

Medical ultrasound imaging

Modern ultrasound imaging

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based on lectures 2008-2023 by J.Kybic,
using plots by J.Hornak¹, E.Dove, P.Stoyen

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¹<http://www.cis.rit.edu/htbooks/mri/>

Doppler ultrasound

US contrast agents

Harmonic imaging

3D US imaging

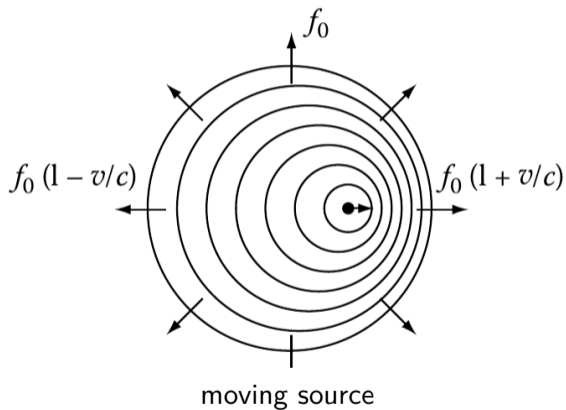
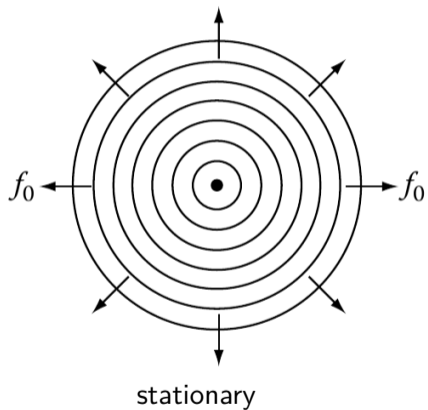
Christian Doppler

1803–1853

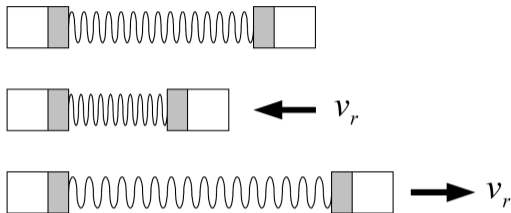


During his time in Prague as a professor he married and had five children, whilst he published over 50 articles on mathematics, physics and astronomy. In 1842 he published his most famous paper *Über das farbige Licht der Doppelsterne* (“Concerning the Colored Light of Double Stars”), which contained his first statement of the Doppler effect.

Doppler frequency shift



Stationary source, moving receiver

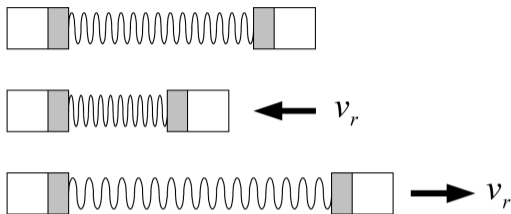


$$f_r = f_s + \frac{v_r}{\lambda_s} = f_s + \frac{v_r}{c} f_s = f_s + f_d, \quad \text{since } \lambda_s = \frac{c}{f_s}$$

Doppler shift

$$f_d = \frac{v_r}{c} f_s$$

Stationary source, moving receiver



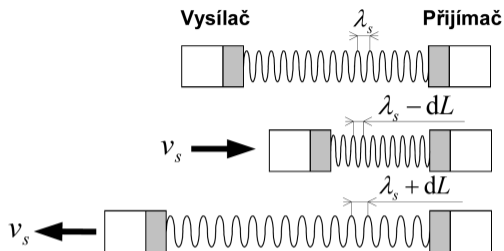
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Doppler shift

$$f_d = \frac{v_r}{c} f_s$$

Example: For $f = 5 \text{ MHz}$, $v_r = 1 \text{ cm/s}$, $f_d = 33 \text{ Hz}$.

Stationary receiver, moving source



Wavelength change

$$\delta\lambda = v_s T_s = \frac{v_s}{f_s}$$

$$\lambda_r = \lambda_s - \delta\lambda = \frac{c}{f_s} - \frac{v_s}{f_s}$$

$$f_r = \frac{c}{\lambda_r} = \frac{c}{c - v_s} f_s$$

Stationary receiver, moving source (2)

$$f_r = \frac{c}{\lambda_r} = \frac{c}{c - v_s} f_s = \frac{1}{1 - \frac{v_s}{c}} f_s$$

Stationary receiver, moving source (2)

$$f_r = \frac{c}{\lambda_r} = \frac{c}{c - v_s} f_s = \frac{1}{1 - \frac{v_s}{c}} f_s$$

