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## Correction to: Plasma Fibrinogen Is a Natural Deterrent to Amyloid β-Induced Platelet Activation and Neuronal Toxicity



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Following publication of the original article (Sonkar et al. 2016), the author reported an error in Fig. 1. The correct version of Fig. 1 is as follows:

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## Reference

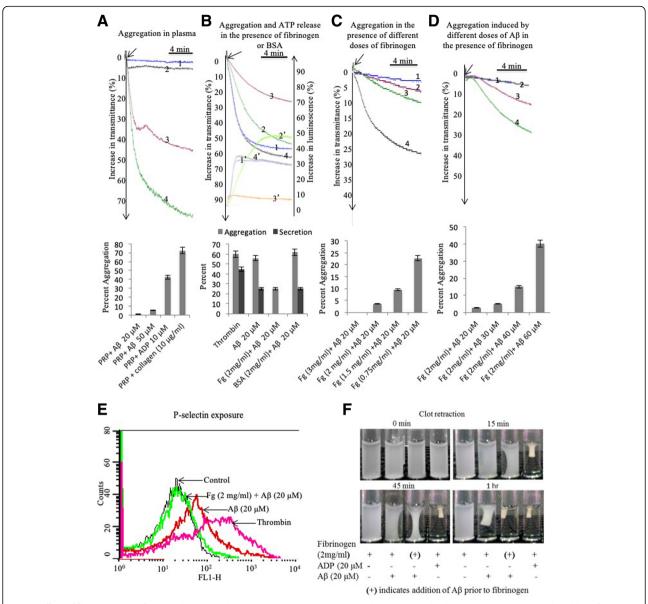
Sonkar VK, et al. Plasma fibrinogen is a natural deterrent to amyloid β-induced platelet activation and neuronal toxicity. Mol Med. 2016;22:224–32. https://doi.org/10.2119/molmed.2016.00003.

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**Fig. 1** Effect of fibrinogen on A $β_{25-35}$ -induced platelet activation. **a** Tracings 1, 2, 3 and 4 represent platelet aggregation in PRP induced by A $β_{25-35}$  (20 and 50 μmol/L), ADP (20 μmol/L) and collagen (10 μg/mL), respectively. The corresponding bar diagram is shown in the lower panel. **b** Tracings 1–4 represent aggregation, and tracings 1'–4' represent corresponding dense granule secretion in washed platelets stimulated with either thrombin (1 U/mL; tracings 2 and 2') or A $β_{25-35}$  (20 μmol/L; tracings 1, 1', 3, 3', 4 and 4'). Tracings 3 and 3' represent platelets pretreated with fibrinogen (2 mg/mL), whereas tracings 4 and 4' represent platelets pretreated with BSA (2 mg/mL). Corresponding bar diagram is shown in the lower panel. **c** Tracings 1, 2, 3 and 4 represent aggregation of washed platelets induced by A $β_{25-35}$  (20 μmol/L) in the presence of 3, 2, 1.5 and 0.75 mg/mL fibrinogen, respectively. Corresponding bar diagram is shown in the lower panel. **d** Tracings 1, 2, 3 and 4 represent aggregation of washed platelets induced by A $β_{25-35}$  (20, 30, 40 and 60 μmol/L, respectively) in the presence of fibrinogen (2 mg/mL). Corresponding bar diagram is shown in the lower panel. **e** Flow cytometric analysis of expression of P-selectin on surface of platelets treated with different reagents and agonists as indicated. **f** Retraction of fibrin clot by platelets stimulated with either A $β_{25-35}$  (20 μmol/L) or ADP (20 μmol/L) in presence of fibrinogen (2 mg/mL) added either before or after exposure to agonists as indicated