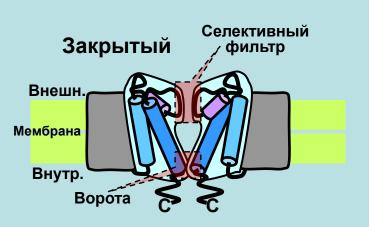
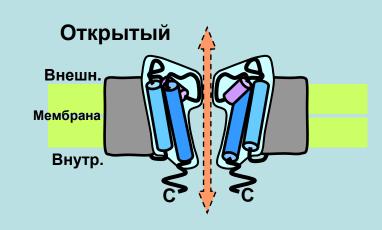


MICTATYT QUIBNOJOTAN MM. AA 5000001548 HARY

Как муха и червяк помогают изучать болезни человека?

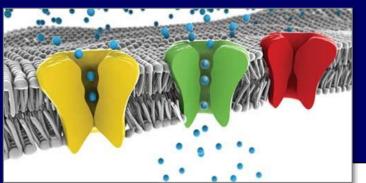
Я.М. Шуба







Классы ионных каналов



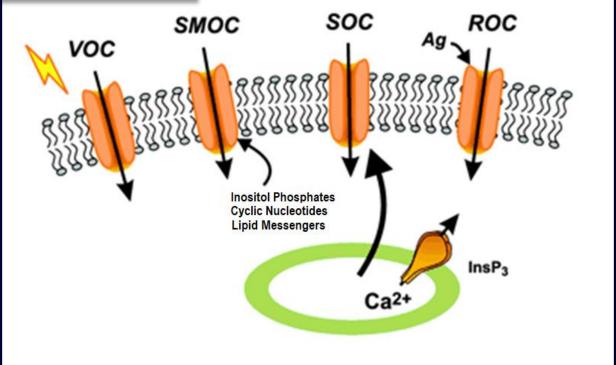
VOC – <u>V</u>oltage-<u>O</u>perated <u>C</u>hannels (потенциал-управляемые);

SMOC – **S**econd **M**essenger-**O**perated **C**hannels

(управляемые вторичными посредниками);

SOC – **S**tore-**O**perated **C**hannels (депо-управляемые);

ROC – Receptor-Operated Channel (рецептор-управляемые).



Каналоспецифичаские лекарственные препараты:

Антиаритмики

I типа: блокаторы натриевых каналов III типа: блокаторы калиевых каналов IV типа: блокаторы кальциевых каналов

Антигипертензивные

блокаторы кальциевых каналов активаторы К_{АТФ}-каналов

Кардиопротекторы

активаторы К_{АТФ}-каналов

Анальгетики

блокаторы натриевых каналов блокаторы кальциевых каналов блокаторы TRPV1 каналов

блокаторы Р2X рецепторов

Антиэпилептики

агонисты ГАМК_А рецепторов

блокаторы натриевых каналов блокаторы НП кальциевых каналов блокаторы ионотропных глутаматных рецепторов

Анестетики

блокаторы К_{АТФ}-каналов (сульфанилмочевины)

Антидиабетики

Локальные: блокаторы натриевых каналов

Общие: активаторы калиевых каналов

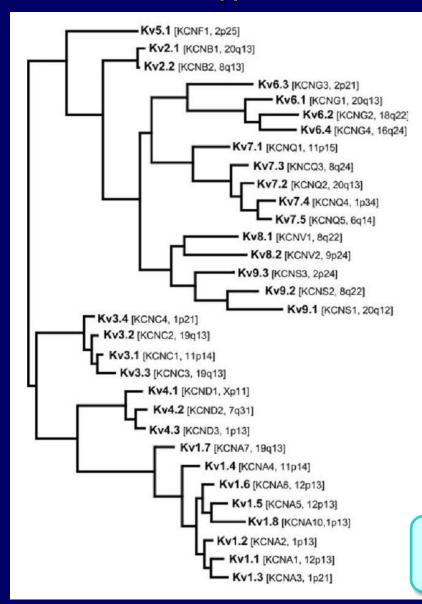
Протираковые

блокаторы
потенциалнезависимых
кальциевых
каналов
(carboxyamidotriazole, CAI)

