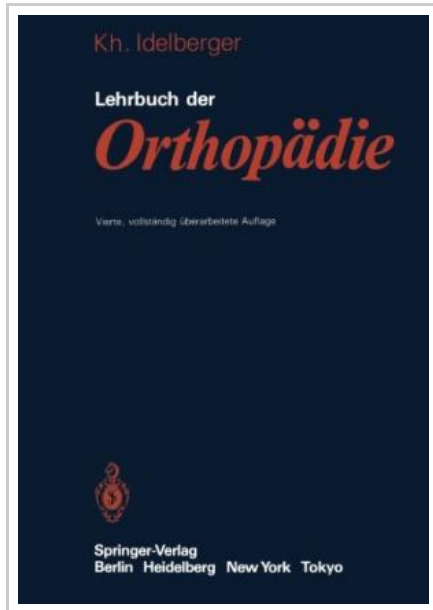




Lehrbuch Der Orthopädie

By K. Idelberger

Springer. Paperback. Book Condition: New. Paperback. 444 pages. Dimensions: 9.6in. x 6.7in. x 1.0in. J. Achondro- und Hypochondroplasie: Erbliche Knorpel- und Knochenwachstumsstörungen, die zu einem dysproportionierten rhizomelen Zwergwuchs führt. Leitsymptome der Achondroplasie: Normal langer Rumpf, kurze Extremitäten, namentlich Oberarm und Oberschenkel, großer Schädel mit eingezogener Nasenwurzel durch Verkrümmung des Os tribasillare, Dreizackhand, aufgetriebene Rippenknochen mit normalen Epiphysen, quadratische Beckenschaukeln, horizontale Pfannendächer, schmale Incisura ischiadica. Leitsymptome der Hypochondroplasie: Normaler Schädel, keine Dreizackhand. Endgröße bis 152 cm gegenüber 135 cm bei der Achondroplasie. 2. Kongenitale spondyloepiphyse Dysplasie: Erbliche, bei der Geburt erkennbare Dysplasie. Leitsymptome: Kurzer Rumpf, kurze Extremitäten, vorspringendes Sternum, Kyphose. Bewegungseinschränkung von Hüft- und Schultergelenken. Endgröße bis 135 cm. 3. Autosomal-dominante Form der metaphysären Chondrodysplasie: Entdeckung erst nach Laufbeginn durch Krümmung der Extremitäten und watschelnden Gang, Coxae varae (in 50 doppelseitig). Endgröße bis 140 cm. 4. Autosomal-rezessive Form der metaphysären Chondrodysplasie: Entdeckung im 2. Lebensjahr durch Minderwuchs. Verbiegungen der langen und kurzen Rippenknochen hängen rachitischen Veränderungen, dünnes, flachschwarzes Haar. Metaphysen im Kniebein reich unregelmäßig in Form und Struktur. Endgröße bis 140 cm. 5. Metaphysäre Chondrodysplasie, Typ Jansen: Selten, schwerer dysproportionierter Zwergwuchs durch verkrümmte Extremitäten.

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