

# Protera's Al technologies in focus

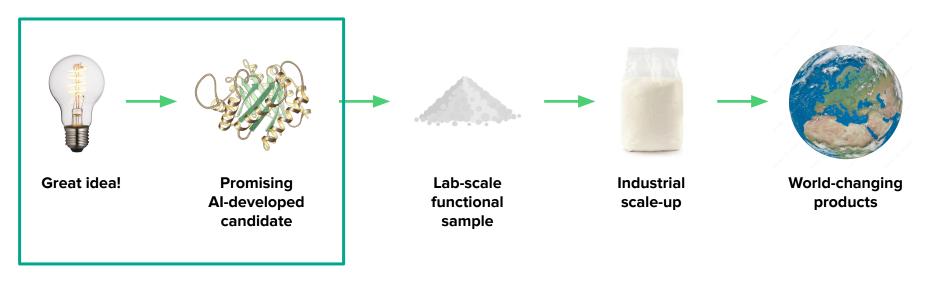
July 2025



# Most protein AI hype is about design

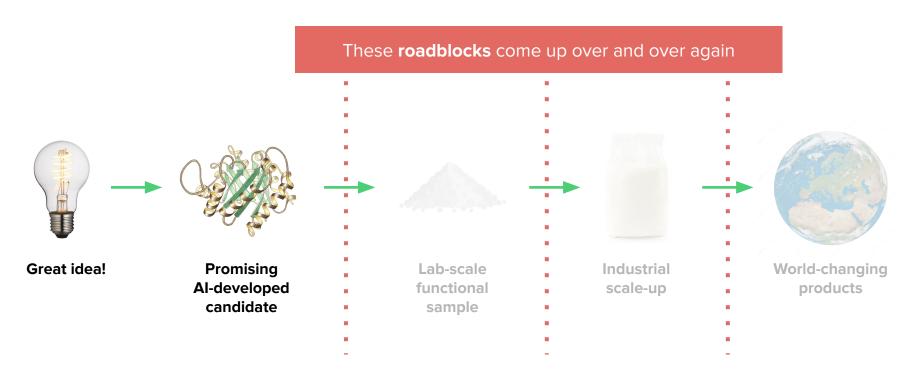


**Hype:** *De novo* design & functional optimisation



#### Most protein solutions fail in production





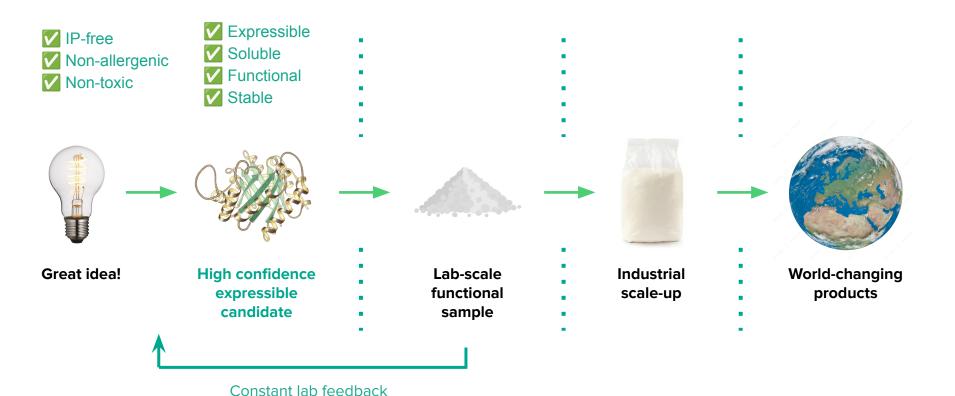
**Poor expression** 

Difficulty scaling up. Poor stability/solubility

Regulation. IP.

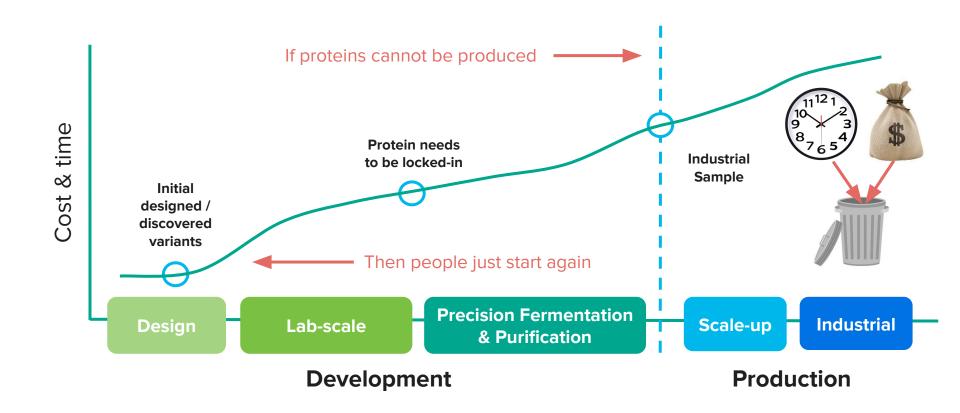
# Our AI enforces producibility & compliance





