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# Easy Links To Essential Pages On The European Commission's Medical Device Website

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New documents and links related to the Medical Device and IVD Regulations are published by the European Commission almost daily. Here is a quick guide to where to find the information you need.

Quickly finding the correct European Commission's webpage to access information relating to the new Medical Device and IVD Regulations is not always easy. The website is now heavily populated with information and forever expanding; navigating it can be like finding your way through a maze.

Here are some quick links to essential information you may need in one handy guide:

| Quick Links To Key European Commission's MDR and IVDR webpages    |
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| <a href="#">Medical Device Regulation (EU) 2017/745 (MDR)</a>     |
| <a href="#">IVD Regulation (EU) 2017/746 (IVDR)</a>               |
| <a href="#">List of notified bodies designated under the MDR</a>  |
| <a href="#">List of notified bodies designated under the IVDR</a> |
| <a href="#">MDR expert panel opinions</a>                         |
| <a href="#">IVDR expert panel opinions</a>                        |
| <a href="#">Implementing and delegated acts</a>                   |
| <a href="#">Guidance documents</a>                                |
| <a href="#">MDR/IVDR Harmonized standards</a>                     |
| <a href="#">Ongoing guidance deliverables</a>                     |
| <a href="#">Implementation rolling plan</a>                       |
| <a href="#">MDR/IVDR MDCG working groups agenda</a>               |

There was also a flurry of documents published by the notified body association, TEAM-NB, in October that are intended to support the implementation of the MDR and IVDR. They are available [here](#).