# ORIGINAL

1 2 3	MARC J. FAGEL MICHAEL S. DICKE TRACY L. DAVIS JENNIFER L. SCAFE STEVEN D. BUCHHOLZ (Conditionally Admitted F	Pursuant to G.R. (2)(c)(2))
4	buchholzs@sec.gov	
5	Attorneys for Plaintiff SECURITIES AND EXCHANGE COMMISSION 44 Montgomery Street, Suite 2600	FILED — ENTERED — RECEIVED
6	San Francisco, California 94104 Telephone: 415-705-2500	SEP 0 8 2009
7 8	UNITED STATES DISTR	CLERK U.S. DISTRICT COURT  WESTERN DISTRICT OF WASHINGTON DEPUTY
9	WESTERN DISTRICT OF W	
10	AT SEATTLE	
11		C09-1263 BAT
12	SECURITIES AND EXCHANGE COMMISSION,	Civil Action No.
13	Plaintiff,	
14	vs.	COMPLAINT
15	CELLCYTE GENETICS CORPORATION and	
16	RONALD W. BERNINGER,	
17	Defendants.	
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19.	Plaintiff Securities and Exchange Commission ("Com	mission" or "SEC") alleges:
20	SUMMARY OF AC	TION
21	During 2007, CellCyte Genetics Corpora	ation ("CellCyte"), a fledgling
22	biotechnology company based in Bothell, Washington, r	repeatedly misled the investing public
23	about its key product, a purported stem cell therapy to tre	eat and repair damaged organs. CellCyte
24	claimed it had received approval from the U.S. Food and	l Drug Administration ("FDA") and was
25	on the verge of beginning human clinical trials with a sp	ecial stem cell compound to repair the
26	heart. Contrary to these claims, CellCyte did not even ke	now how to produce the stem cell
27	compound and had not satisfied any of the FDA requirer	ments to begin human clinical trials.
28	SEC v. CELLCYTE GENETICS CORP.	SECURITIES AND EXCHANGE COMMISSION

COMPLAINT

44 MONTGOMERY STREET, SUITE 2600 SAN FRANCISCO, CA 94104 TELEPHONE: 415-705-2500

- 2. In late 2005, CellCyte paid \$90,000 to license very early-stage stem cell technology. No research had been done using the technology since 2002. CellCyte failed in its efforts to raise money to begin conducting research, so it completed a reverse merger with a shell company controlled by a Canadian stock promoter to become a public company in March 2007. CellCyte agreed that the stock promoter would receive 15 million supposedly "freely tradeable" CellCyte shares as part of an illegal unregistered stock distribution, in exchange for about \$6 million in funding.
- 3. CellCyte then made false and misleading statements in several SEC filings and other materials distributed to potential investors about the Company's purportedly late-stage stem cell research and imminent clinical trials. Ronald Berninger, CellCyte's Chief Scientific Officer and a member of the Board of Directors, originally approved or participated in the drafting of many of the false and misleading statements and knew or was reckless in not knowing that the statements were false and misleading. During fall 2007, the stock promoter conducted a promotional campaign on behalf of CellCyte that included millions of spam emails, blast faxes, and newsletters containing false and misleading statements, some of which originated from CellCyte's own investor materials.
- 4. The misleading stock promotion campaign caused CellCyte's stock price to soar to a high of \$7.50 per share, with a 50-fold increase in trading volume. At its peak, CellCyte had a market capitalization of nearly \$450 million. The stock now trades at less than \$0.08 per share, leaving investors who were deceived by the fraudulent materials with massive losses.
- 5. The Commission seeks an order enjoining Defendants from future violations of the federal securities laws, requiring Berninger to pay a civil monetary penalty, and barring Berninger from serving as an officer or director of a public company.

