Soluble glycoprotein VI (sGPVI) is shed from the platelet surface and is a marker of platelet activation in thrombotic conditions. We assessed sGPVI levels together with patient and clinical parameters in acute and chronic inflammatory conditions, including patients with thermal injury and inflammatory bowel disease and patients admitted to the intensive care unit (ICU) for elective cardiac surgery, trauma, acute brain injury, or prolonged ventilation. Plasma sGPVI was measured by enzyme-linked immunosorbent assay and was elevated on day 14 after thermal injury, and was higher in patients who developed sepsis. sGPVI levels were associated with sepsis, and the value for predicting sepsis was increased in combination with platelet count and Abbreviated Burn Severity Index. sGPVI levels positively correlated with levels of D-dimer (a fibrin degradation product) in ICU patients and patients with thermal injury. sGPVI levels in ICU patients at admission were significantly associated with 28- and 90-day mortality independent of platelet count. sGPVI levels in patients with thermal injury were associated with 28-day mortality at days 1, 14, and 21 when adjusting for platelet count. In both cohorts, sGPVI associations with mortality were stronger than D-dimer levels. Mechanistically, release of GPVI was triggered by exposure of platelets to polymerized fibrin, but not by engagement of G protein-coupled receptors by thrombin, adenosine 5'-diphosphate, or thromboxane mimetics. Enhanced fibrin production in these patients may therefore contribute to the observed elevated sGPVI levels. sGPVI is an important platelet-specific marker for platelet activation that predicts sepsis progression and mortality in injured patients.

Introduction

Beyond its primary role in hemostasis, fibrin also contributes to thrombotic and inflammatory conditions. Excessive fibrin formation occurs in acute and severe inflammation, including trauma, sepsis, disseminated intravascular coagulation (DIC), and deep vein thrombosis. Furthermore, fibrin deposition,
a blotted fibrinolysis, and hypofibrinogenemia are associated with multiple organ failure and mortality. Together with thrombocytopenia, coagulation factor consumption, and acute coagulopathy, fibrin formation and platelet activation are potentially linked in inflammatory settings.

Glycoprotein (GP) VI is the major platelet signaling receptor for collagen and fibrin. GPVI-fibrin engagement may underpin fibrin-related disease pathology. GPVI is expressed only on megakaryocytes and platelets in association with the Fc receptor γ-chain containing an immunoreceptor tyrosine-based activation motif (ITAM). On resting platelets, GPVI is predominantly monomeric, but clusters and dimerizes on activation, triggering ITAM signaling involving Src family kinases. In hemostasis and thrombosis, GPVI supports platelet adhesion and aggregation. GPVI is critical for the maintenance of vessel wall integrity and active factor X (FXa) also trigger GPVI shedding, quantified by an enzyme-linked immunosorbent assay. Plasma sGPVI reflects detection of the soluble ectodomain fragment (sGPVI) in plasma by rapid metalloproteolytic cleavage, and in some cases internalization, on activation. GPVI shedding is induced by GPVI ligands including collagen, collagen-related peptide (CRP), and venom toxins, and by engagement of other platelet ITAM receptors FcγRIIA and CLEC-2. Elevated fluid shear stress and active factor X (FXa) also trigger GPVI shedding, quantified by detection of the soluble ectodomain fragment (sGPVI) in plasma by an enzyme-linked immunosorbent assay. Plasma sGPVI reflects platelet activation in thrombotic conditions including microangiopathy, stroke, DIC, and Alzheimer’s disease and rheumatoid arthritis. However, elevated plasma sGPVI levels in these patient groups is surprising, as only a fraction of platelets would be exposed to collagen, FXa, or elevated shear, and neither FcγRIIA or CLEC-2 plays major roles in hemostasis/thrombosis. The recent finding that fibrin activates GPVI provides a plausible explanation for this increase.

