

# Ainnocence Achieves New Milestone with Protein Foundation Models

Proprietary sequence-first model rivals and in key areas surpasses the leading ProteinGym sequence-only baselines

SAN FRANCISCO, CA, UNITED STATES, August 23, 2025 /EINPresswire.com/ -- Ainnocence, a pioneer in generative Al for biotechnology, today announced the successful internal benchmarking of its protein language model, AINN-P1, trained on UniRef sequences with up to 167 million parameters.

#### Comparative performance on the ProteinGym zero-shot leaderboard

Model (Input Modality)	Avg. ρ	Activity	Binding	Expression	Stability
AINN-P1 (Single seq.)	0.441	0.358	0.390	0.391	0.625
Progen2 M (Single seq.)	0.380	0.393	0.295	0.433	0.396
ESM2 (150 M) (Single seq.)	0.387	0.391	0.326	0.402	0.510
xTrimoPGLM-1B-MLM (Single seq.)	0.379	0.390	0.298	0.406	0.445
Wavenet (MSA)	0.373	0.379	0.325	0.350	0.449
xTrimoPGLM-100B-int4 (Single seq.)	0.364	0.378	0.259	0.380	0.450

The baseline numbers are from the July 2025 ProteinGym zero-shot leaderboard; Ainnocence values are few-shot results reported below.

AINN-P1 is Ainnocence's protein language foundation model, built to understand protein sequence patterns to power next-generation drug discovery. The new model posts [] state-of-the-



Sequence is the new paradigm"

Dr. Lurong Pan

art Spearman correlation scores across four core proteinfitness tasks, reinforcing Ainnocence's sequence-first drugdiscovery platform and underscoring its competitiveness against much larger transformer models from Meta and BioMap.

# Highlights:

- Top overall sequence-only score: With an average  $\rho$  = 0.441, AINN-P1 edges past ProSST (0.438) while using  $\square$  orders-of-magnitude fewer parameters  $\square$  than 100 B-scale transformers.
- Best-in-class stability prediction: A  $\rho$  = 0.625 on stability is the highest of any published sequence-first model, providing critical accuracy for manufacturability screens.
- Closing the binding gap: AINN-P1's binding  $\rho$  = 0.390 exceeds ESM2-150M by 0.064 points, narrowing the historic advantage of structure-aware models.

#### Detailed benchmark results

Recent in-house evaluations of the AINN-P1 performed in a□few-shot embedding □regime for efficiency, produced the following Spearman correlations:□Activity 0.3581, Binding 0.3901, Expression 0.3913, Stability 0.6251.□□These metrics match or exceed many transformer-based

benchmarks, demonstrating that smart architecture and curated data can beat brute-force scale.

Crucially, these benchmarks were achieved without expensive full fine-tuning. Instead, fixed AINN-P1 embeddings feed lightweight regressors – a strategy that slashes compute costs while preserving accuracy, and aligns with emerging best practices when only limited experimental data are available.

## Context within the protein-Al landscape

Meta's ESM family and BioMap's 100 B-parameter xTrimo model have advanced the field dramatically, yet their training budgets remain prohibitive for most labs. Ainnocence's leaner AINN-P1 delivers comparable accuracy without billion-scale parameters or MSAs, validating a sequence-only paradigm for broad adoption.

"Sequence is the new paradigm," said <u>Dr. Lurong Pan, CEO</u>. "By learning the language of proteins, we predict complex properties without relying on structures or wet-lab screens. These results rival Big Tech's best while remaining cost-efficient."

## Integration & impact on Ainnocence's platform

The upgraded model is now being rolled into Sentinus Al® (antibody engineering), Carbon Al® (small-molecule & PROTAC design) and Cellula Al® (cell-therapy optimization). Higher-fidelity predictions of stability and expression accelerate antibody lead-opt studies, while enhanced activity/binding scores improve small-molecule hit selection – compressing discovery timelines from years to weeks.

Ainnocence invites research groups and biopharma partners to leverage its sequence-first Al platform for protein engineering, vaccine work and hard-to-drug targets. Contact service@ainnocence.com\( \text{o}\) or visit\( \text{\text{www.ainnocence.com}}\) for collaboration details.\( \text{\text{0}}\)

#### **About Ainnocence**

Founded in 2021, Ainnocence is a next-generation biotech company whose self-evolving Al platform can virtually screen 10<sup>10</sup> protein sequences or small-molecule candidates for multitarget and multi-objective optimization, optimizing multiple properties simultaneously to deliver high-probability leads with unprecedented speed and cost efficiency.

Lurong Pan, PhD
Ainnocence
+1 205-249-7424
service@ainnocence.com
Visit us on social media:
LinkedIn
YouTube

This press release can be viewed online at: https://www.einpresswire.com/article/842332389

EIN Presswire's priority is source transparency. We do not allow opaque clients, and our editors try to be careful about weeding out false and misleading content. As a user, if you see something we have missed, please do bring it to our attention. Your help is welcome. EIN Presswire, Everyone's Internet News Presswire™, tries to define some of the boundaries that are reasonable in today's world. Please see our Editorial Guidelines for more information.

© 1995-2025 Newsmatics Inc. All Right Reserved.