

Research reports of Biotech on Hub&Spoke Model

Xiang

XBI since Jan-31-2006

XBI SSGA Funds Management Inc

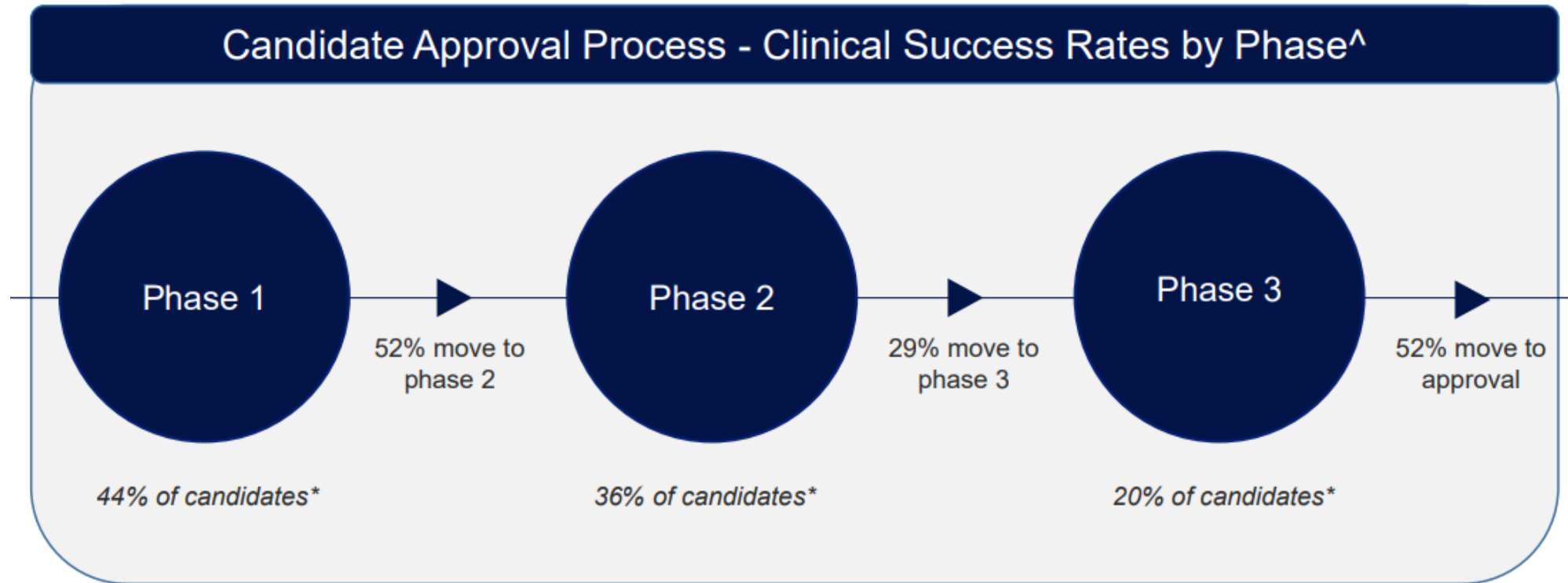
As of Dec-31-2023 \$7.0B

Sep-01-2020 O: 111.31 H: 116.64 L: 100.64 C: 111.43 V: 112.97 M

Events ▾

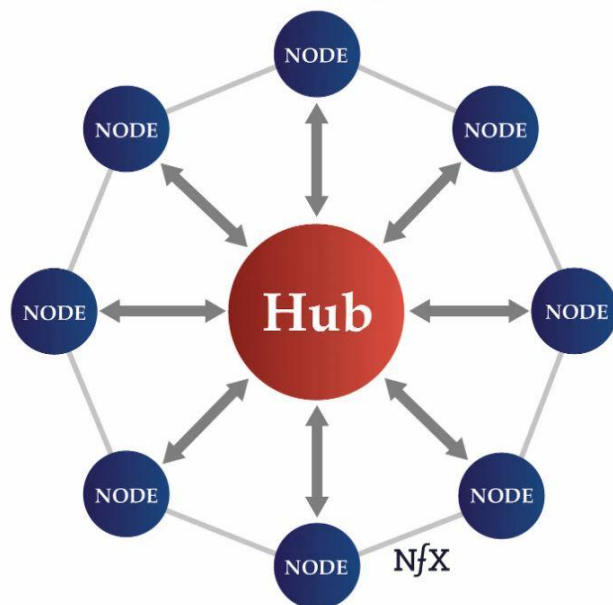


Struggle on drug developments

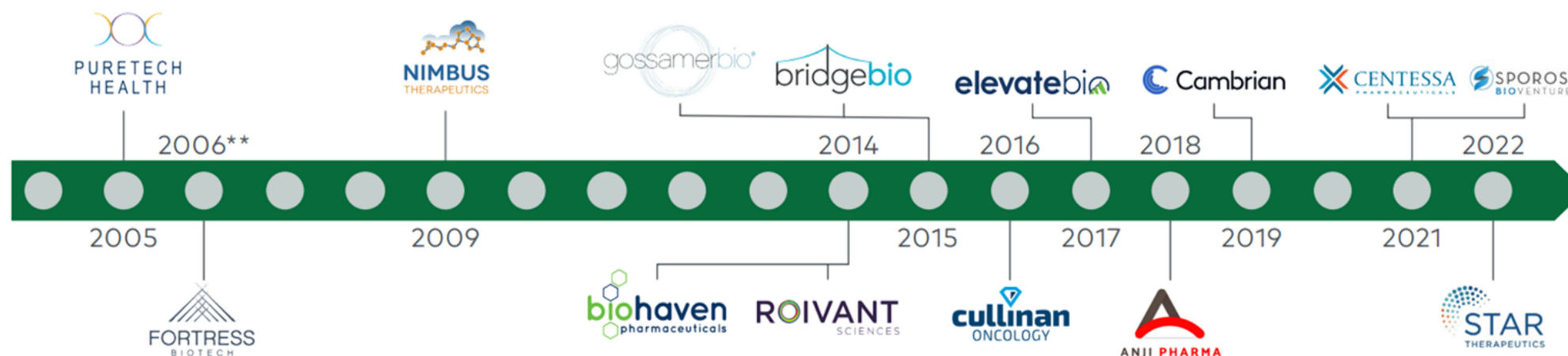


Emerging Biopharma Trend

Hub & Spoke



\$Sticker (Name if private)	Market cap (as 01/31/24, Billion)
\$ROIV (Roivant)	7.95
\$BBIO	5.9
\$BHAVN	3.6
\$CNTA	0.77
\$CGEM	0.76
\$PTCHF	0.65
\$GOSS	0.19
\$FBIO	0.079
ElevateBio	
Nimbus Thx	



FBIO

- **Intro: 10 companies**
- **Ownership:** Market Cap 34 M + 45M FBIOP
 - Chairman 32%, COO 15%, CFO 2.6%, Retail 27.2%,
- **Revenue:** 15M/Q from \$DERM,
- **Cost:** -62M/Q, FTE: 20 (5LT, 6 accountant) + subsidiaries
- Sum: from 80M to 300~400M, >3X increase

Convention:

■ = cost, ■ = income, ■ = alert



From SEC10Q 3Q23

FBIO Pipelines

	Partner Company	Asset(s)	2021	2022	2023	2024	2025	2026	2027	2028	2029	2030+	Potential Peak Sales* (Global)
+60M	Journey (DERM)	Commercial Portfolio (Qbrexza, Amzeeq, etc.)	\$63.1M <i>Net revenue</i>	\$73.7M <i>Net revenue</i>	■	■	■	■	■	■	■	■	● ●
		DFD-29 (small molecule for rosacea)				■	■	■	■	■	■	■	
+10~100M	Checkpoint (CKPT)	Cosibelimab (anti-PD-L1 mAb for cSCC)				■	■	■	■	■	■	■	● ●
		Olafertinib (3rd gen EGFRi)					(Asia)						
+100M	Cyprium^	CUTX-101 (copper histidinate for Menkes)	\$8M <i>Milestone</i>		\$4.5M <i>Milestone</i>	▲ ■	■	■	■	■	■	■	● ● ●
+40M	Caelum**	CAEL-101 (fibril-reactive mAb)	\$56.9M <i>Monetization</i>			■	■	■	■	■	■		● ● ●
-10~50M	Avenue (ATXI)	IV Tramadol (small molecule for post-op pain)					■	■	■	■	■	■	● ●
		AJ201 (small molecule for SBMA)				Ongoing Phase 1b/2a clinical trial in SBMA							● ●
		BAER-101 (GABAA α2/3 PAM)				Potential Phase 2a initiation							● ●
+5M	Mustang (MBIO)	MB-106 (CD20 CAR-T for NHL)							■	■	■	■	● ● ●
		MB-109 (CAR-T + oncolytic virus for GBM)							■	■	■	■	● ●
		MB-117 (gene therapy for XSCID)							■	■	■	■	●
		MB-217 (gene therapy for XSCID)							■	■	■	■	●
		MB-110 (gene therapy for RAG1-SCID)				Ongoing Phase 1/2 clinical trial in Europe							●
+50M	Urica	Dotinurad (URAT1 inhibitor for gout)							■	■	■	■	● ● ●
+50M	Helocyte	Triplex (Cytomegalovirus vaccine)				Ongoing Phase 1 and Phase 2 clinical trials in various indications							● ●

■ = Anticipated product revenue/royalties ▲ = Potential PRV/milestone/monetization proceeds ■ = Potential regulatory approval ● < \$500M | ● ● \$500M - \$1B | ● ● ● > \$1B