From Taylor series, for $x \ll 1$

$$\frac{1}{1 - x} = 1 + x + \frac{x^2}{2} + \dots \approx 1 + x$$

For $v \ll c$

$$f_r \approx \left(1 + \frac{v_s}{c}\right) f_s = f_s + f_d$$

Doppler shift

$$f_d = \frac{v_s}{c} f_s$$

Blood flow speed measurement

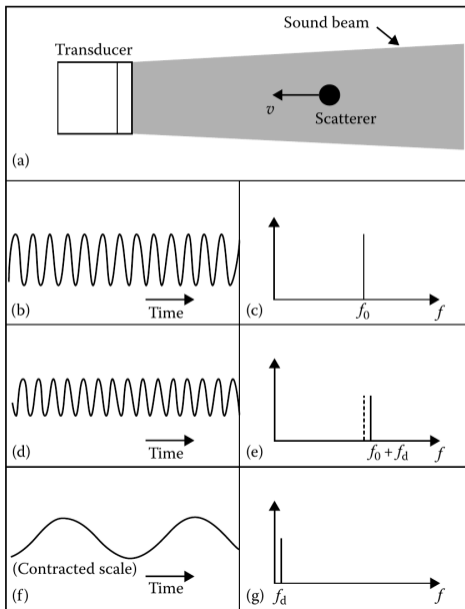
- ▶ Doppler effect: Frequency changes if the source moves with respect to the receiver.
- ▶ Reflection from red blood cells
- ▶ Red blood cells
 - ▶ Moving receiver
 - ▶ Moving source
- ▶ Doppler shift

$$f_r = f_t + f_d$$

$$f_d \approx 2 \frac{v}{c} f_c$$

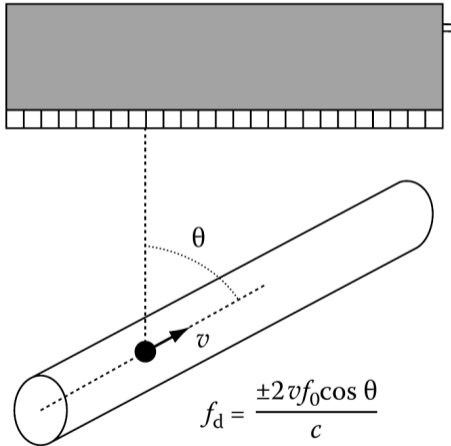
Moving scatterer

time and frequency domains



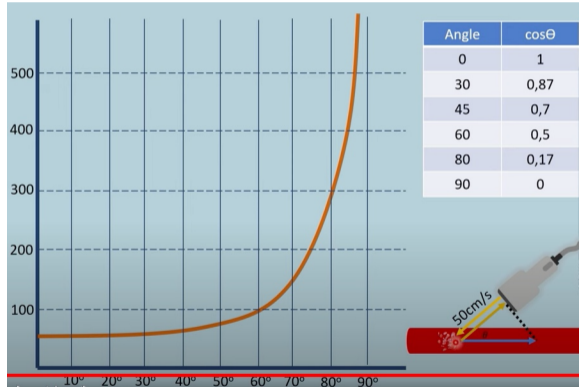
Angle dependency

We only measure the projection along the ray: $v \cos \theta$



$$f_d = \frac{\pm 2vf_0 \cos \theta}{c}$$

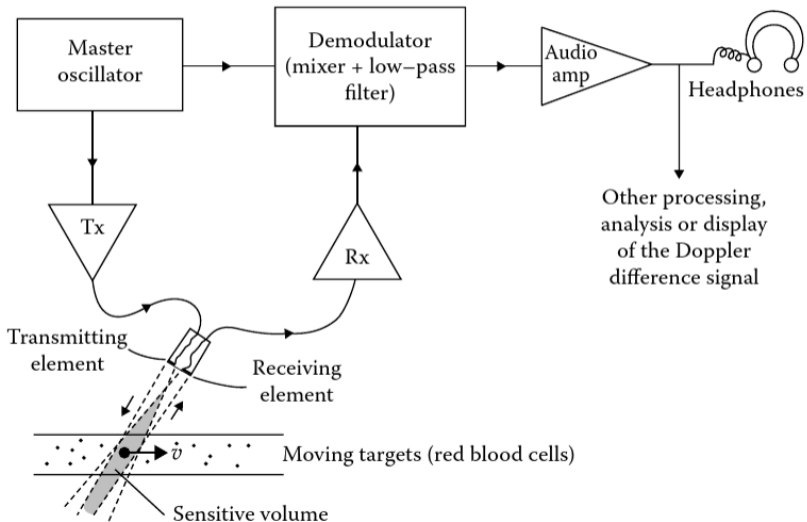
$$f_0 = f_s$$



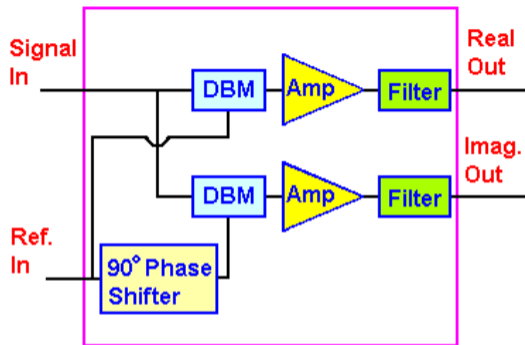
Angle dependency (2)

- ▶ Insonation angle 90° \rightarrow weak or no signal.
- ▶ Known angle \rightarrow angle correction.