Here, we report elevated sGPVI in inflammatory patient cohorts including patients with injury, active inflammatory bowel disease (IBD), and sepsis. sGPVI correlated with D-dimer levels in both acute inflammatory patients admitted to an intensive care unit (ICU) and patients with thermal injury. sGPVI was associated with sepsis progression and mortality, and sGPVI levels enhanced APACHE (Acute Physiology and Chronic Health Evaluation) II score prediction of mortality. Furthermore, we show that fibrin induces GPVI shedding in vitro, possibly explaining sGPVI elevation in patients with acute injury.

**Methods**

**Reagents**

Refer to supplemental Material.

**Blood collection**

Healthy controls. Venous blood was collected from consenting, healthy volunteers into sodium citrate (4%) or 3.2% trisodium citrate vacutainers (Becton Dickinson, Oxford, United Kingdom). Ethical approval was granted by Birmingham University Internal Ethical Review (ERN_11-0175).

ICU patients. Citrated blood was collected from 83 consenting patients admitted to tertiary ICU who underwent cardiac surgery, trauma, invasive ventilation for more than 48 hours, or acute brain injury at University Hospital of Liège, Liège, Belgium. (The experimental protocol was approved by the ethics committee of the University Hospital of Liège [Centre Hospitalier Universitaire; reference number B707201111981]). Sepsis diagnosis based on previous sepsis definitions.

**Thermal injury patients.** Ninety-nine patients with injury affecting up to 95% total body surface area (TBSA) were recruited to the Scientific Investigation of the Biological Pathways Following Thermal Injury Study (SIFTI: REC-12/EM/0432) at the Queen Elizabeth Hospital, Birmingham, United Kingdom. Citrated blood was collected at intervals after injury (day [D] 1, D3, D7, D14, D21, and D28 and month 2, 3, 6, and 12 postinjury). Sepsis diagnosis was made when more than 3 American Burn Association criteria were met, plus positive bacterial culture and/or evidence of antibiotic response. Platelet impedance counts were measured using a Beckman-Coulter UniCel DxH-800 (High Wycombe, United Kingdom) or a Sysmex XN-1000 Analyzer (Milton Keynes, United Kingdom).

Patients with IBD. Citrated blood was collected from 42 consenting patients diagnosed with inactive or active Crohn’s disease and/or ulcerative colitis (UC; Ethics: REC:13/NE/0249).

Patient details can be found in the supplemental Material.

**Plasma preparation**

Platelet-free plasma was isolated from blood by serial centrifugation: 2000g for 20 minutes and then 13 000g for 20 minutes at 4°C for samples from patients with thermal injury and two 15-minute 2500g centrifugations at 37°C for samples from other patient groups and healthy controls (HCs).

**Washed platelet preparation and experiments**

Washed platelets were prepared as described and washed twice with calcium-free Modified Tyrode’s buffer (134 mM NaCl, 0.34 mM Na₂HPO₄, 2.9 mM KCl, 12 mM NaHCO₃, 20 mM HEPES, 5 mM glucose, and 1 mM MgCl₂ at pH 7.3) by centrifugation at 1000g for 10 minutes and resuspended to 500 x 10⁸ platelets/L. Washed platelets were preincubated for 5 minutes with 1 mM CaCl₂ and with/without inhibitors (2 μM Gil254023, 25.7 μM GM6001, 10 μM PRT060318, or 10 μM dasatinib) before addition of 9 μM epifibatide (1×Ibß3 inhibitor). Platelet suspensions were stirred at 1200 rpm/37°C for 1 minute before agonist addition. For fibrin treatment, fibrinogen (100 μg/mL) was added 3 minutes before thrombin (1 U/mL) stimulation. Next, 10 mM Gly-Pro-Arg-Pro peptide (polymerization inhibitor) was added to fibrinogen to achieve monomeric fibrin conditions. Samples were stirred for 1 hour. Levels of intact and proteolyzed GPVI were assessed by western blot, using 1 μg/mL rabbit anti-human GPVI cytoplasmic tail antibody, detecting intact and cleaved GPVI. Densitometry measurements were made using Li-cor Image Studio software.

