



# Tomorrow never dies

**Returns from biotechnology are starting to reflect the sector's spectacular scientific ambition.**

Update  
05 December 2025

The late Steve Jobs observed, "The people who are crazy enough to think they can change the world are the ones who do." Biotech has certainly epitomised that pioneering spirit, from obesity drugs revolutionising global healthcare to gene therapy tackling previously untreatable diseases.

The returns for investors, however, haven't always quite matched the scientific ambition, with the heady euphoria of the pandemic flipping into a protracted bear market as rising interest rates and risk-off sentiment weighed on valuations. Last year saw tentative signs of recovery, fuelled by optimism around a pro-business Trump administration, but subsequently stalled on FDA changes and tariff speculation.

Despite these headwinds, the underlying investment case for biotech has remained as strong as ever, thanks to the powerful structural drivers of ageing populations, a rising burden of chronic diseases and the drive to improve healthcare efficiency. A Darwinian sector shake-out of speculative companies has also left behind a higher-quality universe in its wake.

And it seems that valuations are finally beginning to reflect fundamentals over short-term noise: the NASDAQ Biotechnology Index (NBI) has chalked up an impressive near-40% gain in the last six months, comfortably eclipsing the 16% for the S&P 500. The pressing question is whether this marks the long-awaited start of a sustained recovery for biotech.

## Turning a corner

Political risk hasn't disappeared but greater clarity has helped to allay fears. The US administration's recently-announced 100% tariff on branded pharmaceutical imports appears primarily aimed at high-volume, mass-market drugs (rather than the smaller-scale, specialist treatments typical of biotech) and companies investing in US manufacturing facilities are exempt. The UK has also recently secured a zero tariff deal on pharmaceuticals in return for a 25% price increase on new US drugs.

It's a similar story for the Most Favoured Nation pricing order which benchmarks US drug prices against the lowest price paid in developed countries. To date, this policy has focused on mainstream blockbusters such as Novo Nordisk's Ozempic and Eli Lilly's Zepbound rather than niche biotech treatments.

Regulatory concerns are also easing. While political appointments have been a mixed bag (particularly for vaccine developers), fears of an anti-industry stance are fading. The FDA continues to

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approve drugs at pace, and initiatives such as the National Priority Review Voucher pilot aims to cut approval times from a year to as little as one or two months, potentially adding an additional year of revenue before patent expiry.

Meanwhile, M&A is back with a bang. Big pharma remains heavily reliant on blockbuster drugs, with revenue falling by as much as 80% in the first year after patent expiry. With a projected \$240 billion revenue gap by 2030, pharmaceutical companies have amassed a \$1.3 trillion war chest to replenish their pipelines.

Biotech companies now account for 70% of clinical trials, offering nimble, capital-light R&D that can be scaled through big pharma's global sales infrastructure. M&A is likely to remain a key driver for returns, as acquiring late-stage assets continues to be the most efficient route to near-term revenue growth for major pharma players.

The macro backdrop is also looking more supportive, with the NBI historically trading inversely with interest rates. Sentiment seems to be following suit, with the long-term growth potential of biotech attracting renewed interest from investors looking



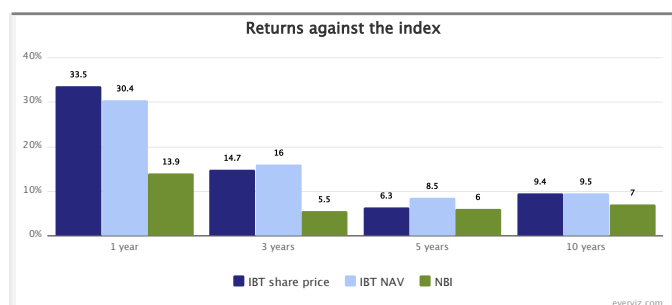
to diversify beyond the concentrated leadership in the technology sector.

In summary, the current rally reflects more than just a short-term rebound from depressed valuations, with genuine momentum driven by innovation, clearer regulatory policy and a healthy M&A pipeline.

## Handle with care

In a sector where the science drives the value, expertise and selectivity remain paramount. **International Biotechnology Trust (IBT)** has exemplified this, navigating multiple market cycles to outperform the NBI over one, three, five and ten years, as shown below.

**Fig.1: IBT Has A Track Record Of Superior Performance**



Source: Schroders factsheet (as at 31/10/2025), based on NASDAQ Biotechnology TR in GBP  
*Past performance is not a reliable indicator of future results*

Asset allocation has played a central role in this success. Managers Ailsa Craig and Marek Poszepczynski's investment process includes a top-down approach, tailoring portfolio exposure to each phase of the biotech cycle. This was demonstrated by their timely shift into more profitable large-caps, such as Gilead, in 2021 ahead of valuations suffering a sharp correction.

As large-cap valuations extended, the managers have pivoted towards clinically de-risked small- and mid-cap companies, building a materially higher weighting than the index. These firms have products which have already launched, are in late-stage trials or under FDA review, offering exposure to the highest-growth phase of the development cycle.

While this segment often commands the highest acquisition premiums, these companies can also generate strong returns by remaining independent thanks to their clear path to commercialisation.

Identifying M&A targets is not a formal objective but IBT has an impressive hit rate on this front, chalking up 34 acquisitions since 2020 and nine this year alone. Notable

deals include Novartis's \$12 billion purchase of Avidity at a 46% premium and Johnson & Johnson's \$15 billion acquisition of Intra-Cellular at a 40% uplift.

The managers have also reduced oncology exposure in recent years: despite attracting significant funding, finding likely winners is difficult in a highly-crowded market, with IBT instead opting for later-stage companies with clearer paths to market.

Around a third of the portfolio is currently allocated to rare diseases, which combines a high unmet medical need, lower clinical trial costs and attractive commercial dynamics. Rare diseases are a key area of focus for the FDA, with regulatory frameworks such as the Orphan Drug Act providing meaningful incentives in the way of market exclusivity and accelerated approval pathways.

And finally, IBT has also put its closed-end structure to good use, increasing gearing to around 20% after Liberation Day to take advantage of the dip in sector valuations and paying its investors a dividend of 4% of NAV.

## In rude health

With cutting-edge innovation, strong secular tailwinds and greater regulatory clarity, biotech's recovery looks increasingly well-supported.

For active investors, the sector may be entering a period where scientific breakthroughs and shareholder returns finally move in tandem, a fitting testament to those visionary mindsets that seek to change the world.

All data as at 03/12/2025 unless stated otherwise.

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