#### JURISDICTION AND VENUE

6. This Court has jurisdiction over this action pursuant to Sections 20(b), 20(d) and 22(a) of the Securities Act of 1933 ("Securities Act") [15 U.S.C. §§ 77t(b), 77t(d) and 77v(a)] and Sections 21(d), 21(e) and 27 of the Securities Exchange Act of 1934 ("Exchange

-2-

-3-

specially formulated compound, when injected into the bloodstream before a dose of stem

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cells, appeared to cause the stem cells to migrate to specific organs and remain there in significant concentrations. Normally, stem cells that are injected into the bloodstream do not remain in any particular organ and are quickly flushed out of the body. The special compound was supplied by a European biotechnology company, which also funded the scientist's research. The biotechnology company and the scientist filed patent applications related to the discovery in 2002.

- 13. In 2002, before the scientist could perform additional research using the compound, the biotechnology company stopped funding the research and supplying the special compound. The scientist performed no further research using the compound after 2002
- 14. Berninger and CellCyte's CEO formed CellCyte as a private company in early 2005 for the purpose of acquiring the rights to the scientist's discovery.
- 15. In January 2005, Berninger and CellCyte's CEO met with the scientist for several days to review the details of her research findings. At the meetings and in discussions thereafter, the scientist told Berninger and CellCyte's CEO that she had only conducted preliminary research.
- 16. The scientist also told Berninger and CellCyte's CEO that she had never conducted research on the compound using mice with injured organs, and therefore had no data showing that the stem cells could repair or improve function in injured organs. The scientist also disclosed that she had not yet done any toxicology studies, and that the mixture of the compound and stem cells had killed some mice during her research. Further, the scientist told Berninger and CellCyte's CEO that CellCyte would need to obtain or develop a supply of the special compound before any further research could begin.
- 17. In October 2005, CellCyte paid \$90,000 to license the scientist's discovery. The license agreement required CellCyte to raise an additional \$5.5 million within one year to demonstrate that it was capable of conducting active research and development of the technology. At least by the time CellCyte entered into the license agreement, Berninger and

CellCyte's CEO were aware that the scientist had only achieved positive results in a small number of mice, and that CellCyte would need to conduct extensive further research and testing on much larger numbers of mice before preparing an investigational new drug ("IND") application for FDA approval to begin any human clinical trials.

## B. CellCyte Merges With a Public Shell Company and Attempts To Conduct Research.

- 18. In late 2006, while unsuccessfully trying to raise \$5.5 million to retain the license to the scientist's discovery and begin research, CellCyte's CEO met a Canadian stock promoter who controlled a public shell company. CellCyte and the stock promoter conducted a reverse merger between CellCyte and the shell company, thereby making CellCyte a public company.
- 19. In connection with the reverse merger, CellCyte received about \$6 million from the stock promoter and other investors, and the stock promoter received about 15 million purportedly "freely tradeable" CellCyte shares. As a result, the stock promoter controlled about 90% of CellCyte's public float (the shares outstanding and available for trading by the public). No registration statement was in effect covering the stock promoter's receipt of the 15 million shares and no exemption from the registration requirements applied.
- 20. CellCyte's stock was first quoted on the Over The Counter (OTC) Bulletin Board on February 16, 2007, and the reverse merger officially closed on March 30, 2007.
- 21. In March 2007, CellCyte began searching for lab space and hiring personnel, including a Vice President of Research & Development who reported to Berninger.
- 22. In May 2007, CellCyte began attempting to formulate the special compound itself for the first time. Within a few months, the Vice President of Research & Development raised concerns to Berninger and CellCyte's CEO that the compound was not properly formulated and could not be used to successfully develop the stem cell technology.
- 23. In October 2007, CellCyte began conducting experiments using the compound in mice. Those experiments failed to produce any of the results necessary to support an IND

