



Basic information	
<p>1995/0013(COD)</p> <p>COD - Ordinary legislative procedure (ex-codecision procedure) Directive</p>	Procedure completed
<p>In vitro diagnostic medical devices: security requirements</p> <p>Repealed by 2012/0267(COD)</p> <p>Subject</p> <p>2.10.03 Standardisation, EC/EU standards and trade mark, certification, compliance 4.20.02 Medical research 4.20.04 Pharmaceutical products and industry 4.60.02 Consumer information, advertising, labelling</p>	

Key events			
Date	Event	Reference	Summary
24/10/1994	Additional information		Summary
19/04/1995	Legislative proposal published	COM(1995)0130 	
15/05/1995	Committee referral announced in Parliament, 1st reading		
06/02/1996	Vote in committee, 1st reading		
06/02/1996	Committee report tabled for plenary, 1st reading	A4-0031/1996	
12/03/1996	Decision by Parliament, 1st reading	T4-0115/1996	
12/03/1996	Debate in Parliament	CRE link	Summary
20/12/1996	Modified legislative proposal published	COM(1996)0643 	Summary
21/05/1997	Debate in Council		
23/03/1998	Council position published	05255/1/1998	Summary
02/04/1998	Committee referral announced in Parliament, 2nd reading		
03/06/1998	Vote in committee, 2nd reading		Summary
03/06/1998	Committee recommendation tabled for plenary, 2nd reading	A4-0225/1998	
17/06/1998	Debate in Parliament	CRE link	Summary
18/06/1998	Decision by Parliament, 1st reading	T4-0362/1998	Summary
05/10/1998	Act approved by Council, 2nd reading		
27/10/1998	Final act signed		
27/10/1998	End of procedure in Parliament		
07/12/1998	Final act published in Official Journal		