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Untreated Laron syndrome patients have symptomatic and asymptomatic hypoglycemia, but with progressing age some develop relative insulin resistance, glucose intolerance, even diabetes mellitus.

[Laron syndrome - Wikipedia](#)

Laron's syndrome, or Laron-type dwarfism, is an autosomal recessive disorder characterized by an insensitivity to growth hormone (GH), usually caused by a mutant growth hormone receptor.

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Laron syndrome (LS), or primary growth hormone (GH) insensitivity, was first described in 1966. Since then, many patients worldwide have been diagnosed with LS, which involves defects in the GH receptor that cause combined congenital deficiency of GH and IGF-I activities.

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For the 64 patients with Laron syndrome (LS) in Israel, we were able to obtain 34 pedigrees. We performed molecular genetic analysis of the GH receptor in 35 patients and 32 family members. We