



Dedication to the IVD industry?

**We I.V.DO that™**

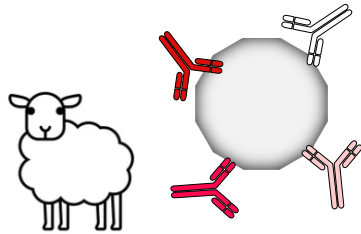


# Antibody engineering for IVD applications

# History of the antibodies

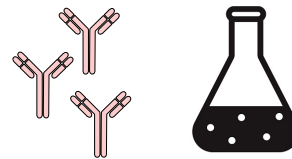
**1800's**  
Nobel Prize 1901

**Polyclonal antibodies**



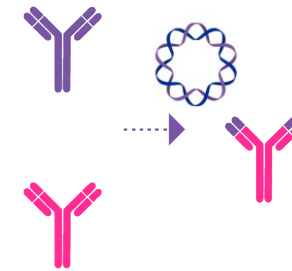
**1970's: Mouse hybridomas**  
Nobel Prize 1984

**Monoclonal antibodies**

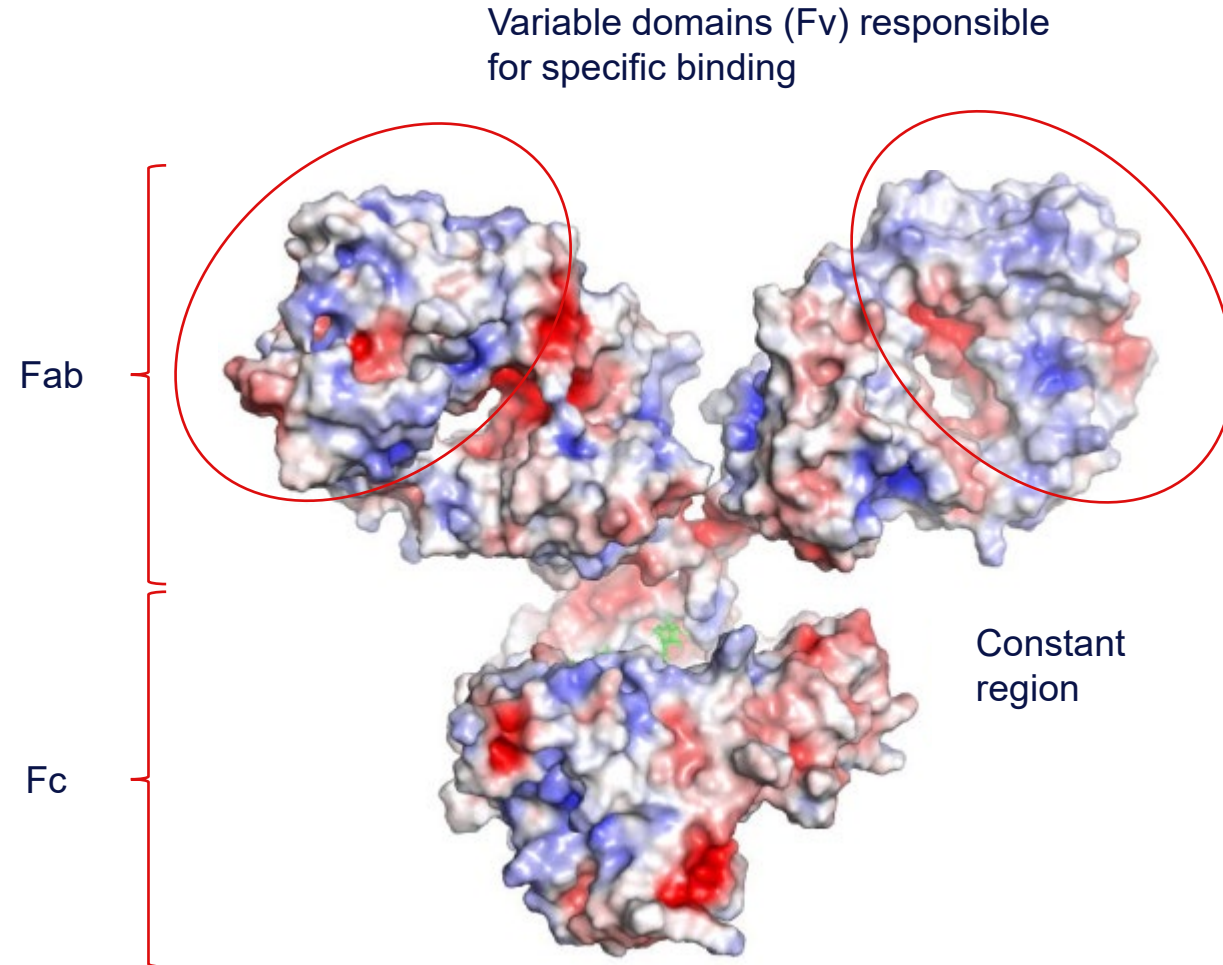


**Now: Recombinant technologies**  
Nobel Prize 2018