блокаторы хлорных каналов (iodinechlorotoxin)

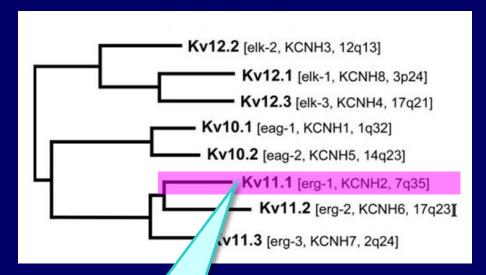


Shaker-подобные

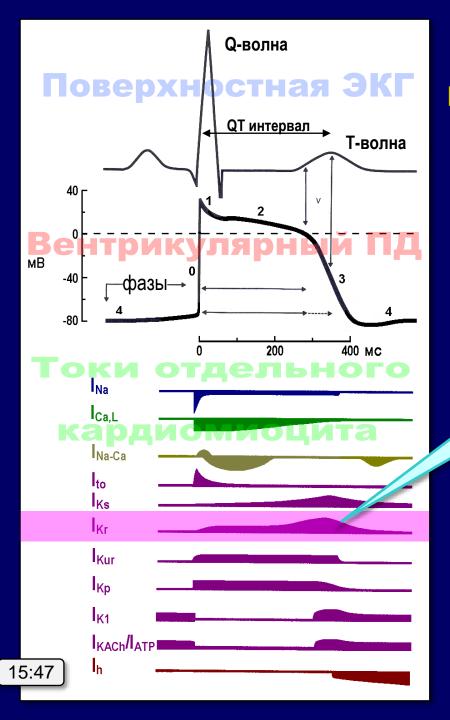


Калиевые каналы млекопитающих

Ether-a-go-go(eag)-подобные



<u>H</u>uman <u>E</u>ther-a-go-go-<u>R</u>elated <u>G</u>ene (HERG)

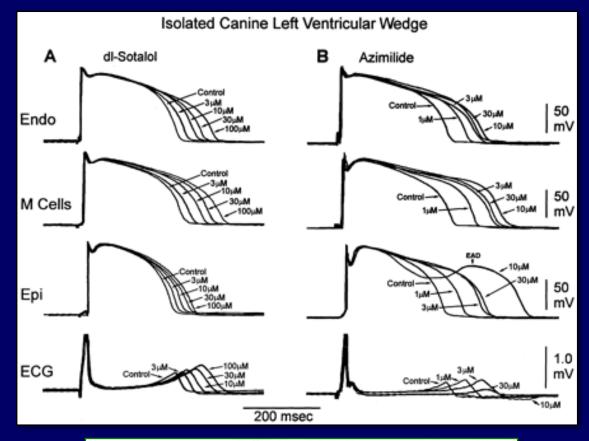


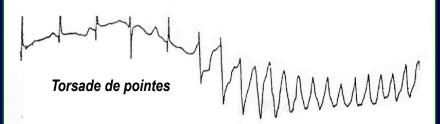
Связь между ЭКГ, потенциалом действия и токами в сердце

HERG-опосредованный **I**_{Kr} - один из основных реполяризирующих калиевых токов в сердце

LQTS (<u>L</u>ong <u>QT</u> <u>S</u>yndrom): синдром удлиненного QT интервала электрокардиограммы - фактор риска и причина возникновения летальных вентрикулярных аритмий *Torsad de Pointes*

