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Domestic Policy Council

Zinsmeister, W. Karl - Subject Files

Stack:	Row:	Sect.:	Shelf:	Pos.:	FRC ID:	Location or Hollinger ID:	NARA Number:	OA Number:
W	31	24	6	7	14303	88293	14062	14165

Folder Title:

Stem Cells 2007: Stem Cell Veto 2007 and Executive Order [3]

Withdrawn/Redacted Material

The George W. Bush Library

DOCUMENT NO.	FORM	SUBJECT/TITLE	PAGES	DATE	RESTRICTION(S)
001	Email	[Page 2] - From: Chris	1	06/06/2007	P6/b6;
002	Draft	[Draft]	2	N.D.	P5;
003	Handwritten Note	[Notes]	1	N.D.	P5;
004	Handwritten Note	[Notes]	6	06/13/2007	P5;
005	Email	Coleman-Isakson: Stem Cells - To: Karl Zinsmeister - From: Christopher Papagianis	2	05/17/2007	P5;
006	Handwritten Note	[Notes]	2	N.D.	P5;
007	Email	Stem Cell Meetings - To: Karl Zinsmeister - From: Christopher Papagianis	1	05/03/2007	P5;

COLLECTION TITLE:

Domestic Policy Council

SERIES:

Zinsmeister, W. Karl - Subject Files

FOLDER TITLE:

Stem Cells 2007: Stem Cell Veto 2007 and Executive Order [3]

FRC ID:

14303

RESTRICTION CODES

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PRM. Personal record misfile defined in accordance with 44 U.S.C. 2201(3).

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008	Email	Stem Cell Meetings--Update - To: Karl Zinsmeister - From: Christopher Papagianis	2	05/02/2007	P5;
009	Handwritten Note	[Notes]	1	N.D.	P5;
010	Email	RE: Update Since We Spoke at 9am re Stem Cells - To: Christopher Papagianis - From: Karl Zinsmeister	1	05/09/2007	P5;
011	Email	More on Stem Cell Meetings - To: Karl Zinsmeister - From: Christopher Papagianis	1	05/08/2007	P5;
012	Email	Update on Stem Cell Meeting/Plan - To: Karl Zinsmeister - From: Christopher Papagianis	1	05/07/2007	P5;
013	Handwritten Note	[Notes]	1	N.D.	P5;

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014	Email	S.30 - To: Candida Wolff - From: Karl Zinsmeister	1	04/26/2007	P5;
015	Email	Update and Proposal on Stem Cells - To: Karl Zinsmeister - From: Christopher Papagianis	2	04/26/2007	P5;
016	Email	Notes for Weldon Call - To: Karl Zinsmeister - From: Christopher Papagianis	1	05/01/2007	P5;
017	Handwritten Note	[Notes]	1	N.D.	P5;
018	Handwritten Note	[Notes]	1	N.D.	P5;
019	Briefing	[Briefing]	5	N.D.	P5;
020	Handwritten Note	[Notes]	1	N.D.	P5;

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DOCUMENT NO.	FORM	SUBJECT/TITLE	PAGES	DATE	RESTRICTION(S)
021	Email	RE: Update and Proposal on Stem Cells - To: Karl Zinsmeister - From: Christopher Papagianis	1	04/27/2007	P5;
022	Briefing	Policy Time - Domestic	2	06/18/2007	P5;
023	Draft	[Draft]	3	N.D.	P5;

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Zinsmeister, Karl

From: Lerner, Daniel A.
Sent: Wednesday, June 06, 2007 8:53 AM
To: Lerner, Daniel A.
Subject: OFFICIAL RELEASE: SAP on H.R. 2560 – Human Cloning Prohibition Act of 2007
Attachments: SAP on HR2560.pdf

**EXECUTIVE OFFICE OF THE PRESIDENT
OFFICE OF MANAGEMENT AND BUDGET
WASHINGTON, D.C. 20503**

6, 2007

STATEMENT OF ADMINISTRATION POLICY

H.R. 2560 – Human Cloning Prohibition Act of 2007

(Rep. DeGette (D) Colorado)

The President unequivocally opposes all forms of human cloning. The Administration is strongly opposed to any legislation that would prohibit human cloning for reproductive purposes but permit the creation of cloned embryos or development of human embryo farms for research, which would require the destruction of nascent human life. Thus, if legislation were presented to the President that permitted human embryos to be created, developed, and destroyed simply for research purposes, his senior advisors would recommend that he veto the bill.

