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**CNS / Psyche-Blood / Immune system / Cells-Acupuncture / Drugs-Electrophysiology and Thermodynamics: The redox potential / charge-,energy transfer is responsible for stress protein or O<sub>2</sub><sup>-</sup>-synthesis / transport and apoptosis/proliferation.**

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The plasma membrane NADPH oxidase is a rather complicated electron transfer system, which resembles probably the most important crossover-, end- and starting point of various signal transduction pathways: Ca<sup>2+</sup>/Mg<sup>2+</sup>-sensitive phosphorylation and dephosphorylation (incl. JAK-STAT-g-protein-pathway) of the complex regulates the electron transfer (incl. thiol/disulfid-interchange-cyt/FeS-protein) between mitochondria / plasma and nucleus / ER (NADH/ATP-NADPH/K<sup>+</sup>-O<sub>2</sub>/O<sub>2</sub><sup>-</sup>-DNA/IgE-cyt P450). Another control is played for instance by arachidonic acid (delivered by PLA<sub>2</sub>). The entire complex is not a permanent entity of the membrane, but is built up during defence conditions: The NADPH oxidase belongs then to enzyme systems like insulinR, nAcChR, adenylylate cyclase, mitochondrial K<sup>+</sup> ATPase/ATP-synthase(CV) presenting a universal principle of nature: The connecting logistics between CNS (adrenal cortex) and blood/immune system/cells is played by charge displacement or separation and charge (energy) transfer (induced by the cation, incl.proton and/or anion pressure) and redox potential and may be visualized by the successful treatment of various diseases and psychological disorders with acupuncture/ homeopathy/ relaxation therapy or drugs. We try to connect natural sciences in order to get involved into this logistic for prediction of effective new treatments, which includes the modeling of interfering drugs for instance.

**References:**

- (01)Kiehl, R. (1976) Dissertation, MPI for Medical Research, Heidelberg; FEBS Letters 61, 68-71; 72, 24-28.
- (02)Kiehl, R. (1980) FEBS Letters 109, 280-282.
- (03)Kiehl, R. and Hanstein, W.G. (1984) Biochim.Biophys.Acta 766,375-385.
- (04)Kiehl, R. and Hanstein, W.G. (1984) 3.EBEC, 323-324.
- (05)Kiehl, R. Varsanyi, M. and Neumann, E. (1987) Biochem.Biophys.Res. Comm.147 ,1251-11258.
- (06)Kiehl, R. (1993) Biol.Chem.Hoppe-Seyler 374, 742.
- (07)Kiehl, R. (1993) Dear Colleague, New in dermatology 2(2),4-6.
- (08)Guengerich, F.P. (1993) American Scientist 81,440-447.
- (09)Kiehl, R. (1994) Int Alk-Ciba Corning Joint Symp, Benzheim.
- (10)Kiehl, R. (1994) Biol.Chem.Hoppe-Seyler 375,S61.
- (11)Kiehl, R. (1995/96) Habilthesis, LMU Munich, Medical Faculty.
- (12)Kiehl, R. (1995) GDCh-Biochemie, Kaiserslautern, P28.
- (13)Kiehl, R. (1995) Amino Acids 9(1), 20.
- (14)Kiehl, R. (1996) GDCH-Biochemie, Giessen, P3.
- (15)Kiehl, R. (1997) BIOforum 12, 686-690.
- (16)Kiehl, R. (1997) Amino Acids 13, 50-51.

- (17)Kiehl, R. (1998) IFCC, Proc. of the 17. Int.Symp. on TheConfl.of CritCareAnal.and NearPatientTesting, Nice,467-482.
- (18)Kiehl, R. (1999) Eur.J.Cell Biology 78 (Suppl.49), 89.
- (19)Kiehl, R. (1999) 2.Joint Meeting, Signal Transduction: Receptors, Mediators and Genes, Langen, P34, p82.
- (20)Kiehl, R.(1999) Biotechnology Int. (Universal Med. Press).