



SCOTTIE
VENTURES

Healthcare and Biotech Industry Report

Associates

Saarthak Gupta

Analysts

Alisha Mehta
Charlie Hubbard
Jacky Sun
Ryan Kumbatovic
Sean Chen

Industry Overview



The biotech and healthcare industry is entering a major inflection point as rising chronic disease burdens and aging populations demand breakthroughs in reshaping medical services and treatments, specifically in their delivery efficiency and quality. Across the core sectors, including digital therapeutics & chronic care, cardiovascular medicine, genetic engineering & gene therapy, AI-driven drug discovery platforms, and medical devices and/or wearables, we're seeing sustained growth driven by increasing demand. This has resulted in increased medical awareness across generations, incentivizing more to adopt medical treatments and medications that were previously unattainable.

Digital therapeutics are becoming essential as employers and insurers shift toward prevention and continuous monitoring. Cardiovascular and longevity treatments remain stable, while gene therapy and embryo genomics are scaling fast as editing and reproductive screening move into wider adoption. At the same time, the foundation of modern healthcare, the AI-enabled virtual care systems and advanced medical devices, is expanding rapidly, pushing the entire system toward proactive and personalized care. The increasing monitoring and treatment programs cater to the growing populations of chronic disease patients. As capital, talent, and research increasingly converge around these high-impact sectors, the industry is positioning itself for faster AI-powered innovation cycles, broader clinical applicability, and improved patient outcomes materially.



Market Growth & CAGR

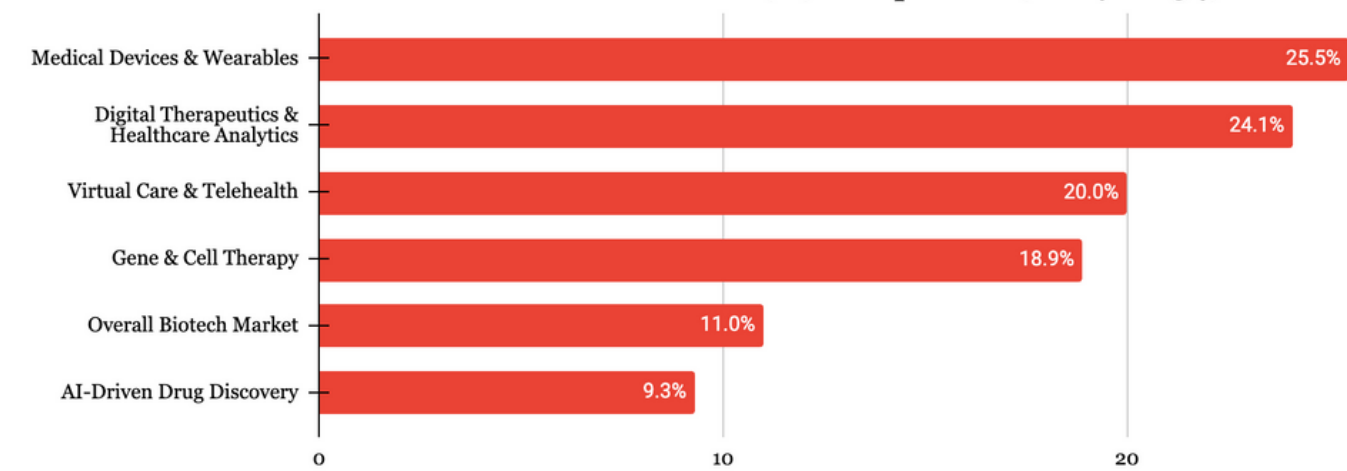
11%
CAGR for Biotech Overall

Between 2024-2034

Subsector	CAGR	Valuation Multiples	Market Size
Medical Devices & Wearables	25.50%	~17x (EV/EBITDA)	Wearables: +26% (LTM), Smartwatches: +218% (2024-2032)
Digital Therapeutics	24.10%	~16x (EV/EBITDA)	Analytics: +457% (2024-2032)
Virtual Care & Telehealth	20%	~15.8x (EV/EBITDA)	Virtual Care: +746% (2024-2032)
Gene & Cell Therapy	18.90%	~5.5 (EV/Revenue)	Gene Therapy: +230% (2023-2030)

