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### **Articles Published in Journals That Entered SCI, SSCI and AHCI Indexes**

- I. **Identification of three novel pathogenic ITGA2B and one novel pathogenic ITGB3 mutations in patients with hereditary Glanzmann's thrombasthenia living in Eastern Turkey**  
Karaman K., Yürektürk E., Geylan H., Yasar A. S. , Karaman S., Aymelek H. S. , Çetin M., Öner A. F.  
PLATELETS, 2020 (Journal Indexed in SCI)
- II. **Wiskott-Aldrich syndrome: Two case reports with a novel mutation.**  
Kamuran K., ÇETIN M., GEYLAN H., KARAMAN S., DEMIR N., YUREKTURK E., YAVUZ İ. H. , YAVUZ G., TUNCER O.  
Pediatric hematology and oncology, vol.34, pp.286-291, 2017 (Journal Indexed in SCI)

### **Refereed Congress / Symposium Publications in Proceedings**

- I. **wiscott aldrich syndrome: new mutaiton in two cases**  
Karaman K., yürektürk e., Geylan H., Çetin M., Yavuz İ. H.  
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