Medicines and Medical Devices Bill

AMENDMENTS

TO BE MOVED

IN GRAND COMMITTEE

Clause 13

LORD LANSLEY

Page 7, line 37, at end insert—

"( ) requirements that, in accordance with any recommendation of the National Institute for Health and Care Excellence, the National Health Service should make a medical device available for use within a specified period,"

After Clause 16

BARONESS BARKER
BARONESS BURT OF SOLIHULL

Insert the following new Clause—

"Introduction of registries for patient safety

(1) Within 6 months of this Act coming into force, the Secretary of State must by regulations introduce provisions enabling the creation of a registry of patients who have undergone a procedure to insert surgical mesh.

(2) The purpose of the registry under subsection (1) is to research and audit the outcomes of such surgeries in terms of the device safety and patient reported outcomes measures.

(3) When creating the registry under subsection (1) the Government must consult—

(a) relevant patient groups;

(b) healthcare professionals; and

(c) anyone else whom the Secretary of State considers appropriate.

(4) Accompanying the regulations the Secretary of State must also lay before Parliament a report outlining the progress towards the establishment of a comprehensive national database of patients who have undergone a procedure involving the insertion of a medical device or implant, and how the database may feed into the creation of further registries.

After Clause 16 - *continued*

(5) A statutory instrument containing regulations under this section may not be made unless a draft of the instrument has been laid before and approved by a resolution of each House of Parliament."

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21 September 2020