The biotech sector is entering a high-growth decade, with the overall market expected to expand at an 11% CAGR from 2024–2034, driven by accelerating demand for next-generation therapeutics and AI-enabled innovation. Sub-industries such as medical devices & wearables (25.5%), digital therapeutics & healthcare analytics (24.1%), and virtual care (20%) are growing the fastest as the healthcare system shifts toward more scalable, data-supported treatment models. Meanwhile, gene and cell therapy (18.9%) continues to gain traction as genetic editing, embryo screening, and CRISPR-based treatments move closer to mainstream clinical adoption. These growth rates reflect how technological convergence such as AI, genomics, software, and diagnostics is reshaping the entire biotech industry and creating opportunities for breakthroughs. Market growth potential is strong due to increasing macro trends and focus as well.

Biotech Subsector CAGR (%) Comparison (2024–2034)





Key Trends

AI as a Catalyst for Discovery and Delivery

Artificial Intelligence has become the backbone of modern biotechnology, driving breakthroughs from molecule discovery to patient care. Platforms like DeepMind's AlphaFold and companies like Recursion Pharmaceuticals have shortened drug discovery cycles by years by slashing R&D costs. Modern AI is now found in diagnostics, medical imaging, and even virtual support tools, embedding itself in every corner of the biotech space.

AI is turning biology into a data problem that can be solved at scale.



Consumerization and Continuous Care

Biotech is shifting from hospital-based treatment to continuous health management. This change is occurring through breakthroughs in telehealth, wearables, and connected diagnostics that allow patients and doctors to monitor conditions in real time. Consumers are now able to track and engage with their personal health data to take a more active role in their well-being. Continuous care is transforming the traditional healthcare model to one based on convenience and prevention rather than spaced intervention.

Healthcare is moving into the hands of consumers.



Preventive and Personalized Medicine

Advances in genomics are enabling medicine to move from reactive to preventative. Gene therapies, embryo screening, and personalized interventions allow treatment (or even avoidance) of disease at the genetic root. Examples in practice include CRISPR and Orchid Health who are creating generalizing personalized medicine at scale.

Biotech is shifting from treatment to prevention.



The Rise of Platform Biotech

Future advancements in biotechnology involve building reusable development engines that enable systematic discovery. Modern biotech platforms are doing this at scale by leveraging a core technology that can be applied across dozens of diseases to create modern healthcare solutions. Platforms enable faster and cheaper therapy discoveries that are not 'one-off' discoveries.

The future belongs to platforms, not one-off drugs.



Tailwinds

Aging & Chronic Disease

Biotech is supported by aging populations and rising rates of diabetes, obesity, heart disease, and other chronic conditions, which drive sustained demand for new therapies, devices, and digital health tools.

AI & Advanced Modalities

AI/ML, gene editing, and next-gen therapies are shortening R&D cycles, enabling more personalized and potentially curative treatments, and attracting strong investment from pharma, VCs, and PE funds.

Supportive Regulation & Funding

Regulators like the FDA and EMA are creating faster approval tracks for breakthrough and rare-disease drugs, while governments continue to fund vaccine programs, pandemic preparedness, and advanced therapies, helping reduce time-to-market and de-risking innovation.

