



## Knowledge Inn (in nature)

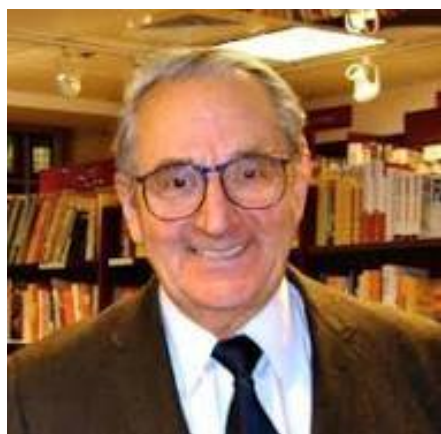
 Research Profile of Elias James Corey (Nobel Laureate)

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### Elias James Corey

Born on 12 July 1928

Methuen, Massachusetts, United States



#### Affiliation

Harvard University, Cambridge, MA, USA

h-index	<b>140</b>
Research papers	246 Up to2016; 1048 High-probability publications From 1950-2011

Nobel Prize in  
Chemistry

Retrosynthetic analysis development of the theory and methodology of  
organic synthesis

**For the greatest benefit to humankind**

1990

### Typical Book titles of Elias James Corey

The Logic of Chemical Synthesis. E. J. Corey and Xue-Min Cheng. John Wiley & Sons Ltd, Chichester, 1989. 436 pp.

Molecules and Medicine E. J. Corey, E. J. Corey, Czakó, Barbara, Lárti, szló, Amazon.com Services LLC

Enantioselective Chemical Synthesis, Methods, Logic, and Practice, Elias J. Corey Laszlo Kurti, 1st Edition, Hardcover ISBN: 9780615395159, Imprint: Academic Press, 15th October 2013, pages: 334

### Academic profile of Elias James Corey

High School	1945	Lawrence High School, Lawrence, MA
<b>Education ( MIT : 1945-1950)</b>		
BS Organic Chemistry	1948	Massachusetts Institute of Technology
PhD Chemistry	1951	Massachusetts Institute of Technology

### Employment. Elias James Corey

#### Faculty member at the University of Illinois (1951 to 1959)

1951	Instructor, University of Illinois at Urbana-Champaign
1954	Assistant professor
1956	Full professor University of Illinois at Urbana-Champaign
1957	Guggenheim Fellowship: Harvard University on sabbatical leave
1958	Fellow: Chemistry, Royal Caroline Institute

#### Faculty member at Harvard University (1959- )

1959-	Professor: Chemistry, Harvard University
1965	Chairman of the department & Sheldon Emory Professor of Organic Chemistry

### Post-Doc career

1948 John C. Sheehan's program on synthetic penicillins, after his graduation

1951 Instructor in chemistry under Roger Adams and Carl S. Marvel at University of Illinois at Urbana-Champaign

He began his work in physical organic chemistry, developing stereo-electronic ideas which are

	<p>still used even today</p> <p>Corey began working on his own at Illinois by applying theories of electron density in molecules and transition states (known as molecular orbital theory) to make predictions about reaction products</p>
	<b>Supervising graduate students</b>
1954	<p>After getting promotion as assistant professor, he started supervising graduate students in experimental projects involving the structure, stereochemistry, and synthesis of complex naturally occurring organic compounds. His success was astounding; promoted to full professor in 1956, at age 27.</p> <p>His research group grew and its activities included new areas of synthesis, metal complexes and enzyme chemistry</p>
	<b>Fellowship &amp; sabbatical leave</b>
1957	<p>During Guggenheim Fellowship period (divided between Harvard and Europe), several key ideas about chemical synthesis emerged. He went first to Robert B. Woodward (on invitation) at Harvard University; then to Switzerland, England, and finally to Sweden. While researching with Sune Bergström, he became intrigued by prostaglandins. Corey synthesized prostaglandins in the mid 1960s.</p>

**Awards (to) Elias James Corey**

<b>Nobel Prize in Chemistry</b>	<b>Retrosynthetic analysis development of the theory and methodology of organic synthesis</b> <b>For the greatest benefit to humankind</b>	<b>1990</b>
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<b>Award</b>	<b>year</b>
ACS Linus Pauling Medal	1973
Benjamin Franklin Medal	1978
Wolf Prize in Chemistry	1986
RSC Robert Robinson Medal	1988
Priestley Medal	2004
.....	70 awards

Press release

No other chemist than Corey has developed such a comprehensive assortment of methods... in the synthesizing laboratory

