

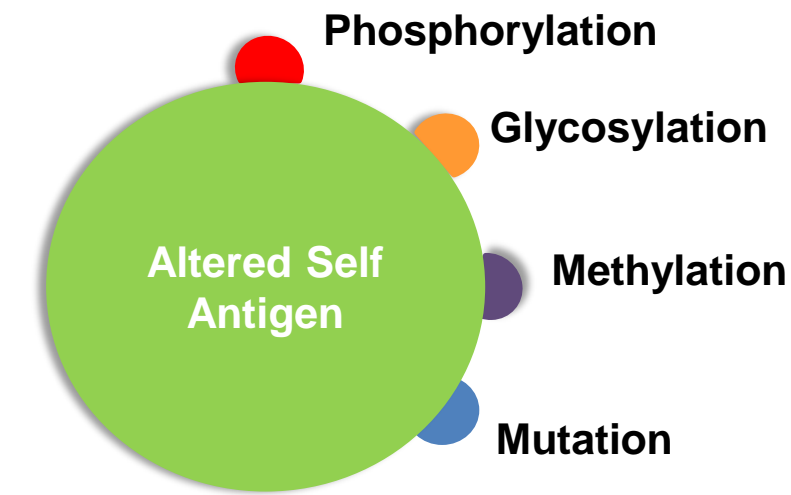
AGENUS' NEXT GENERATION CANCER VACCINE PLATFORMS

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TUMOR SPECIFIC NEO-ANTIGENS

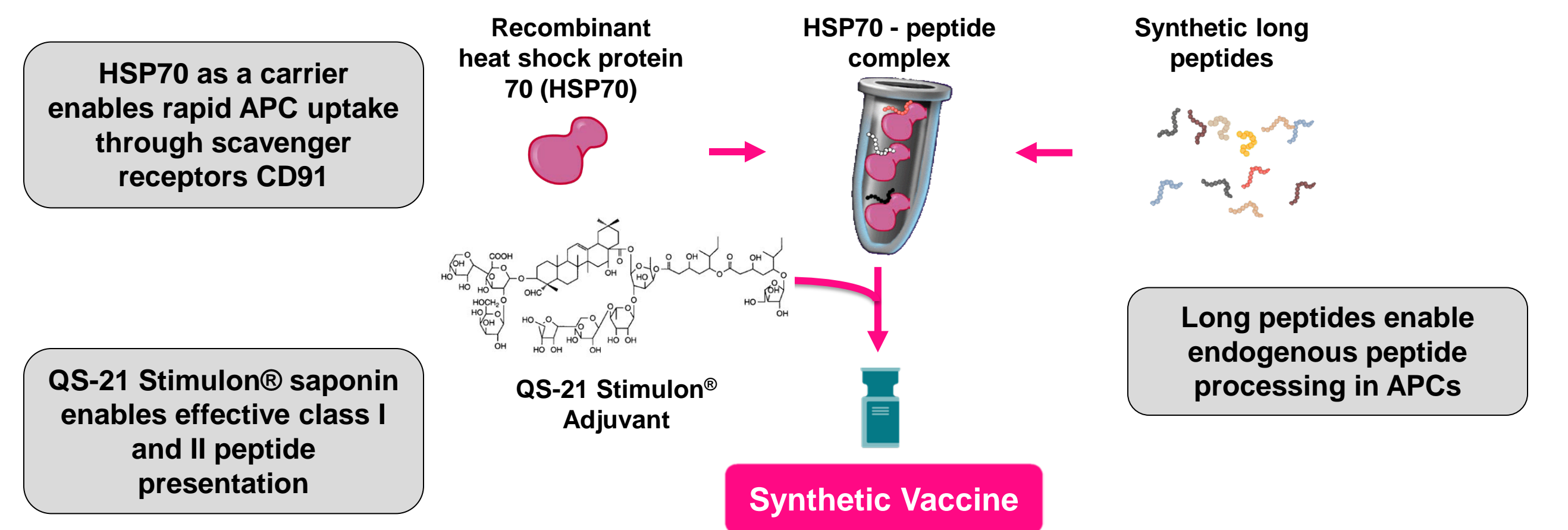
We design our vaccines to target neoantigens that are specifically presented on tumor cells. These include individualized and shared immunogenic DNA mutations and peptides with post-translational modifications (PTMs) specifically presented on tumor cells.



