

Designing a novel biosensor for the detection of infectious pancreatic necrosis virus



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Infectious Pancreatic Necrosis Virus

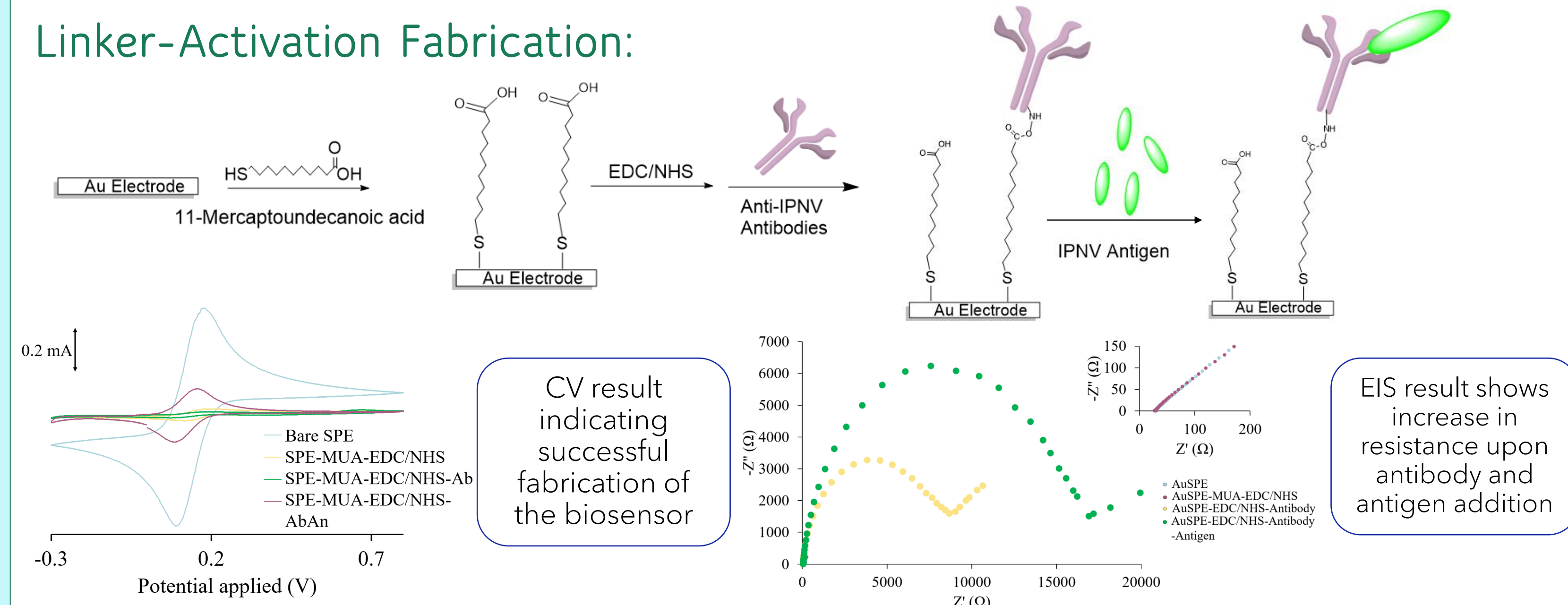
- Infectious Pancreatic Necrosis Virus (IPNV, *Birnaviridae Aquabirnavirus*) causes a life-long, highly contagious disease in Salmonid Fish^{1,2}
- IPNV infection causes necrosis of pancreatic and other tissues
- Effects salmonid fish at all ages but most devastating for young, 70% mortality rate³
- Large implications for the Salmonid industry
- Virus unable to infect human cells in-vitro³
- However, infected salmon consumption correlated with gastrointestinal infections³
- Careful detection needed to alleviate health and economic concerns as no treatment exists⁴



- CV result indicating successful fabrication of the biosensor
- EIS result shows increase in resistance upon antibody and antigen addition

