=perversions 2= Libtayo cemiplimab rwlc Approvedin the U.S. as First and FDAapprovescemiplimab rwlc for metastatic.

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Oct 8 2025 TheFDAevaluatedLibtayounder Priority Review which is reserved for medicines that represent potentially significant improvements in efficacy or safety in thetreatmentof serious conditions

An additional regulatory application is also under review in the European Union with a decision expected by the first half of 2026 The recommendedcemiplimab rwlc dose and schedule is 350 mg as an intravenous infusion over 30 minutes every 3 weeks

View full prescribing information forLIBTAYO LIBTAYO cemiplimab rwlc is a prescription medicine indicated for thetreatmentof adult patients with advanced NSCLC CSCC BCC and as an adjuvanttreatmentfor high risk CSCC

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The Food and Drug Administration FDA hasapprovedLibtayo cemiplimab rwlc for the adjuvanttreatmentof adult patients with cutaneous squamous cell carcinoma CSCC at high risk of recurrence after surgery and radiation On October 8 2025 the Food and Drug Administrationapprovedcemiplimab rwlc Libtayo Regeneron Pharmaceuticals Inc

for the adjuvanttreatmentof adults with cutaneous squamous cell carcinoma CSCC at high risk of recurrence after surgery and radiation

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