Lead development of orally available drug for diabetic retinopathy targeting transglutaminase 2 to inhibit vascular leakage



| OPHTHALMOLOGY | Hit |
|--------------------------|--|
| Product Type | Small Molecule |
| Indication | Diabetic Retinopathy |
| Target | Transglutaminase 2 (TGase 2) |
| MoA(Mechanism of Action) | Inhibition of VEGFR induced vascular leakage by TGase 2 inhibition |
| Competitiveness | Better patient compliance by oral administration, better efficacy in STZ-DR mice model (ivt) than Eylea and imatinib, better vision improvement and TI, wider market positioning throughout NPDR and PDR, applicable to other diabetic complications |
| Development Stage | Hit |
| Route of Administration | oral |

