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A: First of all, you should download the standalone version of the software, from PTC website, and install it. You should also uninstall the one that you have installed, and delete the folder from which you unzipped the standalone version. Then you can install the standalone version using the installer from the downloaded standalone. If you want to install the standalone version using an already installed version, you can do it by running the standalone installer under the version that you have already installed. Also don't forget that you have to install all the required software beforehand, and they have to be compatible with the standalone version. I'm not sure about this one, but you can try this: Evaluation of transactivation of the androgen receptor in prostate cancer cells. To evaluate the transactivation activity of the androgen receptor (AR) in clinical prostate cancer specimens. In contrast to the limited availability of normal human prostate tissue for biochemical and functional studies, a large series of tumor specimens (multiple sites) can be obtained by prostate biopsy. Several AR target genes, including prostate-specific antigen and TMPRSS2-ERG, have been identified as potential prognostic markers in prostate cancer. However, no study has systematically evaluated the transactivation of the AR. The AR transactivation activity of 58 prostate adenocarcinoma specimens (41 nodal metastases and 17 bone metastases) was assessed by using a transient transfection assay. The AR transactivating function of four established prostate cancer cell lines (two basal cell lines, LNCaP and C4-2B, and two luminal cell lines, LAPC-4 and PC3) was also evaluated. High-level expression of TMPRSS2-ERG and prostatespecific antigen was significantly correlated with AR transactivation activity in prostate cancer samples. The prognostic value of AR transactivation was different in metastatic vs nonmetastatic cancers: high-level expression of the AR target genes and the AR transactivating activity were significant predictors of nonmetastatic cancers (pathologic stage pT2 or lower vs pT3-4, hazard ratio, 5.0; and P=0.0001), but were not significant predictors of metastatic cancers (nodal versus bone metastases, hazard ratio, 2.1; P=

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