# Our competitors waste time and money





# Our capabilities span three crucial areas



#### **Protein Discovery**

#### Semantic search

Find novel IP-free proteins

#### **Curated GRAS data sets**

Patent-free, safe proteins

#### Safety filtering

Predict allergenicity & toxicity

#### **Function Prediction**

#### **Zero-shot models**

Data-free protein optimisation

#### **Active supervised learning**

Maximise protein function

#### **Protein-peptide interactions**

Predict binding affinity

#### **Design for Production**

#### **DNA** sequence design

Al expression optimisation

#### **Solubility analysis**

Avoid aggregation

#### **Stability optimisation**

For function & shelf-life



#### **Highlighted technologies:**

- 1 / Semantic protein search
- 2 / Protein-peptide interactions
- 3 / Active learning
- 4 / Predicting expression





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#### **Problem**

Sequence/structure-based search methods miss distant analogs, and often return IP-restricted proteins

#### **Our Solution**

Our semantic search can find IP-free proteins with similar function but completely different sequence/structure.

# Sequence search doesn't always work





Sequence-based search does not always find results with the same meaning.

# Semantic search gives better results





**|** 

Semantic embedding

 The feline lounged on the carpet

The mat had a cat on it

Dave the kitty sprawled on his rug

Mačka je sjedila na prostirci

The cat sat on the flat mat

猫坐在 垫 子上

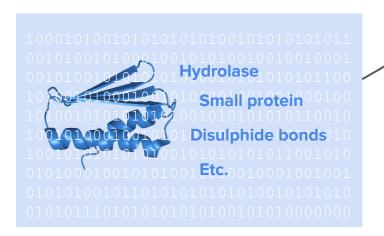
Semantic search can find sentences with the same meaning but **0**% **sequence similarity** 

# Semantic search gives better results



#### **MVLSPADKTNVKAAWGKVGA**

▼ Semantic embedding



MVIAPSDKTEVLAGWGHVGA

**MVLAPADKTNVKAGWGKVGA** 

**MQISTSELHQILGGTAKIAG** 

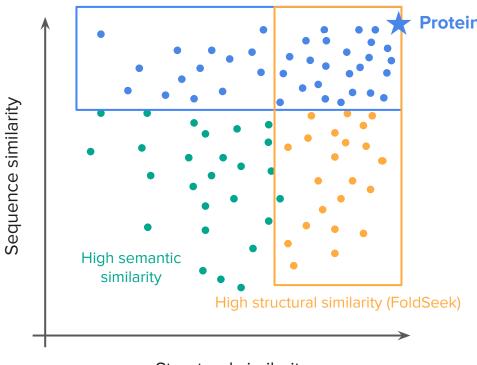
KLPOQRTAVNMLGGTAKIAA

Semantic search can find sentences with the same meaning but **0**% **sequence similarity** 

#### Protera Discovery finds distant analogs







#### **Protein of interest**

If the search protein is IP-restricted, usually the results found by BLAST and FoldSeek will be too.

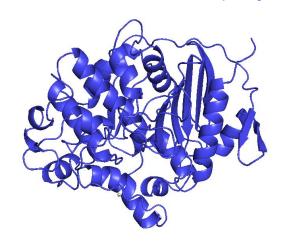
Protein Discovery finds the IP-free analogs that BLAST and FoldSeek miss.

