



Figure S2. Western blot images of PC12 cells pre-treated with the AhR antagonists, (A) α -naphthoflavone (ANF; 1 μ M) and (B) CH223191 (1 μ M), for 1 h and treated for five consecutive days with NGF (25 ng/mL) and ILA, IPA, or Trp (100 nM). Non-pre-treated PC12 cells served as a null control. AhR protein (95 kDa) in PC12 cells was detected by Western blot analysis using a monoclonal antibody specific for AhR. The corresponding β -actin blot served as a loading control. Lane 1 & 7: non-treated control (NTC); Lane 2 & 8: Nerve growth factor (NGF)-treated control; Lane 3 & 9: Cells treated with NGF and papaverine hydrochloride as reference control (RC); Lane 4 & 10: Cells treated with NGF and indole-3-lactic acid (ILA); Lane 5 & 11: Cells treated with substance A; Lane 6 & 12: Cells treated with tryptophan (Trp).