

## Product description & Innovation:

**KSI-044469: a first-in-class oral, BBB-penetrant small molecule targeting a novel Parkinson's Disease (PD)-linked protease to prevent toxic  $\alpha$ -Synuclein ( $\alpha$ -Syn) aggregation.**

- Discovered through AI-powered phenotypic screening of patient-derived dopaminergic neurons.
- Decreases oligomeric and monomeric  $\alpha$ -Synuclein *in vitro* (Figure1).
- Encouraging first *in vivo* experiments in healthy mice (Figure2).
- Strong drug-like properties with clean early safety profile.

## Unmet need & Market:

- > 10 million people affected by PD worldwide<sup>1</sup>.
- No disease modifying treatment options available.
- $\alpha$ -Syn accumulation is described as the biological driver of PD<sup>2</sup>.
- $\alpha$ -Syn is considered as a hardly druggable target.
- Fast-growing market ( $\approx$ 5% CAGR) reaching US \$6 - 8B by 2030<sup>3</sup>.

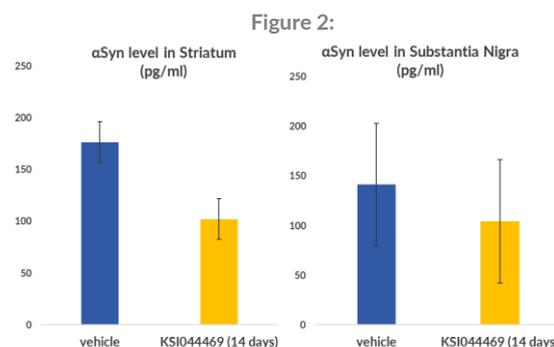
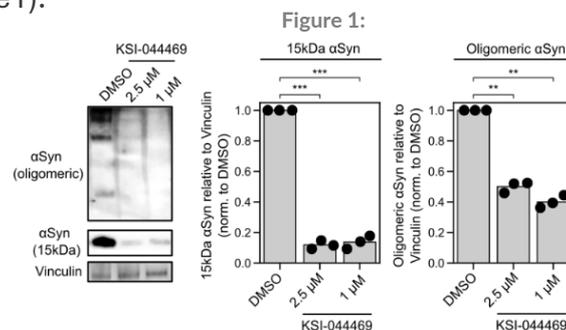
## Positioning – Why invest now:

Our approach is unique to overcome past clinical failures:

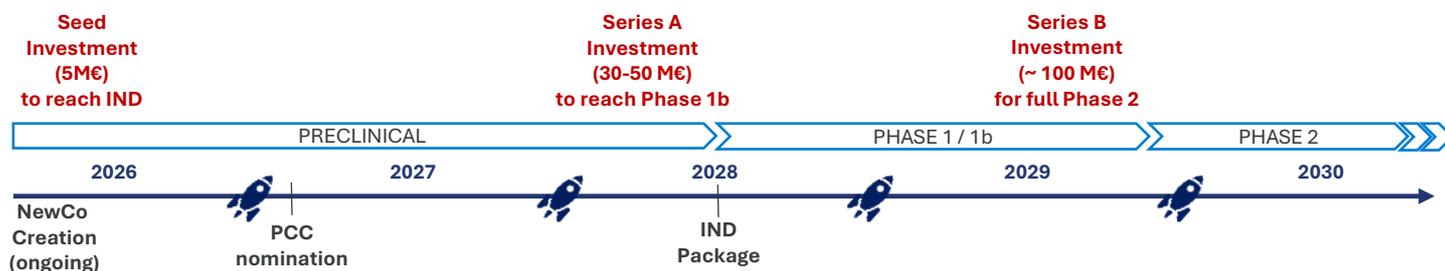
- CNS-penetrant small molecule with confirmed brain exposure
- Upstream regulatory activation with preserved neuronal function, driving catalytic intracellular  $\alpha$ -syn degradation.
- Non-immunogenic, pathway-driven mechanism without antibody/RNA drawbacks.

## Biomarker & Clinical Differentiation:

Our goal is to deliver an oral therapy, enabled by biomarker-based enrichment, for broad synucleinopathy use. The treatment will be fully compatible with symptomatic treatments and potentially preventive as early-detection biomarkers mature.



## Key milestones and development plan 2026 - 2030

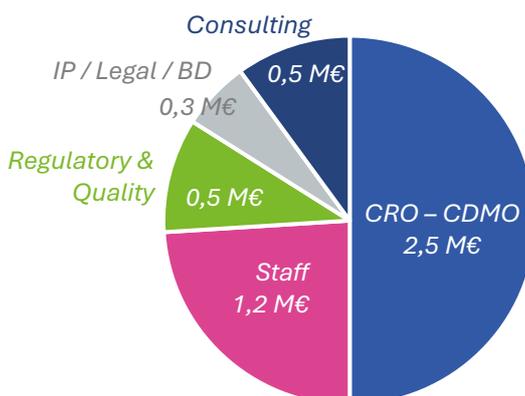


## Fundraising: Seed of 5M€ over 30 months

NewCo planned to raise 5 M€ to nominate PCC & complete IND-enabling studies.

## Exit scenarii:

- Licensing at IND (up to 75M€)
- Series A (30-50M€) to reach Phase 1b



## CONTACT

Mona Boyé  
mona.boyé@ksilink.com  
Mob: +33 6 88 26 64 04