I have a rare, chronic blood disease called PNH

It stands for paroxysmal nocturnal hemoglobinuria. It's rare: it affects literally 1 in a million.

It means my body can't deliver enough oxygen to

- It's caused by a random change in my genes: no one gave it to me, and I can't give it to someone else.
 - my organs. This can make me incredibly tired.
 - I can also get headaches and stomach aches, have trouble breathing and worry about blood clots.
- These symptoms can be hard to see but are very real for me. To learn more, visit ThisisPNH.com.

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