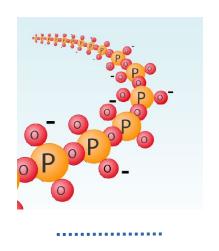


Providing superior treatments for all forms of bleeding



Company Vision: Leading Biotech Focused on the Treatment of Bleeding Disorders



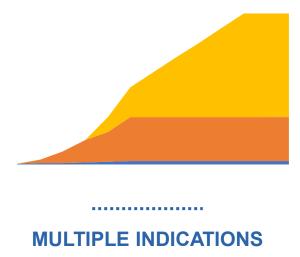
FIRST IN CLASS

SYSTEMIC

CLOTTING ACCELERANT



QUICK PATH TO MARKET





CAY001 Has Potential as a Universal Agent to Treat Multiple Causes of Uncontrolled Bleeding

Congenital Clotting Factor Dysfunction

Hemophilia

Saturated Market

Drug Clotting

Factor Dysfunction

- Coumadin
- FXa
- Direct Thrombin inhibitors
- 5 drugs approved
- Limited effectiveness
- Safety issues

Platelet Dysfunction

(5 MM)

- Antiplatelet drugs (Plavix)
- Congential platelet dysfunction
- Thrombocytopenia

Platelet transfusion

- 44 % Ineffective (Plavix)
- Antigenic (Disorders)
- Limited supply

CAY001 has potential to work across all types of bleeding dysfunction

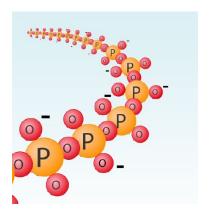
INITIAL FOCUS ON PLATELET DISORDERS

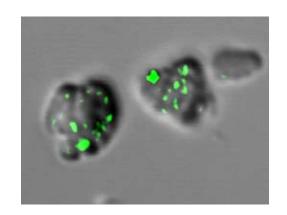


Polyphosphate (PolyP) is a Critical Driver of the Clotting Response. CAY001 is an IV formulation of PolyP

What is PolyP?

- Produced by platelets
- Accelerates clotting
- Unsuccessful hemophilia drug

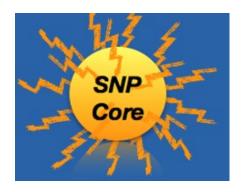




What is CAY001?

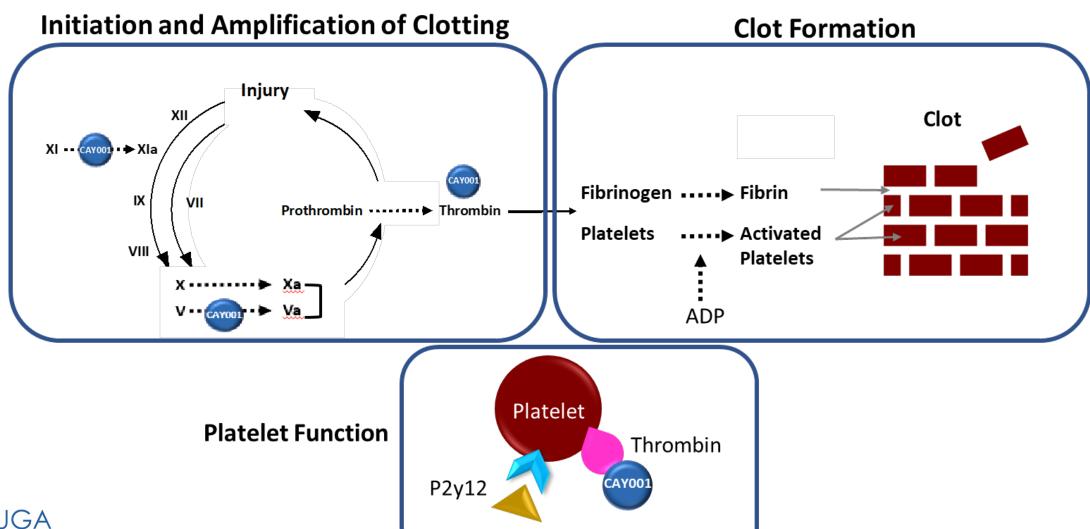
Silica-based nanoparticle (SNP) that improves delivery of PolyP

- Increase half-life
- Lower dose
- IP protection through 2034 (issued patents)





PolyP Accelerates But Does Not Increase Clotting and Can Overcome Platelet Dysfunction

