Synthesis of Sarpagine Alkaloids

Significance: The authors report the enantioselective total synthesis of three sarpagine indole alkaloids which were isolated from the plant family Apocynaceae. The route relies on a common intermediate $G$, which is impressively accessed using key features such as a $[5+2]$ oxidopyridinium cycloaddition and a ring expansion. The three natural products were synthesized in only eight steps starting from known materials (12 steps from commercially available compounds).

Comment: The synthesis commenced with a $[5+2]$ cycloaddition between oxidopyridinium salt $A$ and Aggarwal's chiral ketene equivalent $B$, thus yielding the desired regioisomer $C$ in a 2:1 ratio. Next, ketone $G$ was accessed through an intramolecular palladium-catalyzed enolate coupling of $D$, followed by Wittig reaction, deprotection of the dithiolane, and ring expansion. The indole was introduced in the last step by a Fischer indole synthesis using phenylhydrazines with different substitution patterns to afford the three targets.