

Case Study – Patient Advocate Stories



John Lancaster

Patient advocate, use MY data

I think that although it's the patient's data and in the end it is their decision and their choice, it is also their responsibility to make their data available for the benefit of others.

Data has to be used responsibly and it has to be kept safe.
But it has to be available for research.

We are building an online library, where the patient voice is heard direct, talking about the rewards of using their data



**<http://www.usemydata.org/stories5.shtml>
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“The only source of data is the patient.”