

Novel Molecule Development for the Treatment of Leukemia and Other Cancers

*High Specificity, Three Mechanism, Low
Toxicity, Chimeric & Single Mechanism
Tubulin Molecules*





Leukemia and Brain cancer are the two most prevalent forms of cancer in children+

The two most fatal forms in all
age groups are AML and GBM

+American Cancer Society 2021





Current pediatric treatments are limited by drug toxicity

Many adult therapeutic regimens are not translatable to pediatric standards of care





**M-Life™ Chief
Discovery
Officer**

**A New Approach:
Targeting cancer
but not normal cells**

Peter Crooks, PhD, D.Sci, FRSC, FRPharmS
World renown cancer researcher and drug developer

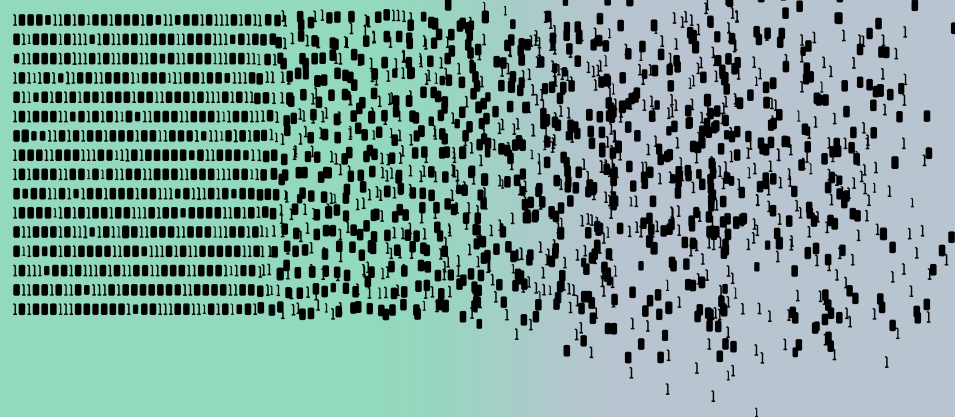


What Makes
Us Different

**Traditional Medicinal
Chemistry**

Directed AI

Big Data Fusion





Results to Date

Tubulin Inhibitors

- ◆ 8 high value assets identified via M-Life™ platforms and entire portfolio licensed (550)
- ◆ Synthesized and invitro tested by the National Cancer Institute NCI60 Panel
- ◆ Low toxicity and high efficacy profiles
- ◆ IP covered by 5 issued patents and licensed to M-Life™

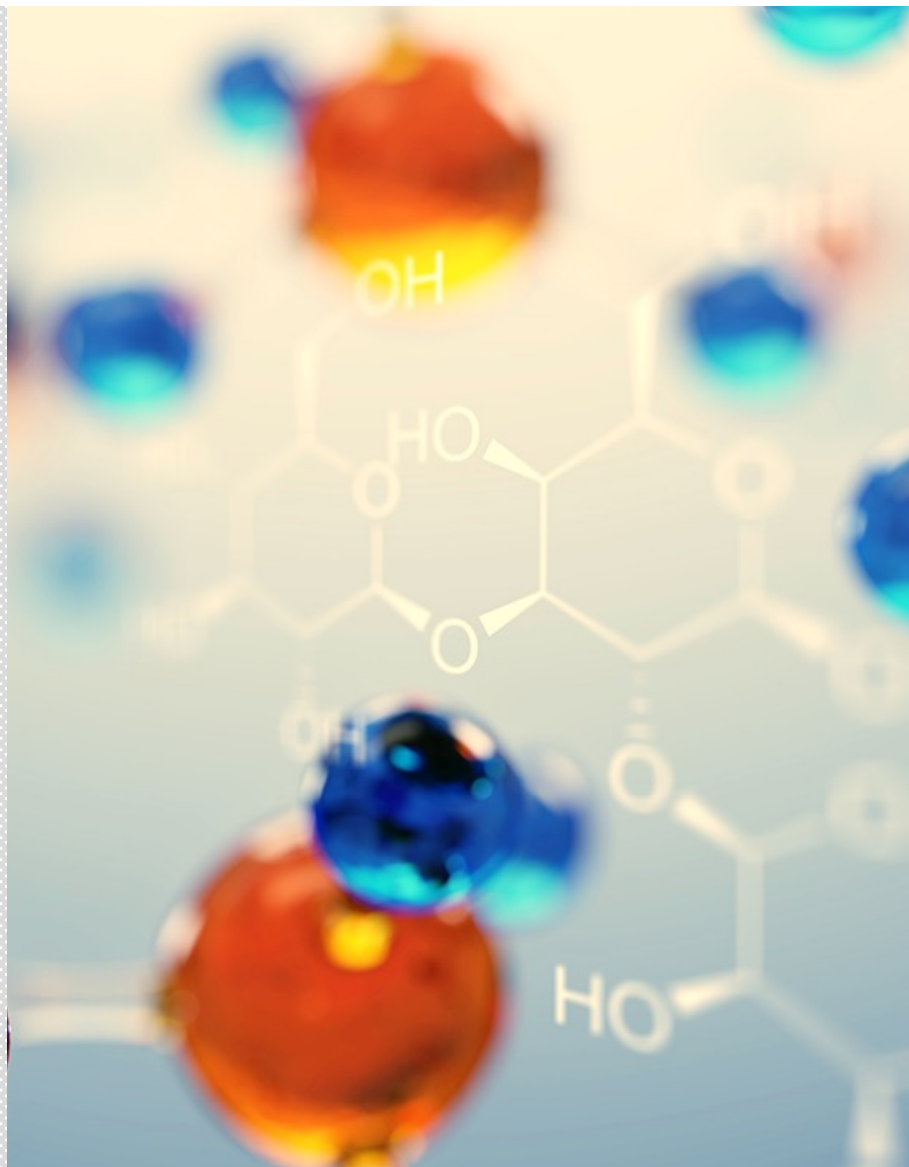
MTA Chimeras (tubulin, glutathione & NFkB)

