

Deafblindness in French Canadians from Quebec: A predominant founder mutation in the *USH1C* gene provides the first genetic link with the Acadian population

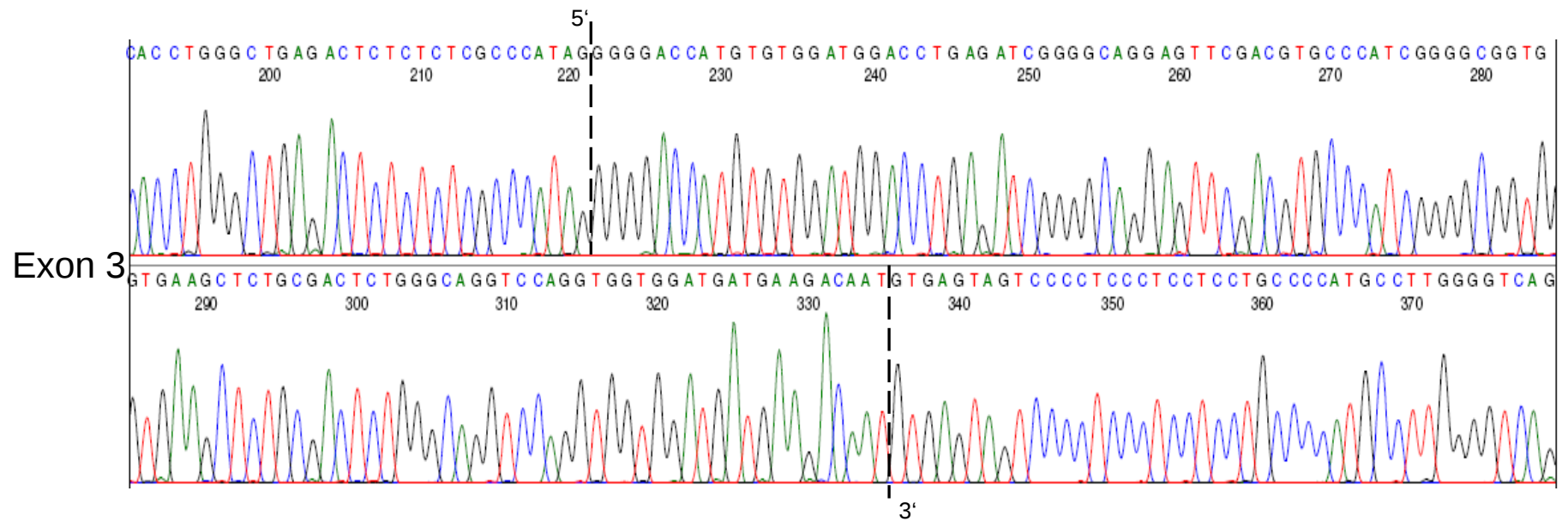
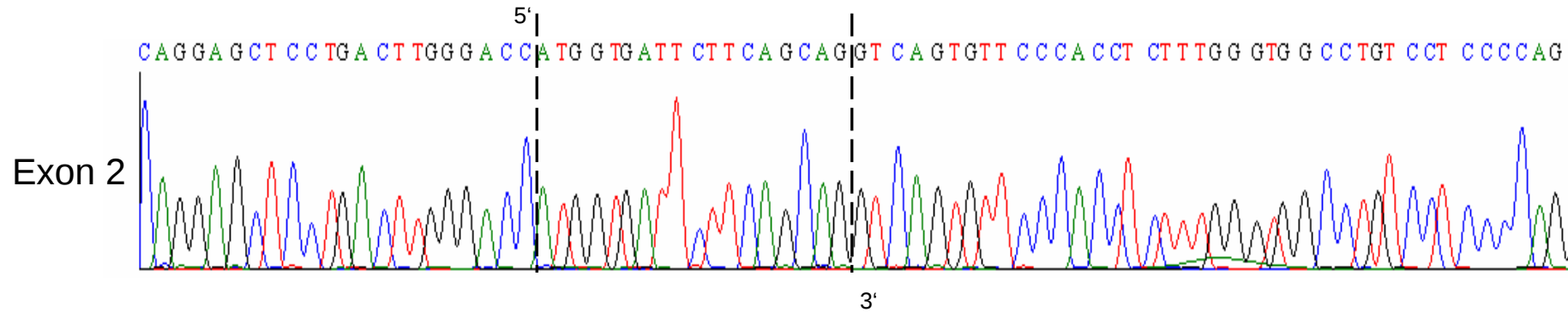
Ebermann et al.

RAW DATA

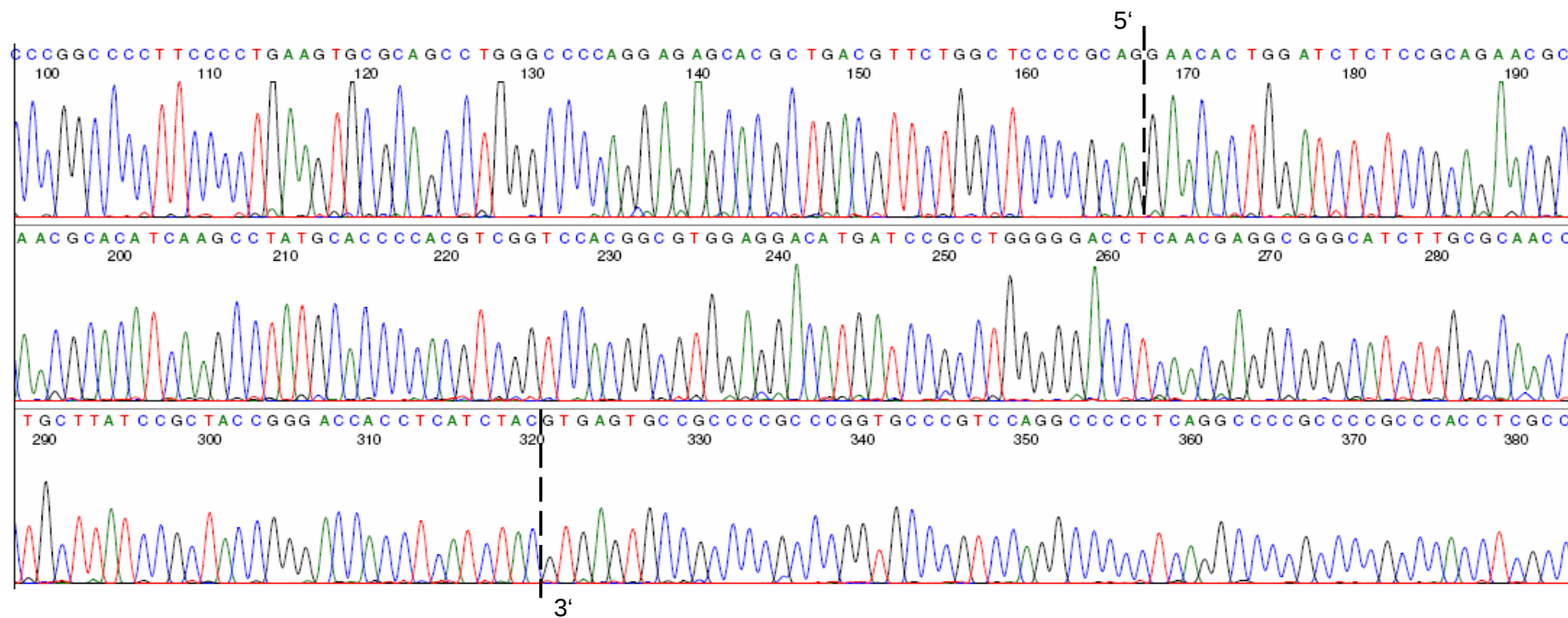
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Mutation screening in USH1 genes in patient 1881.

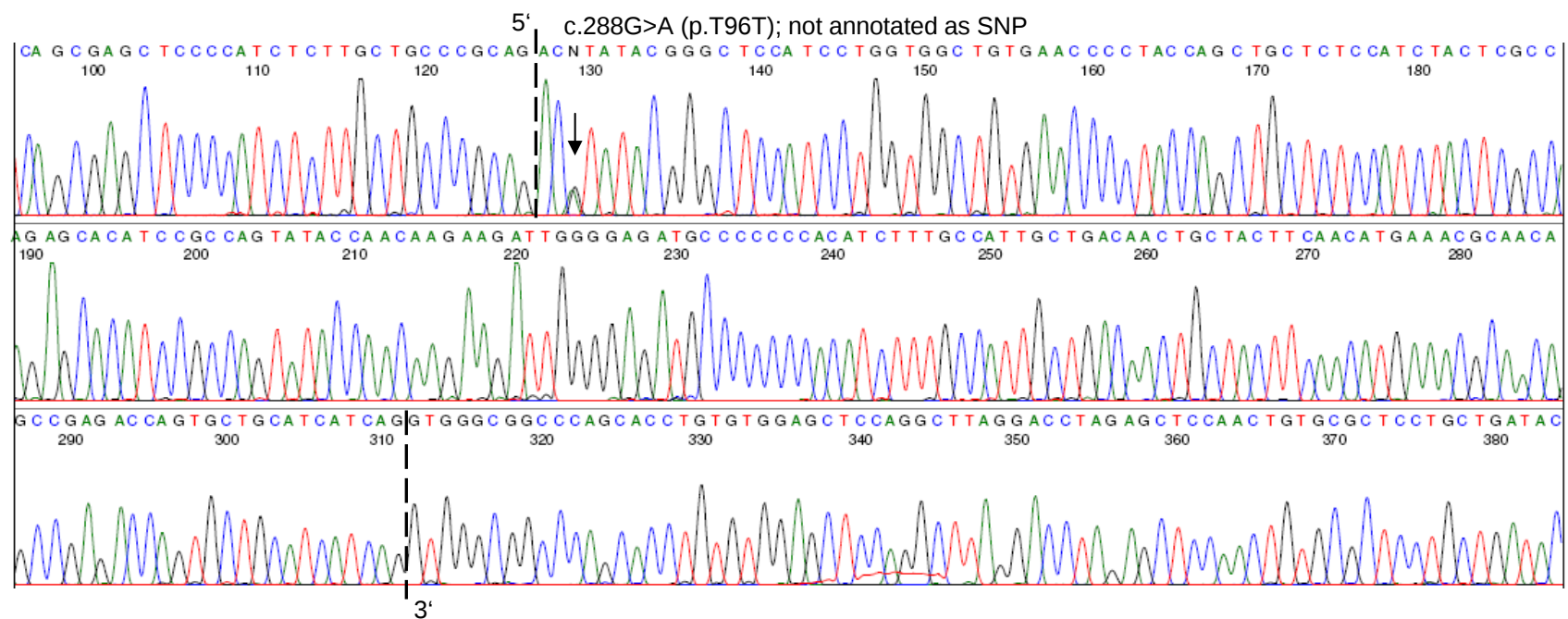
Patient 1881, MYO7A (USH1B)