Sum = 305M – 455M vs 34M Market Cap

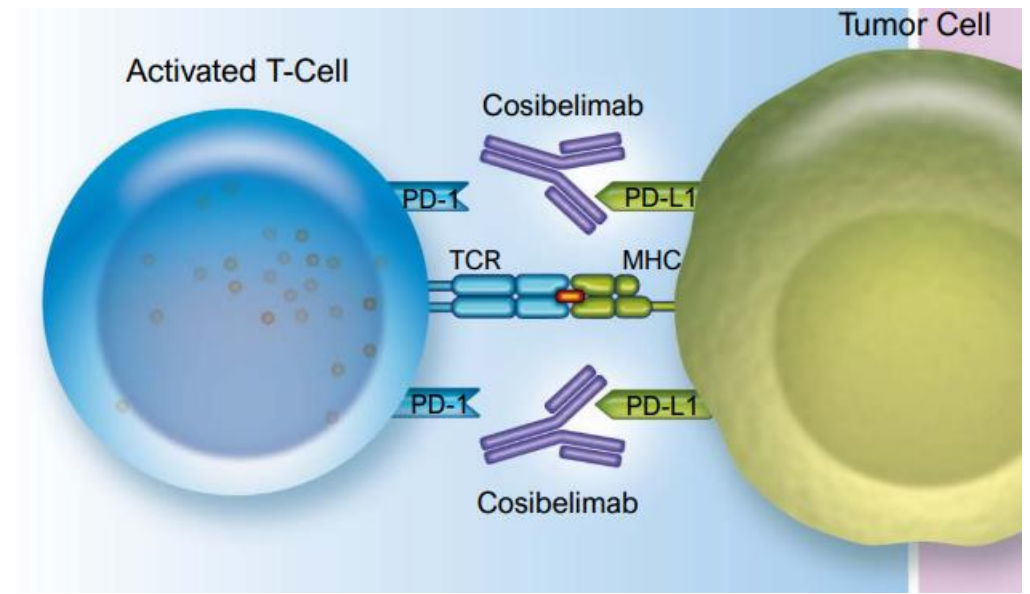
Journey \$DERM for Dermatology

- **Intro:** Treating skin conditions, healing wounds
 - Qbrexza®, Accutane®, Amzeeq®, Zilxi®, Luxamend®,
 - **Legacy:** Targadox®, Ximino®, Exelderm®
- **Ownership:** Market Cap 60M
 - FBIO 53%, CEO 16.8%, Retail %
- **Revenue:** 15M/Q, 15 yrs 0.5M/yr consulting fee to FBIO, 19M deal Maruho
- **Cost:** -22M/Q, 65 FTE (3LT, 3R&D, 2Admin, rest are sales)
- **Pipeline & Market:** *Table
- **FDA:** +50M Approval 2024 DFD-29, BiC for Rosacea vs [Oracea](#), 10~15% royalty to DRL (paid 34M to license from DRL)
- Sum: >110M

Product	2023 Revenue (\$)	2022 Revenue (\$)	Change (%)
Qbrexza®	18,038	19,752	-9%
Accutane®	15,109	14,228	6%
Amzeeq®	4,904	5,892	-17%
Zilxi®	1,567	1,851	-15%
Targadox®	2,386	6,558	-64%
Exelderm®	1,813	3,018	-40%
Ximino®	567	3,775	-85%
Luxamend®	21	—	100%

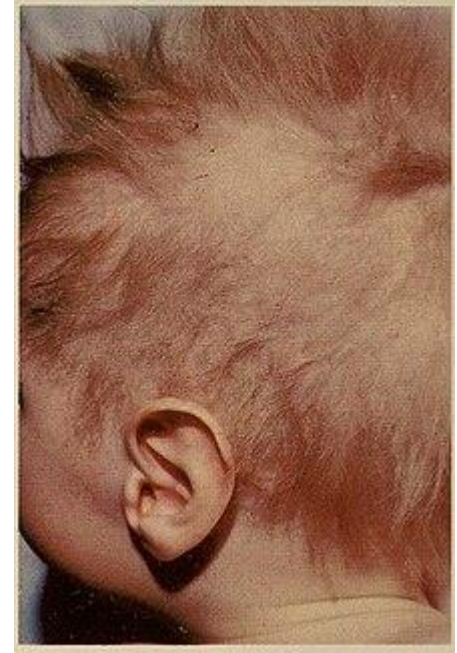
Checkpoint \$CKPT

- **Intro:** Treating Cancer: Cosibelimab(PD1)
- **Ownership: Market Cap~46M**
 - FBIO 20%, CEO 2.4%, Retail 70%
- **Revenue: 0**, 15 yrs 0.5M annual consulting fee to FBIO
- **Cost: -11M/Q**, ~20FTE (2LT, 15CMC, 2Admin)
- **Pipeline:** Cosibelimab
- **Market:** ~1.6B shared cSCC
- **FDA:** 1H2024
- **Sum:** >100M