- ◆ 10 M-Life™ platform designed chimera molecules
- ◆ Lower toxicity than synthesized tubulin molecules with much higher efficacy
- ◆ Pediatric and adult therapeutic use profiles indicating use across most cancer types

Platform Strengths

The ability to:

- ◆ Select 'High Value Molecules' from a sea of possibilities
- ◆ Identify the features responsible for toxicity & efficacy
- ◆ Design optimal molecules...VERY, VERY, VERY FAST & INEXPENSIVELY
- ◆ Identify new modes of action



Strategy

Leverage Out Strengths: (1) Identify Low Tox, High Value Assets (2) Design Novel Molecules

<6-9 mos.

Focus on Pediatric AML & GBM

Development

- Chimera and Tubulin Added IP
- Chimera Small Scale Synthesis
- Chimera NCI60 Testing
- Tubulin In vivo Mouse Testing

CDN Financing

- CDN Approx. \$500k
- Ops Only Exp.
- No Salary or Wages
- Converts at Series A

18-24 mos.

FDA Fast Track Process

Development

- API Scale-up via TCG
- GLP Tox via CRO
- IND
- Possible Phase I

Series A Financing

- \$10M+ Equity
- Institutional
- CDN Conversion

<12 mos.

Exit via Sell-Off

Development

- Expand Use to other cancers
- Select Next Programs

Sale of Cancer Molecules

- > 100M up front
- > 500M back end
- Company Profits Re-Invested in New Programs

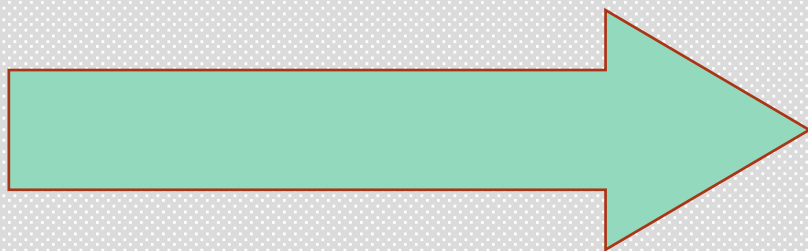


Opportunity

Objective: Drive Favorable Series A Deal Terms

Convertible Debt Financing

- \$10K Minimum
- 3 to 5X Multiplier
- Maximum Offering \$500K
- 4 year Automatic Term Conversion



Use of Proceeds

- 3-5 Chimera Synthesis
- Chimera Molecule NCI 60 Testing
- 8 Licensed Tubulin Molecule *in vivo* Testing
- Chimera molecule IP
- NO SALARIES OR WAGES
- NO OFFICE/BACK-OFFICE SPACE
- NO COMMISSIONS OR BONUSES



For more
information:

Ted Moskal
President, CEO & Founder
Moskal Lifesciences, LLC
tmoskal@m-lifesciences.com
870.680.2141