Headwinds

Costly, Slow R&D

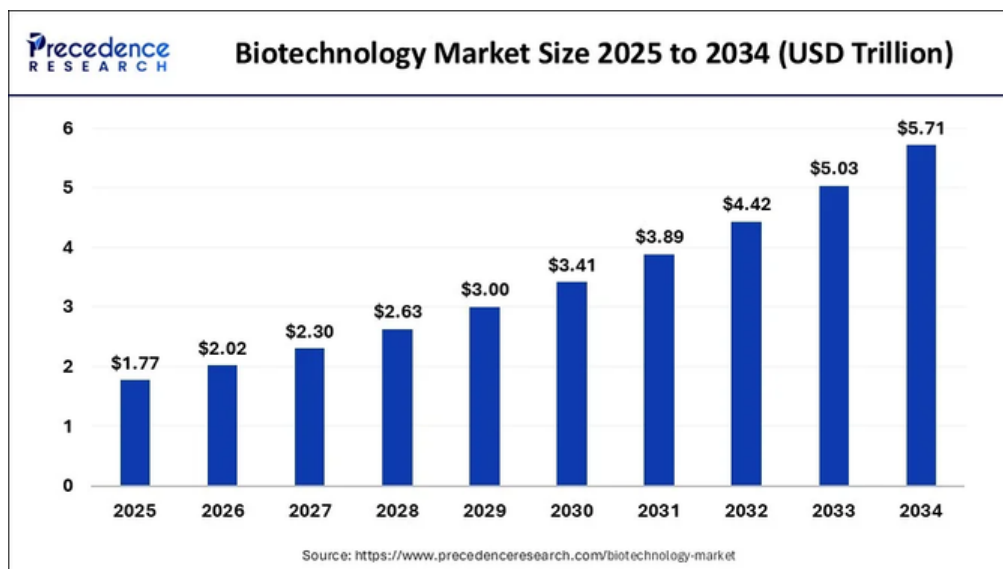
Drug and device development remains extremely capital-intensive and time-consuming, with multi-year clinical trials and strict regulatory review, making failure very expensive and favoring well-funded players.

Pricing & Reimbursement Pressure

Governments and payers are pushing back on high prices and demanding clear cost-effectiveness, while reimbursement delays and market volatility can limit adoption and squeeze margins even for strong products.

Manufacturing & Talent Constraints

Scaling biologics and advanced therapies requires specialized, expensive manufacturing facilities and supply chains, while intense competition for skilled biotech scientists and engineers drives up labor costs and can delay development timelines.



Product Overview

Overall Key Trends:

- Personalization
- AI Integration
- Regulation and New Policy Adaption

The biotechnology industry is undergoing rapid transformation due to the surge of artificial intelligence and breakthroughs in genetic engineering. Such technologies are allowing healthcare solutions to become more precise given to data-integrated systems.

The most prominent area is digital therapeutics as this industry combines medical devices and pharmaceutical interventions with software to manage chronic disease. These platforms help physicians extend care past privileged populations with easy access and to underserved populations through remote monitoring.

In cardiovascular health, scientists are focusing on how to improve imaging and diagnostics using AI to support early detection and prevention methods. Similar to this, gene editing programs are moving past the one fits all model through personalized treatments for individuals. Even though these therapies have high developmental costs, they will deliver long-term savings by removing lifelong management.

In the field of drug discovery, CRISPR and AlphaFold are quickly allowing for the identification of cancer and rare diseases. Tools like these are boosting developmental efficiency and investor confidence. Meanwhile, telemedicine platforms connected to wearables are enabling continuous data collection and supporting the consumerization of health.



Supply Chain

Upstream: Cloud infrastructure by AWS and Google, cybersecurity, CRISPR licenses, gene therapy delivery vectors, sequencing machines, device components, AI tools

Manufacturing: Contracting manufactures to produce products or run clinical trials that meet FDA and regulatory body standards

Downstream: Products reach patients through employers, clinics, doctors, hospitals, and government programs like Medicare

Customers

End Consumer: Patients

Primary Market: Patients through hospitals and private practices

Secondary Market: Labs, Government bulk buyers, Employers, Public Health Programs

Industry Comps

IPO Examples:

- *Hinge Health*: Reached \$3B IPO valuation driven by results in MSK pain reduction, demonstrating that measurable clinical improvement drives valuation in the industry today.
- *Acelyrin*: \$3B IPO valuation after strong clinical results within its immunology pipeline portfolio, also proving how important late-stage clinical success is for biopharma valuation.

SUCCESSFUL EXITS (M&A):

- *One Medical*: Primary Care company acquired by Amazon for \$3.9B, showing that care companies with mature cash flows and delivery models can greatly drive value.
- *Thrive Earlier Detection*: Acquired by Exact Sciences for \$2.15B in an attempt to acquire liquid biopsy data and new technology with high-growth potential.