CUTX-101 Cypritum Therapeutics

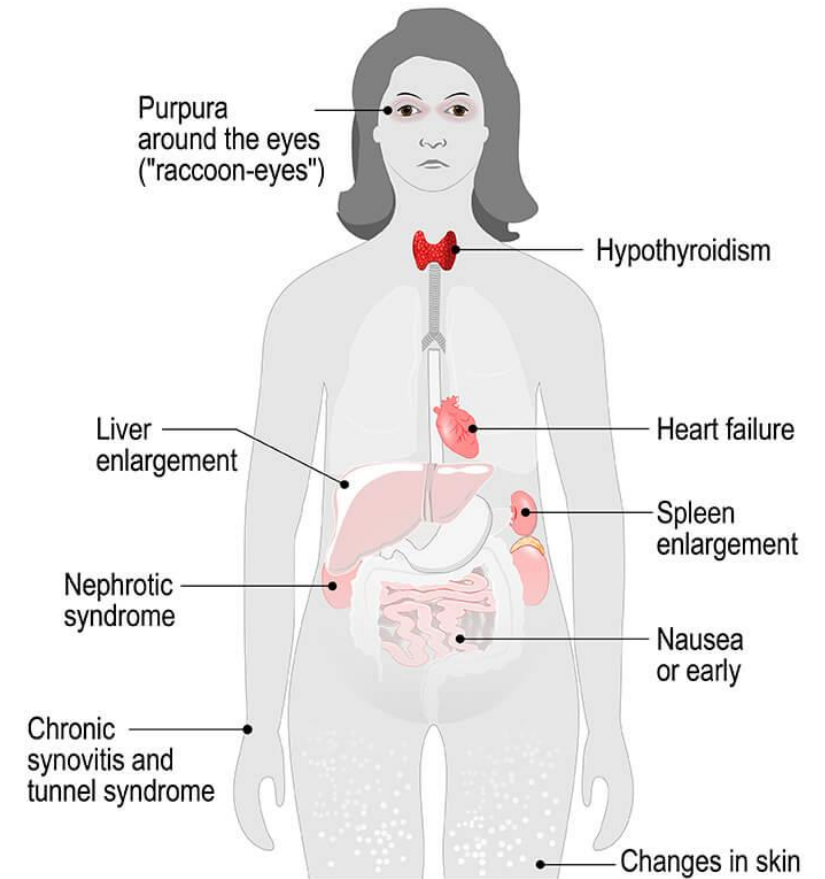
- **Intro: Menkes Disease (ATP7A Mut)**
- **Ownership:**
 - FBIO ~80%,
- **Revenue: 0**
- **Cost: 0 employee**
- **Pipeline: CUTX-101 (injected twice/day)**
- **Market: ~50~225 patients/yr in US**
- **FDA: Approval 2024 + 100M RPV**
- **Sum: >100M**



Caelum Biosciences (AstraZeneca) for light chain (AL) amyloidosis.

- 42% of future proceeds to Caelum from AstraZeneca
- ~\$56.9M 2021, **Another 42M from 100M is normal. Up 147M**
- AstraZeneca's Alexion acquired Caelum Biosciences on 10/5/2021 for up to \$500 million, including \$150 million upfront and up to \$350 million in future contingent milestone payments. FBIO received ~\$56.9 million of such upfront amount and is eligible to receive ~42% of the proceeds from all future milestone payments.

