

## CURRICULUM VITAE ET STUDIORUM

**Name** TERRINONI ALESSANDRO  
**Birth:** Roma, 22-05-1968  
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**Nationality:** Italian

**Languages:** Italian, English

Gender: Male

### **WORK EXPERIENCE**

Present position

Associate Professor at Department of Experimental Medicine, University of Roma "Tor Vergata".

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Previous positions

### **Biographical Sketch**

Alessandro Terrinoni has been, and previously He graduated in Biology at the University of Rome La Sapienza (1993) Science and Human Nutrition at the University of Rome Tor Vergata (2005) and after his PhD at the University of Rome Tor Vergata (2001). Dr Terrinoni spent time working in Dundee (Prof Irwin Mclean), London (prof Martin Raff), University of Leicester, MRC Toxicology Unit Hodgkin Building (prof. Pierluigi Nicotera).

At the scientific level he has worked on cell death, in cancer and dermatological models. It worked on the p53 family, in particular on p63 and p73, of which he identified part of the mechanisms of cell death, of gene transactivation, the biochemical degradation pathways, inhibitors of proteosomal degradation, the physiological regulatory mechanisms (using transgenic mouse models developed in his laboratory). He also identified the biochemical mechanisms of death of keratinocytes in in vitro skin (role of transglutaminase and their substrates) and in vivo, developing transgenic mice and studying human pathologies. These latest research have led to the identification of a new genetic disease and the discovery of the molecular cause of other genetic diseases of the skin. In particular: (1) UNILATERAL PALMOPLANTAR VERRUCOUS NEVUS, new genodermatosis described in OMIM (144200), (2) Marie Unna Hereditary Hypotrichosis (OMIM 146550). He developed vectors for the creation of transgenic animals with MiR expression selectively in the epidermis (MiR24). He developed the system with luciferase for the analysis of the effect of MiR217 on the 3rd minute of Sirt-1.

Currently in the Clinical Biochemistry laboratory he also deals with the role of some MiR in the differentiation of myocytes, colorectal cancer (CRC) and in their analysis in liquid biopsy of CRCpatients.

### **Scientific production**

100 Articles published, 18 First Author, 16 Corresponding Author

Total Citations: 4336 (Scopus)

h-Index: 32 (Scopus)

### **Education**

1993 degree in Biology, with full score.

2001 PhD in Biology and Physiopathology of Epithelia, cum laude, University of Rome "Tor Vergata".

2005 Degree in Human Nutrition Science cum Laude, University of Rome Tor Vergata.

## **Other qualifications**

- National scientific qualification: Full professor of Clinical Biochemistry and Clinical Molecular Biology, Call 2020.
- Biology board qualification (148/150)
- 2<sup>nd</sup> course on apoptosis of the Italian Association for Cell Biology and Differentition. “Gran Sasso” Italy June 3rd to 5th 1998
- Animal models of human Disease: Modelling Human Cancers in the Mouse, a Pratical Issue. European School of Haematology (ESH), Eurolab. Paris, 24-28 January 2002.
- Affymetrix Gene Chip Technology: Sample preparation, Ibridization and Data Mining Tools. IDI-IRCCS, Rome 7 November 2003.

## **Current teaching Activity**

- Degree in Pharmacy (English), University of di Rome Tor Vergata, course of Drug Analysis I, 5 CFU and Clinical Biochemistry, 2CFU
- Degree in Nutrizione Umana, University of di Rome Tor Vergata, course of Clinical Biochemistry 5CFU
- Scuola di Specializzazione in Biochimica e Patologia Clinica. Clinical Biochemistry
- Scuola di Specializzazione in Biochimica e Patologia Clinica Clinical Biochemistry
- Scuola di Specializzazione in Endocrinologia e Malattie del Metabolismo, Clinical Biochemistry
- Board of the PhD program in Molecular Biology, Università di Roma Tor Vergata,

## **Other**

### **Editorial Boards**

Editorial Board for Biochemistry in Encyclopedia of Life Sciences (Wiley)

Editorial Board for Journal of Cytology and Tissue Biology

Editorial Board for International Journal of Molecular Sciences (MDPI)

### **Referee**

Cell Death and Differentiation

Cell Death and Disease

British Journal of Dermatology

Investigative Journal of Dermatology

European Journal of Dermatology

American Journal Of Physiology – Cell Biology

Journal of Molecular Sciences (MDPI)

Nutrients

Others

### **Books**

Alessandro Terrinoni, Southern e Northern Blot. Metodologie Bio-Molecolari nel Nuovo Millennio, Zanichelli 2019

Terrinoni, Alessandro, Melino, Gerry, Serra, Valeria, Alessandrini, Marco, Napolitano, Bianca, and Bruno, Ernesto (Sep 2009) Deafness. In: eLS. John Wiley & Sons Ltd, Chichester. <http://www.els.net> [doi: 10.1002/9780470015902.a0001453.pub2]

Candi, E, McLean, WH, Didona, B, Terrinoni, A, and Melino, G(Dec 2009) Cornification Diseases (Skin Cell Death). In: eLS. John Wiley & Sons Ltd, Chichester. <http://www.els.net> [doi: 10.1002/9780470015902.a0021986]

Melino, G., De Laurenzi, V., Catani, M. V., Terrinoni, A., Ciani, B., Candi, E., Marekov, L. & Steinert, P. M. 1998. The cornified envelope: a model of cell death in the skin. *Results Probl Cell Differ*, 24,

L'involucro corneo un modello di differenziamento e morte cellulare, Eleonora Candi, Alessandro Terrinoni, Gerry Melino, in *Argomenti di biologia molecolare*, SEU, 2006, ISBN: 8889548312 ISBN-13: 9788889548318

### **Grant Scientific Coordinator**

Scientific PI of Ricerca n° 8 - Istituto Dermopatico dell'Immacolata (IDI-IRCCS), afferente al Destinatario Istituzionale n° 1 - Istituto Europeo di Oncologia (IEO), ai fini della realizzazione del progetto “Identificazione di marcatori per la predizione della risposta a nuovi farmaci antitumorali (inibitori di HDAC, tirosina chinasi e pompe ioniche)”

Scientific PI of the following research grant “Ricerca Corrente Italian Ministry of Health”

1. RC-1.1, 2009-2015: “Meccanismi genetici e molecolari che regolano lo sviluppo della cute in condizioni normali e patologiche”.
2. RC-1.3, 2009-2015: “Il differenziamento terminale dell’epidermide: meccanismi molecolari e malattie genetiche correlate”.
3. RC-3.6, 2009-2015: “Ruolo della regolazione trascrizionale e post traduzionale degli elementi della p53 family, nella cancerogenesi e nella resistenza a chemioterapici, in tumori di origine epiteliale. Valutazione di nuovi target terapeutici”.

