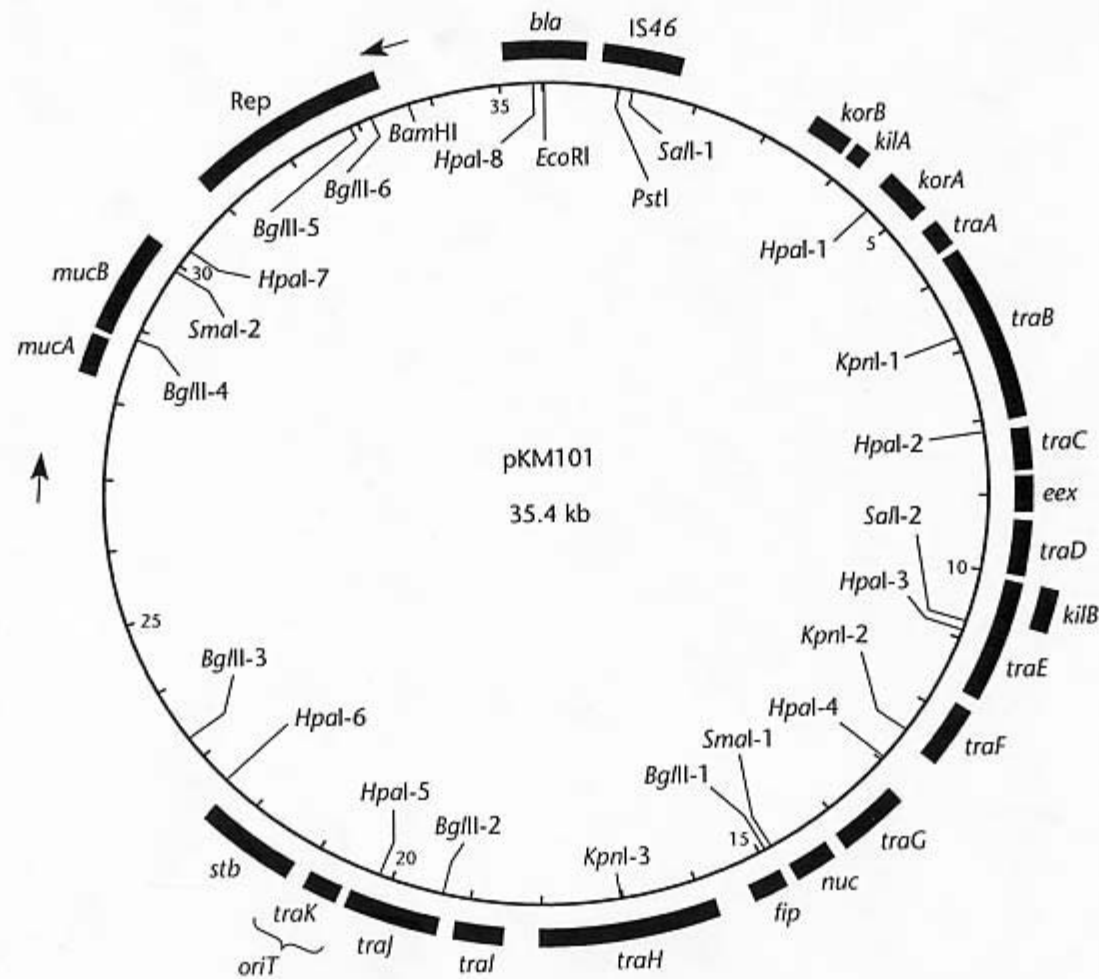
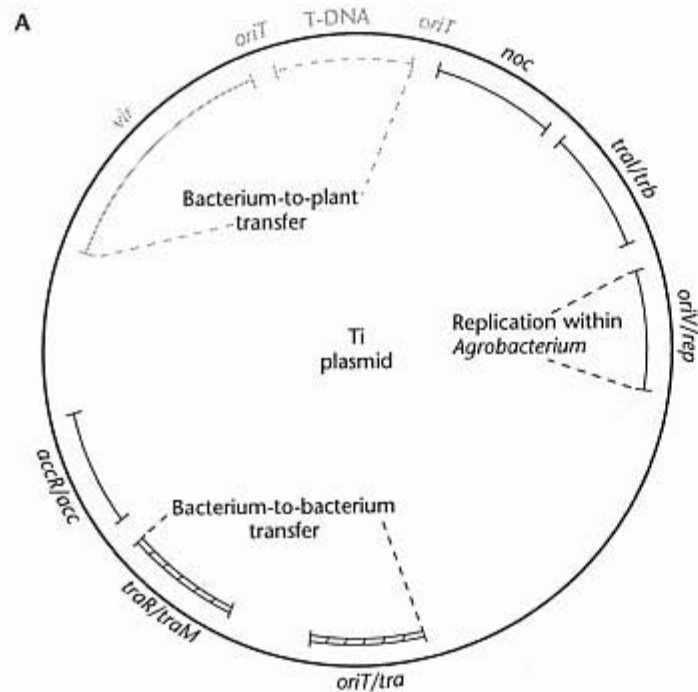