**sGPVI Enzyme-Linked Immunosorbent Assay**

sGPVI levels were measured by sandwich enzyme-linked immunosorbent assay, and concentrations extrapolated from standard curves generated by serial dilution of GPVI ectodomain into 5% vol/vol GPVI-depleted plasma.
D-dimer measurements

D-dimer levels were measured in plasma using the Innovance D-dimer immunoturbidimetric assay (Siemens Healthcare, Eschborn, Germany).

Statistical analysis

Results are reported as mean ± standard deviation, unless stated. D’Agostino-Pearson normality tests determined normality. Student t-tests were performed for normally distributed data; otherwise, Mann-Whitney U tests were performed. For multiple groups, Kruskal-Wallis tests with Dunn’s posttests were performed. Univariate and multivariate logistic regression was used to analyze variable association with 28- and 90-day mortality. Spearman's rank correlation coefficients assessed associations between sGPVI levels and clinical parameters. Longitudinal analysis of sGPVI levels on sepsis, multiple-organ failure, and mortality was performed using linear mixed-effects models (supplemental Material). Statistical analyses were performed using GraphPad Prism (versions 5, 7), SPSS (IBM), and R (version 3.0.3).

Results

We assessed the utility of sGPVI as a biomarker of disease pathology and progression in patients with injury, sepsis, and inflammation by measuring sGPVI in samples from ICU patients, patients with thermal injury, and patients with IBD.

sGPVI levels correlate with clinical and platelet-specific parameters in ICU patients

Plasma sGPVI levels were measured in 83 ICU patients with acute brain injury (n = 12), trauma (n = 13), elective cardiac surgery (n = 53), or prolonged ventilation (n = 5). At T1 (day of ICU admission), sGPVI levels were not significantly elevated above age-matched HCs or at 48 hours postadmission (Figure 1A), although wide ranges of sGPVI levels were observed at both points (5th-95th percentiles: T1, 11.1–43.5 ng/mL; 48 hours, 12.1–39.8 ng/mL). sGPVI at 48 hours increased in 42/83 patients (51%; Figure 1A). In the 15 ICU patients who developed sepsis, sGPVI levels were not elevated above T1 samples, both on day of sepsis diagnosis (Tx) and 7 days after (Tx+7), suggesting no elevations with sepsis onset.
or treatment phases of ICU patients (Figure 1B). sGPVI levels at T1 did not associate with sepsis occurrence, using a simple logistic regression model ($P = .085$).

We compared sGPVI levels in ICU T1 patient samples against biological and clinical parameters (Table 1). As expected, positive correlations were seen with C-reactive protein, white blood cell count, and APACHEII and Sequential Organ Failure Assessment (SOFA) scores. Interestingly, platelet-related parameters, including platelet count and fibrinogen-binding, as well as D-dimer and fibrinogen levels, also correlated significantly with sGPVI. After correction for platelet count, D-dimer levels and platelet fibrinogen-binding remained significantly correlated with sGPVI. After correction for platelet count, D-dimer levels did not increase at 48 hours (Figure 1C). In some patients, D-dimer levels were elevated above 5 mg/L fibrinogen equivalent units (FEU), representing a strong increase.

### Table 1. sGPVI correlations with clinical and biological parameters of ICU patients

<table>
<thead>
<tr>
<th>Variable 1</th>
<th>Variable 2</th>
<th>$r$</th>
<th>$P$</th>
</tr>
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<td>.50</td>
</tr>
<tr>
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<td>Sex</td>
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<td>Stroke</td>
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<td>.11</td>
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<td>Fibrinogen</td>
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<tr>
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<td>Platelet count</td>
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<td>.0008</td>
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<tr>
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<td>D-dimers</td>
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<tr>
<td>C-reactive protein</td>
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<td>C-reactive protein</td>
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<td>D-dimer</td>
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<td>D-dimer</td>
<td>TNF$\alpha$</td>
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<td>IL-17A</td>
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<td>.8</td>
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<td>IL-7</td>
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<td>.024</td>
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<td>IFN$\gamma$</td>
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<td>Platelet-bound fibrinogen (MFI)</td>
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<td>.0009</td>
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<td>Platelet-bound fibrinogen (MFI)</td>
<td>Platelet P-selectin (MFI)</td>
<td>0.14</td>
<td>.22</td>
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</table>

