

**Emily Barron-Casella, Ph.D.**  
**Assistant Professor of Pediatrics, Division of Hematology**  
**The Johns Hopkins University School of Medicine**

**DEMOGRAPHIC INFORMATION**

**Current Appointment:**

Assistant Professor of Pediatrics, Division of Hematology

**Personal Data:**

Address:  
Department of Pediatrics, Division of Hematology  
Ross 1129  
720 Rutland Avenue  
Baltimore, MD 21205

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**Education and Training:**

<u>Degree/Year</u>	<u>Institution</u>	<u>Discipline</u>
B.A./1976	Austin College	Biology
M.S./1981	Texas Tech U. Health Science Center	Anatomy
Ph.D./1988	UT Health Science Center at San Antonio	Cellular and Structural Biology

**Predoctoral Training**

1976-1977	Laboratory Technician Department of Biochemistry Texas Tech University Health Sciences Center Lubbock, Texas
1980-1981	Graduate Assistant Department of Anatomy Texas Tech University Health Sciences Center Lubbock, Texas
1981-1983	Research Assistant Department of Genetics Wadley Institutes of Molecular Medicine

Dallas, Texas

1983-1988            Teaching Assistant  
Department of Cellular and Structural Biology  
University of Texas Health Science Center  
San Antonio, Texas

### Postdoctoral Training

1988-1992            Postdoctoral Fellow  
Department of Molecular Biology and Genetics  
Johns Hopkins University School of Medicine  
Baltimore, Maryland

Postdoctoral Associate  
Howard Hughes Medical Institute  
Baltimore, Maryland

### Professional Experience:

<u>Positions</u>	<u>Institution</u>	<u>Dates</u>
Adjunct Investigator	Pediatric Oncology Branch, Division of Clinical Sciences, NCI, Bethesda, Maryland	1995-2003
Assistant Professor	Johns Hopkins U. School of Medicine	1992-present

## RESEARCH ACTIVITIES

### Publications:

#### Peer reviewed

1. Morrow, J., Stocco, D., and **Barron, E.A.** Spontaneous mutation rate to thioguanine resistance is decreased in polyploid hamster cells. *J. Cell Physiol.* 96: 81-86, 1978.
2. Morrow, J., Sammons, D. and **Barron, E.A.** Puromycin resistance in Chinese hamster cells: Genetic and biochemical studies of partially resistant, unstable clones. *Mutation Res.* 69: 333-346, 1980.
3. Nathan, R.D., Fund, S.J., Stocco, D.M., **Barron, E.A.**, and Markwald, R.R. Sialic Acid: Regulation of electrogenesis in cultured heart cells. *Am. J. Physiol.* 239: C197-207, 1980.

4. **Barron, E.A.**, Markwald, R.R., and Nathan, R.D. Structural analysis on the surface of in vitro embryonic myocardial cell aggregates. J. Mol. Cell. Cardiol. 14: 382-395, 1983.
5. **Barron, E.A.**, Sidhu, R.S., and Bollon, A.P. Construction of a yeast-bacterial DNA cloning vector. J. Clin. Hematol. and Oncol. 13: 53-64, 1983.
6. Cao, Z., **Barron, E.A.**, Carillo, A.J., and Sharp, Z.D. Reconstitution of cell-type-specific transcription of the rat prolactin gene in vitro. Mol. Cell. Biol. 7: 3402-3408, 1986.
7. Cao, Z., **Barron, E.A.**, and Sharp, Z.D. Prolactin upstream factor I mediates cell-specific transcription. Mol. Cell. Biol. 8: 5432-5438, 1988.
8. **Barron, E.A.**, Cao, Z., Schneider, K.E., Carillo, A.J., and Sharp, Z.D. Dual functions of a cis-acting element within the rat prolactin gene promoter. Mol. Cell. Biol. 9:817-819, 1989.
9. Sharp, Z.D., Hesel, S., Cao, A., **Barron, E.A.**, and Sanchez, Y. DNA recognition element required for PUF-I mediated cell-type-specific transcription of the rat prolactin gene. Nucl. Acid Res. 17: 2705-2722, 1989.
10. **Barron, E.A.**, and Corden, J. Conservation of the mammalian RNA polymerase II largest-subunit C-terminal domain. J. Mol. Evol. 35: 405-410, 1992.
11. **Barron-Casella, E.A.**, Kickler, T.S., Rogers, O.C., and Casella, J.F. Expression and purification of functional recombinant epitopes for the platelet antigens, PI<sup>A1</sup> and PI<sup>A2</sup>. Blood 84: 1157-1163, 1994.
12. Casella, J.F., **Barron-Casella, E.A.**, and Torres, M.A. Quantitation of Cap Z in conventional preparations and methods for further purification of actin. Cell Motil. Cytoskele-ton 30: 164-170, 1995.
13. **Barron-Casella, E.A.**, Torres, M.A., Scherer, S.W., Heng, H.H.Q., Tsui, L-C, Casella, J.F. Sequence analysis and chromosomal localization of human Cap Z: Conserved residues within actin-binding domain may link Cap Z to the gelsolin/severin and profilin protein families. J. Biol. Chem., 20: 21472-21479, 1995.
14. Nachmuis, V.T., Golla, R., Casella, J.F., and **Barron-Casella, E.A.** Cap Z, a calcium insensitive protein in resting and activated platelets. FEBS Letters 378: 258-262, 1996.
15. Okamoto, N., Kennedy, S.D., **Barron-Casella, E.A.**, Casella, J.F., Inoko, H., Kickler, T.S. Identification of a human heavy chain antibody fragment directed against human platelet alloantigen 1a by phage display library. Tissue Antigens 51:156-163, 1998.
16. **Barron-Casella, E.A.**, Nebbia, G., Rogers, O.C., King, K.E., Kickler, T.S., Casella, J.F. Construction of a Human Platelet Alloantigen-1a epitope(s) within murine glycoprotein IIIa:

Identification of residues critical to the conformation of the antibody binding site(s). Blood 93:2959-2967, 1999.

17. Lu, J.F., **Barron-Casella, E.**, Deering, R., Heinzer, A.K., Moser, A.B., deMesy Bentley, K.L., Wand, G.S., McGuinness, M., Pei, Z., Watkins, P.A., Pujol, A., Smith, K.D., Powers, J.M. The role of peroxisomal ABC transporters in the mouse adrenal gland: the loss of Abcd2 (ALDR), not Abcd1 (ALD), causes oxidative damage. *Lab Invest.* 87:261-272, 2007.

#### Book Chapter

Bollon, A.P., **Barron, E.A.**, Berent, S.L., Bragg, P.W., Dixon, D., Fuke, M., Hendrix, C., Mahmoudi, M., Sidhu, R.S., Torczynski, R.M. Recombinant DNA techniques: Isolation, Cloning, and Expression of Genes. In: *Recombinant DNA Products: Insulin, Interferon, Growth Hormone*, A.P. Bollon, Ed. (CRC Press, Inc., Boca Raton, FL), 1984.

#### Extramural Sponsorship:

##### Past Awards

Identification of the Epitopes and Antibodies Involved in the HPA-3 (Bak) and HPA-4 (Pen) Alloantigen Systems, 07//01/96-06/30/97, National Blood Foundation grant (E. Barron-Casella, PI)

American Heart Established Investigator Award (E. Barron-Casella, PI)  
Returned/overlap

Modeling of Platelet Alloantigens, 07/01/1998-06/30/2003, NHLBI R29HL59955 (E. Barron-Casella, PI)

Fetal-Maternal Incompatibility of Platelets, 07/01/1999-06/30/02, Advanced Transfusion Practices and Blood Research grant (E. Barron-Casella, PI)

##### Active

U01-NS-042804 (DeBaun M, PI) 10/01/2003-11/30/2012  
5.16 calendar months  
NINDS/NIH  
Silent Cerebral Infarct Multi-Center Clinical Trial

1 R01 HL091759-01 (Casella J, PI) 12/01/2007-11/30/2011  
1.2 calendar months  
NIH/NHLBI  
Longitudinal SIT Trial Plasma Proteomic Biomarker Discovery and Validation in SCI

##### Pending

RO1(Smith KD, Co-PI, E. Barron-Casella, Co-PI) 07/01/08-06/30/13  
6.0 calendar months  
NIH  
XALD: Pathophysiology and Therapy

## EDUCATIONAL ACTIVITIES

### Teaching:

Faculty member of the Cellular and Molecular Medicine graduate program from 10/1999-10/2002.

### Training grant participation:

*Laboratory Research Training in Pediatric Oncology-Hematology*  
8/98-5/03  
National Institutes of Health  
National Cancer Institute  
T32CA60441  
Principal Investigator: Curt Civin, M.D.  
Program Faculty: Emily Barron-Casella, Ph.D.

### Advisees

Swati Jain, MD  
Neil Littman  
Katherine Dix  
Jamey Ingraham  
Anthony Law

## ORGANIZATIONAL ACTIVITIES

### Institutional Administrative Appointments:

1999-2003	Pediatric Core Laboratory Director
2006-present	SITT Repository Supervisor

## RECOGNITION

### Invited talks

March 1994	<i>Fetal-Maternal Incompatibility of Platelets</i> Austin College Lectureship
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Sherman, Texas

December 1997

*Development of an HPA-1a Epitope within Murine GPIIIa: Identification of Residues Necessary for Generating the Human Antigenic Determinant*  
Platform session  
American Society of Hematology meeting  
San Diego, California