

Table 1. Characteristics of the available forty-five Health Technology Assessment Reports on omics-technologies.

Report name [Reference]	Year of publication	Agency	Country	Technology Evaluated	Framework Reported	Health Problem	Description and technical characteristics	Safety	Clinical effectiveness	Costs and economic valuation	Ethical analysis	Organisational aspects	Social aspects
Clinical Applications of Proteomic Techniques [64]	2006	Agencia de Evaluación de Tecnologías Sanitarias de Andalucía	Spain	Proteomics	Not Reported	1	1	0	1	0	0	0	0
Prostate cancer gene 3 (ProgenSA PCA3) assay in the diagnosis of prostate cancer [72]	2006	National Horizon Scanning Center	UK	The ProgenSA PCA3	Not reported	0	1	1	1	1	0	0	0
Genetic screening for familial hypercholesterolaemia [70]	2007	Australian Safety and Efficacy Register of New Interventional Procedures-Surgical	Australia	Genetic screening	Not Reported	1	1	1	1	1	1	0	0
Impact of gene expression profiling tests on breast cancer outcomes [69]	2008	Agency for Healthcare Research and Quality	USA	Oncotype DX; MammaPrint; Breast Cancer Profiling test	EGAPP	1	1	1	1	1	0	0	0
A comparison of Gene Expression Profiling tests for Breast cancer [65]	2009	Health Services Assessment Collaboration	New Zealand	OncotypeDx, MammaPrint	Not Reported	1	1	0	1	1	0	0	0
MammaPrint®: in vitro test for the evaluation of individual risk of metastasis in surgically treated women for breast cancer [67]	2011	Agencia sanitaria e sociale regionale- Regione Emilia-Romagna	Italy	MammaPrint	Not Reported	0	1	1	1	1	1	1	1

Economic assessment of genetic tests in mammary carcinoma cancer treatment [51]	2011	Health Technology Assessment Unit of Madrid	Spain	OncotypeDx, MammaPrint	Not Reported	0	0	0	1	1	0	0	0
Genetic testing for hereditary mutations in the VHL gene that cause von Hippel-Lindau syndrome [68]	2012	Adelaide Health Technology Assessment	Australia	Genetic testing	Diagnostic assessment framework	1	1	1	1	1	1	0	0
Allomap™. Genetic test for cardiac transplant rejection. [66]	2012	Agencia de Evaluación de Tecnologías Sanitarias de Andalucía	Spain	Allomap™	Not Reported	1	0	1	1	1	0	0	0
Gene expression profiling and expanded immunohistochemistry tests to guide the use of adjuvant chemotherapy in breast cancer management: a systematic review and cost-effectiveness analysis [43]	2013	National Institute for Health and Care Excellence (University of Sheffield)	UK	OncotypeDx, MammaPrint, Endopredict, Prosigna	EGAPP	1	0	0	1	1	0	0	0
Importance of gene profiling tests in the choice of treatment for breast cancer [59]	2013	finCCHTA	Finland	OncotypeDx, MammaPrint, Endopredict	ACCE	1	1	0	1	1	0	1	0
Genetic testing for hereditary mutations in the RET gene [42]	2013	Adelaide Health Technology Assessment	Australia	Genetic testing	Diagnostic assessment framework	1	1	1	1	1	1	1	0
Use of OncotypeDX for guiding adjuvant chemotherapy decisions in early stage invasive breast cancer patients in Alberta [53]	2013	Health Technology Assessment Unit of University of Alberta	Canada	OncotypeDx	Not Reported	1	1	1	1	1	0	0	1