- III класс антиаритмики,
- антигистамины,
- антималярийные



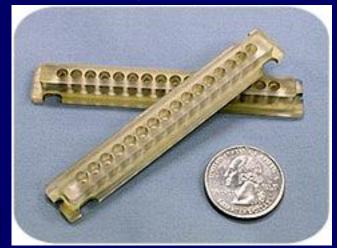


Автоматизированная электрофизиология

Patch-clamp



16-ти лунковые чипы



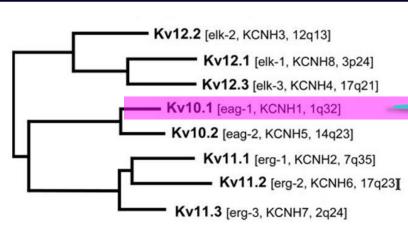
Микроэлектродная



16-ти электродный робот



Ether a go-go(eag)-подобные К+ каналы



Eag1 – онкогенный калиевый канал

Asher et al. World Journal of Surgical Oncology 2010, 8:113 http://www.wjso.com/content/8/1/113



REVIEW Open Access

Eag and HERG potassium channels as novel therapeutic targets in cancer

Viren Asher^{1*}, Heidi Sowter², Robert Shaw³, Anish Bali⁴, Raheela Khan⁵

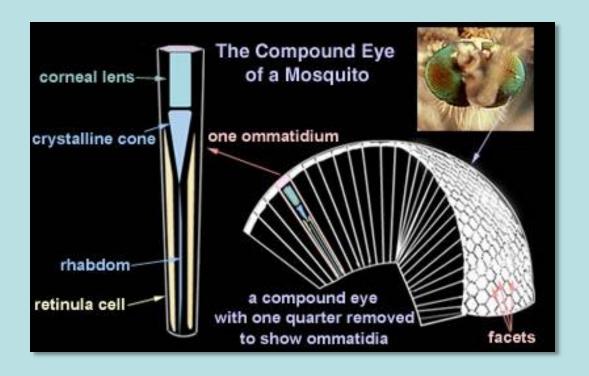
Abstract

Voltage gated potassium channels have been extensively studied in relation to cancer. In this review, we will focus on the role of two potassium channels, Ether à-go-go (Eag), Human ether à-go-go related gene (HERG), in cancer and their potential therapeutic utility in the treatment of cancer. Eag and HERG are expressed in cancers of various organs and have been implicated in cell cycle progression and proliferation of cancer cells. Inhibition of these channels has been shown to reduce proliferation both in vitro and vivo studies identifying potassium channel modulators as putative inhibitors of tumour progression. Eag channels in view of their restricted expression in normal tissue may emerge as novel tumour biomarkers.

Еад1 в норме присутствует только в мозге и в незначительных количествах в периферических тканях. Его экспрессия сильно увеличивается в распространенных опухолях человека. Присутствие Eag1 связано с усилением пролиферации.



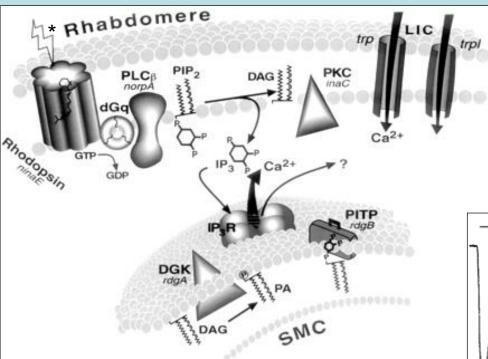
Compound eye (фасеточный глаз) of arthropods



The type of <u>eye</u> comonly found in <u>arthropods</u>. A compound eye has a mesh-like appearance because it consists of hundreds or thousands of tiny lens-capped optical units called <u>ommatidia</u>. Each ommatidium has its own <u>cornea</u>, <u>lens</u>, and photoreceptor cells for distinguishing brightness and color. Individual ommatidia guide light through a lens and cone into a channel, known as a <u>rhabdom</u>, which contains light-sensitive cells. These are connected to optical nerve cells to produce the image. The ommatidia are separated from each other by varying degrees of <u>pigment</u>.

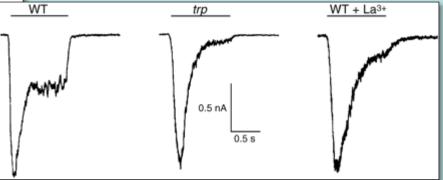
Transient Receptor Potential (TRP) channel in *Drosophila* vision

Phototransduction in the compound eye of arthropods



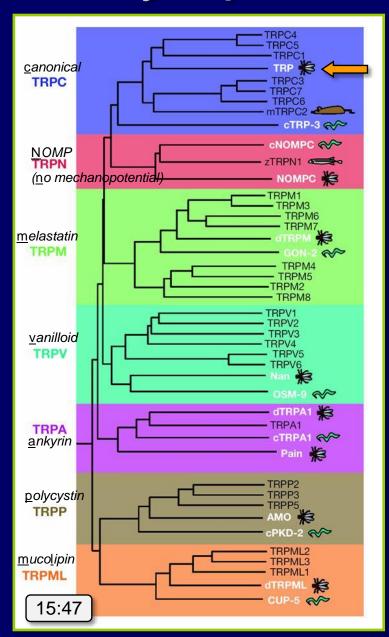
The phosphoinositide cascade of insect's vision. Cloned genes (for all of which mutants are available) are shown in italics, alongside their corresponding proteins. Upon absorption of light, rhodopsin (ninaE gene) is converted to the active metarhodopsin state, which activates a heterotrimeric G protein (dGq). This leads to activation of phospholipase C (PLC, norpA gene) and subsequent opening of two classes of light-sensitive channels encoded at least in part by trp and trpl genes, by an as yet unknown mechanism. TRP and TRPL opening leads to the light-induced current (LIC). Deactivation of channel activity is regulated by protein kinas C (PKC, inaC gene).

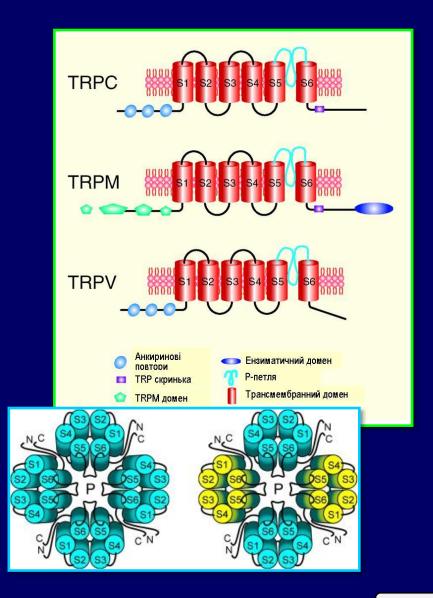
A spontaneous *Drosophila* phototransduction mutant identified in 1977 (Minke, 1977) displayed transient receptor potentials (trp) in response to continuous light. Identification of the gene product underlying that mutation and recognition of its function as an ion channel (Montel and Rubin, 1989) gave rise to the awareness of a new class of cation channels that differed significantly from the canonical voltage-dependent channels.



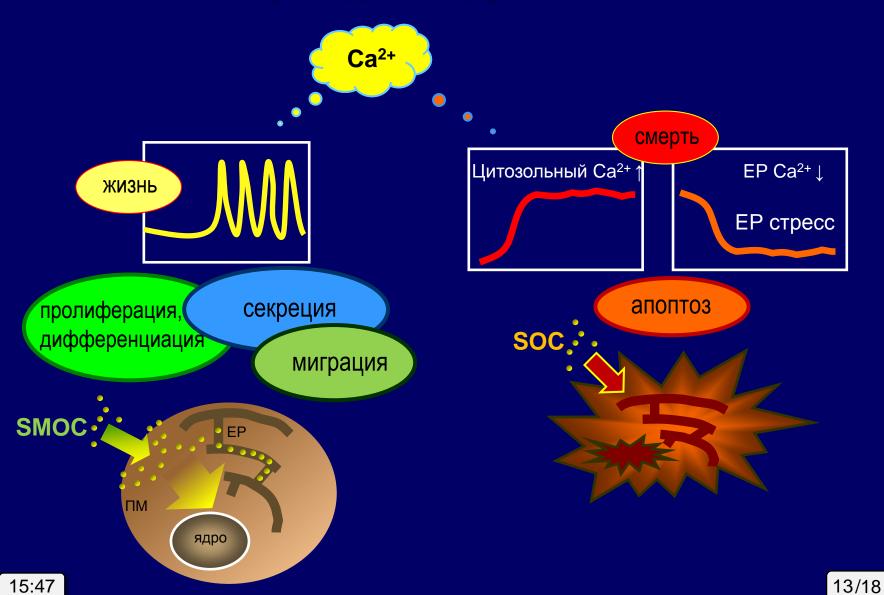
The *trp* phenotype. Light-induced currents in response to prolonged intense orange lights were recorded in voltage-clamped photoreceptors of wild type (WT), the *trp* mutant, and WT treated with La3. A peak response and a plateau characterize the light response of WT. The rapid peak-plateau transition is a manifestation of Ca2-dependent light adaptation. The response of the *trp* photoreceptor decays close to baseline during light due to exhaustion of excitation. A similar decay of the light response close to baseline is obtained by application of 10 M La3 to the extracellular medium of WT photoreceptors.

Суперсемейство TRP каналов





TRP-каналы в кальциевой сигнализации: от нормы до канцерогенеза



TRP-каналы как терморецепторы и сенсоры химических раздражителей...

Identification of a cold receptor reveals a general role for TRP channels in thermosensation

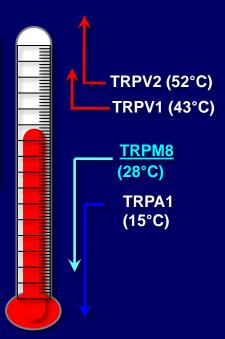
NATURE [VOL 416 | 7 MARCH 2002 | WWW.nature.com]

David D. McKerny*†, Werner M. Neuhausser*† & David Julius*

* Department of Cellular and Molecular Pharmacology, University of California, San Francisco, California 94143-0450, USA

† These authors contributed equally to this work





TRP-канали как детерминанты заболеваний человека...

TRP channels in cancer.

Prevarskaya N, Zhang L, Barritt G. Biochim Biophys Acta. 2007 Aug;1772(8):937-46.

TRP channels: functions and involvement in **neurologic disease**.

Benarroch EE.

Neurology. 2008 Feb 19;70(8):648-52.

TRP channel and cardiovascular disease.

Watanabe H, Murakami M, Ohba T, Takahashi Y, Ito H. Pharmacol Ther. 2008 Jun;118(3):337-51.

TRP channels as emerging targets for pain therapeutics.

Broad LM, Mogg AJ, Beattie RE, Ogden AM, Blanco MJ, Bleakman D. Expert Opin Ther Targets. 2009 Jan;13(1):69-81.

TRP channels: emerging targets for respiratory disease.

Banner KH, Igney F, Poll C.

Pharmacol Ther. 2011 Jun;130(3):371-84.



... возможности генной терапии рака

Маркеры типов рака

- 1) carcinoembryonic antigen (CEA, CEACAM gene family), which is produced by tumors of gastrointestinal system as well as by some other cancers such as lung, breast, ovarian;
- **2) prostate specific antigen** (PSA, also known as kallikrein III, gene *KLK3*);
- 3) bladder tumor antigen (BTA);
- **4) cacrcinoma antigen 15-3** (CA 15-3, gene *MUC1*) characteristic of advanced breast cancer;
- **5) cancer antigen 125** (CA 125, gene *MUC16*), which is typical of ovarian cancer:
- **6) carbohydrate antigen 19-9** (CA 19-9, synthesis involves three genes *FUT2*, *FUT3*, *ST3GAL1*), which is the most useful marker of pancreatic cancer, and is also elevated in colon cancer.

