# CHEMICAL HERITAGE FOUNDATION

# **ROGER E. KARESS**

The Pew Scholars Program in the Biomedical Sciences

Transcript of an Interview Conducted by

Andrea R. Maestrejuan at

Centre de Génétique Moléculaire Gif-sur-Yvette, France

on

23, 24, 25, and 26 April 1996

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# ACKNOWLEDGEMENT

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# **ROGER E. KARESS**

1955	Born in New York City, New York, on 28 March
	Education
1976 1980	BS, Biochemistry, Yale University PhD, Biochemistry, Rockefeller University
	Professional Experience
1980-1983	Carnegie Institution of Washington, Department of Embryology Postdoctoral Fellow
1983-1986	Imperial College of Science and Technology, University of London Research Fellow, Department of Biochemistry
1986-1992	New York University School of Medicine, Department of Biochemistry, Assistant Professor
1992-present	Centre de Génétique Moléculaire, Centre National de la Recherche Scientifique, Directeur de recherche

## Selected Publications

- Karess, R. et al., 1979. Cellular information in the genome of recovered avian sarcoma virus directs the synthesis of transforming protein. *Proceedings of the National Academy of Sciences USA* 76:3154-58.
- Rettenmier, C. et al., 1979. Tryptic peptide analysis of avian oncovirus *gag* and *pol* gene products. *Journal of Virology* 32:102-13.
- Karess, R. and H. Hanafusa, 1981. Viral and cellular *src* genes contribute to the structure of recovered avian sarcoma virus transforming protein. *Cell* 24:155-64.
- Poirier, F. et al., 1982. Role of p60<sup>src</sup> kinase activity in the induction of neuroretinal cell proliferation by Rous sarcoma virus. *Journal of Virology* 42:780-89.
- Karess, R. and G. Rubin, 1982. A small tandem duplication is responsible for the unstable *white-ivory* mutation in *Drosophila*. *Cell* 30:63-69.
- Karess, R.E. and D.M. Glover, 1984. Analysis of P transposable element functions in *DrosophilaCell* 38:135-46.
- Karess, R. and D.M. Glover, 1989. *Rough-deal:* A gene required for proper mitotic segregation of chromosomes in *Drosophila. Journal of Cell Biology*109:2951-61.

- Medina-Acosta, E. et al., 1989. The promastigote surface protease (gp63) of *Leishmania* is expressed but differentially processed and localized in the amastigote stage. *Molecular and Biochemical Parasitology* 37:263-74.
- Karess, R. et al., 1991. The regulatory light chain of nonmuscle myosin is encoded by *spaghetti-squash*, a gene required for cytokinesis in *Drosophila*. *Cell* 65:1177-89.
- Llamazares, S. et al., 1991. *Polo* encodes a protein kinase homolog required for mitosis in *Drosophila. Genes and Development* 5:2153-65.
- Medina-Acosta, E. et al., 1993. Structurally distinct genes for the surface protease *Leishmania mexicana* are developmentally regulated. *Molecular and BiochemicalParasitology* 57:31-45.
- Gomes, R. et al., 1993. Abnormal anaphase resolution (aar): A locus required for progression through mitosis. *Journal of Cell Science* 104:583-93.
- Wheatley, S. et al., 1995. *Drosophila* nonmuscle myosin II is required for rapid cytoplasmic transport during oogenesis and for axial nuclear migration in early embryos. *Development* 121:1937-46.
- Jordan, P. and R. Karess. Site-directed mutagenesis of *Drosophila* nonmuscle myosin light chain suggests a requirement for myosin phosphorylation during oogenesis. (Submitted)

### ABSTRACT

**Roger E. Karess** and his two older sisters grew up in Great Neck, New York. Their grandparents were Jews from Eastern Europe, and their neighborhood consisted of other family members and people with similar backgrounds. Karess's father had a law degree but did not practice; he worked in a family business for many years and then was in insurance. His mother was a homemaker. Both parents were adamant that all three children would go to college. The older sister is a chemist in industry; the younger, after a dancing career, became a social worker. Karess discusses his upbringing as a Reform Jew; Europeans' attitudes toward Americans; racism and anti-Semitism in Europe; Karess' Jewish identity; and Roman Catholic influences on contemporary France.

Karess cannot remember not being interested in science. He enjoyed the experiments in elementary school and reading the life stories of great scientists in Paul de Kruif's *Microbe Hunters*. His fourth-grade teacher noted his "passion" for science. In high school he took advanced science courses, and he attended a summer program for high school students at Jackson Laboratory; there he studied the effects of heavy metals on mouse embryo development and was introduced to reading scientific articles. He also attended a high-school science program at Columbia University.

He was accepted at Yale University where he worked in David Ward's lab studying paroviruses. He talks about the difference between liking science and doing science; about his regret at not having taken more lab classes at Yale and about having taken courses in medieval Latin and art history. He developed an interest in tumor viruses and wrote a class paper on host virus restriction. He talks more about working in the Ward lab; about having worked on reverse transcriptase in the Ted Reid lab; and about letters of recommendation he received from Yale professors.

