
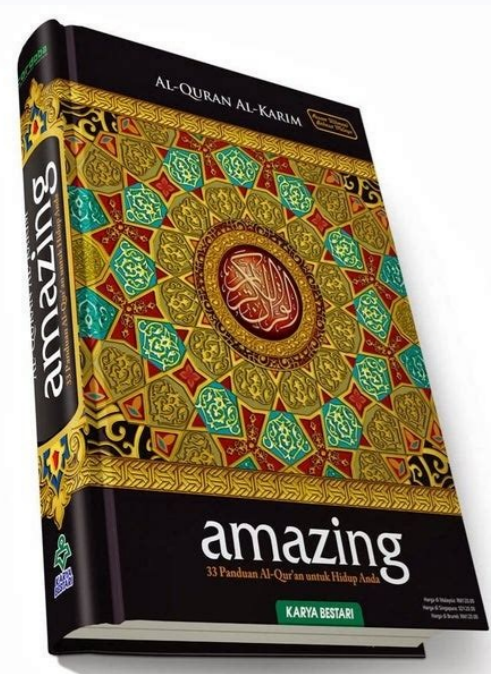


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Kisah Budak-budak Nakal

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Pada zaman dahulu, ada sekumpulan budak nakal yang sering bermain di dalam hutan. Di dalam hutan itu, terdapat (sebuah pondok kecil yang **didiami** oleh seorang lelaki tua.

Budak-budak nakal itu sering **mengejek** dan **mempersendakan** lelaki tua itu. Mereka langsung tidak menghormati lelaki tua itu.

Sikap mereka yang **keterlaluan** menyebabkan lelaki tua itu merasakan budak-budak itu perlu diberi pengajaran. Maka, lelaki tua itu **berdoa** kepada Tuhan supaya budak-budak nakal itu berubah sikap.

Pada suatu hari, selepas budak-budak nakal itu mengejek lelaki tua, mereka berasa sangat dahaga. Mereka terus pergi ke sungai untuk minum. (Ketika berada di **tebing sungai**, langkah terkejutnya (mereka -tatkala melihat bayangan muka mereka yang sangat tua dan buruk di permukaan sungai.)

Setelah itu, barulah (mereka sedar akan kesilapan mereka yang selalu mengejek lelaki tua tersebut.) Kemudian, budak-budak nakal itu terus pergi berjumpa dengan lelaki tua itu untuk meminta maaf atas kesilapan yang telah mereka lakukan.

Semenjak hari itu, mereka tidak lagi mengejek lelaki tua tersebut.

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|-------------|-------------------|-------------------|
| didiami 居住 | mempersendakan 作弄 | berdoa 祈祷 |
| mengejek 嘲笑 | keterlaluan 过分 | tebing sungai 河岸边 |



بِسْمِ اللَّهِ الرَّحْمَنِ الرَّحِيمِ

تفسير سورة التين

بجانب ملایو

ترجمه بلغة الملايو

محمد ادريس عبد الرؤوف المروري الأزهرى

من

نسخ التفسير الطبع بين في الرواية والمدنية من علم التفسير

للإمام محمد بن علي بن محمد الشوكاني

حقوق الطبع والنقل محفوظة

طبعة مطبعة دار الفکر

١٣٤٢ هـ - ١٩٢٠ م - ١٩٦٦

Kitab tafsir al-quran pertama dalam bahasa melayu di malaysia. Kitab tafsir al-quran pertama dalam bahasa melayu. Kitab tafsir al-quran pertama dalam bahasa melayu dikenali sebagai. Tafsir al-quran dalam bahasa melayu. Tafsir al quran dalam bahasa melayu pdf.

It first gained national attention as Lou Gehrig's disease, named after the famous baseball player who was diagnosed with ALS in 1939. I want to help people learn about the National ALS Registry, clinical trials, and other resources that can provide support and hope." The National ALS Registry helps gather information from those who are living with this terrible disease. Ed Tessaro was diagnosed with ALS in 2009. It is important to include as many people as possible living with the disease to get the most accurate information. This disease makes the nerve cells stop working and die. And every personal story can help contribute more to fighting this terrible disease. The Registry has found that almost 16,000 persons with ALS live in the United States, as of 2014. The nerves lose the ability to trigger specific muscles, which causes the muscles to become weak and leads to paralysis. In the future, scientists may find that many factors together cause ALS. Below are the latest news from the Registry such as research notifications, new publications, and events. If you or someone you care about has ALS, you can learn more about the National ALS Registry here. New! New ALS Research Notifications: Click here to see clinical trials and epidemiological research studies that are using the Registry to recruit and how you can potentially participate. ALS: Much-Needed Help for a Disease With No Cure Medscape Commentarystatic external icon New! Harrison D, Mehta P, van Es M, Stommel E, Drory V, Nefussy B, van den Berg L, Crayle J, & Bedlack R. To date, the cause of ALS is unknown, and there is still no known cure. Dr. Mehta is the principal investigator leading the National ALS Registry, the only national population-based registry in the U.S. that collects information to help scientists learn more about who gets ALS and what may cause it. We need to help better understand the causes of this terrible disease and empower patients and researchers," states Dr. Paul Mehta, a medical epidemiologist at the Centers for Disease Control and Prevention (CDC)/Agency for Toxic Substances and Disease Registry (ATSDR). Amyotroph Lateral Scler Frontotemporal Degener; 2018; DOI: 10.1080/21678421.2018.1457059. Researchers from all around the world can access the Registry data to help scientists learn more about what causes this disease. "ALS reversals": demographics, disease characteristics, treatments, and co-morbiditiesexternal icon. Almost 16,000 people in the United States were living with ALS in 2014; every year doctors diagnose about 5,000 more people. I'm happy just to stall this thing and get all of us to another chapter in our life." "Place a link to this article on your organization's website." New! Another fatal case of ALS: What can we do? When patients join, it helps give researchers more information. In my case, it's quite literal, but with everything that goes on in my life, I realize that every family has had a crisis." One of the things that helps him and others with ALS is reaching out for community support and resources. ALS is a disease that affects the nerve cells that make muscles work in both the upper and lower parts of the body. Because everyone's diagnosis is personal. Although no cause has been found for most cases of ALS, a number of inherited factors have been found to cause familial ALS. But every person with ALS has an individual story, and understanding these stories will help researchers ultimately piece together clues about this disease. As he learned to cope with this diagnosis, he reflects, "I've never considered myself a victim of the disease because I believe in my heart all of us have a wheelchair. ATSDR scientists have been studying many factors that could be linked with ALS such as heredity and environmental exposures. New! Amyotrophic lateral sclerosis (ALS) is a fatal neurological disease that attacks the nerve cells. The disease strikes quickly, usually leading to death within 2-5 years of diagnosis. This could lead to a better understanding of the causes of ALS, and could help offer a better future for people with ALS in all communities Concludes Ed, "It's tough, but maybe we can fling open a door or two in the next couple of years of research. No one knows what causes most cases of ALS. Everyone's story is different, and everyone's piece of the puzzle is essential. Other scientists have looked at diet or injury, including head trauma from sports. Note, by enrolling in the Registry, you can receive these notifications automatically in your registered email address. (Atlanta, GA) "Losing beloved football player Dwight Clark is just another reminder of how important the National ALS Registry is to support research. Many people know ALS as Lou Gehrig's disease, named after the famous baseball player who got the illness and had to retire in 1939 because of it. "The ongoing struggle is to encourage newly diagnosed people to come in and talk about their condition.

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