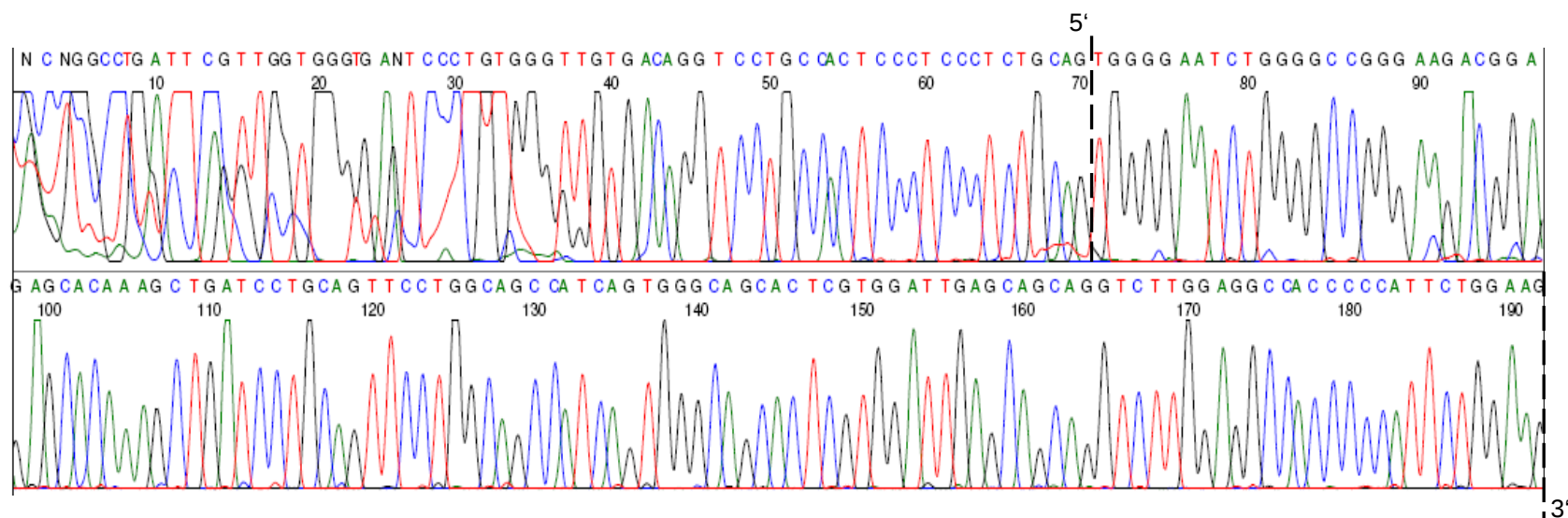
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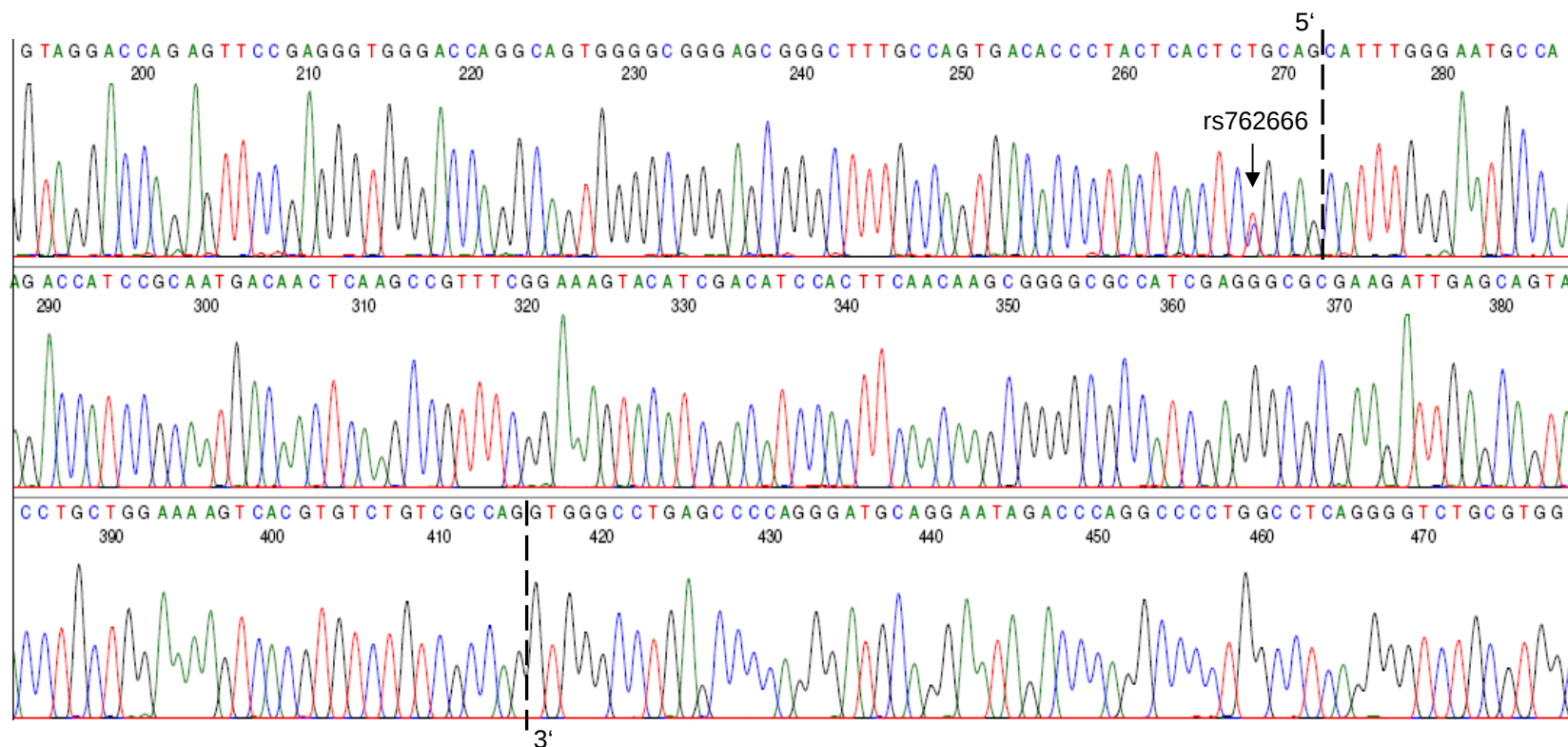
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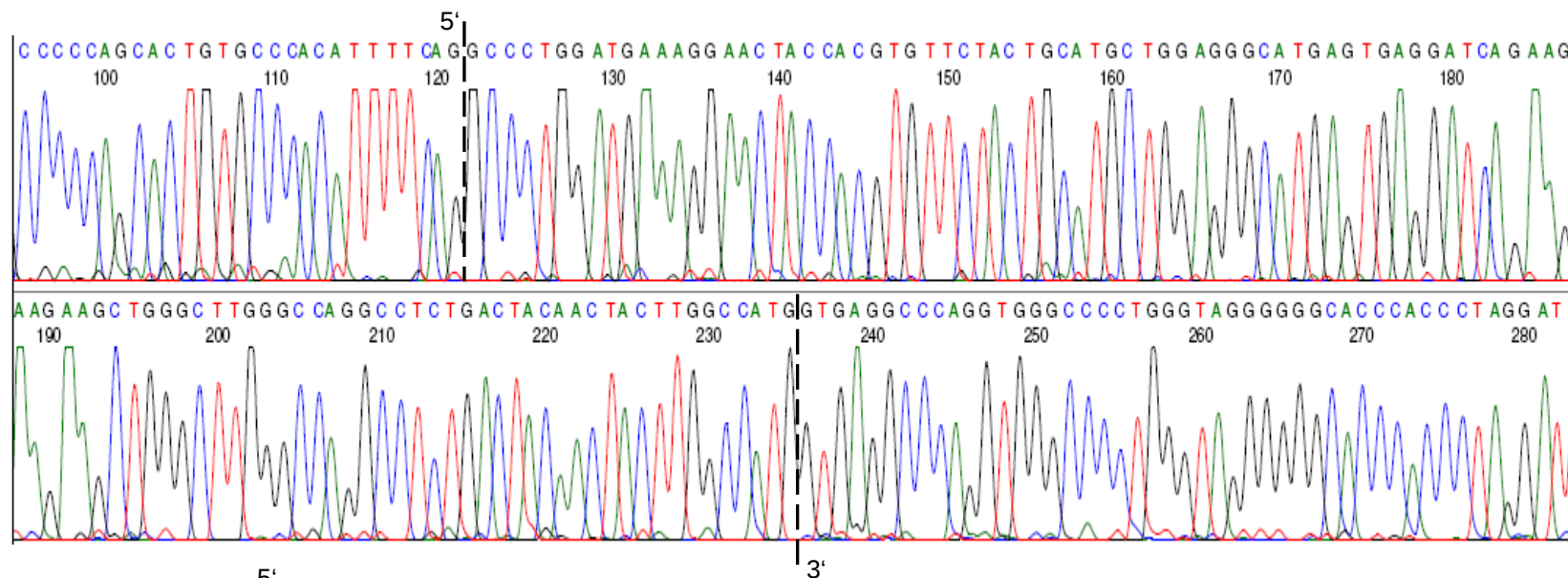
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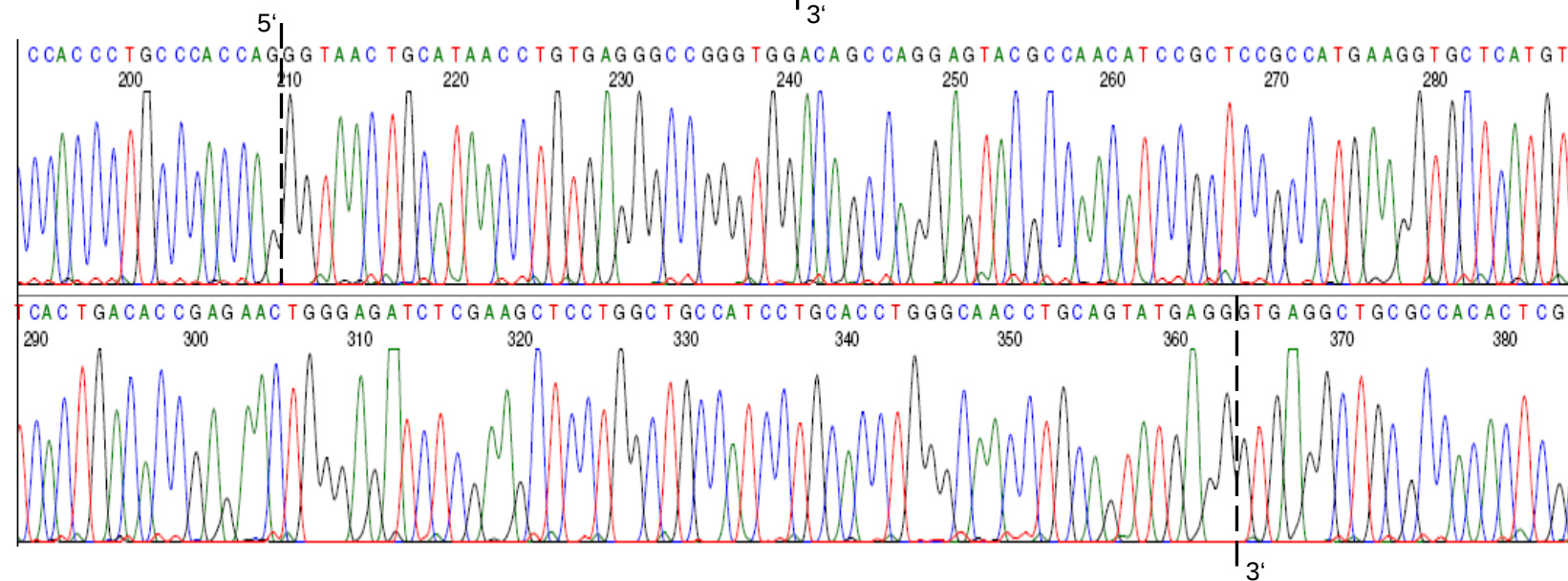
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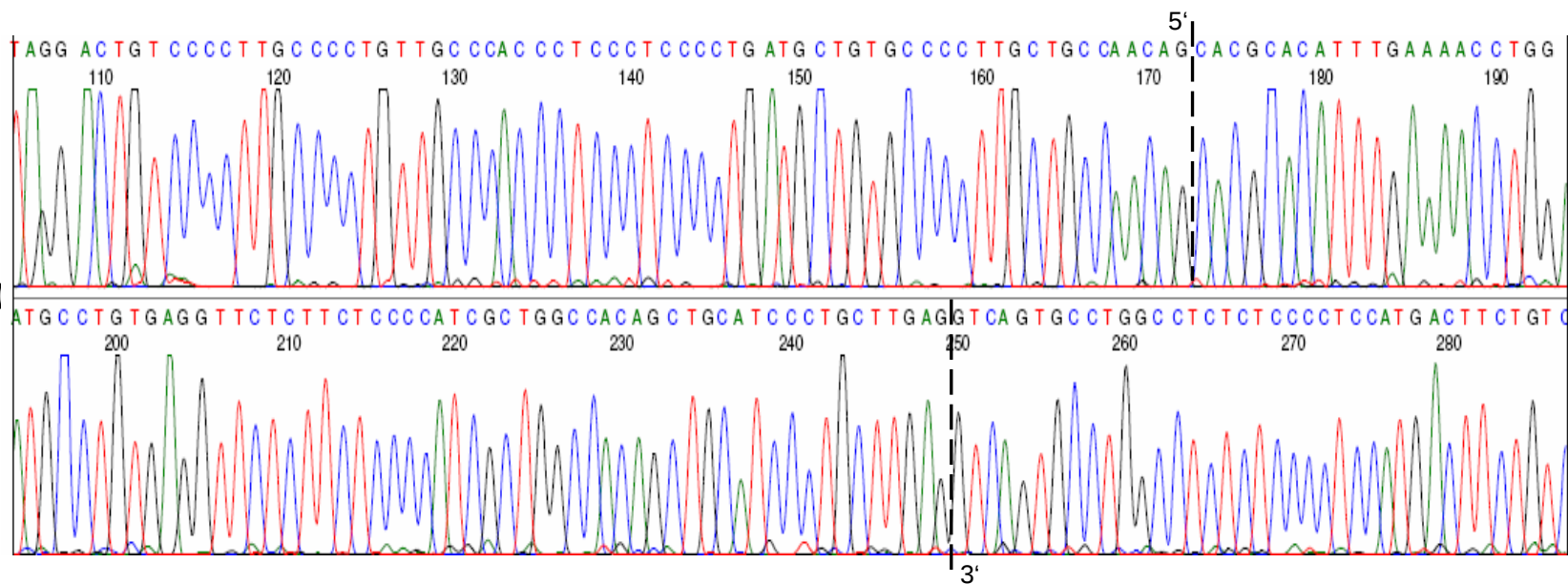
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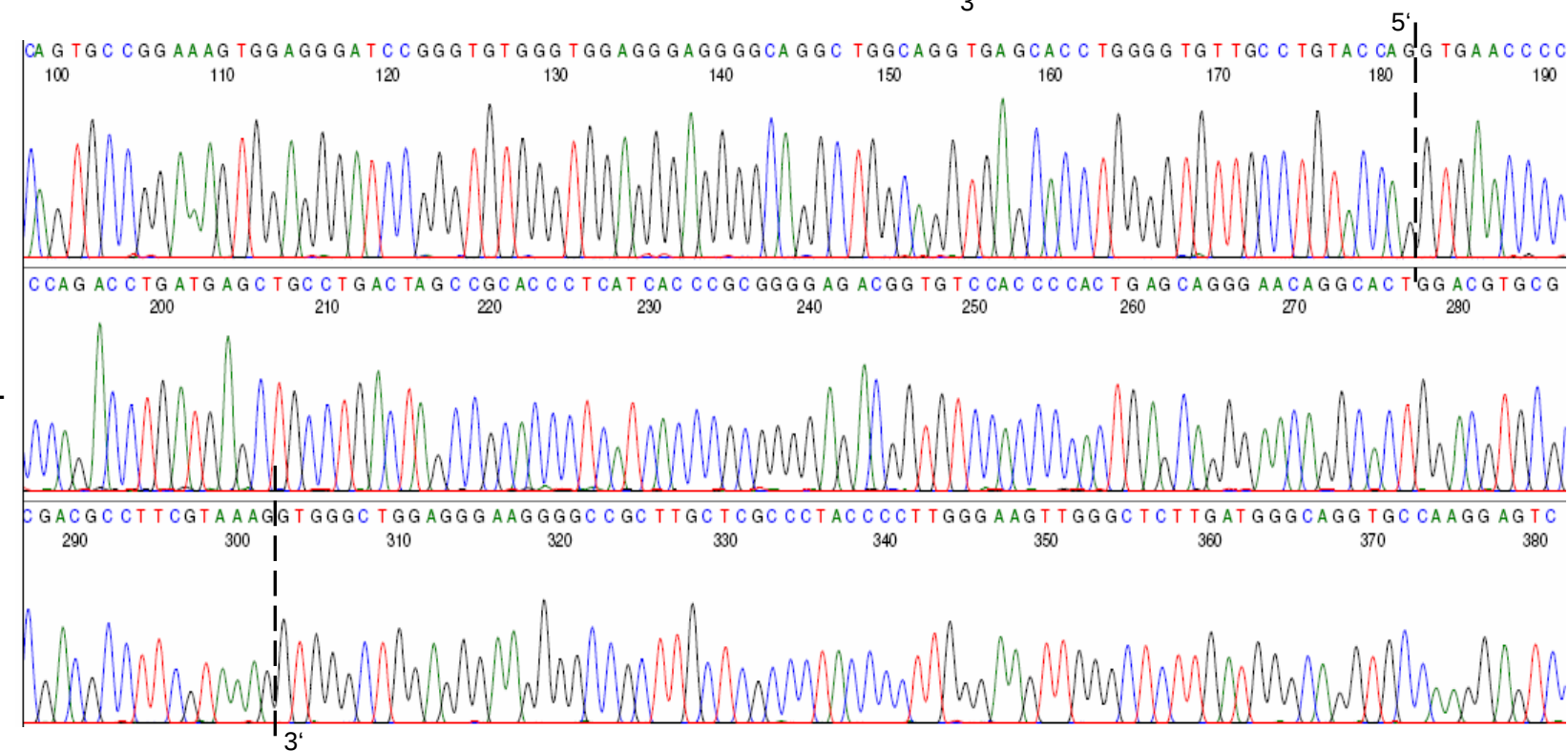
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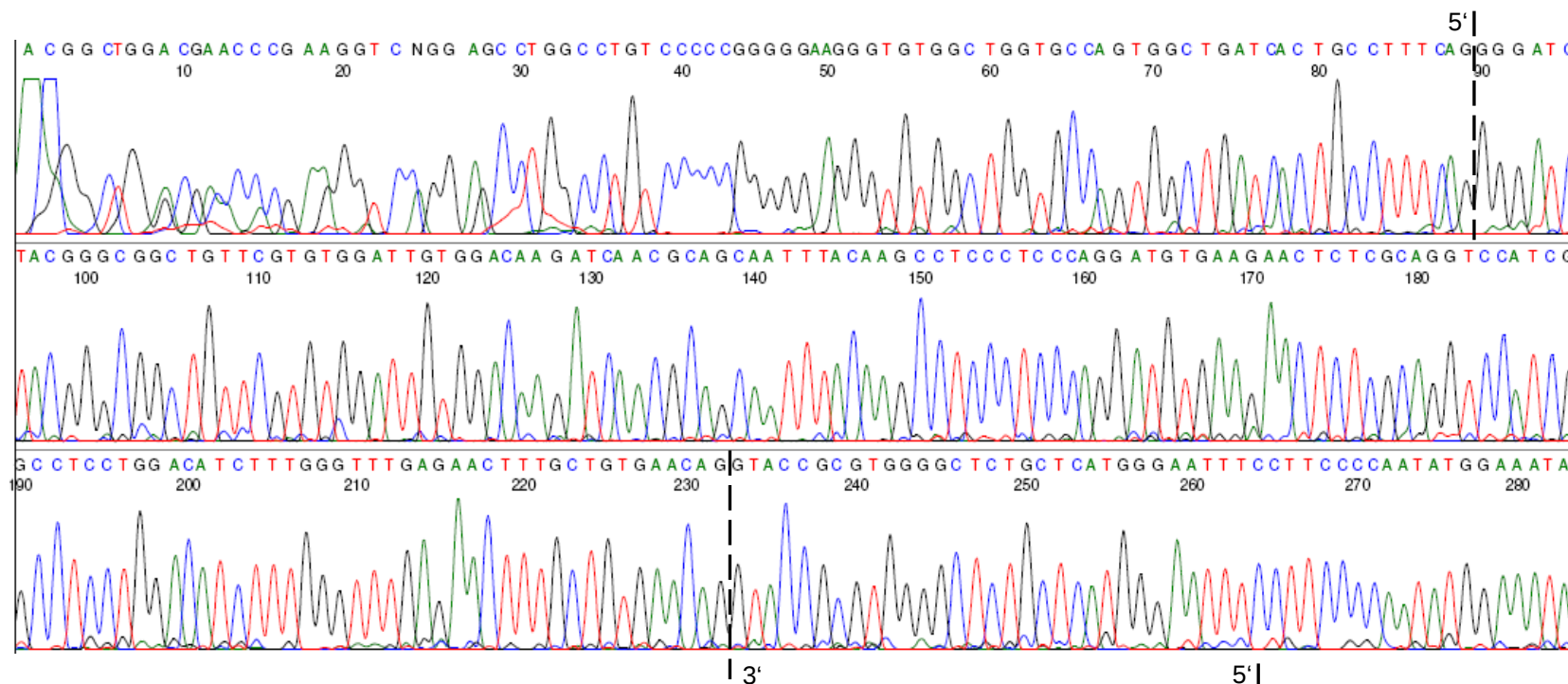
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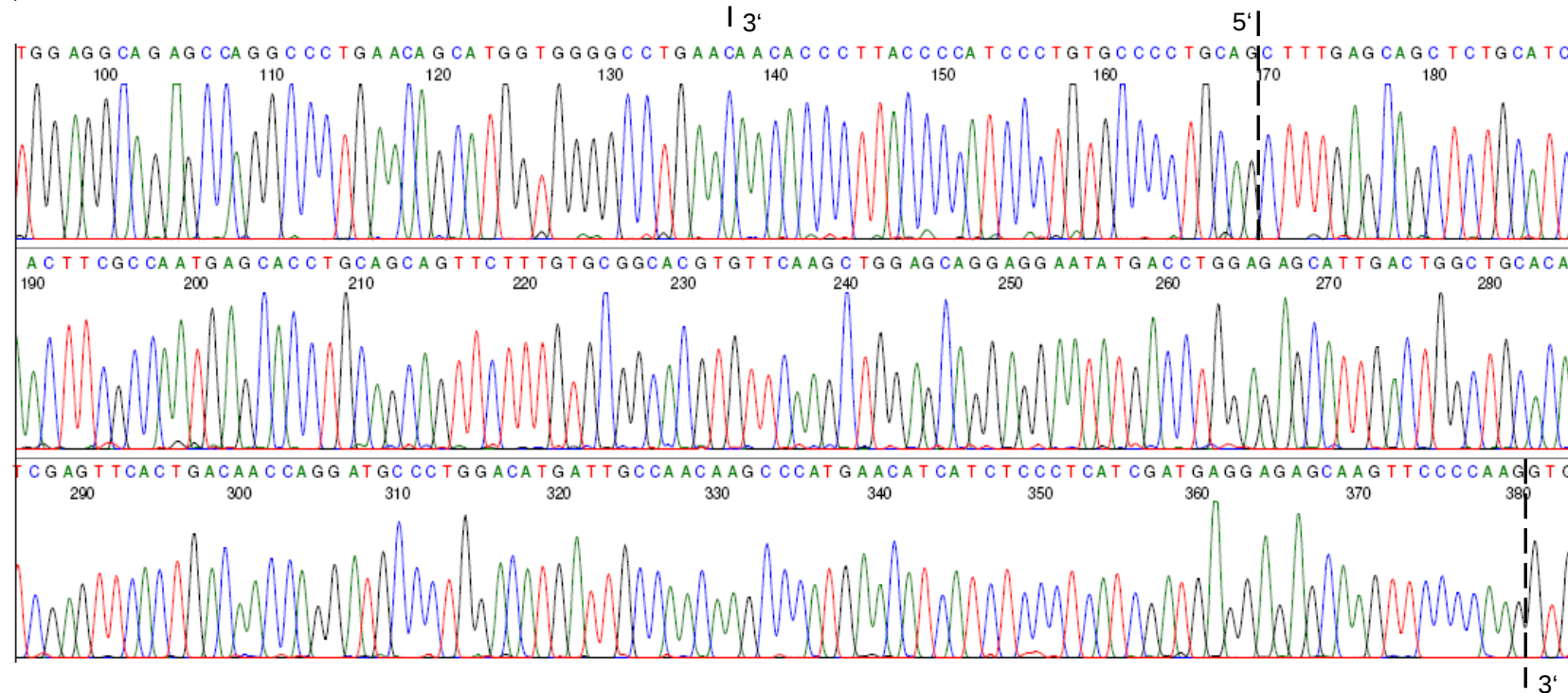
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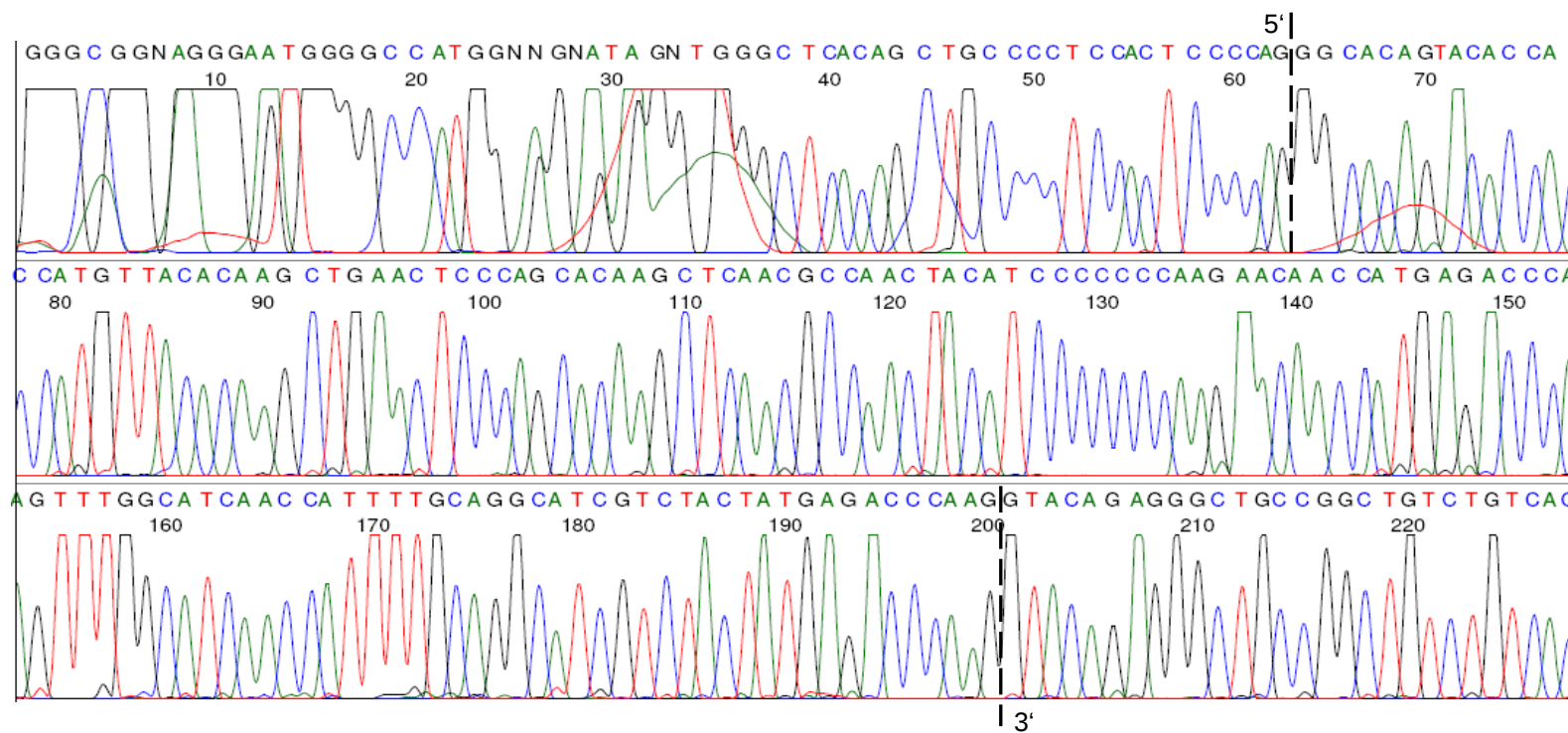
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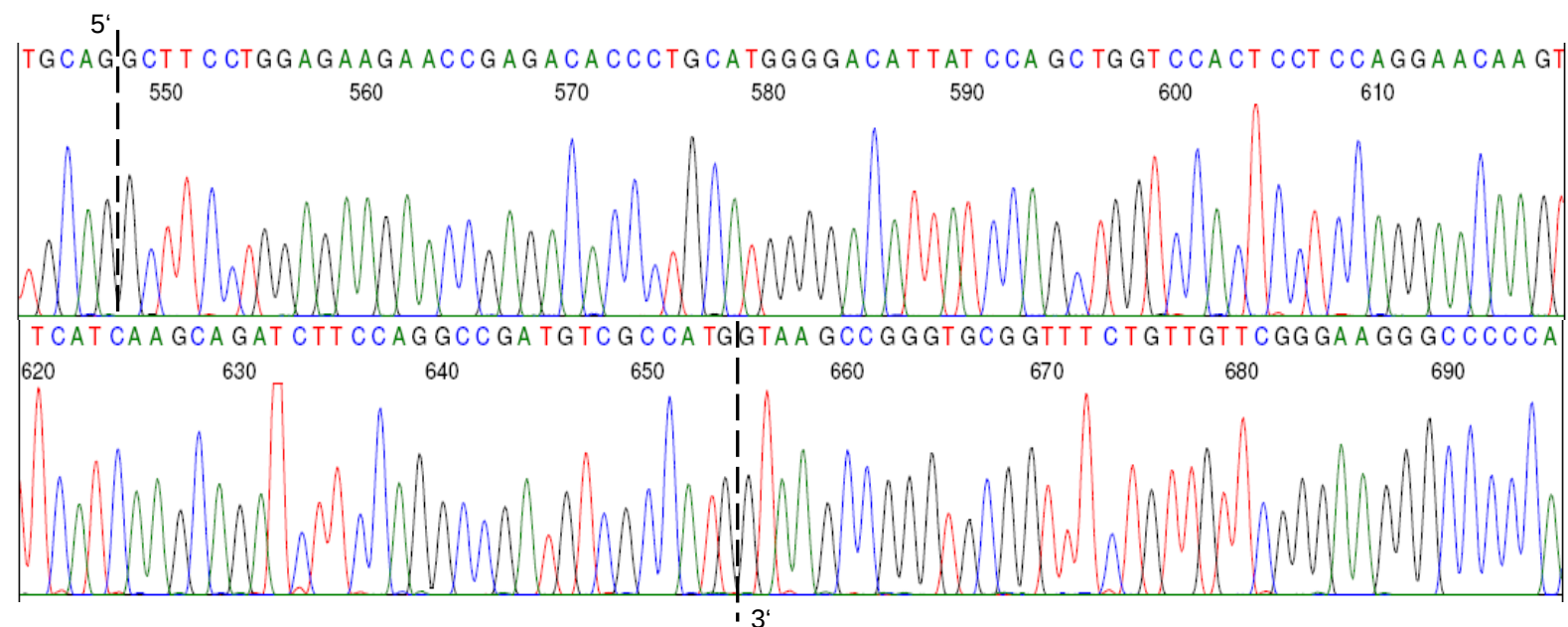
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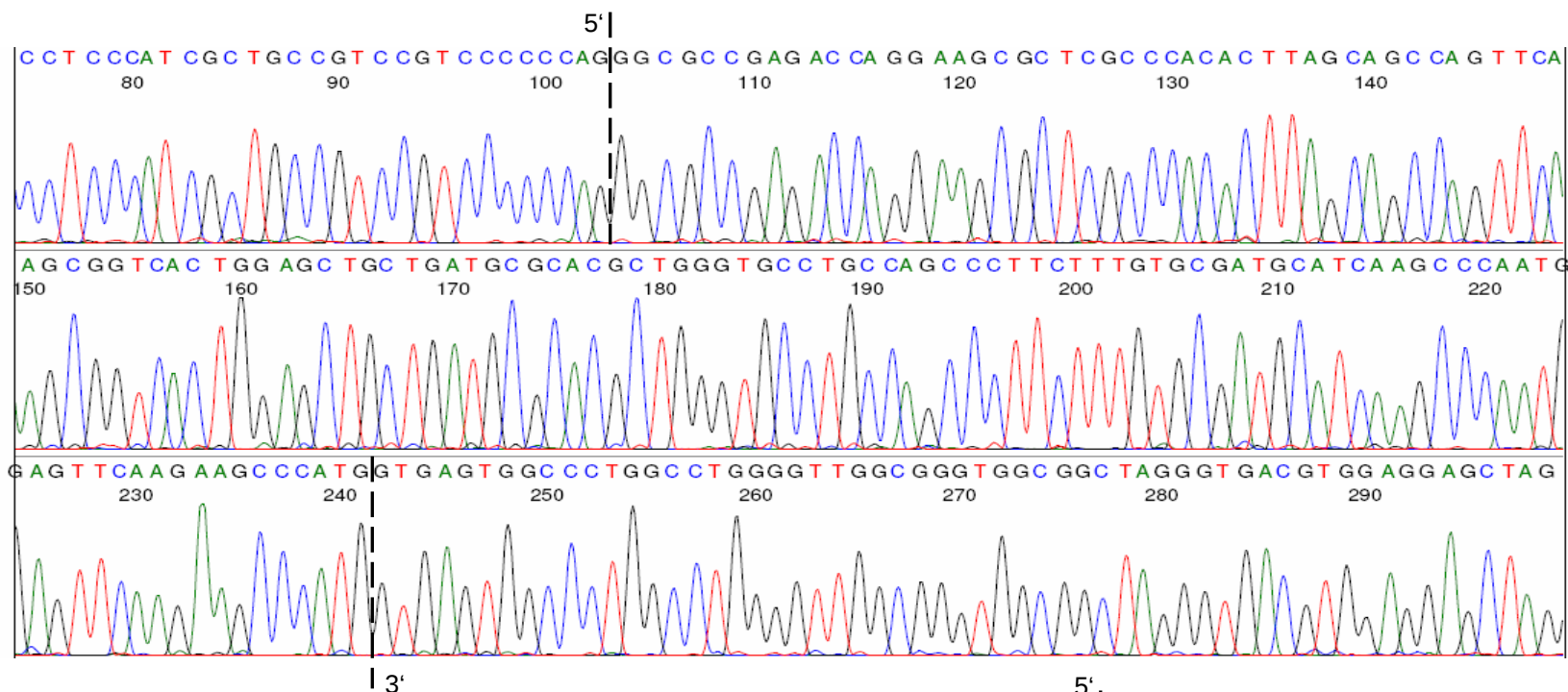
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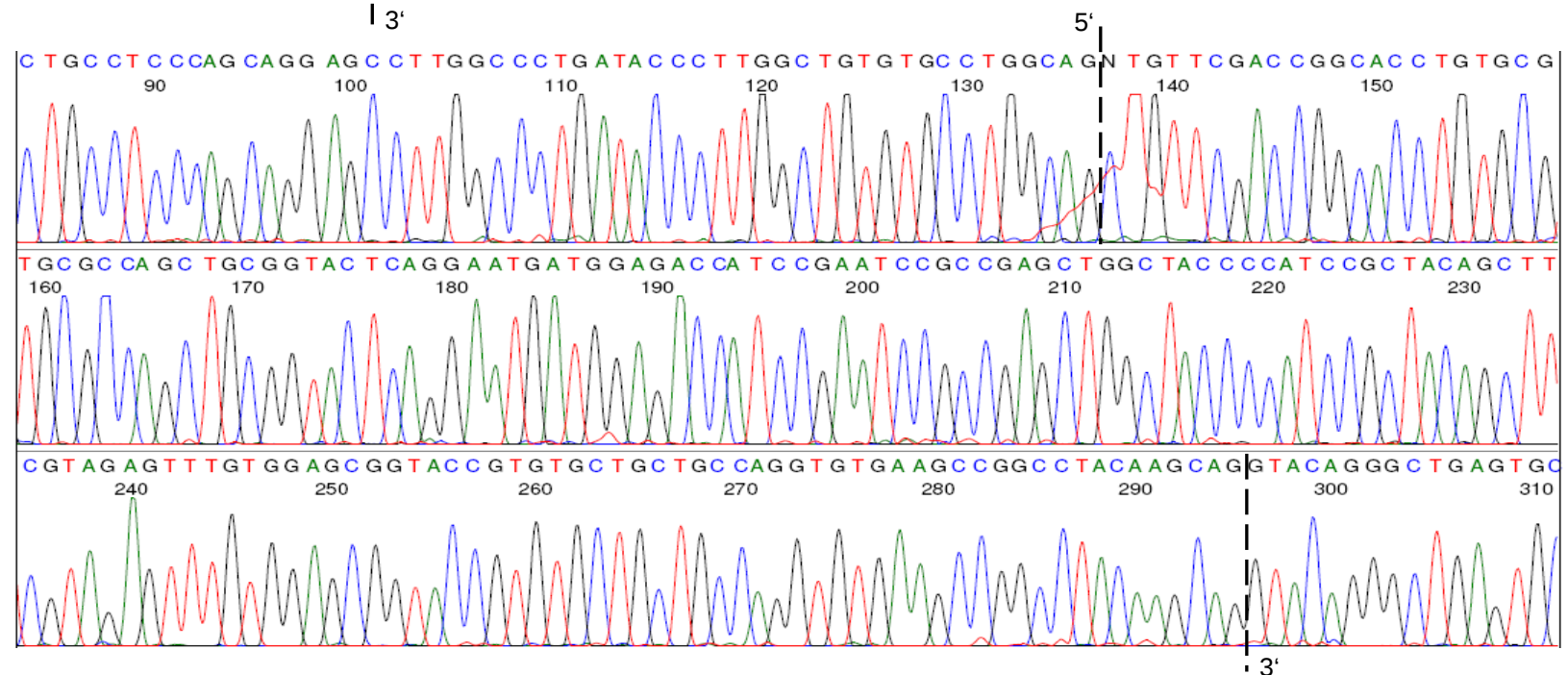
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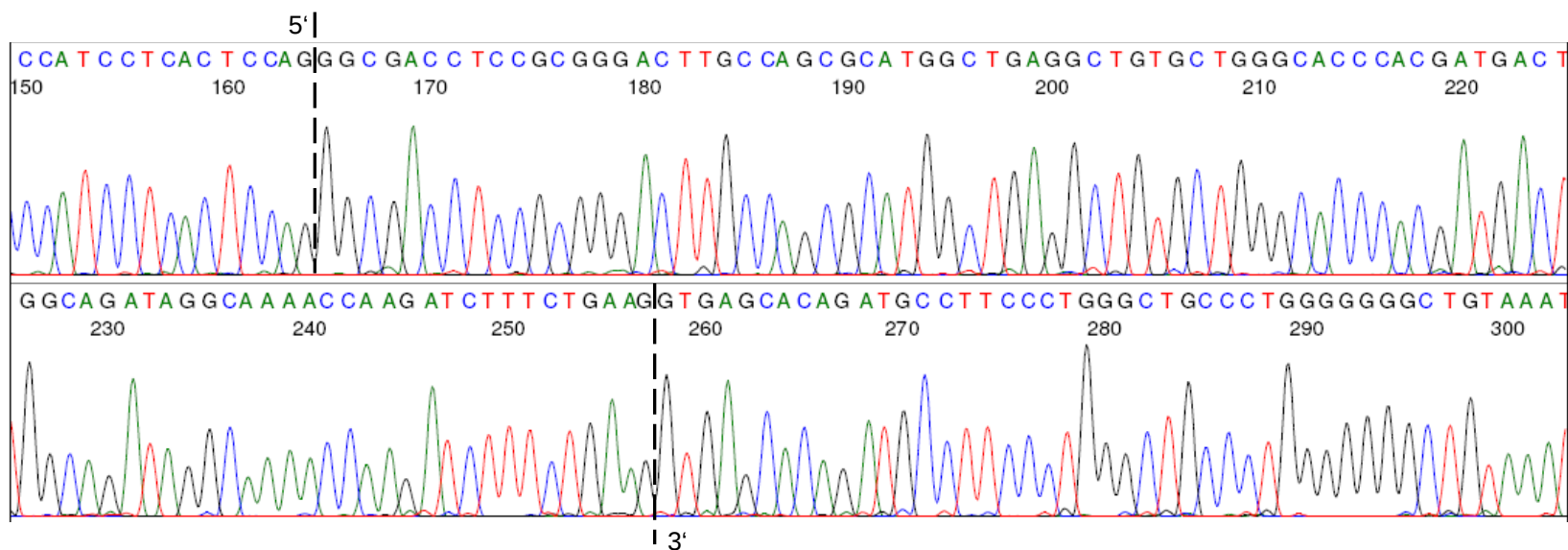
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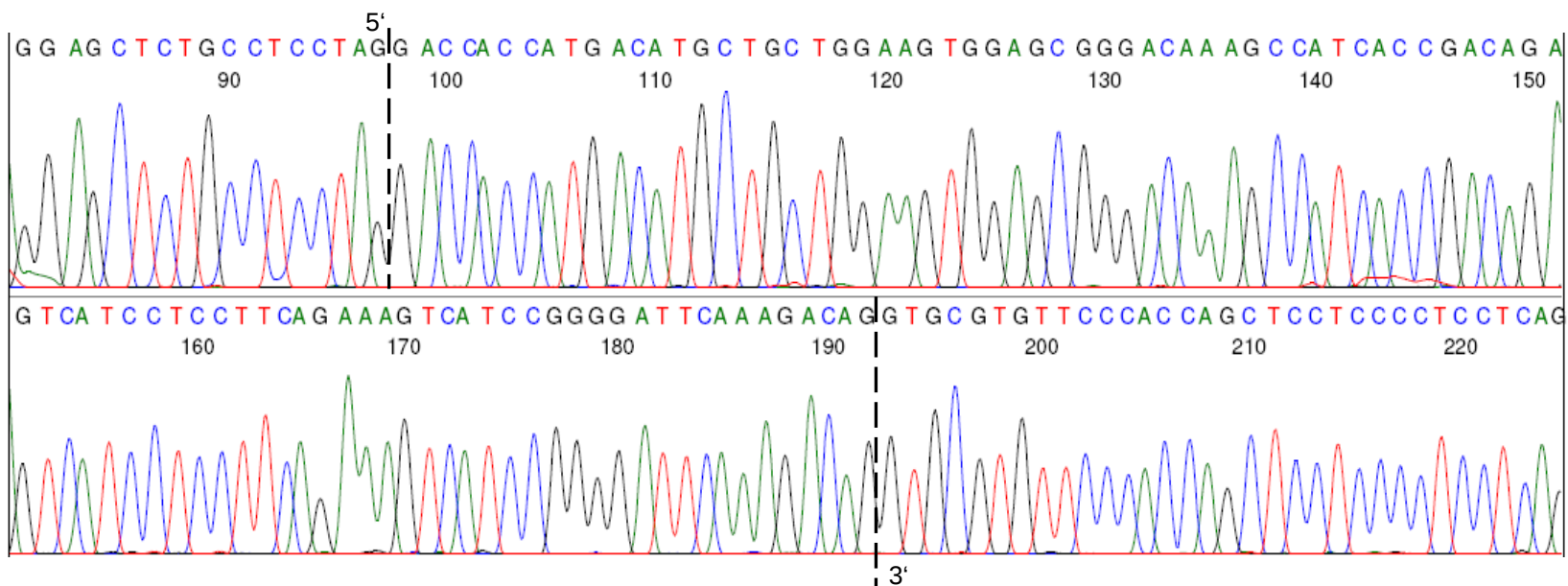
