The European Association Medical Devices - Notified Bodies

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Team-NB Notified Bodies members' intention to submit application to be designated against MDR and/or IVDR

In the framework of the new Medical Device Regulations, you will find below the Team-NB members' intention to submit their application to be designated against MDR and/or IVDR.
The Team-NB notified bodies members have a double advantage to pass their designation audit:

- ✓ The compliance with the Code of Conduct whose requirements are in advance of the current MDD and IVDD and is a big progression towards complying with the new MDR and IVDR.
- ✓ The Team-NB working groups reviewing the new texts of the regulations to help the members to identify and interpret the impact of the new requirements on the procedures and processes of the Notified Bodies. The aim is that the working group will provide guidelines to support the members in submitting their application in the quickest time and be ready to pass their designation audit against the new regulations.

Id. Number	Notified Body	Country	MDR	IVDR
0473	Intertek AMTAC Certification Services	UK	✔	
0086	BSI Product Certification	UK	✔	✔
1912	DARE!! Medical Certifications	NL	✔	
0124	DEKRA Certification GmbH	D	✔	
0344	DEKRA Certification B.V.	NL	✔	✔
0653	EKAPTY SA	GR	✔	
1282	Ente Certificazione Macchine Srl.	IT	✔	✔
2460	DNV Nemko Presafe AS	NO	✔	
0297	DQS Medizinprodukte	D	✔	
0413	Intertek Semko	S	✔	
1984	Kiwa	TR	✔	
0459	LNE/G-MED	F	✔	✔
0088	LRQA	UK	✔	✔
0483	mdc medical device certification	D	✔	✔
0482	MEDCERT	D	✔	
0050	NSAI	IRL	✔	✔
0543	Presafe Denmark A/R	DK	✔	✔
0120	SGS	UK	✔	✔
1304	SIQ Ljubljana	SL	✔	
2195	SZUTEST	TR	✔	
0044	TÜV Nord Cert.	D	✔	
0197	TÜV Rheinland LGA Products	D	✔	✔
0123	TÜV SÜD	D	✔	✔
0843	UL International Limited	UK	✔	✔