MDEG мутация: Glycine430Phenylalanine

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www.nature.com/cet

ture.com/cgt

A novel strategy for cancer therapy by mutated mammalian degenerin gene transfer

Masayoshi Horimoto,¹ Yutaka Sasaki,¹,² Shinya Ugawa,³,⁴ Shigeo Wada,¹ Takashi Toyama,¹ Kenya Iyoda,² Takayuki Yakushijin,² Yuki Minami,³ Toshifumi Ito,¹ Taizo Hijioka,⁵ Akiko Eguchi,⁶ Mahito Nakanishi,⁶ Shoichi Shimada,⁴ Masaya Tohyama,³ Norio Hayashi,² and Masatsugu Hori¹

Departments of ¹Internal Medicine and Therapeutics, ²Molecular Therapeutics, and ³Anatomy and Neuroscience, Osaka University Graduate School of Medicine, Osaka, Japan; ⁴Department of Anatomy II, Nagoya City University Medical School, Nagoya, Japan; ⁵Department of Gastroenterology, Osaka-minami National Hospital, Osaka, Japan; and ⁶Department of Neurovirology, Research Institute for Microbial Diseases, Osaka University, Osaka, Japan.

Mammalian degenerin (MDEG) is a member of the amiloride-sensitive sodium ion channel family, and its site-directed active mutant (MDEG-G430F) induces massive Na⁺ influx into cells, leading to cell ballooning and cell bursting. We attempted a novel therapeutic approach for gastric cancers by transferring MDEG-G430F into cancer cells using tumor-specific promoters. In carcinoembryonic antigen (CEA)-producing gastric cancer cells, the level of cell death observed when MDEG-G430F was used with a CEA promoter was similar to that observed when using a potent nonspecific promoter such as the cytomegalovirus promoter. In an *in vivo* study, fusogenic liposome complexes containing MDEG-G430F driven by the CEA promoter were injected intraperitoneally into CEA-producing gastric cancer cells in a mouse peritoneal dissemination model. Although all 15 of the control mice were dead by 50 days postinoculation, 13 of the 15 mice treated with MDEG-G430F survived. These results indicate that transfering MDEG-G430F into cancer tissues using tumor-specific promoters can achieve striking and selective cancer cell death

15:47 ve of the transcriptional efficiency of the promoters used *in vivo*, and suggest that this approach is a promising new strategy gene therapy. Cancer Gene Therapy (2000) 7, 1341–1347

промотор маркера рака

Рекомбінантна плазміда

кодирующая последовательность канала

кодирующая последовательность маркера экспрессии

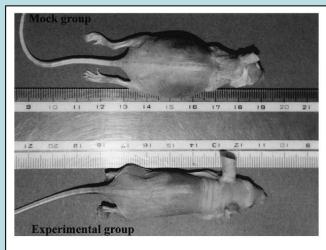


Figure 6. Mice with peritoneal dissemination of MKN45-P cells. Top: Representation of a mouse in the mock group at 30 days postinoculation. Mice inoculated with 1 × 10⁶ MKN45-P cells were injected i.p. with fusogenic liposomes containing CEApCDNA3 plasmid each week. All mice in the mock and control groups exhibited substantial peritoneal dissemination with massive ascites. Bottom: Representative demonstration of a mouse in the experimental group at 30 days postinoculation. Mice inoculated with 1 × 10⁶ MKN45-P cells were injected i.p. with fusogenic liposomes containing CEApCDNA3 MDEG-G430F plasmid each week. Mice in the experimental group exhibited substantially less peritoneal dissemination.

Key words: Mutated ion channel; degenerin; carcinoembryonic antigen promoter; fusogenic liposome; peritoneal dissemination.



Спасибо за внимание!



Київ, "Наукова думка", 2010