Vengono indicale le VEQ per le quali sono già disponibili i risultati.
Le VEQ dell'anno in corso verranno riportate dopo valutazione.
I risultati delle VEQ indicate sono disponibili su richiesta.

VEQ CEQAS - GenQA	
Tipo	**Anno**
Cytogenetic - Amniotic Fluid	2009, 2010, 2011, 2012, 2013, 2014, 2015, 2016, 2017, 2018, 2019, 2021
Cytogenetic -Blood (Postnatal karyotyping)	2009, 2010, 2011, 2012, 2013, 2014, 2015, 2016, 2017, 2018, 2019, 2022, 2023
Cytogenetic -Chorionic Villus	2011, 2012, 2013, 2014, 2015, 2016, 2017, 2018, 2019, 2021
Cytogenetic -Myeloid (AML/MDS/CML)	2014, 2015, 2016, 2017, 2018, 2019, 2021, 2022, 2023, 2024, 2025, 2026
Haematological Technical FISH	2022
Myeloma	2023, 2024, 2026
Constitutional microarray analysis (CMA) (postnatal)	2011, 2012, 2013, 2014, 2017, 2025
Prenatal constitutional Copy Number Variant (CNV) detection	2016, 2020, 2021, 2023, 2024
Pathogenicity of germline postnatal copy number variants (CNV) (Classification only)	2020
Genetic Counselling -Dysmorphology	2015, 2016, 2017
Clinical Genetic -Dysmorphology	2022, 2025
Genetic Counselling -Monogenic disorder	2015, 2016
Genetic Counselling -Oncogenetics	2015, 2016, 2017
Genetic counselling	2021, 2022, 2024
Clinical Genetics - Oncogenetics	2021, 2022, 2023, 2024, 2025, 2026
Clinical Genetics - Monogenic Disorders	2021, 2023, 2026
Familial Colorectal Cancer and Polyposis	2022, 2025
BRCA testing for ovarian and prostate cancer - somatic	2022, 2024
Next Generation Sequencing (NGS) germline NGS hot spot testing	2023
Neurofibromatosis and rasopathies	2025
Trio Sequencing – Postnatal (neonatal and paediatric)	2026
Variant validation	2026

VEQ EMQN	
Tipo	**Anno**
Postnatal Constitutional CNV Detection	2018, 2019, 2022
BRCA (familial breast cancer) - full scheme	2012, 2014, 2016, 2017
BWS (Beckwith-Wiedemannsyndrome)	2015, 2016, 2019, 2021
DNA sequencing- Sanger	2012, 2014, 2015, 2016, 2017, 2018, 2019, 2020, 2021, 2023, 2024, 2025
Familial Adenomatous Polyposis (FAP/APC/MAP)	2016, 2018, 2019, 2020
FRAX (Fragile X syndrome) - full scheme	2012, 2014, 2015, 2016, 2017, 2019, 2022
DNA sequencing - NGS (v Germline)	2017, 2018, 2019, 2020, 2021, 2024
Huntington Disease (HD)	2017, 2018, 2019
Lynch sindrome	2018, 2019, 2020, 2021
Ovarian Cancer (v Somatic)	2019, 2020, 2021
PWAS Prader-Willi e Angelman syndromes)	2012, 2014, 2016, 2019, 2020, 2021, 2022
SMA (Atrofia muscolare spinale)	2012, 2014, 2015, 2016, 2017, 2019, 2020
Y-Chromosome microdelections	2012, 2014, 2015, 2016, 2017
Hereditary breast/ovarian cancer (panel testing)	2022
Charcot-Marie-Tooth disease / Hereditary Neuropathy with liability for pressure palsies	2023

VEQ REGIONE LOMBARDIA	
Tipo	**Anno**
Citogenetica prenatale costituzionale	2019, 2021, 2022, 2023, 2024, 2025, 2026
Citogenetica postnatale	2019, 2020, 2022,2023, 2024, 2025, 2026
Citogenetica oncoematologia	2020, 2021,2022, 2023, 2024, 2025, 2026
A-CGH post-natale	2018, 2021, 2022, 2023, 2024, 2025, 2026
BRCA1-BRCA2 germinale	2019, 2021, 2022, 2023, 2024, 2025, 2026
BRCA1-BRCA2 (somatico)	2024, 2025, 2026
CFTR (fibrosi cistica)	2019, 2021, 2023, 2024, 2026
FRAXA (Sindrome X-fragile)	2022, 2023, 2025, 2026
HFE (emocromatosi)	2020, 2022, 2025, ?2026
Fattore V di Leiden	2021, 2022, 2023, 2024, ?2026
Fattore II	2022, 2023, 2024, ?2026

VEQ CEQAS - GENie	
Tipo	**Anno**
BRCA1, BRCA2 and HRR gene variant classification assessment	2025
HGVS nomenclature Basic	2025
ISCN Neoplasia Karyotyping Basic	2025
ISCN Constitutional Karyotyping Basic	2025
ISCN Constitutional FISH Basic	2025

UK NEQAS	
Tipo	**Anno**
UK NEQAS BCR::ABL1 Major Quantification	2026

	In corso

Data 15/01/2026