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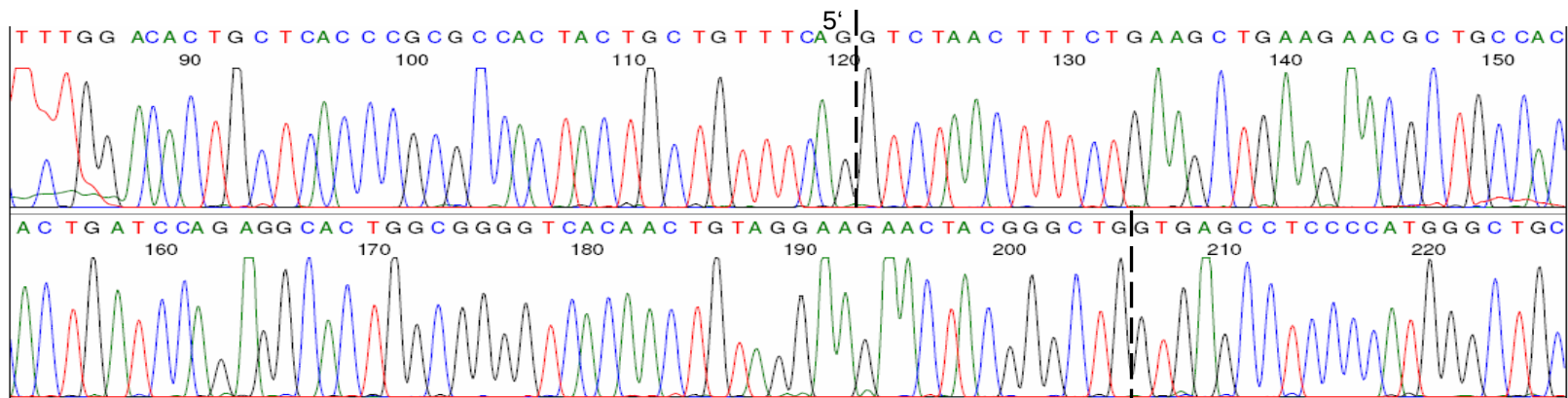
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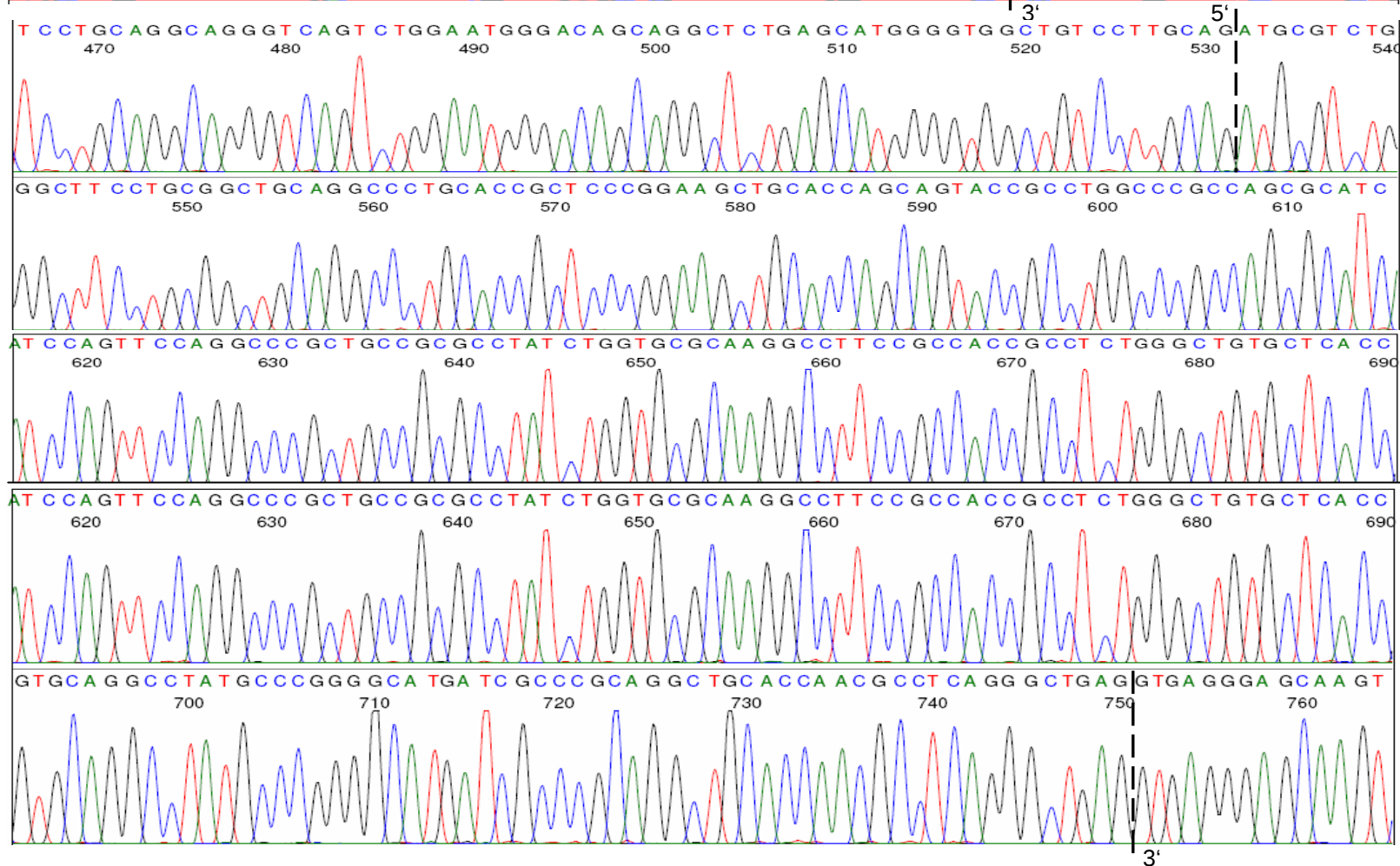
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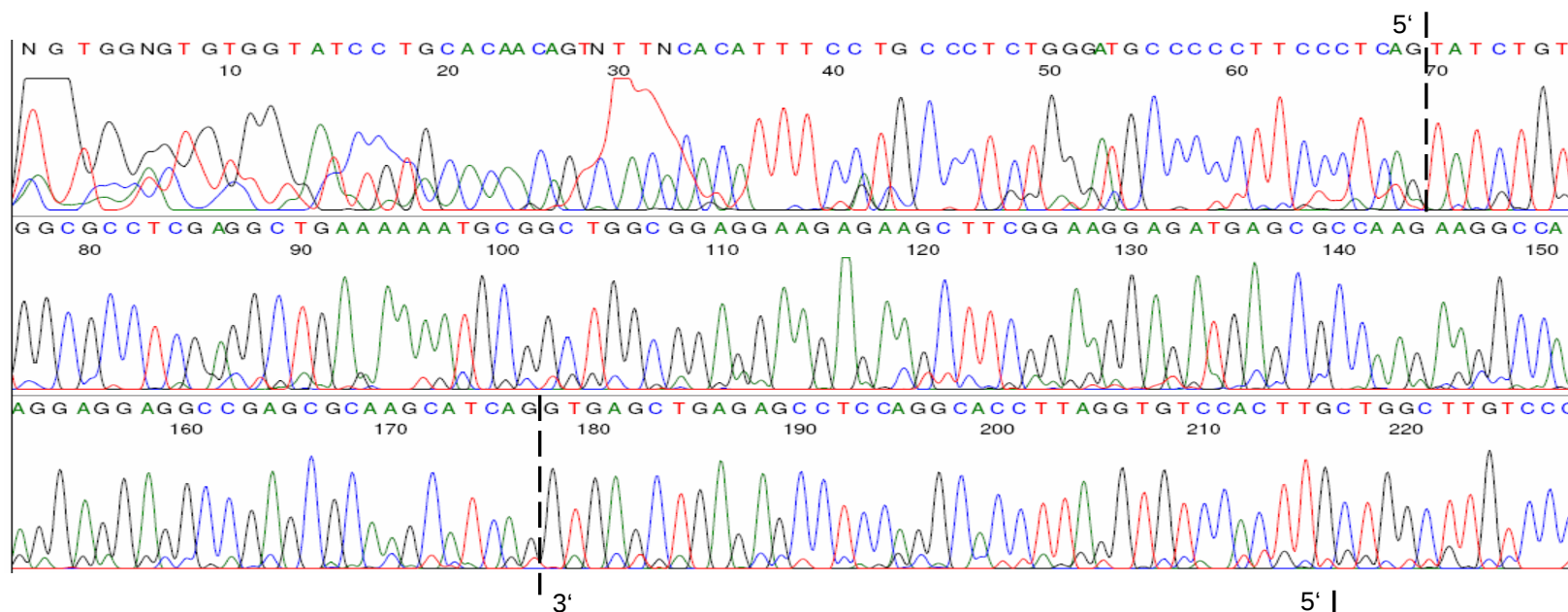
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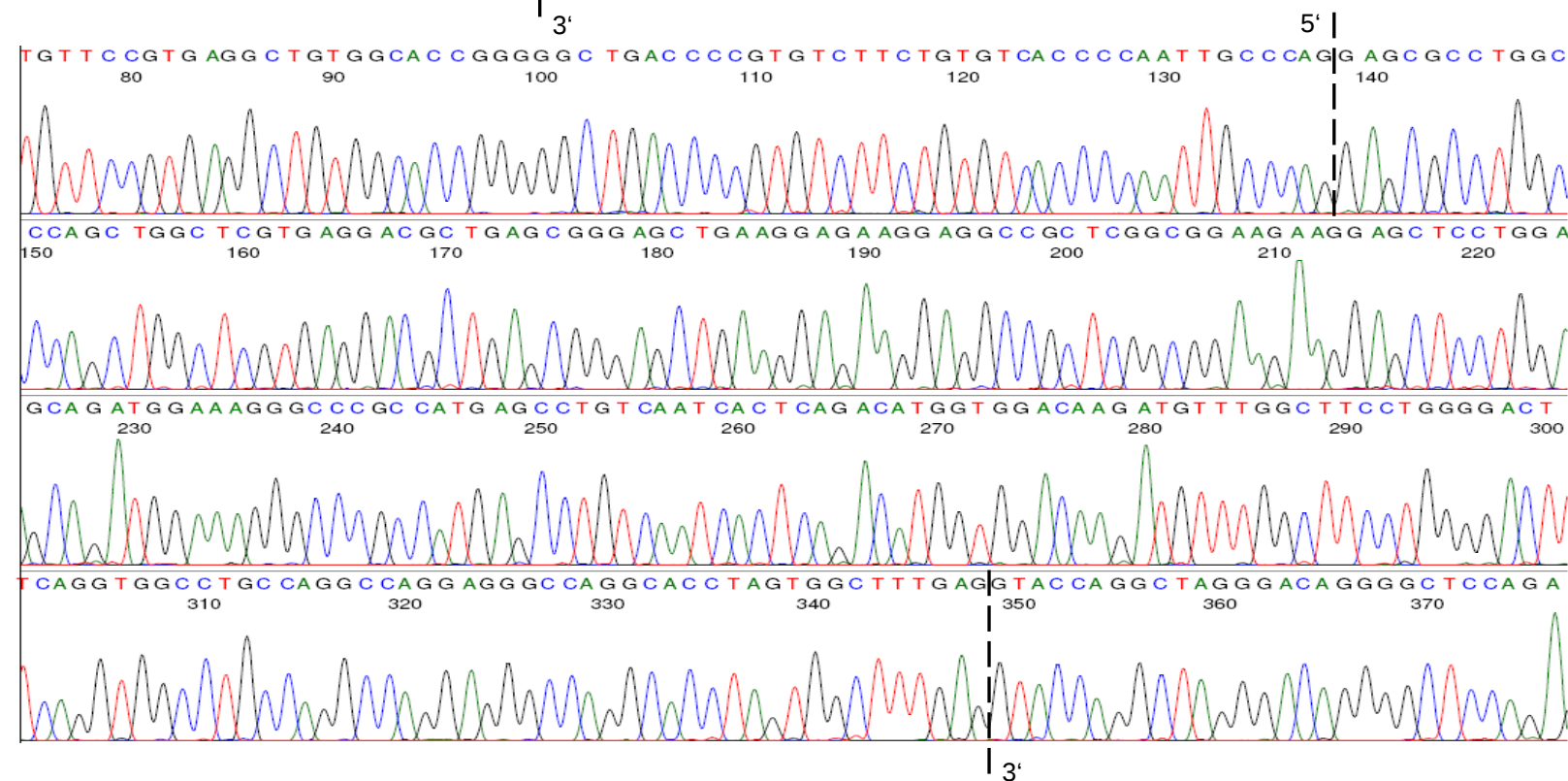
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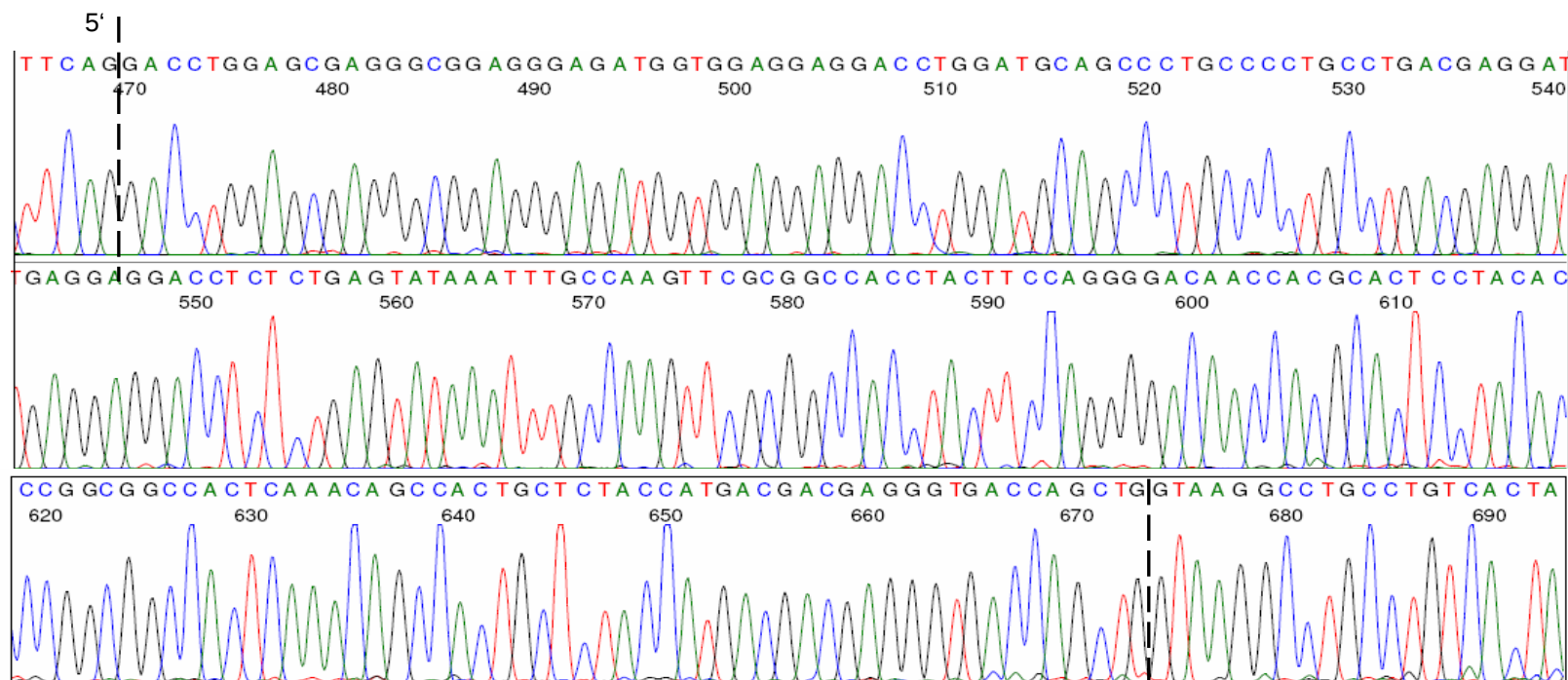
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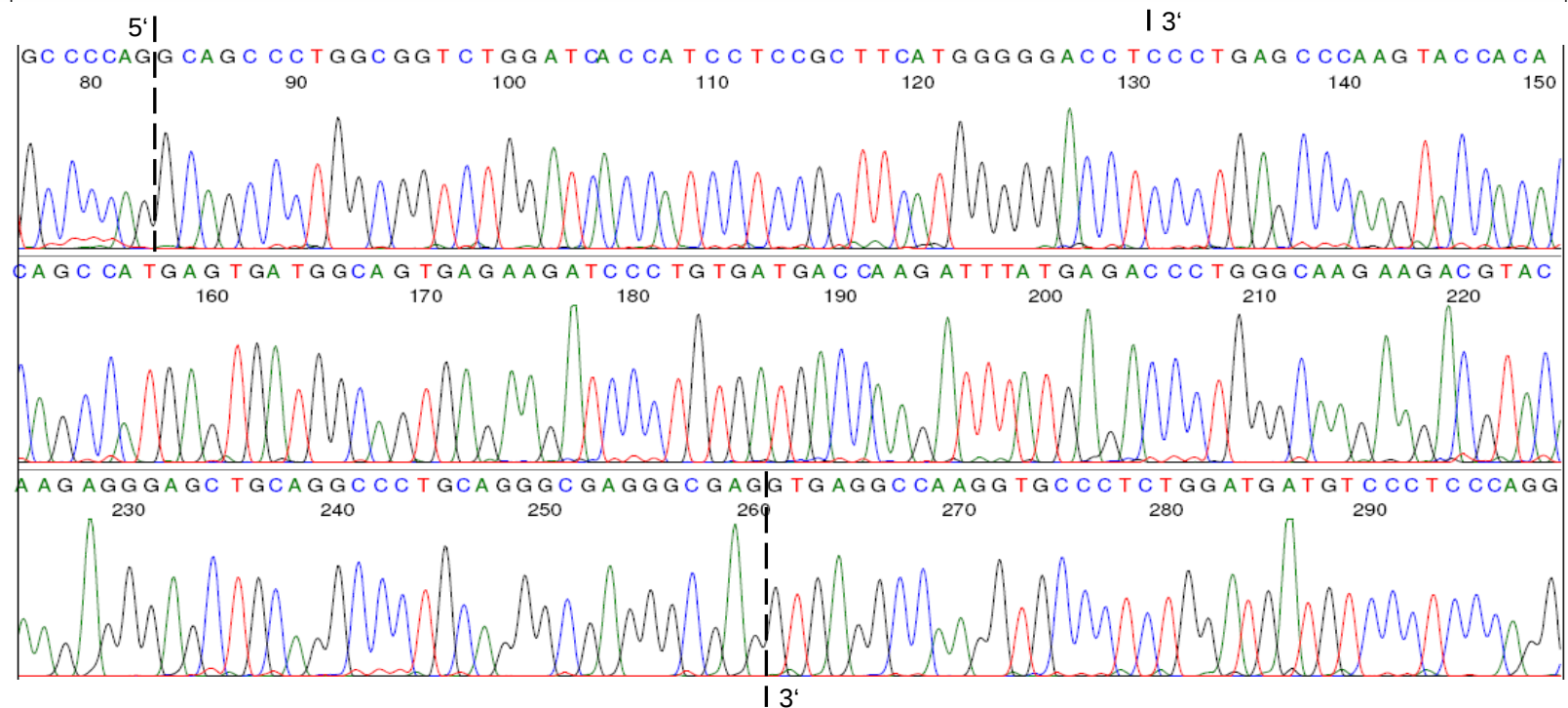
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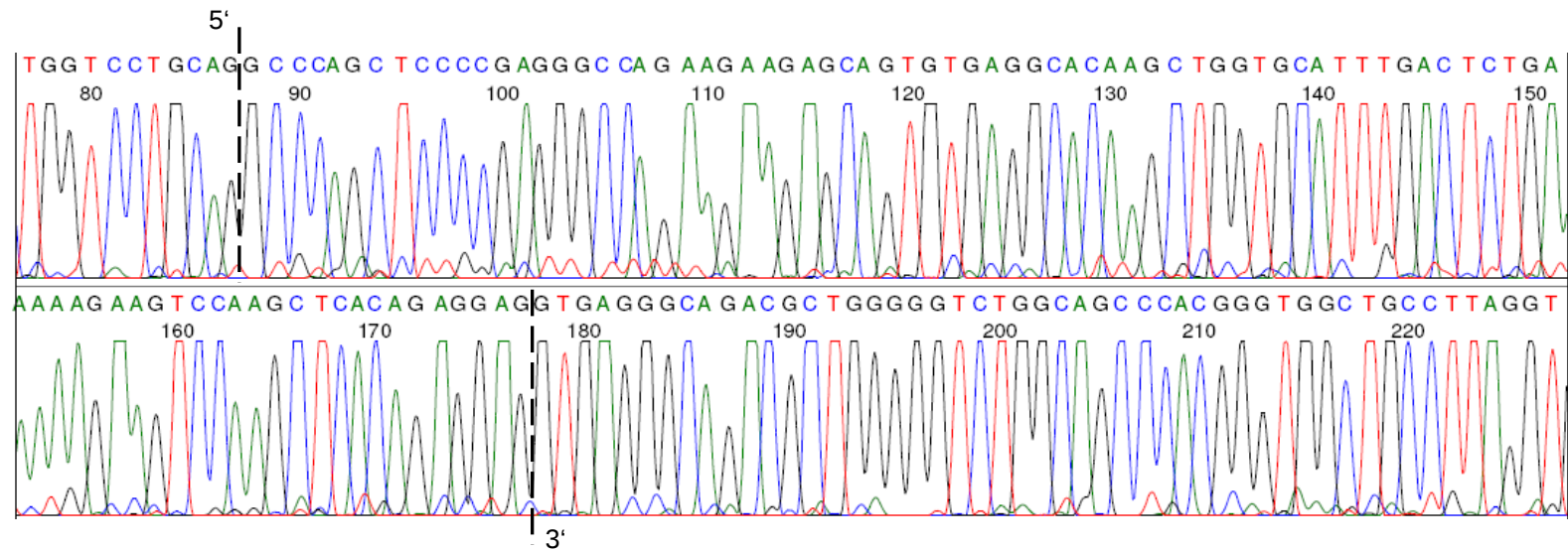
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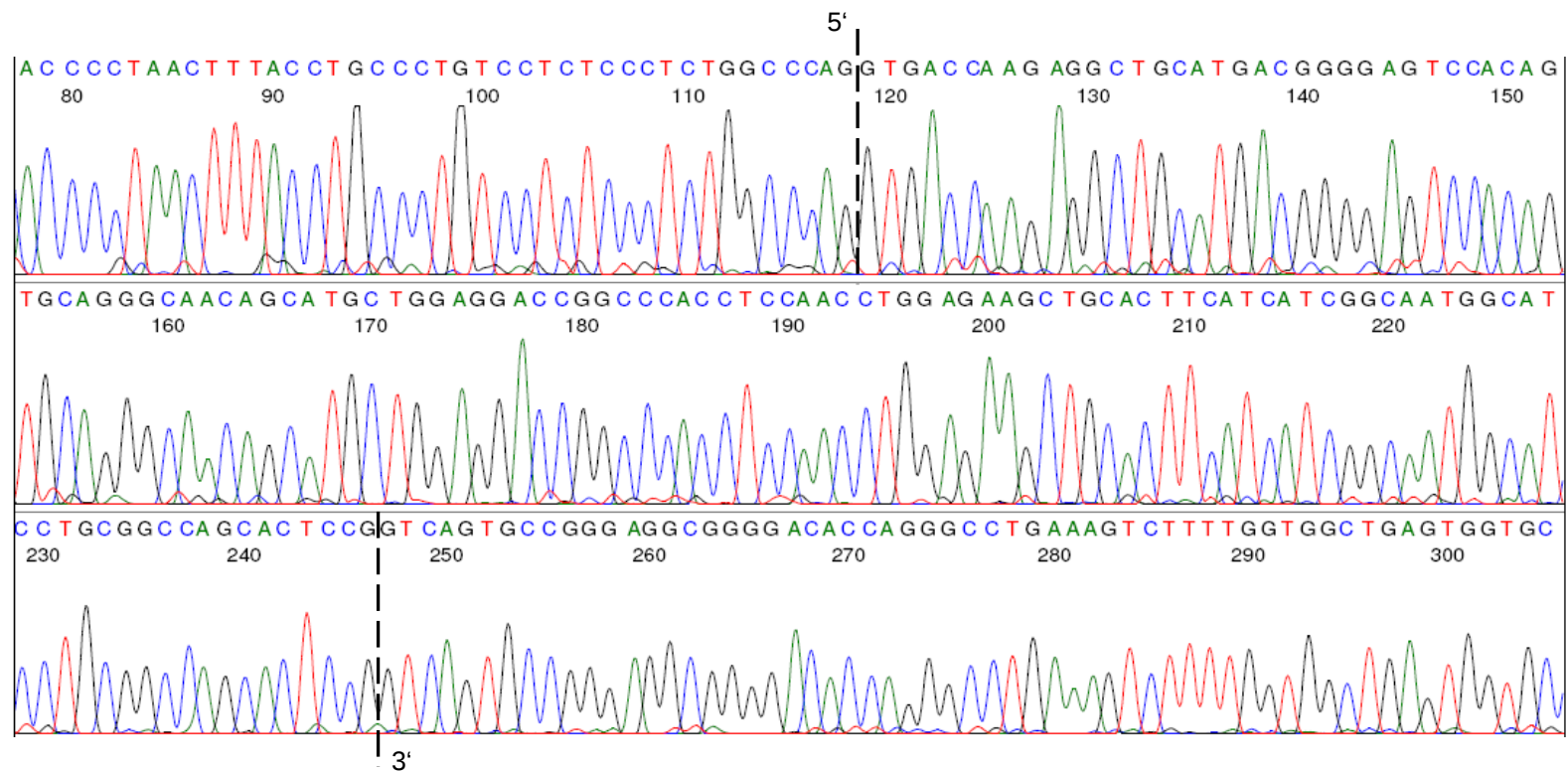
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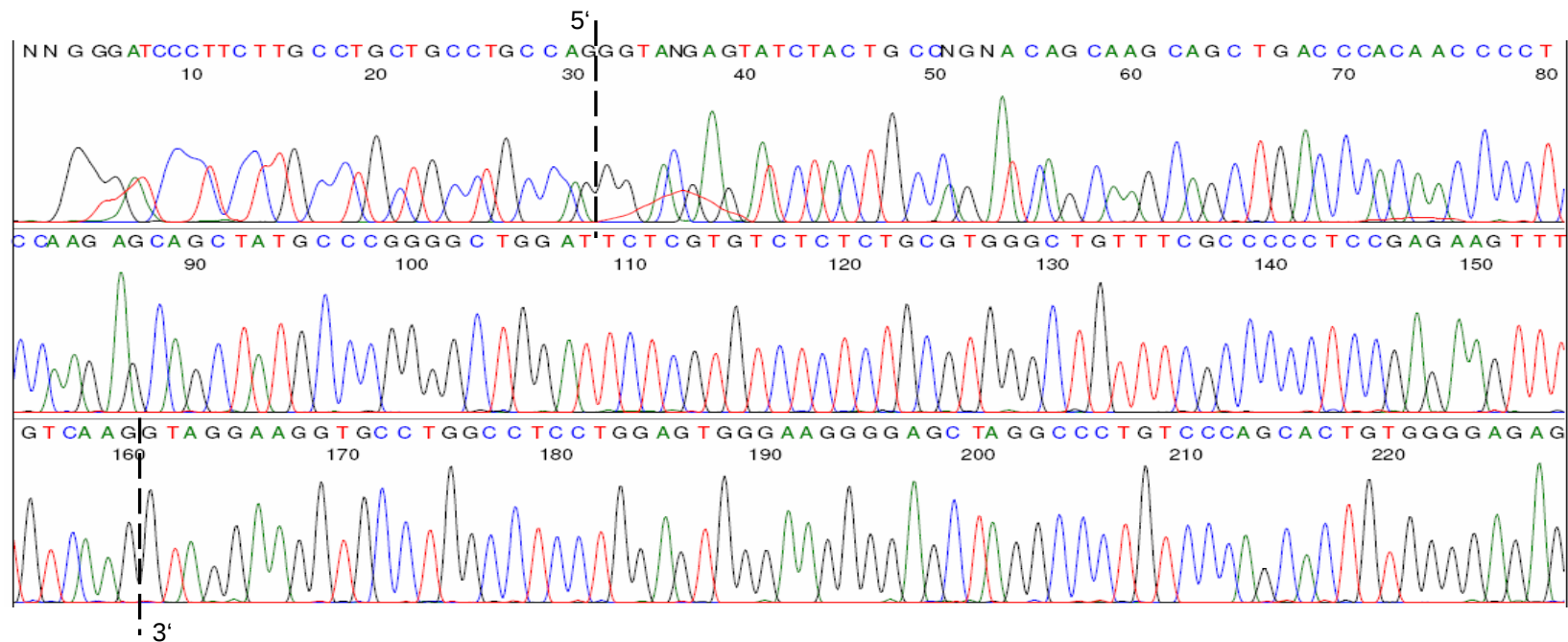
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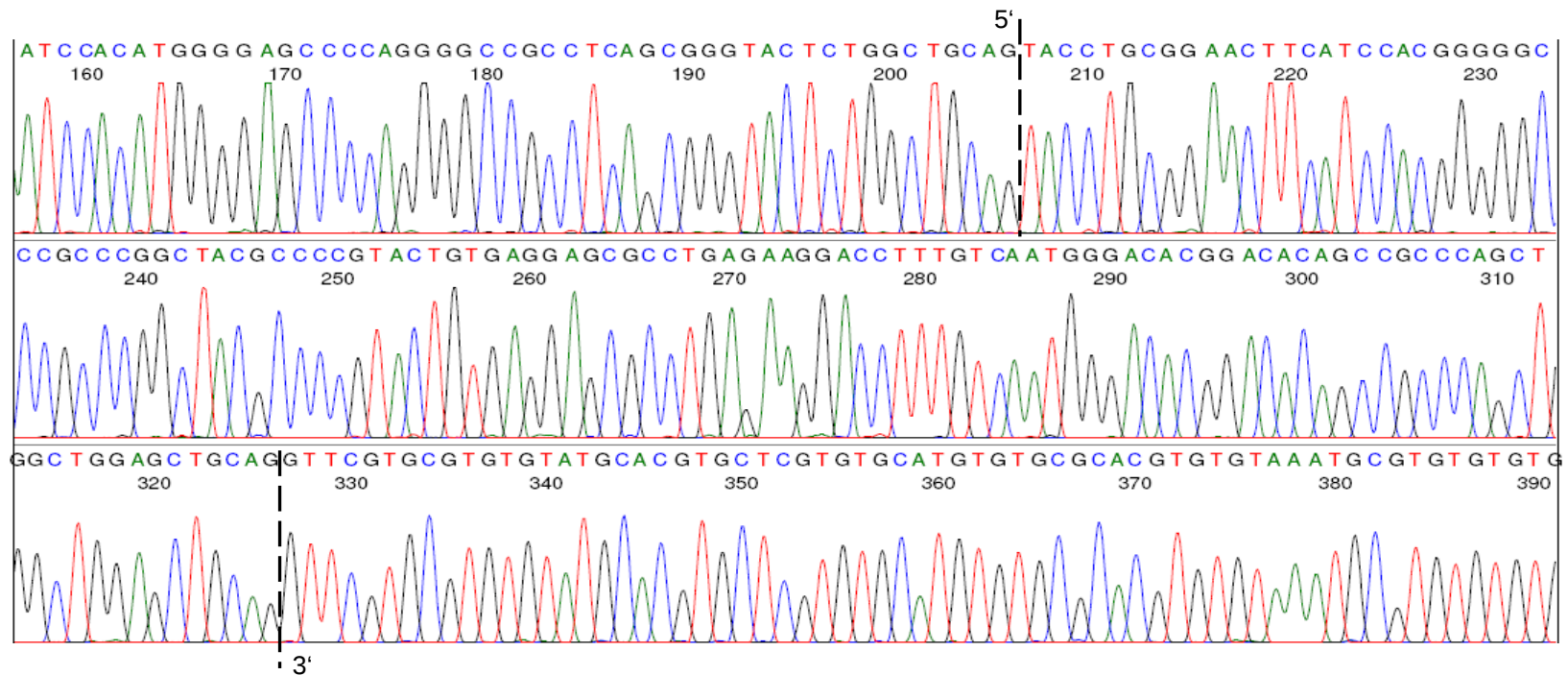
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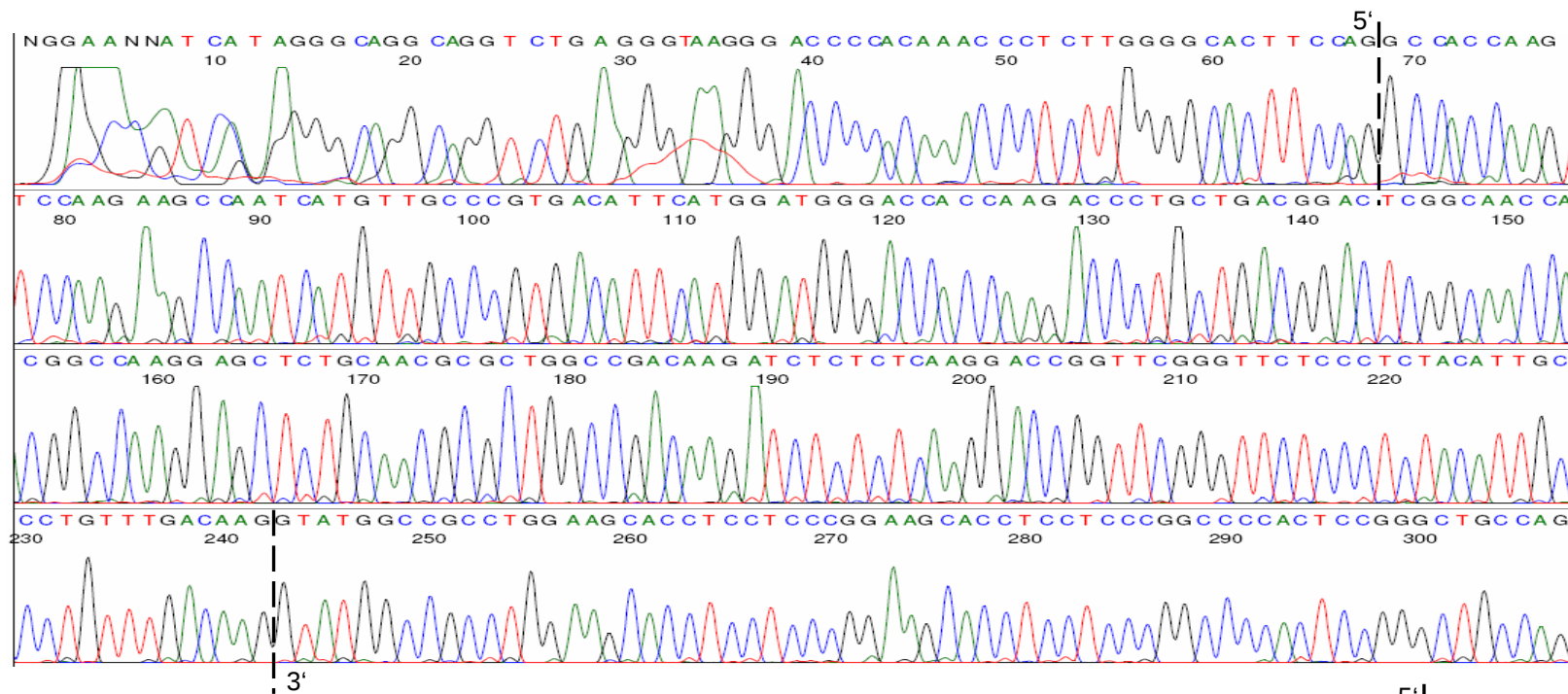
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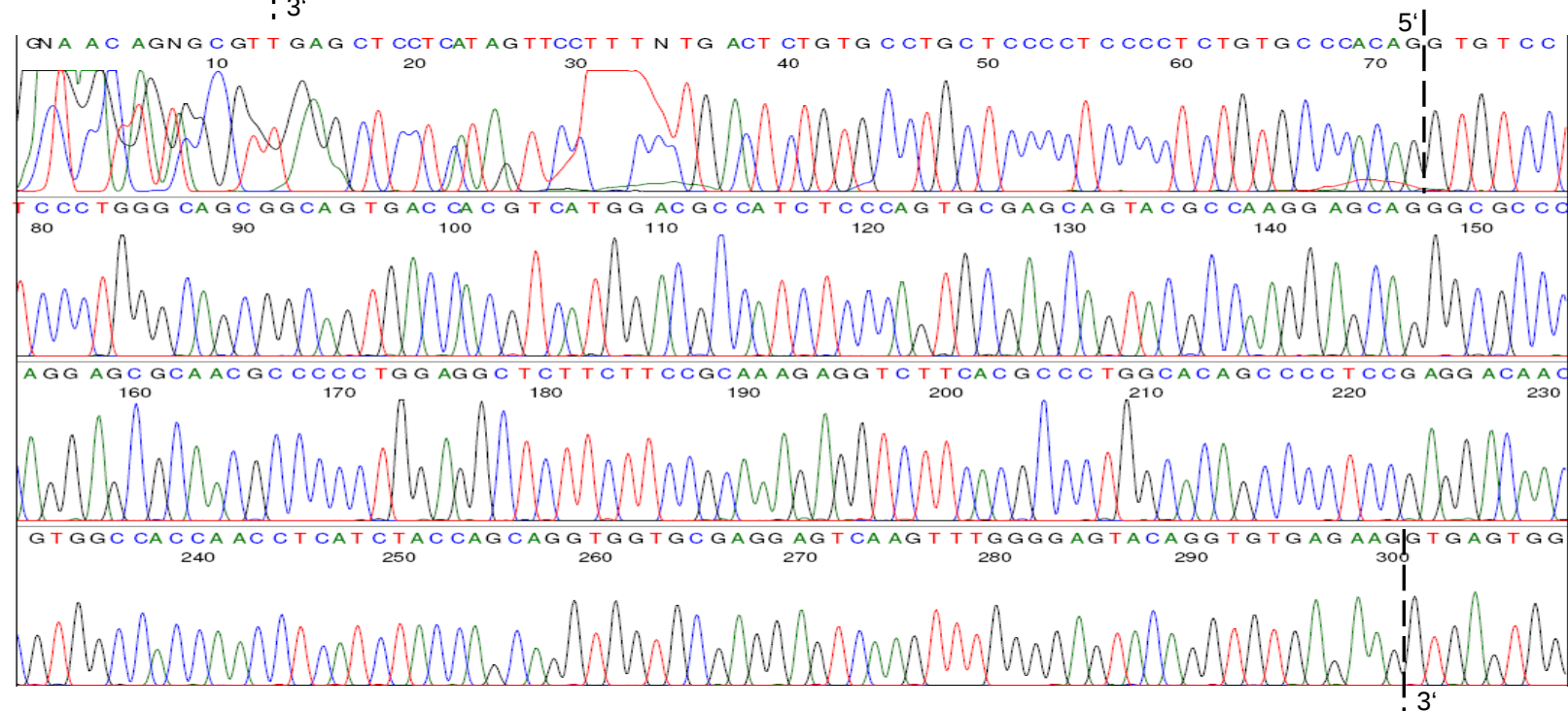
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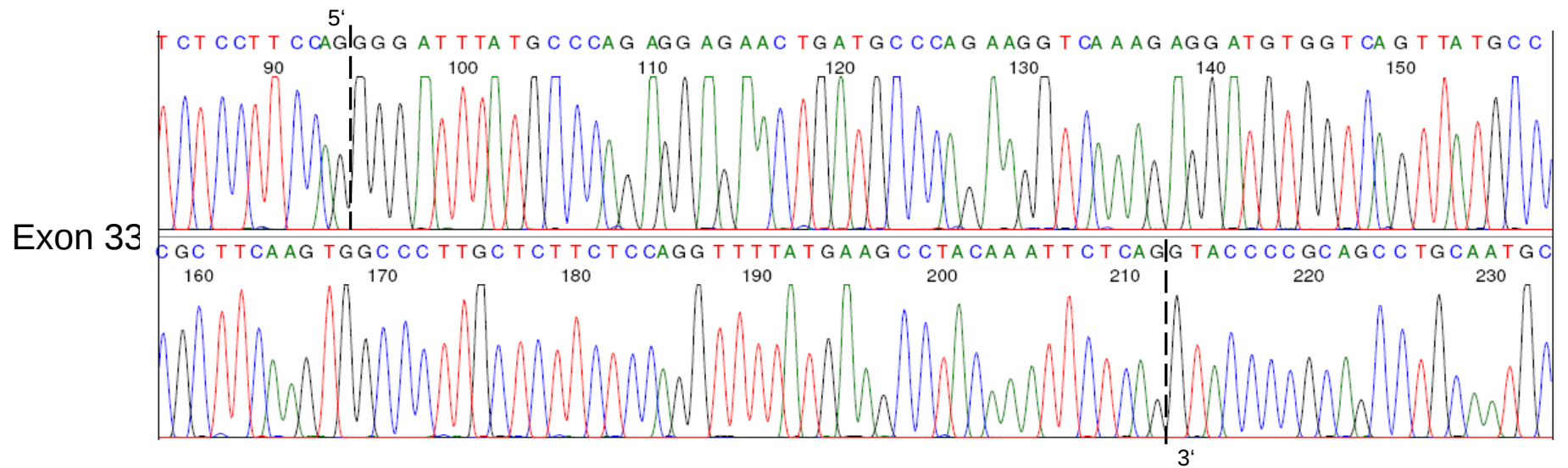
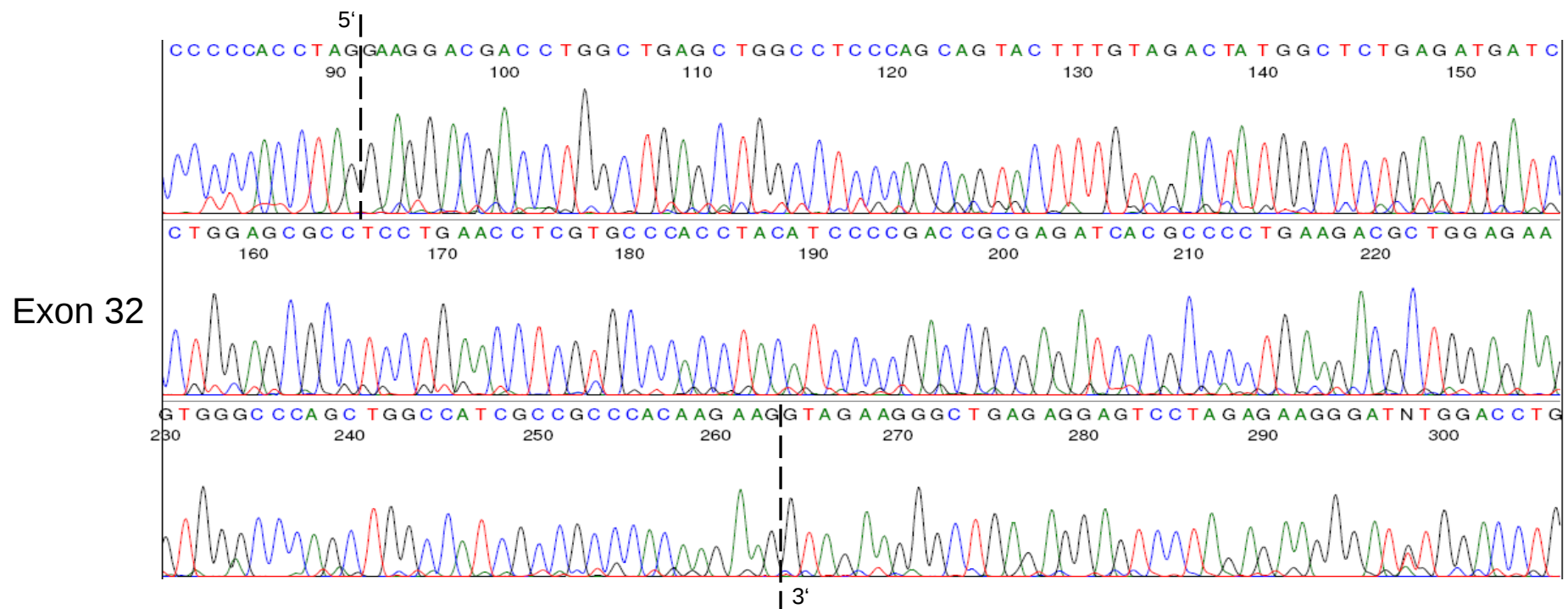