* * * * *

Zinsmeister, Karl

From: Lerner, Daniel A.
Sent: Wednesday, June 06, 2007 10:09 AM
To: Lerner, Daniel A.
Subject: OFFICIAL RELEASE: S. 5 – Stem Cell Research Enhancement Act of 2007 (House Rules)
Attachments: SAP on S5 (House Rules).pdf

**EXECUTIVE OFFICE OF THE PRESIDENT
OFFICE OF MANAGEMENT AND BUDGET
WASHINGTON, D.C. 20503**

6, 2007

Rules)

STATEMENT OF ADMINISTRATION POLICY

S. 5 – Stem Cell Research Enhancement Act of 2007

(Sen. Reid (D) NV and 41 cosponsors)

The Administration strongly opposes House passage of S. 5, which would use Federal taxpayer dollars to support and encourage the destruction of human life for research. The bill would compel all American taxpayers to pay for research that relies on the intentional destruction of human embryos for the derivation of stem cells, overturning the President's policy that funds research without promoting such ongoing destruction. If S. 5 were presented to the President, he would veto the bill.

The President strongly supports medical research and has worked with Congress to increase resources for the National Institutes of Health. This Administration is the first to provide Federal funds for human embryonic stem cell research and has done so without encouraging the destruction of human embryos. The President's policy permits the Federal funding of research using embryonic cell lines created prior to August 9, 2001, the date his policy was announced, along with stem cell research using other kinds of cell lines. Scientists can therefore explore the potential applications of such cells, but the Federal government does not offer incentives or encouragement for the destruction of human life.

Over the past six years, more than \$130 million in taxpayer dollars has been devoted to human embryonic stem cell research consistent with the President's policy. Overall, more than \$3 billion has gone to innovative research on all forms of stem cells, contributing to proven medical treatments that use human stem cells from adult and other non-embryonic sources. S. 5 seeks to replace the Administration's policy by providing Federal funding for the first time for a line of research that involves the intentional destruction of living human embryos for the derivation of their cells. Destroying nascent human life for research raises serious ethical problems, and millions of Americans consider the practice immoral. The Administration believes that government has a duty to use the people's money responsibly, both supporting important public purposes and respecting moral boundaries.

Alternative types of human stem cells – drawn from adults, children, umbilical-cord blood, and other non-embryonic sources, without doing harm to the donors – have already achieved therapeutic results in thousands of patients with many different diseases. Moreover, a recent series of encouraging research reports offer hope that stem cells drawn from non-embryonic sources may possess characteristics like those of embryonic stem cells. In addition, researchers are now working to develop promising new techniques to produce stem cells similar in nature to those derived from human embryos, but without

6/6/2007

harming or destroying embryos.

S. 5 clearly acknowledges the potential of these new advances and the importance of supporting them. The provisions in the bill to promote research leading to new techniques for deriving the functional equivalent of embryonic stem cells, without harming or destroying human embryos, encourage science that is not in conflict with ethics.

Unfortunately, the bill then crosses that same ethical line, by proposing to use taxpayer dollars to encourage destruction of nascent human life for research. The Administration believes that research on alternative sources of stem cells is extremely promising and provides robust opportunities to advance science without compelling American taxpayers to participate in ongoing destruction of human embryos.

Moreover, private sector support and public funding by several States for this line of research, which will add up to several billion dollars in the coming few years, argue against the notion of any urgent shortfall of research funding. Whatever one's view of the ethical issues or the state of the research, the future of this field does not require a policy of Federal subsidies that is offensive to the moral principles of millions of Americans.

The President believes that by enacting appropriate policy safeguards while encouraging the development of novel scientific techniques, it is possible to advance scientific and medical frontiers without violating moral principles. S. 5 fails to find that balance and is therefore deeply troubling to millions of Americans. The President urges Congress to pass an alternative bill that would advance stem cell research without encouraging destruction of human life.

* * * * *

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OA Num.:

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NARA Num.:

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FOIA IDs and Segments:

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2014-0503-F

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110TH CONGRESS
1ST SESSION

S. 30

AN ACT

To intensify research to derive human pluripotent stem cell lines.

