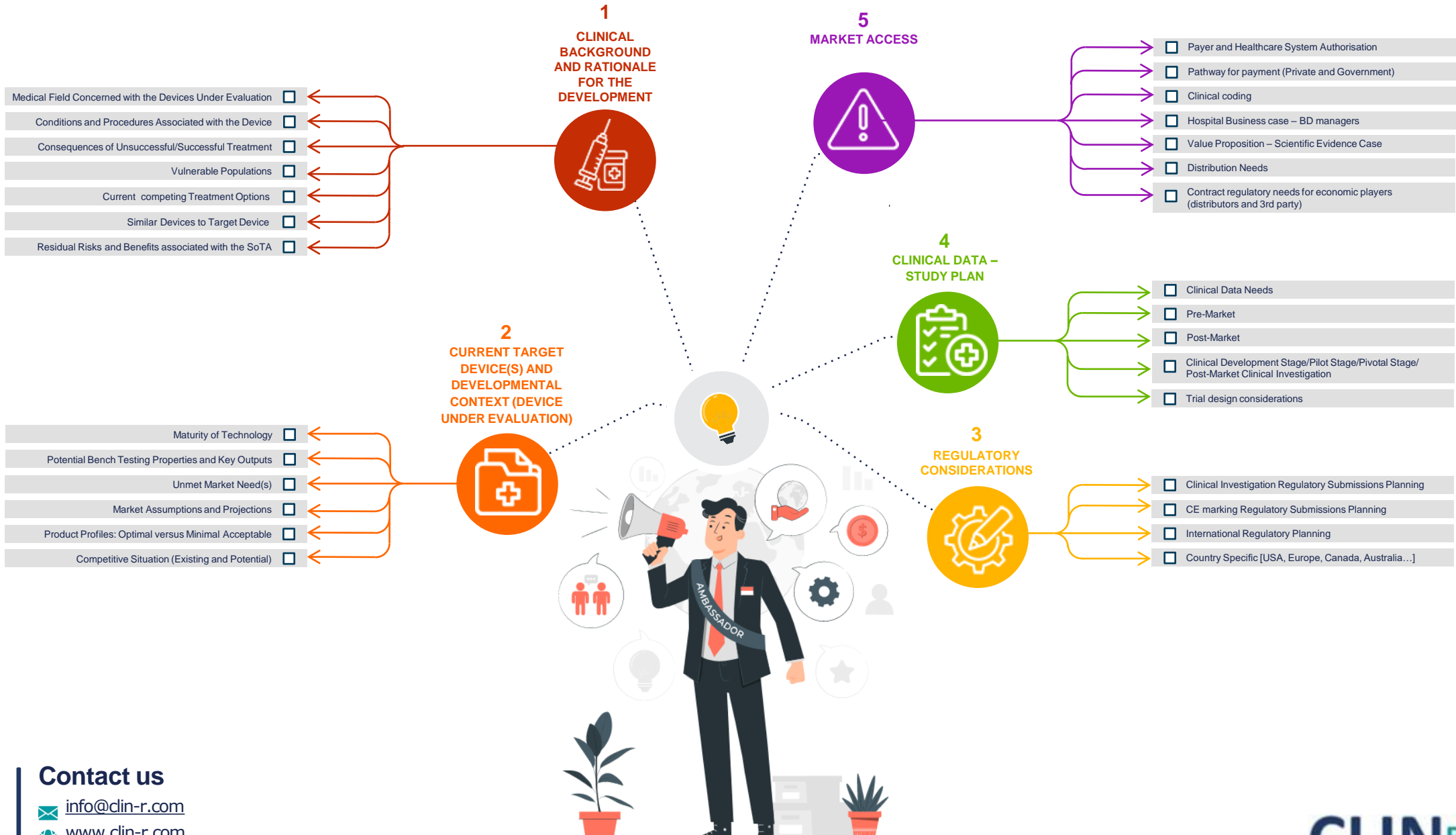


CLINICAL DEVELOPMENT PLAN

TO MEET MEDTECH INTERNATIONAL MARKET PLANNING AND INVESTMENT NEEDS

“Annex XIV of the Medical Devices Regulation (EU) 2017/745 (MDR), section 1(a), 8th indent, requires a Clinical Development Plan (CDP). Some manufacturers may need more clarity, though, in order to meet CDP requirements as part of their broader MDR compliance effort



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