COMMUNICATION SIGNALS PRODUCED BY TUMOR-INDUCING AND TARTRATE-CATABOLIC PLASMIDS OF AGROBACTERIUM VITIS. Nathanial Lowe and Michael A. Savka*. Biological Sciences, Rochester Institute of Technology. massbi@rit.edu

Agrobacterium vitis, a Gram-negative bacterium, causes a disease known as crown gall on grape plants. Crown gall is characterized by the development of plant cell tumors after infection with tumorigenic strains of A. vitis. Tumorigenic A. vitis strains contain a large extrachromosomal DNA element known as tumor-inducing plasmid (pTi) and may also contain a tartrate-catabolic plasmid (pTr). The Ti and Tr plasmids may allow A. vitis strains to interact in alternative lifestyles with host plants and competing microorganisms. Because bacterial signals called Nacyl-homoserine lactones (acyl-HSLs) are important in the ecology of plant-associated bacteria, we determined the acyl-HSLs produced by Ti and Tr plasmids from nopaline-, octopine- and vitopine-type strains of A. vitis. An Agrobacterium tumefaciens derivative, UBAPF2, which alone does not produce acyl-HSLs, was conjugated with wild type A. vitis strains to assemble a collection of UBAPF2 strains carrying a Ti or Tr plasmid. This collection allowed for the identification of acyl-HSL signal(s) produced from pTi or pTr plasmids. Transconjugates (UBAPF2 containing a single Ti or Tr element) were tested for signal production using an acyl-HSL-specific A. tumefaciens biosensor strain NTL4 in diffusion well assays. To further characterize acyl-HSLs, signal preparations were separated by thin layer chromatography (TLC) and detected using biosensor strain NTL4. Fourteen of the sixteen transconjugates produced a short-chain acyl-HSL signal identified as 3-oxo-C8-HSL by mobility and detection characteristics. Transconjugates containing pTrAB3 and pTrTm4 did not produce a detectable acyl-HSL signal. UBAPF2 transconjugates harboring pTiNi1, pTrAB4, pTrRr4 or pTrAT6 produced the characteristic short-chain signal and, interestingly, an additional long-chain acyl-HSL signal absent in TLC signal profiles of corresponding wild type strains.