**Engineered antibodies**

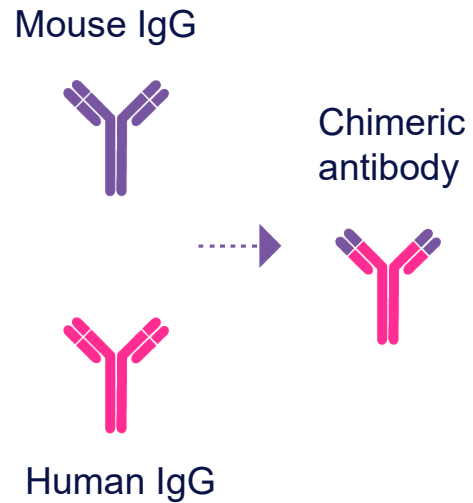


# Antibody engineering means modification of antibody sequence to enhance or reduce wanted properties

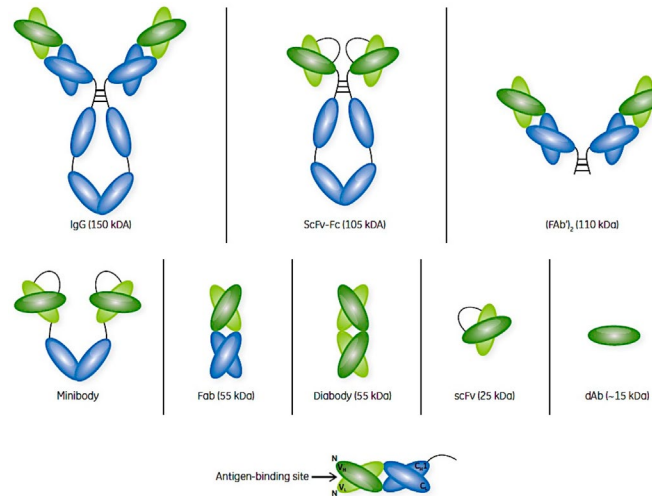


- ✓ Enhancing specificity or affinity
- ✓ Improving stability or reducing aggregation
- ✓ Directed evolution through structure analysis and rational design

# Antibody engineering for IVD applications

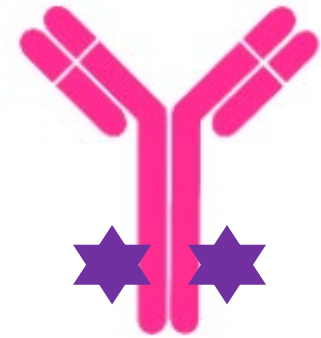


- ✓ Chimeric antibodies to reduce HAMA effect and others interferences in IVD assays