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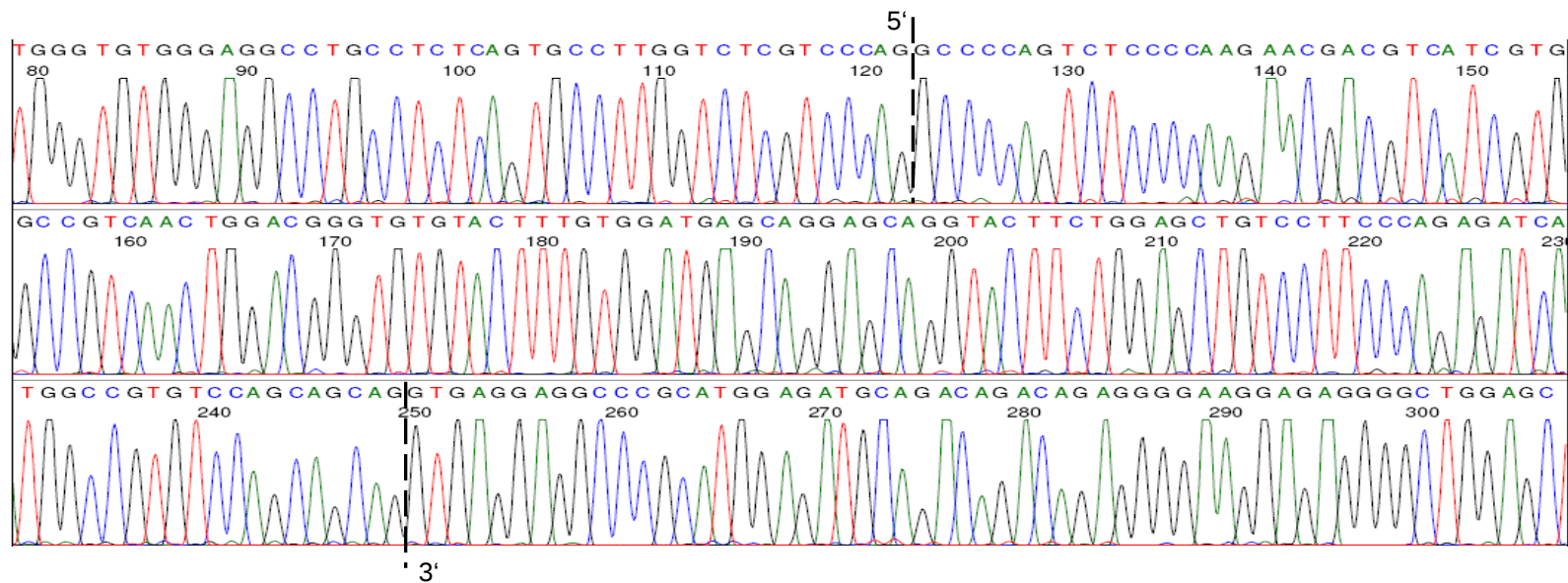