1 *Be it enacted by the Senate and House of Representa-*
2 *tives of the United States of America in Congress assembled,*

3 **SECTION 1. SHORT TITLE.**

4 This Act may be cited as the “Hope Offered through
5 Principled and Ethical Stem Cell Research Act” or the
6 “HOPE Act”.

1 **SEC. 2. PURPOSES.**

2 It is the purpose of this Act to—

3 (1) intensify research that may result in im-
4 proved understanding of or treatments for diseases
5 and other adverse health conditions; and

6 (2) promote the derivation of pluripotent stem
7 cell lines without the creation of human embryos for
8 research purposes and without the destruction or
9 discarding of, or risk of injury to, a human embryo
10 or embryos other than those that are naturally dead.

11 **SEC. 3. HUMAN PLURIPOTENT STEM CELL RESEARCH.**

12 Part H of title IV of the Public Health Service Act
13 (42 U.S.C. 289 et seq.) is amended by inserting after sec-
14 tion 498C the following:

15 **“SEC. 498D. HUMAN PLURIPOTENT STEM CELL RESEARCH.**

16 “(a) IN GENERAL.—The Secretary shall conduct and
17 support basic and applied research to develop techniques
18 for the isolation, derivation, production, or testing of stem
19 cells, including pluripotent stem cells that have the flexi-
20 bility of embryonic stem cells (whether or not they have
21 an embryonic source), that may result in improved under-
22 standing of or treatments for diseases and other adverse
23 health conditions, provided that the isolation, derivation,
24 production, or testing of such cells will not involve—

25 “(1) the creation of a human embryo or em-
26 bryos for research purposes; or

1 “(2) the destruction or discarding of, or risk of
2 injury to, a human embryo or embryos other than
3 those that are naturally dead.

4 “(b) GUIDELINES.—Not later than 90 days after the
5 date of the enactment of this section, the Secretary, after
6 consultation with the Director of NIH, shall issue final
7 guidelines that—

8 “(1) provide guidance concerning the next steps
9 required for additional research, which shall include
10 a determination of the extent to which specific tech-
11 niques may require additional animal research to en-
12 sure that any research involving human cells using
13 these techniques would clearly be consistent with the
14 standards established under subsection (a);

15 “(2) prioritize research with the greatest poten-
16 tial for near-term clinical benefit;

17 “(3) consistent with standards established
18 under subsection (a), take into account techniques
19 outlined by the President’s Council on Bioethics and
20 any other appropriate techniques and research; and

21 “(4) in the case of research involving stem cells
22 from a naturally dead embryo, require assurances
23 from grant applicants that no alteration of the tim-
24 ing, methods, or procedures used to create, main-
25 tain, or intervene in the development of a human

1 embryo was made solely for the purpose of deriving
2 the stem cells.

3 “(c) REPORTING REQUIREMENTS.—Not later than
4 January 1 of each year, the Secretary shall prepare and
5 submit to the appropriate committees of the Congress a
6 report describing the activities carried out under this sec-
7 tion during the fiscal year, including a description of the
8 research conducted under this section.

9 “(d) RULE OF CONSTRUCTION.—Nothing in this sec-
10 tion shall be construed as altering the policy in effect on
11 the date of enactment of this section regarding the eligi-
12 bility of stem cell lines for funding by the National Insti-
13 tutes of Health.

14 “(e) AUTHORIZATION OF APPROPRIATIONS.—There
15 is authorized to be appropriated such sums as may be nec-
16 essary to carry out this section.

17 “(f) DEFINITIONS.—In this section:

18 “(1) NATURALLY DEAD.—The term ‘naturally
19 dead’ means having naturally and irreversibly lost
20 the capacity for integrated cellular division, growth,
21 and differentiation that is characteristic of an orga-
22 nism, even if some cells of the former organism may
23 be alive in a disorganized state.

24 “(2) HUMAN EMBRYO OR EMBRYOS.—The term
25 ‘human embryo or embryos’ includes any organism,

1 not protected as a human subject under part 46 of
2 title 45, Code of Federal Regulations, as of the date
3 of enactment of this section, that is derived by fer-
4 tilization, parthenogenesis, cloning, or any other
5 means from one or more human gametes or human
6 diploid cells.