Continuous wave Doppler



Quadrature detector



- ▶ *Input:* $g_a = \cos(at)$, $g_b = \cos(bt)$
- ▶ *Output:* $g = g_a g_b = \frac{1}{2} \cos((a+b)t) + \frac{1}{2} \cos((a-b)t)$
- ▶ Signal $\cos((a+b)t)$ can be filtered (low-pass filter)
- ▶ Difference frequency signal $s_r = \cos((a-b)t)$
- ▶ “Imaginary” signal s_i shifted by 90° : $\sin((a-b)t)$

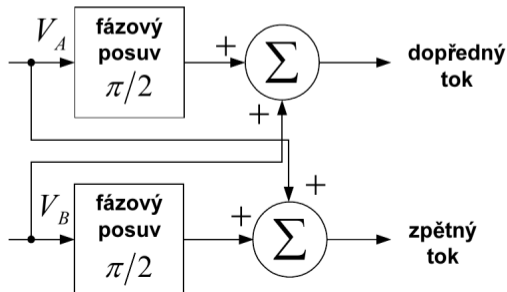
Directional demodulation

To distinguish positive/negative flow direction, $\pm f_d$.

Method 1: Phase-domain processing

- ▶ Quadrature mixer with f_s
- ▶ Phase offset $\angle s_r = \angle s_i = \pm 90^\circ$

	$f_d > 0$	$f_d < 0$
$s_r + T_{90}s_i$	0	$2s_r$
$T_{90}s_r + s_i$	$2s_r$	0



Directional demodulation

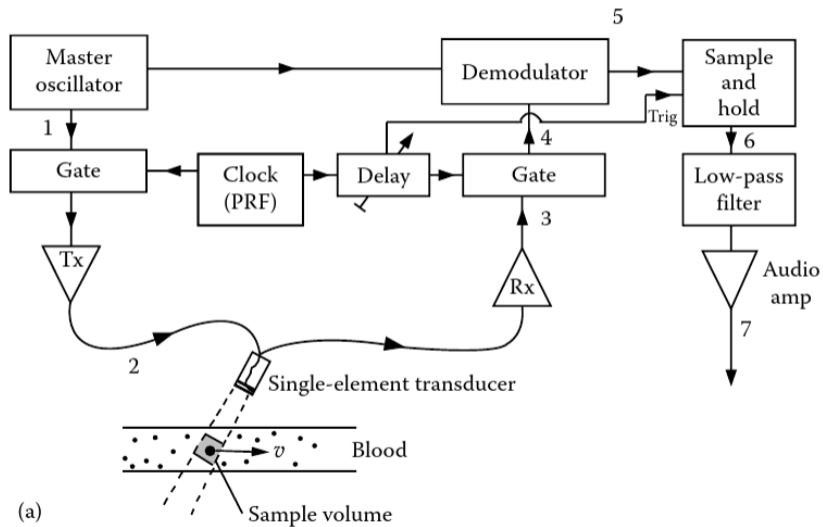
To distinguish positive/negative flow direction, $\pm f_d$.

Method 2: Frequency shift

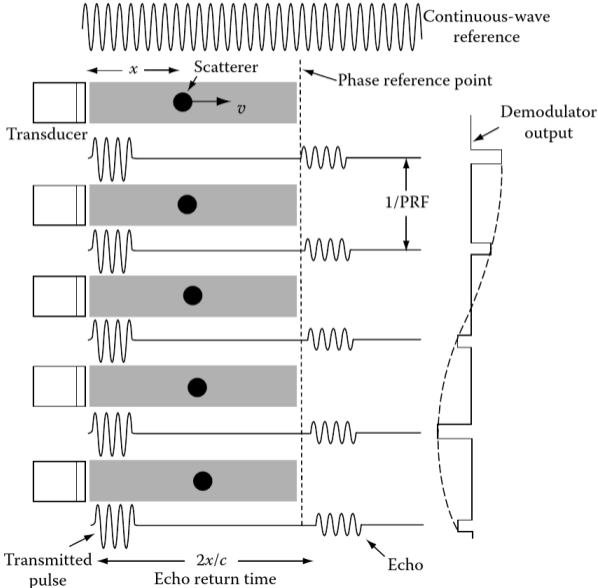
- ▶ Quadrature mixer with $f_s + f_o$
- ▶ $f_d = 0 \rightarrow$ mixer output f_o
- ▶ $f_d = \text{freq}(s_r) - f_o$