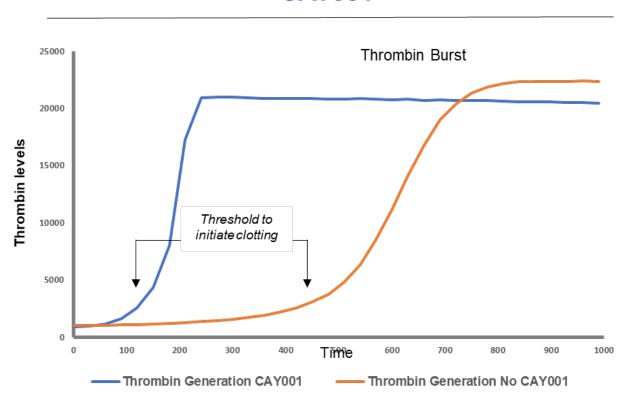


ADP



CAY001 Accelerates Thrombin Generation 4X. No Increase in the Amount of Thrombin Produced

CAY001



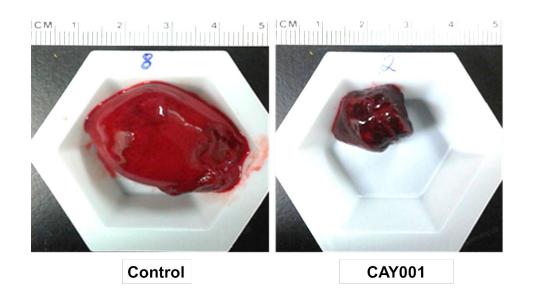
Angew.Chem. Int. Ed. 2015, 54,4018 -4022

Faster and better clotting at injury sites without increased clotting risk

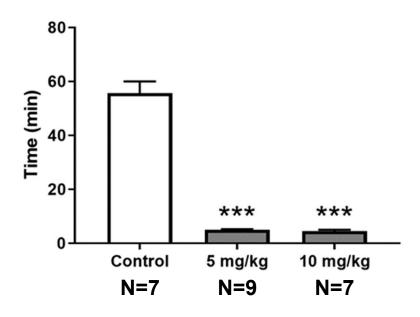


CAY001 in Rat Model: Improved Clot Formation, Reduced Bleeding Time and Blood Loss

Clots formed with CAY001 had greater density and localized at the injury site in Rat



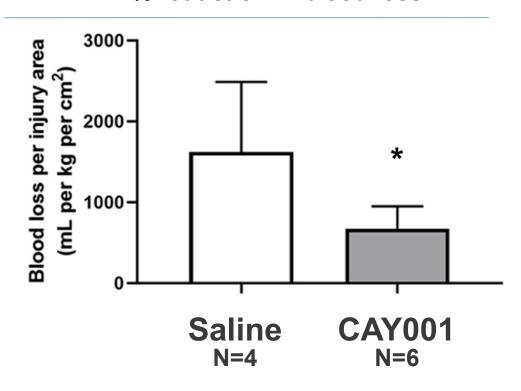
Bleeding time reduced by 90% 31% reduction in blood loss in Rat



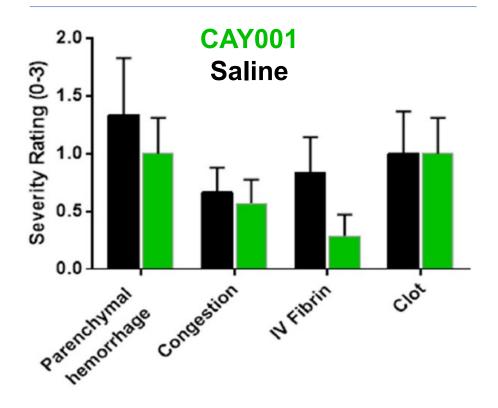


CAY001 in Pig Model: Reduced Blood Loss Without Increased Clotting

42 % reduction in blood loss



No increase in adverse events





Plavix Reversal is a Large Unmet Need







0.150 MM patients

0.250 MM patients

1.12 MM patients (\$5 B)



Plavix Reversal Provides a Quick Path to Market

Commercial Fast to Market Strategy Leveraging FDA's Accelerated Approval Pathway



The FDA instituted its Accelerated Approval Program to allow for earlier approval of drugs that treat serious conditions, and that fill an unmet medical need based on a surrogate endpoint. A surrogate endpoint is a marker, such as a laboratory measurement,



Commercial Fast to Market Strategy Leveraging FDA's Accelerated Approval Pathway



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Precedent in bleeding indication: Andexxa

- Reversal of FXa anticoagulants
- Approval based on 71 healthy volunteers
- Black box warning
- Portola (PTLA): Acquired by Alexion for \$1.4 B (May 2020)

Implications for CAY001

- ✓ Treats life-threatening or uncontrolled bleeding
- Development path focused on high unmet need populations (No treatment for anti-platelet reversal)
- ✓ Potential surrogate endpoints identified



Clinical Proof of Concept Possible with \$10M Raise Initial Raise of \$4M To Get to First in Human

Pre-IND meeting
May 2020

IND-enabling work
Now - Dec 2021

IND Dec 2021 First in human studies 2022 -





Investment to Secure IND \$4 MM

Clinical Stage \$6 MM



The Cayuga Team



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Nancy Stagliano, PhD CEO Neuron23, Former CEO True North Therapeutics

