

# Winning BD Pitch Decks

---

**June-2025**

**BioPing - "Ping the Right Partner"**

# Disclaimer



This presentation contains information compiled from publicly available sources, industry reports, and third-party data believed to be reliable. It is intended solely for general informational and strategic guidance purposes within the life sciences and biotech business development context. The analysis and commentary reflect the presenter's personal views and do not constitute scientific, clinical, legal, or any sort of professional advice. No representation or warranty is made as to the accuracy or completeness of the information provided. Recipients are encouraged to perform their own due diligence and consult qualified professionals before acting on any insights shared. The presenter, BioPing and its owners disclaim any liability for decisions made or actions taken based on this content.

# Suggested Content

(Keep ~15–20 slides for introductory meeting)

## 1. Company Snapshot

- Who you are, what you do, and your vision
- Founders, team, platform/approach, location, IP
- Optional: tagline that sums up your differentiation

## 2. Platform / Pipeline Snapshot

- Put the asset in context of broader platform (if applicable)
- Optional: show platform potential with other assets

## 3. Unmet Need / Disease Background

- What's the medical need, why it matters now
- High-level market context
- Opportunity for differentiation

## 4. Mechanism of Action / Biology Rationale

- How your program works, biological rationale
- Why the target matters (and is validated)
- Competitive advantage (vs. SOC or failed approaches)

## 5. Preclinical Data Summary (for preclinical stage assets)

- Proof-of-concept (in vitro, in vivo, PK/PD, safety)
- Disease models used and why they're relevant
- Benchmark vs. SOC (if possible)

## 5. Clinical Overview (if in clinic)

- Trial design, patient pop, key endpoints
- Summary of safety and efficacy data (topline if ongoing)
- Any early signs of differentiation

## 7. Competitive Landscape

- Who else is working on this?
- Your edge: why you're different and better
- Avoid full grid — just key players and how you win

## 8. IP / CMC / Manufacturing

- Patent status (granted/pending) + coverage years
- Any differentiation from a formulation, delivery, CMC
- Optional: freedom to operate

## 9. Development Plan & Milestones

- Preclinical/clinical roadmap with timing
- What inflection points are upcoming
- Cost/effort estimate to get to next value step

## 10. What You're Looking For

- Licensing? Co-development? Option deal?
- Flexibility or open to structure
- What makes now the right time to engage

# Additional Suggestions

## For Preclinical vs Clinical Stage

Element	Preclinical Asset	Clinical Asset
Strong emphasis on	MoA, in vivo models, biomarker strategy, target rationale	Safety, efficacy signals, study design, patient outcomes
Must-have data	Dose-response, tox, PK/PD, target engagement, reproducibility	Summary table of trial data, adverse events, comparator data
Optional	Companion diagnostic plan, IND-ready status	KOL feedback, patient experience data, extension plans

### Final Tips:

- Keep visuals clean, not data-dump slides
- Highlight 2–3 differentiation points consistently
- Avoid overpromising — just enough to **build interest**
- Have **backup slides** with detailed data if asked
- Tailor to the audience (scientific vs. BD vs. commercial)



## Contact

CEO and Founder

Gaurav (Vik) Vij

[gvij@cdslifescigroup.com](mailto:gvij@cdslifescigroup.com)