<https://doi.org/10.3390/antib4030259>

- ✓ Smaller size enable high-density binding and improved accessibility

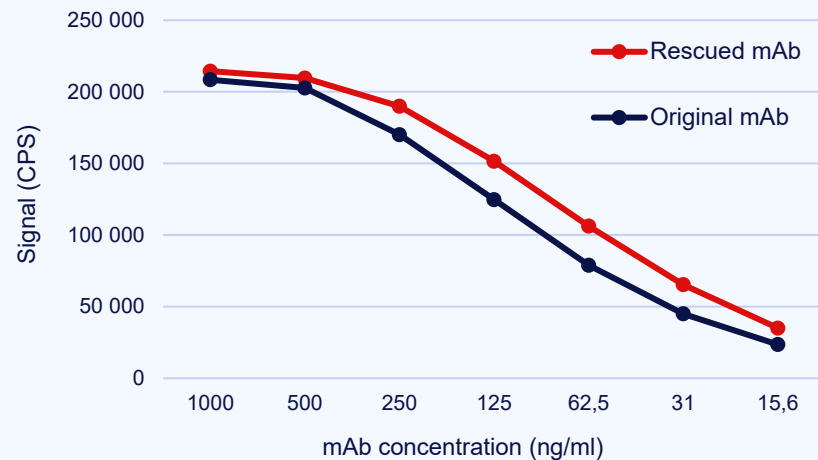


- ✓ Add-on functionalities for purification and site directed conjugation

# Case Example 1 - Antibody rescue for non-viable hybridomas

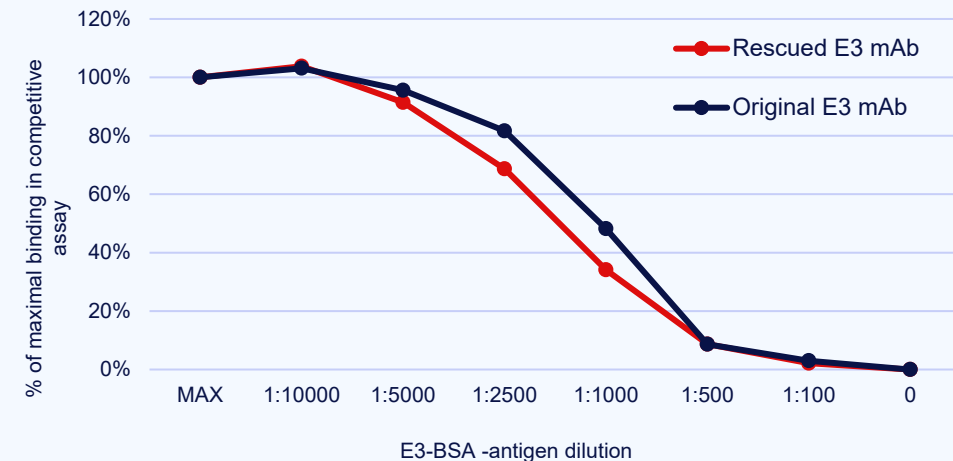
Drugs of Abuse: **LSD antibody** rescued and expressed in mouse IgG<sub>1</sub> framework

- ✓ Secured the supply for a high-performing mAb with minimal changes

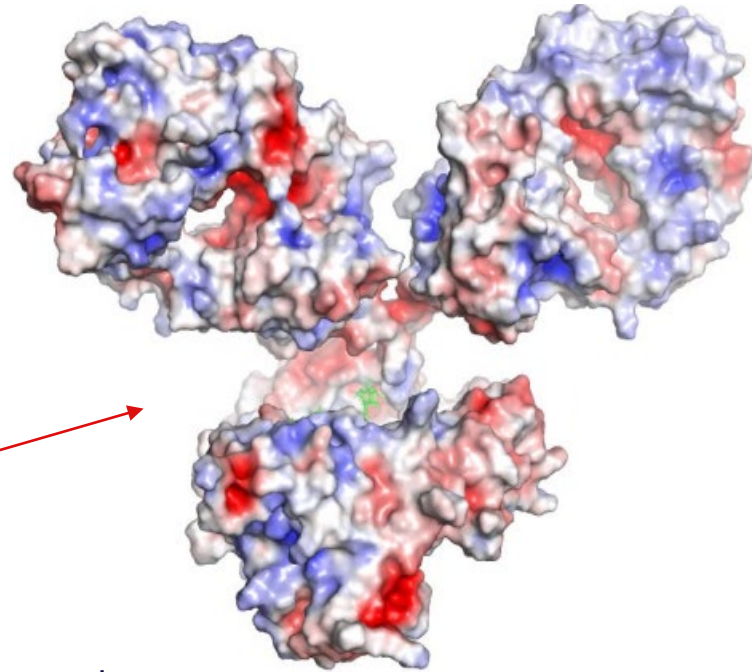


Womens' Health: **E3 antibody** rescued and re-engineered into human IgG<sub>1</sub> framework

- ✓ Secured the supply, and provided a new mAb version with reduced HAMA effect risk



# Case Example 2 - Targeted antibody engineering for aggregation prone mAb



## Framework switch

- Recombinant variants with varying constant regions
- Reduced aggregation tendency observed

## Point mutations in aggregation-prone regions

- Computational modelling for several mAbs with varying solubility properties
- Aggregation-prone regions identified for the aggregating mAb
- Suggested point mutations currently studied