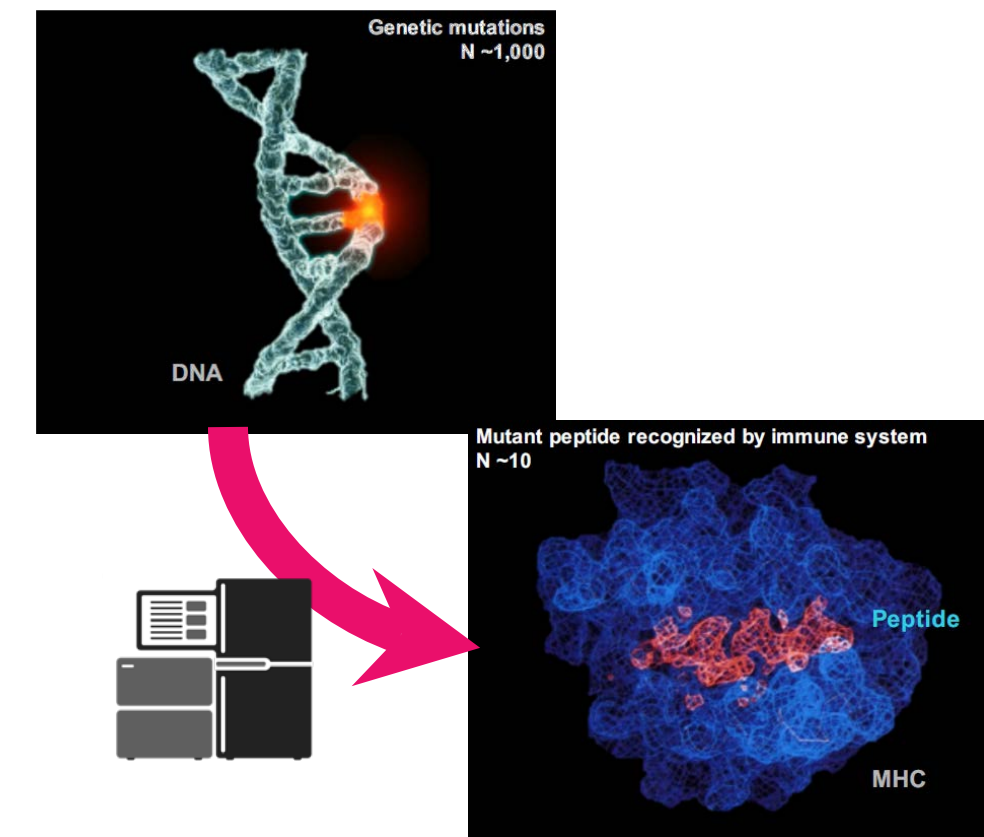
VACCINE STRATEGY



VACCINE FORMAT



AIM™ IDENTIFIES IMMUNOGENIC MUTATIONS

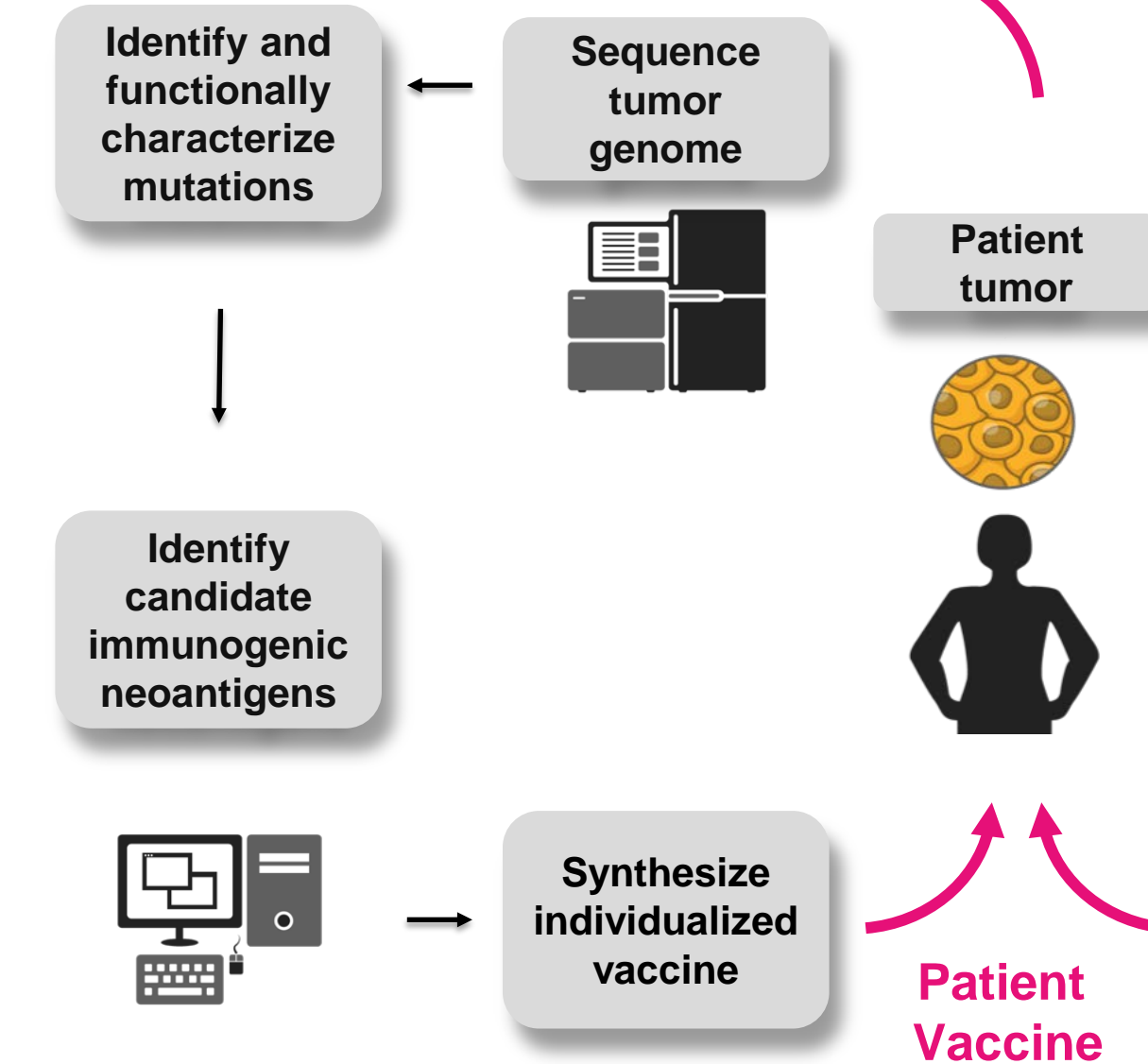


The Agenus Immunogenic Mutation (AIM™) workflow comprises rapid multi-sample NGS profiling followed by advanced bioinformatics and computational immunology algorithms for mutation characterization and selection.

The AIM™ algorithm incorporates best-in-class mutation identification and HLA binding algorithms. Further, AIM™ integrates proprietary findings using MS ligandome data, immunological response in humans and mice, and TCR functionality.