-5-

1	application. CellCyte conducted additional experiments using mice in November 2007 and
2	March 2008, which also failed.
3	C. CellCyte Makes False and Misleading Statements About Its Research.
4	24. After CellCyte became a public company in March 2007, it made false and
5	misleading statements in SEC filings about its research and development efforts and business
6	prospects. Berninger originally approved or participated in the drafting of many of these
7	statements for use in materials that were distributed to potential investors before and after
8	CellCyte became a public company.
9	25. CellCyte made false and misleading statements in four different SEC filings,
10	including a Form 8-K current report filed April 5, 2007, a Form 10-Q quarterly report filed
11	May 18, 2007, a Form SB-2 registration statement filed June 29, 2007, and a Form 10-Q
12	quarterly report filed August 14, 2007. CellCyte's Board, including Berninger, approved the
13	Form SB-2 filing, and CellCyte's CEO emailed Berninger a draft of the SB-2 filing that
14	contained the false and misleading statements and asked Berninger to review it. Berninger
15	also was present at Board meetings during which the May and August 2007 Form 10-Q
16	reports were approved or discussed.
17 18	<ul> <li>i. CellCyte Falsely Claimed That Its Stem Cell Drugs Were Already in FDA- Approved Clinical Trials.</li> </ul>
19	26. In the SEC filings listed in paragraph 25 above and other materials distributed
20	to potential investors, CellCyte stated that its stem cell discoveries "are the first stem cell
21	enabling drugs to enter Investigational New Drug ('IND') supported by the United States
22	Food and Drug Administration ('FDA') clinical trials."
23	27. In reality, Berninger knew that CellCyte never filed an IND application for the
24	stem cell technology and therefore never received FDA approval to begin clinical trials.
25	Berninger knew or was reckless in not knowing that the statement was materially false and
26	misleading.
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-6-

1	ii. CellCyte Misled Investors About the Advanced Stage of Its Research and
2	<u>Development.</u>
3	28. In the SEC filings listed in paragraph 25 above and other materials distributed
4	to potential investors, CellCyte portrayed its stem cell research as having been "proven in
5	extensive late-stage animal studies."
6	29. In reality, Berninger knew that the research had not been proven in extensive
7	late-stage animal studies. Berninger knew or was reckless in not knowing that the statement
8	was materially false and misleading.
9	iii. CellCyte Falsely Stated That It Was Within Months of Starting Clinical Trials To Repair the Heart.
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11	30. In the SEC filings listed in paragraph 25 above and other materials distributed
12	to potential investors, CellCyte claimed that "the company is advancing [the heart compound]
13	into human trials for repair of the heart with an IND submission scheduled for the second half
14	of 2007."
15	31. In reality, Berninger knew that CellCyte never attempted any research nor
16	achieved any results to prove that its stem cell technology could repair the heart in mice, a
17	prerequisite to beginning any clinical trial to repair the heart in humans. Indeed, in June 2006
18	Berninger told CellCyte's CEO that the Company's target was to complete research and
19	testing for an IND submission involving the heart in two years, i.e. by June 2008. Berninger
20	knew or was reckless in not knowing that the statement was materially false and misleading.
21	iv. CellCyte Falsely Stated That It Was Working Closely With Swedish
22	Medical Center.
23	32. In the SEC filings listed in paragraph 25 above and other materials distributed
24	to potential investors, CellCyte claimed to be "working closely with Swedish Medical Center
25	in Seattle and their internationally recognized organ transplant group" on the stem cell
26	technology.
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-7-

- 33. In reality, CellCyte's CEO had merely spoken to an acquaintance at Swedish Medical Center, a leading health care provider in the Pacific Northwest, about the possibility of conducting joint research in the future. Berninger knew that CellCyte was not actually working with Swedish Medical Center. Berninger knew or was reckless in not knowing that the statement was materially false and misleading.
  - v. <u>CellCyte Falsely Stated That Its Drugs Had Been Shown To Improve Bone Marrow Engraftment.</u>
- 34. In the SEC filings listed in paragraph 25 above and other materials distributed to potential investors, CellCyte claimed that during its research using the special compound, "the stem cells migrated directly to the bone marrow, therefore increasing the effective dose of stem cells available for engraftment."
- 35. In reality, Berninger knew that the preliminary research on the compound had not shown stem cells migrating to the bone marrow. CellCyte never conducted any research involving bone marrow, and it had no data showing that the compound caused stem cells to localize in bone marrow or improved bone marrow engraftment. Berninger knew or was reckless in not knowing that the statement was materially false and misleading.

