

**BIOGRAPHICAL SKETCH**

Provide the following information for the key personnel in the order listed on Form Page 2.  
Follow this format for each person. **DO NOT EXCEED FOUR PAGES.**

NAME Ken D. McCarthy, Ph.D.		POSITION TITLE Professor	
EDUCATION/TRAINING <i>(Begin with baccalaureate or other initial professional education, such as nursing, and include postdoctoral training.)</i>			
INSTITUTION AND LOCATION	DEGREE <i>(if applicable)</i>	YEAR(s)	FIELD OF STUDY
California State University at Fullerton	B.A.	1971	Biology
University of Utah	Ph.D.	1975	Pharmacology

**A. Positions and Honors:****Research and/or Professional Experience**

1975 -1978 U.S.P.H.S. Fellow, Division of Developmental Biology, University of California, Los Angeles  
 1979 -1984 Assistant Professor, Department of Pharmacology, University of North Carolina at Chapel Hill, Chapel Hill, NC  
 1985 - 1990 Associate Professor, Department of Pharmacology, University of North Carolina at Chapel Hill, Chapel Hill, NC  
 1990 - Professor, Department of Pharmacology, University of North Carolina at Chapel Hill, Chapel Hill, NC

**Professional Service**

1985 -1986 NIH Study Section, Neurobiology 2; Adhoc Member  
 1986 -1987 NIH Study Section, Neurobiology 2; Member  
 1987 - 1991 NIH Study Section, Neurological Science 2; Member  
 1987 - 1988 Program Committee member for 1988 and 1989 Winter Conference on Brain Research  
 1988 NIH Site Visit - Baltimore, MD.  
 1989 Symposium Organizer, Tissue Culture Association, entitled "In Vitro Models for Evaluating Mechanisms of Neurotoxicity".  
 1992 Program Chairman for 1993 American Society for Neurochemistry  
 1992 Premeeting Workshop Co-organizer for 1993 ASN Meeting entitled "Knockout Strategies In Molecular Neurobiology"  
 2004 NIH Study Section, Neurodegeneration and Biology of Glia; Member

**B. Selected peer-reviewed publications:**

McCarthy KD and de Vellis J. -Adrenergic receptor modulation of -adrenergic, adenosine and PGE1 increased adenosine 3':5'-cyclic monophosphate levels in primary cultures of glia. J.Cyclic Nuc.Res. 4: 15-26, 1978.  
 McCarthy KD and de Vellis J. Preparation of separate astroglial and oligodendroglial cell cultures. J.Cell Biol. 85: 890-902, 1980.  
 McCarthy KD. An autoradiographic analysis of -adrenergic receptors on immunocytochemically defined astroglia. J.Pharmacol.Exp.Ther. 226: 282-290, 1983.  
 Ingraham C and McCarthy KD. Plasticity of process-bearing glial cell cultures from neonatal rat cerebral cortical tissues. J.Neurosci. 9:63-69,1989.  
 Lerea L and McCarthy KD. Astroglial cells in vitro are heterogeneous with respect to expression of the alpha1-adrenergic receptor. Glia 2:135-147, 1989.  
 Salm AK and McCarthy KD. Expression of  $\beta$ -adrenergic receptors by astrocytes isolated from adult rat cortices. Glia 2: 346-352, 1989.