Multi-Round Success Stories:

- *Hims & Hers*: Multiple rounds of funding that led to a SPAC of \$1.6B and a multi-billion dollar public company today, backed by a proven model of direct-to-consumer virtual care and pharmacy system.
- *Omada Health*: Has currently raised through Series E funding, having a clinical success background in diabetes and chronic care, once again proving that ongoing funding is typically a byproduct of clinical results and late-stage success.

Industry Comps

Acquisition Examples:

- *Flatiron Health*: Oncology data platform acquired by Roche for \$1.9B, which paid a premium for Flatiron's validated data infrastructure and recurring SaaS-like revenue base.
- *Grail*: Liquid Biopsy Diagnostics company that was acquired by Illumina for \$8B, with a valuation supported by compelling multi-cancer early detection results and a strong pipeline of clinical studies.

Failure/Distress Stories:

- *Zymergen*: IPO'd at \$3B, collapsed due to unproven core technology, weak product pipeline, and unrealistic revenue projections, and was sold for \$300M and eventually went bankrupt
- *Theranos*: Blood diagnostics company that was valued at \$9B but collapsed after extensive fraud and inability to validate its blood-testing technology hurt the company and drove it to bankruptcy as well.

Takeaways/In-Summary:

Early-stage biotech and digital health companies still raise sizable rounds, but valuations now hinge heavily on late-stage clinical data or proven commercial traction. Investors and acquirers increasingly prioritize Phase II/III results and real-world performance, making strong late-stage validation essential for avoiding down-rounds, discounted exits, or complete failure.



Industry Overview: EnsiliTech develops silica-based stabilization that allows biologics to remain viable without refrigerated transport. The biopharmaceutical cold-chain sector is experiencing sustained demand growth as biologics, mRNA vaccines, and gene therapies become a larger share of global drug pipelines: they each require tightly controlled temperatures to maintain efficacy. The cold-chain logistics industry Ensilitech operates within is growing at 11–13% CAGR as UK and broader Europe underscores a regulatory environment that prioritises quality assurance across supply networks, where cold-chain failures have historically caused critical delays and a loss of 26–30 billion annually. As customers including vaccine manufacturers, logistics providers, and gene therapy developers seek to reduce dependence on fragile refrigerated systems, EnsiliTech aligns directly with this rising demand.

Market Growth & CAGR: Pharmaceuticals must stay within specific temperature-controlled ranges when transported through the supply chain. The global biopharma cold-chain logistics market is valued at \$18.23 billion in 2024 and is expected to reach \$51.58 billion by 2034, indicating a 10.98% CAGR. The U.S. cold-chain packaging market is valued at \$1.23 billion in 2024 and projected to grow 14.6% annually from 2025–2030. Additionally, over 50% of new drug approvals in the UK and EU are now temperature-sensitive biologics, meaning cold-chain spending is rising not only from volume growth but from a structural shift in modern therapeutics. Given that EnsiliTech operates across these regions, it is reasonable to situate its surrounding industry CAGR at 12–13%, positioning its technology to capture various rapidly expanding markets.

Product Overview: Ensilication is a technology used for preventing vaccine doses from being wasted by thermally protecting biomolecules. This product establishes a silica network to prevent degradation from extreme cold and hot temperatures, -80° C and 50°C serving as the bounds respectively in order to improve shelf-life. Unlike other stabilization technologies, ensilication does not rely on lyophilization and is designed for pre-existing and new biopharmaceutical formulations ([Ensili Tech](#)).

Supply Chain & Customers: Ensilitech's technology disrupts the current supply chain by reducing cold chain dependency on biopharmaceutical supply. Their primary customers are large biopharmaceutical companies like Pfizer, Merck, Moderna, and Astrogenca, who produce vaccines and gene therapies that require temperature sensing devices. Ensilitech uses specialized raw material suppliers for its key component, the silica based encapsulation process that couples closely with pharmaceutical manufacturers to implement this technology. Distribution takes place by working with logistics providers whose companies want to optimize temperature controlled transport and lower overhead ([Fleming](#)).