A father of modern organic synthesis







## Expertise





- 📖 Molecular synthesis
- 📖 Development of several new synthetic reagents
- 📖 Methodology of organic synthesis
- 📖 Significantly improved on the science of organic synthesis
- 📖 Corey-Bakshi-Shibata reaction for the reduction of ketones to secondary alcohols
  
- 📖 **Computer software + Synthesis**
- 📖 One among the first few chemists to use computer analysis to design chemical syntheses
- 📖 Corey went on to pioneer the use of computers in retrosynthetic analysis

Retrosynthetic Analysis (October 1957 to 1970)	
Traditional syntheses	<ul style="list-style-type: none"><li>○ Starts with simple (or at least readily available) molecules i.e. building blocks</li><li>○ Assembled by a sequence of reactions</li><li>○ Final step leading to formation of the desired (complicated) target molecule</li></ul>
Retrosynthetic analysis	<p><b>Phase I</b></p> <ul style="list-style-type: none"><li>○ Breaking a target molecule theoretically into simpler precursor structures</li><li>○ Continuing the disassembly process until simple starting materials were arrived at</li></ul> <p><b>Phase II</b></p> <ul style="list-style-type: none"><li>○ Stepwise synthesis</li><li>○ A combination of intelligent selection step based on known reactions of related molecules</li><li>○ Experimental trial and error method tried until successful</li></ul>
Outcome	<ul style="list-style-type: none"><li>📖 This synthetic strategy subsequently became the standard norm for organic synthesis</li><li>📖 Developed more than one hundred synthetic reactions and transformations by himself</li><li>📖 Development of numerous medicinal drugs, including prostaglandins and ginkgolide</li><li>📖 Generation of potential synthetic pathways</li><li>📖 Computer-assisted synthetic analysis grown into a new paradigm; widespread; now indispensable omnipotent tool</li></ul>

## Elias James Corey (Open-heart musings)

- 📖 I have always been guided by a desire to be a worthy son to the father and courageous mother who raised me
- 📖 Though my given name was William James Corey, my mother (Fatima Hasham Corey b. 1900,

d. 1970 ) renamed me Elias after my father's death (businessman, d. 1930) due to pneumonia just eighteen months after my birth.
 My aunt taught me to be efficient and to take pleasure in a job well done, no matter how mundane.
 In 1945, I entered MIT in engineering. I was a keen mathematician. But, soon became a convert to organic chemistry, with its "intrinsic beauty and great relevance to human health"
 Wife: Claire Higham Corey (Graduate from University of Illinois; m. Sep-1961)
 Son: David Reid Corey (Biochemist, b. 1963)
 Son: John Corey (b. 1965)
 Daughter: Susan Corey (Anthropologist, b. 1967)
b: born; d: dead; m: married

<b>Source: Elias J. Corey's speech at the Nobel Banquet, December 10, 1990</b>
There is a children's poem by Julia Carney entitled "Little Things" ; one verse of which reads:
<p>Little drops of water, little grains of sand          Make the mighty ocean and the pleasant land          Thus the little minutes, humble though they be          Make the mighty ages of eternity.</p>
 It is by countless small steps, like the little grains of sand, that we endeavor to understand ourselves and our universe, though confronted by endless complexity  Whatever our field – science, technology, humanities, art – we are excited by the challenge of research at an endless frontier  And also humbled by the vastness of our ignorance  We are heartened by the belief that knowledge, if wisely used, can benefit mankind
<ul style="list-style-type: none"> <li>○ Naturally occurring organic substances are the basis of all life on earth</li> <li>○ Science at the molecular level defines a fundamental language of life</li> <li>○ Most of today's medicines are synthetic</li> <li>○ Really, majority of tomorrow's will be conceived and produced by synthetic chemists</li> <li>○ The knowledge of synthetic chemistry is a probe for the attainment of a deep understanding of matter, chemical change and life</li> </ul>

<b>Select titles of research output of Elias James Corey high impact (# citations) publications</b>	
<b>Year</b>	<b>Publication</b>
2016	Mahender Reddy K, Bhimireddy E, Thirupathi B, Breitler S, Yu S, <b>Corey EJ</b> . Cationic Chiral Fluorinated Oxazaborolidines. More Potent, Second Generation Catalysts for Highly Enantioselective Cycloaddition Reactions. <i>Journal of the American Chemical Society</i> . PMID <a href="https://pubmed.ncbi.nlm.nih.gov/26812167/">26812167</a> DOI: <a href="https://doi.org/10.1021/jacs.6b00100">10.1021/jacs.6b00100</a>
2015	Rajendar G, <b>Corey EJ</b> . A systematic study of functionalized oxiranes as initiating groups for