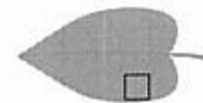
Figure 5.1



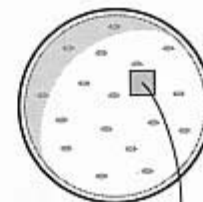
Box 5.1

**B**

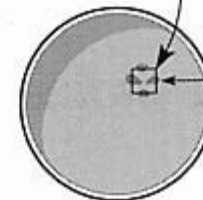
Cut out a piece of the leaf



Float in plate containing an *Agrobacterium* strain with engineered Ti plasmid

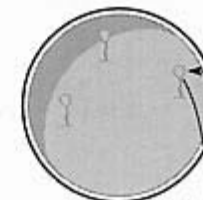


Incubate on plate containing plant regeneration medium



Regenerating plants

Excise germinating shoots and transplant to plates containing kanamycin



Kanamycin-resistant shoots

Transgenic plant



Box 5.1 cont.

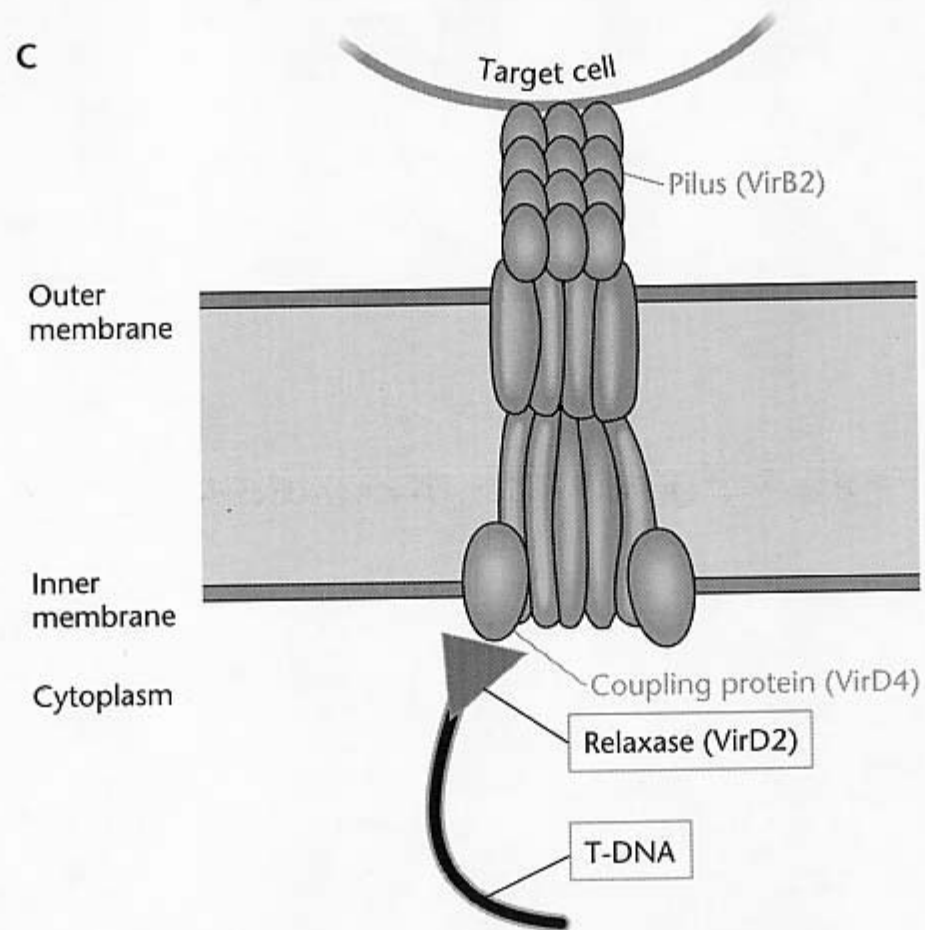
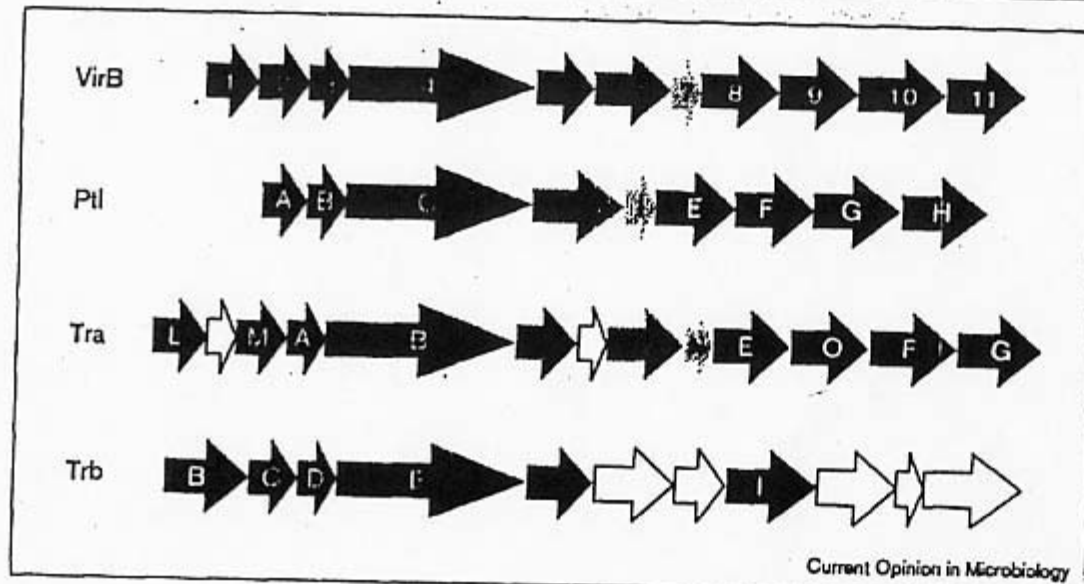