Structural similarity

# **Example: Beta lactamase**



Industrial  $\beta$ -lactamase enzyme from bacteria used as query:



Returned experimentally confirmed **B-lactamase** enzymes from fungi

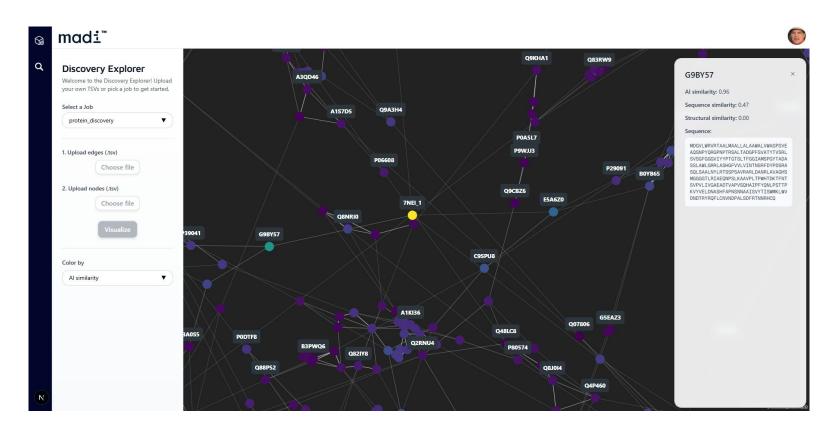


E. g. 94.6% Al similarity

No sequence or structure similarity

# Interactively explore protein space







#### **Highlighted technologies:**

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2 / Protein-peptide interactions

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4 / Predicting expression





#### **Problem**

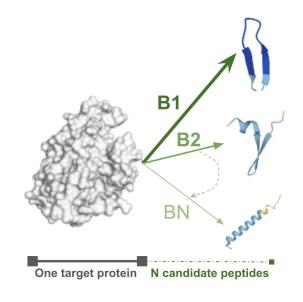
Understanding peptide-protein interactions (PePIs) is vitally important. Quantitative data to understand their strengths is lacking.

#### **Our Solution**

Our unsupervised ranking method can rank peptide-protein interactions without needing labelled data.

#### **Peptide-protein interaction strength matters**





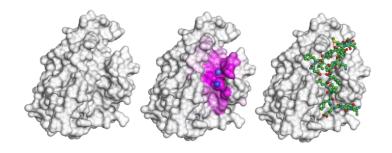
Binding affinities can be ranked

Identifying the **best protein-peptide matches** is key for bioengineering and therapeutic applications

# Unsupervised Als overcome data scarcity



1 protein-peptide measurement costs \$1000s and takes weeks.



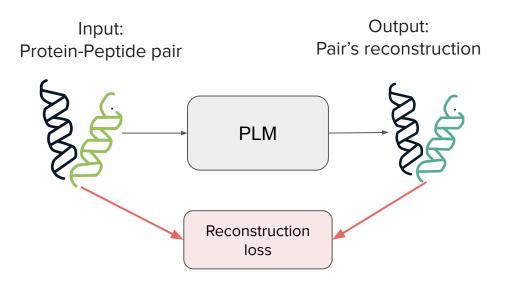
"Supervised" Als need large amounts of **labelled data**.

We would need thousands of measurements to train a supervised Al.

Unsupervised methods recognise universal patterns and require **0 measurements**.

#### PLMs understand peptide interactions





If PLMs (Protein Language Models) understand intra-sequence interactions, why not inter-sequence?

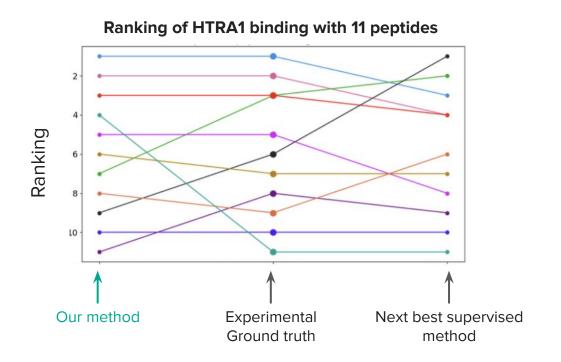
Our **PLM** ranks **protein-peptide interactions** in an **unsupervised** manner using reconstruction loss



# Our AI can quickly rank binding affinities



We validated the efficacy of our method across **5 different case** studies crucial in cancer, Alzheimer's and HIV research.



in neurodegenerative
diseases like Alzheimer's

Our method can screen thousands of candidates in minutes.





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#### **Problem**

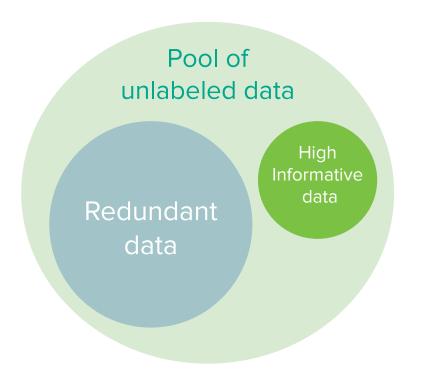
Al models need real-world data in order to understand proteins. Data is expensive.