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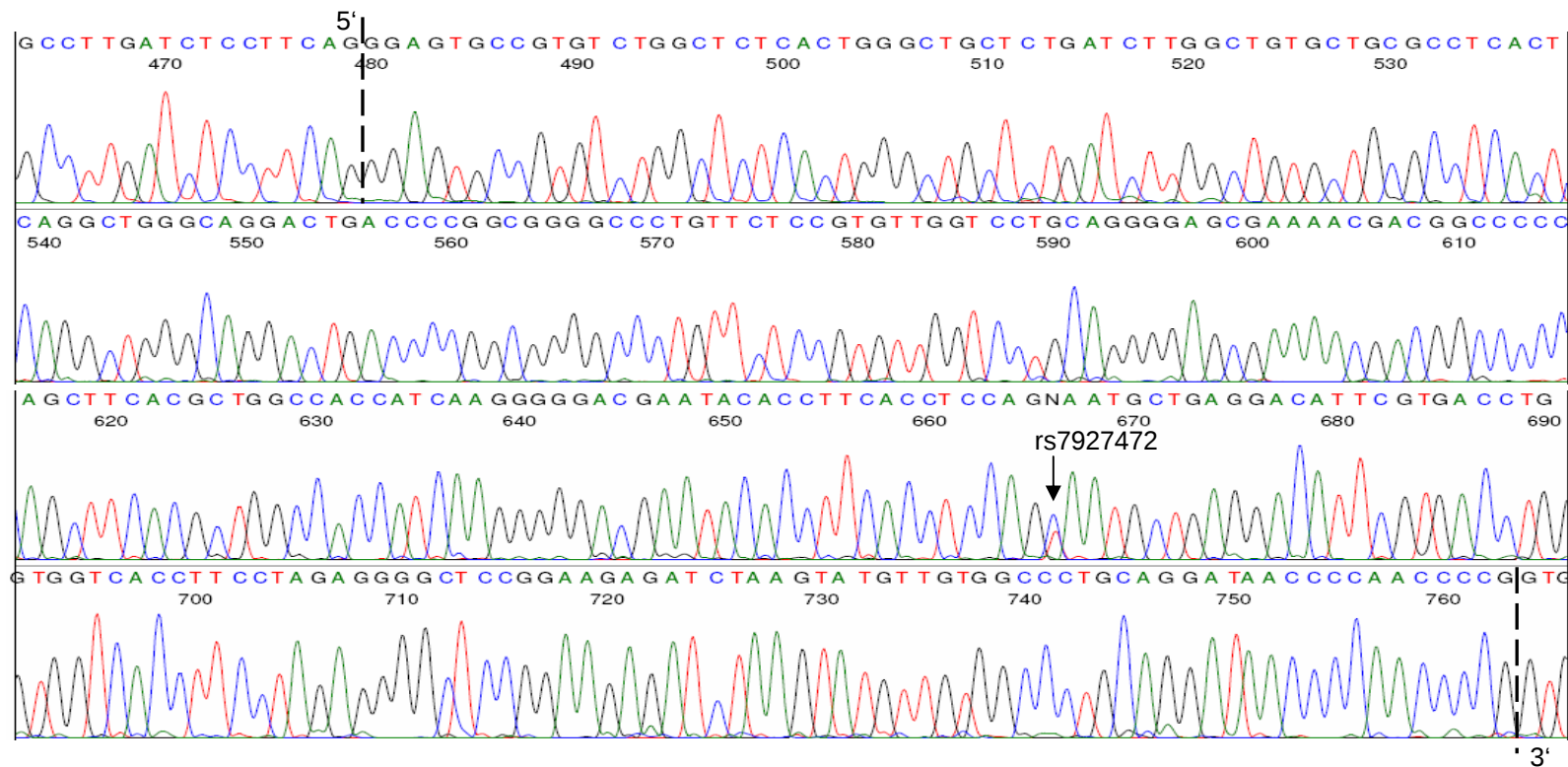




