



Tony Hunter *** Brian Druker *** John Mendelsohn



Bridging science and clinical applications



put forth the possibility TKs could be targeted to inhibit cancer cell



target TKs, leading to successful targeted therapies.

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Developed the first tyrusine kinase-targeted



Developed the first tyrosine kinase-targeted

Discovered tyrosine kinase and opened up a new field of cancer research





His discovery of protein tyrosine phosphorylation and tyrosine kinases (TKs) in 1979 shed light on the signaling mechanisms which cancer cells depend upon and set the foundation for the development of over 29 TK inhibitor cancer drugs.







Brian Druker

Turned a chronic cancer into a treatable disease



① Druker participated in the development and led the successful clinical trial of the miracle drug, Gleevec*, which is a targeted therapy that can effectively treat chronic myelogenous leukemia (CML), turning a fatal cancer into a treatable disease.



2 CML used to be incurable and most patients died within two years. Gleevec® increased survival rate from 50% to 90%.



3 Gleevec* is known as the most successful targeted cancer therapy in the 21st century and was approved for clinical use by the US FDA in 2001

Led the successful development of antibody-based EGFR therapy



- Mendelsohn and his team developed the first clinically approved cancer therapy using an antibody (Erbitux*) to target the TK of an epidermal growth factor receptor (EGFR).
- In liver metastases from colorectal cancer, the 3-year survival rate of patients receiving Erbitux* plus chemotherapy treatment is 41% compared with 18% for those receiving chemotherapy alone. (Ye et al., 168, 189)
- 3 EGFR plays a critical role among many cancers, such as breast, lung, colorectal, ovarian, bladder, esophageal, head and neck cancer, etc. Many other successful targeted therapies now focus on EGFR.