Spearman’s correlation coefficients, represented as $r$ values, between sGPVI levels and clinical/biological parameters measured from samples taken on day 1 (T1) on admission to ICU. Significance observed when $P < .05$. Bold indicates significant correlation. MFI, mean fluorescence intensity; PTT, partial thromboplastin time.

sGPVI levels in ICU patients are associated with mortality

sGPVI association with ICU patient mortality at T1 was assessed by simple logistic regression, using a sGPVI cutoff based on median levels found in healthy donors to stratify this cohort into low/normal ($\leq 22.3$ ng/mL) and elevated ($> 22.3$ ng/mL) sGPVI groups. The elevated sGPVI group showed reduced survival probability (Figure 1D). Significant associations between T1 sGPVI with 28- and 90-day mortality were also observed (supplemental Table 4). This significant association remained when using multiple logistic regression alongside platelet count ($P = .036$ and $P = .014$). SOFA and APACHEII scores formed significant associations with mortality (supplemental Table 4); however, sGPVI levels no longer predicted mortality in patients with sepsis from this ICU cohort after correction for disease severity. Interestingly, D-dimer levels and platelet count formed no significant associations with 28- and 90-day mortality at T1 ($P = .059$ and .058 and $P = .220$ and .077, respectively; supplemental Table 4).

sGPVI is raised in thermal injury patients with sepsis

Although the range of sGPVI levels in HCs is tightly maintained, sGPVI in longitudinal samples from 99 patients with thermal injury with burns afflicting up to 95% TBSA burn (mean, 24%; IQR, 8%-38%) showed a broad distribution (Figure 2A). sGPVI levels increased from D1 postinjury, with a significant peak observed at around D14 before slowly returning to HC levels by month 2 postinjury (Figure 2A).

sGPVI levels weakly correlated with platelet count in this cohort (Spearman’s rank coefficient $r = 0.119$; $P = .015$). Consistent with other reports of significant platelet count reductions at D3 followed by rebound thrombocytosis at D14 postinjury,42 here platelet counts within septic and nonseptic patients reached a nadir at D3, followed by rebound thrombocytosis peaking at D14 for nonseptic patients and D21 for septic patients (Figure 2B).

A 58% incidence of sepsis occurred in this cohort, with onset between D4 and D8 (median, 5.5 days).37 Septic patients had higher peak sGPVI levels compared with nonseptic patients and HCs (Figure 2C). After normalization for platelet count, sGPVI was significantly elevated in septic patients compared with nonseptic patients at D3, D7, D14, and D28 (Figure 2D), coincident with sepsis onset and progression.

D-dimer levels in patients with thermal injury with burns of 15% TBSA or more ($n = 61$) increased from D1 postthermal injury to a peak at D14, gradually normalizing during M3 (Figure 2E), echoing sGPVI levels (Figure 2A). Significant positive correlation with D-dimer and sGPVI levels were observed when assessing samples from all points ($r = 0.35$; Table 2). Alongside this, significant positive correlations were seen at D14, D28, and month 2, with D14 giving the strongest correlation ($r = 0.48$; Table 2, sGPVI). D-dimer and sGPVI correlations also strengthened after correction for platelet count (Table 2, normalized sGPVI).

sGPVI levels in patients with thermal injury are associated with sepsis and mortality

sGPVI associations with sepsis, multiple-organ failure, and mortality were assessed in patients with thermal injury. $\chi^2$ statistical analysis demonstrated significant associations with sGPVI and the proportion of patients developing sepsis and patient outcome ($P < .001$), with overrepresentation of patients bearing high sGPVI developing sepsis across all times (supplemental Table 5). Significant sGPVI associations with sepsis patient outcome at D7 and D14 postinjury were observed when examining set times ($P = .023$ and .043, respectively). Interestingly, sGPVI association with sepsis is
at D7 and D14 was strengthened when corrected for platelet count (supplemental Table 5).

