

**DEVELOPING A SCALABLE ROUTE TO THE PROLINE CORE OF  
VOXILAPREVIR: AN ACTIVE INGREDIENT IN VOSEVI® FOR THE  
PAN-GENOTYPIC TREATMENT OF HEPATITIS C**

Dunetz, J.; Chan, J.; Bonderoff, S.<sup>1</sup>; Bringley, D.; Cagulada, A.; Chan, L.; Duo, T.<sup>1</sup>;  
Gligoric-Trkulja, O.<sup>1</sup>; Huang, Z.; Keaton, K.; Phoenix, J.<sup>1</sup>; Ross, B.; Shah, N.;  
Shapiro, N.; Siler, D.; Tang, D.; Tiong, E.<sup>1</sup>; Williams, B.; Xu, D.; Yu, L.

Gilead Sciences, 333 Lakeside Drive, Foster City, CA 94404

<sup>1</sup>Gilead Alberta ULC, 1021 Hayter Road, Edmonton, Alberta, T6S 1A1, Canada  
*joshua.dunetz@gilead.com*

Vosevi® is a fixed-dose combination of sofosbuvir, velpatasvir, and voxilaprevir for the pan-genotypic treatment of hepatitis C infection. Voxilaprevir has a complex structure highlighted by eight stereocenters, three cyclopropanes, and a proline core. Key developments of a process route will be discussed with focus on the assembly of the proline fragment.

