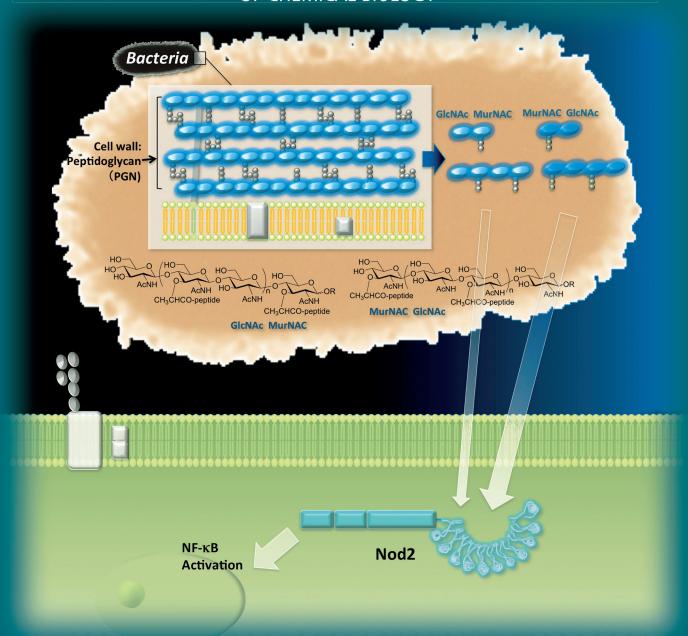
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The inside cover picture shows glycan-sequence-dependent human Nod2 recognition of PGN fragments. On p. 482 ff., Y. Fujimoto, K. Fukase, et al. report how they chemically synthesized a range of di- or tetrasaccharide fragments of PGN, and showed that $(MurNAc\beta(1{\longrightarrow}4)GlcNAc)_2\text{-containing structures}$ exhibited stronger activity than those containing $(GlcNAc\beta(1{\longrightarrow}4)MurNAc)_2$, produced with different enzymatic degradation process.

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