EISAI RECEIVES APPROVAL FOR NEW INDICATION FOR ANTICANCER AGENT HALAVEN® FOR T

For patients with liposarcoma (143 patients), Halaven demonstrated a statistically significant improvement in OS over dacarbazine (Halaven, median OS: 15.6 months vs dacarbazine, median OS: 8.4 months; HR 0.51 [95% CI=0.35-0.75]).

In this study, the most common treatment-emergent adverse events (incidence greater than or equal to 25%) in patients treated with Halaven were fatigue, neutropenia, nausea, alopecia, constipation, peripheral neuropathy, abdominal pain, and pyrexia, which was consistent with the known side-effect profile of Halaven.

3. About Soft Tissue Sarcoma