	cationic polycyclization reactions. <i>Journal of the American Chemical Society</i> . 137: 5837-44. PMID <a href="#">25871500</a> DOI: <a href="#">10.1021/jacs.5b03229</a>
2014	Han Y, Ma Y, <a href="#">Keresztes I</a> , <a href="#">Collum DB</a> , <b>Corey EJ</b> . Preferential geminal bis-silylation of 3,4-benzothiophane is caused by the dominance of electron withdrawal by R3Si over steric shielding effects. <i>Organic Letters</i> . 16: 4678-9. PMID <a href="#">25157594</a> DOI: <a href="#">10.1021/ol502348y</a>
2014	Surendra K, <b>Corey EJ</b> . Diiodoindium(III) cation, InI <sub>2</sub> <sup>+</sup> , a potent yneophile. Generation and application to cationic cyclization by selective π-activation of C≡C. <i>Journal of the American Chemical Society</i> . 136: 10918-20. PMID <a href="#">25095905</a> DOI: <a href="#">10.1021/ja506502p</a>
2013	Li JJ, <b>Corey EJ</b> . Drug Discovery: Practices, Processes, and Perspectives <i>Drug Discovery: Practices, Processes, and Perspectives</i> . DOI: <a href="#">10.1002/9781118354483</a>
2013	<b>Corey EJ</b> , <a href="#">Kurti L</a> . Enantioselective Chemical Synthesis: Methods, Logic, and Practice <i>Enantioselective Chemical Synthesis: Methods, Logic, and Practice</i> . 1-328.
2012	<a href="#">Newhouse TR</a> , Li X, <a href="#">Blewett MM</a> , Whitehead CM, <b>Corey EJ</b> . A tetradentate ligand for the enantioselective Ti(IV)-promoted oxidation of sulfides to sulfoxides: origin of enantioselectivity. <i>Journal of the American Chemical Society</i> . 134: 17354-7. PMID <a href="#">23046346</a> DOI: <a href="#">10.1021/ja305991k</a>
2012	Surendra K, <b>Corey EJ</b> . Highly enantioselective proton-initiated polycyclization of polyenes. <i>Journal of the American Chemical Society</i> . 134: 11992-4. PMID <a href="#">22780430</a> DOI: <a href="#">10.1021/ja305851h</a>
2011	Qiu WW, Surendra K, Yin L, <b>Corey EJ</b> . Selective formation of six-membered oxa- and carbocycles by the In(III)-activated ring closure of acetylenic substrates. <i>Organic Letters</i> . 13: 5893-5. PMID <a href="#">21992657</a> DOI: <a href="#">10.1021/ol202621g</a>
2011	Liang H, Hu L, <b>Corey EJ</b> . Di-tert-butylisobutylsilyl, another useful protecting group. <i>Organic Letters</i> . 13: 4120-3. PMID <a href="#">21744824</a> DOI: <a href="#">10.1021/ol201640y</a>
2011	Li JJ, <b>Corey EJ</b> . Name Reactions in Heterocyclic Chemistry II <i>Name Reactions in Heterocyclic Chemistry II</i> . DOI: <a href="#">10.1002/9781118092828</a>
2010	<a href="#">Shenvi RA</a> , <b>Corey EJ</b> . Synthetic access to bent polycycles by cation-π cyclization. <i>Organic Letters</i> . 12: 3548-51. PMID <a href="#">20670018</a> DOI: <a href="#">10.1021/ol101410g</a>
2009	<a href="#">Baqi Y</a> , Giroux S, <b>Corey EJ</b> . A study of the epoxidation of cycloolefins by the t-BuOH copper-permanganate system. <i>Organic Letters</i> . 11: 959-61. PMID <a href="#">19182889</a> DOI: <a href="#">10.1021/ol802923n</a>
2008	Giroux S, <b>Corey EJ</b> . Enantioselective synthesis of a simple benzenoid analogue of glycinoclepin A. <i>Organic Letters</i> . 10: 5617-9. PMID <a href="#">19007174</a> DOI: <a href="#">10.1021/ol8024633</a>
2007	Ryu DH, Kim KH, Sim JY, <b>Corey EJ</b> . Catalytic enantioselective Diels-Alder reactions of furans and 1,1,1-trifluoroethyl acrylate <i>Tetrahedron Letters</i> . 48: 5735-5737. DOI: <a href="#">10.1016/j.tetlet.2007.06.097</a>
2006	Mushti CS, Kim JH, <b>Corey EJ</b> . Total synthesis of antheliolide A. <i>Journal of the American Chemical Society</i> . 128: 14050-2. PMID <a href="#">17061886</a> DOI: <a href="#">10.1021/ja066336b</a>
2005	Reddy LR, <b>Corey EJ</b> . Facile air oxidation of the conjugate base of rofecoxib (Vioxx™), a possible contributor to chronic human toxicity <i>Tetrahedron Letters</i> . 46: 927-929. DOI: <a href="#">10.1016/j.tetlet.2004.12.055</a>

