Dear NCCN Guidelines Panel:

On behalf of Genentech, Inc., please find data from the GO30140 study enclosed for your review. This study evaluated the combination of Tecentriq\textsuperscript{\textregistered} (atezolizumab) and Avastin\textsuperscript{\textregistered} (bevacizumab) for the treatment of advanced or metastatic hepatocellular carcinoma (HCC).

**Request:**
Please consider the inclusion of Tecentriq and Avastin as an initial first-line systemic treatment for patients with advanced or metastatic HCC.

**Rationale:**
Patients with advanced HCC continue to have limited treatment options. Data from the GO30140 study demonstrate activity of Tecentriq and Avastin in this patient population.\textsuperscript{1}

- In patients with unresectable or advanced HCC (n=73), treatment with Tecentriq plus Avastin resulted in an ORR of 32\% per INV-assessed RECIST and 34\% per IRF-assessed HCC mRECIST. Responses were observed in all assessed patient subgroups, including etiology, region, baseline alpha-fetoprotein levels and tumor burden.
- Responses were durable with 52\% of patients having responses lasting ≥ 6 months and 26\% lasting for ≥ 12 months.
- The safety profile of Tecentriq and Avastin was consistent with previously reported safety risks of the individual medicines.

**FDA Clearance:**
The U.S. Food and Drug Administration granted Breakthrough Therapy Designation for Tecentriq in combination with Avastin as first-line treatment for patients with advanced or metastatic HCC.\textsuperscript{2}

- Tecentriq is not FDA-approved for the treatment of HCC. Please refer to the product prescribing information for the full FDA-approved indications and safety information of Tecentriq and Avastin, available at:

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Thank you for your consideration. If you have any questions, please contact us at the phone number and email provided above.

Respectfully submitted,
Patrice Esser, PharmD, MPH
References:
