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氧化应激与DNA损伤

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Oxidative Stress and DNA Injury

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摘要 人和动物机体细胞在遭受氮氧化合物、钙和病原体等体内外环境的刺激后,氧化和抗氧化系统之间的平衡被破坏,从而促进细胞内活性氧分子(ROS)的大量产生和积累,最终导致机体产生氧化应激。氧化应激可以导致DNA链断裂、DNA位点突变、DNA双链畸变和原癌基因与肿瘤抑制基因突变等形式的DNA损伤;同时,DNA也在遭受脱嘌呤和脱嘧啶、X射线、紫外线、烷化剂和嵌入剂等体内外物理或化学因素的刺激下造成DNA损伤,DNA损伤也能诱导机体产生氧化应激。本文主要对氧化应激与DNA损伤之间的联系作一综述,以期为后续的相关研究提供参考。

关键词: 氧化应激 ROS DNA损伤

Abstract: After challenged by intra- and extracellular environmental factors, such as nitric oxide, calcium and pathogenic organisms, the balance between oxidative and antioxidant defense systems in human and animal cells is broken, which results in the accumulation of reactive oxygen species (ROS) within cells, and finally oxidative stress occurs. Oxidative stress can cause DNA damage, such as DNA strand breaks, point mutations, aberrant DNA cross-linking, and mutations in proto-oncogenes and tumor suppressor genes; meanwhile, when challenged by physical or chemical factors, such as depurination and apyrimidinic factors, X-ray, ultraviolet, alkylating agent and intercalator, DNA also could be injured, furthermore, DNA damage is a source to oxidative stress in body. This paper mainly reviewed oxidative stress, DNA damage, and their relationship, which could provide a theoretical basis for a further research.

Keywords: oxidative stress, ROS, DNA damage

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- [1] KANNINEN K,WHITE A R,KOISTINAHO J,et al.Targeting glycogen synthase kinase-3 β for therapeutic benefit against oxidative stress in Alzheimer's disease:involvement of the Nrf2-ARE Pathway[J].International Journal of Alzheimer's Disease,2011,2011:985085.
- [2] PANTANO C,REYNAERT N L,VAN DER VLIET A,et al.Redox-sensitive kinases of the nuclear factor- κ B signaling pathway[J].Antioxidants & Redox Signaling,2006,8(9/10):1791-1806.
- [3] MEIRA L B,BUGNI J M,GREEN S L,et al.DNA damage induced by chronic inflammation contributes to colon carcinogenesis in mice[J].The Journal of Clinical Investigation,2008,118(7):2516-2525.
- [4] ROWE L A,DEGTYAREVA N,DOETSCH P W.DNA damage-induced reactive oxygen species (ROS) stress response in *Saccharomyces cerevisiae* [J].Free Radical Biology and Medicine,2008,45:1167-1177.
- [5] HAMANAKA R B,CHANDEL N S.Mitochondrial reactive oxygen species regulate cellular signaling and dictate biological outcomes[J].Trends Biochemical Sciences,2010,35:505-513.
- [6] BRAGADO P,ARMESILLA A,SILVA A,et al.Apoptosis by cisplatin requires p53 mediated p38 α MAPK activation through ROS generation [J].Apoptosis,2007,12:1733-1742.
- [7] LIU B,CHEN Y M,ST CLAIR D K.ROS and p53:a versatile partnership[J].Free Radical Biology and Medicine,2008,44:1529-1535.
- [8] KLAUNIG J E,KAMENDULIS L M.The role of oxidative stress in carcinogenesis[J].Annual Review of Pharmacology and Toxicology,2004,44:239-267.
- [9] VALKO M,IZAKOVIC M,MAZUR M,et al.Role of oxygen radicals in DNA damage and cancer incidence[J].Molecular and Cellular Biochemistry,2004,266:37-56.
- [10] RICQUIER D,BOUILLAUD F.The uncoupling protein homologues:UCP1,UCP2,UCP3,StUCP and AtUCP[J].The Biochemical Journal,2000,345:161-179.
- [11] AFFOURTIT C,CRICHTON P G,PARKER N,et al.Novel uncoupling proteins[J].Novartis Foundation Symposium,2007,287:70-80.
- [12] MCCUBREY J A,LAHAIR M M,FRANKLIN R A.Reactive oxygen species-induced activation of the MAP kinase signaling pathways[J].Antioxidants & Redox Signaling,2006,8(9/10):1775-1789.
- [13] LEVONEN A L,LANDAR A,RAMACHANDRAN A,et al.Cellular mechanisms of redox cell signalling:role of cysteine modification in controlling antioxidant defences in response to electrophilic lipid oxidation products[J].Biochemical Journal,2004,378:373-382.
- [14] 蒋满荣.DNA损伤对哺乳动物细胞周期和凋亡的影响[D].博士学位论文.上海:中国科学院研究生院,2006.
- [15] BARTEK J,LUKAS J.DNA damage checkpoints:from initiation to recovery or adaptation[J].Current Opinion in Cell Biology,2007,19:238-245.
- [16] HARRISON J C,HABER J E.Surviving the breakup:the DNA damage checkpoint[J].Annual Review of Genetics,2006,40:209-235.
- [17] MATSUOKA S,BALLIF B A,SMOGORZEWSKA A,et al.ATM and ATR substrate analysis reveals extensive protein networks responsive to DNA damage[J].Science,2007,316:1160-1166.
- [18] SHILOH Y.ATM (ataxia telangiectasia mutated):expanding roles in the DNA damage response and cellular homeostasis[J].Biochemical Society Transactions,2001,29:661-666.
- [19] YEUNG M,DUROCHER D.Engineering a DNA damage response without DNA damage[J].Genome Biology,2008,9(7):227.
- [20] AGAMI R,BERRARDS R.Distinct initiation and maintenance mechanisms cooperate to induce G1 cell cycle arrest in response to DNA damage [J].Cell,2000,102:55-66.
- [21] BUSINO L,DONZELLI M,CHIESA M,et al.Degradation of Cdc25A by β -TrCP during S phase and in response to DNA damage [J].Nature,2003,426:87-91.
- [22] LUKAS J,LUKAS C,BARTEK J.Mammalian cell cycle checkpoints:signalling pathways and their organization in space and time[J].DNA Repair,2004,3(8/9):997-1007.
- [23] NOWSHEEN S,YANG E S.The intersection between DNA damage response and cell death pathways[J].Experimental Oncology,2012,34(3):243-254.
- [24] GOLDING S E,MORGAN R N,ADAMS B R,et al.Pro-survival AKT and ERK signaling from EGFR and mutant EGFRvIII enhances DNA double-strand break repair in human glioma cells[J].Cancer Biology & Therapy,2009,8:730-738.
- [25] MUKHERJEE B,MCELLIN B,CAMACHO C V,et al.EGFRvIII and DNA double-strand break repair:a molecular mechanism for radioresistance in glioblastoma[J].Cancer Research,2009,69:4252-4259.
- [26] BRANDSMA I,VAN GENT D C.Pathway choice in DNA double strand break repair:observations of a balancing act[J].Genome Integrity,2012,3(1):9.
- [27] COUSSENS L M,WERB Z.Inflammation and cancer[J].Nature,2002,420:860-867.
- [28] LIANG F Q,GODLEY B F.Oxidative stress-induced mitochondrial DNA damage in human retinal pigment epithelial cells:a possible mechanism for RPE aging and age-related macular degeneration[J].Experimental Eye Research,2003,76(4):397-403. 
- [29] ISHIKAWA K,TAKENAGA K,AKIMOTO M,et al.ROS-generating mitochondrial DNA mutations can regulate tumor cell metastasis [J].Science,2008,320:661-664.
- [30] LI H,HONG Z H.Mitochondrial DNA mutations in human tumor cells[J].Oncology Letters,2012,4(5):868-872.

