

Curriculum Vitae Professor Dr Patrick A. Baeuerle

Name: Patrick Alexander Baeuerle

Date of birth: 24 November 1957

Research Priorities: Development of cancer drugs, tumour immunology, cell biology, protein engineering, therapeutic antibodies

Patrick Baeuerle is a biologist and carries out research in the field of applied immunology. His focus is the development of immune-based cancer drugs. He has worked on the development of novel cancer therapies based on antibodies, fusion proteins, cytokines and gene-modified immune cells for over 20 years. This work has led to the worldwide approval of the leukaemia drug Blincyto and the foundation of so far seven biotech companies in the USA and one in Germany, currently with more than ten clinical programmes testing novel therapies for cancer patients.

Academic and Professional Career

since 2015	Founder of biotech companies Harpoon, TCR^2 , $iOmx$, Maverick, Cullinan, Werewolf and Aktis
since 2015	Executive Partner, MPM Capital LLC, Cambridge, USA
since 2016	Chief Scientific Officer (CSO, Biologics), Cullinan Oncology Inc., Cambridge, USA
2012 - 2015	Vice President of Research, Biotech Company AMGEN Inc., Munich, Germany and Thousand Oaks, USA
since 2000	Professor h.c. Immunology, Faculty of Medicine, Ludwig Maximilian University of Munich (LMU), Munich, Germany
1998 - 2012	Scientific Board of Directors, Micromet GmbH AG and Inc., Munich, Germany and Rockville, USA
1996 - 1998	Director of Drug Discovery, Tularik Inc., South San Francisco, USA

1992 - 1996	Ordinary Professor of Biochemistry and Molecular Biology, Faculty of Medicine, Albert Ludwig University of Freiburg, Freiburg im Breisgau, Germany
1991	Habilitation in Biochemistry, LMU Munich, Munich, Germany
1989 - 1993	Principal Investigator, Research Group, Gene Center Martinsried, LMU Munich, Munich, Germany
1987 - 1989	Postdoctoral Fellow, Whitehead Institute, Cambridge, USA
1982 - 1987	Graduate and Doctoral thesis, Max Planck Institute (MPI), Martinsried and European Molecular Biology Laboratory (EMBL), Heidelberg, Germany
1978 - 1982	Degree in Biology, University of Konstanz, Konstanz, Germany

Functions in Scientific Societies and Committees

since 2021	Scientific Advisory Board, BIOSS Centre for Biological Signalling Studies, Albert Ludwig University of Freiburg, Freiburg im Breisgau, Germany
2014 - 2015	Member, Scientific Panel of Health, European Commission
2009 - 2012	Member, Scientific Review Board, Qatar National Research Fund, Doha, Qatar
2006 - 2007	Member, Health Science Roadmap Committee for Cancer, Federal Ministry of Education and Research (BMBF), Germany
1999 - 2004	Member, Scientific Advisory Board, Istituto Scientifica San Raffaele University Hospital, Milan, Italy
1993 - 1995	Member, Scientific Advisory Board, Paul Ehrlich Center, Hadassah Medical School, University of Jerusalem, Jerusalem, Israel

Project Coordination, Membership in Collaborative Research Projects

1989 - 1996	Member, Collaborative Research Centre, (SFB) 217, 190, 364, 388 and 505, German
	Research Foundation (DFG), Germany
1989 - 1996	Board of Directors, SFB 364, 388 and 505, DFG, Germany

Honours and Awarded Memberships

since 2021	Member, German National Academy of Sciences Leopoldina, Germany
2022	Hirsch index ranking of 137 with over 82,000 citations
2019	Lennart Philipson Award for Contributions to Immunotherapy for Cancer, EMBL,
	Heidelberg, Germany

2019	Entrepreneur of the Year, Boston, USA
2019	Among the 0.01% most cited scientists worldwide, according to study by loannides et al. (Stanford University), USA $$
2002	Member, European Molecular Biology Organization (EMBO), Heidelberg, Germany
1991 - 2000	Germany's most cited biomedical scientist, according to a study of the Institute for Scientific Information (ISI), Philadelphia, USA
1993	First prize winner, Prix Européen de l'Avenir
1991 - 1993	Karl Winnacker Award and Scholarship, Hoechst AG, Frankfurt am Main, Germany
1987 - 1989	Fellowship Abroad, DFG, Germany
1983 - 1987	Scholarship, Max Planck Society, Munich, Germany and EMBL, Heidelberg, Germany

Research priorities

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Blincyto is known as a bispecific antibody, which briefly links the T immune cells of cancer patients with cancer cells and thus initiates their elimination. A common feature of all cancer therapies developed by Patrick Baeuerle is that they activate the immune cells of patients against cancer cells, using novel proteins tailor-made for this purpose with recombinant methods.

Baeuerle's initial research focused on relaying pathogenic signals from the cell membrane into the cell nucleus. The inducible transcription factor NF-kappaB plays a key role here. The latter is activated by various pathogenic stimuli in many diseases and initiates the expression of defence genes and immune signalling proteins. His research has led to the discovery of new subunits of NF-kappaB (I-kappaB and p65/RelA) and has explained the canonical activation mechanism of NF-kappaB, which is based on the degradation of the inhibitory subunit I-kappaB by the ubiquitin system.

Since 1998 Patrick Baeuerle has developed novel cancer drugs, primarily in biotech companies in the USA. As Head of Research at Micromet, he was responsible for developing the bispecific antibody blinatumomab, which was approved in the USA in 2014 and in Europe in 2015 as the leukaemia treatment "Blincyto". Blinatumomab is not only the first, but until recently was also the only approved antibody of its kind able to eliminate the patient's cancer cells by using their T immune cells. Numerous other cancer drugs that Baeuerle has helped to develop are currently

undergoing clinical trials. For example, T cells that have been genetically modified with a synthetic TRuC receptor are currently being tested in mesothelioma patients. For patients with other cancers the next generation of bispecific antibodies is being developed which are only activated after penetrating into the tumour. Preclinical research includes work on novel cytokine fusion proteins that link interleukins-2 and -12. Many cancer therapies have been co-developed by Patrick Baeuerle and promise to have both improved tolerability and increased efficacy.