### D. CellCyte Makes Material Omissions About Its Research and Operations.

- 36. CellCyte also omitted critical information from its public statements about its research. Most significantly, CellCyte failed to disclose that it was unable to obtain the specially formulated compound from the European biotechnology company that had originally funded the stem cell research, and that CellCyte had not determined how to properly formulate the special compound from material that CellCyte obtained from other sources. CellCyte needed to have a sufficient supply of the special compound before it could begin conducting the extensive additional research that was required to determine whether an IND application could be filed to begin human clinical trials.
- 37. CellCyte touted a lack of safety and toxicity concerns about the special compound, but failed to state that no safety or toxicology studies had been done or that some

mice had died during the scientist's preliminary research, suggesting that some doses of the compound and stem cells were in fact toxic and even fatal.

- 38. CellCyte also failed to disclose that its experiments in mice between October 2007 and March 2008 were unsuccessful.
- 39. The information omitted by CellCyte, and the false and misleading statements described in paragraphs 26 to 34 above, were material to investors because they concerned the Company's ultimate likelihood of success in developing stem cell technology to repair damaged organs.

### E. The Stock Promoter Conducts a Widespread Promotional Campaign.

- 40. In April 2007, Berninger and CellCyte's CEO met with the Canadian stock promoter to discuss investor relations, a marketing budget, and the stock promoter's efforts to raise additional capital for CellCyte. CellCyte gave the stock promoter investor materials that contained the false and misleading statements described in paragraphs 26 to 34 above, to use in preparing promotional materials.
- 41. In August 2007, CellCyte's CEO told Berninger that the stock promoter was ready to launch a promotional campaign for CellCyte and would spend \$2 million on the campaign in exchange for additional stock in a future offering. Between August and December 2007, the stock promoter distributed millions of spam emails, blast faxes, and newsletters that contained false and misleading statements about CellCyte, some of which originated from CellCyte's own investor materials.
- 42. During the promotional campaign, CellCyte's stock price rose from around \$4.00 to \$7.50, and its daily trading volume increased from 2,000 to more than 100,000 shares. At one point during the campaign, CellCyte's market capitalization reached nearly \$450 million.
- 43. By the end of January 2008, after the stock promotion campaign had concluded (and after much of the Canadian promoter's stock had been dumped into the market), CellCyte's stock price declined below a dollar. The stock currently trades at around \$0.07 per share.

-9-

TELEPHONE: 415-705-2500

1	FIRST CLAIM FOR RELIEF	
2	Violations of Securities Act Sections 5(a) and 5(c) by CellCyte	
3	44. The Commission realleges and incorporates by reference paragraphs 1 through	
4	43.	
5	45. By engaging in the conduct described above, CellCyte, directly or indirectly,	
6	made use of means or instruments of transportation or communication in interstate commerce	
7	or of the mails to offer or to sell securities through the use or medium of a prospectus or	
8	otherwise when no registration statement had been filed or was in effect as to such securities	
9	and no exemption from registration was available.	
10	46. By reason of the foregoing, CellCyte has violated and, unless restrained and	
11	enjoined, will continue to violate Sections 5(a) and 5(c) of the Securities Act [15 U.S.C.	
12	§§ 77e(a) and 77e(c)].	
13	SECOND CLAIM FOR RELIEF	
14	Violations of Exchange Act Section $10(b)$ and Rule $10b$ -5 Thereunder by All Defendants	
15	47. The Commission realleges and incorporates by reference paragraphs 1 through	
16	46.	
17	48. By engaging in the conduct described above, CellCyte and Berninger, directly	
18	or indirectly, in connection with the purchase or sale of securities, by the use of means or	
19	instrumentalities of interstate commerce or of the mails, with scienter:	
20	(a) employed devices, schemes, or artifices to defraud;	
21	(b) made untrue statements of material facts or omitted to state material	
22	facts necessary in order to make the statements made, in the light of the	
23	circumstances under which they were made, not misleading; and	
24	(c) engaged in acts, practices, or courses of business which operated or	
25	would operate as a fraud or deceit upon other persons, including purchasers	
26	and sellers of securities.	
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-10-

