



Phase II - USAISR Platelet Results

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Key Highlights

USAISR results confirm Vitrafy cryopreservation ecosystem outperforms regulatory and quality standards across all platelet protocols tested.

94%

Post thaw recovery - no-wash protocol, with high functionality across all quality measures

Independent
US Army
Validation

Largest blood platelet testing program in commercial volumes, using the Vitrafy cryopreservation ecosystem

Category First
Opportunity

Position the Company to address this critical market gap

Phase II In-Vitro Study Background & Aims



Study Aim

Assess and determine Vitrafy's cryopreservation ecosystem **capability** to successfully cryopreserve critical response **blood platelets**.



CRADA Framework

Conducted under the Cooperative Research and Development Agreement (**CRADA**) between Vitrafy and the U.S. Army.



Independent Validation

Conduct independent, **U.S. Army-authored validation** by the U.S. Army Institute of Surgical Research ahead of FDA Medical Device registration.



Rapid Deployment Capability

Establish a practical operational solution for a rapidly deployable cryopreserved platelet product (**decentralised no wash** protocol).



Largest Testing Program to Date

Apheresis platelets from **20 donors / 60+ samples** tested via the Vitrafy ecosystem using 3 protocols.



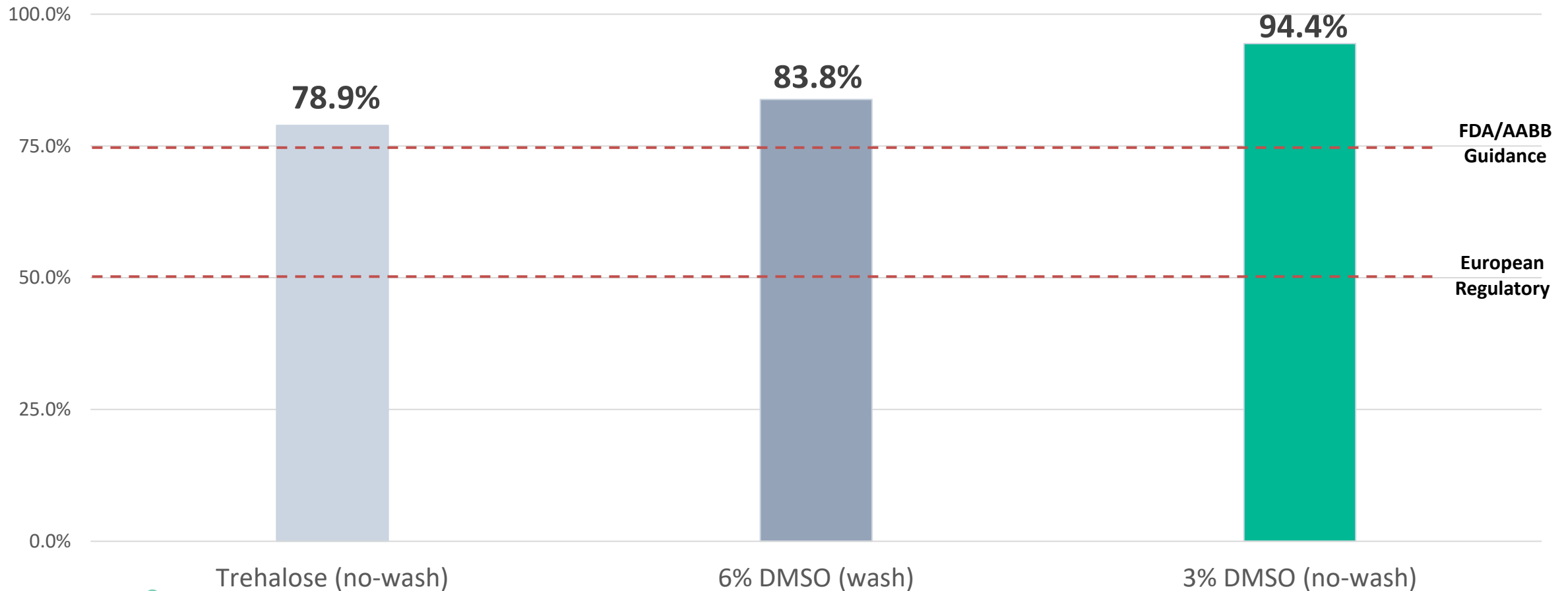
IPO Commitment Delivered

USAISR Phase II validation was a **core deliverable** identified at Vitrafy's November 2024 IPO — **now completed**.