- Brooks RC, McCarthy KD and Morell P. Receptor-stimulated phospholipase  $\alpha_2$  activation is coupled to influx of external calcium and not to mobilization of intracellular calcium in C62B glioma cells. *J.Biol.Chem.* 264: 20147-20153, 1989.
- Lerea LS and McCarthy KD. Neuron-associated astroglial cells express  $\beta$ - and  $\alpha_1$ -adrenergic receptors *in vitro*. *Brain Res.* 521:7-14, 1990.
- Salm AK and McCarthy KD. Norepinephrine-evoked calcium transients in cultured type 1 astroglia. *Glia* 3:529-538, 1990.
- McCarthy KD and Salm AK. Pharmacologically distinct subsets of astrocytes identified by their calcium response to neuroligands. *Neurosci.* 41: 325-333, 1990.
- Enkvist MOK and McCarthy KD. Activation of protein kinase C blocks astroglial gap junction communication and inhibits the spread of calcium waves. *J.Neurochem.* 59:519-526, 1992.
- Shao Y and McCarthy KD. Regulation of astroglial responsiveness to neuroligands in primary culture. *Neurosci.* 55:991-1001, 1993.
- Shao Y and McCarthy KD. Quantitative relationship between  $\alpha_1$ -adrenergic receptor density and the receptor-mediated calcium response in individual astroglial cells. *Mol.Pharmacol.* 44:247-254, 1993.
- Enkvist MOK and McCarthy KD. Astroglial gap junction communication is increased by treatment with either glutamate or high  $K^+$  concentration. *J.Neurochem.* 62:489-495, 1994.
- Shao YP, Enkvist MOK and McCarthy KD. Glutamate blocks astroglial stellation: Effect of glutamate uptake and volume changes. *Glia* 11:1-10, 1994.
- Lyons S, Morrell P and McCarthy KD. Schwann cells exhibit  $P_2Y$  purinergic receptors that regulate intracellular calcium and are upregulated by cyclic AMP analogs. *J.Neurochem.* 63:552-560, 1994.
- He M and McCarthy KD. Oligodendroglial signal transduction systems are developmentally regulated. *J.Neurochem.* 63:501-508, 1994.
- Shao Y and McCarthy KD. Plasticity of astrocytes. *Glia* 11:147-155, 1994.
- Shao Y, Porter JT and McCarthy KD. Neuroligand receptor heterogeneity among astroglia. *Perspect. Dev.Neurobiol.* 2:205-215, 1994.
- Porter JT and McCarthy KD. GFAP-positive hippocampal astrocytes *in situ* respond to glutamatergic neuroligands with increases in  $[Ca^{2+}]_i$ . *Glia* 13: 101-112, 1995.
- Shao Y and McCarthy KD. Receptor-mediated calcium signals in astroglia: multiple receptors, common stores and all-or-nothing responses. *Cell Calcium* 17:187-196, 1995.
- Porter JT and McCarthy KD. Adenosine receptors modulate  $[Ca^{2+}]_i$  in hippocampal astrocytes *in situ*. *J.Neurochem.* 65:1515-1523, 1995.
- Giaume C and McCarthy KD. Control of gap-junctional communication in astrocytic networks. *Trends Neurosci.* 19:319-325, 1996.
- Porter JT and McCarthy KD. Hippocampal astrocytes *in situ* respond to glutamate released from synaptic terminals. *J.Neurosci.* 16:5073-5081, 1996.
- He M, Howe DG and McCarthy KD. Oligodendroglial signal transduction systems are regulated by neuronal contact. *J.Neurochem.* 67:1491-1499, 1996.
- Porter JT and McCarthy KD. Astrocytic neurotransmitter receptors *in situ* and *in vivo*. *Prog.Neurobiol.* 51:439-455, 1997.
- Shao Y and McCarthy KD. Responses of Bergmann glia and granule neurons *in situ* to N-methyl-Daspartate, norepinephrine, and high potassium. *J.Neurochem.* 68:2405-2411, 1997.
- Howe DG and McCarthy KD. Analysis of neuron-Schwann cell interactions at the single cell level using dicisronic retroviral vectors. *J. Neurosci. Methods*, 83: 133-142, 1998.
- Shelton MK and McCarthy KD. Mature hippocampal astrocytes exhibit functional metabotropic and ionotropic glutamate receptors *in situ*. *Glia*, 26(1): 1-11, 1999.
- Shelton MK and McCarthy KD. Hippocampal astrocytes exhibit  $Ca^{2+}$ -elevating muscarinic cholinergic and histaminergic receptors *in situ*. *J.Neurochem.* 74: 555-563, 2000.
- Howe DG and McCarthy KD. Retroviral inhibition of cAMP-dependent protein kinase inhibits myelination but not Schwann cell mitosis stimulated by interaction with neurons. *J.Neurosci.* 15: 3513-3521, 2000
- Nett, WJ, Oloff SH, and McCarthy KD. Hippocampal astrocytes *in situ* exhibit calcium oscillations that occur independent of neuronal activity. *J. Neurosci.* 22: 528-537, 2002.
- Fiacco TA and McCarthy KD. Astrocytic  $Ca^{++}$  waves increase spontaneous neuronal AMPA currents. *J. Neurosci.* 24:722-732, 2004.
- Lin, W., Kemper A, McCarthy KD, Pytel P, Wang JP, Campbell, IL, Utset MF, Popko B. Interferon-gamma induced medulloblastoma in the developing cerebellum. *J Neurosci.* 2004 Nov 10;24(45):10074-83

Fiacco TA, McCarthy KD. Intracellular astrocyte calcium waves in situ increase the frequency of spontaneous AMPA receptor currents in CA1 pyramidal neurons. *J Neurosci.* 2004 Jan 21;24(3):722-32.

Ye, P, Popken, GJ, Kemper A, McCarthy, K, Popko, B., and D'Ercole, AJ. Astrocyte-specific overexpression of insulin-like growth factor-1 promotes brain overgrowth and glial fibrillary acidic protein expression. *J Neuroscience Res.* 78:472-484, 2004.

Pascual, O, Casper, K, Kubera, C, Zhang, J, Revilla-Sanchez R, Sul, J-Y, Takano, H, Moss, SJ, McCarthy, KD, and Haydon, PG. Astrocyte Purinergic Signaling Coordinates Synaptic Networks, *Sci*, 310 (5745):113-116, 2005

Casper KB and McCarthy, KD. GFAP-positive progenitor cells produce neurons and oligodendrocytes throughout the CNS. *Mol Cell Neurosci.* Feb. 2006

### **C. Current Grant Support**

NS033938-06                      4/04 – 3/09

NIH

Priority Score: 2.8%

Title: Astrocyte Regulation of Neuronal Activity In Vivo

Thrust: To develop and use conditional gene knockouts to assess the role of astrocytes in regulating neuronal activity.

NS020212                      12/05 – 11/09

NIH

Priority Score: 11%

Title: Neuronal-Astrocytic Communications In Vivo

Thrust: To use multiphoton imaging, patch clamp electrophysiology, caged molecules and genetically-modified mice to examine the role of astrocytes in synaptic transmission.