Figure 1



Alignment of genes encoding type IV transporter systems. These systems include the VirB system of *A. tumefaciens*, the Ptl system of *B. pertussis*, the Tra system of the IncN plasmid pKM101 and the Trb genes of the Tra2 region of the IncP plasmid RP4. Arrows of the same color represent homologous genes. Unfilled arrows represent genes that have no homologue.

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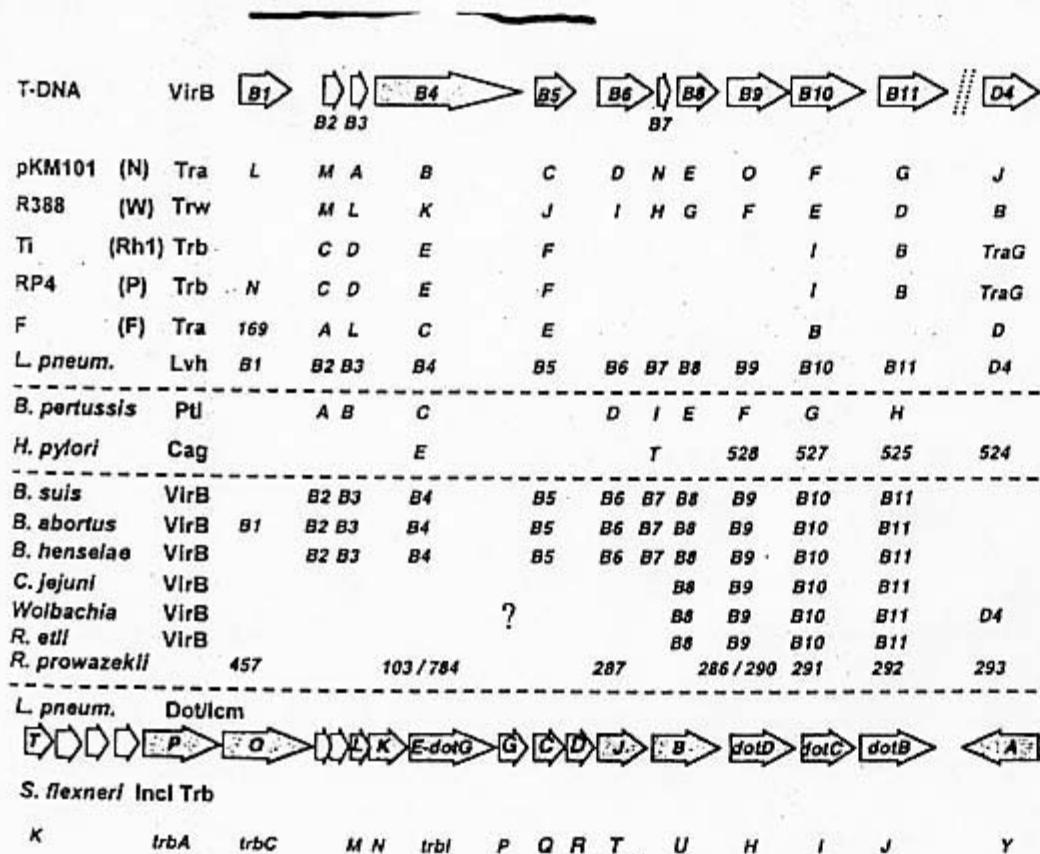


Fig. 2. The type IV systems known or postulated to translocate macromolecular substrates intercellularly. The *A. tumefaciens* *virB* gene products shown across the top assemble as the T-DNA transfer system. The next three groups of type IV systems, separated by dashed lines, are composed of homologues of some or all of the VirB proteins. The top group corresponds to systems shown to transfer DNA between bacteria, the *B. pertussis* and *H. pylori* systems deliver known substrates (PT and CagA respectively) to mammalian cells, and the third group corresponds to systems whose substrates are presently unknown but are postulated to be effector proteins. The symbol (?) denotes the absence of sequence information in the database for other *virB* genes in these bacteria. The *L. pneumophila* *dot/icm* gene products shown across the bottom are homologues of the *Shigella flexneri* Collb-P9 (Incl) transfer proteins. This system can conjugally transfer DNA, but its proposed role in virulence is to export effector proteins.