



Introduction to EndNote

Dr. Ghasem Azarian

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What is EndNote? Reference management

- Mendeley: a free reference manager and an academic social network
- Zotero: a powerful, easy-to-use research tool
- **Refworks**: online bibliographic management program
- EndNote: an industry standard software tool
 - On the Windows
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J Pediatr Hematol Oncol. 2017 Jul;39(5):332-336. doi: 10.1097/MPH.000000000000790.

Health-related Quality of Life and its Predictors Among Transfusion-depende Patients.

Tuysuz G1, Tayfun F.

Author information

Abstract

PURPOSE: With regular blood transfusions and modern medical management strategies, transfusion-depender are currently living into adulthood, but they still suffer from the chronicity of the disease and its complications. T the health-related quality of life (HRQoL) scores in pediatric TDT patients compared with healthy controls.

PATIENTS AND METHODS: The PedsQL 4.0 Generic Core Scales were administered to 80 pediatric TDT patients (aged 5 to 10 y, child selfreport) and their parents (parent-proxy report), as well as age-matched and sex-matched healthy controls (80 participants and their parents).

RESULTS: Patients with TDT and their parents rated lower HRQoL scores in all domains (physical, emotional, social, and school functioning) compared with the healthy population (P<0.01). The univariate analysis indicated that the total HRQoL score for children's self-reports was negatively predicted by high ferritin levels and the presence of any complication. The only negative predictor of the total HRQoL score in multivariate analyses was a high ferritin level (>1800 ng/dL).

CONCLUSIONS: This study showed that thalassemia has a negative impact on HRQoL when compared with a healthy population. For a better quality of life, intensive treatment should be applied for complications and high ferritin levels.

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Parent proxy-report of their children's healthrelated quality [Health Qual Life Outcomes. 2007]

Health related quality of life in Middle Eastern children with beta-thala [BMC Blood Disord. 2012]

Review Meta-analysis of quality of life in children and adolescents with ADI [Res Dev Disabil. 2016]

