Synthesis of Prostaglandin D<sub>2</sub> Receptor Antagonist

**Significance:** An efficient kilogram-scale synthesis of the target prostaglandin D<sub>2</sub> receptor antagonist features a Friedel–Crafts cyclization of an iminopyrrole to generate the azaindole core in D. Key steps are (1) a very efficient asymmetric hydrogenation to install the single stereogenic center (G → H) and (2) a mild sulfenylation using the shelf-stable N-arylsulfonfimidamide I.

**Comment:** The high yield of the hydrogenation was surprisingly insensitive to solvent, but it was sensitive to the E/Z ratio. Thus, batches of G that contained 9% of the Z-isomer afforded H in only 81% ee, whereas batches of G containing 1% of the Z-isomer gave H in 96% ee. The E/Z ratio of the Horner–Wadsworth–Emmons reaction (14:1) could be upgraded to 1000:1 by crystallizing the phosphate salt of G.