Anandamide (N-arachidonoylethanolamide) is a lipid signal molecule that was the first endogenous agonist for cannabinoid receptors to be discovered. Cannabinoid receptor type 1 (CB1) is widely distributed in neurons and nonneuronal cells in brain and peripheral organs including sperm, eggs, and preimplantation embryos. A study by Wang and colleagues in this issue of the JCI demonstrates that a critical balance between anandamide synthesis by N-acylphosphatidylethanolamine–selective phospholipase D (NAPE-PLD) and its degradation by fatty acid amide hydrolase (FAAH) in mouse embryos and oviducts creates locally an appropriate “anandamide tone” required for normal embryo development, oviductal transport, implantation, and pregnancy (see the related article beginning on page 2122). Adverse effects of elevated levels of anandamide on these processes resulting from FAAH inactivation are mimicked by administration of (-)-Δ9-tetrahydrocannabinol (THC; the major psychoactive constituent of marijuana), due to enhanced signaling via CB1. These findings show that exogenous THC can swamp endogenous anandamide signaling systems, thereby affecting multiple physiological processes.

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Attempts to understand the mechanisms responsible for the psychoactive properties of tetrahydrocannabinol (THC) in marijuana led to the discovery of canna-

Nonstandard abbreviations used: 2-AG, 2-arachidonoylglycerol; CB1, cannabinoid receptor type 1; FAAH, fatty acid amide hydrolase; NAPE-PLD, N-arachidonoylthanolamine–selective phospholipase D; THC, (−)-Δ⁹-tetrahydrocannabinol.

Conflict of interest: The author has declared that no conflict of interest exists.

Citation for this article: J Clin Invest. 2006;116:2087–2090 (2006). doi:10.1172/JCI29424.

N-acylphosphatidylethanolamine–selective phospholipase D (NAPE-PLD) when neurons and other cells are stimulated by neurotransmitters and hormones. Released anandamide is quickly eliminated by membrane-bound fatty acid amide hydrolase (FAAH), indicative of a role for this lipid in cell signaling. Anandamide signaling via cannabinoid receptors is not restricted to the central nervous system. We now know that CB1, CB2, NAPE-PLD, and FAAH are widely distributed in nonneuronal somatic cells of peripheral organs including those of the reproductive system, as well as in germ cells of vertebrates and invertebrates (1–8). This may account for the effects of marijuana and THC on multiple aspects of reproductive physiology, including secretion of gonadotrophic hormones by the pituitary gland, secretion of sex steroids by gonads, sperm production, ovulation, mating behavior, sperm capacitation, fertilization, early embryonic development, implantation of blastocysts into the uterine endometrium, placental functions, fetal growth, number of pregnancies carried to term, lactation, sucking behavior of newborns, postnatal development, and growth of malignant breast and prostate cells.
FAAH is the metabolic gatekeeper for anandamide signaling

Endocannabinoid signaling via CB1 regulates cleavage of mouse eggs, oviductal transport of preimplantation embryos, blastocyst hatching from the zona pellucida, and implantation of blastocysts into the uterine mucosa (3, 6, 7). In the study by Wang et al. in this issue of the JCI (9), NAPE-PLD was detected in mouse embryos at the stage of the fertilized egg through to the blastocyst stage, while FAAH first appeared in 2-cell embryos (Figure 1). FAAH expression was upregulated in the morula and blastocyst. FAAH was mostly expressed in trophoderm of blastocysts, suggesting an active role of embryonic FAAH in reducing anandamide levels at potential implantation sites in the uterine mucosa (6, 9). An inverse distribution of these enzymes was present in mouse oviducts on days 1–4 of pregnancy, with higher levels of Nape-pld in epithelium of the isthmus than of the ampulla (9). In contrast, FAAH was expressed at higher levels in ampulla than in isthmus epithelium. Suppression of FAAH activity in embryos and oviducts pharmacologically or by genetic ablation elevated anandamide levels in situ, which resulted in inhibition of embryonic development and expression of cell lineage genes required for differentiation, retention of embryos within the oviduct, impaired implantation, and reduced fertility. Administration of exogenous THC similarly affected early pregnancy events. Furthermore, the adverse effects on initiation of pregnancy resulting from inactivation of FAAH and by addition of THC were blocked by the CB1 selective antagonist SR141716. Unlike endogenous neurotransmitters/neuromodulators, which are rapidly released and degraded locally on demand in the body, exogenous drugs such as THC persist and swamp endogenous anandamide signaling systems, thereby exerting long-term effects on multiple physiological processes. The findings reported by Wang et al. (9) describe an extremely provocative model for the biological effects of THC and marijuana on functions of other peripheral organs as well as the brain.

