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The interaction between heparin and its primary receptor, the glycosaminoglycan-binding protein (GRP, PG-GP). The interaction between heparin and a recombinant form of the GRP, the glycosaminoglycan-binding protein (GRP, PG-GP), has been investigated using two alternative assays, an ELISA for GRP and a blotting assay for heparin-GRP complex formation. The assays were highly specific for each other as shown by the similar range of concentrations over which concentrations of each GRP or heparin are able to bind to each other, the mean K_d values calculated from these results being 34 nM (range, 2 to 285 nM) for GRP binding to heparin, 37 nM (range, 2 to 210 nM) for heparin binding to GRP, and 2 nM (range, 1 to 4.5 nM) for GRP binding to heparin. Experiments with a variety of experimental conditions suggested that the interaction between heparin and GRP does not require the presence of divalent cations or a protease-sensitive cleavage site in the GRP. Antibodies raised against GRP were used to investigate possible interactions between GRP and heparin in various preparations of heparin-binding protein (HBP, PG-LB, PG-GP, PG-GLA). A variety of preparations of heparin-binding protein from human umbilical cord, human liver, and sheep skin were investigated using blotting and an ELISA, which showed that all the preparations tested contain a GRP-like polypeptide that interacted with heparin but not with hyaluronate. The interactions were found to be dependent on the presence of divalent cations but also to be sensitive to proteases and sulfation of heparin. Taken together, these results suggest that the interaction between heparin and

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