

Book chapters and review articles:

1. Fassihi H, JA McGrath. Preimplantation genetic diagnosis for inherited skin disorders: from bench to bedside to birth: Thought Leader Commentary. *Dermquest (Research Updates) 2006*
2. Fassihi H, White I. Medicine and Surgery: An Integrated Textbook (Chapter 14: Disease of the Skin). Editor E Lim (Churchill Livingstone) 2007
3. Fassihi H, Mellerio JE, McGrath JA. Prenatal diagnosis for severe genetic skin disorders. *Ichthyosis Support Group Newsletter 2007*
4. Fassihi H, McGrath JA. Life with Epidermolysis Bullosa (Chapter 1.4.2.3: Prenatal and Preimplantation Diagnosis in Epidermolysis Bullosa). Editors Fine and Hinter (Springer Wien New York) 2008
5. Fassihi H. Clinical Editor. Current Medical Literature in Dermatology

Peer review publications:

1. **Fassihi H**, McGrath JA. [Prenatal diagnosis of epidermolysis bullosa.](#) *Dermatol Clin* 2010; 28: 231-7.
2. **Fassihi H**, Liu L, Renwick PJ, Braude PR, McGrath JA. [Development and successful clinical application of preimplantation genetic haplotyping for Herlitz junctional epidermolysis bullosa.](#) *Br J Dermatol* 2010 Jun;162(6):1330-6.
3. Purdie KJ, Pourreyaon C, **Fassihi H**, Cepeda-Valdes R, Frew JW, Volz A, Weissenborn SJ, Pfister H, Proby CM, Bruckner-Tuderman L, Murrell DF, Salas-Alanis JC, McGrath JA, Leigh IM, Harwood CA, South AP. [No Evidence That Human Papillomavirus Is Responsible for the Aggressive Nature of Recessive Dystrophic Epidermolysis Bullosa-Associated Squamous Cell Carcinoma.](#) *J Invest Dermatol* 2010; 130: 2853-5.
4. **Fassihi H**, Iqbal K, Sarkany R, Scarisbrick J, Novaković L. UVA1 phototherapy in the treatment of sclerodermatoid GVHD. *Serbian J Dermatology and Venereology* 2009; 4: 147-152.
5. Arita K, Wessagowit V, Inamadar AC, Palit A, **Fassihi H**, Lai-Cheong JE, Pourreyaon C, South AP, McGrath JA. Unusual clinical and molecular findings in Kindler Syndrome. *Br J Dermatol* 2007; 157: 1252-6.
6. **Fassihi H**, Sarkany R. Vitiligo and phototherapy. *BMJ Minerva* 2007; 334: 1064.
7. Pourreyaon C, Cox G, Mao X, Volz A, Baksh N, Wong T, **Fassihi H**, Arita K, O'Toole EA, Ocampo-Candiani J, Chen M, Hart IR, Bruckner-Tuderman L, Salas-Alanis JC, McGrath JA, Leigh IM, South AP. Patients with Recessive Dystrophic Epidermolysis Bullosa Develop Squamous-Cell Carcinoma Regardless of Type VII Collagen Expression. *J Invest Dermatol* 2007; 127: 2438-44.
8. Renwick PJ, Trussler J, Ostad-Saffari E, **Fassihi H**, Braude P, Ogilvie CM, Abbs S. Proof of principle and first cases using preimplantation genetic haplotyping - a paradigm shift for embryo diagnosis. *Reprod Biomed Online* 2006; 13: 110-9.

9. Lu L, Choy YS, Wessagowit V, Ozoemena L, Dopping-Hepenstal P, **Fassihi H**, McGrath JA. Single nucleotide polymorphism in a commonly utilised *LAMB3* Primer Sequence: Implications for mutation detection and haplotype analysis in junctional epidermolysis bullosa. *J Invest Dermatol* 2006; 44: 48-51.
10. **Fassihi H**, Lu L, Wessagowit V, Ozoemena L, Jones C, Dopping-Hepenstal P, Denyer J, Atherton DJ, Mellerio JE, McGrath JA. Complete maternal isodisomy of chromosome 3 in a child with recessive dystrophic epidermolysis bullosa but no other phenotypic abnormalities. *J Invest Dermatol* 2006; 126: 2039-43.
11. Evans SE, Erkin G, **Fassihi H**, Chan I, Paller AS, Sürücü S, McGrath JA. Ectodermal dysplasia-skin fragility syndrome resulting from a new homozygous mutation, 888delC, in the desmosomal protein, plakophilin 1. *J Am Acad Dermatol* 2006; 55: 157-61.
12. [Burch JM, Fassihi H, Jones CA, Mengshol SC, Fitzpatrick JE, McGrath JA.](#) Kindler syndrome: a new mutation and new diagnostic possibilities. *Arch Dermatol* 2006; 142: 620-4.
13. **Fassihi H**, Renwick PJ, Black C, McGrath JA. Single cell PCR amplification of microsatellites flanking the *COL7A1* gene and suitability for preimplantation genetic diagnosis of Hallopeau-Siemens recessive dystrophic epidermolysis bullosa. [J Dermatol Sci.](#) 2006; 42: 241-8.
14. **Fassihi H**, Eady RAJ, Mellerio JE, Ashton GHS, Dopping-Hepenstal PJC, Denyer J, Nicolaides KH, Rodeck CH, McGrath JA. Prenatal diagnosis for severe inherited skin disorders: 25 years' experience. *Br J Dermatol* 2006; 154: 106-113.
15. **Fassihi H**, Grace J, Lashwood A, Whittock NV, Braude PR, Pickering SJ, McGrath JA. Preimplantation genetic diagnosis of skin fragility-ectodermal dysplasia syndrome: birth of a healthy baby four years after embryo diagnosis and following two frozen embryo replacement cycles. *Br J Dermatol* 2006; 154: 546-550.
16. **Fassihi H**, Wong T, Wessaowit V, McGrath JA, Mellerio JE. Target proteins in inherited and acquired blistering skin disorders. *Clin Exp Dermatol* 2006; 31: 252-259.
17. **Fassihi H**, Wessagowit V, Ashton GHS, Moss C, Ward R, Denyer J, Mellerio JE, McGrath JA. Complete paternal uniparental isodisomy of chromosome 1 resulting in Herlitz junctional epidermolysis bullosa. *Clin Exp Dermatol* 2005; 30: 71-74.
18. **Fassihi H**, Ashton GHS, Denyer J, Mellerio JE, Mason G, McGrath JA. Prenatal diagnosis of Herlitz junctional epidermolysis bullosa in non-identical twins. *Clin Exp Dermatol* 2005; 30: 180-182.
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20. Ozanic Bulic S, **Fassihi H**, Mellerio JE, McGrath JA, Atherton DJ. Thalidomide in the management of epidermolysis bullosa pruriginosa. *Br J Dermatol* 2005; 152: 1332-1334.
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22. **Fassihi H**, Diba V, Jones CA, Dopping-Hepenstal PJC, Burrows N, McGrath JA. Transient bullous dermolysis of the newborn in three generations. *Br J Dermatol* 2005; 153: 1058-1063.
23. Lai Cheong JE, Wessagowit V, **Fassihi H**, McGrath JA. Clinical and molecular aspects of Kindler syndrome. *G Ital Dermatol Venereol* 2005; 140: 531-538.