7 “(3) RISK OF INJURY.—The term ‘risk of in-
8 jury’ means subjecting a human embryo or embryos
9 to risk of injury or death greater than that allowed
10 for research on fetuses in utero under section
11 46.204(b) of title 45, Code of Federal Regulations,
12 and section 498(b) of this Act.”.

13 **SEC. 4. NATIONAL AMNIOTIC AND PLACENTAL STEM CELL**
14 **BANK.**

15 (a) IN GENERAL.—The Secretary of Health and
16 Human Services shall enter into a contract with the Insti-
17 tute of Medicine for the conduct of a study to recommend
18 an optimal structure for an amniotic and placental stem
19 cell bank program and to address pertinent issues to maxi-
20 mize the potential of such technology, including collection,
21 storage, standards setting, information sharing, distribu-
22 tion, reimbursement, research, and outcome measures. In
23 conducting such study, the Institute should receive input
24 from relevant experts including the existing operators of

1 federal tissue bank programs and the biomedical research
2 programs within the Department of Defense.

3 (b) REPORT.—Not later than 180 days after the date
4 of enactment of this Act, the Institute of Medicine shall
5 complete the study under subsection (a) and submit to the
6 Secretary of Health and Human Services and the appro-
7 priate committees of Congress a report on the results of
8 such study.

Passed the Senate April 11, 2007.

Attest:

Secretary.



EXECUTIVE OFFICE OF THE PRESIDENT
OFFICE OF MANAGEMENT AND BUDGET
WASHINGTON, D.C. 20503

April 10, 2007
(Senate)

STATEMENT OF ADMINISTRATION POLICY

S. 30 – Hope Offered through Principled and Ethical Stem Cell Research Act

(Sens. Coleman (R) MN and Isakson (R) GA)

The Administration strongly supports Senate passage of S. 30, a bill to support and intensify research into techniques of deriving pluripotent stem cells without harming or destroying human embryos.

Pluripotent stem cells are cells that can be differentiated into nearly all the cell types in the human body. At present, the means of obtaining such cells involve the destruction or endangerment of living human embryos, raising serious moral concerns. Ongoing research is investigating whether stem cells derived from the tissues of donors, umbilical cord blood, amniotic fluid, placentas, and other non-embryonic sources may be pluripotent. Numerous techniques are being explored by researchers, and significant scientific advances have been reported in the past few years.

This bill authorizes the Department of Health and Human Services to support the use and further development of techniques for producing pluripotent cells like those derived from human embryos, but without harming or destroying the embryos. Research using pluripotent stem cells is still at a very early stage, and it will be years before researchers know how much promise lies in therapeutic applications. This early stage is precisely when it is most important to develop ethically responsible techniques, so that the potential of pluripotent stem cells can be explored without violating human dignity and life.

This bill also directs the Department, under contract with the Institute of Medicine, to study and recommend the optimal structure for an amniotic and placental stem cell bank. While research in this area is still developing, recent publications suggest that a new and readily available source of pluripotent stem cells may exist in the amniotic fluid that cushions babies in the womb. The Administration supports further investigation in this area.

The Administration does not believe science and ethics need be at odds. Scientists have shown they have the ingenuity and skill to pursue the potential benefits of stem cell research without endangering nascent human life in the process. By seeking intensified support for non-destructive alternatives, we believe we can advance medical research in valuable ways while respecting ethical boundaries that are vital to millions of American taxpayers.

* * * * *

110TH CONGRESS
1ST SESSION

S. 5

AN ACT

To amend the Public Health Service Act to provide for
human embryonic stem cell research.

1 *Be it enacted by the Senate and House of Representa-*
2 *tives of the United States of America in Congress assembled,*

3 **SECTION 1. SHORT TITLE.**

4 This Act may be cited as the “Stem Cell Research
5 Enhancement Act of 2007”.

1 **SEC. 2. HUMAN EMBRYONIC STEM CELL RESEARCH.**

2 Part H of title IV of the Public Health Service Act
3 (42 U.S.C. 289 et seq.) is amended by inserting after sec-
4 tion 498C the following:

5 **“SEC. 498D. HUMAN EMBRYONIC STEM CELL RESEARCH.**

6 “(a) IN GENERAL.—Notwithstanding any other pro-
7 vision of law (including any regulation or guidance), the
8 Secretary shall conduct and support research that utilizes
9 human embryonic stem cells in accordance with this sec-
10 tion (regardless of the date on which the stem cells were
11 derived from a human embryo) .