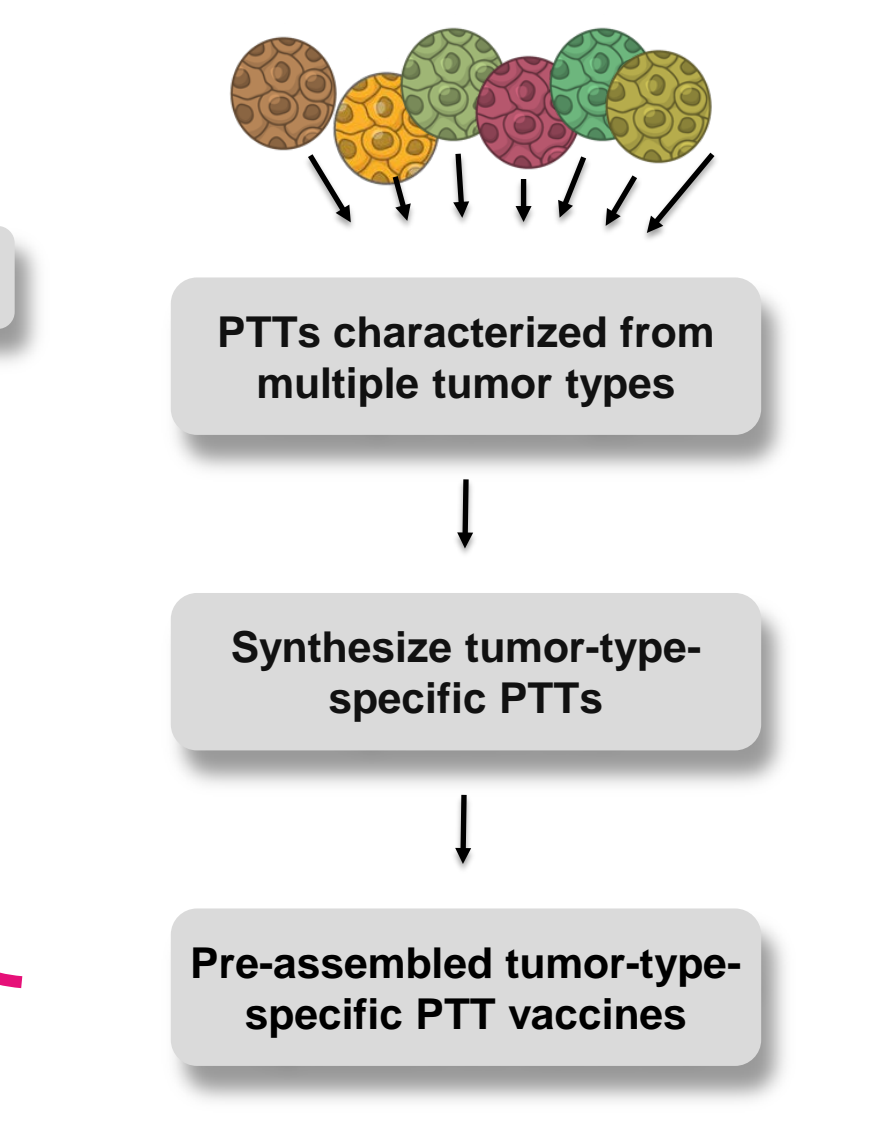
AutoSynVax™

On-demand manufacture



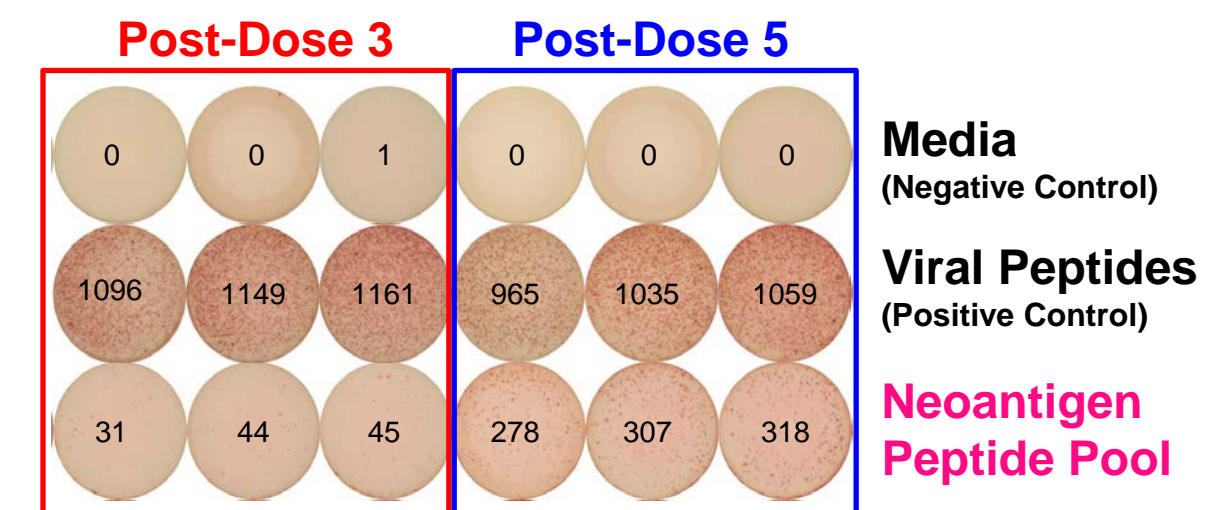
PhosphoSynVax™

Ready-to-ship

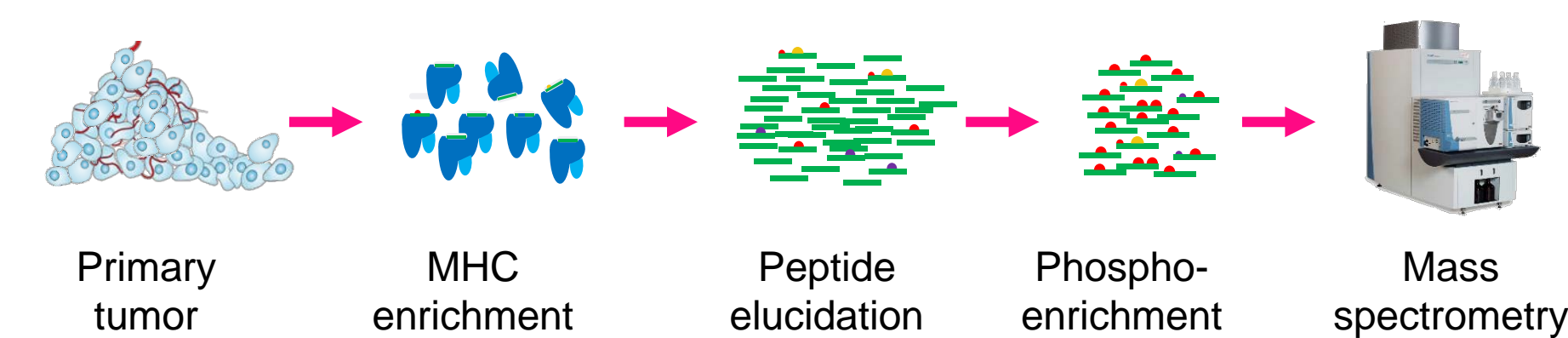


CLINICAL IMMUNE RESPONSE

Under compassionate use, we treated a metastatic hepatocellular carcinoma (HCC) patient. AIM™ identified 24 non-synonymous mutations and 13 prioritized for vaccine inclusion. AutoSynVax™ long peptides were manufactured. The patient was dosed every two weeks. The T-cell response was measured by ELISpot using pooled short mutation-containing peptides



SHARED PHOSPHO-NEOANTIGENS

