Data Sharing Statement		
Article Info	https://dx.doi.org/10.21037/apm-22-714	
Item	**Question**	**Authors' Response (place "-" if not applicable)**
1	Would you like to share data collected for your study to others?	Yes
2	If not, would you like to share the reason for your decision?	-
3	What data in particular will be shared?	The data on patient demographic, intraoperative characteristics, blood pressure, EuroSCORE II, Vasoactive Inotropic Score (VIS) will be shared.
4	Any other documents will be shared? Such as study protocol, statistical analysis plan, informed consent form, clinical study report, analytic code.	Statistical analysis plan, informed consent form, and clinical study report can also be shared if requested.
5	When will data availability begin?	From the publication date
6	When will data availability end?	Two years within the publication date, since risk score and protamine dose may be updated over time.
7	To whom will you share the data?	Anesthesiologists, surgeons and physicians who are interested in the studies related to protamine side effects.
8	For what type of analysis or purpose?	For systematic review or meta-analysis on pretreatment medication for prevention of protamine side effects.
9	How or where can the data/documents be obtained?	Emails could be sent to the address below to obtain the shared data: nophanan.chi@mahidol.edu
10	Any other restrictions?	We may balance the potential benefits and risks for each request and then provide the data that could be shared.