Longitudinal statistical prediction models were used to evaluate sGPVI as a predictive marker of sepsis (Figure 2F). The sGPVI discriminatory value for predicting sepsis in patients with thermal injury, represented as an area under the receiver operating characteristic (AUROC) curve value, was determined using a linear mixed-effects model of sGPVI and time and identified that D14 provided the strongest predictive value for sepsis (AUROC, 0.73) and multiple-organ failure (AUROC, 0.77; supplemental Table 8).

Multivariate analysis of sGPVI with platelet count strengthened the sepsis predictive value at D3, D7, and D14 ($P = .75-.78$; supplemental Table 9).

Logistic regression was performed to assess sGPVI associations with mortality in the patients with thermal injury with 15% or more TBSA burns and compared with D-dimer levels. Significant associations of sGPVI levels at D1 and 28-day mortality were observed ($P < .05$; supplemental Table 6). After correcting for platelet count, D1 sGPVI mortality association was not significant, but it was at D14 and D21 (supplemental Table 7). Interestingly,
D-dimer association with 28-day mortality was not significant (supplemental Table 7). D7, D14, and D28 samples postinjury gave the best sGPVI discriminatory predictive values for predicting mortality at 1 year (represented as AUROC, 0.68–0.75; supplemental Table 8). Mortality predictive values of sGPVI were also improved when adjusted for platelet count (supplemental Table 7). Multivariate analysis of sGPVI with platelet count improved the predictive value at D3, D7, and D14 for predicting mortality (P = .71–.83; supplemental Table 9). sGPVI and ABSI values on D1 enhanced or maintained predictive power for sepsis (AUROC, 0.80; compared to AUROC, 0.77 for ABSI alone) and mortality (AUROC, 0.80; unchanged from ABSI alone). sGPVI and APACHE(II) together demonstrated improved prediction of mortality from D3 to D28 (supplemental Table 10). Together, these data suggest sGPVI could be a useful addition to other clinical parameters for predicting sepsis and mortality.

Table 2. Correlations of sGPVI with D-dimers in patients with thermal injury

<table>
<thead>
<tr>
<th>Time</th>
<th>HC</th>
<th>D1</th>
<th>D3</th>
<th>D7</th>
<th>D14</th>
<th>M2</th>
<th>M3</th>
<th>M6</th>
<th>M12</th>
<th>All times</th>
</tr>
</thead>
<tbody>
<tr>
<td>sGPVI</td>
<td>n</td>
<td>r</td>
<td>P</td>
<td>n</td>
<td>r</td>
<td>P</td>
<td>n</td>
<td>r</td>
<td>P</td>
<td>n</td>
</tr>
<tr>
<td>0</td>
<td>12</td>
<td>.03</td>
<td>.943</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>33</td>
<td>.39</td>
<td>.011*</td>
<td>341</td>
</tr>
<tr>
<td>1</td>
<td>35</td>
<td>.04</td>
<td>.808</td>
<td>100</td>
<td>.30</td>
<td>.503</td>
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</tr>
<tr>
<td>2</td>
<td>20</td>
<td>.28</td>
<td>.239</td>
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<td>.908</td>
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<tr>
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<td>.188</td>
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<td>.27</td>
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<td>.03</td>
<td>.943</td>
<td>341</td>
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</tbody>
</table>

sGPVI values, between sGPVI levels and normalized sGPVI corrected for platelet count and D-dimer levels in patients with thermal injury at different points at injury. Bold indicates significant correlation.