#### **Our Solution**

Our active learning pipeline reduces costs by telling us which experiments will help our models learn best.

#### Not all data points are equally informative





Enriching our training database must involve a careful selection process

Active Learning allows our Al to determine the data it wants to learn from

#### Our AI knows best what it needs



# Our Al detects the key data points that boost performance, and requests labelling

Performance of our AI when selectively adding highly informative data



10X cost reduction

Boosts project feasibility and success



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#### **Problem**

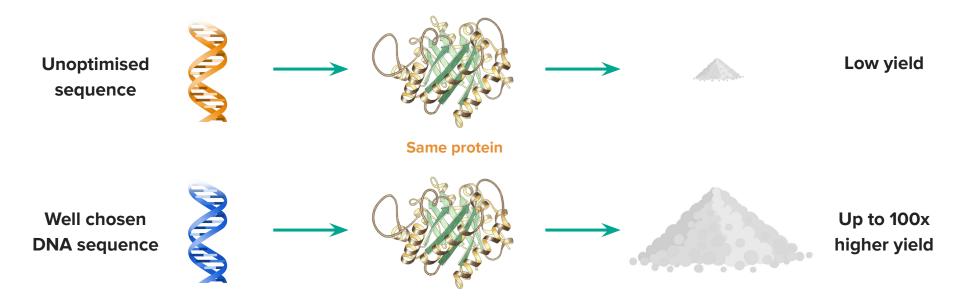
Standard methods to maximise protein expression are not good enough.

#### **Our Solution**

Our proven, best-in-class expression predictor offers precise control over protein expression levels.

#### **DNA** sequence choice affects expression





The choice of DNA sequence can **profoundly affect protein production efficiency**.

#### Codon "optimisation" tools are blunt



```
Fixed protein sequence:

Met-Ser-Thr-Pro-Gly-Leu-Lys...

ATG TCC ACC CCT GGT CTC AAA...

ATG TCT ACA CCG GGA CTG AAG...

ATG AGC ACT CCC GGG CTT AAG...

ATG AGC ACT CCT GGC CTT AAG...

ATG TCA ACC CCA GGC CTC AAG...
```

Tools like GenScript's **GenSmart** deliver a single "best" sequence.

- ★ no context ★ can't compare existing sequences
- ★ no fine control ★ only a single option each time

#### Our AI quantifies expression



Fixed protein sequence: Met-Ser-Thr-Pro-Gly-Leu-Lys...

**Essentially infinite possible RNA** encodings: ATG TCC ACC CCT GGT CTC AAA... 6.2 ATG TCT ACA CCG GGA CTG AAG... 34.2 ATG AGC ACT CCC GGG CTT AAG... 92.3

ATG AGC ACT CCT GGG CTT AAG... 65.2 ATG TCA ACC CCA GGC CTC AAG... 0.9



contextual

can be used to compare known sequences

tune down expression to keep cells happy can try multiple options in vitro

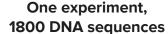
# Our AI model performs extremely well

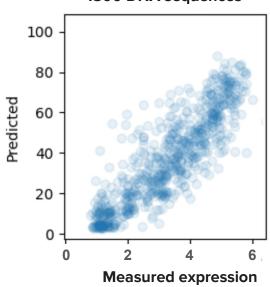


validated on 120k sequences across 72 experiments.