**Clinical implications**

Aberrant functioning of anandamide signaling systems in embryos and oviducts in women may lead to ectopic pregnancy in the oviduct and/or impaired fertility (3, 7, 9). Similar adverse effects may be associated with abuse of marijuana by women of reproductive age. This possibility is consistent with studies showing that women who smoke tobacco show an increased incidence of ectopic pregnancy (10), which correlates with the direct inhibitory effects of tobacco smoke and nicotine on oviductal transport of embryos in hamsters (11). Faah-null female mice are less fertile than wild-type females (9). This observation is clinically relevant, since reduced peripheral FAAH activity is associated with spontaneous abortion in women (4). Oviductal transport of embryos in mice was shown to be regulated by crosstalk between anandamide signaling via CB1 and β-adrenergic receptors in the oviductal muscularis (7). Acetaminophen is a popular over-the-counter analgesic and antiinflammatory agent. Following decylation, it is converted in the body to the bioactive N-acylphenolamine (also known as AM404) via FAAH-dependent arachidonic acid conjugation (12). AM404 is a ligand for CB1 receptors and is a potent inhibitor of anandamide reuptake, the inhibition of which leads to elevated anandamide levels in the body. Do women using β-adrenergic medications or...
acetaminophen show elevated incidences of ectopic pregnancy and impaired fertility?

There are provocative similarities between anandamide signaling in embryos and oviductal epithelia and signaling pathways associated with sperm capacitation in the female reproductive tract. Studies on human and porcine sperm suggested the possible existence of an anandamide concentration gradient in the oviduct that regulates sperm capacitation (2, 3, 8, 9, 13, 14). A mechanism underlying this postulated anandamide gradient is indicated by the differences in the distribution of Nape-pld and Faah in mouse oviduct (9). Sperm and preimplantation mouse embryos express functional CB1 receptors as well as key enzymes (Nape-PLD and FAAH) for the release of anandamide and its subsequent degradation (9). Hence, the presence of other endocannabinoids in the microenvironment surrounding the oviduct might orchestrate subtle differences in their physiology.

Non-CB1 and non-CB2 receptors for anandamide

Anandamide also can produce biological responses via mechanisms that do not involve cannabinoid receptors (8, 17). It is an agonist for transient receptor potential channel vanilloid receptor subunit 1 (TRPV1) and can directly inhibit calcium channels, directly activate glutamate ionotropic receptors, and directly block activation of nicotinic α-polypeptide 7 receptor by acetylcholine. Significantly, anandamide inhibits spontaneous acrosome reactions in boar sperm during capacitation by activating TRPV1 receptors (13). Thus, alternate pathways for the actions of anandamide and other endocannabinoids in reproductive physiology and other bodily functions warrant careful examination.

Drug development

Anandamide signaling is directly involved in a myriad of physiological and pathological processes and presents many potential targets for the development of novel therapeutic drugs (18). Unfortunately, cannabinergic ligands affect almost every physiological system investigated (1). Thus, drugs acting on anandamide signaling may produce a wide variety of side effects that would limit their utility. For example, possible medicinal uses of THC are limited by its psychoactive properties. One strategy for drug development targets specific cannabinoid receptors. The CB1 antagonist SR141716 (rimonabant; trade name Acomplia) suppresses appetite (19). The European Medicines Agency recently recommended approval of Acomplia for use as a weight-loss drug. An alternate approach targets removal and degradation of anandamide (18). Basal levels of anandamide in the brain and peripheral organs are quite low (15). It is rapidly released and degraded locally. Thus, drugs that prevent anandamide reuptake and hydrolysis might be useful clinically (18). Given the results presented in the current report (9), such drugs may need to be carefully evaluated to judge their effects on women of reproductive age and those that are pregnant.