### **Meetings (speaker)**

Alessandro Terrinoni et al. The role of TGase3 in ultraviolet photodamage. 11th EWCD European Workshops on Cell Death. DEATH NEVER DIES FIUGGI May 6th to 11th 2018

Alessandro Terrinoni et al, Luteolin-7-glucoside inhibits IL-22/STAT3 pathway, reducing proliferation, acanthosis and inflammation in keratinocytes and in mouse psoriatic model. EWCD European Workshops on Cell Death. DEATH NEVER DIES FIUGGI APRIL 3rd to 8th 2016

Palombo R, Porta G, Bruno E, Provero P, Serra V, Neduri K, Viziano A, Alessandrini M, Micarelli A, Ottaviani F, Melino G, Terrinoni A. OTX2 regulates the expression of TAp63 leading to macular and cochlear neuroepithelium development. P63 international, Ricerca e confronti Padova, 24-25 November 2015 (Speaker)

Serra V, Codispoti A, Bruno E, van Bokhoven H, Candi E, Melino G, Terrinoni A (2012) The molecular and physiological mechanism responsible of sensorineural deafness in EEC syndrome: role of the p63 transcription factor. *Journal of Investigative Dermatology* 132:S104-S104.

2nd Annual Meeting of the European-Society-for-Dermatological-Research (ESDR) Location: Venice, ITALY Date: SEP 07-10, 2012 (Speaker)

Alessandro Terrinoni. Involvement of p63 apoptosis/differentiation pathway in cochlear neuroepithelium. and sensorineural deafness. 20th Euroconference on Apoptosis “From Death to Eternity”. (speaker)

Valeria Serra, Alessandro Terrinoni Ernesto Bruno, Hans van Bohkoven. The role of p63 in cochlea development and in sensorineural deafness in EEC syndrome (Speaker). 5th p63/p73 Workshop in Lyon 11-14 September 2011

Terrinoni A, Serra V, van Bohkoven H, Codispoti A, Melino G (2011) The role of p63 in cochlea development and in sensorineural deafness in EEC syndrome. Journal of Investigative Dermatology 131:S67-S67. 41st Annual Meeting of the European-Society-for-Dermatological-Research Location: Barcelona, SPAIN Date: SEP 07-10, 2011

Candi E, Cipollone R, Codispoti A, Melino G, Terrinoni A (2008) Transcriptional study of p63alpha mutants found in ectodermal dysplasia syndromes. Journal of Investigative Dermatology 128:S128-S128. International Investigative Dermatology Meeting Location: Kyoto, JAPAN Date: MAY 12-17, 2008

Terrinoni A, Codispoti A, Paradisi M, Didona B, Melino G (2007) Functional characterization of different Connexin 26 mutants identified in KID Syndrome patients: impairment of membrane trafficking and connexon assembly. Journal of Investigative Dermatology 127:S83-S83. 37th Annual Meeting of the European-Society-for-Dermatological-Research Location: Zurich, SWITZERLAND Date: SEP 05-08, 2007.

Terrinoni A, Didona B, Codispoti A, Zocchi L, Melino G (2006) The novel Connexin 26 Gly11Glu heterozygous mutation is responsible of a case of KID Syndrome, via the impairment of membrane trafficking and connexon assembly. Journal of Investigative Dermatology 126:81-81. 67th Annual Meeting of the Society-for-Investigative-Dermatology Location: Philadelphia, PA Date: MAY 03-06, 2006

## **Selected publications**

1. Rapanotti, M.C.; Cugini, E.; Nuccetelli, M.; Terrinoni, A.; Di Raimondo, C.; Lombardo, P.; Costanza, G.; Cosio, T.; Rossi, P.; Orlandi, A., et al. MCAM/MUC18/CD146 as a Multifaceted Warning Marker of Melanoma Progression in Liquid Biopsy. *Int J Mol Sci* 2021, 22, doi:10.3390/ijms222212416.
2. Caporali, S.; Didona, B.; Paradisi, M.; Mauriello, A.; Campione, E.; Falconi, M.; Iacobelli, F.; Minieri, M.; Pieri, M.; Bernardini, S., et al. Post Zygotic, Somatic, Deletion in KERATIN 1 V1 Domain Generates Structural Alteration of the K1/K10 Dimer, Producing a Monolateral Palmar Epidermolytic Nevus. *Int J Mol Sci* 2021, 22, doi:10.3390/ijms22136901.
3. Caporali, S.; Calabrese, C.; Minieri, M.; Pieri, M.; Tarantino, U.; Marini, M.; D'Ottavio, S.; Angeletti, S.; Mauriello, A.; Cortese, C., et al. The miR-133a, TPM4 and TAp63gamma Role in Myocyte Differentiation Microfilament Remodelling and Colon Cancer Progression. *Int J Mol Sci* 2021, 22, doi:10.3390/ijms22189818.
4. Diluvio, L.; Caporali, S.; Lozzi, F.; Campione, E.; Mazzilli, S.; Lanna, C.; Bianchi, L.; Bernardini, S.; Minieri, M.; Mauriello, A., et al. Birt-Hogg-Dube syndrome, from non-invasive dermatologic assessment to gene testing, molecular and ultrastructural histologic analysis. *J Eur Acad Dermatol Venereol* 2020, 34, e206-e209, doi:10.1111/jdv.16168.
5. Terrinoni, A.; Palombo, R.; Pitolli, C.; Caporali, S.; De Berardinis, R.; Ciccarone, S.; Lanzillotta, A.; Mauramati, S.; Porta, G.; Minieri, M., et al. Role of the TAp63 Isoform in Recurrent Nasal Polyps. *Folia Biol (Praha)* 2019, 65, 170-180.
6. Terrinoni, A.; Calabrese, C.; Basso, D.; Aita, A.; Caporali, S.; Plebani, M.; Bernardini, S. The circulating miRNAs as diagnostic and prognostic markers. *Clin Chem Lab Med* 2019, 57, 932-953, doi:10.1515/cclm-2018-0838.
7. Palombo, R.; Caporali, S.; Falconi, M.; Iacobelli, F.; Morozzo Della Rocca, B.; Lo Surdo, A.; Campione, E.; Candi, E.; Melino, G.; Bernardini, S., et al. Luteolin-7-O-beta-d-Glucoside Inhibits Cellular Energy Production Interacting with HEK2 in Keratinocytes. *Int J Mol Sci* 2019, 20, doi:10.3390/ijms20112689.
8. Terrinoni, A.; Didona, B.; Caporali, S.; Chillemi, G.; Lo Surdo, A.; Paradisi, M.; Annichiarico-Petruzzelli, M.; Candi, E.; Bernardini, S.; Melino, G. Role of the keratin 1 and keratin 10 tails in the pathogenesis of ichthyosis hystrix of Curth Macklin. *PLoS ONE* 2018, 13, e0195792, doi:10.1371/journal.pone.0195792.
9. Frezza, V.; Terrinoni, A.; Pitolli, C.; Mauriello, A.; Melino, G.; Candi, E. Transglutaminase 3 Protects against Photodamage. *J Invest Dermatol* 2017, 137, 1590-1594, doi:10.1016/j.jid.2017.02.982.
10. Palombo, R.; Savini, I.; Avigliano, L.; Madonna, S.; Cavani, A.; Albanesi, C.; Mauriello, A.; Melino, G.; Terrinoni, A. Luteolin-7-glucoside inhibits IL-22/STAT3 pathway, reducing proliferation, acanthosis, and inflammation in keratinocytes and in mouse psoriatic model. *Cell Death Dis* 2016, 7, e2344, doi:10.1038/cddis.2016.201.
11. Palombo, R.; Porta, G.; Bruno, E.; Provero, P.; Serra, V.; Neduri, K.; Viziano, A.; Alessandrini, M.; Micarelli, A.; Ottaviani, F., et al. OTX2 regulates the expression of TAp63 leading to macular and cochlear neuroepithelium development. *Aging (Albany NY)* 2015, 7, 928-936, doi:10.18632/aging.100839.
12. Terrinoni, A.; Serra, V.; Bruno, E.; Strasser, A.; Valente, E.; Flores, E.R.; van Bokhoven, H.; Lu, X.; Knight, R.A.; Melino, G. Role of p63 and the Notch pathway in cochlea development and sensorineural deafness. *Proc Natl Acad Sci U S A* 2013, 110, 7300-7305, doi:10.1073/pnas.1214498110.
13. Tucci, P.; Agostini, M.; Grespi, F.; Markert, E.K.; Terrinoni, A.; Vousden, K.H.; Muller, P.A.; Dotsch, V.; Kehrloesser, S.; Sayan, B.S., et al. Loss of p63 and its microRNA-205 target results in enhanced cell migration and metastasis in prostate cancer. *Proc Natl Acad Sci U S A* 2012, 109, 15312-15317, doi:10.1073/pnas.1110977110.
14. Amelio, I.; Lena, A.M.; Viticchie, G.; Shalom-Feuerstein, R.; Terrinoni, A.; Dinsdale, D.;

