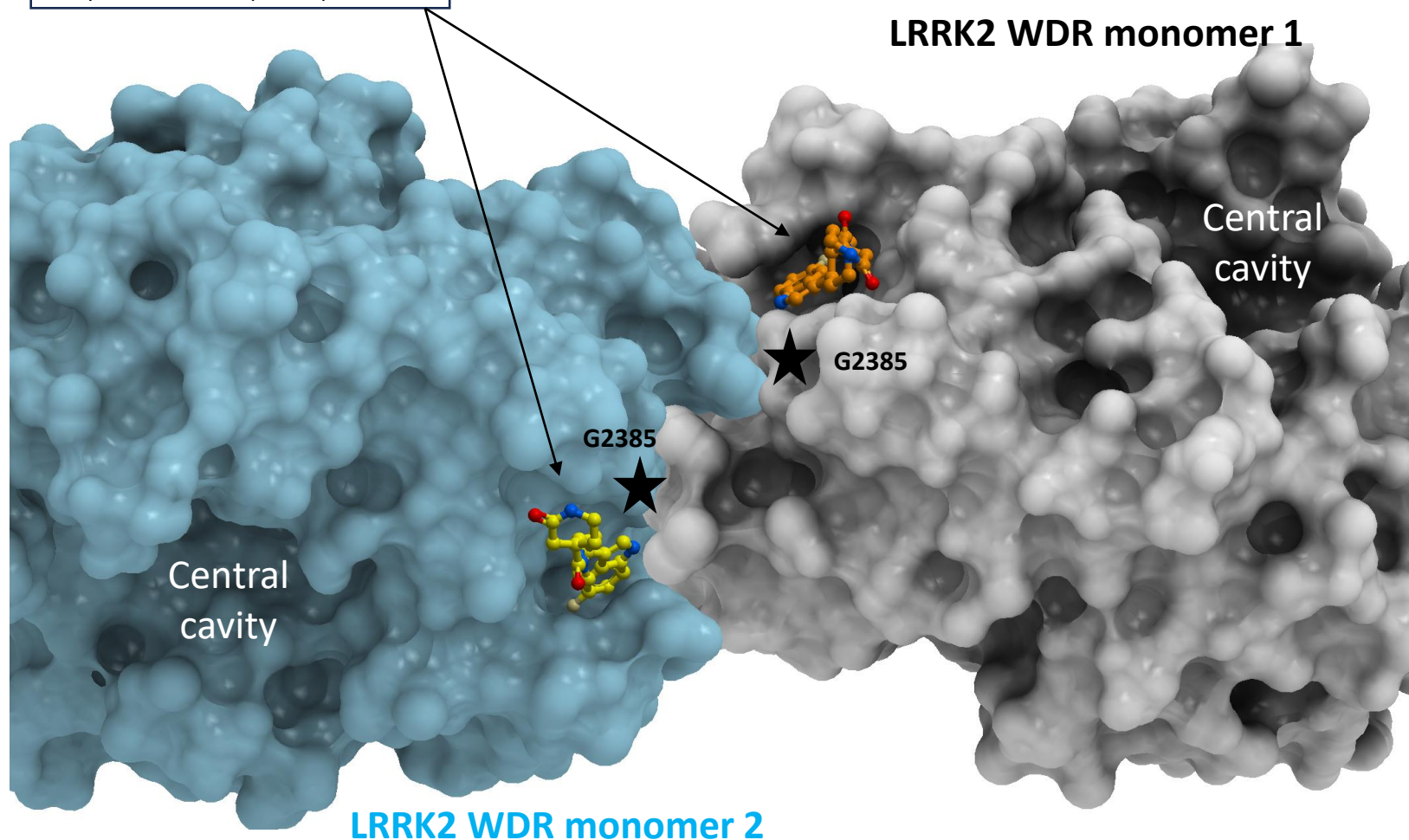


CACHE\_1193\_26  
SPR LRRK2 48 $\mu$ M – 58% binding  
SPR NSD2 – NA – 36% binding  
100% solub /no agg. at 200  $\mu$ M  
Co-crystallized at unexpected pocket



- Pathogenic, dimerization-defective LRRK2 mutants such as G2385R have enhanced kinase activity
- CACHE\_1193\_26 occupies a side pocket next to G2385
- CACHE\_1193\_26 still binds in solution to the G2385R mutant, suggesting a different binding mode in solution.
- Nevertheless, the structure reveals a potential mechanism to stabilize the dimeric form of LRRK2 WDR

Structure solved by Aiping Dong, Hong Zeng and Levon Halebelian, Structural Genomics Consortium