Prognostic genomic tests in early breast: MammaPrint® and Oncotype DX® [49]	2014	Agencia de Evaluación de Tecnologías Sanitarias de Andalucía	Spain	MammaPrint; Oncotype DX	Not Reported	1	1	0	1	1	0	0	0
Technology Assessment of Molecular Pathology Testing for the Estimation of Prognosis for Common Cancers [58]	2014	Agency for Healthcare Research and Quality	USA	Molecular Pathology Testing	Not Reported	1	1	1	1	1	1	1	1
Gene expression profiling of 21 genes in breast cancer to quantify the risk of disease recurrence and predict adjuvant chemotherapy benefit [47]	2014	Medical Services Advisory Committee	Australia	OncotypeDx	Not Reported	1	0	1	1	1	0	1	0
Oncotype Dx in Women and Men with ER-Positive Her2 Negative Early Stage Breast Cancer Who are Lymph Node-Positive: A review of Clinical Effectiveness and Guidelines[54]	2014	Canadian Agency for Drugs and Technologies in Health	Canada	OncotypeDx	Not Reported	0	0	0	1	0	0	0	0
Next Generation DNA Sequencing, A review of the cost-effectiveness and Guidelines [63]	2014	Canadian Agency for Drugs and Technologies in Health	Canada	Next Generation Sequencing	Not Reported	0	0	0	1	1	0	0	0
Second generation prognostic genomic tests in early breast cancer: EndoPredict® & Prosigna™ [62]	2015	Agencia de Evaluación de Tecnologías Sanitarias de Andalucía	Spain	EndoPredict®; Prosigna™	Not Reported	1	1	0	1	0	0	0	0

Proteomanalyse im Urin zur Erkennung einer diabetischen Nephropathie bei Patientinnen und Patienten mit Diabetes mellitus und arteriellem Hypertonus [52]	2015	Institut für Qualität und Wirtschaftlichkeit im Gesundheitswesen	Germany	Proteomics	Not Reported	1	1	0	1	0	0	0	0
Next Generation Sequencing Gene panels for Targeted Therapy in Oncology and Haemato-Oncology [40]	2015	Belgian Health Care Knowledge Centre	Belgium	Next Generation Sequencing / In (Hemato)-Oncology	Not reported	0	1	1	1	1	0	1	1
Gene Expression Profiling And Immunohistochemistry Tests For Personalised Management of Adjuvant Chemotherapy Decisions in Early Breast Cancer [71]	2015	Belgian Health Care Knowledge Centre	Belgium	OncotypeDx, Mammprint	Not Reported	1	1	1	1	1	0	0	0
Next Generation sequencing in diagnostiek [50]	2015	Health Council of the Netherlands	The Netherlands	Next Generation Sequencing	Not Reported	1	1	0	0	1	1	1	0
Gene sequencing of tumours: clinical validity and utility of molecular profiles obtained by next-generation sequencing technologies [56]	2015	l'Institut national d'excellence en santé et en services sociaux	Canada	Next Generation Sequencing	ACCE	0	1	0	1	0	0	0	0

Using The Oncotype DX ® Assay For Therapeutic Decision Making In The Context Of Invasive Breast Cancer Treatment [57]	2016	l'Institut national d'excellence en santé et en services sociaux	Canada	Oncotype DX	Not Reported	0	1	0	1	1	0	0	0
Biomarker-based tests for the decision for or against adjuvant systemic chemotherapy in primary breast cancer [77]	2016	Institut für Qualität und Wirtschaftlichkeit im Gesundheitswesen	Germany	MAAA	Not Reported	1	0	0	1	0	0	0	0
Prenatal Diagnosis through Next Generation Sequencing [74]	2016	Swedish Agency for Health Technology Assessment and Assessment of Social Services	Sweden	Next Generation Sequencing / In prenatal Screening	Not Reported	0	1	0	0	0	1	0	1
HTA Report Next Generation Sequencing (NGS) [44]	2017	Agenas	Italy	Next Generation Sequencing	Not Reported	1	1	1	1	1	1	1	1
Genexpressionstest Mammaprint® [78]	2017	EUnetHTA Collaboration	EU	Mammaprint	EUnetHTA Core model	1	1	1	1	0	0	0	0
Utility of exome sequencing for diagnosed dysmorphic syndromes, with or without intellectual disabilities. Literature review Executive summary [39]	2017	Agencia de Evaluación de Tecnologías Sanitarias de Andalucía	Spain	Exome Sequencing	Not Reported	0	0	0	0	0	1	1	0