12 “(b) ETHICAL REQUIREMENTS.—Human embryonic
13 stem cells shall be eligible for use in any research con-
14 ducted or supported by the Secretary if the cells meet each
15 of the following:

16 “(1) The stem cells were derived from human
17 embryos that have been donated from in vitro fer-
18 tilization clinics, were created for the purposes of
19 fertility treatment, and were in excess of the clinical
20 need of the individuals seeking such treatment.

21 “(2) Prior to the consideration of embryo dona-
22 tion and through consultation with the individuals
23 seeking fertility treatment, it was determined that
24 the embryos would never be implanted in a woman
25 and would otherwise be discarded.

1 “(3) The individuals seeking fertility treatment
2 donated the embryos with written informed consent
3 and without receiving any financial or other induce-
4 ments to make the donation.

5 “(c) GUIDELINES.—Not later than 60 days after the
6 date of the enactment of this section, the Secretary, in
7 consultation with the Director of NIH, shall issue final
8 guidelines to carry out this section.

9 “(d) REPORTING REQUIREMENTS.—The Secretary
10 shall annually prepare and submit to the appropriate com-
11 mittees of the Congress a report describing the activities
12 carried out under this section during the preceding fiscal
13 year, and including a description of whether and to what
14 extent research under subsection (a) has been conducted
15 in accordance with this section.”.

16 **SEC. 3. ALTERNATIVE HUMAN PLURIPOTENT STEM CELL**
17 **RESEARCH.**

18 Part H of title IV of the Public Health Service Act
19 (42 U.S.C. 284 et seq.), as amended by section 2, is fur-
20 ther amended by inserting after section 498D the fol-
21 lowing:

22 **“SEC. 498E. ALTERNATIVE HUMAN PLURIPOTENT STEM**
23 **CELL RESEARCH.**

24 “(a) IN GENERAL.—In accordance with section 492,
25 the Secretary shall conduct and support basic and applied

1 research to develop techniques for the isolation, derivation,
2 production, or testing of stem cells that, like embryonic
3 stem cells, are capable of producing all or almost all of
4 the cell types of the developing body and may result in
5 improved understanding of or treatments for diseases and
6 other adverse health conditions, but are not derived from
7 a human embryo.

8 “(b) GUIDELINES.—Not later than 90 days after the
9 date of the enactment of this section, the Secretary, after
10 consultation with the Director, shall issue final guidelines
11 to implement subsection (a), that—

12 “(1) provide guidance concerning the next steps
13 required for additional research, which shall include
14 a determination of the extent to which specific tech-
15 niques may require additional basic or animal re-
16 search to ensure that any research involving human
17 cells using these techniques would clearly be con-
18 sistent with the standards established under this sec-
19 tion;

20 “(2) prioritize research with the greatest poten-
21 tial for near-term clinical benefit; and

22 “(3) consistent with subsection (a), take into
23 account techniques outlined by the President’s Coun-
24 cil on Bioethics and any other appropriate tech-
25 niques and research.

1 “(c) REPORTING REQUIREMENTS.—Not later than
2 January 1 of each year, the Secretary shall prepare and
3 submit to the appropriate committees of the Congress a
4 report describing the activities carried out under this sec-
5 tion during the fiscal year, including a description of the
6 research conducted under this section.

7 “(d) RULE OF CONSTRUCTION.—Nothing in this sec-
8 tion shall be construed to affect any policy, guideline, or
9 regulation regarding embryonic stem cell research, human
10 cloning by somatic cell nuclear transfer, or any other re-
11 search not specifically authorized by this section.

12 “(e) DEFINITION.—

13 “(1) IN GENERAL.—In this section, the term
14 ‘human embryo’ shall have the meaning given such
15 term in the applicable appropriations Act.

16 “(2) APPLICABLE ACT.—For purposes of para-
17 graph (1), the term ‘applicable appropriations Act’
18 means, with respect to the fiscal year in which re-
19 search is to be conducted or supported under this
20 section, the Act making appropriations for the De-
21 partment of Health and Human Services for such
22 fiscal year, except that if the Act for such fiscal year
23 does not contain the term referred to in paragraph
24 (1), the Act for the previous fiscal year shall be
25 deemed to be the applicable appropriations Act.

