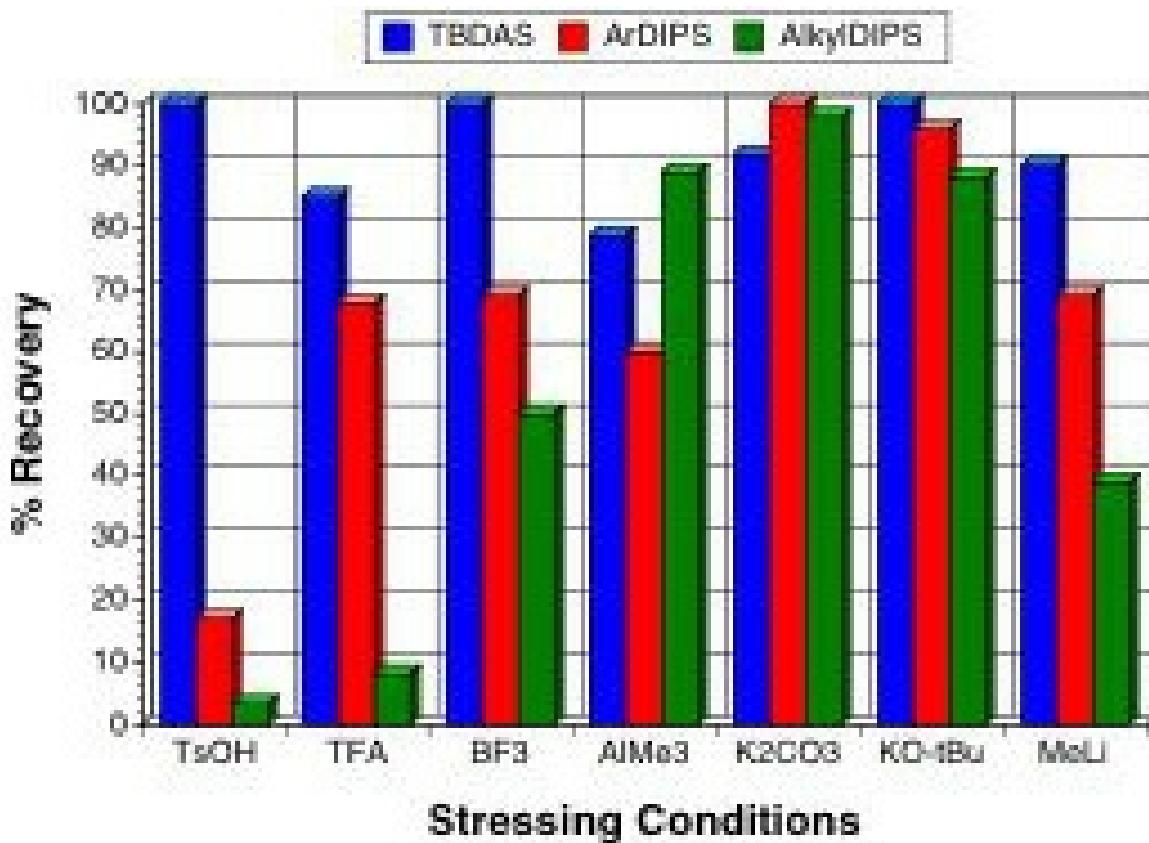
### A Robust Linker for Solid Phase Diversity-Oriented Synthesis

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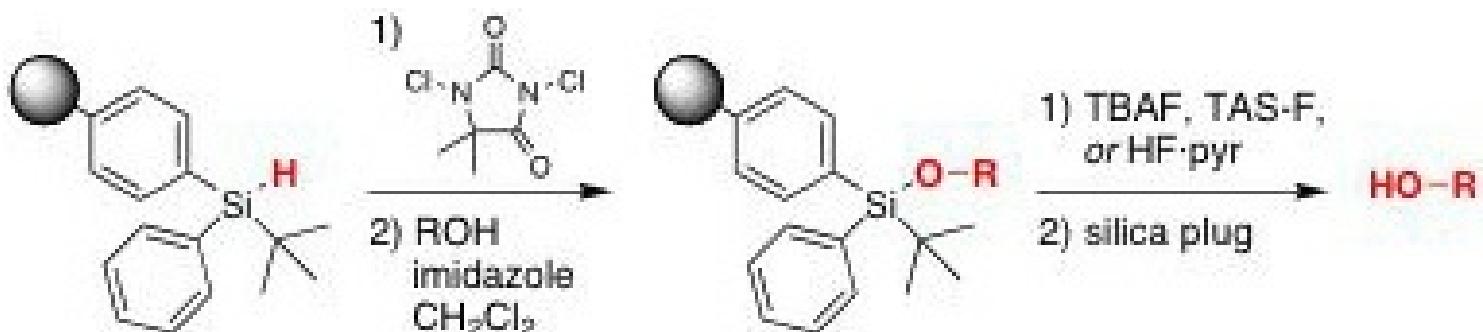


# The TBDAS Linker

The *t*-butyldiarylsilyl (TBDAS) linker was developed by Drs. [Christine DiBlasi](#) and [Daniel Macks](#) in our group for solid phase organic synthesis. This robust linker is stable to a variety of reaction conditions, but can be cleaved with relatively mild, chemoselective fluoride reagents to yield alcohol products.



As depicted graphically here, the TBDAS linker (blue) is stable to a range of acidic and basic reaction conditions commonly used in organic synthesis. In contrast, previously reported arylidisiopropylsilyl (ArDIPS, red) and alkylidisiopropylsilyl (AlkyIDIPS) linkers are generally less stable to these conditions.



The TBDAS linker is activated by chlorination with dichlorodimethylhydantoin, followed by loading of the desired alcohol-containing substrate. After synthetic transformation, the product is cleaved from the solid

support by treatment with TBAF, TAS-F, or HF·pyr. The product can be conveniently separated from the cleavage reagents by passage through a short plug of reverse phase over normal phase silica gel.

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*For full details on the synthesis and use of this linker, see:*

DiBlasi, C. M.; Macks, D. E.; Tan, D. S.\* “An acid-stable *tert*-butyldiarylsilyl (TBDAS) linker for solid-phase organic synthesis.” *Org. Lett.* 2005, 7, 1777-1780.

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