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The glomerular basement membrane: not gone, just forgotten

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The glomerular capillaries function as the filtration barrier that retains albumin and other plasma proteins in the circulation. The unresolved question that has been asked for more than 50 years is, Which structural component of these capillaries constitutes the main molecular sieve that normally retains albumin and allows its passage in diseases associated with proteinuria? There is considerable evidence implicating both the glomerular basement membrane (GBM) and the epithelial filtration slits as the barrier.

However, the prevailing point of view at present is that the slit diaphragms bridging the filtration slits are responsible for this important function, and evidence implicating the GBM is largely ignored or forgotten. In this issue of the JCI, Jarad et al. show that in laminin β2-deficient (Lamb2−/−) mice, proteinuria can be directly attributed to the altered composition of the GBM (see the related article beginning on page 2272). Changes in the permeability of the GBM and its organization were primary to changes in the epithelium, as they preceded foot process effacement and loss of slit diaphragms.

It has been more than 50 years since the early days of EM, when the ultrastructural organization of the glomerular capillary wall was defined. Yet there is still no consensus concerning which component—the slit diaphragms bridging the filtration slits or the glomerular basement membrane (GBM)—represents the primary glomerular filtration barrier. Over the intervening years, the pendulum has swung back and forth according to the interpretation of the evidence available. Since the discovery in 1999 of nephrin, a slit diaphragm protein, and its identification as the defective gene in patients with congenital nephrosis of the Finnish type (CNF) (1), the prevailing view has been that the slit diaphragms located in the filtration slits that attach the adjoining foot processes to one another contain pores responsible for the molecular sieving that prevents passage of albumin across the capillary wall. The work of Jarad et al. (2) in this issue of the JCI clearly demonstrates that defects in the composition and integrity of the GBM meshwork can lead to proteinuria, thus emphasizing the crucial role of the GBM in filtration and lending support to the concept that the fibrillar meshwork of the GBM functions as the molecular sieve that retains albumin.

The swinging of the pendulum
The first direct experimental evidence aimed at identification of the glomerular filtration barrier was based on the use of the electron-dense tracer ferritin (3, 4) and later, graded dextrans of varying sizes (MW, 32,000–125,000 Da) (5), which pinpointed the GBM as the layer that prevents passage of both of these tracers. Ferritin and dextrans of the size of albumin or greater accumulated against the luminal surface of the GBM. Later on, quite a number of different tracers were used, and most accumulated against the GBM rather than in the epithelial slits. This work was superseded by findings obtained largely by Karnovsky and coworkers using enzymatic tracers of different sizes such as catalase (MW, 220,000 Da) and HRP (MW, 40,000 Da), which were detected by a histochemical reaction (reviewed in ref. 6). These studies suffered from serious technical limitations (6), but they were interpreted as demonstrating that the filtration slits form the selective filter. Based on these studies, it was proposed that the GBM is a coarse pre-filter and the epithelial slits function as the crucial molecular sieving layer. This idea was reinforced by the discovery of a periodic structure interpreted as “slit-pores” in the slit membranes bridging the filtration slits (7). Later, Ryan and Karnovsky (8) observed that when renal blood flow was normal, plasma albumin, like ferritin and dextrans, was absent from the GBM and urinary spaces, but large amounts could be detected by immunocytochemistry in both sites if the tissue was immersion fixed or renal blood flow was interrupted by ligation of the renal artery. These studies underlined the importance of the GBM as the protein filter under normal flow conditions and helped to explain some of the authors’ earlier results that had been obtained from kidney tissue fixed by immersion. Yet the view that the GBM is a coarse pre-filter and the epithelial slits serve as a fine filter restricting the passage of albumin has lived on and remained predominant in the field.

Based on the work of Brenner and associates on the importance of charge as a determinant of glomerular filtration, interest in the 1970s and 1980s shifted to defining the charge barrier (9). A sialic acid–rich, highly negatively charged cell-surface coat was found on podocytes, and regularly spaced,