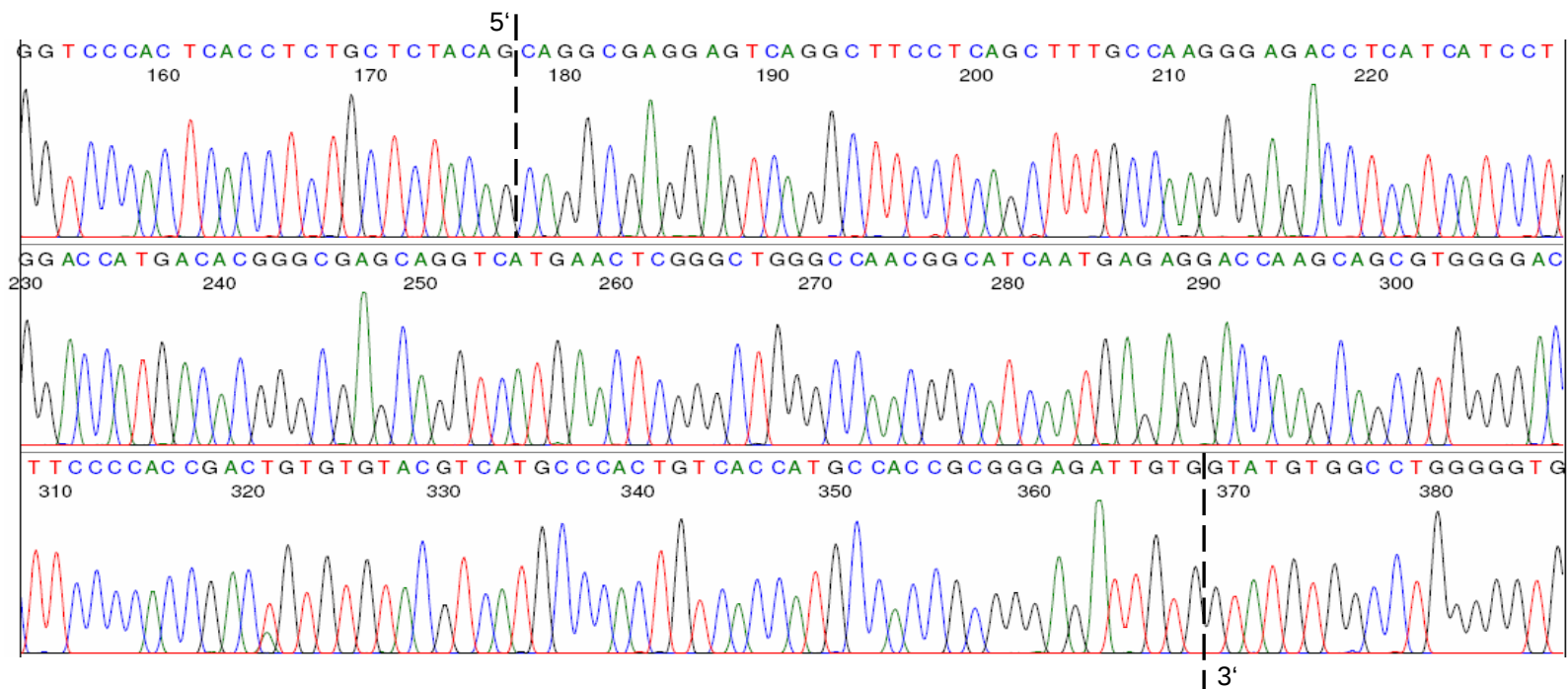
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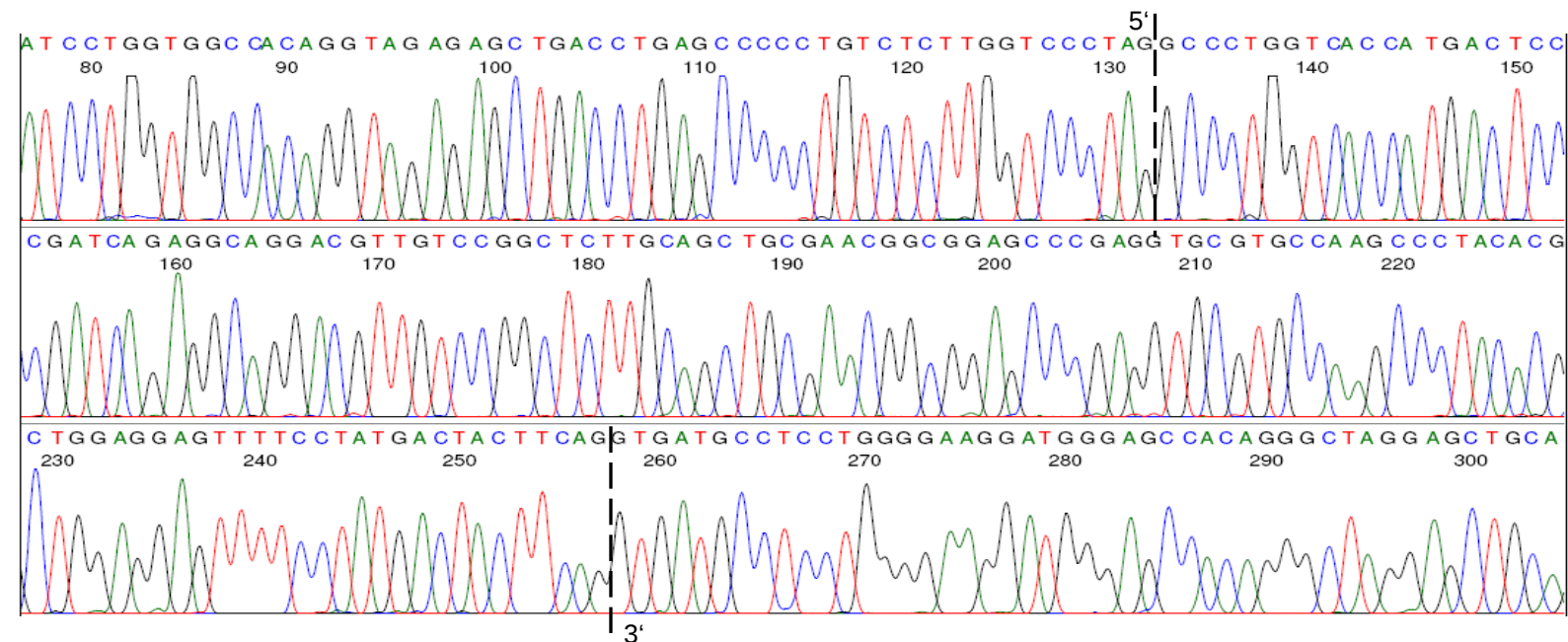
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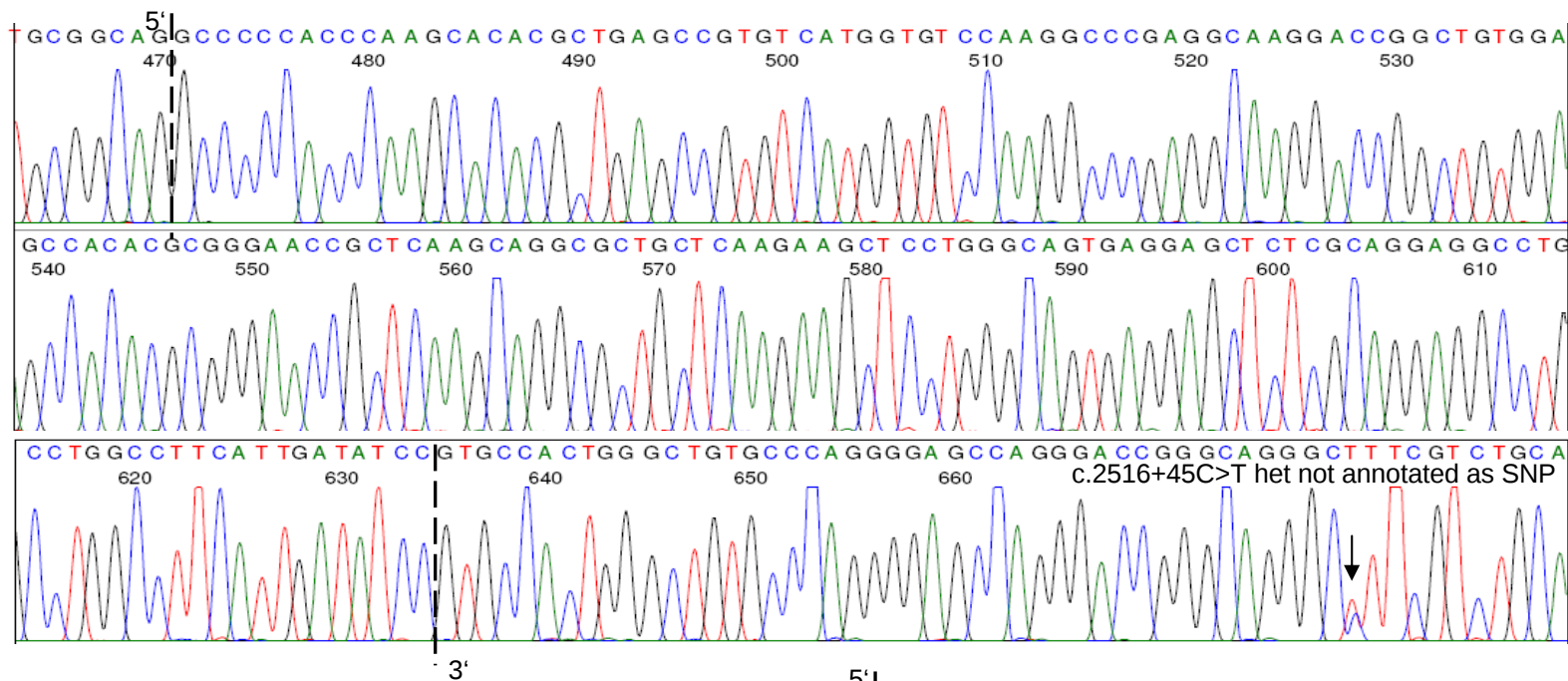
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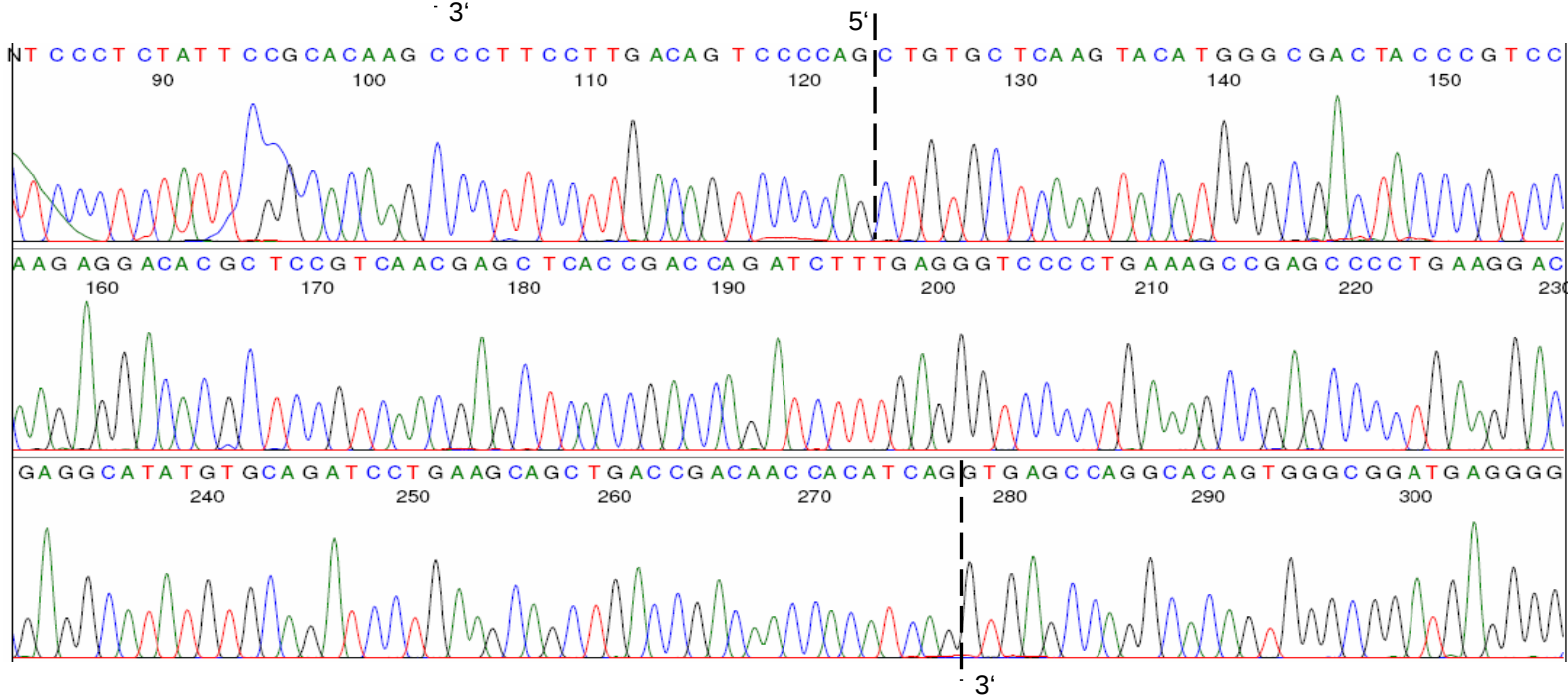
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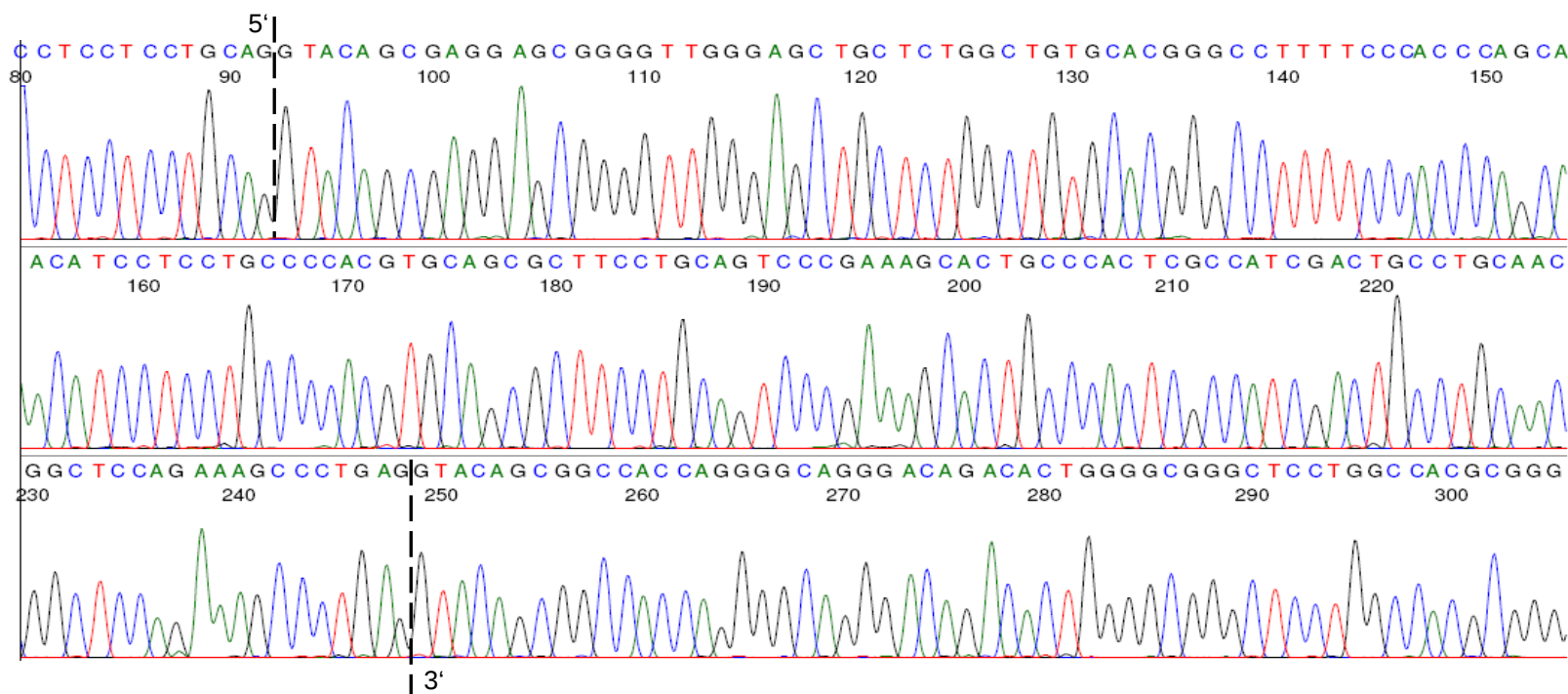
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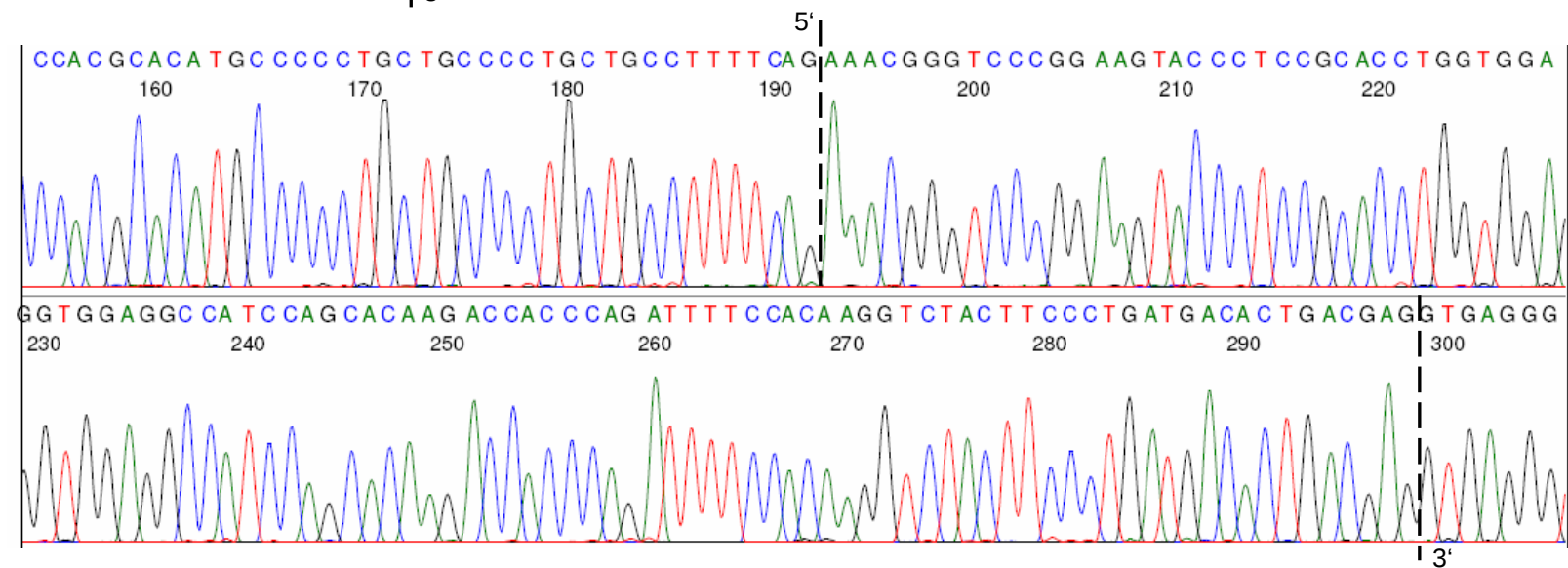
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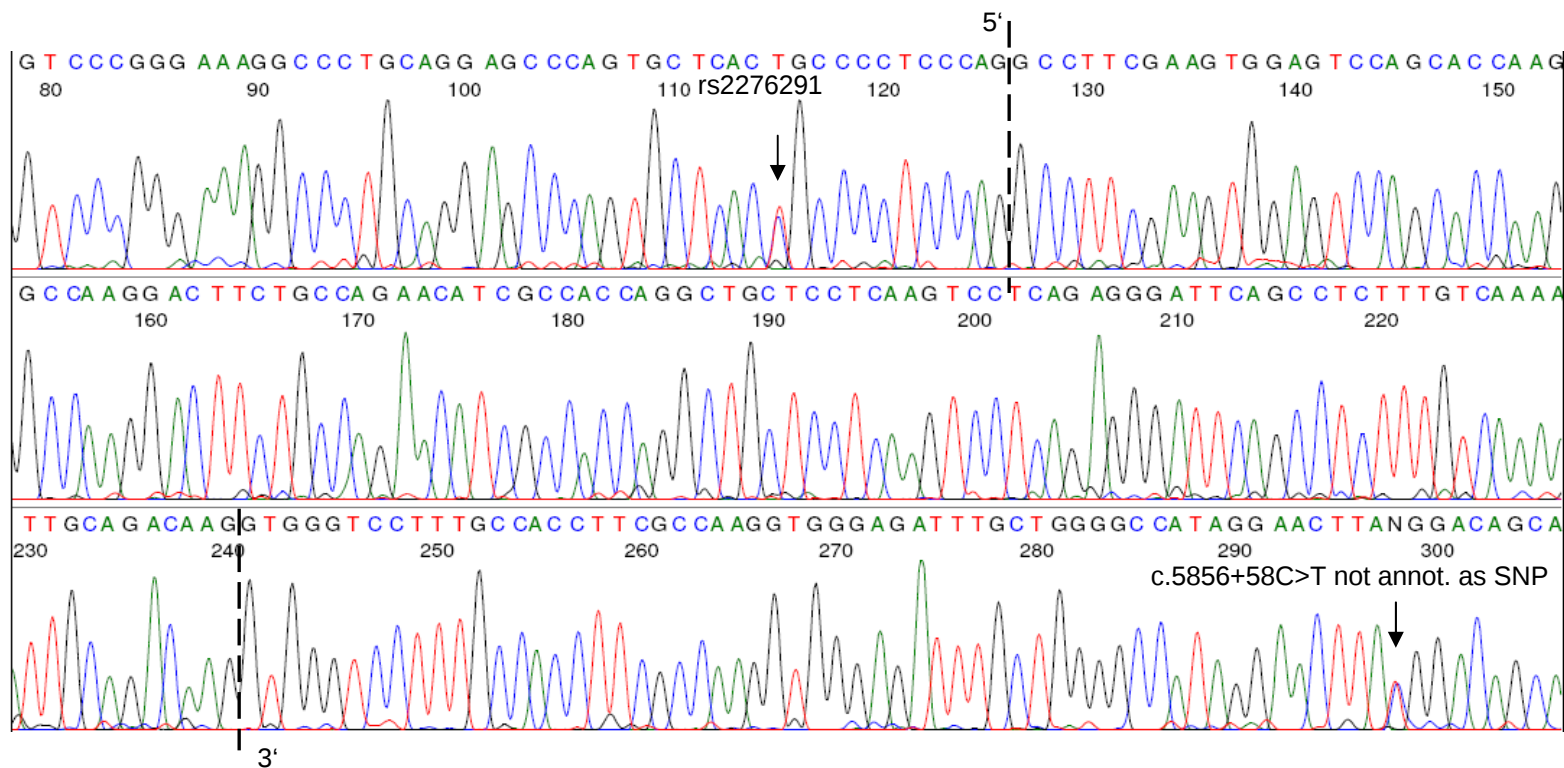
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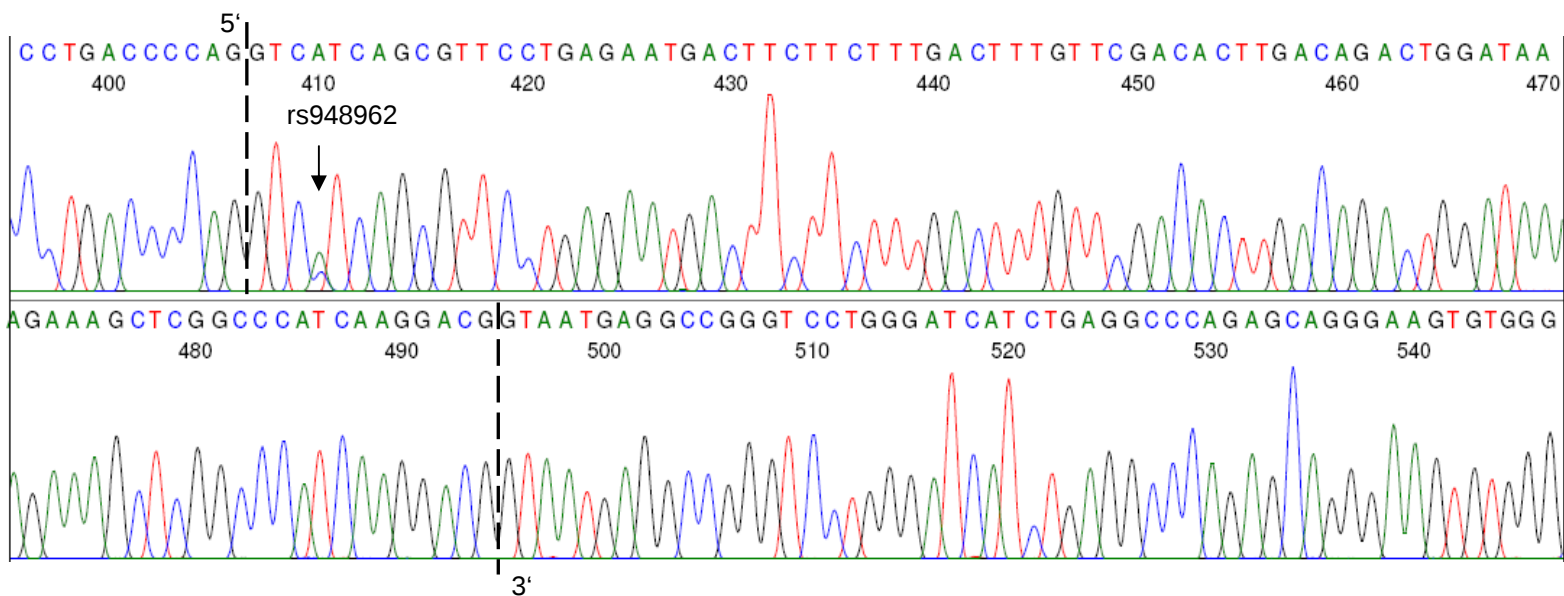
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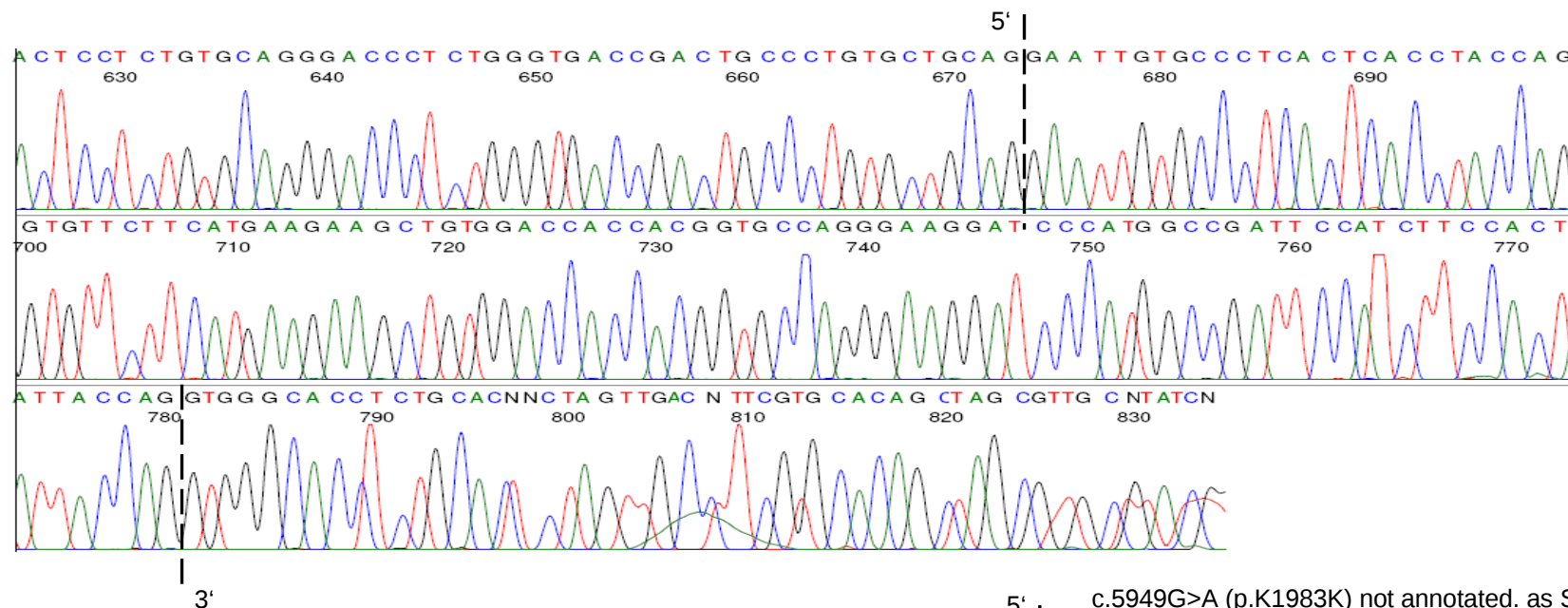
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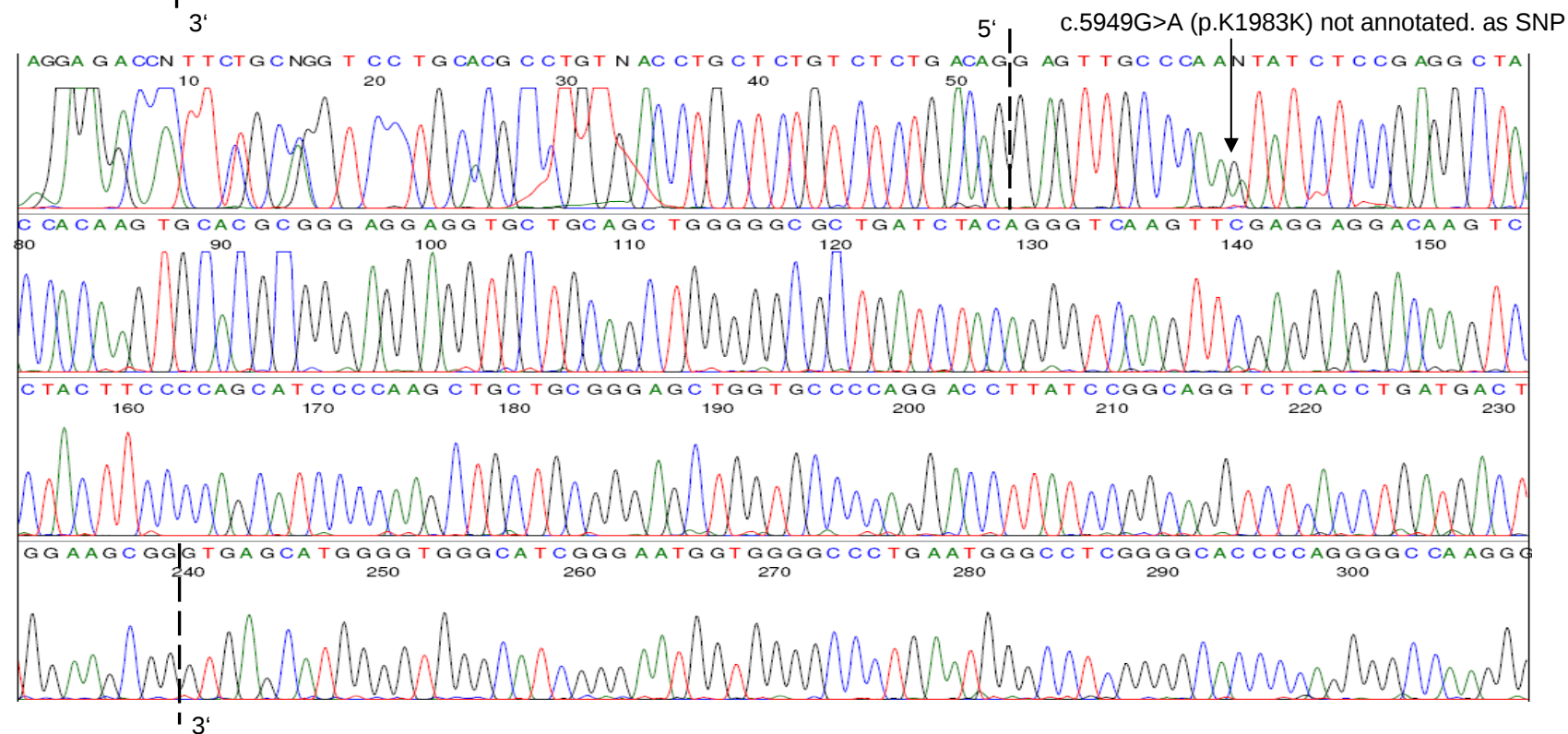
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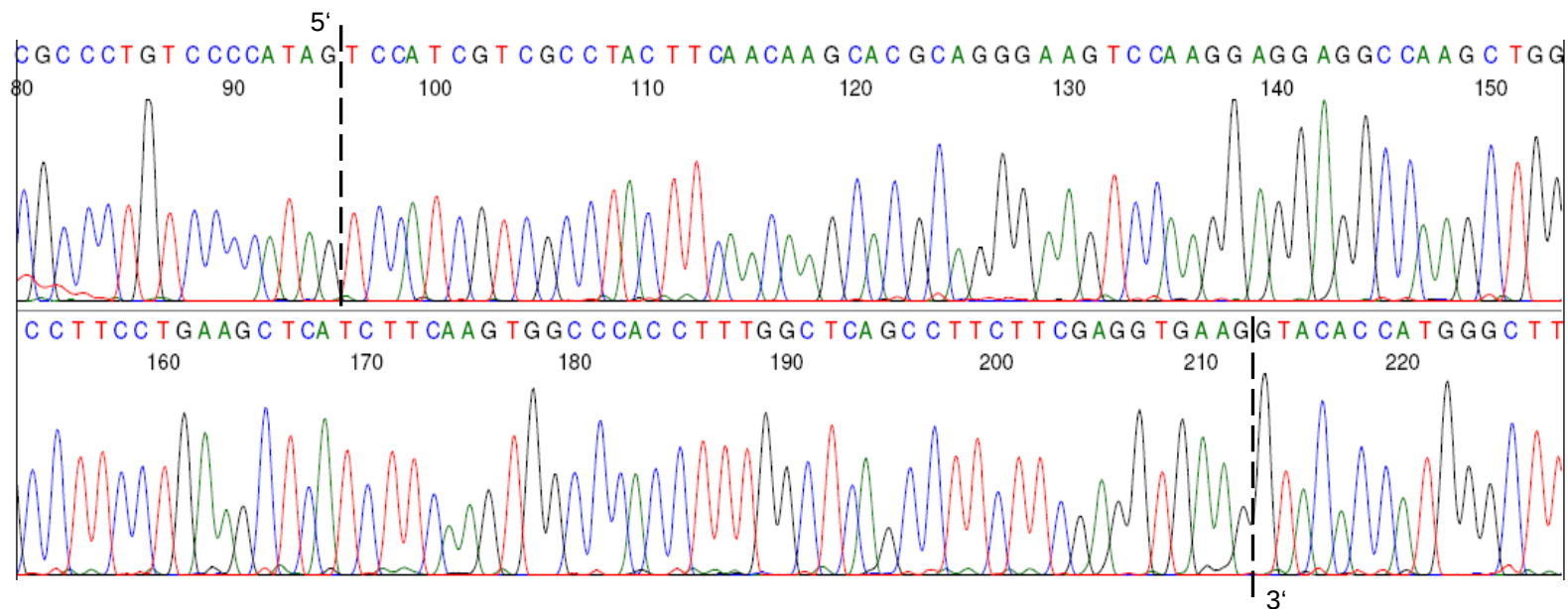
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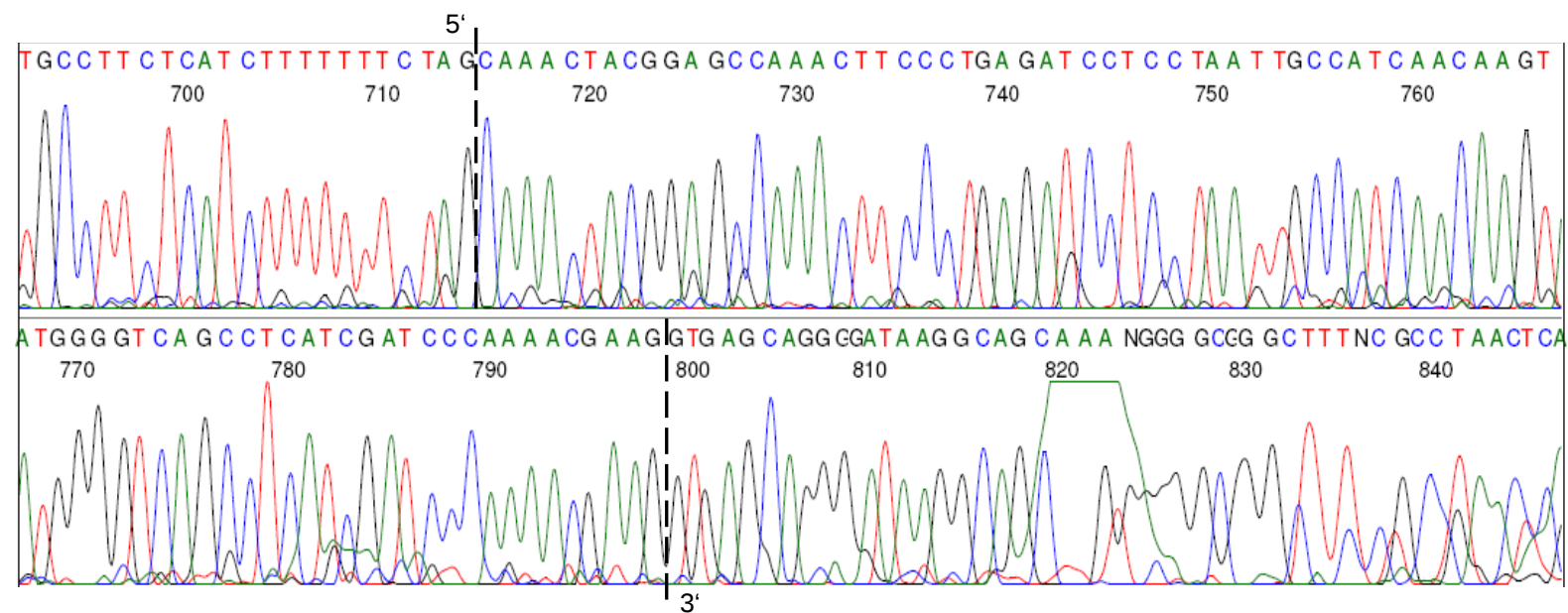
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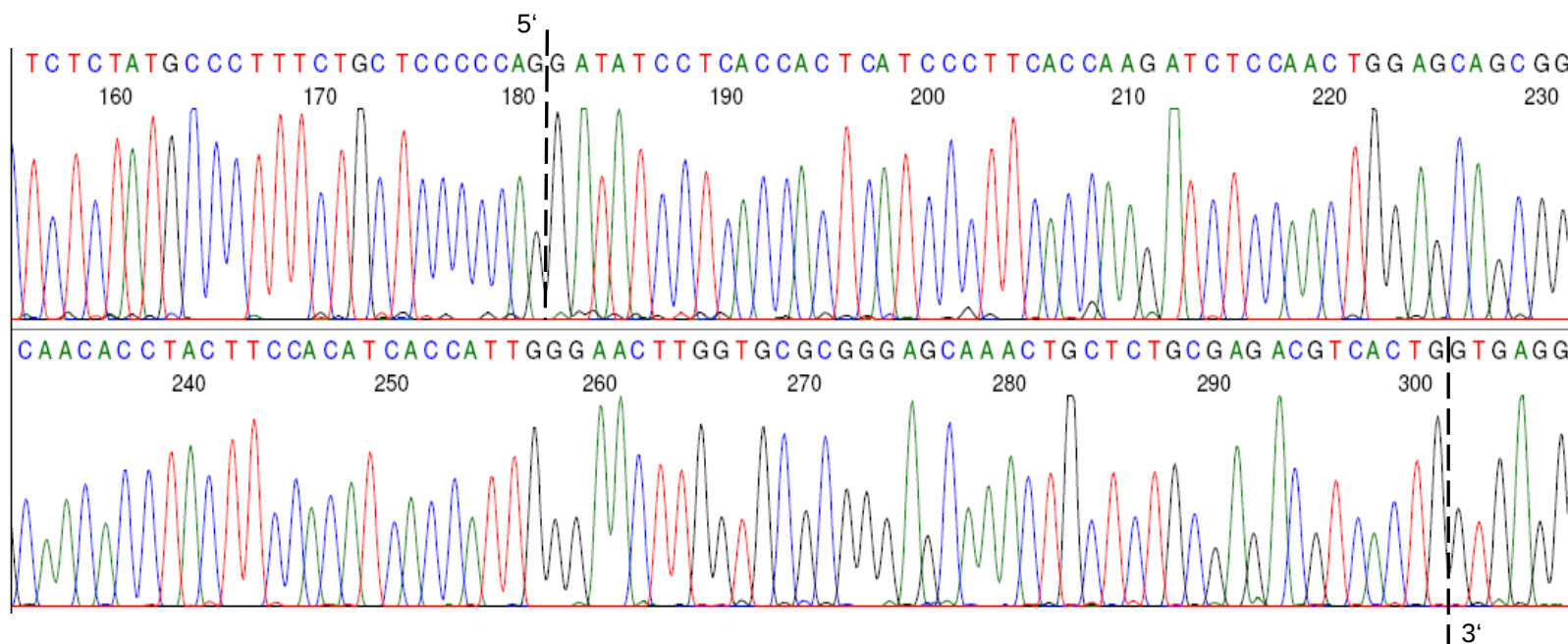
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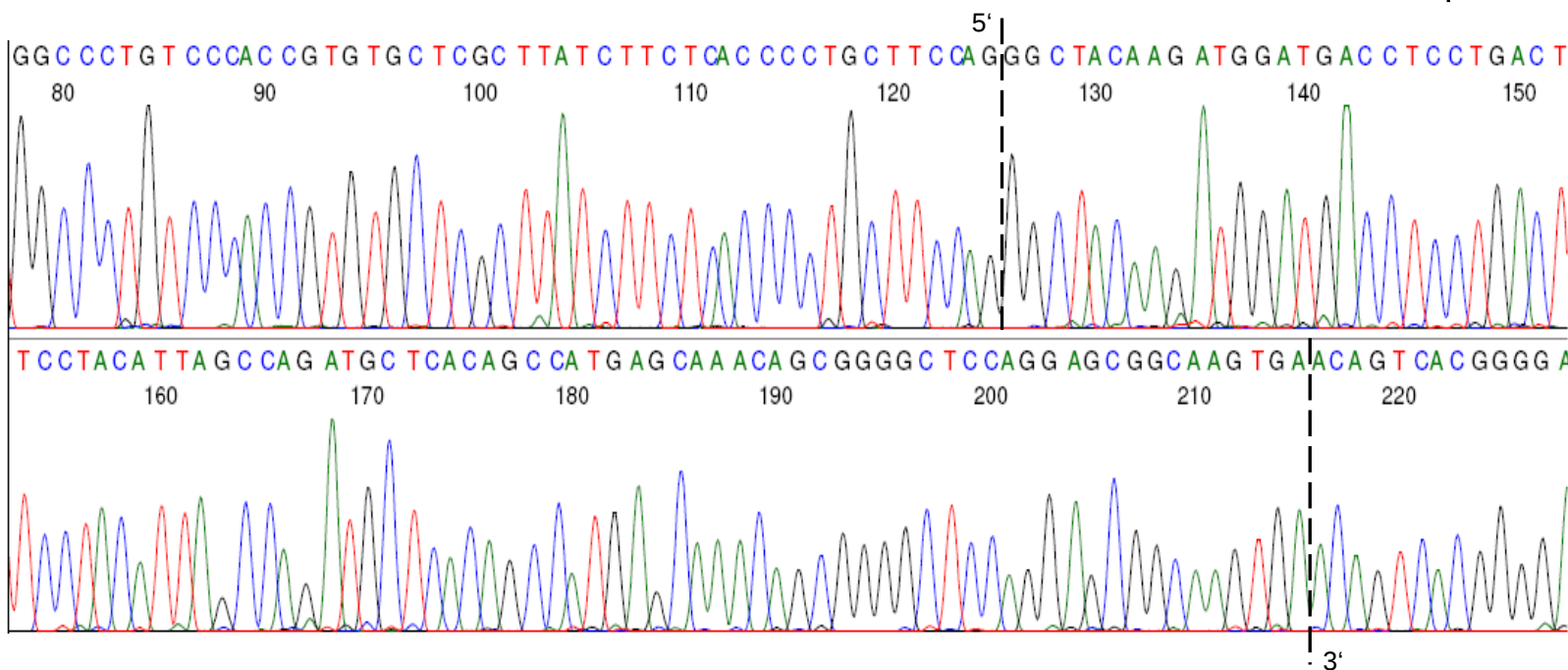
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Exon 48