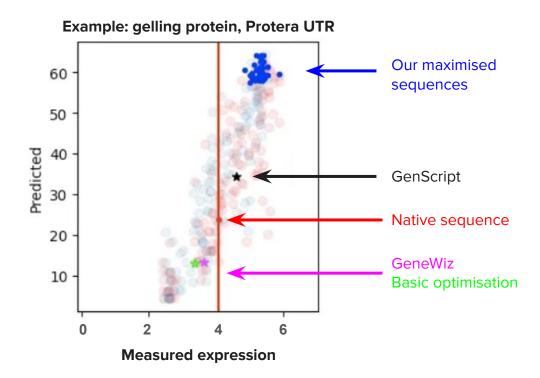


# Spearman ρ: 0.88

Largest ever DNA expression study

# **Our AI outperforms competition**









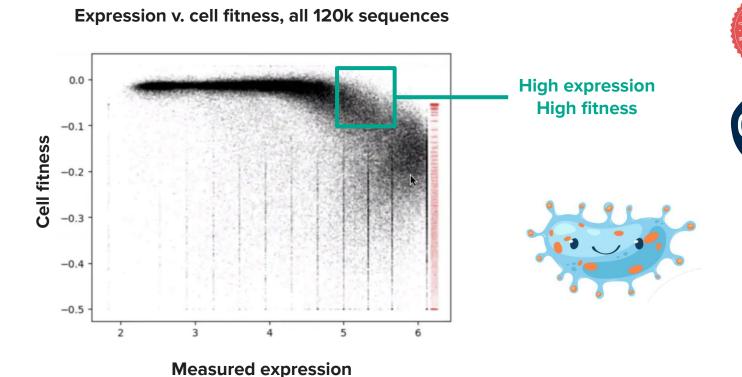
Our Al picked higher-expressing sequences than **GenScript** in

81%

of cases

#### Our AI helps find the expression sweet spot





# Our AI helps us compare & contextualise

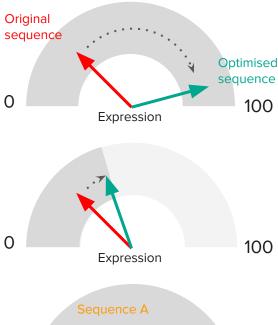


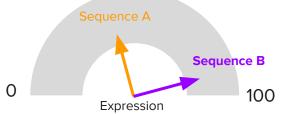
Everyone **assumes** expression optimisation works like this:

In reality, many proteins don't have **any** high-expressing DNA sequences.

Traditional methods will not tell you this.

Our method can also compare the **sequences** you already have in stock



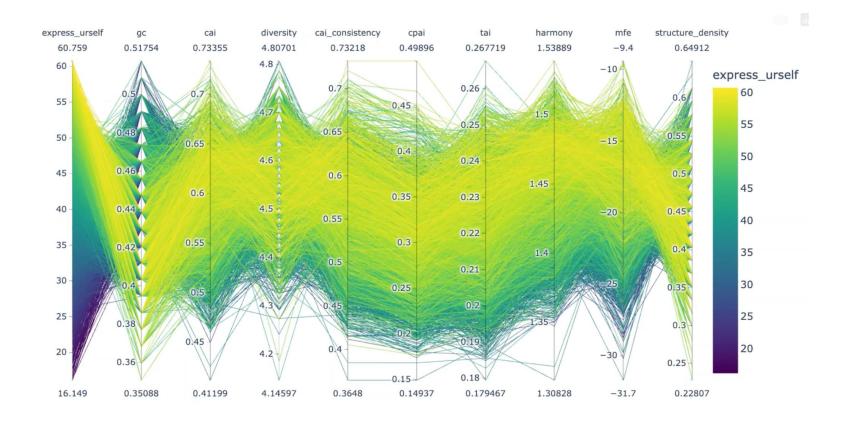






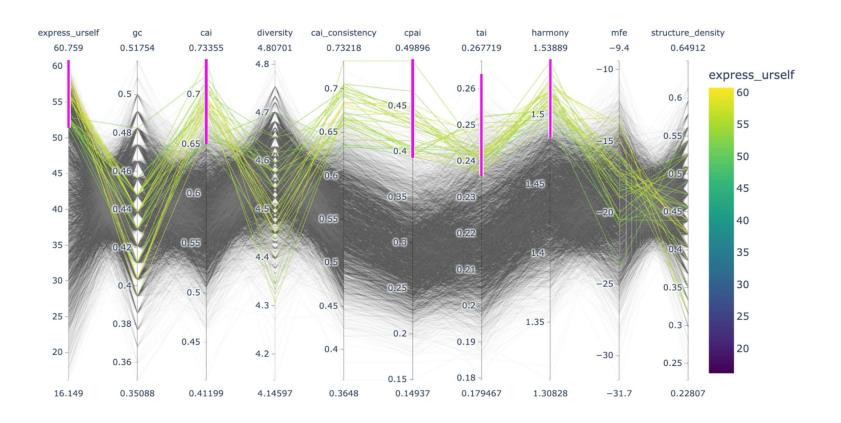
# Our tool lets you interactively optimise