Spearman’s correlation coefficients, represented as r values, between sGPVI and normalized sGPVI.

D-dimer values, between sGPVI levels and normalized sGPVI corrected for platelet count and D-dimer levels in patients with thermal injury at different points at injury. Bold indicates significant correlation.

**P < .05.
***P < .01.
****P < .005.

sGPVI is detectable in patients with IBD. (A) sGPVI levels of patients with IBD with inactive and active Crohn’s disease (n = 13 and n = 12) and inactive and active UC (n = 12 and n = 13) compared with HC (n = 20). Mann-Whitney U test was performed to compare sGPVI levels of different IBD patient groups and HC (median shown). (B) sGPVI levels of patients with IBD with hemoglobin (Hgb) levels of above or below 120 g/L (n = 28, n = 13, respectively) compared with HC (n = 20). A Kruskal-Wallis statistical test with Dunn’s multiple comparisons was performed to compare the 3 groups of patients (median shown).

Exposure of platelets to fibrin, but not GPCR stimulation, induces GPVI shedding

Engagement of GPVI by collagen, CRP, or convulxin leads to ITAM-signaling dependent metallopeptolytic release of a 55-kDa sGPVI fragment, leaving a 10- to 15-kDa membrane-bound remnant.23,24,26,27,43 To ascertain whether fibrin could induce GPVI shedding, we first assessed the effect of thrombin or other GPCR ligands on GPVI levels. GPVI shedding was calculated as a percentage of GPVI detected in unstimulated samples compared with GPVI levels after stimulation. Figure 4A shows that 1-hour treatment of washed platelets with thrombin induced a loss of full-length GPVI that was comparable to shedding achieved by collagen or CRP and similar loss to the potent GPVI shedder, A23187 (Figure 4C). These treatments induced the appearance of the 10- to 15-kDa remnant in ~50% of donors. In contrast, treatment with adenosine 5’-diphosphate (ADP), the PAR-1 or PAR-4 peptide agonists, or thromboxane mimetic U46619 did not induce loss of intact GPVI (Figure 4B-C).

As engagement of PAR-1 or PAR-4 via activating peptides did not induce GPVI cleavage, and GPVI does not carry a recognized...
thrombin-cleavage site, we considered whether GPVI shedding was secondary to GPVI/fibrin engagement. Platelets were treated with thrombin in the presence of fibrinogen to produce fibrin. This treatment triggered loss of intact GPVI (Figure 5A) and was comparable to A23187-induced shedding. Fibrinogen alone did not induce GPVI shedding (Figure 5A), and shedding required fibrin polymerization as inclusion of GPRP ablated GPVI proteolysis (Figure 5A-B), demonstrating that fibrin polymers were more effective than monomers at inducing shedding.

To assess whether fibrin-induced GPVI shedding required ITAM signaling, platelet suspension s were pretreated with maximally effective concentrations of Src and Syk inhibitors, dasatinib, and PRT060318. GPVI shedding induced by thrombin-fibrinogen in the presence of either inhibitor was not different to fibrin alone (Figure 5), indicating that an active ITAM signaling pathway was not crucial for fibrin-induced GPVI shedding. The role of multiple metalloproteinases in fibrin-induced GPVI shedding was determined by inclusion of GM6001, a broad metalloproteinase inhibitor, or GI254023, an inhibitor of A Disintegrin and Metalloproteinase (ADAM) 10, to platelet suspensions, with both having minimal effect on reducing fibrin-mediated shedding (Figure 5B).

