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**Janssen Submits Application to U.S. FDA Seeking Approval of Amivantamab for the Treatment of Patients with Metastatic Non-Small Cell Lung Cancer with EGFR Exon 20 Insertion Mutations**

As a trusted partner, we are excited to share that, today, Janssen announced the submission of a Biologics License Application (BLA) to the U.S. Food and Drug Administration (FDA) for amivantamab, an investigational, fully-human epidermal growth factor receptor (EGFR) and mesenchymal epithelial transition factor (MET) bispecific antibody with immune cell-directing activity that targets tumors with activating and resistance EGFR and MET mutations and amplifications.

The application marks the first-ever regulatory submission for the treatment of patients with NSCLC with EGFR exon 20 insertion mutations. Therefore, Janssen hopes that the filing of amivantamab can continue to progress treatment options for those suffering from one of the most prevalent forms of cancer.

**To learn more about the amivantamab, please view the press release issued by Janssen [here](#).**