

Eisai and Merck collaborate for cancer cure

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Eisai and [Merck](#) has announced a clinical trial collaboration. The deal is to evaluate the safety, tolerability and efficacy of Merck's anti-PD-1 therapy, pembrolizumab (marketed in the US under the brand name KEYTRUDA), in combination with Eisai oncology compounds lenvatinib mesylate (a multi-targeting RTK inhibitor marketed in the US under the brand name LENVIMA, "lenvatinib") and eribulin mesylate (a microtubule dynamics inhibitor marketed in nearly 60 countries HALAVEN, "eribulin") in multiple clinical trials.

The planned studies include a multicenter, open-label Phase 1b/2 study of lenvatinib plus pembrolizumab in select solid tumors and an open-label, single-arm, multicenter Phase 1b/2 study to evaluate the efficacy and safety of eribulin in combination with pembrolizumab in metastatic triple-negative breast [cancer](#). Eisai and Merck will establish a Joint Development Committee to oversee clinical development activities.

"This collaboration could be a major step in the direction of developing combination regimens in different types of cancer, potentially maximizing the value of eribulin and lenvatinib," said Dr Kenichi Nomoto, president, oncology product creation unit, Eisai Product Creation Systems. He added, "Together, Eisai and Merck seek to explore combination regimens that have the potential to create synergistic effects between lenvatinib and pembrolizumab as well as between eribulin and pembrolizumab. Our hope is that we will bring treatments to market that make a difference in the lives of people battling cancer."

"Cancer is a complex disease that often requires different approaches to help patients achieve the best possible outcome. The collaboration with Eisai exemplifies Merck's focus on advancing breakthrough science in immuno-oncology. We look forward to evaluating pembrolizumab in combination with eribulin and also with lenvatinib in different tumor types," said Dr Eric Rubin, therapeutic area head, oncology early-stage development, Merck Research Laboratories.

The studies are expected to begin in the second half of 2015. Financial terms of the agreement were not disclosed.