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## Rhabdomyolysis: Review of the literature | Request PDF

[https://www.researchgate.net/publication/262526190\\_Rhabdomyolysis\\_Review\\_of\\_the\\_literature](https://www.researchgate.net/publication/262526190_Rhabdomyolysis_Review_of_the_literature)

Background **Duchenne muscular dystrophy (DMD)** is the most common **childhood muscular dystrophy** that anesthesiologists can encounter in the operation room, and patients with **DMD** are **susceptible** to ...

## Design for the sacubitril/valsartan (LCZ696) compared with ...

<https://www.sciencedirect.com/science/article/pii/S0002870317302053>

**Sacubitril/valsartan (LCZ696)** is an **angiotensin receptor neprilysin inhibitor** approved for the treatment of **adult heart failure (HF)**; however, the benefit of **sacubitril/valsartan** in **pediatric HF patients** is unknown.

**Cited by:** 8

**Author:** Robert Shaddy, Charles Canter, Nancy H...

**Publish Year:** 2017

## 2018 ACVIM Forum Research Abstract Program - 2018 ...

<https://onlinelibrary.wiley.com/doi/full/10.1111/jvim.15319>

Oct 25, 2018 · Golden retriever **muscular dystrophy (GRMD)** is a genetically homologous model that has been used increasingly to study pathogenesis and potential treatments for **DMD**. **GRMD** dogs develop **cardiomyopathy** similar to **DMD**, but the disease progression has not been well defined.

## Dilated cardiomyopathy - The Lancet

[https://www.thelancet.com/journals/lancet/article/PIIS0140-6736\(16\)31713-5/fulltext](https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(16)31713-5/fulltext)

Jul 22, 2017 · In addition to Emery-Dreifuss **muscular dystrophy**, **Duchenne muscular dystrophy** and **Becker muscular dystrophy** commonly present in childhood, with a predominant musculoskeletal phenotype, in the setting of **coexistent dilated cardiomyopathy**. These two dystrophies are **caused** by mutations in the **DMD** gene.

**Name of Journal:** *World Journal of Clinical Cases*

**Manuscript NO:** 49766

**Manuscript Type:** CASE REPORT

**Recurrent hypotension induced by sacubitril/valsartan in cardiomyopathy  
secondary to Duchenne muscular dystrophy: A case report**

Li JM *et al.* Sacubitril/ valsartan induced hypotension in DMD cardiomyopathy

Jia-Min Li, Han Chen

**Abstract**

**BACKGROUND**

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Recurrent hypotension induced by sacubitril/valsartan in cardi



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## Diseases | Free Full-Text | A Rare Adverse Event of ...

<https://www.mdpi.com/2079-9721/7/2/38/htm> ▼

In this case report, we describe sacubitril/valsartan as the sole cause of rhabdomyolysis which is a rare adverse drug event. The patient presented with the two cardinal features of rhabdomyolysis namely muscle weakness and dark colored urine which subsequently led to hyperthermia and acute renal failure.

Author: Prashanth Rawla, Jeffrey Pradeep Raj... Publish Year: 2019

## Cardiac Management of the Patient With Duchenne ...

[https://www.researchgate.net/publication/328003270\\_Cardiac\\_Management\\_of\\_the\\_Patient...](https://www.researchgate.net/publication/328003270_Cardiac_Management_of_the_Patient...)

Duchenne muscular dystrophy (DMD) is a genetic, X-linked recessive disease with an associated cardiomyopathy characterized by myocardial fibrosis leading to heart failure, arrhythmias, and death.

## Dilated Cardiomyopathy | Circulation Research

<https://www.ahajournals.org/doi/10.1161/CIRCRESAHA.116.309396>

Less common forms of primary cardiomyopathies are peripartum, tachycardia-induced, stress-provoked Takotsubo cardiomyopathy and myocarditis, according to the 2006 American Heart Association (AHA) definition and classification. 2 Interestingly, myocarditis and peripartum cardiomyopathy can occur in a familial setting and are believed to have a genetic component. 22 – 24 Secondary forms of cardiomyopathies, in which cardiomyopathy ...

## Dilated Cardiomyopathy - Medscape

<https://img.medscape.com/pi/iphone/medscapeapp/html/A152696-business.html> ▼

Dilated cardiomyopathy is characterized by ventricular chamber enlargement and systolic dysfunction with greater left ventricular (LV) cavity size with little or no wall hypertrophy. Hypertrophy can be judged as the ratio of LV mass to cavity size; this ratio is decreased in persons with dilated



4,210 Results

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## Treatment of dystrophin cardiomyopathies

[https://www.researchgate.net/.../259720307\\_Treatment\\_of\\_dystrophin\\_cardiomyopathies](https://www.researchgate.net/.../259720307_Treatment_of_dystrophin_cardiomyopathies)

Treatment of dystrophin cardiomyopathies. ... monitoring approaches in the field of muscular dystrophy and cardiomyopathy. ... transplantation in Duchenne muscular dystrophy: A case report. Article.

Author: Josef Finsterer, Linda Crine