Results - Platelet Recovery vs Fresh

All protocols tested using Vitrafy's cryopreservation ecosystem exceeded the regulatory standard and quality guidelines. 3% DMSO no-wash protocol delivered the highest performance with a 94% post-thaw recovery

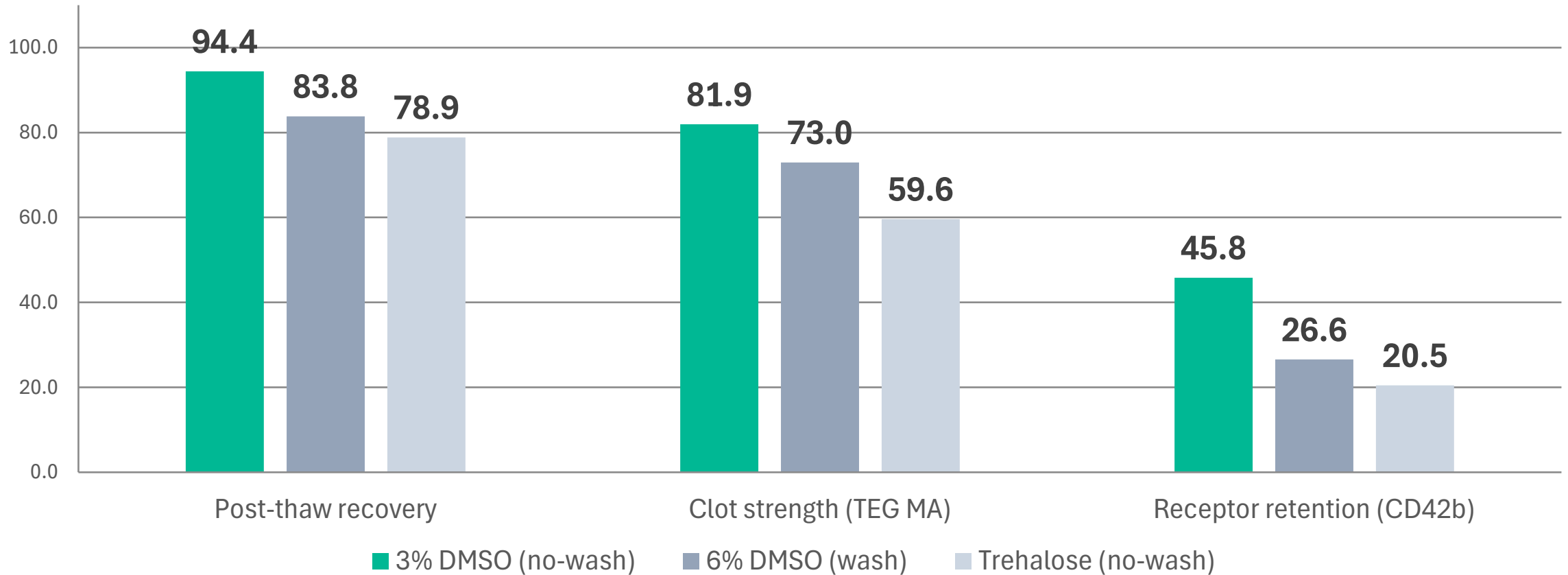
Phase II Post-Thaw Platelet Recovery - % of Pre-Freeze (Fresh) Baseline



Results - Platelet Functionality vs Fresh

Key platelet functionality metrics were measured post thaw – including clot strength and the retention of key surface receptors.

Functionality Markers — % of Pre-Freeze (Fresh) Baseline



Outlook

In Vitro validation milestone complete— multiple pathways now activated.

FDA medical device registration (GUARDION) – H1 FY27

Phase II Results Publication (military & civilian) – H1 FY27

FDA, Military & Civilian blood network engagement