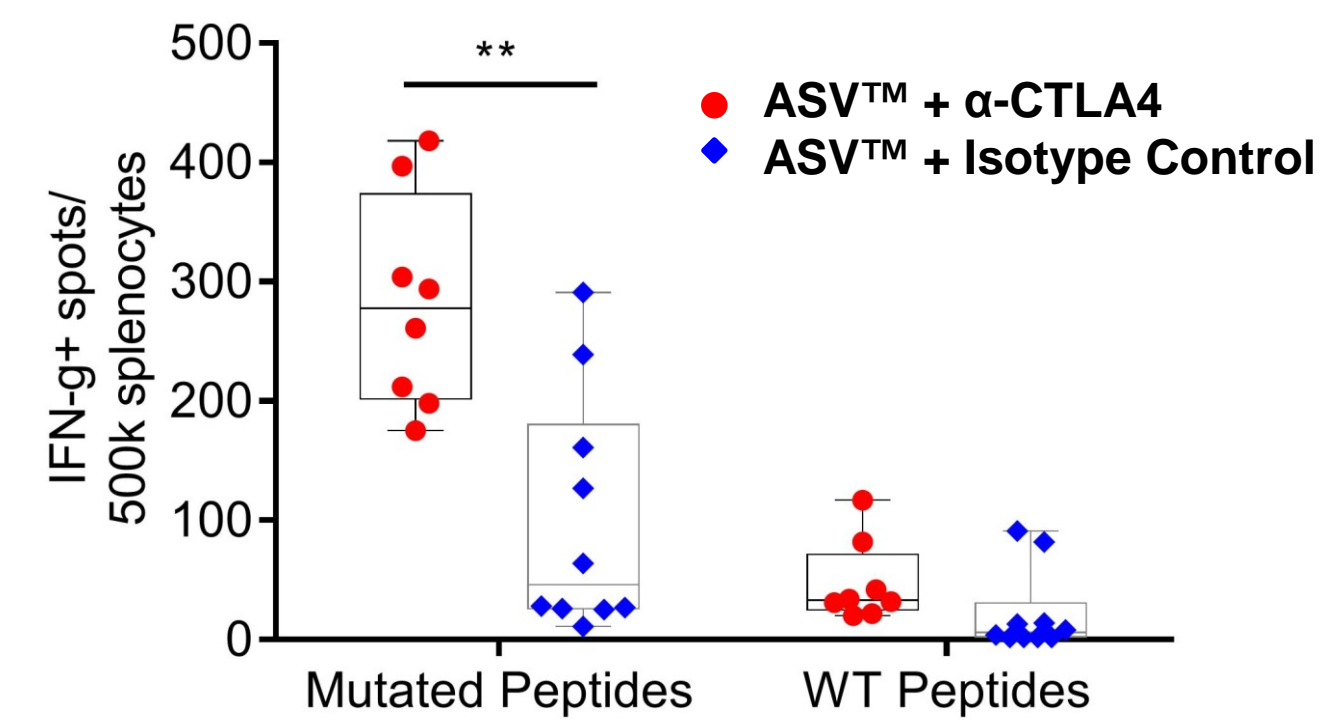


We identified 100's of HLA ligands that are shared, specifically presented on tumor cells, and recognized by central memory T cells from healthy donors.

PHASE 1 AutoSynVax™ TRIAL

- Multiple sites
- Enrollment completed
- First patient successfully dosed
- Safety and immunological readout expected 2017
- Planned combination with immunomodulatory agents

SYNERGIZES WITH mAbs



PRECLINICAL IMMUNOLOGICAL & TUMOR REPOSE

Agenus vaccines mediates tumor control (left), immune memory (middle) and phospho-peptide recognition (right)

