| Durang d Action   | D         | T           | Nat Luculan auto d | Planning to | Not Planning to |
|---|-----------|-------------|--------------------|-------------|-----------------|
| Proposed Action   | Responses | impiementea | Not Implemented    | Implement   | Implement       |
| Eliminate annual protocol renewals for non-USDA species and non-DOD protocols.  | 94        | 44 (47%)    | 50 (53%)           | 11 (12%)    | 39 (41%)        |
| Discontinue the USDA pain and distress<br>classifications for non-Animal Welfare<br>Act regulated species   | 93        | 14 (15%)    | 79 (85%)           | 10 (11%)    | 69 (74%)        |
| As the default, implement Designated<br>Member Review rather than Full<br>Committee Review  | 94        | 62 (66%)    | 32 (34%)           | 3 (3%)      | 29 (31%)        |
| Reduce IACUC requirements for experimental details that are unrelated to the health and safety of animals   | 93        | 65 (70%)    | 28 (30%)           | 7(8%)       | 19 (20%)        |
| Adopt NIH OLAW's allowance for<br>"expediting" protocol amendments via<br>a new Veterinary Verification and<br>Consultation (VVC) process, thereby<br>reducing/eliminating full IACUC<br>involvement  | 94        | 72 (77%)    | 22 (23%)           | 10 (11%)    | 12 (13%)        |
| Expand the scope of administrative approval authority by allowing small changes to protocols to be handled administratively (by IACUC staff or via Veterinary Verification and Consultation process)  | 94        | 87 (93%)    | 7 (7%)             | 5 (5%)      | 2 (2%)          |
| Simplify the IACUC protocol form with standardized language and content requirements. Examples include 1) A standard rodent, pre-approved, pre-operative preparation plan and post-operative recovery plan template 2) A built in commonly used drug formulary that will provide the pre-approved dosage, route and frequency of administration 3) Convert formatted text boxes to simple drop down lists for fast data selection/entry and enhanced reporting capabilities 4) Drop down list of euthanasia methods/dosages that are pre-approved for a given species |           |             |                    | 45 (48%)    | 12 (13%)        |
| Does your institution provide template language?  | 94        | 50 (53%)    | 44 (47%)           |             |                 |
| Does your institution provide procedural libraries?   | 86        | 32 (37%)    | 54 (63%)           |             |                 |
| Replace required documentation on how a proposed protocol was not unnecessarily duplicative with a simple attestation.  | 94        | 63 (67%)    | 31 (33%)           | 14 (15%)    | 17 (18%)        |
| Standardize veterinary review procedures and communications to investigators.   | 92        | 69 (75%)    | 23 (25%)           | 7 (8%)      | 16 (17%)        |
| Review SOPs on a less frequent basis<br>(e.g. every two to three years) based on<br>potential risk (e.g., skill of investigative<br>team, outcomes of PAM reporting)  | 93        | 73 (78%)    | 20 (22%)           | 10 (11%)    | 8 (9%)          |
| Allow investigators to provide an approximate number or range of animals needed over the course of a research project rather than an exact number   | 94        | 41 (44%)    | 53 (56%)           | 9 (10%)     | 40 (43%)        |

| Allow investigators to provide a range of time for post op observation rather than an exact time (e.g., 4-6 hours) instead of every 4 hours) to allow flexibility and avoid findings on deviations from the language where actions were appropriate Replace mandatory triennial regulatory refresher seminar with an array of instructional sessions to streamline protocol writing and review. Solicit feedback on how the institution can assist investigators Adopt standardized models for training and documentation.  Allow use of ad hoc consultants in place of IACUC members (e.g. environmental health and safety or physical plant personnel trained to assess facilities for semianual inspections of non USDA species  Eliminate the requirement for a literature search for category D and E procedures for non-USDA species  Eliminate the requirement for a literature search for category C patch and the procedures for all species and funding source per protocol  Eliminate the requirement to describe facilities and husbandry in the protocol  Eliminate the requirement to describe facilities and husbandry in the protocol  Eliminate the requirement to describe facilities and centralized staff support.  Do not require a protocol re-write for PISI triennial review if your process allows modifications to be included in the latest version of the protocol rather.   |   |     |                  |             |                         |            |
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| if a protocol issuing central facilities and centralized staff support.  Do not require a protocol re-write for PHS triennial review if your process allows modifications to be included in the latest version of the protocol rather  31 (33%)  62 (67%)  12 (13%)  48 (52%)  | 1   |     |                  |             |                         |            |
| and centralized staff support.  Do not require a protocol re-write for PHS triennial review if your process allows modifications to be included in the latest version of the protocol rather   |   | 93  | 80 (86%)         | 13 (14%)    | 6 (7%)                  | 6 (7%)     |
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| then as an attachment  | the latest version of the protocol rather |     |                  |             |                         |            |
| unan as an attachment.   | than as an attachment.                    |     |                  |             |                         |            |

<sup>\*</sup>Percentage of Total Responses