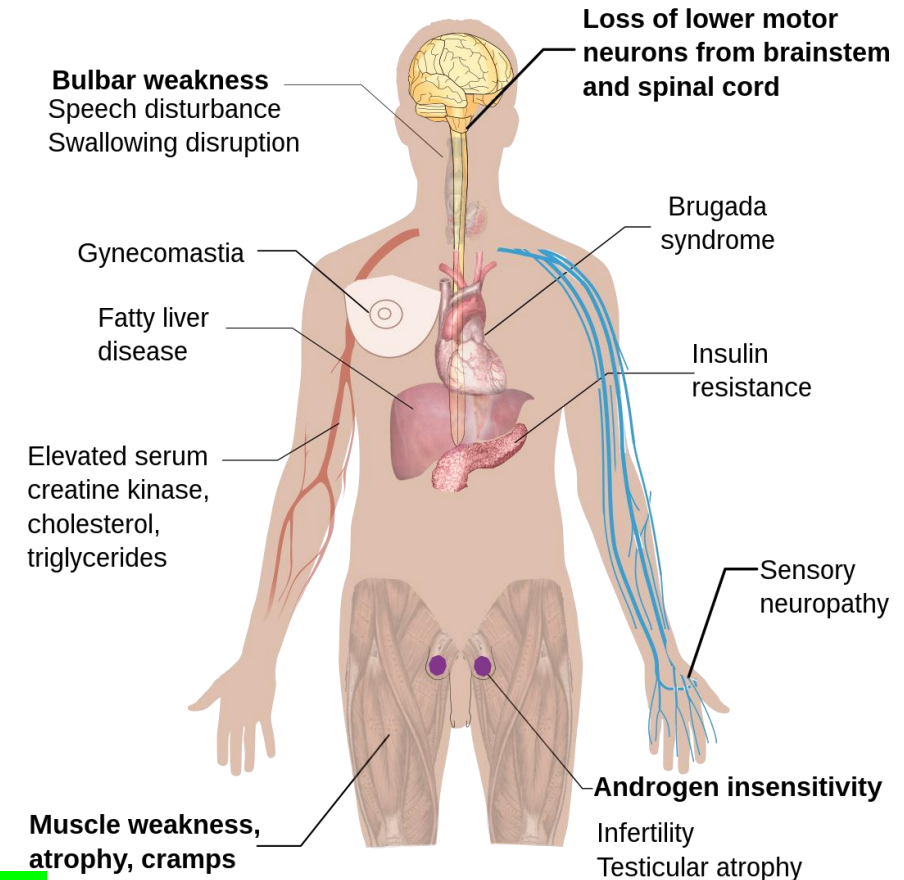
AMYLOIDOSIS signs and symptoms



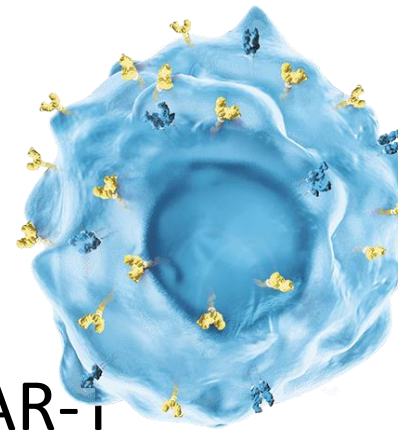
Avenue \$ATXI AJ201

- **Intro:** SBMA/Kennedy's Disease: AJ201 (ODD)
- **Ownership:** Market Cap 5M
 - FBIO 20%, CEO <1%, Retail 88% (Robert.D 10%)
- **Revenue:** 0, 15 yrs 0.5M annual consulting fee to FBIO
- **Cost:** -3.6M/Q, 10 FTE(5LT, 2CMC)
 - To FBIO, annual, $-3.6 * 4 * 12\% + 0.5 = -1.23M$
 - For all IND development, ~3-4yrs. -5M
- **Pipeline:** AJ201/BAER101/IV Tranmadol
- **Market:** ~shared 1.5B
- **FDA:** 2025 IV Tranmadol, rest 28/29
- **Sum:** Due hub&spoke model, this can be at least 0.5M/yr.