1 “(f) AUTHORIZATION OF APPROPRIATIONS.—There
2 is authorized to be appropriated such sums as may be nec-
3 essary for each of fiscal years 2008 through 2010, to carry
4 out this section.”.

Passed the Senate April 11, 2007.

Attest:

Secretary.



EXECUTIVE OFFICE OF THE PRESIDENT
OFFICE OF MANAGEMENT AND BUDGET
WASHINGTON, D.C. 20503

April 10, 2007
(Senate)

STATEMENT OF ADMINISTRATION POLICY

S. 5 – Stem Cell Research Enhancement Act of 2007

(Sen. Reid (D) NV and 39 cosponsors)

The Administration strongly opposes Senate passage of S. 5, which would use Federal taxpayer dollars to support and encourage the destruction of human life for research. The bill would compel all American taxpayers to pay for research that relies on the intentional destruction of human embryos for the derivation of stem cells, overturning the President's policy that funds research without promoting such ongoing destruction. If S. 5 were presented to the President, he would veto the bill.

The President strongly supports medical research and has worked with Congress to increase resources for the National Institutes of Health. This Administration is the first to provide Federal funds for human embryonic stem cell research and has done so without encouraging the destruction of human embryos. The President's policy permits the Federal funding of research using embryonic cell lines created prior to August 9, 2001, the date his policy was announced, along with stem cell research using other kinds of cell lines. Scientists can therefore explore the potential applications of such cells, but the Federal government does not offer incentives or encouragement for the destruction of human life.

Over the past six years, more than \$130 million in taxpayer dollars has been devoted to human embryonic stem cell research consistent with the President's policy. Overall, more than \$3 billion has gone to innovative research on all forms of stem cells, contributing to proven medical treatments that use human stem cells from adult and other non-embryonic sources. S. 5 seeks to replace the Administration's policy by providing Federal funding for the first time for a line of research that involves the intentional destruction of living human embryos for the derivation of their cells. Destroying nascent human life for research raises serious ethical problems, and millions of Americans consider the practice immoral. The Administration believes that government has a duty to use the people's money responsibly, both supporting important public purposes and respecting moral boundaries.

Alternative types of human stem cells – drawn from adults, children, umbilical-cord blood, and other non-embryonic sources, without doing harm to the donors – have already achieved therapeutic results in thousands of patients with many different diseases. Moreover, a recent series of encouraging research reports offer hope that stem cells drawn from non-embryonic sources may possess characteristics like those of embryonic stem cells. In addition, researchers are now working to develop promising new techniques to produce stem cells similar in nature to those derived from human embryos, but without harming or destroying embryos.

S. 5 clearly acknowledges the potential of these new advances and the importance of supporting

them. The provisions in the bill to promote research leading to new techniques for deriving the functional equivalent of embryonic stem cells, without harming or destroying human embryos, encourage science that is not in conflict with ethics.

Unfortunately, the bill then crosses that same ethical line, by proposing to use taxpayer dollars to encourage destruction of nascent human life for research. The Administration believes that research on alternative sources of stem cells is extremely promising and provides robust opportunities to advance science without compelling American taxpayers to participate in ongoing destruction of human embryos.

Moreover, private sector support and public funding by several States for this line of research, which will add up to several billion dollars in the coming few years, argue against the notion of any urgent shortfall of research funding. Whatever one's view of the ethical issues or the state of the research, the future of this field does not require a policy of Federal subsidies that is offensive to the moral principles of millions of Americans.

The President believes that by enacting appropriate policy safeguards while encouraging the development of novel scientific techniques, it is possible to advance scientific and medical frontiers without violating moral principles. S. 5 fails to find that balance and is therefore deeply troubling to millions of Americans. The President urges Congress to pass an alternative bill that would advance stem cell research without encouraging destruction of human life.

