

K220820 Parky AppNov 17, 2022
241 days to decisionK220820 · Product code: **GYD** · Neurology
Source: <https://www.510kdatabase.net/k220820/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Tremor (GYD)
Date received	Mar 21, 2022
Decision date	Nov 17, 2022
Days to decision	241 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	H2o Bilisim Yazilim
Location	Cankaya, TR
Contact	Yagmur Selin Gulmus-Kolay
510(k) history	1 submissions · 1 cleared · 2022-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k220820/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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