Impacts of SBMA



Mustang Bio \$MBIO



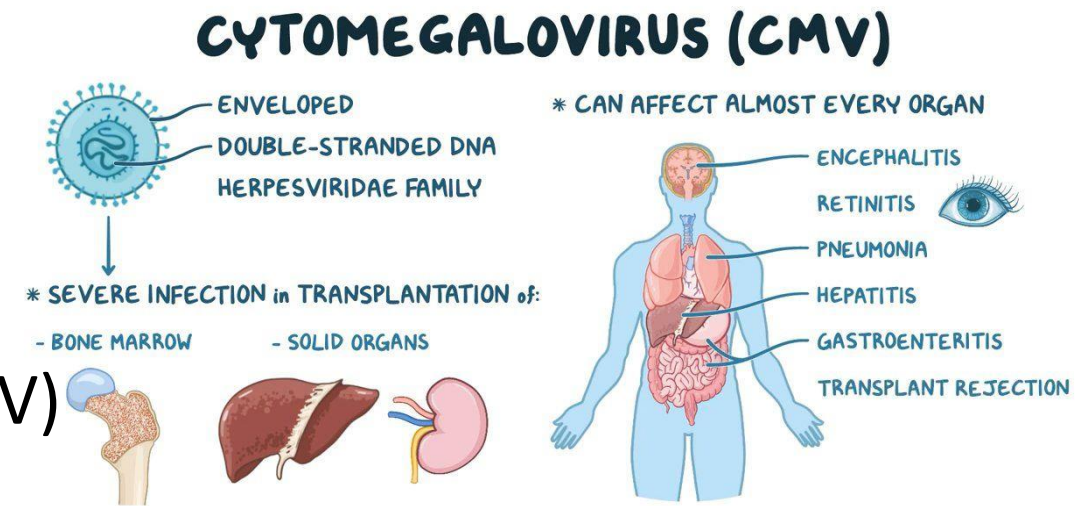
CAR T Cell

The T cell with the CAR added helps find and fight specific targeted cells

- Intro: Non-Hodgkin Lymphoma/CLL CAR-T
- **Ownership:** Market Cap 12.8M
 - FBIO ~30%, retail ~69.9%, CEO 1.2%
- **Revenue:** 0, 15 yrs, 1M annual MSA fee to FBIO
- **Cost:** -14.1M/Q, ~100 FTE (9LT, ~25R&D, ~25clinical),
 - To FBIO, annual, $-14.1 \times 4 \times 20\% + 1 = -10.8\text{M}$
 - For all IND development, ~5yrs. -55M
- **Sum:** Because of hub&spoke model, at least 0.5-1M/yr.

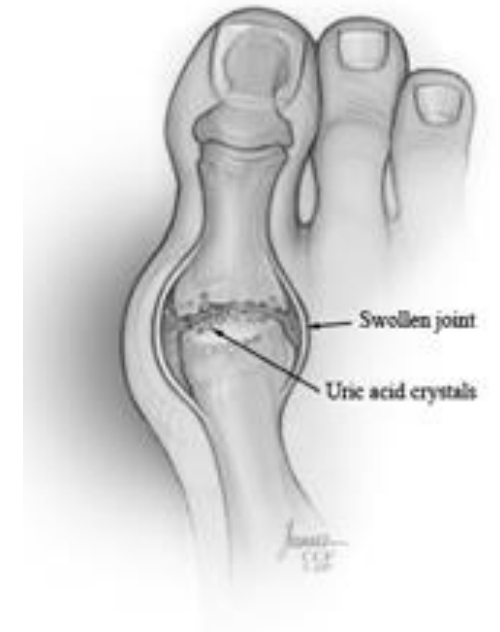
Helocyte Inc

- Immunotherapies for cytomegalovirus (CMV)
- **Ownership:**
 - FBIO ~90% effective
- **Pipeline:**
 - Phase2, 3.22 M Grant from NIH, Phase1b Funded by City of Hope
 - Phase 2: Liver Transplant (2024), funded up 20M from NIH
- **Cost:** 0 employee
- **Market:** ~1.5B shared SOT and HSCT
- **FDA:** ~potential Approval 2027
- Sum: >50M



Urica

- **Intro:** Treat gout and others associated with hyperuricemia
- **Ownership:** Market Cap M
 - FBIO 75%, CEO %, Retail %
- **Revenue:** 0, 15 yrs 0.5M annual consulting fee to FBIO
- **Cost:** 3 FTE (2.5LT),
- **Pipeline:** Dotinurad (approved in Japan, FUJI), oral URAT1 inhibitor
- **Market:** ~4M US gout patients, ~1.4B
- **FDA:** ~potential Approval 2027
- **Sum:** >50M



Summary of FBIO vs FBIOP

- Series (1 vs 1.09375^{yr}) Dividends accrue daily (effective 9.8%)
- **1 vs 1.098^{yr}**

Share Class	Common	9.375% Series A Preferred
# of shares	8.942M	3.427M
Price	\$1.8	\$13
If holding 3 yrs.	1	1.32
5yrs		1.60
10yrs		2.55
15yrs		4.06