**Discussion**

The incidence, management, and knowledge of pathophysiological processes relating to sepsis have improved during the last 20 years; however, sepsis remains a significant public health problem across the world, with more than 31 million cases presenting annually and sepsis-related fatality occurring in 1 in 5 cases.44,45 Biomarkers with high sensitivity and specificity that rapidly and accurately differentiate sepsis from noninfectious conditions including systemic inflammatory response syndrome are therefore in demand to help implement the correct therapeutic regime. Diagnosis of sepsis is a subjective clinical judgment focused on assessing the roots of sepsis, using tools such as the SOFA or APACHE scoring systems,46-48 with both systems including evaluation of patient platelet count. Blood rheology and platelet function are progressively and severely altered in patients with severe sepsis.49,50 Evidence suggests a prominent role for inappropriate platelet activation and aggregation during sepsis,51 and platelet indices beyond platelet count are useful to evaluate illness severity and prospectively identify critically ill patients.52,53 In this regard, sGPVI represents an excellent candidate marker of pathological platelet activation in the setting of sepsis, as it is a platelet/megakaryocyte-specific membrane protein that is stable on circulating platelets but
Figure 5. Fibrin stimulation of platelets induces GPVI shedding. (A) Western blot for GPVI after stimulation of platelets in suspension by fibrinogen, fibrin, or fibrin in the presence of inhibitors. Washed platelets (500 × 10^6/L) were stimulated with fibrinogen (100 μg/mL) alone, thrombin (1 U/mL) in the presence of fibrinogen (polymerized fibrin), fibrin in the presence of Src and Syk inhibitors (dasatinib, 10 μM; PRT-060318, 10 μM), fibrin in the presence of G254023 (2 μM, ADAM10 inhibitor) and GM6001 (25.7 μM, broad matrix metalloproteinase inhibitor), and monomeric fibrin in the presence of GPRP (10 μM) under stirring conditions for 1 h at 37°C in the presence of epifibatide (9 μM) and CaCl2 (1 mM). GPRP was added with fibrinogen 3 minutes before thrombin stimulation to prevent fibrin polymerization. G254023 and GM6001 were added 5 minutes before fibrinogen and thrombin stimulation. Membranes were blotted for GPVI and GPVI remnant after shedding. (B) Quantitation analysis of GPVI shedding induced by different forms of fibrin and fibrin-induced shedding in the presence of inhibitors. Percentage of GPVI represents levels calculated as a percentage of GPVI in unstimulated samples compared with GPVI levels after stimulation. Results are shown as mean ± SEM. A 1-way ANOVA was performed, with Tukey's posttest, to compare fibrin-induced shedding in the presence of GPRP and the presence of inhibitors and fibrinogen alone to unstimulated samples. N.S., nonsignificant; n = 7 donors.

released on ligand engagement of ITAM receptors, exposure to FXa, or abnormal fluid shear rates. sGPVI levels are not elevated as a consequence of ablated platelet production.

We have investigated the utility of sGPVI as a marker of platelet activation in these pathological settings and explored the value of sGPVI to aid diagnosis and predict patient outcomes in conjunction with standard clinical parameters, including platelet count and injury severity scores. We have also investigated mechanisms that may drive release of sGPVI in trauma/inflammation patients. Elevated sGPVI levels were detected in the plasma of patients with thermal injury and patients with IBD with active disease. Although there were no significant sGPVI elevations in patients with ICU, we found significant correlations with sGPVI levels and other biological parameters including ISTH DIC score, C-reactive protein, and injury severity scores (APACHEII, SOFA). sGPVI levels no longer predicted mortality in ICU sepsis patients after correction for disease severity.

sGPVI was not elevated during the acute thermal injury phase but became significantly elevated with sepsis development. Disseminated intravascular coagulation is commonly observed in septic patients. Platelets are activated and form fibrin-rich thrombi, leading to further platelet activation. Fibrin accumulates during acute inflammation and tissue injury and underlies sepsis pathology. D-dimers, a fibrin byproduct of coagulation and clot resolution produced by the action of plasmin, are an indirect index of fibrin formed in these patients. D-dimer levels correlated with sGPVI levels in ICU patients and patients with thermal injury, even when correcting for platelet count, suggesting a potential link between sGPVI and fibrin in multiple patient cohorts. In support of this, sGPVI levels significantly correlated with platelet-bound fibrinogen in samples from ICU patients, indicating both fibrin formation and platelet activation. Synergy between D-dimer and sGPVI levels in both ICU and thermal injury patient cohorts implies a role for fibrin in GPVI shedding observed in these patients.