He entered graduate school at Rockefeller University; he began early in Vincent Allfrey's lab so as to gain more lab experience. He then transferred to the Hidesaburo Hanafusa lab to study retroviruses. Here he discusses changes in his confidence as a scientist over time; his evaluation of himself as an undergraduate researcher; undergraduates in his own lab; his performance on his senior exam; his reasons for selecting Rockefeller for graduate school; Rockefeller's unstructured program; and playing softball at Rockefeller with Mark Rieman and jogging with Michael Greenberg. He goes on to describe Hanafusa as a teacher and a mentor and Hanafusa's research on tumor viruses. Karess himself sought to identify the RNA binding site for retroviruses but was thwarted by technical difficulties. Karess then talks about how William Hayward distinguishes between transformation-competent and transformationdefective virus cells; how Peter Duesberg's radiolabeling of viral RNA helps demonstrate the existence of an oncogene; Hanafusa's research on proto-oncogenes; how Karess seeks to isolate the src protein; and Raymond Erikson's discovery that src is a kinase. Karess was challenged in his attempt to identify the first known kinase and unable at the time to discover the fps oncogene. This leads to an explanation of the factors involved in scientific breakthroughs and the need to interpret data with fresh, objective eyes. He evaluates his self-confidence at the end of his doctorate.

Here Karess gives his opinion on the constructive and destructive effects of competition in science and the need to take risks in research. He goes on to compare the structures of scientific research in France and the United States; the advantages and disadvantages of doing research in France; and the relative prestige of publishing in American and European journals.

Karess accepted a position as a principal investigator at the Centre de Génétique Moléculaire (CGM) near Paris. When he published an article in *Cell* he encountered the politics of scientific publishing. He goes on to describe funding in France; the Centre National de la Recherche Scientifique (CNRS) and setting up a lab at CNRS; and his own funding. More discussion of the funding of scientific research in France leads to a discussion of Karess's funding in the United States and his opinion about the need for reforms in the way science is done in both France and the United States.

Karess's research interests shifted from oncogenes to *Drosophila* genetics, and he developed an interest in transposable elements. He accepted a postdoc position in the Gerald Rubin lab at the Carnegie Institution of Washington, where he studied unstable alleles in *Drosophila*. Rubin's discovery of P elements revolutionized *Drosophila* genetics. Karess analyzed P transposable element function. He then accepted a second postdoc position in David Glover's lab at Imperial College of Science and Technology, University of London. He talks about trends in assigning names to *Drosophila* genes and the names Karess and others created.

Karess applied for his first academic position and accepted an offer from New York University. Here he discusses the people in his own lab. He took up studying *Leishmania*. Karess moved his lab to the CGM in Paris, where he has been studying the *rough-deal* gene. Karess concludes with an assessment of his scientific research.

### UCLA INTERVIEW HISTORY

#### **INTERVIEWER:**

Andrea R. Maestrejuan, Interviewer, UCLA Oral History Program. B.A., History, University of CaliforniaIrvine, 1988; B.S., Biological Sciences, University of California, Irvine, 1988; C.Phil., History, University of California, Riverside.

### TIME AND SETTING OF INTERVIEW:

Place: Karess's office, Centre de Génétique Moléculaire, Gif-sur-Yvette, France.

**Dates, length of sessions:** April 23, 1996 (150 minutes); April 24, 1996 (157) ; April 25, 1996 (170) ; April 26, 1996 (118).

### **Total number of recorded hours: 9.9**

## Persons present during interview: Karess and Maestrejuan.

### CONDUCT OF INTERVIEW:

This interview is one in a series with Pew scholars in the biomedical sciences conducted by the UCLA Oral History Program in conjunction with the Pew Charitable Trusts's Pew Scholars in the Biomedical Sciences Oral History and Archives Project. The Project has been designed to document the backgrounds, education, and research of biomedical scientists awarded four-year Pew scholarships since 1988.

To provide an overall framework for Project interviews, the director of the UCLA Oral History Program and three UCLA faculty project consultants developed a topic outline. In preparing for this interview, Maestrejuan held a telephone preinterview conversation with Karess to obtain written background information (curriculum vitae, copies of published articles, etc.) and to agree on an interviewing schedule. She also reviewed prior Pew scholars' interviews and the documentation in Karess's file at the Pew Scholars Program office in San Francisco, including his proposal application, letters of recommendation, and reviews by Pew Scholars Program national advisory committee members.

For general background on the recent history of the biological sciences, Maestrejuan consulted J.D. Watson et al., *Molecular Biology of the Gene*. 4th ed. Menlo Park, CA: Benjamin/Cummings, 1987, and Bruce Alberts et al., *Molecular Biology of the Cell*. 3rd ed. New York: Garland, 1994.

The interview is organized chronologically, beginning with Karess's childhood in New York and continuing through his education at Yale University and Rockefeller University, his postdocs at Carnegie Institution of Washington and the Imperial College of Science and Technology, and his positions at New York University and the Centre de Génétique Moléculaire. Major topics discussed include the molecular biology of oncogenes, *Drosophila* genetics, and the infrastructures of scientific research in France and the United States.

### ORIGINAL EDITING:

Gregory M. Beyrer, editorial assistant, edited the interview. He checked the verbatim transcript of the interview against the original tape recordings, edited for punctuation, paragraphing, and spelling, and verified proper names. Words and phrases inserted by the editor have been bracketed.

Karess reviewed the transcript. He verified proper names and made minor corrections and additions.

Steven J. Novak, senior editor, prepared the table of contents and interview history.

Beyrer assembled the biographical summary and index.

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