Remarks on Parity-Violation in Atomic Physics

- 1) The selection rules we have just derived, viz
- 1 STARK: (Us) = e E · Sd3x[r Ψf Ψ;] s couples, or drives transitions Ψ; > Ψf, only when Ψf is opposite parity to Ψ; (i.e. ΔJ=±1); (1)
- ②ZEEMAN: (Uz) = B. Sd3x [m Ψf Ψi] S couples, or drives transitions Ψi > Ψf, only when Ψf is same parity as Ψi (i.e. ΔI = 0); (2)

 depend on the assumptions: (1) [E & B have definite parities (-) & (+) resp. [so

then It & on have definite parities (-) & (+) resp.], (2) the quantum states \(\); & \(\) \(\) have definite parities [Usu.(-)\(^1\), for state of orbital \(\) momentum \(\) \(\).

- 2) Bound states 4: & 4 in atoms are generated principally by electromagnetic couplings (mainly Contomb) between the proton (nucleus) and its electron(s). Then if parity P is a "good" (conserved) quantum # for EM couplings, P will also be good for the atomic states 4: & 4, and above rules are absolute.
- 3) BUT, suppose the P-conserving EM coupling between proton of electron has a small admixture of a P-nonconserving interaction... this is the case for the modern "electroweak" theory. (Weinberg, Glashow, Salam; 1979) which unifies EM & weak interactions into one (combined) field. Then parity P is almost, but not quite, a good quantum # for a tornic states \$\psi\$, and \$\psi\$ becomes a parity-mixed state. To lowest order in the parity-mixing, we write \[\Psi \times \wideta = \psi + \kappa \in \times \times \text{the nonimal state parity, \$\Psi\$ is opp. parity to \$\Psi\$; (3) \kspace \kappa = \psi + \kappa \in \text{\$\psi}\$ \text{\$\te

Here $K \sim (\text{weak coupling strength})/(\text{EM coupling strength})$ is very small; for a Single proton-single electron interaction: $|K| \sim 10^{-10}$. But $K \neq 0$ violates parity for the State since: $P\widetilde{\Psi} = \pm (\widetilde{\Psi} - 2K\varphi)$, when Ψ has (\pm) parity.

4) Parity-violation in atoms can be searched for as follows. The wavefunction combinations which occur in the Stark & Zeeman matrix elements above are

$$\left[\begin{array}{c} \Psi_f^* \, \Psi_i \, \rightarrow \, \widetilde{\Psi}_f^* \, \widetilde{\Psi}_i \, = \, \Psi_f^* \Psi_i \, + \, \text{Ki} \big[\Psi_f^* \, \varphi_i \big] \, + \, \text{K}_f^* \big[\varphi_f^* \, \Psi_i \big] \, , \, \, \text{to} \, \, \Theta(\kappa) \, . \quad \ \, (4) \\ \text{these terms have parity opposite to} \, \, \Psi_f^* \, \Psi_i \, \end{array} \right]$$

The Stark matrix element (w.r.t. parity-mixed V's) picks up new terms ...

$$\rightarrow \langle U_s \rangle = e \mathbb{E} \cdot \int d^3x \left[r \psi_f^* \psi_i \right] + \kappa_i e \mathbb{E} \cdot \int d^3x \left[r \psi_f^* \psi_i \right] + \kappa_f^* e \mathbb{E} \cdot \int d^3x \left[r \psi_f^* \psi_i \right], \tag{5}$$

and likewise the Zeeman matrix element $\langle U_z \rangle$ acquires terms in K. Now, consider driving the transition $\psi_i \rightarrow \psi_f$ by an electric field E, when the States $i \notin f$ have the <u>Same</u> "parity". For parity-puve states, Such a transition is <u>forbidden</u> by the selectron rule in Eq.(1); Same parity i.e. the matrix element $eE \cdot \int d^3x \left[\pi \psi_f^* \psi_i^* \right] = 0$. But for the parity parity-mixed states, the terms in K in Eq.(5) are non-zero, so we have

$$\rightarrow \langle U_s \rangle = e \mathbb{E} \cdot \left\{ \kappa_i \int d^3x \left[r \Psi_f^* \varphi_i \right] + \kappa_f^* \int d^3x \left[r \varphi_f^* \Psi_i \right] \right\}, \qquad (6)$$

as a transition amplitude for an otherwise forbidden transition $i \rightarrow f^{3/4} \Delta J = 0$. So, if we see a <u>violation</u> of the selection rule $\Delta J = \pm 1$ for E-field driven transitions (i.e. we detect a $\Delta J = 0$ transition driven by E), we can blame it on electroweak parity-nonconserving (PNC) effects. Likewise, a violation of $\Delta J = 0$ for B-field driven transitions implies PNC for an atom.

5) Atomic φ expts have in fact shown the existence of forbidden transitions and PNC effects in atoms. They are ferociously difficult, because the measurable violation rates go as |⟨U⟩|² α |κ|² ⟨⟨< 1. See R.T.Robiscoe & W. L. Williams, Nucl. Instr. Methods 197, 567 (June 1982).