# Our tool lets you interactively optimise







# **Case studies**



# **BASF** — Optimizing a commercial enzyme





Client optimised an enzyme for **over a decade** 

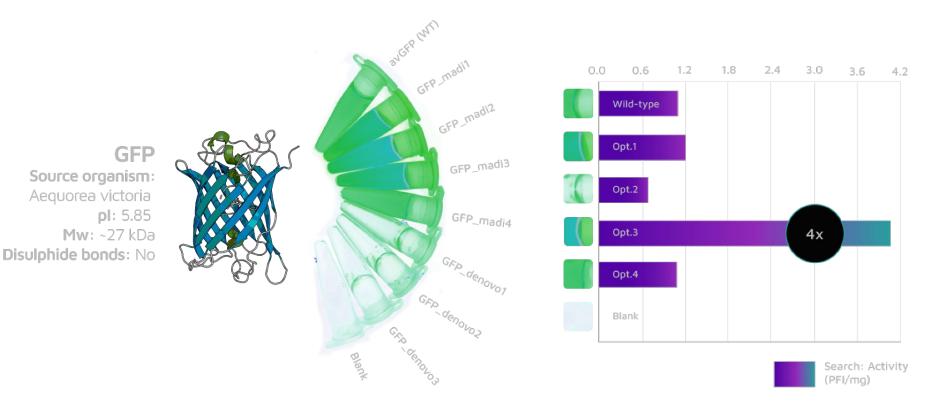
#### Target:

Protera to optimise application performance

Protera created fine-tuned models and delivered IP-ready enzymes with 4x higher performance

# Our optimised GFP – 4x brighter





Models are trained on **single point mutations to predict high-order mutations** (i.e. more than 2-3 mutations per variant)



# Our platform on antimicrobials



#### Our new focus:



# Developing <u>antimicrobials</u>, leveraging our expertise and progress in <u>bakery antifungals</u>

4 years of R&D, lab testing, consumer insights and machine learning give us a competitive advantage in antimicrobials:

- Safeguard against bacteria, fungi, and viruses
- **£113 billion** market in food, personal care, crop protection, and healthcare

#### And a promising candidate for personal care:

Key microorganisms	Protera Activity
Candida albicans	<b>V</b>
Escherichia coli	V
Staphylococcus aureus	V
Pseudomonas aeruginosa	V
Aspergillus brasiliensis	=

#### Our first candidate product ready for testing in application:

- Outperforms existing preservatives in just one Al iteration
- Low allergenicity + high purity
- High performance in neutral pH aqueous solutions
- Production scaled to 40L fermenter, yielding **50g in one month**

# **Antimicrobials for personal care:**



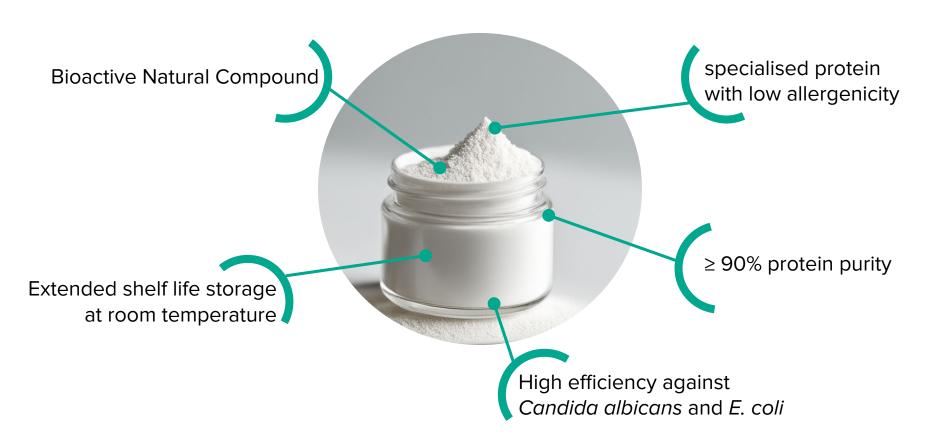
# a public health issue

- Preservatives are critical ingredients, yet most are harmful chemicals
- Negative consumer perception about Parabens → Proven Endocrine Disruptors
   ⇒ will probably be banned in EU in 2029
- Current replacements create irritation, allergic reaction, and endocrine disruption and natural ones are considered hazardous (CLP classification).
  - → Replacement of harmful chemical preservatives
  - → Low allergenicity product
  - → Non-toxic product
  - → Product compatible with skin needs

#### Protera product,

#### • protera

# antimicrobial plant based protein





# **Conclusion**



#### A quick recap



1 /

Our semantic protein search finds IP-free candidates with completely distinct sequences.

3/

Our **active learning** pipeline reduces costs by telling us which experiments will help our models learn best.

2/

Our **protein-peptide ranking** method lets us quantify interactions to find beneficial variants.

4 /

Our proven, best-in-class expression predictor offers innovators precise control over protein expression levels.



# Thanks!

