

# Information Regarding Clinical Trial

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| <b>Title</b>                       | Part 1 Run In for first 15 patients<br><br>“A Phase II, multicenter, randomized, two-arm study of Capmatinib (INC280, an oral MET inhibitor) combination therapy vs docetaxel in pretreated adult patients with EGFR wild-type, ALK rearrangement negative locally advanced/metastatic NSCLC”  |
| <b>Protocol Number</b>             | CINC280D2201   |
| <b>Sponsor</b>                     | Novartis   |
| <b>Investigator</b>                | J. Thaddeus Beck, MD<br>3232 N. North Hills Blvd.<br>Fayetteville, AR 72703  |
| <b>Study Related Phone Numbers</b> | 479-587-1700 (24 hours)  |
| <b>Key Points for Study</b>        | Part 1: Run In for first 15 patients<br>Must be stage IIIB or IV, EGFR and ALK neg, prior platinum- based chemotherapy and PD L1 inhibitor therapy with PD L1 to be last drug given, must have at least 1 measurable lesion (Recist 1.1). Part 1 will treat with Capmatinib and Spartalizumab combination. Review will take place after all subjects have at least 24 wks of follow up before start of Part 2. |