Spontaneous aggregation of platelets and hyperfibrinogenemia are enhanced in the acute period (<48 hours) after thermal trauma. We found that sGPVI levels were significantly elevated in patients with burn injuries at D14 postinjury. Low sGPVI levels were observed at earlier points, suggesting that only a fraction of platelets were activated by the injury and related collagen exposure, or that low sGPVI levels reflected low platelet counts. When sGPVI levels were adjusted for platelet count, there was significant elevation at D3 postinjury in septic patients and sGPVI remained elevated at D7 and D14. As minimal collagen exposure is expected at these times, an alternative mechanism for platelet activation and GPVI shedding is likely.

We observed associations between sepsis and high sGPVI levels in patients with thermal injury at all points when stratifying sGPVI levels into low (≤12.7 ng/mL), medium (>12.7 to ≤37.7 ng/mL), and high (>37.7 ng/mL) levels. These associations were improved when sGPVI levels were corrected for platelet count. Significant associations with sGPVI and sepsis were observed at D7, potentially correlating to sepsis onset. A longitudinal statistical prediction of sGPVI levels in septic and nonseptic patients over time indicated sGPVI having moderate value for predicting sepsis at early points with a score of 0.68 at D7 and 0.73 at D14, which improved...
Fibrin-induced shedding was assessed whether fibrin stimulation induced GPVI shedding. Independent of fibrinogen interaction, ADAM10 plays a major role in GPVI shedding, with fibrin generation being a potential consequence of disease for most patients with IBD, with around 17% of patients with IBD having iron deficiency anemia, increasing in prevalence to around 60% when studying hospitalized patients. Although the molecular basis is unclear, there are numerous links between iron deficiency and platelet activation. Fibrin levels may reflect platelet activation in iron-deficient patients with IBD, which could also aid clinical management of these patients. In the chronic IBD inflammatory cohort, sGPVI levels were elevated in patients with active UC and correlated inversely with ferritin and hemoglobin levels. Iron deficiency anemia is commonly associated with IBD, with around 17% of patients with IBD having iron deficiency anemia, increasing in prevalence to around 60% when studying hospitalized patients. Although the molecular basis is unclear, there are numerous links between iron deficiency and platelet activation. Fibrin levels may reflect platelet activation in iron-deficient patients with IBD, which could also aid clinical management of these patients.

Fibrin was recently reported to activate platelets via GPVI, with the interaction contributing to thrombus stabilization. Here we demonstrate for the first time that fibrin-exposed platelets shed GPVI to an extent comparable to GPVI ligands. GPVI shedding was not a result of PAR engagement, as neither PAR-1 nor PAR-4-activating peptides induced GPVI shedding. Treatment with ADP and U46619 also failed to induce shedding, suggesting activation of platelets via GPCRs is not sufficient to induce GPVI shedding. Thrombin treatment variably stimulated GPVI shedding between donors, consistent with other studies, which may depend on sufficient fibrinogen storage and release from platelets and amount of fibrin formed. GPVI lacks a recognized cleavage site for thrombin, arguing against direct GPVI cleavage by the protease.

Fibrin generation is a potential consequence of disease for most patients and is minimal exposure to collagen. Plasma sGPVI strongly associates with onset of sepsis and with mortality, supporting the relevance of sGPVI as a clinical platelet activation marker.

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Authorship

Contribution: S.J.M designed and performed research; collected, analyzed, and interpreted data; made the figures; and wrote the manuscript. R.J.D. and C.D. collected, analyzed, and interpreted data and processed samples. C.S.-M.L., J.B., and N.M., analyzed and interpreted data. C.L. N.L., P. Hampson, C.M.W., A.B.,
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