TELEPHONE: 415-705-2500

1	49. By reason of the foregoing, CellCyte and Berninger have violated and, unless
2	restrained and enjoined, will continue to violate Section 10(b) of the Exchange Act [15 U.S.C.
3	§ 78j(b)] and Rule 10b-5 [17 C.F.R. § 240.10b-5].
4	THIRD CLAIM FOR RELIEF
5	Violations of and Aiding and Abetting Violations of Exchange Act Section 13(a)
6	and Rules 12b-20, 13a-11 and 13a-13 Thereunder by Defendants
7	50. The Commission realleges and incorporates by reference paragraphs 1 through
8	49.
9	51. By engaging in the conduct described above, CellCyte violated Section 13(a)
10	of the Exchange Act [15 U.S.C. § 78m(a)] and Rules 12b-20, 13a-11 and 13a-13 [17 C.F.R.
11	§§ 240.12b-20, 240.13a-11 and 240.13a-13], which obligate issuers of securities registered
12	pursuant to Section 12 of the Exchange Act [15 U.S.C. § 78l] to file with the Commission
13	accurate quarterly and current reports.
14	52. By reason of the foregoing, CellCyte has violated and, unless restrained and
15	enjoined, will continue to violate Section 13(a) of the Exchange Act [15 U.S.C. § 78m(a)] and
16	Rules 12b-20, 13a-11 and 13a-13 [17 C.F.R. §§ 240.12b-20, 240.13a-11 and 240.13a-13].
17	53. By engaging in the conduct described above, Berninger knowingly provided
18	substantial assistance to CellCyte's filing of materially false and misleading reports with the
19	Commission.
20	54. By reason of the foregoing, Berninger aided and abetted CellCyte's violations
21	of Section 13(a) of the Exchange Act [15 U.S.C. § 78m(a)] and Rules 12b-20, 13a-11 and
22	13a-13 [17 C.F.R. §§ 240.12b-20, 240.13a-11 and 240.13a-13]. Unless restrained and
23	enjoined, Berninger will continue to aid and abet such violations.
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1	PRAYER FOR RELIEF
2	WHEREFORE, the Commission respectfully requests that this Court:
3	I.
4	Issue an order permanently restraining and enjoining CellCyte and its agents, servants
5	employees, attorneys, and assigns, and those persons in active concert or participation with
6	them, from violating, directly or indirectly, Sections 5(a) and 5(c) of the Securities Act [15
7	U.S.C. §§ 77q(a) and 77q(c)]; Sections 10(b) and 13(a) of the Exchange Act [15 U.S.C. §§
8	78j(b) and 78m(a)]; and Rules 10b-5, 12b-20, 13a-11 and 13a-13 thereunder [17 C.F.R. §§
9	240.10b-5, 240.12b-20, 240.13a-11 and 240.13a-13]; and permanently restraining and
10	enjoining Berninger and his agents, servants, employees, attorneys, and assigns, and those
11	persons in active concert or participation with them, from violating, directly or indirectly,
12	Section 10(b) of the Exchange Act [15 U.S.C. § 78j(b)] and Rule 10b-5 thereunder [17 C.F.R.
13	§ 240.10b-5] and from aiding and abetting violations of Section 13(a) of the Exchange Act
14	[15 U.S.C. § 78m(a)] and Rules 12b-20, 13a-11 and 13a-13 thereunder [17 C.F.R. §§
15	240.12b-20, 240.13a-11 and 240.13a-13].
16	II.
17	Issue an order directing Berninger to pay a civil monetary penalty under Section
18	21(d)(3) of the Exchange Act [15 U.S.C. § 78u(d)(3)].
19	III.
20	Issue an order barring Berninger from serving as an officer or director of any public
21	company, pursuant to Section 21(d)(2) of the Exchange Act [15 U.S.C. § 78u(d)(2)].
22	IV.
23	Retain jurisdiction of this action in accordance with the principles of equity and the
24	Federal Rules of Civil Procedure in order to implement and carry out the terms of all orders
25	and decrees that may be entered, or to entertain any suitable application or motion for
26	additional relief within the jurisdiction of this Court.
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SEC V. CELLCYTE GENETICS CORP. COMPLAINT

SECURITIES AND EXCHANGE COMMISSION 44 MONTGOMERY STREET, SUITE 2600 SAN FRANCISCO, CA 94104 TELEPHONE: 415-705-2500

-13-