Building a healthier future



Cell & Gene Therapy

Bayer's strategy

We are on the cusp of a new wave of innovation in healthcare

For decades, many diseases have been labeled as "intractable", meaning medicine could not provide an answer for patients beyond addressing their symptoms to a certain extent. By targeting diseases at the cellular or genetic level, we have the potential to shift from treating symptoms to stopping or even reversing diseases.



Heart failure, neurodegenerative diseases like **Parkinson's,** or genetic diseases like **Huntington's** are just a few examples of where **cell and gene therapies may make a difference** in the lives of patients.

A pharmaceutical (r)evolution: cell therapy and gene therapy

Bayer's journey began over 125 years ago with small molecule pharmaceuticals like Aspirin[™]. Large molecules and targeted therapies like Eylea[™] for age-related macular degeneration came next. These treatments opened new doors and improved the lives of millions of patients.

Researchers are already working on the next wave of innovation: cell therapy and gene therapy. These could be a breakthrough for patients living with diseases for which limited or no options are available today.

Three waves of innovation in the pharmaceuticals space: 3 Our next step Cell & Gene Therapy

Our biologics

Large molecules & targeted therapies

1) Our stronghold Small molecules MOSTLY SYMPTOMATIC

TREATMENT, UNDERLYING DISEASE NOT REVERSED

POTENTIAL

DISEASE

TO REVERSE



Between 2019 and 2023, Bayer has invested more than € 3.5 billion in the build-up of a cell and gene therapy platform. This includes the acquisition of BlueRock and AskBio, as well as a collaboration with Mammoth Biosciences.

Pioneering scientific advancement

We believe that a healthier future can be achieved by leveraging science, passion and collaboration. We are combining our internal expertise with external collaborations and acquisitions, jointly advancing the potential of the next wave of innovation in the pharmaceuticals space for the ultimate benefit of patients.





Did you know that there are already multiple approved **cell therapies and gene therapies** for the treatment of various types of cancer, certain blood disorders like hemophilia A and B, as well as other rare diseases such as spinal muscular atrophy?^{1,2}

We are shifting treatment paradigms...

First-generation cell and gene therapies are already impacting the lives of patients around the world. The future is enormously exciting – cell therapy and gene therapy could lead to far more targeted, personalized treatments. It could also mean finding a way to finally defeat diseases that we once thought untreatable.



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... across multiple areas



- 1 Source: American Society of Gene and Cell Therapy, and Citeline. Gene, Cell, & RNA Therapy Landscape: Q3 2022 Quarterly Data Report, https://asgct.org/global/documents/asgct-citeline-q3-2022-report.aspx. Accessed Dec. 2022.
- 2 Source: OCT) JX Yu et al., Nature vol 19, September 2020; AAV) Cortellis; GE) Cortellis; iPSC) JY Kim et al., Stem cell reviews and reports, Springer, September 2021