Molecular testing for Lynch syndrome in people with colorectal cancer: systematic reviews and economic evaluation [75]	2017	National Horizon Scanning Center	UK	Microsatellite instability testing	EGAPP	1	1	1	0	1	0	0	0
Pharmacogenomic Testing for Psychotropic Medication Selection: A Systematic Review of the Assurex GeneSight Psychotropic Test [83]	2017	Ontario Health Quality	Canada	Pharmacogenomic testing	Not Reported	1	1	0	1	0	0	0	0
Prognostic genomic tests in early breast cancer: up-date of evidence. Executive summary [76]	2018	Agencia de Evaluación de Tecnologías Sanitarias de Andalucía	Spain	MammaPrint, Oncotype DX, EndoPredict and Prosigna	Not reported	0	0	1	1	0	0	0	0
Use Of Endopredict And Prosigna In Early Invasive Breast Cancer [46]	2018	l'Institut national d'excellence en santé et en services sociaux.	Canada	Endopredict, Prosigna	EGAPP	0	1	0	1	1	0	0	0
Mammaprint Test for personalised management of adjuvant chemotherapy decisions in early breast cancer [45]	2018	Belgian Health Care Knowledge Centre	Belgium	Mammaprint	Not Reported	1	0	1	1	1	0	1	0
The use of whole genome sequencing in clinical practice: Challenges and organisational considerations for Belgium [41]	2018	Belgian Health Care Knowledge Centre	Belgium	Whole Genome Sequencing	Not Reported	0	1	0	0	1	0	1	0

FoundationOne® CDx: genetic profiling of solid tumours [82]	2019	Ludwig Boltzmann Institut	Austria	Next-Generation Sequencing	Not Reported	1	0	0	0	1	0	0	0
Rapid Genome-wide Testing: Clinical Utility and Cost-Effectiveness [48]	2019	Canadian Agency for Drugs and Technologies in Health	Canada	Genome-wide Testing	Not Reported	0	0	0	1	1	0	0	0
Complete sequencing of the Exome for Etiological Investigation of intellectual disability of Indeterminate cause [73]	2019	Conitec	Brasil	Sequenciamento de nova geração – sequenciamento completo do exoma	Not Reported	0	1	0	1	1	0	0	0
Clinical utility of genomic signature in Early-stage breast cancer [47]	2019	Haute autorité de santé	France	MAAA	Not Reported	0	1	0	1	0	0	0	1
Prosigna Gene Signature to Assess Expected Benefit from Chemotherapy in Breast Cancer [60]	2019	Norwegian Institute of Public Health	Norway	Prosigna	Not Reported	1	1	1	1	1	0	1	0
Three biomarker tests to help diagnose preterm labour: a systematic review and economic evaluation [80]	2019	National Institute for Health Research	UK	Diagnostic tests PartoSur; Actim Partus; Rapid Fetal Fibronectin 10Q Cassette Kit	Not Reported	1	1	1	1	1	0	0	0
Gene Expression Profiling Tests for Early-Stage Invasive Breast Cancer: A Health Technology Assessment [61]	2020	Canadian Agency for Drugs and Technologies in Health AND Ontario Health Technology Assessment Series	Canada	OncotypeDx, MammaPrint, Endopredict, Prosigna	Not Reported	1	0	1	1	1	0	1	0

Genetic risk prediction test for cardiovascular disease [79]	2020	Agencia de Evaluación de Tecnologías Sanitarias de Andalucía	Spain	Genetic risk prediction test	Not reported	1	1	1	1	1	1	1	0
Genome-Wide Sequencing for Unexplained Developmental Disabilities or Multiple Congenital Anomalies: A Health Technology Assessment [81]	2020	Ontario Health Quality	Canada	Genome-Wide Sequencing	Not Reported	1	1	1	1	1	0	0	0
Total (45)						28	31	22	38	33	10	14	7

Abbreviations: ACCE: Analytic validity, Clinical validity, Clinical utility, Ethical, legal and social implications; EGAPP: Evaluation of Genomic Applications in Practice and Prevention; MAAA: multianalyte assays with algorithmic analysis;