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Articles Case law	The thalassemia syndromes: molecular basis and prenatal diagnosis in 1990. HH Kazazian Jr - Seminars in hematology, 1990 - europepmc.org Abstract In this review I have outlined the molecular basis and prenatal diagnosis of alpha- thalassemia and then concentrated on the state of our knowledge of the molecular basis of
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Any time Since 2017 Since 2016 Since 2013 Custom range	[HTML] Complications of β-thalassemia major in North America , AR Cohen, Thalassemia Clinical Research Network - Blood, 2004 - Am Soc Hematology Treatment of patients with β-thalassemia major has improved dramatically during the past 40 years; however, the current clinical status of these patients remains poorly characterized. We performed a cross-sectional study of 342 patients in the Registry of the National Institutes of Cited by 409 Related articles All 7 versions Cite Save
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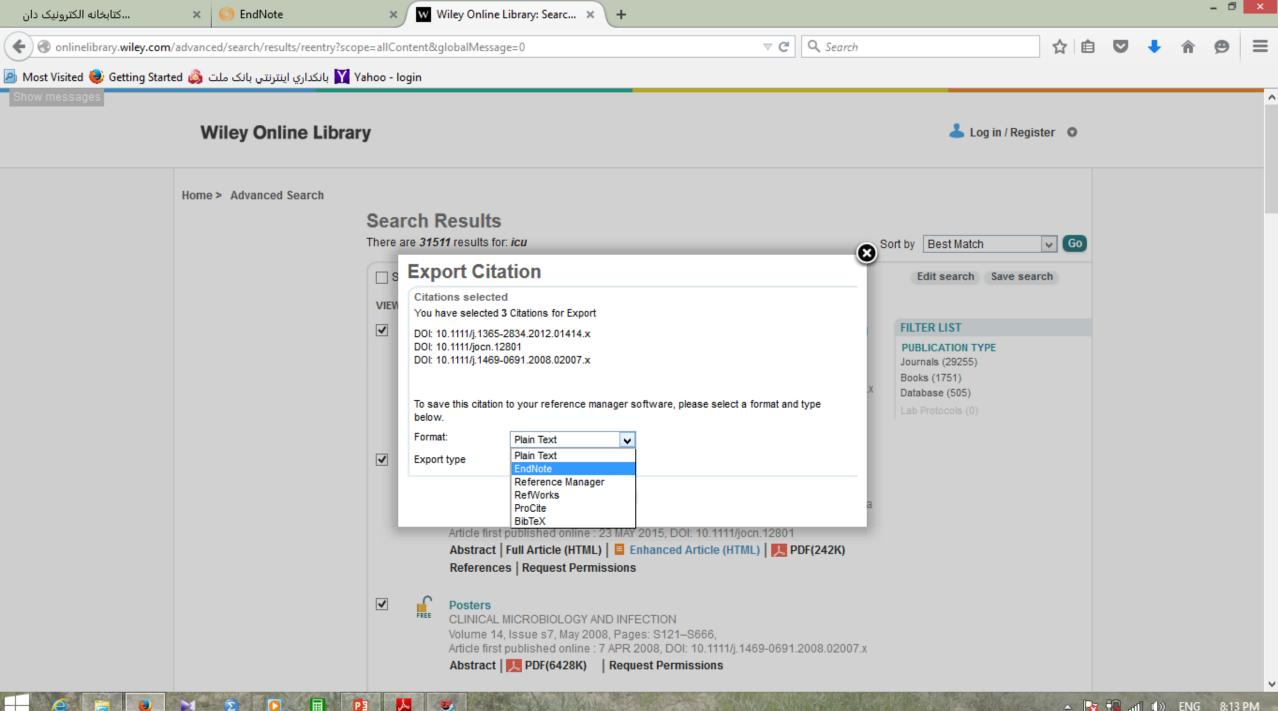
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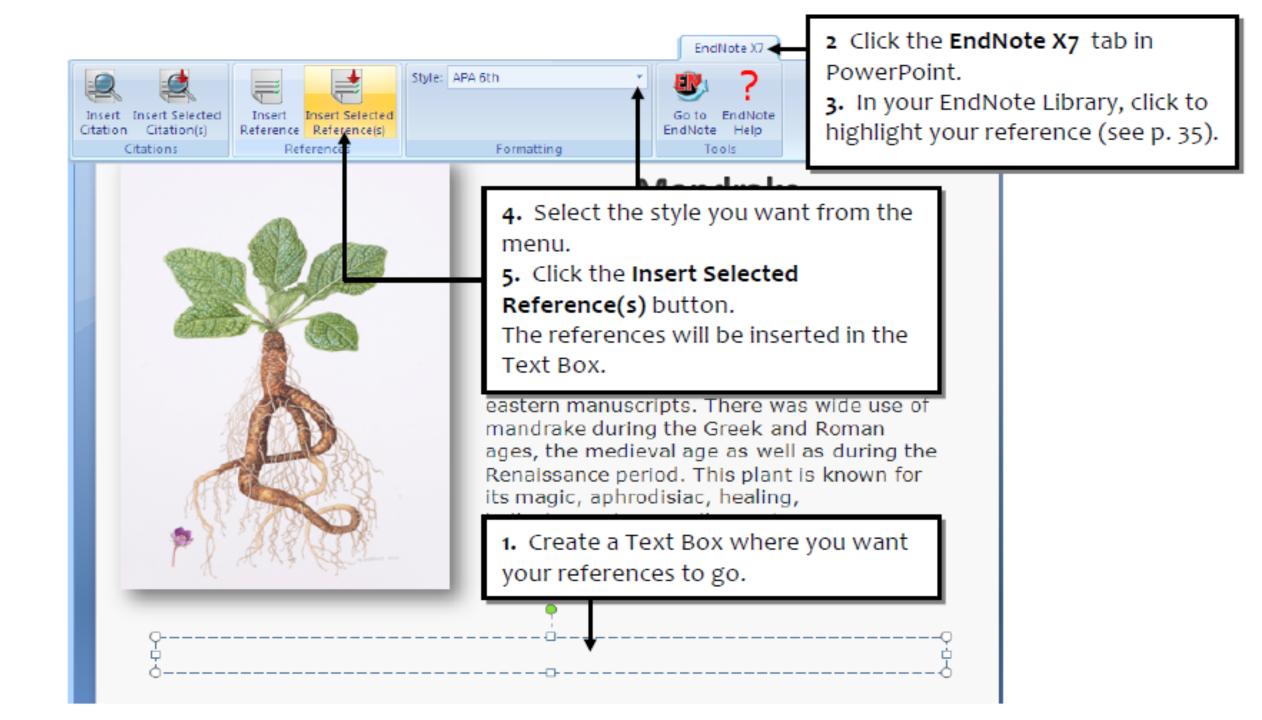
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