2005	Li JJ, <b>Corey EJ</b> . <b>Name Reactions in Heterocyclic Chemistry</b> <i>Name Reactions in Heterocyclic Chemistry</i> . 1-558. DOI: <a href="https://doi.org/10.1002/0471704156">10.1002/0471704156</a>
2000	Busch-Petersen J, <b>Corey EJ</b> . Lithium N-trityl-N-(R)-1-phenethylamide, a readily available and useful base for the enantioselective formation of chiral enolates from achiral ketones <i>Tetrahedron Letters</i> . 41: 6941-6944.
1999	Han X, <b>Corey EJ</b> . A catalytic enantioselective total synthesis of (-)-wodeshiol. <i>Organic Letters</i> . 1: 1871-2. PMID <a href="https://pubmed.ncbi.nlm.nih.gov/10836046/">10836046</a>
1996	<b>Corey EJ</b> , Matsuda SP, <a href="#">Baker CH</a> , <a href="#">Ting AY</a> , Cheng H. Molecular cloning of a Schizosaccharomyces pombe cDNA encoding lanosterol synthase and investigation of conserved tryptophan residues. <i>Biochemical and Biophysical Research Communications</i> . 219: 327-31. PMID <a href="https://pubmed.ncbi.nlm.nih.gov/8604986/">8604986</a> DOI: <a href="https://doi.org/10.1006/bbrc.1996.0232">10.1006/bbrc.1996.0232</a>
1995	<b>Corey EJ</b> , <a href="#">Roberts JD</a> . John Clark Sheehan - September 23, 1915-March 21, 1992 <i>Biographical Memoirs. National Academy of Sciences (U.S.)</i> . 68: 291-302. PMID <a href="https://pubmed.ncbi.nlm.nih.gov/11616354/">11616354</a>
1983	Systemic hemodynamic effects of leukotrienes C4 and D4 in the rat <i>American Journal of Physiology - Heart and Circulatory Physiology</i> Published 1 April 1983 Vol. 244 no. 4, H628-H633
1982	Drazen JM, Venugopalan CS, Austen KF, <a href="#">Brion F</a> , <b>Corey EJ</b> . Effects of leukotriene E on pulmonary mechanics in the guinea pig. <i>The American Review of Respiratory Disease</i> . 125: 290-4. PMID <a href="https://pubmed.ncbi.nlm.nih.gov/6279000/">6279000</a>
1981	Levine L, Morgan RA, Lewis RA, Austen KF, Clark DA, <a href="#">Marfat A</a> , <b>Corey EJ</b> . Radioimmunoassay of the leukotrienes of slow reacting substance of anaphylaxis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> . 78: 7692-6. PMID <a href="https://pubmed.ncbi.nlm.nih.gov/6950409/">6950409</a>
1980	Rådmark O, Malmsten C, Samuelsson B, Clark DA, Goto G, <a href="#">Marfat A</a> , <b>Corey EJ</b> . Leukotriene A: stereochemistry and enzymatic conversion to leukotriene B. <i>Biochemical and Biophysical Research Communications</i> . 92: 954-61. PMID <a href="https://pubmed.ncbi.nlm.nih.gov/6244821/">6244821</a> DOI: <a href="https://doi.org/10.1016/0006-291X(80)90795-0">10.1016/0006-291X(80)90795-0</a>
1976	<b>Corey EJ</b> , <a href="#">Narasaka K</a> , <a href="#">Shibasaki M</a> . A direct, stereocontrolled total synthesis of the 9,11-azo analogue of the prostaglandin endoperoxide, PGH <sub>2</sub> . <i>Journal of the American Chemical Society</i> . 98: 6417-8. PMID <a href="https://pubmed.ncbi.nlm.nih.gov/965655/">965655</a>
1975	<b>Corey EJ</b> , <a href="#">Ensley HE</a> . Letter: Preparation of an optically active prostaglandin intermediate via asymmetric induction. <i>Journal of the American Chemical Society</i> . 97: 6908-9. PMID <a href="https://pubmed.ncbi.nlm.nih.gov/1184891/">1184891</a>
1972	<b>Corey EJ</b> , <a href="#">Snider BB</a> . A total synthesis of ( )-fumagillin. <i>Journal of the American Chemical Society</i> . 94: 2549-50. PMID <a href="https://pubmed.ncbi.nlm.nih.gov/5016935/">5016935</a>
1972	<b>Corey EJ</b> , <a href="#">Becker KR</a> , Varma RK. <b>Efficient generation of the 15S configuration</b> in prostaglandin synthesis. Attractive interactions in stereochemical control of carbonyl reduction. <i>Journal of the American Chemical Society</i> . 94: 8616-8. PMID <a href="https://pubmed.ncbi.nlm.nih.gov/4638997/">4638997</a>
1969	<b>Corey EJ</b> , <a href="#">Wipke WT</a> . Computer-assisted design of complex organic syntheses. <i>Science (New York, N.Y.)</i> . 166: 178-92. PMID <a href="https://pubmed.ncbi.nlm.nih.gov/17731475/">17731475</a> DOI: <a href="https://doi.org/10.1126/science.166.3902.178">10.1126/science.166.3902.178</a>
1969	<b>Corey EJ</b> , Lin K, Yamamoto H. Separation of the cyclization and rearrangement processes of lanosterol biosynthesis. Enzymic conversion of 20,21-dehydro-2,3-oxidosqualene to a dehydroprotosterol. <i>Journal of the American Chemical Society</i> . 91: 2132-4. PMID <a href="https://pubmed.ncbi.nlm.nih.gov/5784177/">5784177</a>