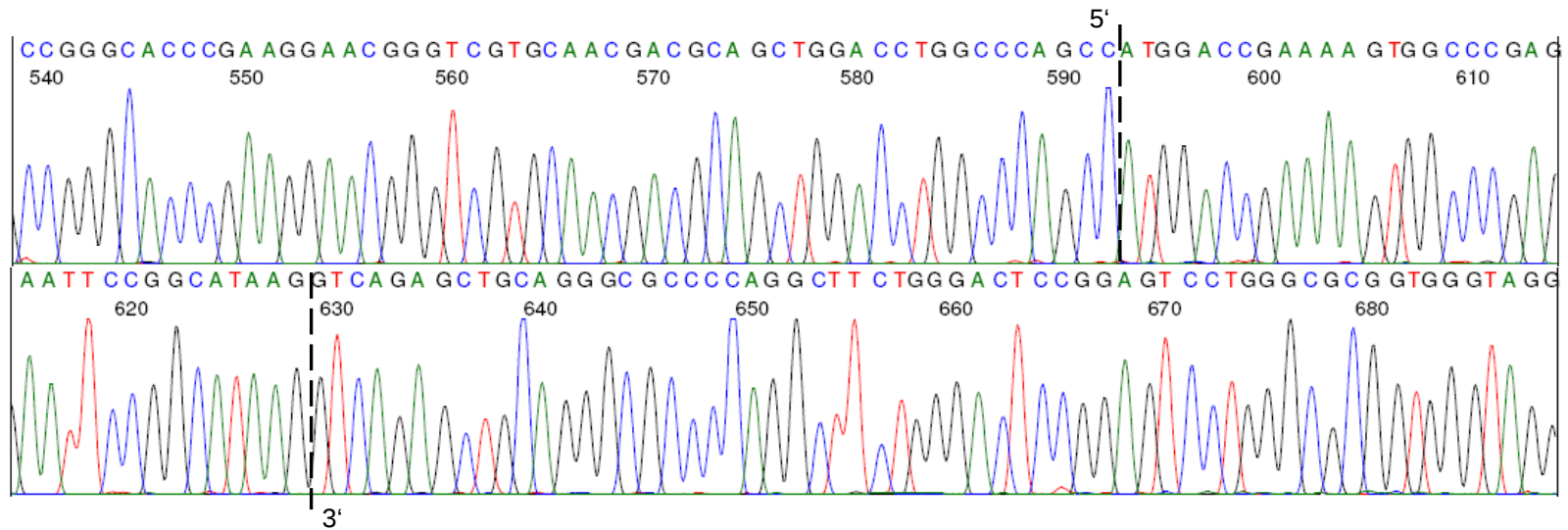


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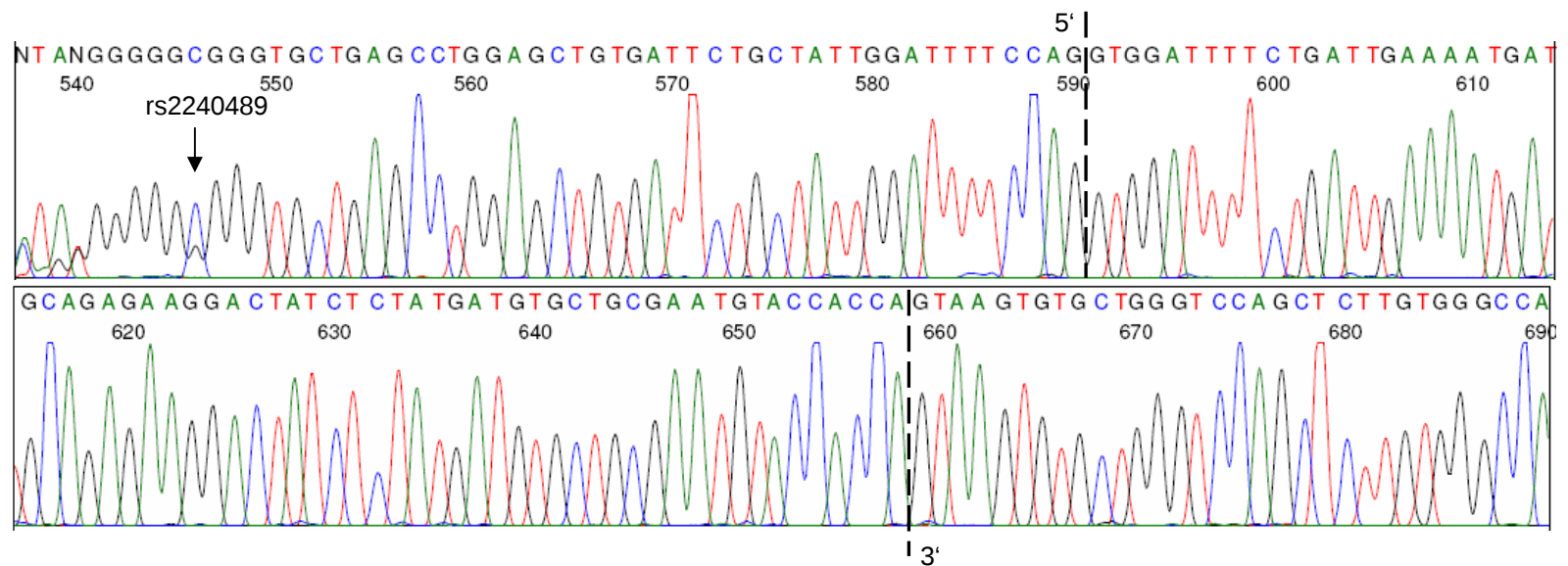


