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| Questions | Stakeholder response |
| * Should the ST PASA account for any additional input factors beyond those listed in new NER 3.7.3(g)(1) to (3)? Please outline how any suggested additions would further the PASA objective.
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| * Should the Procedures include any other information, including any additional ST PASA outputs, not canvassed in the consultation paper? Please outline how any suggested additions would further the PASA objective.
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| * Are there any material adverse impacts of the proposal for registered participants, relative to the current ST PASA requirements?
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| * Is the proposed 168-hour maximum recall period appropriate for PASA availability, or should it be longer or shorter?
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| * Are any additional clarifications needed from AEMO about the ST PASA process?
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| * Are there any adverse impacts for particpants if LRC runs are discontinued as proposed?
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| * Do participants have any other observations/comments?
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