NAMCP Medical Technologies Dossier Template

PURPOSE: The purpose is to provide medical directors and manufacturers with a medical device dossier format that accounts for evidence development approaches and unique aspects of these technology types (versus simply following drug-based formats that may not be ideal for other types of health technologies).

This format has been developed with Medical Directors and device providers working collaboratively to ensure that the format is the most useful, practical, and succinct for US payer decision making.

Executive Summary (1-2 pages total)
Disease Clinical and Economic Burden Overview – half a page
Intended Indication and Use - one paragraph
Core Value and Evidence Supporting TECHNOLOGY (including key value messages/themes/510K status) 1-2 pages

List of Tables
List of Figures
List of Abbreviations

1 Burden of Illness (section to be succinct)
1.1 Clinical Characteristics and Presentation of Medical Condition
1.2 Epidemiology
  1.2.1 Incidence of Medical Condition
  1.2.2 Prevalence of Medical Condition
  1.2.3 Risk Factors for Medical Condition
1.3 Clinical
  1.3.1 Major Adverse Health Outcomes Associated with Medical Condition
  1.3.2 Key Comorbidities and Increased Risk for Adverse Health Outcomes

1.4 Unmet Need
  1.4.1 Description of Unmet Need
    • Listing of Alternative Tests/Treatments including pharmacological, and their outcomes, adverse effects, in market place currently etc.
    • Description and characteristics of device/technology
    • Procedure codes to include J Code, CPT & ICD-10
  1.4.2 How the New Technology Will Address Unmet Need
    • Indicated Patients
    • Fit in Clinical Pathway

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2 Product Information
2.1 Technology Description and Characteristics
2.2 Device Classification and Approval Status
2.3 Procedure Codes if applicable to include JCode, CPT and ICD-10
2.4 Device Components and Specifications

3 Value Evidence Supporting TECHNOLOGY
3.1 Clinical Efficacy and Patient Outcomes
3.1.1 Core Value Drivers - (i.e., Value Messages) with supporting information/references including description of associated clinical (pre-clinical, feasibility, pivotal studies) registry and post market data
3.1.2 Role of TECHNOLOGY in Improving Patient Management and Health Outcomes

3.2 Quality of Life Outcomes
3.2.1 Core Value Drivers - (i.e., Value Messages) with supporting information/references
3.2.2 Role of TECHNOLOGY in Improving Patient Quality of Life
3.2.3 Device Safety Profile

3.3 Economic Outcomes
3.3.1 Core Value Drivers - (i.e., Value Messages) with supporting information/references including description of associated clinical, registry and post-market data and models
3.3.2 Role of TECHNOLOGY in Improving Economic Outcomes/Cost Offsets

3.4 Econometrics
3.4.1 Direct Costs
3.4.2 Indirect Costs

4 Current Payer Coverage
4.1 Describe current coverage status among payers.

5 Future Directions and Applications
5.1 Emerging Clinical Applications – limit to 1 page if applicable

Appendix A: Literature Search Methodology
- Provide a list of publications on the product for committee review

Appendix B: References

Appendix C: Additional Information (e.g., evidence tables)