- [31] WANG D,KREUTZER D A,ESSIGMANN J M.Mutagenicity and repair of oxidative DNA damage:insights from studies using defined lesions [J].Mutation Research/Fundamental and Molecular Mechanisms of Mutagenesis,1998,400: 99- 115.
- [32] EVERT B A,SALMON T B,SONG B W,et al.Spontaneous DNA damage in *Saccharomyces cerevisiae* elicits phenotypic properties similar to cancer cells[J].Journal of Biological Chemistry,2004,279: 22585- 22594.
- [33] 吴东明.14-3-3zeta蛋白多克隆抗体的制备与鉴定 [D].硕士学位论文.合肥:安徽大学,2007.
- [34] KANG M A,SO E Y,SIMONS A L,et al.DNA damage induces reactive oxygen species generation through the H2AX-Nox1/Rac1 pathway[J].Cell Death and Disease,2012,3:e249.
- [1] 安清聪,张春勇,李美荃,陈克磷,郭荣富.谷氧还蛋白1和硫氧还蛋白1基因在高黎贡山猪不同组织中表达规律及维生素E对其在氧化应激细胞中表达的影响[J].动物营养学报,2013,25(8): 1825-1835
- [2] 阳坦,孙志洪,李晓敏.在早期断奶过程中氧化应激的产生机制及其影响[J].动物营养学报,2013,25(4): 705-714
- [3] 吴文旋,段永邦,李胜利.饲料阴阳离子差对围产期奶牛酸碱平衡、血浆钙浓度及抗氧化应激的影响[J].动物营养学报,2013,25(4): 856-863
- [4] 马思聪,李磊,李海燕,敖远扬,唐志如.瞬时受体电位2通道介导动物氧化应激的机理及其营养调控[J].动物营养学报,2013,25(10): 2231-2237
- [5] 张春勇,陈克磷,黄金昌,郭荣富.谷氧还蛋白1和硫氧还蛋白1基因在云南乌金猪不同组织中的表达特点及L-组氨酸对其在氧化应激细胞中表达的影响[J].动物营养学报,2012,24(12): 2415-2423
- [6] 魏炳栋,陶浩,于维,陈群,李林.黄芪多糖对氢化可的松氧化应激状态下肉仔鸡生长性能、脏器指数及抗氧化能力的影响[J].动物营养学报,2012,24(10): 1939-1945
- [7] 徐歆,毛予龙,李雅丽,胡胜兰,吴红照,李卫芬.枯草芽孢杆菌B1对饲喂高脂饲料小鼠抗氧化功能的影响[J].动物营养学报,2012,24(10): 2067-2072
- [8] 杨小军,高泽,刘凯,王益兵,覃定奎,姚军虎*.谷氨酰胺对肉仔鸡肠道黏膜淋巴细胞增殖活性、氧化应激和免疫应激的调控作用[J].动物营养学报,2011,23(02): 274-279
- [9] 崔志文,黄琴,黄怡,吴红照,文静,李卫芬*.枯草芽孢杆菌对Caco-2细胞抗氧化功能的影响研究[J].动物营养学报,2011,23(02): 293-298
- [10] 黄金昌,郭荣富*.谷氧还蛋白和硫氧还蛋白对动物抗氧化应激生物学效应的研究进展[J].动物营养学报,2010,22(04): 845-850
- [11] 袁施彬,陈代文.不同氧化应激模式下仔猪血细胞参数变化的比较研[J].动物营养学报,2008,20(06): 617-623
- [12] 刘永祥,芮于明.日粮镁对肉仔鸡腿肌中活性氧产量的影响(英文)[J].动物营养学报,2007,19(03): 204-210