- Russo, G.; Fortunato, C.; Bonanno, E.; Spagnoli, L.G., et al. miR-24 triggers epidermal differentiation by controlling actin adhesion and cell migration. *J Cell Biol* 2012, 199, 347-363, doi:10.1083/jcb.201203134.
15. Terrinoni, A.; Pagani, I.S.; Zucchi, I.; Chiaravalli, A.M.; Serra, V.; Rovera, F.; Sirchia, S.; Dionigi, G.; Miozzo, M.; Frattini, A., et al. OTX1 expression in breast cancer is regulated by p53. *Oncogene* 2011, 30, 3096-3103, doi:10.1038/onc.2011.31.
  16. Terrinoni, A.; Codispoti, A.; Serra, V.; Didona, B.; Bruno, E.; Nistico, R.; Giustizieri, M.; Alessandrini, M.; Campione, E.; Melino, G. Connexin 26 (GJB2) mutations, causing KID Syndrome, are associated with cell death due to calcium gating deregulation. *Biochem Biophys Res Commun* 2010, 394, 909-914, doi:10.1016/j.bbrc.2010.03.073.
  17. Campione, E.; Terrinoni, A.; Orlandi, A.; Codispoti, A.; Melino, G.; Bianchi, L.; Mazzotta, A.; Garaci, F.G.; Ludovici, A.; Chimenti, S. Cerebral cavernomas in a family with multiple cutaneous and uterine leiomyomas associated with a new mutation in the fumarate hydratase gene. *J Invest Dermatol* 2007, 127, 2271-2273, doi:10.1038/sj.jid.5700851.
  18. Gressner, O.; Schilling, T.; Lorenz, K.; Schulze Schleithoff, E.; Koch, A.; Schulze-Bergkamen, H.; Lena, A.M.; Candi, E.; Terrinoni, A.; Catani, M.V., et al. TAp63alpha induces apoptosis by activating signaling via death receptors and mitochondria. *EMBO J* 2005, 24, 2458-2471, doi:10.1038/sj.emboj.7600708.
  19. Terrinoni, A.; Ranalli, M.; Cadot, B.; Leta, A.; Bagetta, G.; Vousden, K.H.; Melino, G. p73-alpha is capable of inducing scotin and ER stress. *Oncogene* 2004, 23, 3721-3725, doi:10.1038/sj.onc.1207342.
  20. Billon, N.; Terrinoni, A.; Jolicoeur, C.; McCarthy, A.; Richardson, W.D.; Melino, G.; Raff, M. Roles for p53 and p73 during oligodendrocyte development. *Development (Cambridge)* 2004, 131, 1211-1220, doi:10.1242/dev.01035.

## TOTAL PUBBLICATIONS

[1-100]

1. Rapanotti, M.C.; Cugini, E.; Nuccetelli, M.; Terrinoni, A.; Di Raimondo, C.; Lombardo, P.; Costanza, G.; Cosio, T.; Rossi, P.; Orlandi, A., et al. MCAM/MUC18/CD146 as a Multifaceted Warning Marker of Melanoma Progression in Liquid Biopsy. *Int J Mol Sci* 2021, 22, doi:10.3390/ijms222212416.
2. Pieri, M.; Pignalosa, S.; Perrone, M.A.; Russo, C.; Noce, G.; Perrone, A.; Terrinoni, A.; Massoud, R.; Bernardini, S. Evaluation of the Diesse Cube 30 touch erythrocyte sedimentation method in comparison with Alifax test 1 and the manual Westergren gold standard method. *Scand J Clin Lab Invest* 2021, 81, 181-186, doi:10.1080/00365513.2021.1881996.
3. Perna, A.; Passiatore, M.; Massaro, A.; Terrinoni, A.; Bianchi, L.; Cilli, V.; D'orio, M.; Proietti, L.; Taccardo, G.; De Vitis, R. Skin manifestations in COVID-19 patients, state of the art. A systematic review. *Int J Dermatol* 2021, 60, 547-553, doi:10.1111/ijd.15414.
4. Nicolai, E.; Nuccetelli, M.; Sarubbi, S.; Basile, V.; Perrone, M.A.; Terrinoni, A.; Minieri, M.; Pieri, M.; Bernardini, S. Performance evaluation of the new Chemiluminescence Immunoassay CL-1200i Thyroid Panel. *J Immunoassay Immunochem* 2021, 10.1080/15321819.2021.2017301, 1-13, doi:10.1080/15321819.2021.2017301.
5. Minieri, M.; Leoni, B.D.; Bellincampi, L.; Bajo, D.; Agnoli, A.; De Angelis, A.M.; Pieri, M.; Equitani, F.; Rossi, V.; Valente, F., et al. Serum iPTH range in a reference population: From an integrated approach to vitamin D prevalence impact evaluation. *Clin Chim Acta* 2021, 521, 1-8, doi:10.1016/j.cca.2021.06.004.
6. De Stefano, A.; Caporali, S.; Di Daniele, N.; Rovella, V.; Cardillo, C.; Schinzari, F.; Minieri, M.; Pieri, M.; Candi, E.; Bernardini, S., et al. Anti-Inflammatory and Proliferative

- Properties of Luteolin-7-O-Glucoside. *Int J Mol Sci* **2021**, *22*, 1-19, doi:10.3390/ijms22031321.
- 7. Caporali, S.; Didona, B.; Paradisi, M.; Mauriello, A.; Campione, E.; Falconi, M.; Iacovelli, F.; Minieri, M.; Pieri, M.; Bernardini, S., et al. Post Zygotic, Somatic, Deletion in KERATIN 1 V1 Domain Generates Structural Alteration of the K1/K10 Dimer, Producing a Monolateral Palmar Epidermolytic Nevus. *Int J Mol Sci* **2021**, *22*, doi:10.3390/ijms22136901.
  - 8. Caporali, S.; Calabrese, C.; Minieri, M.; Pieri, M.; Tarantino, U.; Marini, M.; D'Ottavio, S.; Angeletti, S.; Mauriello, A.; Cortese, C., et al. The miR-133a, TPM4 and TAp63gamma Role in Myocyte Differentiation Microfilament Remodelling and Colon Cancer Progression. *Int J Mol Sci* **2021**, *22*, doi:10.3390/ijms22189818.
  - 9. Campione, E.; Lanna, C.; Cosio, T.; Rosa, L.; Conte, M.P.; Iacovelli, F.; Romeo, A.; Falconi, M.; Del Vecchio, C.; Franchin, E., et al. Lactoferrin as Antiviral Treatment in COVID-19 Management: Preliminary Evidence. *Int J Environ Res Public Health* **2021**, *18*, doi:10.3390/ijerph182010985.
  - 10. Campione, E.; Lanna, C.; Cosio, T.; Rosa, L.; Conte, M.P.; Iacovelli, F.; Romeo, A.; Falconi, M.; Del Vecchio, C.; Franchin, E., et al. Lactoferrin Against SARS-CoV-2: In Vitro and In Silico Evidences. *Front Pharmacol* **2021**, *12*, 666600, doi:10.3389/fphar.2021.666600.
  - 11. Spoto, S.; Legramante, J.M.; Minieri, M.; Fogolari, M.; Terrinoni, A.; Valeriani, E.; Sebastiani, C.; Bernardini, S.; Ciccozzi, M.; Angeletti, P.S. How biomarkers can improve pneumonia diagnosis and prognosis: procalcitonin and mid-regional-pro-adrenomedullin. *Biomark Med* **2020**, *14*, 549-562, doi:10.2217/bmm-2019-0414.
  - 12. Rapanotti, M.C.; Viguria, T.M.S.; Spallone, G.; Terrinoni, A.; Rossi, P.; Costanza, G.; Campione, E.; Lombardo, P.; Pathirannehalage, C.D.; Orlandi, A., et al. Minimal Residual Disease in Melanoma:molecular characterization of in transit cutaneous metastases and Circulating Melanoma Cells recognizes an expression panel potentially related to disease progression. *Cancer Treat Res Commun* **2020**, *25*, 100262, doi:10.1016/j.ctarc.2020.100262.
  - 13. Perrone, M.A.; Viola, F.G.; Minieri, M.; Caporali, S.; Copponi, A.; Sancesario, G.; Angeletti, S.; Massoud, R.; Romeo, F.; Bernardini, S., et al. The Von Willebrand Factor Antigen Plasma Concentration: a Monitoring Marker in the Treatment of Aortic and Mitral Valve Diseases. *Folia Biol (Praha)* **2020**, *66*, 133-141.
  - 14. Diluvio, L.; Caporali, S.; Lozzi, F.; Campione, E.; Mazzilli, S.; Lanna, C.; Bianchi, L.; Bernardini, S.; Minieri, M.; Mauriello, A., et al. Birt-Hogg-Dube syndrome, from non-invasive dermatologic assessment to gene testing, molecular and ultrastructural histologic analysis. *J Eur Acad Dermatol Venereol* **2020**, *34*, e206-e209, doi:10.1111/jdv.16168.
  - 15. Ciotti, M.; Ciccozzi, M.; Terrinoni, A.; Jiang, W.C.; Wang, C.B.; Bernardini, S. The COVID-19 pandemic. *Crit Rev Clin Lab Sci* **2020**, *57*, 365-388, doi:10.1080/10408363.2020.1783198.
  - 16. Terrinoni, A.; Palombo, R.; Pitelli, C.; Caporali, S.; De Berardinis, R.; Ciccarone, S.; Lanzillotta, A.; Mauramati, S.; Porta, G.; Minieri, M., et al. Role of the TAp63 Isoform in Recurrent Nasal Polyps. *Folia Biol (Praha)* **2019**, *65*, 170-180.
  - 17. Terrinoni, A.; Calabrese, C.; Basso, D.; Aita, A.; Caporali, S.; Plebani, M.; Bernardini, S. The circulating miRNAs as diagnostic and prognostic markers. *Clin Chem Lab Med* **2019**, *57*, 932-953, doi:10.1515/cclm-2018-0838.
  - 18. Spoto, S.; Fogolari, M.; De Florio, L.; Minieri, M.; Vicino, G.; Legramante, J.; Lia, M.S.; Terrinoni, A.; Caputo, D.; Costantino, S., et al. Procalcitonin and MR-proAdrenomedullin combination in the etiological diagnosis and prognosis of sepsis and septic shock. *Microb Pathog* **2019**, *137*, 103763, doi:10.1016/j.micpath.2019.103763.
  - 19. Palombo, R.; Caporali, S.; Falconi, M.; Iacovelli, F.; Morozzo Della Rocca, B.; Lo Surdo, A.; Campione, E.; Candi, E.; Melino, G.; Bernardini, S., et al. Luteolin-7-O-beta-d-

- Glucoside Inhibits Cellular Energy Production Interacting with HEK2 in Keratinocytes. *Int J Mol Sci* **2019**, *20*, doi:10.3390/ijms20112689.
20. Lozzi, F.; Lanna, C.; Mazzeo, M.; Garofalo, V.; Palumbo, V.; Mazzilli, S.; Diluvio, L.; Terrinoni, A.; Bianchi, L.; Campione, E. Investigational drugs currently in phase II clinical trials for actinic keratosis. *Expert Opin Investig Drugs* **2019**, *28*, 629–642, doi:10.1080/13543784.2019.1636030.
21. Terrinoni, A.; Didona, B.; Caporali, S.; Chillemi, G.; Lo Surdo, A.; Paradisi, M.; Annichiarico-Petruzzelli, M.; Candi, E.; Bernardini, S.; Melino, G. Role of the keratin 1 and keratin 10 tails in the pathogenesis of ichthyosis hystrix of Curth Macklin. *PLoS ONE* **2018**, *13*, e0195792, doi:10.1371/journal.pone.0195792.
22. Pitolli, C.; Pietroni, V.; Marekov, L.; Terrinoni, A.; Yamanishi, K.; Mazzanti, C.; Melino, G.; Candi, E. Characterization of TG2 and TG1-TG2 double knock-out mouse epidermis. *Amino Acids* **2017**, *49*, 635–642, doi:10.1007/s00726-016-2356-3.
23. Frezza, V.; Terrinoni, A.; Pitolli, C.; Mauriello, A.; Melino, G.; Candi, E. Transglutaminase 3 Protects against Photodamage. *J Invest Dermatol* **2017**, *137*, 1590–1594, doi:10.1016/j.jid.2017.02.982.
24. Palombo, R.; Savini, I.; Avigliano, L.; Madonna, S.; Cavani, A.; Albanesi, C.; Mauriello, A.; Melino, G.; Terrinoni, A. Luteolin-7-glucoside inhibits IL-22/STAT3 pathway, reducing proliferation, acanthosis, and inflammation in keratinocytes and in mouse psoriatic model. *Cell Death Dis* **2016**, *7*, e2344, doi:10.1038/cddis.2016.201.
25. Palombo, R.; Giannella, E.; Didona, B.; Annichiarico-Petruzzelli, M.; Melino, G.; Terrinoni, A. Cutaneous mosaicism, in KRT1 pI479T patient, caused by the somatic loss of the wild-type allele, leads to the increase in local severity of the disease. *J Eur Acad Dermatol Venereol* **2016**, *30*, 847–851, doi:10.1111/jdv.13153.
26. Palombo, R.; Porta, G.; Bruno, E.; Provero, P.; Serra, V.; Neduri, K.; Viziano, A.; Alessandrini, M.; Micarelli, A.; Ottaviani, F., et al. OTX2 regulates the expression of TAp63 leading to macular and cochlear neuroepithelium development. *Aging (Albany NY)* **2015**, *7*, 928–936, doi:10.18632/aging.100839.
27. Giamboi-Miraglia, A.; Cianfarani, F.; Cattani, C.; Lena, A.M.; Serra, V.; Campione, E.; Terrinoni, A.; Zambruno, G.; Odorisio, T.; Di Daniele, N., et al. The E3 ligase Itch knockout mice show hyperproliferation and wound healing alteration. *FEBS J* **2015**, *282*, 4435–4449, doi:10.1111/febs.13514.
28. Campione, E.; Paterno, E.J.; Candi, E.; Falconi, M.; Costanza, G.; Diluvio, L.; Terrinoni, A.; Bianchi, L.; Orlandi, A. The relevance of piroxicam for the prevention and treatment of nonmelanoma skin cancer and its precursors. *Drug Des Devel Ther* **2015**, *9*, 5843–5850, doi:10.2147/DDDT.S84849.
29. Terrinoni, A.; Giardina, E.; Pertusi, G.; Cascella, R.; Serra, V.; Bornacina, C.; Palombo, R.; Tiberio, R.; Gattoni, M.; Novelli, G., et al. Absence of filaggrin mutation in a patient affected by pachyonychia congenita and mild atopic dermatitis. *Eur J Dermatol* **2014**, *24*, 703–704, doi:10.1684/ejd.2014.2446.
30. Diluvio, L.; Torti, C.; Terrinoni, A.; Candi, E.; Piancatelli, R.; Piccione, E.; Paterno, E.J.; Chimenti, S.; Orlandi, A.; Campione, E., et al. Dermoscopy as an adjuvant tool for detecting skin leiomyomas in patient with uterine fibroids and cerebral cavernomas. *BMC Dermatol* **2014**, *14*, 7, doi:10.1186/1471-5945-14-7.
31. Campione, E.; Diluvio, L.; Terrinoni, A.; Orlandi, A.; Latino, M.P.; Torti, C.; Pietroleonardo, L.; Botti, E.; Chimenti, S.; Bianchi, L. Severe erythrodermic psoriasis in child twins: from clinical-pathological diagnosis to treatment of choice through genetic analyses: two case reports. *BMC Res Notes* **2014**, *7*, 929, doi:10.1186/1756-0500-7-929.
32. Terrinoni, A.; Serra, V.; Bruno, E.; Strasser, A.; Valente, E.; Flores, E.R.; van Bokhoven, H.; Lu, X.; Knight, R.A.; Melino, G. Role of p63 and the Notch pathway in cochlea development and sensorineural deafness. *Proc Natl Acad Sci U S A* **2013**, *110*, 7300–7305, doi:10.1073/pnas.1214498110.

33. Giacobbe, A.; Bongiorno-Borbone, L.; Bernassola, F.; Terrinoni, A.; Markert, E.K.; Levine, A.J.; Feng, Z.; Agostini, M.; Zolla, L.; Agro, A.F., et al. p63 regulates glutaminase 2 expression. *Cell Cycle* **2013**, *12*, 1395-1405, doi:10.4161/cc.24478.
34. Conforti, F.; Yang, A.L.; Piro, M.C.; Mellone, M.; Terrinoni, A.; Candi, E.; Tucci, P.; Thomas, G.J.; Knight, R.A.; Melino, G., et al. PIR2/Rnf144B regulates epithelial homeostasis by mediating degradation of p21WAF1 and p63. *Oncogene* **2013**, *32*, 4758-4765, doi:10.1038/onc.2012.497.
35. Campione, E.; Diluvio, L.; Terrinoni, A.; Di Stefani, A.; Orlandi, A.; Chimenti, S.; Bianchi, L. Progressive late-onset of cutaneous angiomas as possible sign of cerebral cavernous malformations. *Dermatol. Online J.* **2013**, *19*.
36. Yis, U.; Terrinoni, A. Sjogren-Larsson syndrome: report of monozygote twins and a case with a novel mutation. *Turk J Pediatr* **2012**, *54*, 64-66.
37. Tucci, P.; Agostini, M.; Grespi, F.; Markert, E.K.; Terrinoni, A.; Vousden, K.H.; Muller, P.A.; Dotsch, V.; Kehrloesser, S.; Sayan, B.S., et al. Loss of p63 and its microRNA-205 target results in enhanced cell migration and metastasis in prostate cancer. *Proc Natl Acad Sci U S A* **2012**, *109*, 15312-15317, doi:10.1073/pnas.1110977109.
38. Terrinoni, A.; Serra, V.; Codispoti, A.; Talamonti, E.; Bui, L.; Palombo, R.; Sette, M.; Campione, E.; Didona, B.; Annicchiarico-Petruzzelli, M., et al. Novel transglutaminase 1 mutations in patients affected by lamellar ichthyosis. *Cell Death Dis* **2012**, *3*, e416, doi:10.1038/cddis.2012.152.
39. Amelio, I.; Lena, A.M.; Viticchie, G.; Shalom-Feuerstein, R.; Terrinoni, A.; Dinsdale, D.; Russo, G.; Fortunato, C.; Bonanno, E.; Spagnoli, L.G., et al. miR-24 triggers epidermal differentiation by controlling actin adhesion and cell migration. *J Cell Biol* **2012**, *199*, 347-363, doi:10.1083/jcb.201203134.
40. Vernole, P.; Muzi, A.; Volpi, A.; Terrinoni, A.; Dorio, A.S.; Tentori, L.; Shah, G.M.; Graziani, G. Common fragile sites in colon cancer cell lines: role of mismatch repair, RAD51 and poly(ADP-ribose) polymerase-1. *Mutat Res* **2011**, *712*, 40-48, doi:10.1016/j.mrfmmm.2011.03.00610.1016/j.mrfmmm.2011.04.006.
41. Terrinoni, A.; Pagani, I.S.; Zucchi, I.; Chiaravalli, A.M.; Serra, V.; Rovera, F.; Sirchia, S.; Dionigi, G.; Miozzo, M.; Frattini, A., et al. OTX1 expression in breast cancer is regulated by p53. *Oncogene* **2011**, *30*, 3096-3103, doi:10.1038/onc.2011.31.
42. Serra, V.; Castori, M.; Paradisi, M.; Bui, L.; Melino, G.; Terrinoni, A. Functional characterization of a novel TP63 mutation in a family with overlapping features of Rapp-Hodgkin/AEC/ADULT syndromes. *Am J Med Genet A* **2011**, *155A*, 3104-3109, doi:10.1002/ajmg.a.34335.
43. Rufini, S.; Lena, A.M.; Cadot, B.; Mele, S.; Amelio, I.; Terrinoni, A.; Desideri, A.; Melino, G.; Candi, E. The sterile alpha-motif (SAM) domain of p63 binds in vitro monoasialoganglioside (GM1) micelles. *Biochem Pharmacol* **2011**, *82*, 1262-1268, doi:10.1016/j.bcp.2011.07.087.
44. Browne, G.; Cipollone, R.; Lena, A.M.; Serra, V.; Zhou, H.; van Bokhoven, H.; Dotsch, V.; Merico, D.; Mantovani, R.; Terrinoni, A., et al. Differential altered stability and transcriptional activity of DeltaNp63 mutants in distinct ectodermal dysplasias. *J Cell Sci* **2011**, *124*, 2200-2207, doi:10.1242/jcs.079327.
45. Terrinoni, A.; Codispoti, A.; Serra, V.; Didona, B.; Bruno, E.; Nistico, R.; Giustizieri, M.; Alessandrini, M.; Campione, E.; Melino, G. Connexin 26 (GJB2) mutations, causing KID Syndrome, are associated with cell death due to calcium gating deregulation. *Biochem Biophys Res Commun* **2010**, *394*, 909-914, doi:10.1016/j.bbrc.2010.03.073.
46. Terrinoni, A.; Codispoti, A.; Serra, V.; Bruno, E.; Didona, B.; Paradisi, M.; Nistico, S.; Campione, E.; Napolitano, B.; Diluvio, L., et al. Connexin 26 (GJB2) mutations as a cause of the KID syndrome with hearing loss. *Biochem Biophys Res Commun* **2010**, *395*, 25-30, doi:10.1016/j.bbrc.2010.03.098.

47. Pagani, I.S.; Terrinoni, A.; Marenghi, L.; Zucchi, I.; Chiaravalli, A.M.; Serra, V.; Rovera, F.; Sirchia, S.; Dionigi, G.; Miozzo, M., et al. The mammary gland and the homeobox gene Otx1. *Breast J* **2010**, *16 Suppl 1*, S53-56, doi:10.1111/j.1524-4741.2010.01006.x.
48. Keller, M.A.; Watschinger, K.; Golderer, G.; Maglione, M.; Sarg, B.; Lindner, H.H.; Werner-Felmayer, G.; Terrinoni, A.; Wanders, R.J.; Werner, E.R. Monitoring of fatty aldehyde dehydrogenase by formation of pyrenedecanoic acid from pyrenedecanal. *J Lipid Res* **2010**, *51*, 1554-1559, doi:10.1194/jlr.D002220.
49. Wen, Y.; Liu, Y.; Xu, Y.; Zhao, Y.; Hua, R.; Wang, K.; Sun, M.; Li, Y.; Yang, S.; Zhang, X.J., et al. Loss-of-function mutations of an inhibitory upstream ORF in the human hairless transcript cause Marie Unna hereditary hypotrichosis. *Nat Genet* **2009**, *41*, 228-233, doi:10.1038/ng.276.
50. Vernole, P.; Muzi, A.; Volpi, A.; Dorio, A.S.; Terrinoni, A.; Shah, G.M.; Graziani, G. Inhibition of homologous recombination by treatment with BVDU (brivudin) or by RAD51 silencing increases chromosomal damage induced by bleomycin in mismatch repair-deficient tumour cells. *Mutat Res* **2009**, *664*, 39-47, doi:10.1016/j.mrfmmm.2009.02.005.
51. Menghini, R.; Casagrande, V.; Cardellini, M.; Martelli, E.; Terrinoni, A.; Amati, F.; Vasa-Nicotera, M.; Ippoliti, A.; Novelli, G.; Melino, G., et al. MicroRNA 217 modulates endothelial cell senescence via silent information regulator 1. *Circulation* **2009**, *120*, 1524-1532, doi:10.1161/CIRCULATIONAHA.109.864629.
52. Diluvio, L.; Campione, E.; Paterno, E.J.; Orlandi, A.; Terrinoni, A.; Chimenti, S. Peculiar clinical and dermoscopic remission pattern following imiquimod therapy of basal cell carcinoma in seborrhoeic areas of the face. *J Dermatolog Treat* **2009**, *20*, 124-129, doi:10.1080/09546630802441226.
53. Codispoti, A.; Colombo, E.; Zocchi, L.; Serra, V.; Pertusi, G.; Leigheb, G.; Tiberio, R.; Bornacina, G.; Zuccoli, R.; Ramponi, A., et al. Knuckle pads, in an epidermal palmoplantar keratoderma patient with Keratin 9 R163W transgrediens expression. *Eur J Dermatol* **2009**, *19*, 114-118, doi:10.1684/ejd.2008.0575.
54. Campione, E.; Diluvio, L.; Paterno, E.J.; Di Marcantonio, D.; Francesconi, A.; Terrinoni, A.; Orlandi, A.; Chimenti, S. Kaposi's sarcoma in a patient treated with imatinib mesylate for chronic myeloid leukemia. *Clin Ther* **2009**, *31*, 2565-2569, doi:10.1016/j.clinthera.2009.11.018.
55. Zocchi, L.; Bourdon, J.C.; Codispoti, A.; Knight, R.; Lane, D.P.; Melino, G.; Terrinoni, A. Scotin: A new p63 target gene expressed during epidermal differentiation. *Biochem Biophys Res Commun* **2008**, *367*, 271-276, doi:10.1016/j.bbrc.2007.12.115.
56. Zocchi, L.; Terrinoni, A.; Candi, E.; Ahvazi, B.; Bagetta, G.; Corasaniti, M.T.; Lena, A.M.; Melino, G. Identification of transglutaminase 3 splicing isoforms. *J Invest Dermatol* **2007**, *127*, 1791-1794, doi:10.1038/sj.jid.5700768.
57. Porzio, O.; Massa, O.; Cunsolo, V.; Colombo, C.; Malaponti, M.; Bertuzzi, F.; Hansen, T.; Johansen, A.; Pedersen, O.; Meschi, F., et al. Missense mutations in the TGM2 gene encoding transglutaminase 2 are found in patients with early-onset type 2 diabetes. Mutation in brief no. 982. Online. *Hum Mutat* **2007**, *28*, 1150, doi:10.1002/humu.9511.
58. Nucci, C.; Gasperi, V.; Tartaglione, R.; Cerulli, A.; Terrinoni, A.; Bari, M.; De Simone, C.; Agro, A.F.; Morrone, L.A.; Corasaniti, M.T., et al. Involvement of the endocannabinoid system in retinal damage after high intraocular pressure-induced ischemia in rats. *Invest Ophthalmol Vis Sci* **2007**, *48*, 2997-3004, doi:10.1167/iovs.06-1355.
59. Didona, B.; Codispoti, A.; Bertini, E.; Rizzo, W.B.; Carney, G.; Zambruno, G.; Dionisi-Vici, C.; Paradisi, M.; Pedicelli, C.; Melino, G., et al. Novel and recurrent ALDH3A2 mutations in Italian patients with Sjogren-Larsson syndrome. *J Hum Genet* **2007**, *52*, 865-870, doi:10.1007/s10038-007-0180-z.
60. Candi, E.; Rufini, A.; Terrinoni, A.; Giambò-Miraglia, A.; Lena, A.M.; Mantovani, R.; Knight, R.; Melino, G. DeltaNp63 regulates thymic development through enhanced

- expression of FgfR2 and Jag2. *Proc Natl Acad Sci U S A* **2007**, *104*, 11999-12004, doi:10.1073/pnas.0703458104.
61. Campione, E.; Terrinoni, A.; Orlandi, A.; Codispoti, A.; Melino, G.; Bianchi, L.; Mazzotta, A.; Garaci, F.G.; Ludovici, A.; Chimenti, S. Cerebral cavernomas in a family with multiple cutaneous and uterine leiomyomas associated with a new mutation in the fumarate hydratase gene. *J Invest Dermatol* **2007**, *127*, 2271-2273, doi:10.1038/sj.jid.5700851.
62. Ponassi, R.; Terrinoni, A.; Chikh, A.; Rufini, A.; Lena, A.M.; Sayan, B.S.; Melino, G.; Candi, E. p63 and p73, members of the p53 gene family, transactivate PKCdelta. *Biochem Pharmacol* **2006**, *72*, 1417-1422, doi:10.1016/j.bcp.2006.07.031.
63. Candi, E.; Terrinoni, A.; Rufini, A.; Chikh, A.; Lena, A.M.; Suzuki, Y.; Sayan, B.S.; Knight, R.A.; Melino, G. p63 is upstream of IKK alpha in epidermal development. *J Cell Sci* **2006**, *119*, 4617-4622, doi:10.1242/jcs.03265.
64. Candi, E.; Rufini, A.; Terrinoni, A.; Dinsdale, D.; Ranalli, M.; Paradisi, A.; De Laurenzi, V.; Spagnoli, L.G.; Catani, M.V.; Ramadan, S., et al. Differential roles of p63 isoforms in epidermal development: selective genetic complementation in p63 null mice. *Cell Death Differ* **2006**, *13*, 1037-1047, doi:10.1038/sj.cdd.4401926.
65. Barcaroli, D.; Bongiorno-Borbone, L.; Terrinoni, A.; Hofmann, T.G.; Rossi, M.; Knight, R.A.; Matera, A.G.; Melino, G.; De Laurenzi, V. FLASH is required for histone transcription and S-phase progression. *Proc Natl Acad Sci U S A* **2006**, *103*, 14808-14812, doi:10.1073/pnas.0604227103.
66. van Steensel, M.A.; van Geel, M. Does recessive EKV exist? *J Invest Dermatol* **2005**, *124*, 268-269; author reply 270-261, doi:10.1111/j.0022-202X.2004.23517.x.
67. Ramadan, S.; Terrinoni, A.; Catani, M.V.; Sayan, A.E.; Knight, R.A.; Mueller, M.; Krammer, P.H.; Melino, G.; Candi, E. p73 induces apoptosis by different mechanisms. *Biochem Biophys Res Commun* **2005**, *331*, 713-717, doi:10.1016/j.bbrc.2005.03.156.
68. Gressner, O.; Schilling, T.; Lorenz, K.; Schulze Schleithoff, E.; Koch, A.; Schulze-Bergkamen, H.; Lena, A.M.; Candi, E.; Terrinoni, A.; Catani, M.V., et al. TA $\beta$ p63alpha induces apoptosis by activating signaling via death receptors and mitochondria. *EMBO J* **2005**, *24*, 2458-2471, doi:10.1038/sj.emboj.7600708.
69. Cassidy, A.J.; van Steensel, M.A.; Steijlen, P.M.; van Geel, M.; van der Velden, J.; Morley, S.M.; Terrinoni, A.; Melino, G.; Candi, E.; McLean, W.H. A homozygous missense mutation in TGM5 abolishes epidermal transglutaminase 5 activity and causes acral peeling skin syndrome. *Am J Hum Genet* **2005**, *77*, 909-917, doi:10.1086/497707.
70. Terron-Kwiatkowski, A.; Terrinoni, A.; Didona, B.; Melino, G.; Atherton, D.J.; Irvine, A.D.; McLean, W.H. Atypical epidermolytic palmoplantar keratoderma presentation associated with a mutation in the keratin 1 gene. *Br J Dermatol* **2004**, *150*, 1096-1103, doi:10.1111/j.1365-2133.2004.05967.x.
71. Terrinoni, A.; Ranalli, M.; Cadot, B.; Leta, A.; Bagetta, G.; Vousden, K.H.; Melino, G. p73-alpha is capable of inducing scotin and ER stress. *Oncogene* **2004**, *23*, 3721-3725, doi:10.1038/sj.onc.1207342.
72. Terrinoni, A.; Leta, A.; Pedicelli, C.; Candi, E.; Ranalli, M.; Puddu, P.; Paradis, M.; Angelo, C.; Bagetta, G.; Melino, G. A novel recessive connexin 31 (GJB3) mutation in a case of erythrokeratoderma variabilis. *J Invest Dermatol* **2004**, *122*, 837-839, doi:10.1111/j.0022-202X.2004.22311.x.
73. Terrinoni, A.; Cocuroccia, B.; Gubinelli, E.; Zambruno, G.; Candi, E.; Melino, G.; Girolomoni, G. Identification of the keratin K9 R162W mutation in patients of Italian origin with epidermolytic palmoplantar keratoderma. *Eur J Dermatol* **2004**, *14*, 375-378.
74. Rossi, M.; Sayan, A.E.; Terrinoni, A.; Melino, G.; Knight, R.A. Mechanism of induction of apoptosis by p73 and its relevance to neuroblastoma biology. In *Annals of the New York Academy of Sciences*, New York Academy of Sciences: 2004; Vol. 1028, pp 143-149.
75. Munarriz, E.; Barcaroli, D.; Stephanou, A.; Townsend, P.A.; Maisse, C.; Terrinoni, A.; Neale, M.H.; Martin, S.J.; Latchman, D.S.; Knight, R.A., et al. PIAS-1 is a checkpoint

- regulator which affects exit from G1 and G2 by sumoylation of p73. *Mol Cell Biol* **2004**, *24*, 10593-10610, doi:10.1128/MCB.24.24.10593-10610.2004.
76. Candi, E.; Paradisi, A.; Terrinoni, A.; Pietroni, V.; Oddi, S.; Cadot, B.; Jogini, V.; Meiyappan, M.; Clardy, J.; Finazzi-Agro, A., et al. Transglutaminase 5 is regulated by guanine-adenine nucleotides. *Biochem J* **2004**, *381*, 313-319, doi:10.1042/BJ20031474.
77. Billon, N.; Terrinoni, A.; Jolicoeur, C.; McCarthy, A.; Richardson, W.D.; Melino, G.; Raff, M. Roles for p53 and p73 during oligodendrocyte development. *Development (Cambridge)* **2004**, *131*, 1211-1220, doi:10.1242/dev.01035.
78. Marulli, G.C.; Campione, E.; Chimenti, M.S.; Terrinoni, A.; Melino, G.; Bianchi, L. Type I lamellar ichthyosis improved by tazarotene 0.1% gel. *Clin Exp Dermatol* **2003**, *28*, 391-393, doi:10.1046/j.1365-2230.2003.01318.x.
79. Pucci, B.; Claudio, P.P.; Masciullo, V.; Bellincampi, L.; Terrinoni, A.; Khalili, K.; Melino, G.; Giordano, A. pRb2/p130 promotes radiation-induced cell death in the glioblastoma cell line HJC12 by p73 upregulation and Bcl-2 downregulation. *Oncogene* **2002**, *21*, 5897-5905, doi:10.1038/sj.onc.1205750.
80. Candi, E.; Paradisi, A.; Terrinoni, A.; Cadot, B.; Rufini, A.; Puddu, P.; Melino, G. Role of transglutaminase 5 in epidermis. *Minerva Biotechnol.* **2002**, *14*, 155-158.
81. Candi, E.; Oddi, S.; Paradisi, A.; Terrinoni, A.; Ranalli, M.; Teofoli, P.; Citro, G.; Scarpato, S.; Puddu, P.; Melino, G. Expression of transglutaminase 5 in normal and pathologic human epidermis. *J Invest Dermatol* **2002**, *119*, 670-677, doi:10.1046/j.1523-1747.2002.01853.x.
82. Bernassola, F.; Federici, M.; Corazzari, M.; Terrinoni, A.; Hribal, M.L.; De Laurenzi, V.; Ranalli, M.; Massa, O.; Sesti, G.; McLean, W.H., et al. Role of transglutaminase 2 in glucose tolerance: knockout mice studies and a putative mutation in a MODY patient. *FASEB J* **2002**, *16*, 1371-1378, doi:10.1096/fj.01-0689com.
83. Terrinoni, A.; Smith, F.J.; Didona, B.; Canzona, F.; Paradisi, M.; Huber, M.; Hohl, D.; David, A.; Verloes, A.; Leigh, I.M., et al. Novel and recurrent mutations in the genes encoding keratins K6a, K16 and K17 in 13 cases of pachyonychia congenita. *J Invest Dermatol* **2001**, *117*, 1391-1396, doi:10.1046/j.0022-202x.2001.01565.x.
84. Terrinoni, A.; Rugg, E.L.; Lane, E.B.; Melino, G.; Felix, D.H.; Munro, C.S.; McLean, W.H. A novel mutation in the keratin 13 gene causing oral white sponge nevus. *J Dent Res* **2001**, *80*, 919-923, doi:10.1177/00220345010800031401.
85. Terrinoni, A.; Dell'Arciprete, R.; Fornaro, M.; Stella, M.; Alberti, S. Cyclin D1 gene contains a cryptic promoter that is functional in human cancer cells. *Genes Chromosomes Cancer* **2001**, *31*, 209-220, doi:10.1002/gcc.1137.
86. Candi, E.; Oddi, S.; Terrinoni, A.; Paradisi, A.; Ranalli, M.; Finazzi-Agro, A.; Melino, G. Transglutaminase 5 cross-links loricrin, involucrin, and small proline-rich proteins in vitro. *J Biol Chem* **2001**, *276*, 35014-35023, doi:10.1074/jbc.M010157200.
87. Vernole, P.; Terrinoni, A.; Didona, B.; De Laurenzi, V.; Rossi, P.; Melino, G.; Grimaldi, P. An SRY-negative XX male with Huriez syndrome. *Clin Genet* **2000**, *57*, 61-66, doi:10.1034/j.1399-0004.2000.570109.x.
88. Terrinoni, A.; Puddu, P.; Didona, B.; De Laurenzi, V.; Candi, E.; Smith, F.J.; McLean, W.H.; Melino, G. A mutation in the V1 domain of K16 is responsible for unilateral palmoplantar verrucous nevus. *J Invest Dermatol* **2000**, *114*, 1136-1140, doi:10.1046/j.1523-1747.2000.00983.x.
89. Terrinoni, A.; Candi, E.; Oddi, S.; Gobello, T.; Camaione, D.B.; Mazzanti, C.; Zambruno, G.; Knight, R.; Melino, G. A glutamine insertion in the 1A alpha helical domain of the keratin 4 gene in a familial case of white sponge nevus. *J Invest Dermatol* **2000**, *114*, 388-391, doi:10.1046/j.1523-1747.2000.00890.x.
90. Richard, G.; Brown, N.; Smith, L.E.; Terrinoni, A.; Melino, G.; Mackie, R.M.; Bale, S.J.; Uitto, J. The spectrum of mutations in erythrokeratodermias--novel and de novo mutations in GJB3. *Hum Genet* **2000**, *106*, 321-329, doi:10.1007/s004390051045.

91. De Laurenzi, V.; Rossi, A.; Terrinoni, A.; Barcaroli, D.; Levrero, M.; Costanzo, A.; Knight, R.A.; Guerrieri, P.; Melino, G. p63 and p73 transactivate differentiation gene promoters in human keratinocytes. *Biochem Biophys Res Commun* **2000**, *273*, 342-346, doi:10.1006/bbrc.2000.2932.
92. De Laurenzi, V.D.; Catani, M.V.; Terrinoni, A.; Corazzari, M.; Melino, G.; Costanzo, A.; Levrero, M.; Knight, R.A. Additional complexity in p73: induction by mitogens in lymphoid cells and identification of two new splicing variants epsilon and zeta. *Cell Death Differ* **1999**, *6*, 389-390, doi:10.1038/sj.cdd.4400521.
93. Melino, G.; De Laurenzi, V.; Catani, M.V.; Terrinoni, A.; Ciani, B.; Candi, E.; Marekov, L.; Steinert, P.M. The cornified envelope: a model of cell death in the skin. *Results Probl Cell Differ* **1998**, *24*, 175-212, doi:10.1007/978-3-540-69185-3\_9.
94. Junakovic, N.; Terrinoni, A.; Di Franco, C.; Vieira, C.; Loevenbruck, C. Accumulation of transposable elements in the heterochromatin and on the Y chromosome of *Drosophila simulans* and *Drosophila melanogaster*. *J Mol Evol* **1998**, *46*, 661-668, doi:10.1007/pl00006346.
95. De Laurenzi, V.; Costanzo, A.; Barcaroli, D.; Terrinoni, A.; Falco, M.; Annicchiarico-Petruzzelli, M.; Levrero, M.; Melino, G. Two new p73 splice variants, gamma and delta, with different transcriptional activity. *J Exp Med* **1998**, *188*, 1763-1768, doi:10.1084/jem.188.9.1763.
96. Chalvet, F.; di Franco, C.; Terrinoni, A.; Pelisson, A.; Junakovic, N.; Bucheton, A. Potentially active copies of the gypsy retroelement are confined to the Y chromosome of some strains of *Drosophila melanogaster* possibly as the result of the female-specific effect of the flamenco gene. *J Mol Evol* **1998**, *46*, 437-441, doi:10.1007/pl00006323.
97. Terrinoni, A.; Franco, C.D.; Dimitri, P.; Junakovic, N. Intragenomic distribution and stability of transposable elements in euchromatin and heterochromatin of *Drosophila melanogaster*: non-LTR retrotransposon. *J Mol Evol* **1997**, *45*, 145-153, doi:10.1007/pl00006214.
98. Mostaccioli, S.; DeLaurenzi, V.; Terrinoni, A.; Richard, G.; Didona, B.; Cavalieri, R.; Melino, G. White sponge nevus is caused by mutations in mucosal keratins. *EUR. J. DERMATOL.* **1997**, *7*, 405-408.
99. Junakovic, N.; Di Franco, C.; Terrinoni, A. Evidence for a host role in regulating the activity of transposable elements in *Drosophila melanogaster*: the case of the persistent instability of Bari 1 elements in Charolles stock. *Genetica* **1997**, *100*, 149-154, doi:10.1023/a:1018325427405.
100. Di Franco, C.; Terrinoni, A.; Dimitri, P.; Junakovic, N. Intragenomic distribution and stability of transposable elements in euchromatin and heterochromatin of *Drosophila melanogaster*: elements with inverted repeats Bari 1, hobo, and pogo. *J Mol Evol* **1997**, *45*, 247-252, doi:10.1007/pl00006227.