1	State of Arkansas	A Bill	
2	89th General Assembly	A DIII	
3	Regular Session, 2013		HOUSE BILL 1267
4			a
5	By: Representatives Kerr, Branscum, Carnine, Clemmer, Cozart, Ferguson, Gillam, Gossage, Hobbs,		
6	Leding, Linck, Lowery, Magi		
7	By: Senators J. Dismang, Holland, J. Hutchinson, J. Key, Rapert		
8		For An Act To Be Entitled	
9	AN ACT TO REQUIRE A PRIOR APPROVAL PROCESS FOR		, DOD
10		·	
11		AL AND INVESTIGATIONAL SURGICAL	
12	AND MEDICA	L DEVICES; AND FOR OTHER PURPOSE	iS.
13 14			
14 15		Subtitle	
16	AN AC	CT TO REQUIRE A PRIOR APPROVAL	
10 17		ESS FOR EXPERIMENTAL AND	
17		STIGATIONAL SURGICAL PROCEDURES A	AND
10 19		CAL DEVICES.	AND
20	нды	AL DEVICES.	
21			
22	RE IT ENACTED BY THE G	ENERAL ASSEMBLY OF THE STATE OF	ΔΡΚΔΝSΔS.
23	DI II IMMOTID DI IIII O		industry.
24	SECTION 1. Arka	nsas Code Title 23, Chapter 86,	Subchapter 1, is
25		tional section to read as follow	- · · · · ·
26	23-86-122. Prio	r approval process for experimen	tal and investigational
27	surgical procedures an	d medical devices.	
28	(a) "Health car	rier" means a health maintenance	organization, hospital
29	medical service corpor	ation, or a disability insurance	company.
30	(A)	"Health carrier" includes a sel	f-insured governmental
31	or church plan and thi	rd-party administrators that adm	inister or adjust
32	disability benefits for a disability insurer, hospital medical service		
33	corporation, health maintenance organization, self-insured governmental plan		
34	or self-insured church	plan.	
35	<u>(B)</u>	"Health carrier" does not inclu	de:
36		(i) An automobile insurer pay	ing medical or hospital

T	benefits under § 23-89-202(1) or a self-insured employer health benefits	
2	plan; or	
3	(ii) A person, company, or organization licensed or	
4	registered to issue or who issues any insurance policy or insurance contract	
5	in this state as described in §§ 23-62-102 and 23-62-104 — 23-62-107	
6	providing medical or hospital benefits for accidental injury or disability.	
7	(b) A health carrier that denies coverage for a specific surgical	
8	procedure or medical device cleared or approved for marketing by the United	
9	States Food and Drug Administration as experimental, investigational, or both	
10	shall develop a process by which a surgeon, before providing the service, may	
11	present medical evidence to obtain an exception for the individual patient	
12	for coverage of the surgical procedure or medical device, or both.	
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