Patient 1881, *USH1C*

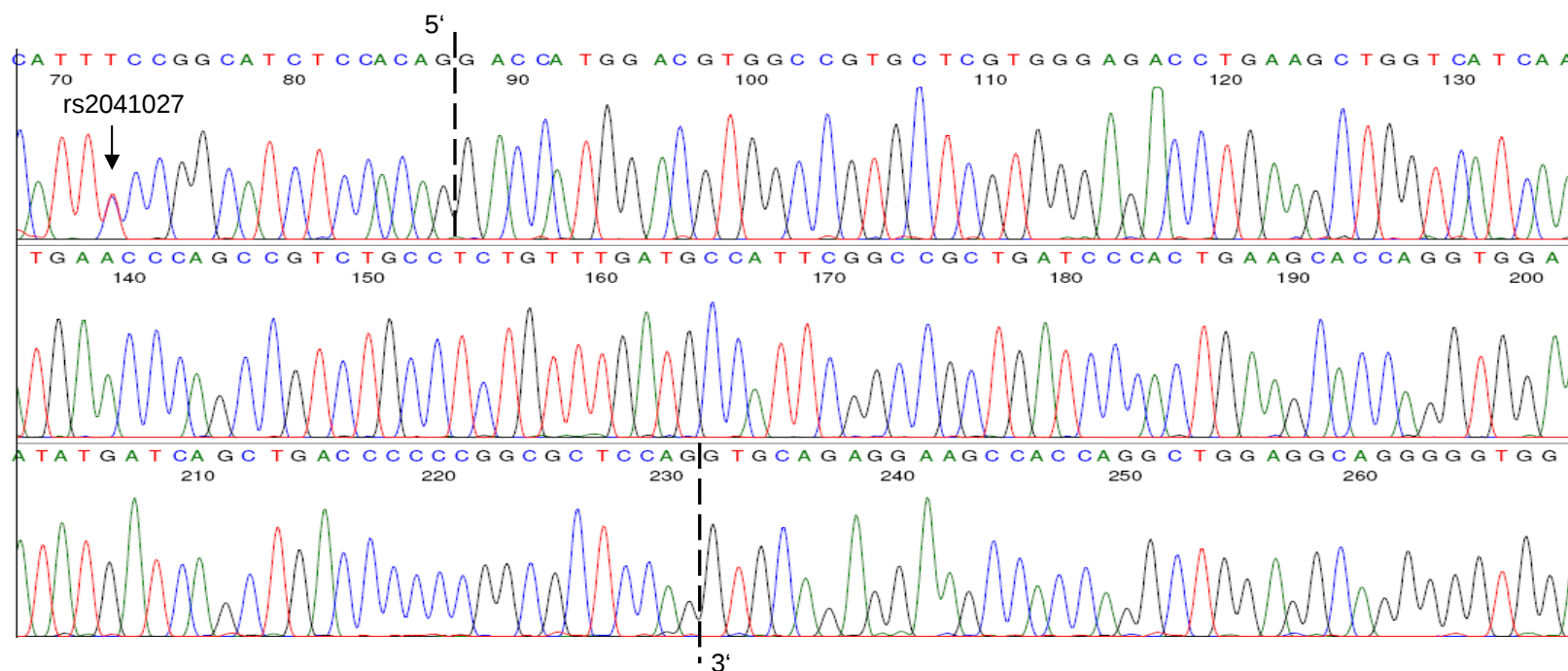
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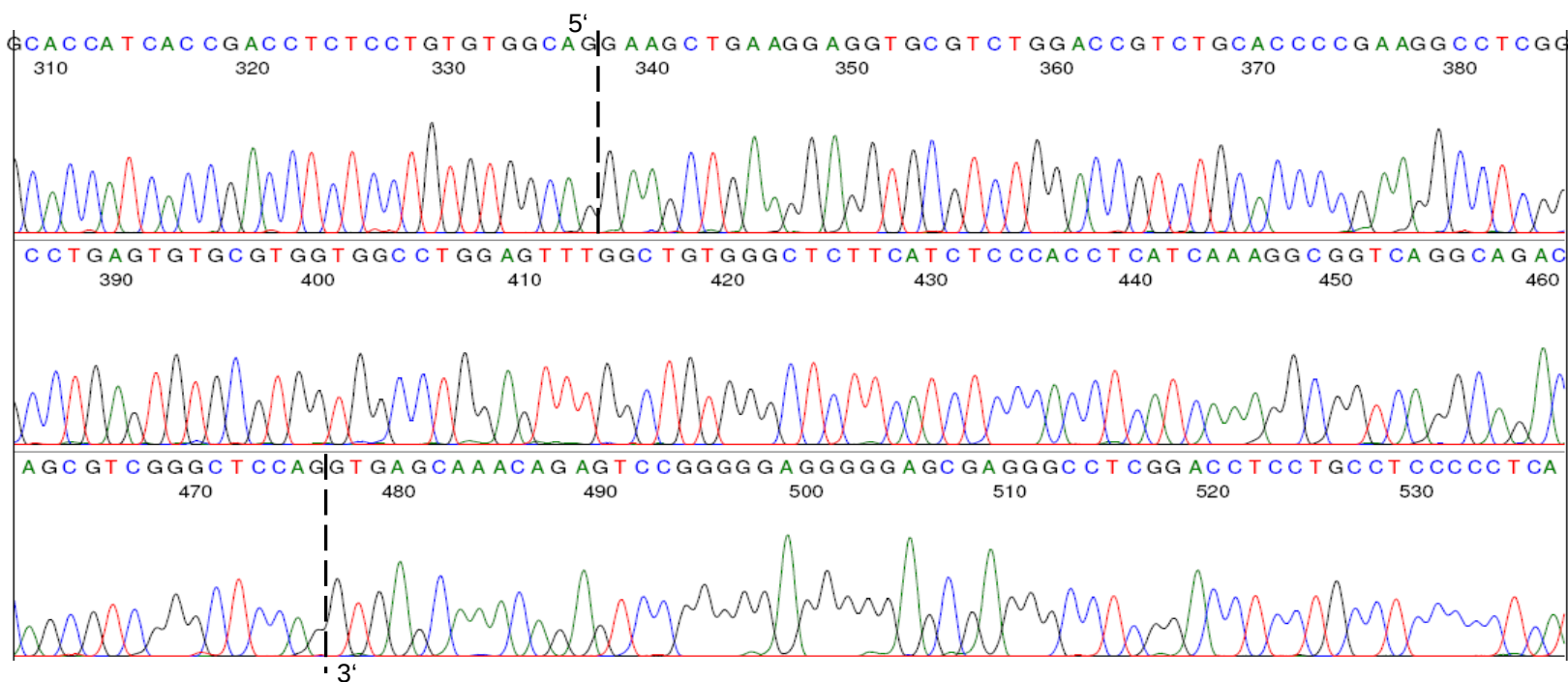
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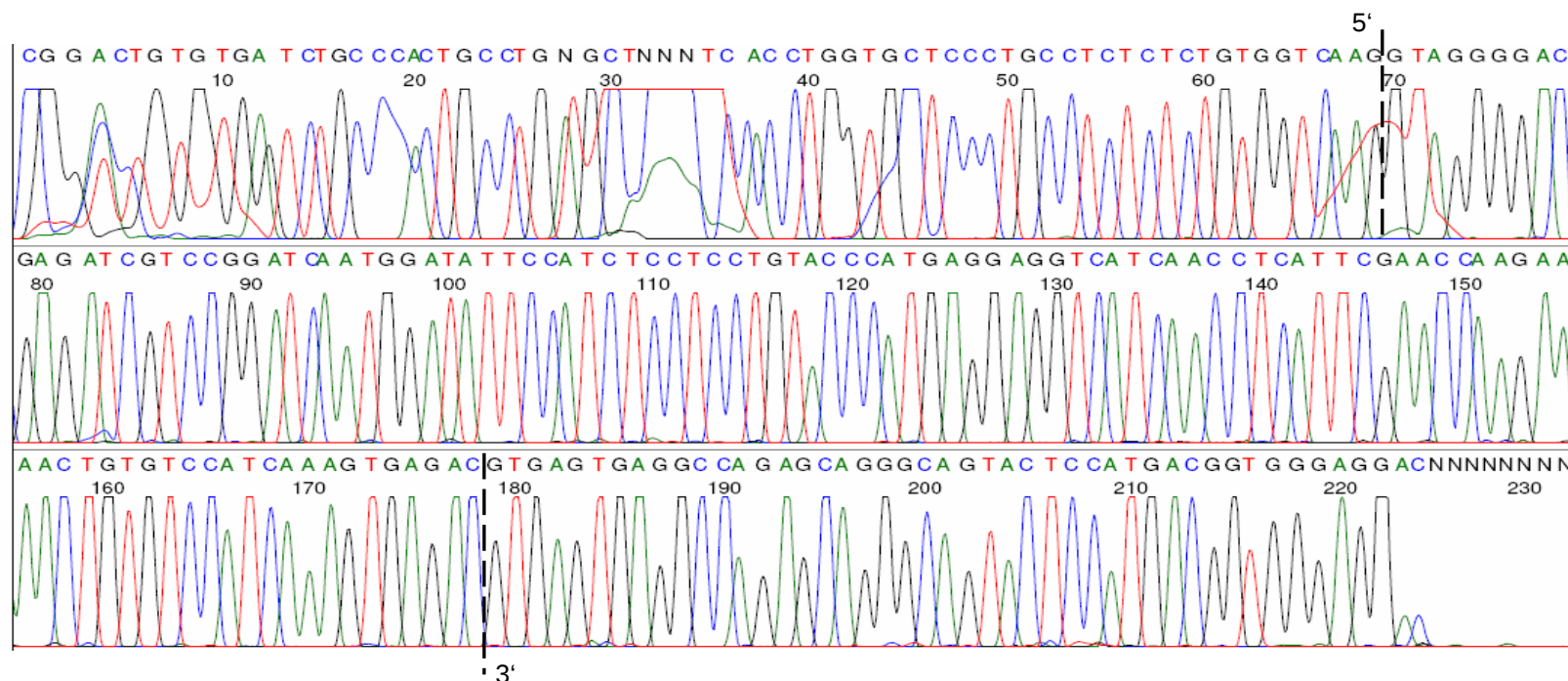
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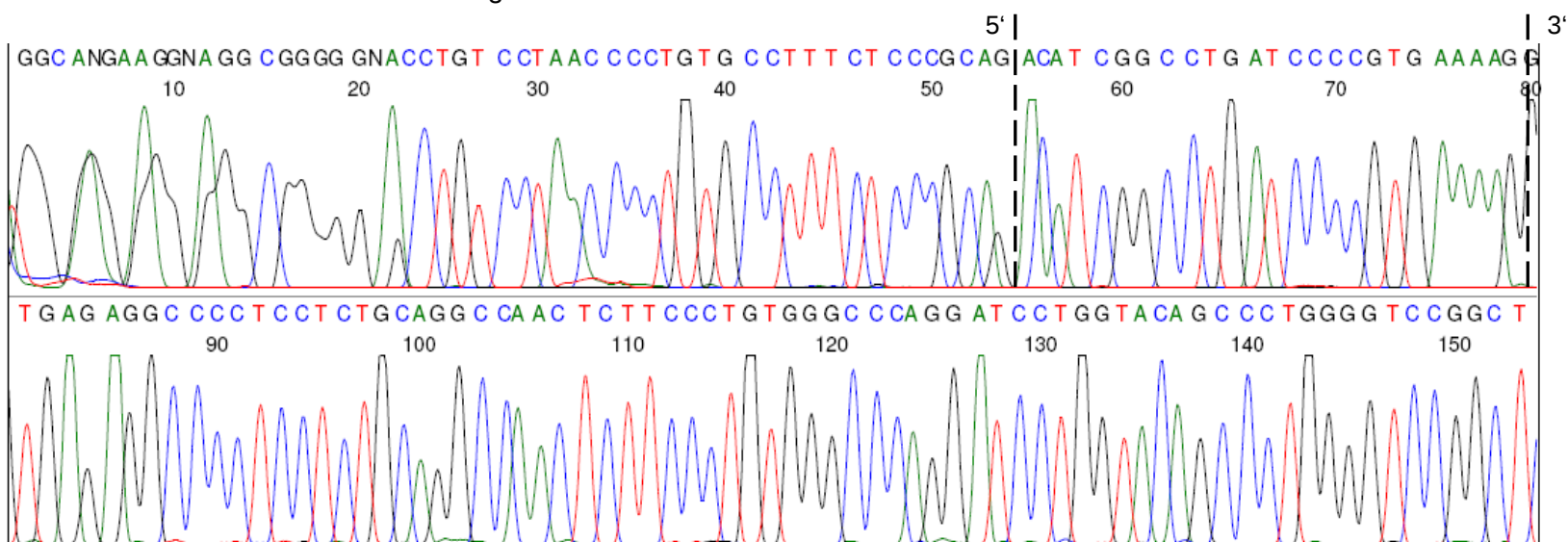
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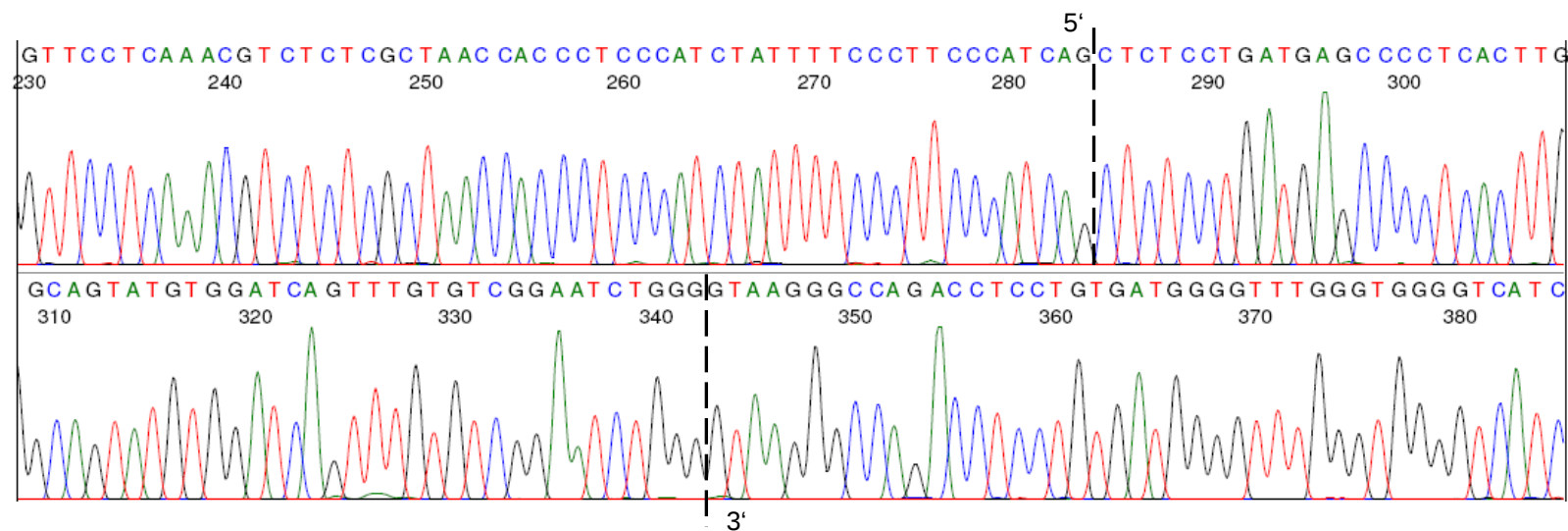
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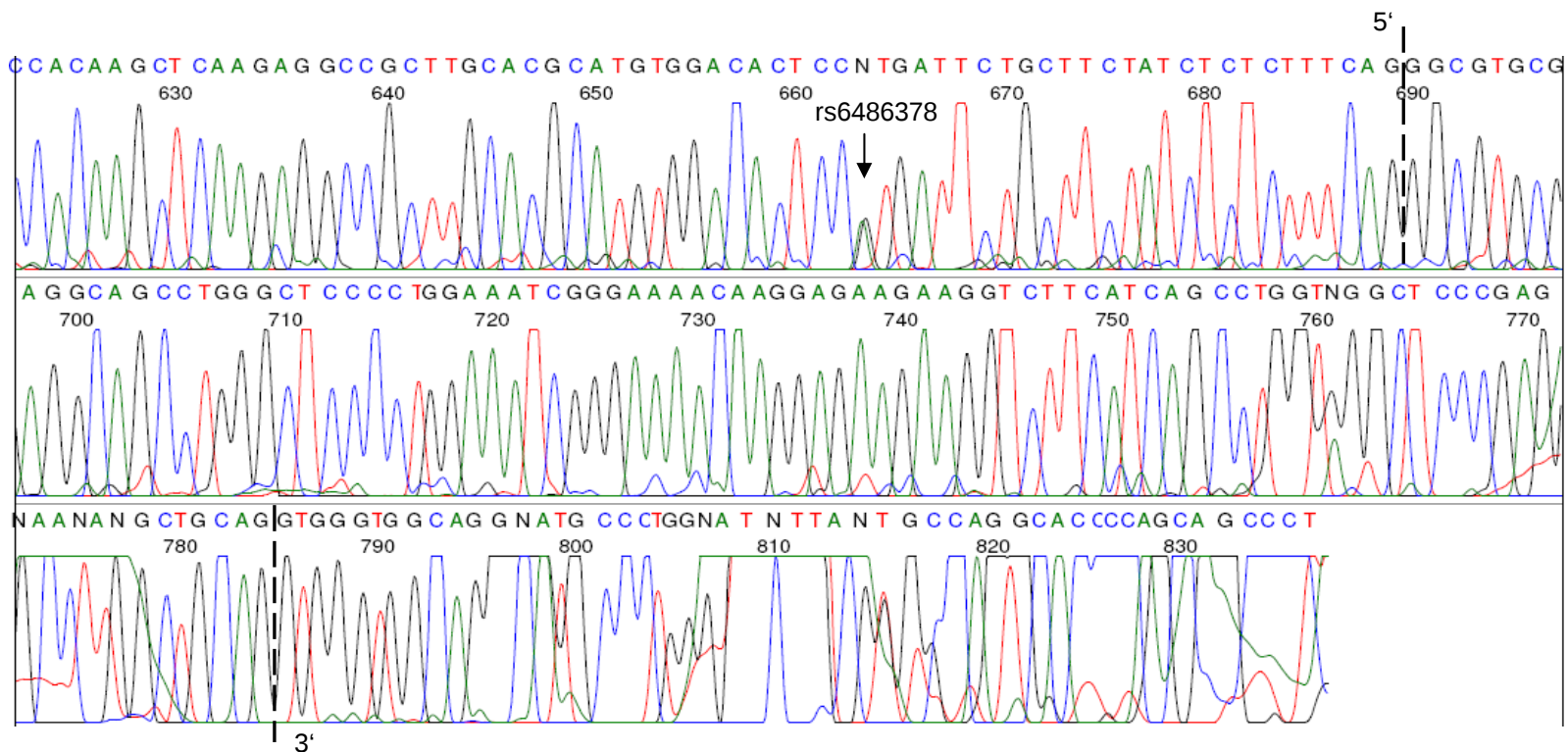
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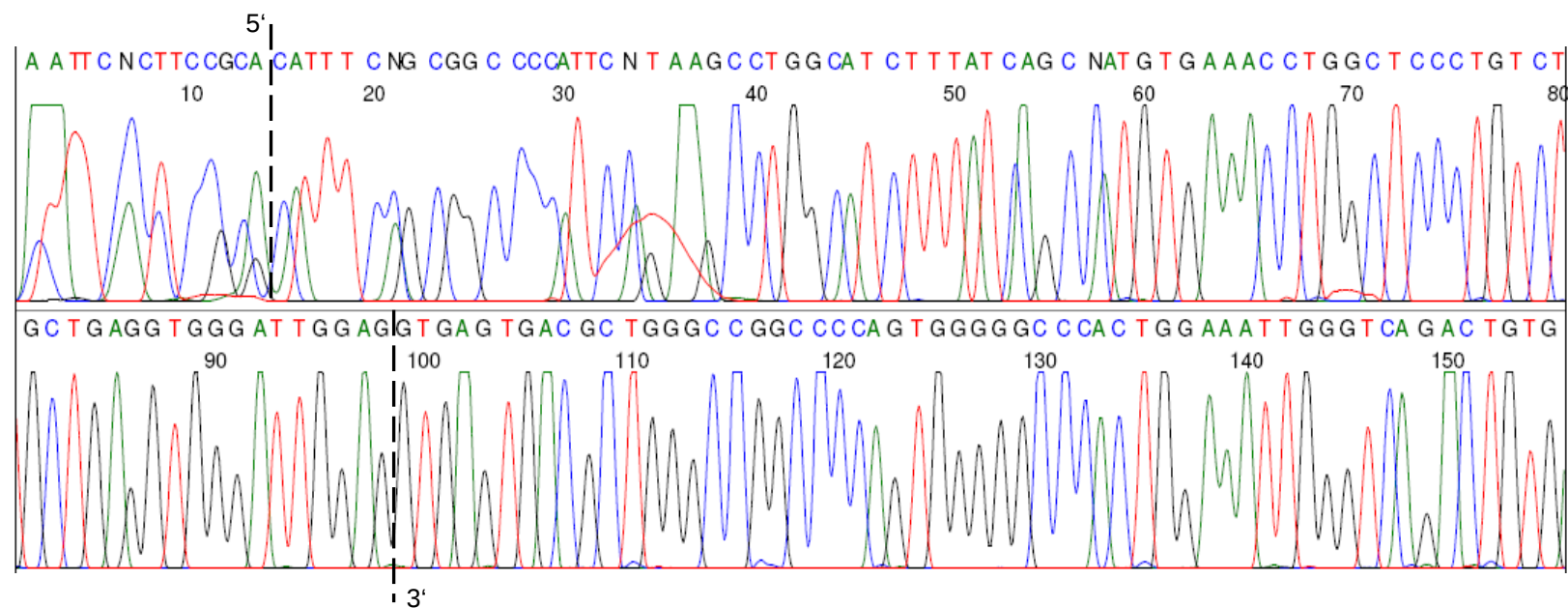
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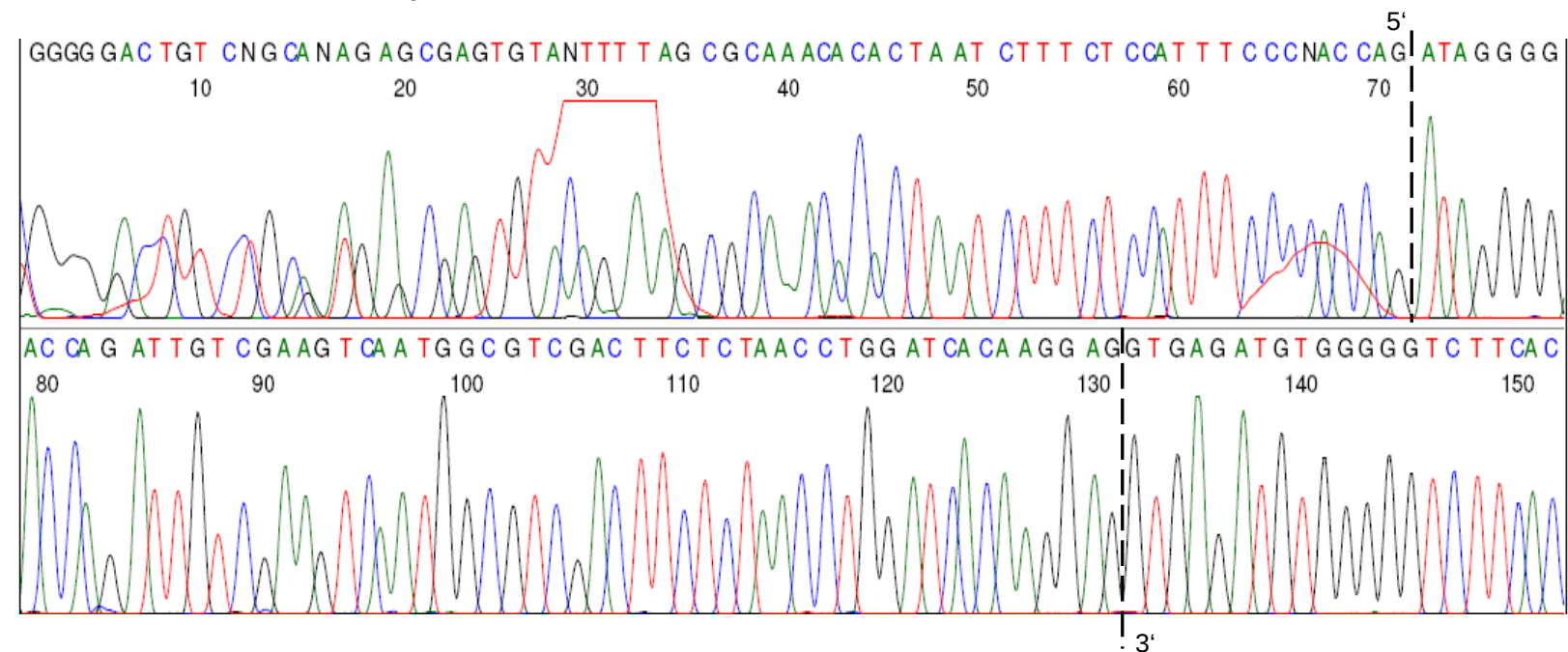
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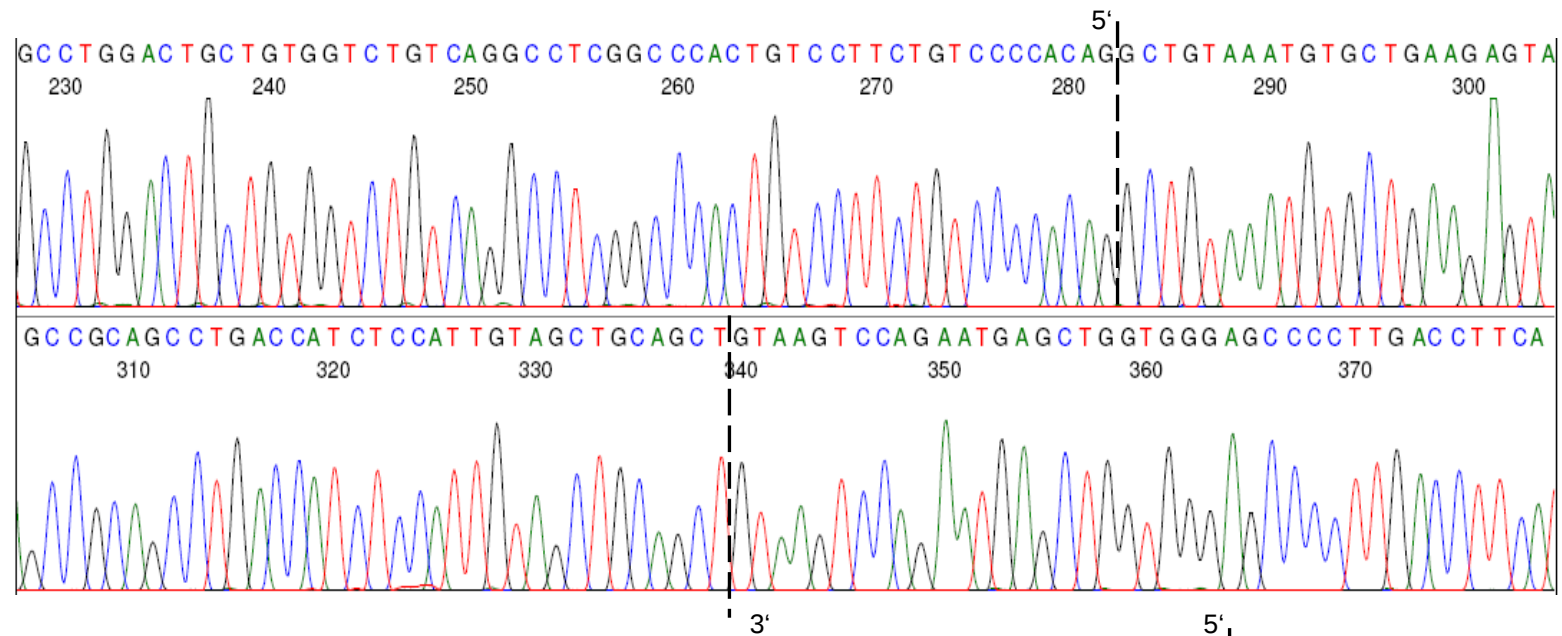
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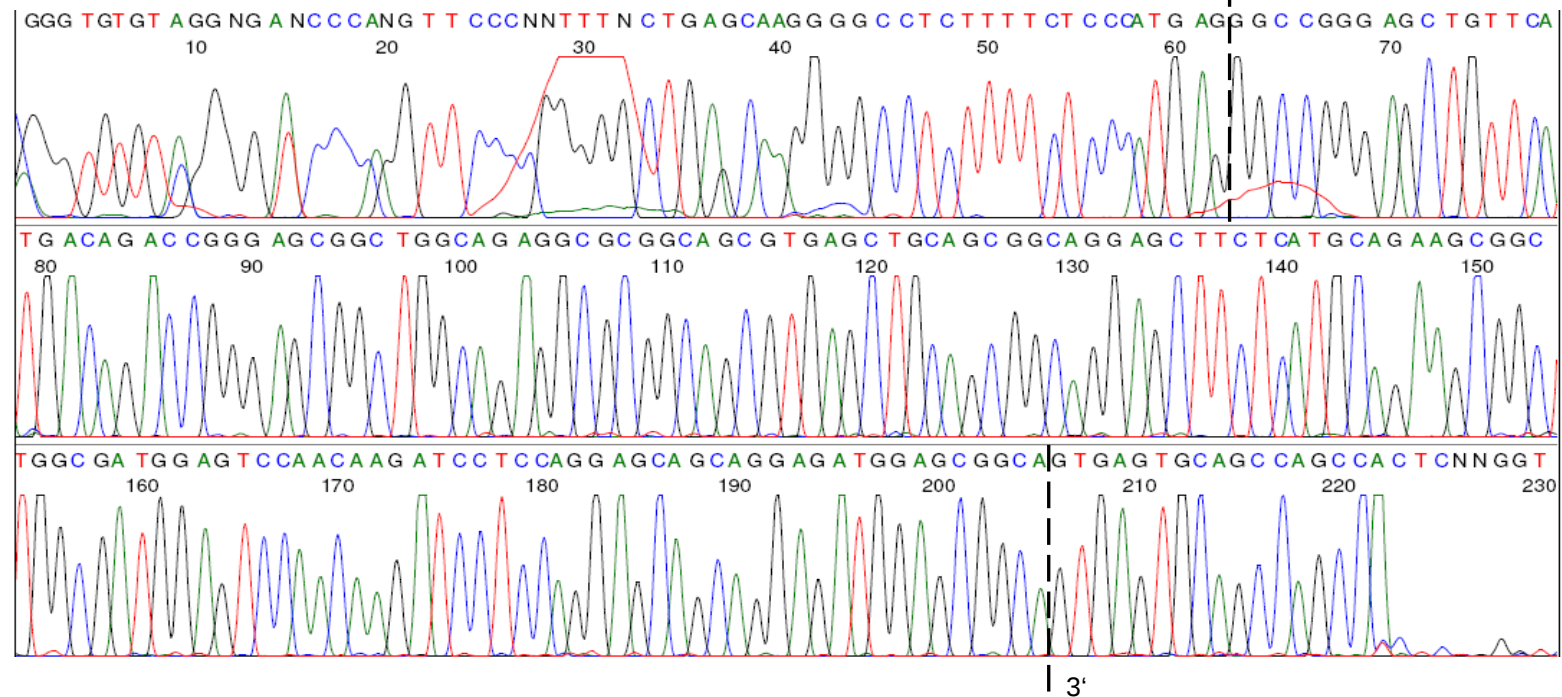
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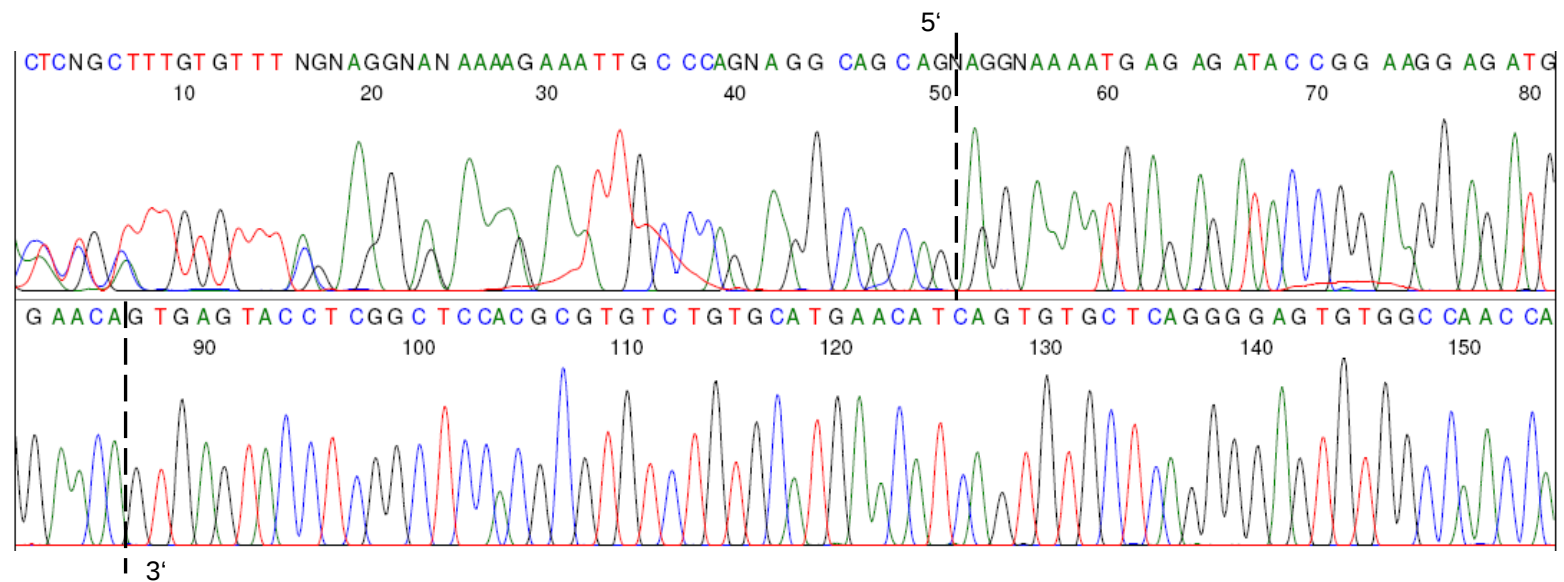
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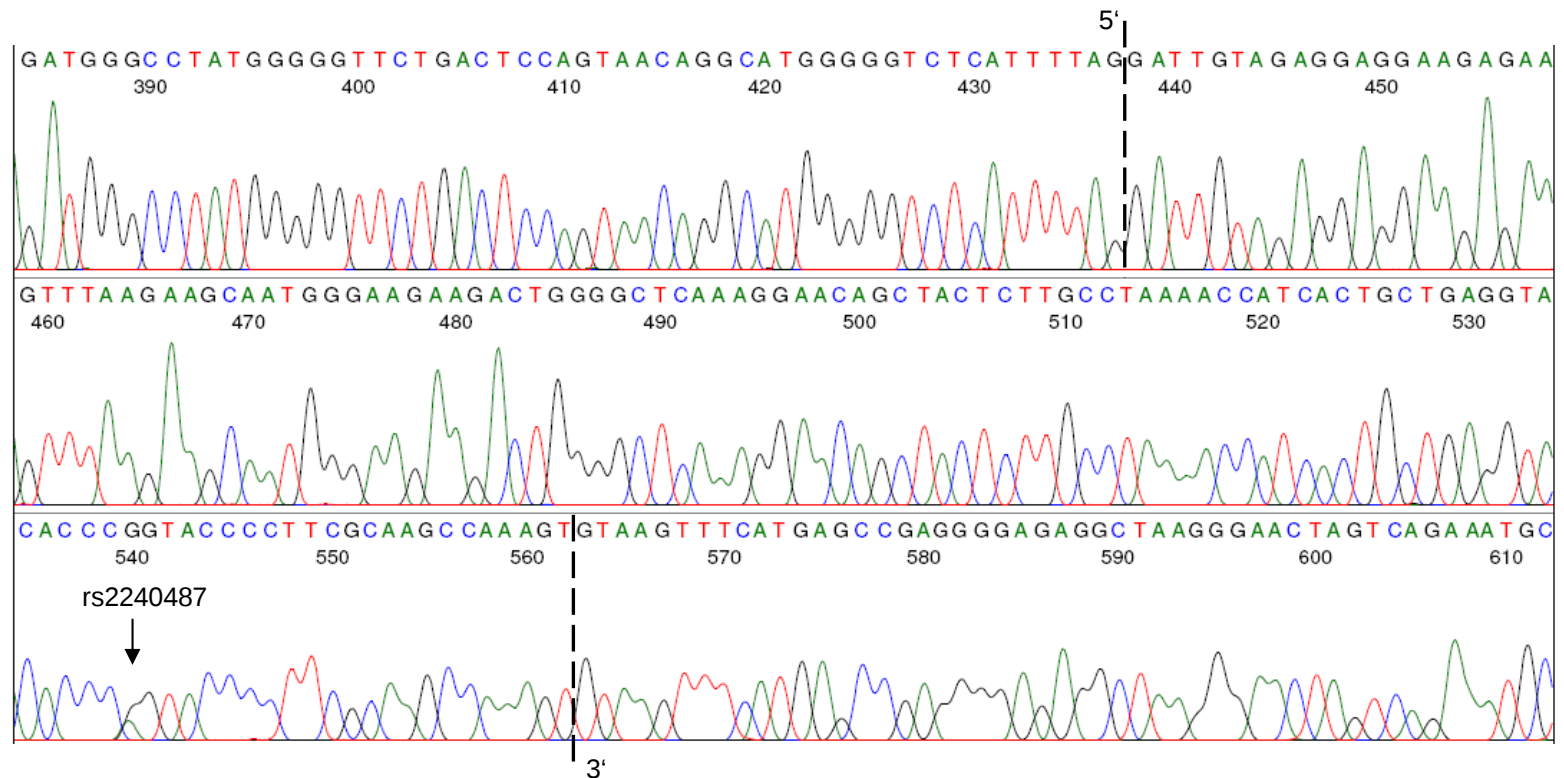
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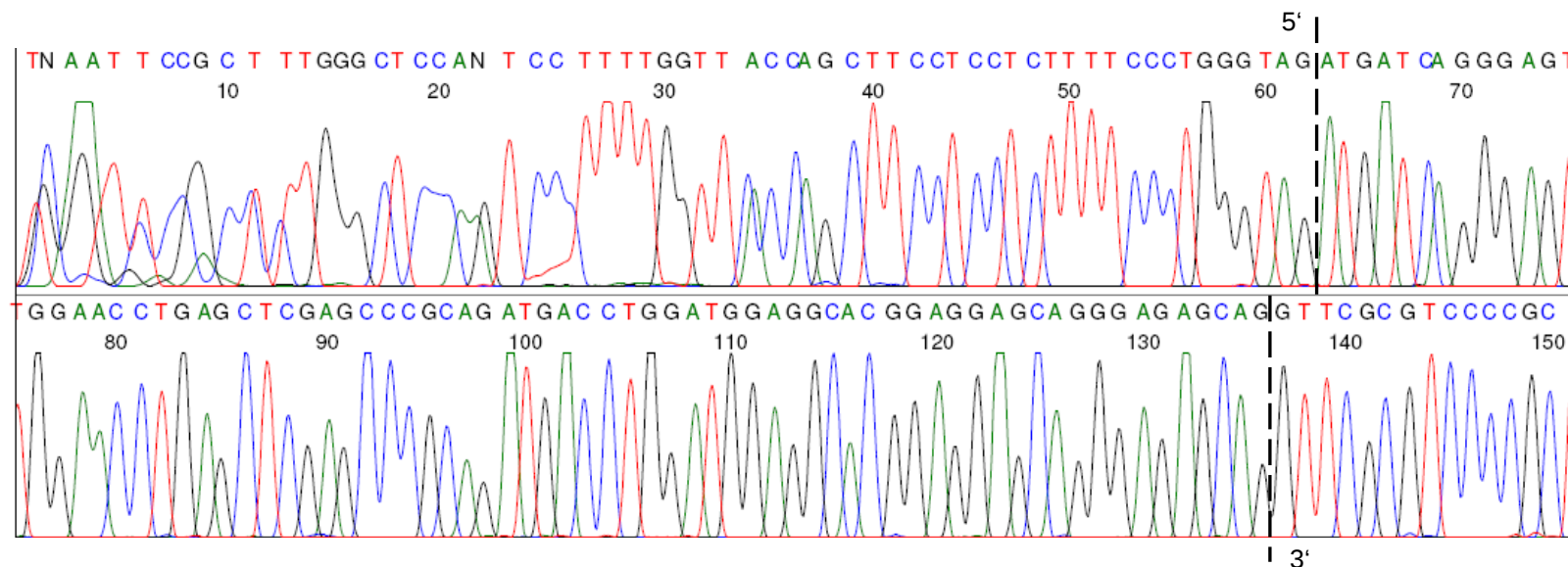
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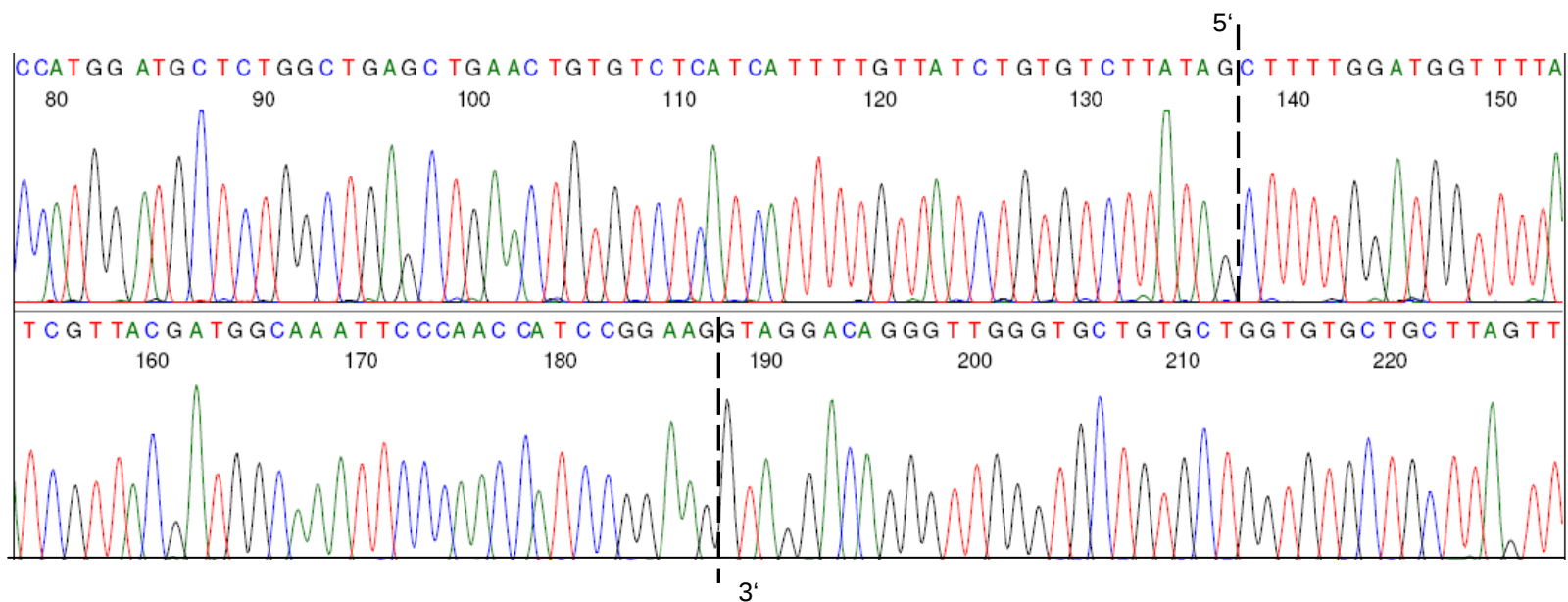
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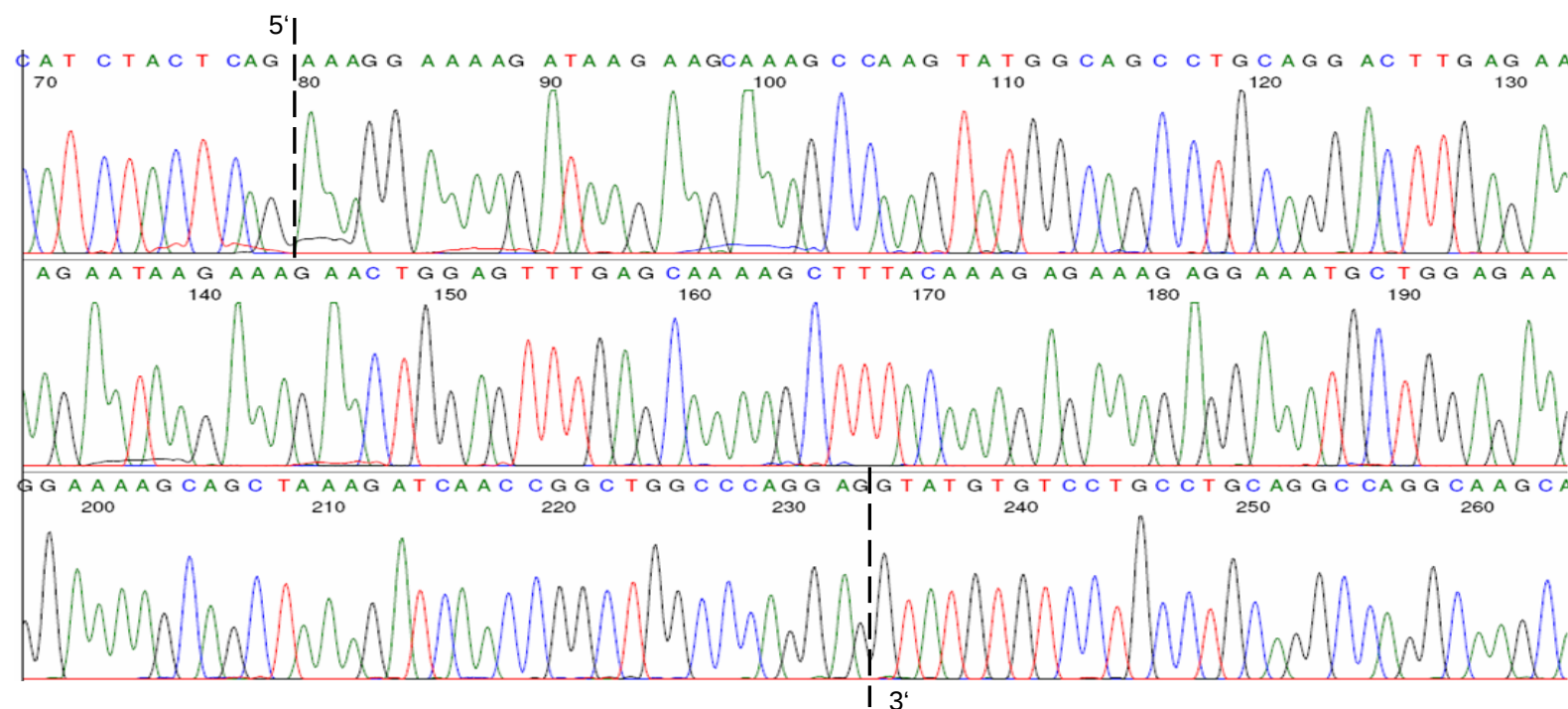
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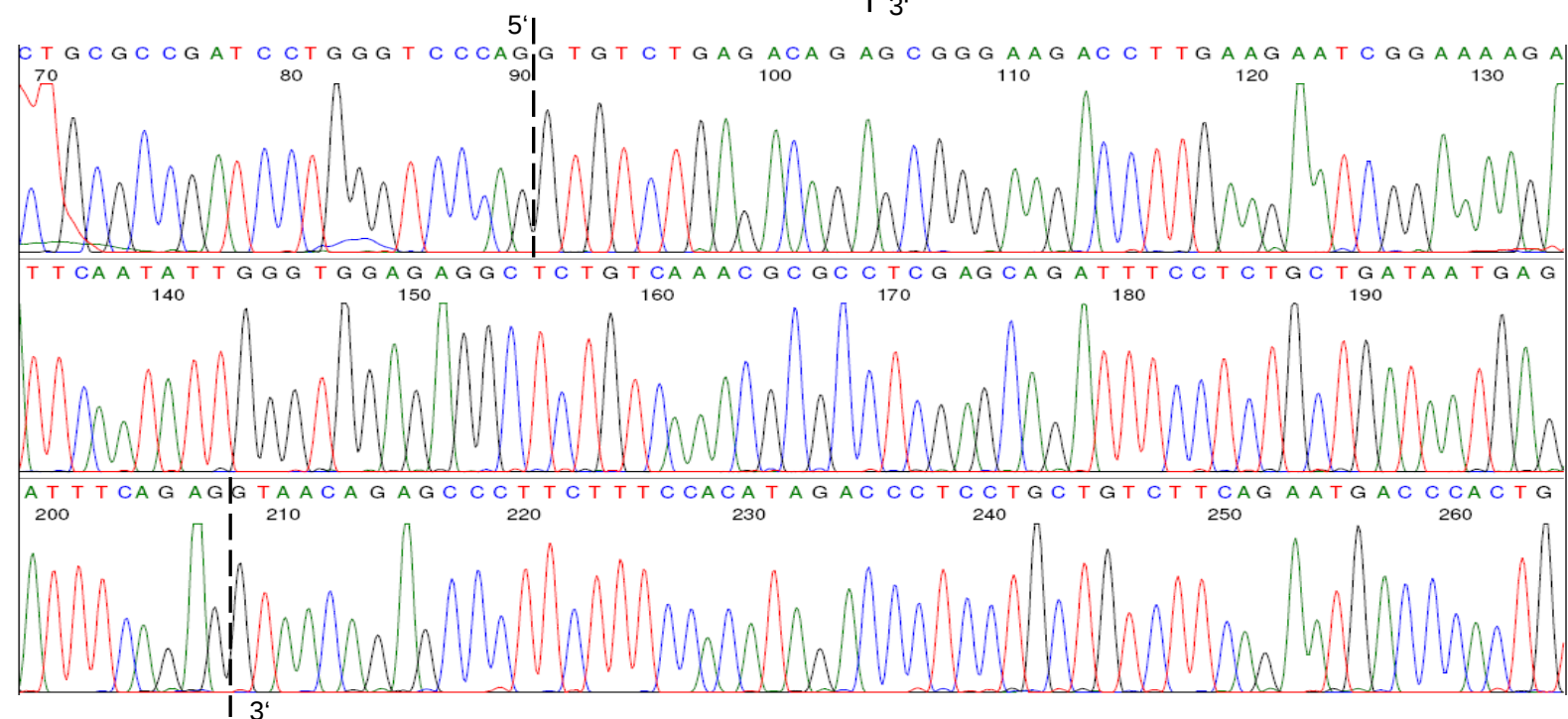
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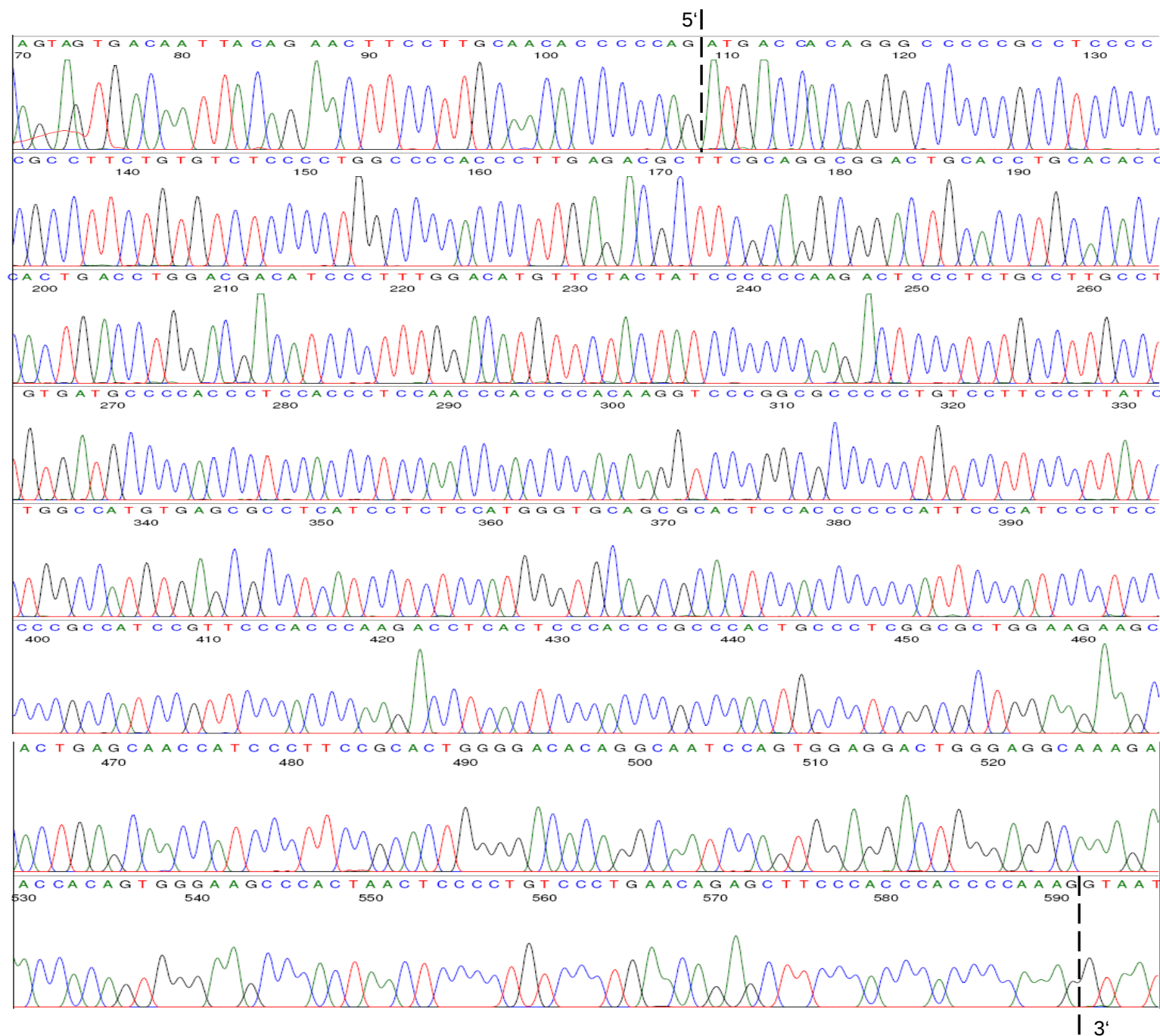
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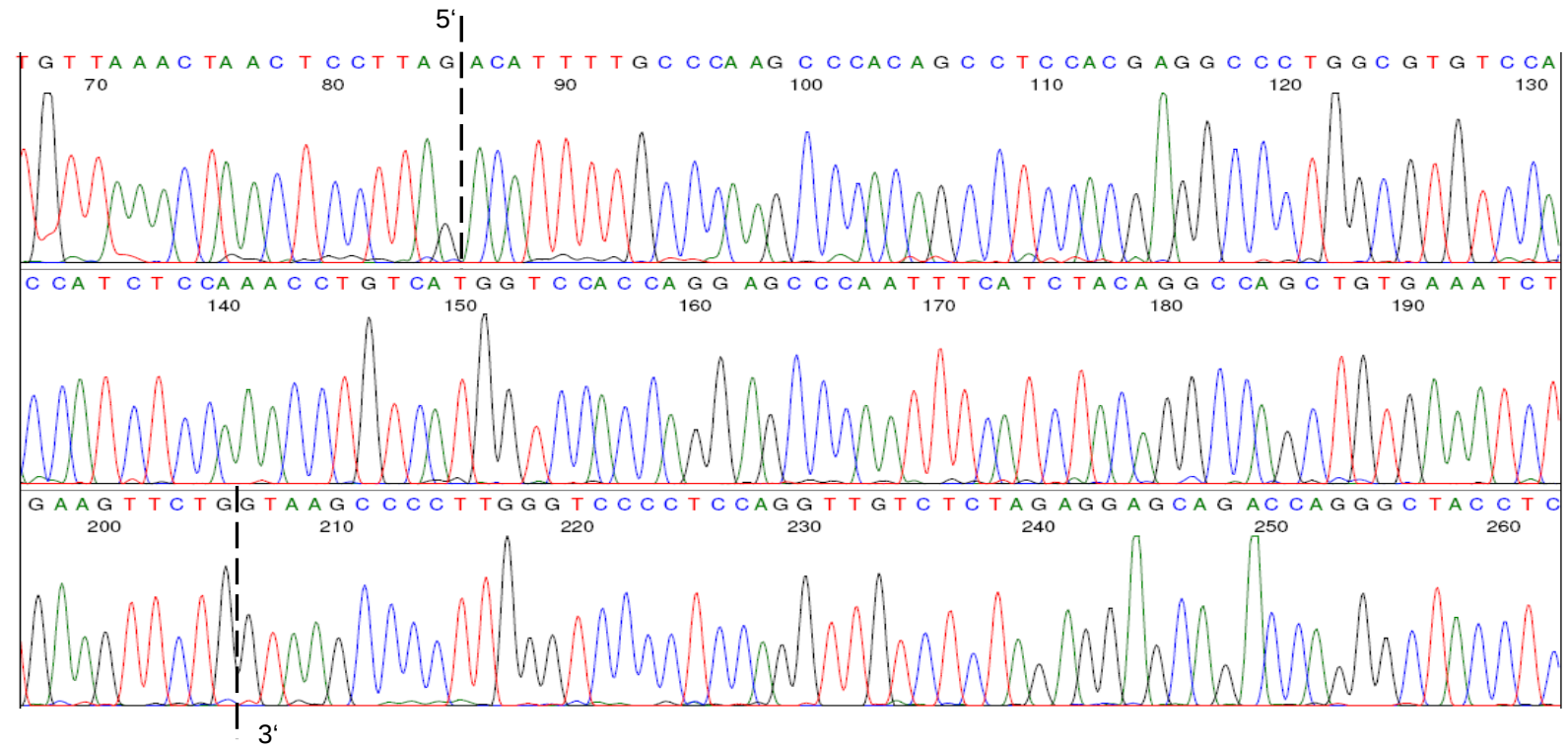
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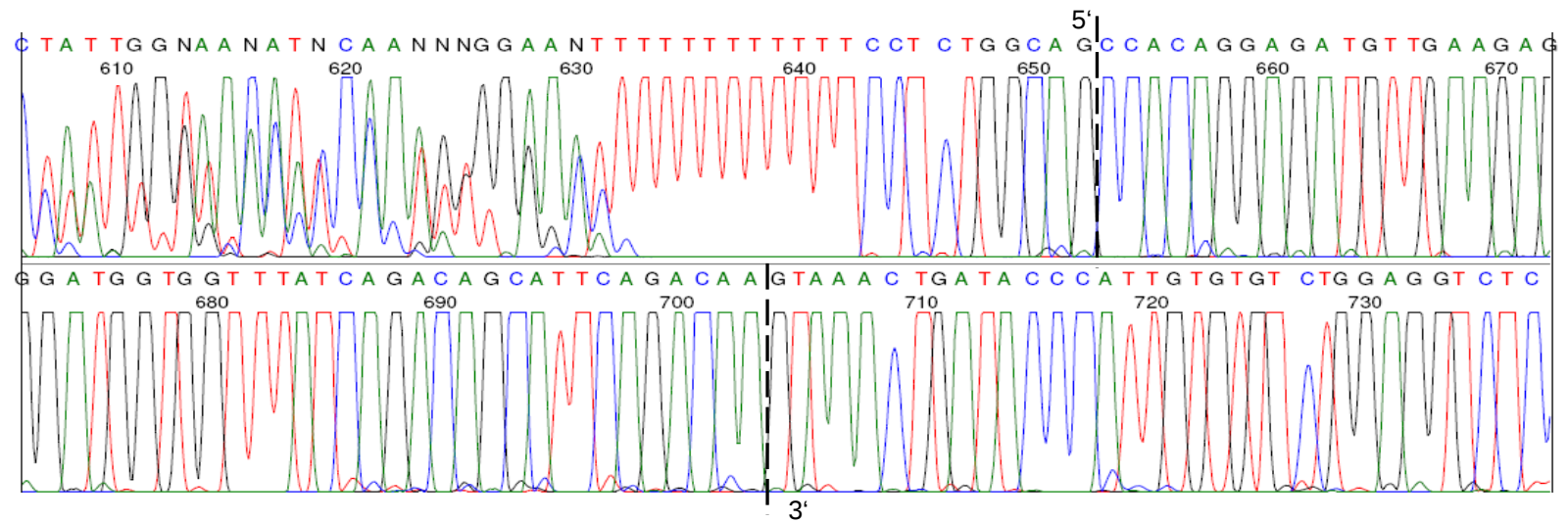
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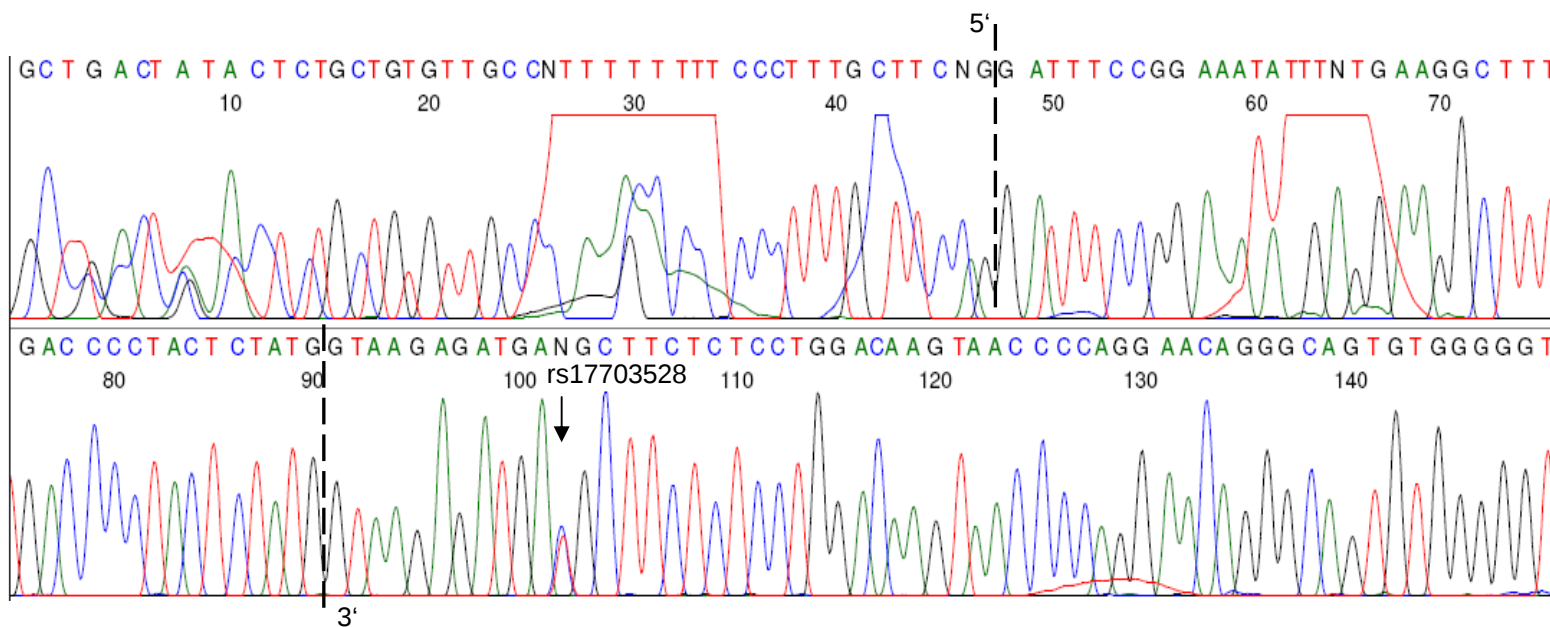
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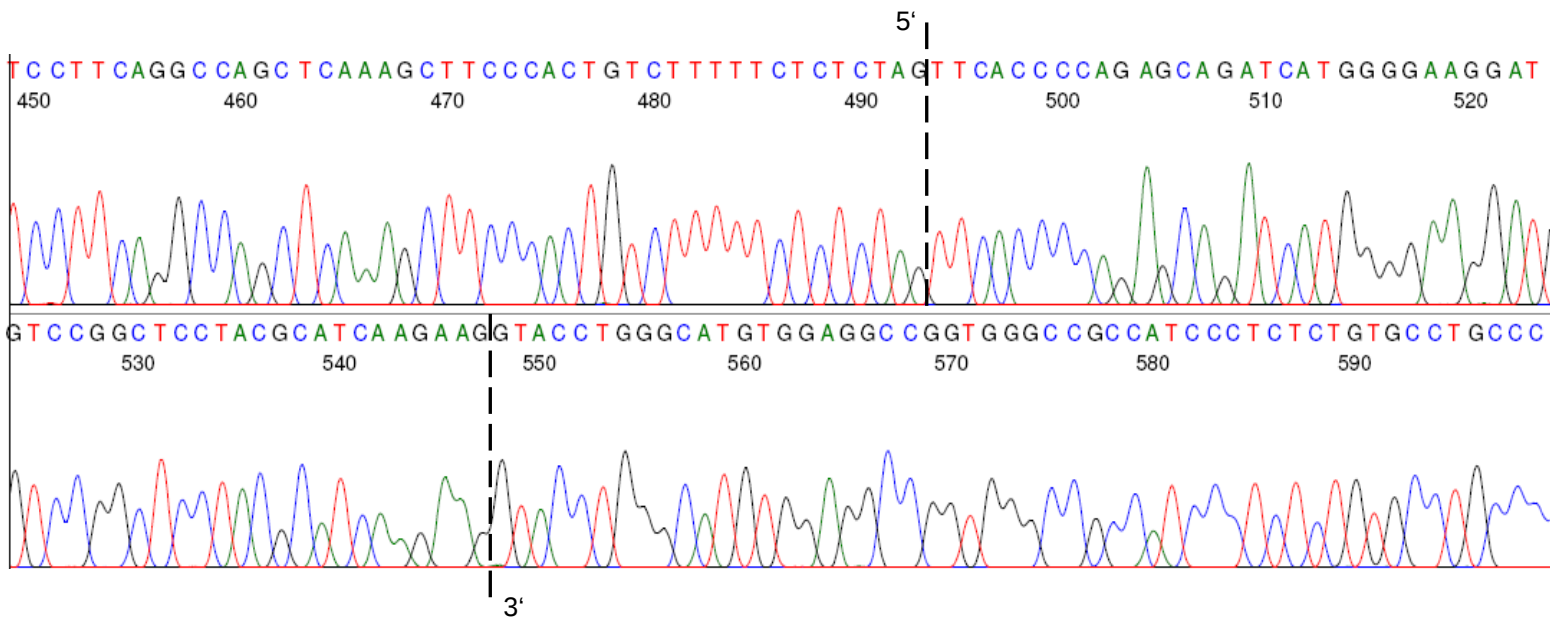
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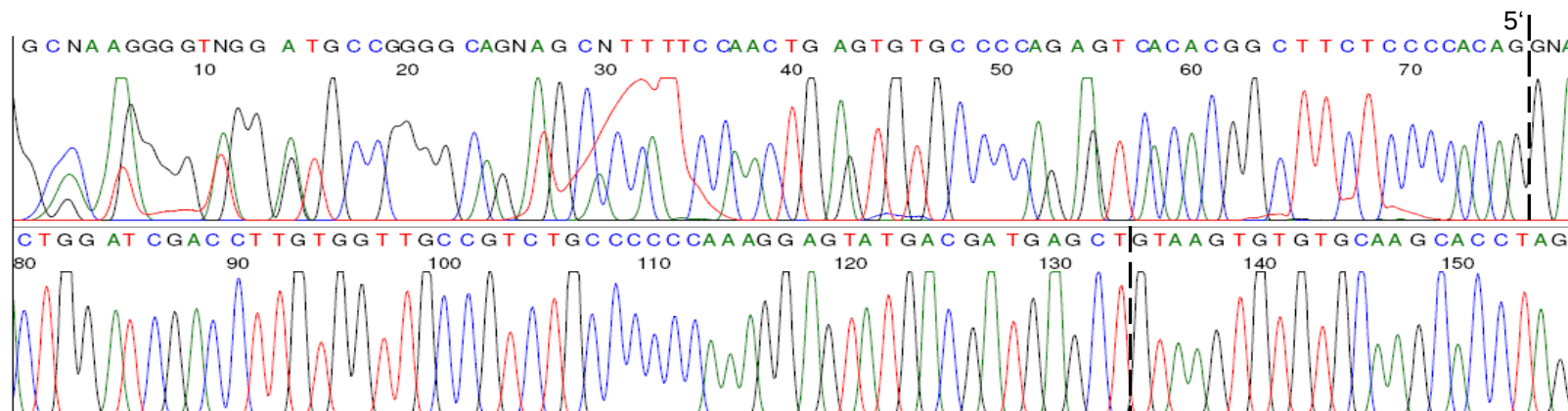
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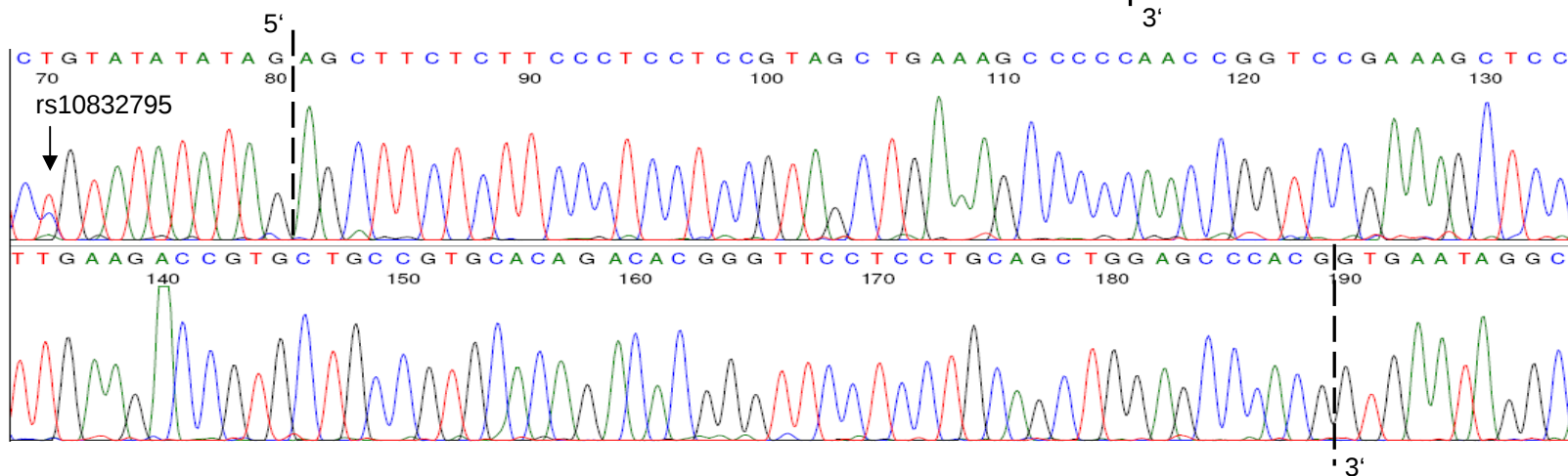
Exon 17



Exon 20



Exon G



Exon 21