1967	<b>Corey EJ, Ortiz de Montellano PR</b> , Lin K, Dean PD. 2,3-iminosqualene, a potent inhibitor of the enzymic cyclization of 2,3-oxidosqualene to sterols. <i>Journal of the American Chemical Society</i> . 89: 2797-8. PMID <a href="#">6043808</a>
1967	<b>Corey EJ, Ortiz de Montellano PR</b> . Enzymic synthesis of beta-amyrin from 2,3-oxidosqualene. <i>Journal of the American Chemical Society</i> . 89: 3362-3. PMID <a href="#">6042774</a>

<b>Year</b>	<b>Publication</b>
1959	Proof of the Structure and Stereochemistry of $\alpha$ -Amyrin by Synthesis from a $\beta$ -Amyrin Derivative, Glycyrrhetic Acid. <b>Corey, E. J.</b> ; Cantrall, E. W. Department of Chemistry, University of Illinois, Urbana, IL, USA. <i>Journal of the American Chemical Society</i> (1959), 81(7), pp. 1745-1751.
1957	Total Synthesis of Pentacyclosqualene. <b>Corey, E. J.</b> ; Sauers, R. R. Department of Chemistry, University of Illinois, Urbana, IL, USA. <i>Journal of the American Chemical Society</i> (1957), 79(14), pp. 3925-3926.
1954	The Stereochemistry of $\alpha$ -Haloketones. V. Prediction of the Stereochemistry of $\alpha$ -Brominated Ketosteroids. <b>Corey, Elias J.</b> Department of Chemistry, University of Illinois, Urbana, IL, USA. <i>Journal of the American Chemical Society</i> (1954), 76(1), pp. 175-179.
1953	Prediction of the Stereochemistry of Alpha-brominated Ketosteroids. <b>Corey, E. J.</b> Department of Chemistry, University of Illinois, Urbana, IL, USA. <i>Experientia</i> (became Cellular and Molecular Life Sciences in 1997) (1953), 9(9), pp. 329-331.
1952	Syntheses and Reactions of Acyclic N,N-diacylglycines. Sheehan, John C.; <b>Corey, Elias J.</b> Department of Chemistry, Massachusetts Institute of Technology, Cambridge, MA, USA. <i>Journal of the American Chemical Society</i> (1952), 74(18), pp. 4555-4559.
1951	The Synthesis of Substituted Penicillins and Simpler Structural Analogs. VI. The Synthesis of a 6-Phenylacetyl-amino- $\beta$ -lactam-thiazolidine. Sheehan, John C.; <b>Corey, Elias J.</b> Department of Chemistry, Massachusetts Institute of Technology, Cambridge, MA, USA. <i>Journal of the American Chemical Society</i> (1951), 73(10), pp. 4756-4759.
1950	The Total Synthesis of a 5-Phenylpenicillin: Methyl 5-Phenyl-(2-carbomethoxyethyl)-penicillinate. Sheehan, John C.; Buhle, Emmett L.; <b>Corey, Elias J.</b> ; Laubach, Gerald D.; Ryan, James J. Department of Chemistry, Massachusetts Institute of Technology, Cambridge, MA, USA. <i>Journal of the American Chemical Society</i> (1950), 72(8), pp. 3828-3829

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