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14303

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14165

NARA Num.:

14062

FOIA IDs and Segments:

2017-0029-F

2014-0503-F

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Email	Coleman-Isakson: Stem Cells - To: Karl Zinsmeister - From: Christopher Papagianis	2	05/17/2007	P5;

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Email	Stem Cell Meetings - To: Karl Zinsmeister - From: Christopher Papagianis	1	05/03/2007	P5;

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Email	RE: Update Since We Spoke at 9am re Stem Cells - To: Christopher Papagianis - From: Karl Zinsmeister	1	05/09/2007	P5;

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COLLECTION:

Domestic Policy Council

SERIES:

Zinsmeister, W. Karl - Subject Files

FOLDER TITLE:

Stem Cells 2007: Stem Cell Veto 2007 and Executive Order [3]

FRC ID:

14303

OA Num.:

14165

NARA Num.:

14062

FOIA IDs and Segments:

2017-0029-F

2014-0503-F

RESTRICTION CODES

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Withdrawal Marker

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FORM	SUBJECT/TITLE	PAGES	DATE	RESTRICTION(S)
Email	More on Stem Cell Meetings - To: Karl Zinsmeister - From: Christopher Papagianis	1	05/08/2007	P5;

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FORM	SUBJECT/TITLE	PAGES	DATE	RESTRICTION(S)
Email	Update on Stem Cell Meeting/Plan - To: Karl Zinsmeister - From: Christopher Papagianis	1	05/07/2007	P5;

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FORM	SUBJECT/TITLE	PAGES	DATE	RESTRICTION(S)
Email	S.30 - To: Candida Wolff - From: Karl Zinsmeister	1	04/26/2007	P5;

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FORM	SUBJECT/TITLE	PAGES	DATE	RESTRICTION(S)
Email	Update and Proposal on Stem Cells - To: Karl Zinsmeister - From: Christopher Papagianis	2	04/26/2007	P5;

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FORM	SUBJECT/TITLE	PAGES	DATE	RESTRICTION(S)
Email	Notes for Weldon Call - To: Karl Zinsmeister - From: Christopher Papagianis	1	05/01/2007	P5;

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—
Sununu

call

Sununu

(Scientists do this
mid May)

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Records Not Subject to FOIA

Court Sealed - The document is withheld under a court seal and is not subject to the Freedom of Information Act.

Withdrawal Marker

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FORM	SUBJECT/TITLE	PAGES	DATE	RESTRICTION(S)
Email	RE: Update and Proposal on Stem Cells - To: Karl Zinsmeister - From: Christopher Papagianis	1	04/27/2007	P5;

**This marker identifies the original location of the withdrawn item listed above.
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COLLECTION:

Domestic Policy Council

SERIES:

Zinsmeister, W. Karl - Subject Files

FOLDER TITLE:

Stem Cells 2007: Stem Cell Veto 2007 and Executive Order [3]

FRC ID:

14303

OA Num.:

14165

NARA Num.:

14062

FOIA IDs and Segments:

2017-0029-F

2014-0503-F

RESTRICTION CODES

Presidential Records Act - [44 U.S.C. 2204(a)]

- P1 National Security Classified Information [(a)(1) of the PRA]
- P2 Relating to the appointment to Federal office [(a)(2) of the PRA]
- P3 Release would violate a Federal statute [(a)(3) of the PRA]
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- P6 Release would constitute a clearly unwarranted invasion of personal privacy [(a)(6) of the PRA]

PRM. Personal record misfile defined in accordance with 44 U.S.C. 2201(3).

Deed of Gift Restrictions

- A. Closed by Executive Order 13526 governing access to national security information.
- B. Closed by statute or by the agency which originated the document.
- C. Closed in accordance with restrictions contained in donor's deed of gift.

Freedom of Information Act - [5 U.S.C. 552(b)]

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Zinsmeister, Karl

From: Zinsmeister, Karl
Sent: Wednesday, April 11, 2007 6:46 PM
To: Zinsmeister, Karl
Subject: Fw: S 30, alternatives, passed 70-28

S.5. 63-34
Castle

-----Original Message-----

From: Papagianis, Christopher C.
To: Zinsmeister, Karl; Troy, Tevi D.
Sent: Wed Apr 11 18:41:36 2007
Subject: S 30, alternatives, passed 70-28

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FORM	SUBJECT/TITLE	PAGES	DATE	RESTRICTION(S)
Briefing	Policy Time - Domestic	2	06/18/2007	P5;

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COLLECTION:

Domestic Policy Council

SERIES:

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FOLDER TITLE:

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FORM	SUBJECT/TITLE	PAGES	DATE	RESTRICTION(S)
Draft	[Draft]	3	N.D.	P5;

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