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| 1. Record Nr.           | UNINA9910300235603321  |
| Titolo                  | Craniofacial and Dental Developmental Defects [[electronic resource] ] :<br>Diagnosis and Management // edited by J Timothy Wright   |
| Pubbl/distr/stampa      | Cham : , : Springer International Publishing : , : Imprint : Springer ,<br>2015  |
| ISBN                    | 3-319-13057-9  |
| Edizione                | [1st ed. 2015.]  |
| Descrizione fisica      | 1 online resource (131 p.)   |
| Disciplina              | 599935<br>611.01816<br>617.522059<br>617.6<br>618.92   |
| Soggetti                | Dentistry<br>Oral surgery<br>Maxilla - Surgery<br>Pediatrics<br>Human genetics<br>Oral and Maxillofacial Surgery<br>Human Genetics   |
| Lingua di pubblicazione | Inglese  |
| Formato                 | Materiale a stampa   |
| Livello bibliografico   | Monografia   |
| Note generali           | Description based upon print version of record.  |
| Nota di bibliografia    | Includes bibliographical references at the end of each chapters and<br>index.  |
| Nota di contenuto       | Introduction -- Failure of Tooth Eruption -- Non-Syndromic<br>Hypodontia: Diagnosis and Treatment -- Syndromic Hypodontia --<br>Ectodermal Dysplasias -- Conditions Associated with Premature Tooth<br>Exfoliation -- Defects of Enamel Development -- Defects of Dentin<br>Development -- Managing the Cleft Patient -- Future Directions in<br>Diagnosis and Treatment.  |
| Sommario/riassunto      | This book is intended as a reference that will provide the practitioner<br>with a framework for establishing a diagnosis and developing a suitable<br>treatment plan in patients presenting with a range of developmental<br>defects of the teeth. The conditions covered include failure of tooth<br>eruption, hypodontia, premature tooth exfoliation, defects of enamel |

development, and defects of dentin development, with full consideration of both syndromic and non-syndromic defects. In each case the phenotype and genotype are first described, followed by diagnostic information, including the availability of genetic testing, and treatment options. Summarizing tables are used to highlight the key diagnostic features, and helpful illustrated case presentations are included. Cleft palate is also addressed, with details on etiology, phenotypes, treatment timing and approaches, and dental management. The closing chapter provides stimulating reflections on potential future directions in the diagnosis and treatment of these disorders, encompassing changes in management related to environmental–genetic interactions, tissue engineering, and materials.

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