Extremely Preterm Severely Small for Gestational Age Newborns Are At Risk For Motor Developmental Impairment at 3 Years of Age.

Postnatal Development of Endogenous Free Fatty Acids and Its Effects on Unbound Bilirubin in Newborn Infants ≥ 37 Weeks Gestational Age.

Incidence of Short Stature at 3 Years of Age in Japanese Late-Preterm Infants.

Association between renin-angiotensin systems gene polymorphisms and intrauterine growth restriction in a Japanese population.

A vascular endothelial growth factor polymorphism is associated with the development of chronic lung disease in Japanese premature newborns.

The relation between steroid responsiveness and immunofluorescence findings in infant idiopathic nephrotic syndrome.

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The carrier frequency in the mothers of 158 Japanese cases with Duchenne/Becker muscular dystrophy.
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Multicenter, Double-Blind, Placebo-Controlled, Randomized Trial of Rituximab for the Treatment of Childhood-Onset Refractory nephrotic syndrome.
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A case of C3 glomerulonephritis in a 6-month-old infant accompanied by glomerular endothelial injury: the expanding spectrum of complement-related kidney diseases.
Natsuki Matsunoshita, Hiroshi Kaito, Shigee Hara, Shingo Ishimori, Takeshi Ninchoji, Kiyoshi Hamahira, Norishige Yoshikawa, Kazumoto Iijima.

Actual outcome and problem of cyclosporine treatment in children with steroid-sensitive nephrotic syndrome.
Takeshi Ninchoji, Hiroshi Kaito, Natsuki Matsunoshita, Shingo Ishimori, Norishige Yoshikawa, Kazumoto Iijima.

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A prostaglandin D2 metabolite is elevated in the urine samples of patients with Duchenne muscular dystrophy.

Three-dimensional gait analysis of Duchenne muscular dystrophy; a trial to evaluate the therapeutic effect of RNA/ENA chimera antisense oligonucleotide that induces dystrophin exon 45 skipping.

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Assessment of Unbound Bilirubin in Newborns After Abdominal Surgery.
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Heterozygous microdeletion of 16q covering SALL1 and RPGRIP1L could be a novel contiguous gene syndrome with renal impairment.

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Natural History and Protein Expression Pattern in Autosomal Recessive Alport Syndrome

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Standard value of urine HVA/VMA ratio in neonates to screen for Menkes disease
Mariko Yagi, Noriko Kusunoki, Tomoko Lee, Ichiro Morioka, Yasuhiro Takeshima

The 3rd Asian Congress for Inherited Metabolic Diseases / The 55th Annual Meeting of The Japanese Society for Inherited Metabolic Diseases Chiba 2013.11.27-29
A Japanese family with citrin deficiency : a daughter with NICCD and a mother without symptoms of CTLN2.
Mariko Yagi, Noriko Kusunoki, Tomoko Lee, Atsuo Kikuchi, Shigeo Kure, Yasuhiro Takeshima.

Outcome of newborn screening for inborn errors of metabolism using tandem mass spectrometry in Hyogo.