**REFERINȚE SELECTATE**

Următoarele baze de date pot fi utile specialiștilor pentru a afla detalii despre frecvența, structura și asocierea variațiilor genetice cu anumite afecțiuni, precum și despre alternativele de tratament existente:

**dbSNP** – bază de date conținând localizarea, frecvența și structura variațiilor genetice localizate ( <https://www.ncbi.nlm.nih.gov/projects/SNP/> ).

**ClinVar** – bază de date care asociază variațiile genetice patogenice cu detalii asupra gradului de patogenicitate și cu dovezile științifice curente (<https://www.ncbi.nlm.nih.gov/clinvar/> ).

**PharmGKB** – bază de date care indică interacțiile dintre variații genetice și medicamente (<https://www.pharmgkb.org/> ).

**GeneCards** – oferă informații despre rolul genelor și a proteinelor codificate (<http://www.genecards.org/> ).

**Specific Genetic Disorders** – oferă informații de diagnostic și tratament pentru boli rare cu etiologie genetică (<https://www.genome.gov/10001204/> ).

**Susan G Komen Foundation** – oferă informații despre rolul mutațiilor BRCA1/BRCA2 în patogeneza cancerelor la femei și bărbați (<http://ww5.komen.org/BreastCancer/InheritedGeneticMutations.html> ).

*Lista de referințe de mai jos reprezintă o selecție a studiilor și informațiilor utilizate pentru alcătuirea acestui raport. Această listă nu reprezintă întregul set de informații care au fost folosite pentru generarea acestui raport, ci doar în măsura în care pot fi utile specialiștilor și la care dumneavoastră puteți apela.*

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