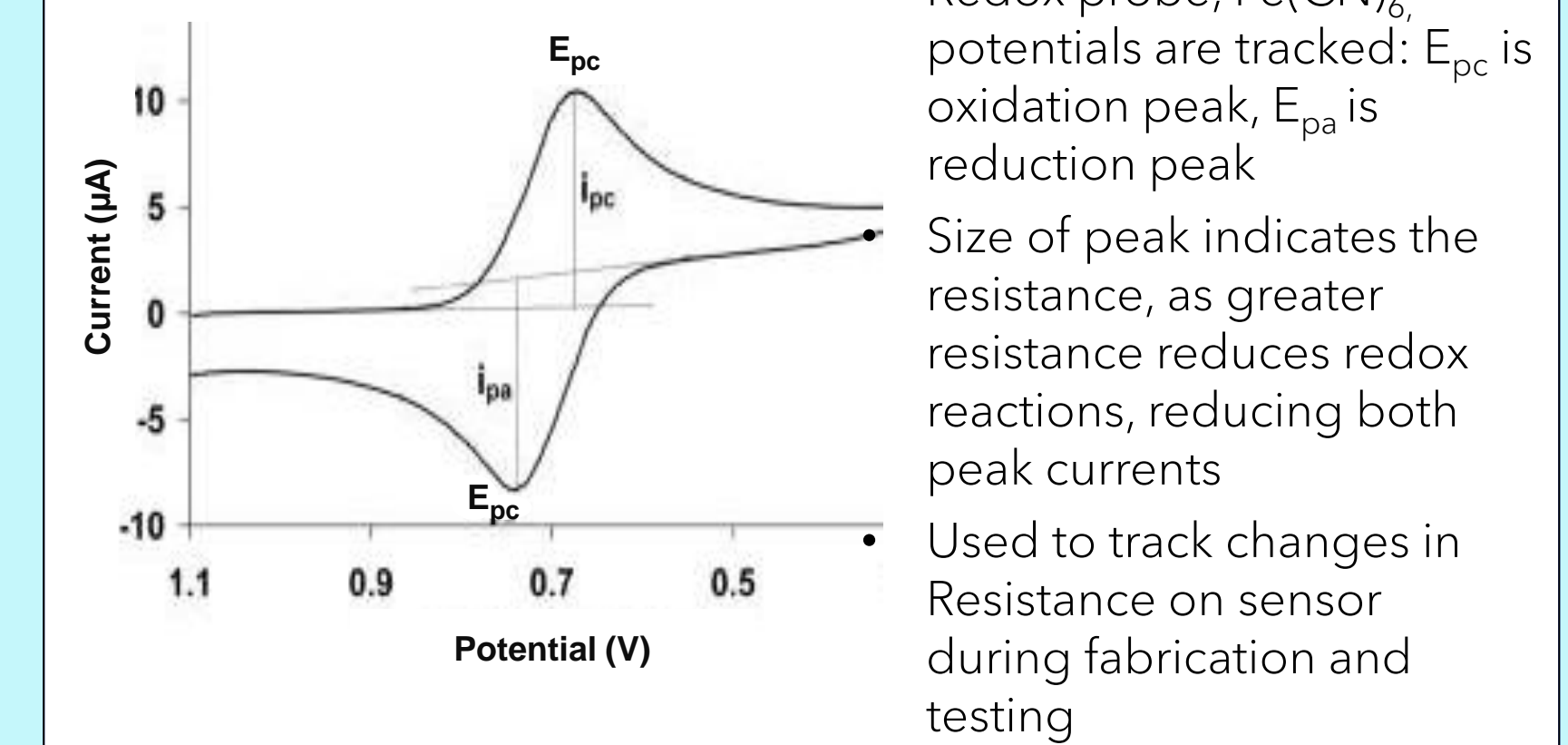
Methods & Results

Linker-Activation Fabrication:

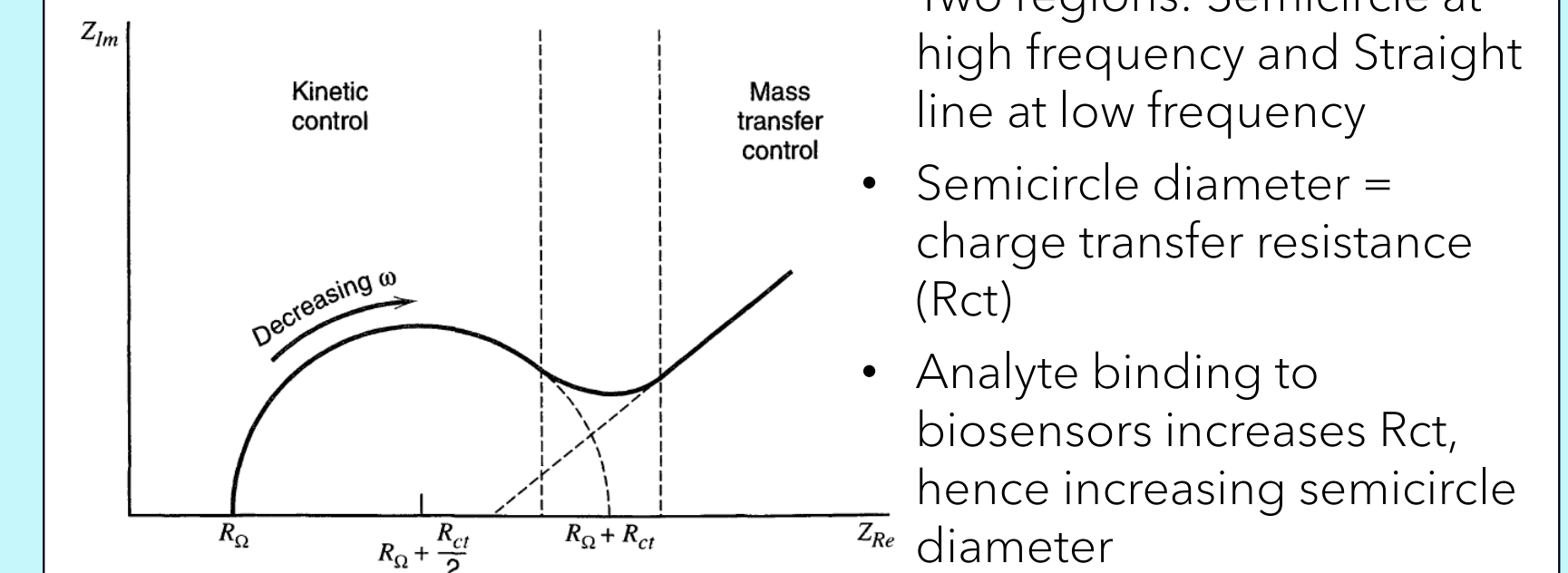


Decoding the Graphs

Cyclic Voltammetry (CV) plots:

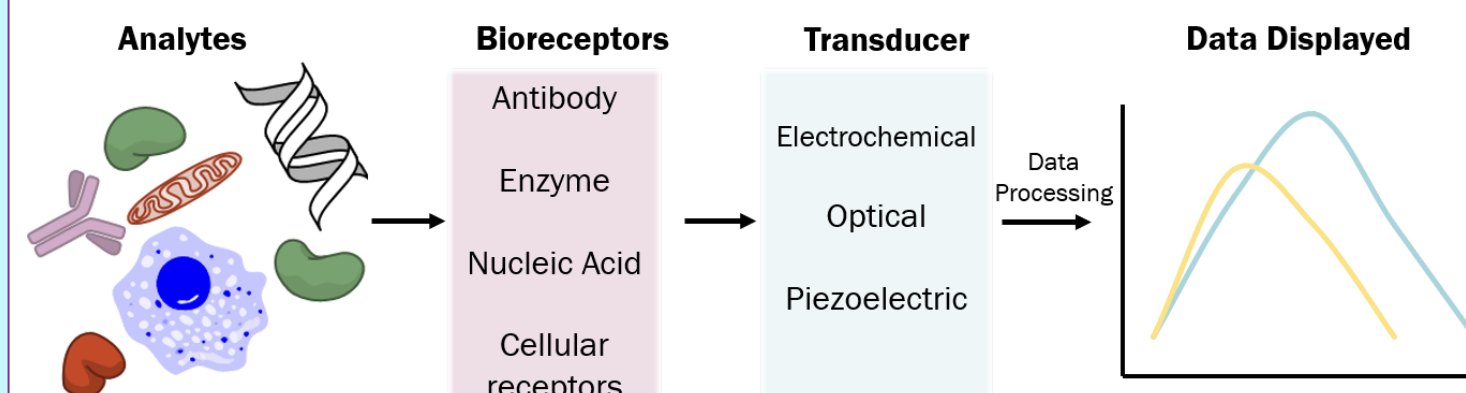


EIS Nyquist plots:



Biosensors

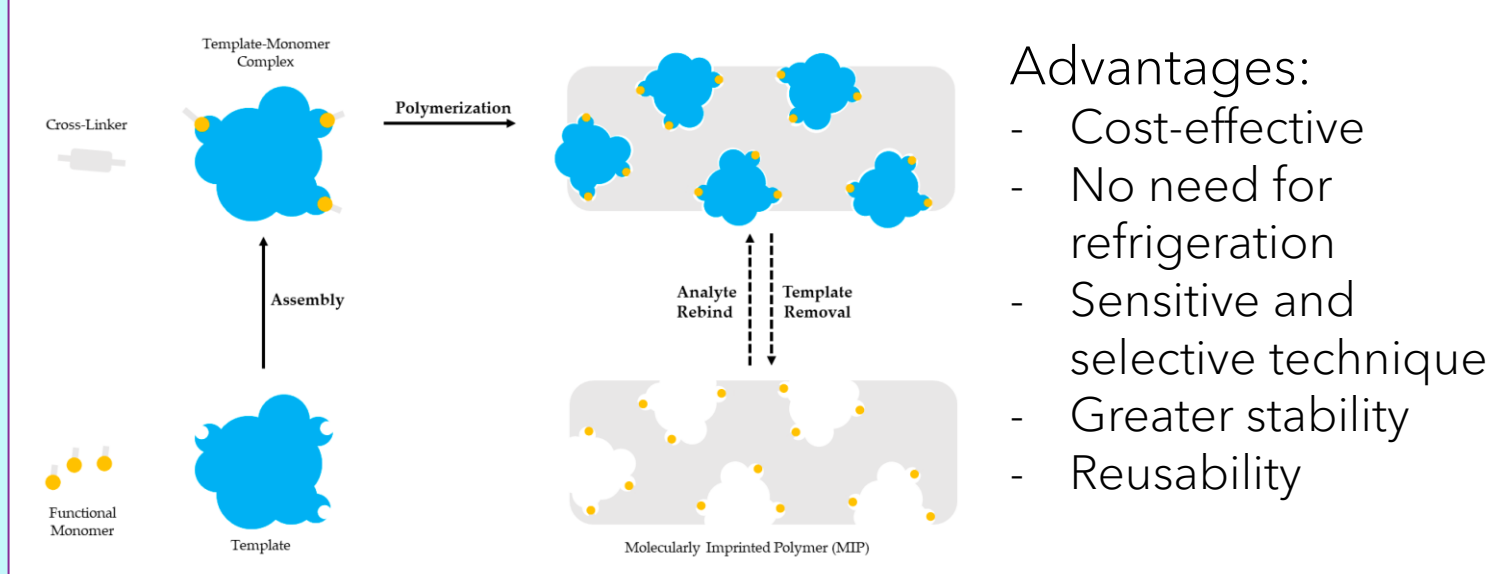
Used as diagnostic tools to detect analytes in a matrix
Composed of following:



Biosensors offer multiple advantages for IPNV detection⁴:

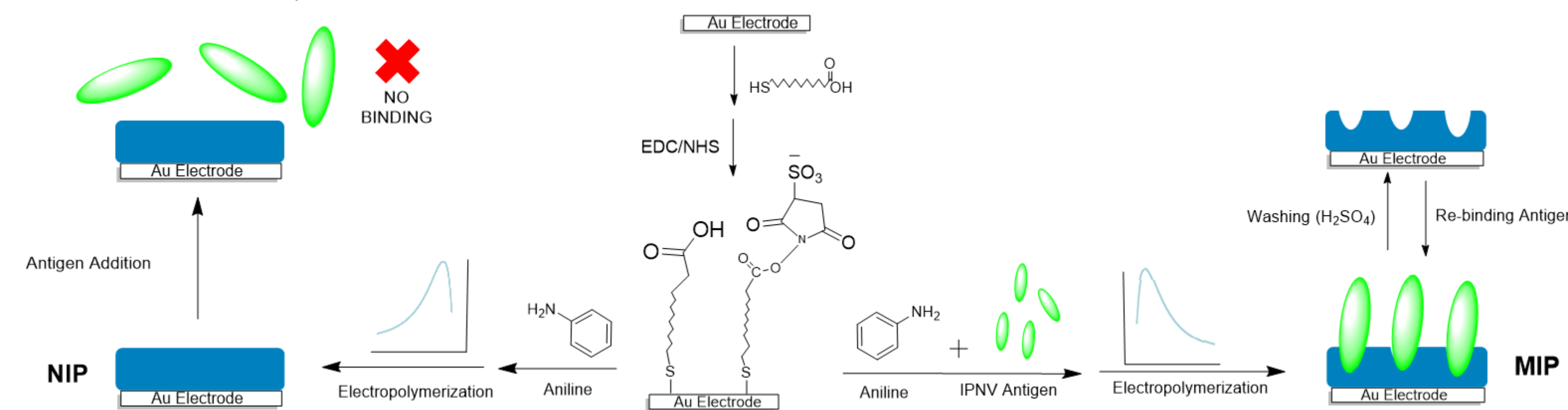
- Aquaculture farmers can screen for IPNV themselves
- Cost-effective
- Modification of surface to improve bioreceptor immobilization, detection, and sensitivity of detection

Molecular Imprinting polymer (MIP) modification, creates an artificial bioreceptor from a template⁵



- Advantages:
- Cost-effective
 - No need for refrigeration
 - Sensitive and selective technique
 - Greater stability
 - Reusability

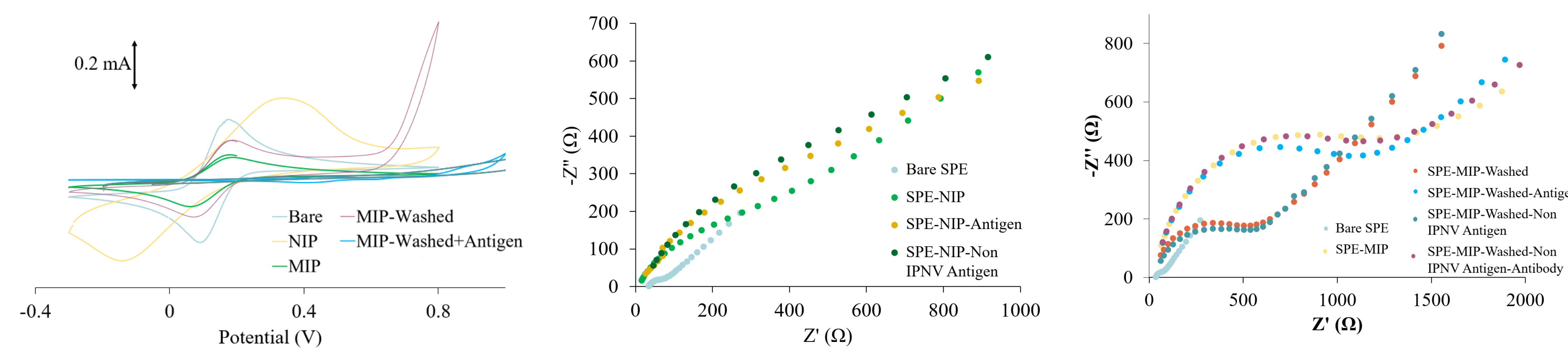
Molecular Imprinting Polymer Fabrication:



CV used to track MIP formation, showed successful formation

Non-imprinting polymer (NIP) showed no sites for antigen binding as expected

MIP showed specific sites where only Antigen bound, while non-antigen proteins did not



Conclusion and Next Steps

IPNV Antigen was successfully detected using a variety of biosensor fabrications techniques

- MIP Fabrication showed promising results
 - Continue with fabrication to generate a calibration curve (determine detection limits)

Experiment successful on pure antigens, future experimentation on real samples (e.g. Salmon meat)



References:

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