INTERACTION OF THE FIRST COMPONENT OF COMPLEMENT (C1) IN PLATELET ADHESION AND AGGREGATION INDUCED BY COLLAGEN. J.L. Wautier, H. Souchon, A.P. Peltier and J.P. Caen, Hôpital Lariboisière, Paris, France

The attachment of C1 to human platelets through C1s and the inhibition of platelet aggregation induced by collagen by purified C1q prompted us to investigate a possible interaction between collagen-induced C1 and collagen. C1q is a molecule composed of two moieties: a collagen like region and a globular region composed of polypeptides which may be isolated following hydrolysis by collagenase and pepsin respectively. Pure C1 subcomponents (C1q, C1r, C1s), fractions of C1q (pepsin or collagenase treated) and specific antisera (anti C1q, anti C1s) were tested for their effect on collagen-induced aggregation and adhesion of gel filtered platelets. The pepsin derived fragment of C1q containing the collagen like region inhibited platelet aggregation and adhesion to collagen while collagenase treated C1q (globular moiety) did not modify the reaction. Preincubation of collagen with C1s inhibited platelet adhesion and aggregation. In other experiments platelets modified by anti C1s did not react with collagen while anti C1q, anti C3 or anti fibrinogen did not influence the reaction with collagen. These results suggest that the collagen like region of C1q binds to C1s with a possible reaction between C1s associated on the platelet or a platelet structure similar to C1s and collagen.

THE INVOLVEMENT OF FACTORS XII & XI IN THE PLATELET COAGULANT ACTIVITY. J.M. Connellan, S. Bowden, and P.A. Castaldi. Austin Hospital and University of Melbourne, Melbourne, 3084, Australia.

Platelets provide coagulant activities, other than the Phospholipid Platelet factor 3, which can be released from the platelet. In Platelet Poor Plasma, the initiation of the intrinsic pathway is proposed to proceed via factors XII & XI as well as 2 other factors, Prekallekrein and Kininogen. The role of these factors in the initiation of coagulation in Platelet Rich Plasma is under investigation. Walsh, (Br.J.Haemat.22.393,1972) and Muntz (Haemostasis in press 1977), have shown that platelets devoid of factor XII will still produce a coagulant activity when Platelet Rich Plasma is incubated with collagen. However, we have recently shown that this reaction requires the presence of factor XI. Factor XII has been purified by the method of Saito (Circ.Res.34,641,1974), and factor XI has been purified by the method of Connellan, Castaldi and Muntz. (Haemostasis, in press 1977).

Specific antibodies were developed in rabbits and the IgG from the antiserum was coupled to Sepharose. These factors could be extracted from plasma by the immobilized antibodies and could then be eluted by the use of alkali or guanidine HCL, thus resulting in a rapid purification of these factors.

Platelets were washed free of loosely bound clotting factors and provided significant amounts of coagulant activity when ruptured by either freezing and thawing or by glycerol treatment. Platelets from a factor XI deficient patient also provided this activity. The platelet activity was extracted by passage through both Sepharose-heparin column and Sepharose-anti-factor XI column; the activity appeared to be identical to plasma factor XI.

DIFFERENT REQUIREMENTS FOR INTRINSIC FACTOR-Xa FORMING ACTIVITY AND PLATELET FACTOR 3 ACTIVITY AND THEIR RELATIONSHIP TO PLATELET AGGREGATION AND RELEASE. P.N. Walsh and M.S. Lipscomb. Temple University School of Medicine, Philadelphia, Pennsylvania, U.S.A.

Platelets provide coagulant activity in part by making available membrane phospholipids which promote the reactions of: (1) factors XIIa, VIII, IX and calcium to activate factor X (intrinsic factor-Xa forming activity or XaFA); and, (2) factors Xa, V and calcium to activate prothrombin (platelet factor 3 activity or PF3A). We have studied the availability of XaFA and PF3A, platelet aggregation and [¹⁴C]-5HT release in washed platelet suspensions, incubated with collagen. XaFA developed rapidly (100% activity at 15-30 sec) well before maximum [¹⁴C]-5HT release (2-4 min) and decayed to <10% at 5-10 min. PF3A developed slowly (100% activity after 20 min stirring with collagen) well after maximal aggregation and release. Maximal XaFA availability occurred in unstirred platelet suspensions incubated with collagen and not undergoing aggregation or release, and was not enhanced by stirring, whereas PF3A did not become available unless platelet suspensions were stirred with resultant aggregation and release. PF3A, XaFA, aggregation and release were inhibited by pretreating platelet suspensions with p-chloro-mercuribenzenesulfonate, concanavalin A or a combination of 2-deoxyglucose and antimycin A, whereas pretreatment with indomethacin inhibited aggregation, release and PF3A availability but not XaFA availability. These results indicate that the biochemical determinants of XaFA and PF3A are different. PF3A developed only when aggregation and release occurred whereas XaFA was independent of aggregation and release.