

MILLIMETER/SUBMILLIMETER SPECTRUM OF THE CCP ($X^2\Pi_r$) RADICAL

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The pure rotational spectrum of the CCP radical ($X^2\Pi_r$) has been measured using millimeter-wave direct absorption techniques. This work is the first high-resolution data for this radical. This species was created by the reaction of gas-phase phosphorus and acetylene in the presence of argon carrier gas and an AC glow discharge. Twenty-four rotational transitions were measured in the frequency range of 120 to 413 GHz. Each transition is separated into two spin-orbit components; however, only the $\Omega = 1/2$ component has been observed to date. This spin component is additionally split into lambda-doublets. Below 209 GHz, hyperfine interactions due to the phosphorus nuclear spin of $I = 1/2$ are evident, splitting each lambda-doublet into two lines. The data were fit with a case (c) Hamiltonian, and effective rotational, lambda-doubling, and phosphorus hyperfine constants were determined. Additional searches are being conducted for the $\Omega = 3/2$ component. These data prove that CCP has a linear geometry in its ground state. This structure is similar to other MC_2 species, such as CCS and CCl, in contrast to NaC_2 , MgC_2 , AlC_2 , and SiC_2 . The change in structure from T-shaped to linear is caused by a increase in the covalent bonding interactions between the C_2 moiety and the heteroatom.