

MAINE STATE LEGISLATURE

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FIRST REGULAR SESSION

ONE HUNDRED AND THIRTEENTH LEGISLATURE

Legislative Document

NO. 665

H.P. 495 House of Representatives, March 6, 1987
Reference to the Committee on State and Local Government
suggested and ordered printed.

EDWIN H. PERT, Clerk
Presented by Representative MCGOWAN of Canaan.
Cosponsored by Representatives CARROLL of Gray, MICHAUD
of East Millinocket and Senator USHER of Cumberland.

STATE OF MAINE

IN THE YEAR OF OUR LORD
NINETEEN HUNDRED AND EIGHTY-SEVEN

AN ACT to Reform Regulatory Proceedings
under Rulemaking of the Maine
Administrative Procedure Act.

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5 Be it enacted by the People of the State of Maine as
6 follows:

7 Sec. 1. 5 MRSA §8052, sub-§4, as amended by PL
8 1981, c. 524, §3, is further amended to read:

9 4. Relevant information considered. The agency
10 shall consider all relevant information available to
11 it, including statements and arguments filed, before
12 adopting any rule.

13 In considering the adoption of regulations, the agen-
14 cy shall adhere to the following rules:

1 A. Administrative decisions shall be based on
2 adequate information concerning the need for and
3 consequences of proposed rulemaking;

4 B. Potential benefits of regulation must out-
5 weigh potential cost to society;

6 C. Objectives shall be chosen to maximize net
7 benefits to society;

8 D. Among alternative approaches to any given
9 regulatory objective, the alternative involving
10 the least net cost to society shall be chosen un-
11 less there are legal or other compelling reasons
12 for choosing a different option; and

13 E. The agency shall seek to reduce dispropor-
14 tionate economic impacts of major rules on small
15 businesses through different compliance or re-
16 porting requirements or timetables, consolidating
17 or simplifying compliance and design standards
18 and exempting from requirements where appropri-
19 ate.

20 During the process of adoption of a rule, including
21 the public hearing and the post-hearing comment peri-
22 od, persons are to be encouraged to testify about and
23 comment on the cost and benefits of the proposed rule
24 and the analysis of alternative regulatory mechanisms
25 described in the regulatory impact analysis.

26 Sec. 2. 5 MRSA §8052, sub-§7, as amended by PL
27 1985, c. 680, §2, is further amended to read:

28 7. Adoption of rule. The agency shall, in adopt-
29 ing rules, be consistent with the terms of the pro-
30 posed rule, except to the extent it determines neces-
31 sary to address concerns raised in comments and make
32 specific findings supporting those changes. No rule
33 may be adopted which contains requirements that could
34 not reasonably have been anticipated from the notice
35 of rulemaking pursuant to section 8053, subsection 2
36 and from the terms of the proposed rule. No rule may
37 become effective unless:

38 A. The agency adopts it within 120 days of the
39 final date by which data, views or arguments may

1 be submitted to the agency for consideration in
2 adopting the rule; and

3 B. This adopted rule is approved by the Attorney
4 General as to form and legality, as required by
5 section 8056, within 150 days of the final date
6 by which those comments may be submitted.

7 The final date for comments may be extended if notice
8 of doing so is published before that final date, in
9 the consolidated notice referred to in section 8053.

10 Sec. 3. 5 MRSA §8053, sub-§3-B is enacted to
11 read:

12 3-B. Regulatory impact analysis. All agencies
13 shall prepare and make available to all interested
14 parties at the time of publication of the notice re-
15 ferred to in subsection 2, a statement describing in
16 clear terms the expected impact of the proposed rule
17 on benefited and regulated entities. These statements
18 shall provide as comprehensive an analysis as possi-
19 ble under the particular circumstances and shall, at
20 a minimum, contain the following:

21 A. A specific citation of the law to be imple-
22 mented by the proposed rule;

23 B. A description of the objective sought to be
24 achieved by the rule;

25 C. A description of the potential benefits, in-
26 cluding those that cannot be quantified in mone-
27 etary terms, and of those likely to receive these
28 benefits;

29 D. A description of the potential costs, includ-
30 ing those that cannot be quantified in monetary
31 terms and of those likely to bear these costs;

32 E. A description of the potential net benefits
33 of the rule, including an evaluation of effects
34 that cannot be quantified in monetary terms;

35 F. A description of the available alternative
36 regulatory approaches that could achieve substan-
37 tially the same regulatory objective, at lower

1 cost, together with a statement of the potential
2 benefits and costs and a brief description of the
3 reasons which prevent the alternatives from being
4 substituted for the one proposed;

5 G. A description of the reporting, record keep-
6 ing and other compliance requirements, including
7 costs of equipment, supplies, labor, legal, con-
8 sulting and accounting services that may be in-
9 curring; and

10 H. A description of the types of businesses that
11 may be affected, using standard industrial clas-
12 sification codes, together with information on
13 the number of small, medium and large businesses
14 within those types.

15 Sec. 4. 5 MRSA §8057-A is enacted to read:

16 §8057-A. Periodic meetings

17 State agencies shall conduct periodic informal
18 meetings with parties affected by agency regulatory
19 decision making to discuss the efficiency of the rule
20 development process, the use and utility of negotia-
21 tion techniques in rule development, the direct and
22 indirect costs and benefits, the intended impact of
23 agency rules generally and ways in which the rule de-
24 velopment process can be improved. The agency shall
25 report to the Governor on steps it has taken as a
26 consequence of this meeting, including plans for
27 periodic future meetings for this same purpose. Meet-
28 ings held pursuant to this provision shall be sched-
29 uled at least once a year and notice shall be pro-
30 vided to those persons listed in section 8053, sub-
31 section 1, paragraph A.

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STATEMENT OF FACT

2 The purpose of this bill is to implement the
3 terms of Executive Order 13, fiscal year 1985-86
4 dealing with regulatory reform. As noted in the Exec-
5 utive Order, "decisions that reflect a balanced ap-
6 proach to protecting the public health and welfare
7 are inherently beneficial."

8 In addition to implementing the Executive Order,
9 section 2 of this bill also imposes a prohibition on
10 extensive changes from proposed to final rules, with-
11 out allowing opportunity for involvement by those who
12 did not participate in the process because they may
13 not have been affected by a proposed rule, but are
14 affected by the final rule, adopted after the record
15 has closed.

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