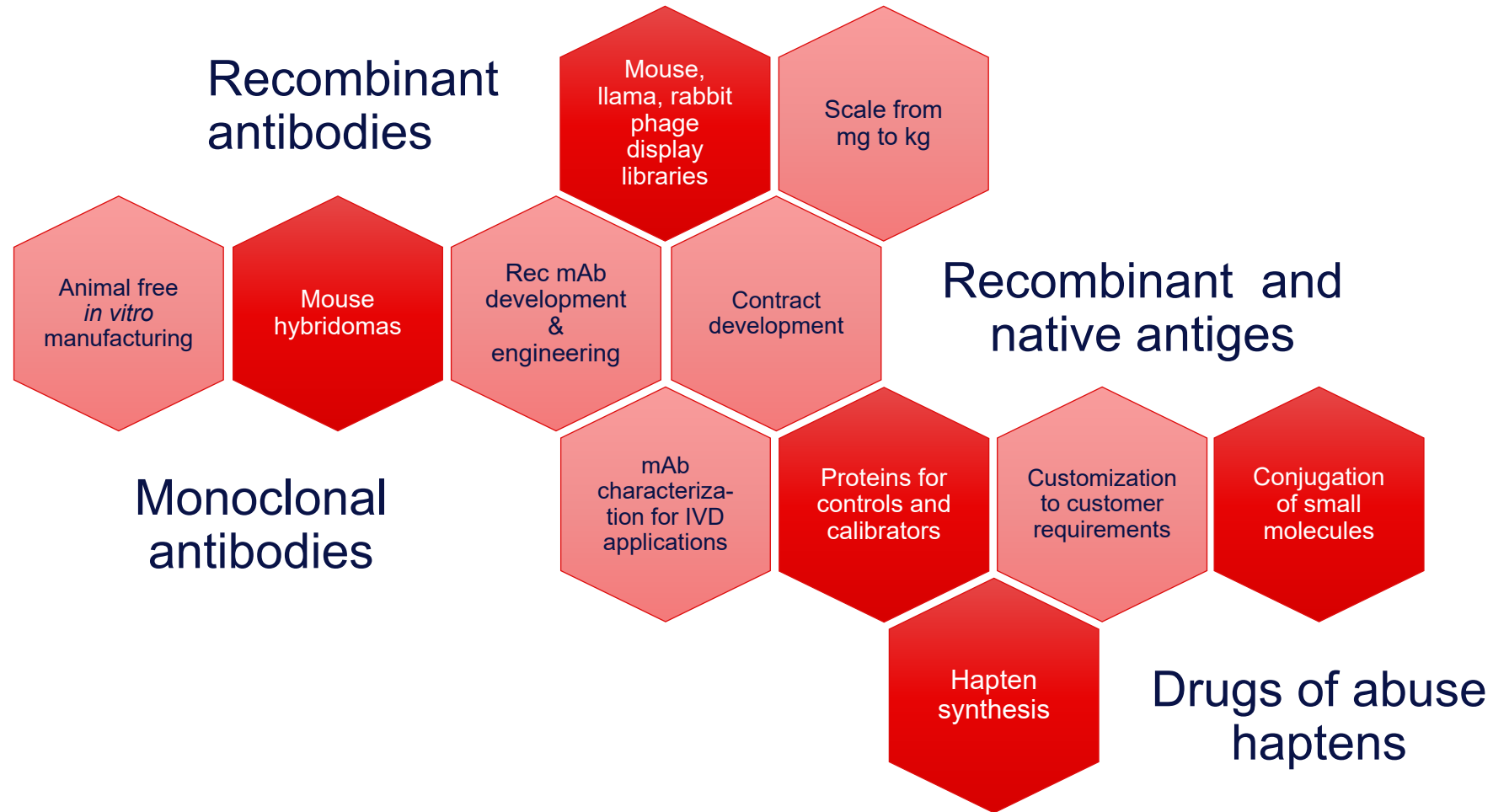
Historically, antibody discovery limited by the immune response repertoire of animals – mostly mice.

**In the future, novel mAbs can be discovered computationally and any mAb can be optimized.**

How can the IVD industry leverage this technology disruption?



# Our R&D capabilities



R&D innovations that  
keep pace with yours?

**We I.V.DO that™**



Thank you for **y**our time