Pulsed wave Doppler

(PW)



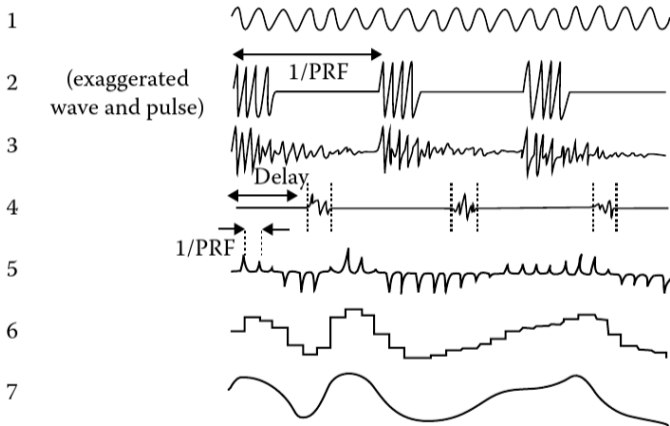
Sampled Doppler shift signal



PW Doppler shift signals

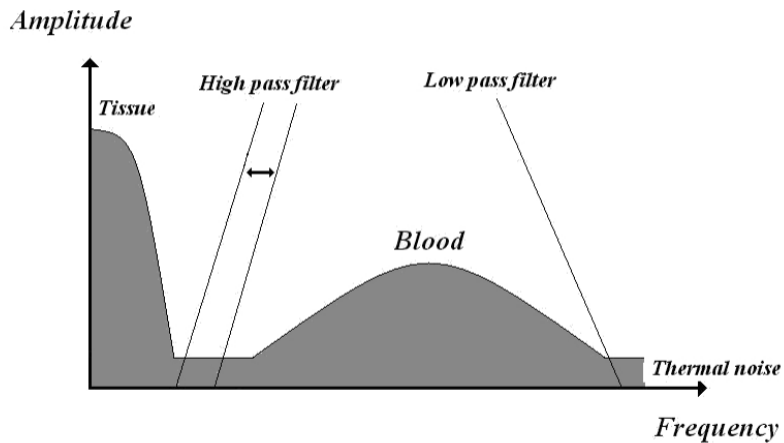
Point on diagram

Signal representation at this point

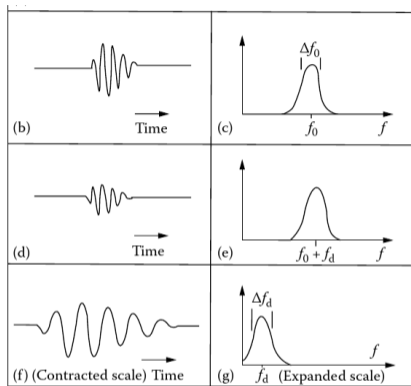


PRF: pulse repetition frequency, 2: transmitted signal, 3: received signal, 4: gated signal, 5: demodulated signal, 6: interpolated signal, 7: output

PW Doppler spectrum



Speed uncertainty

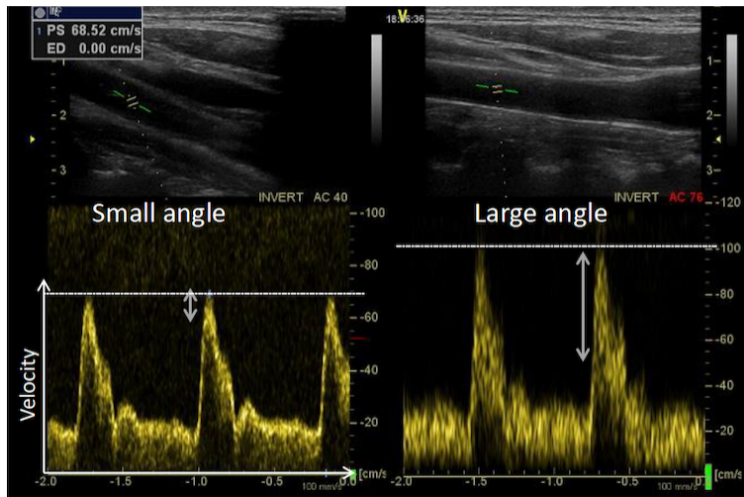


$$\Delta f_d = \frac{2v}{c\tau} = \frac{f_d}{\underbrace{f_s \tau}_Q}, \quad \Delta v = \frac{v}{Q}$$

τ — pulse length, Q — quality factor, number of cycles in a pulse

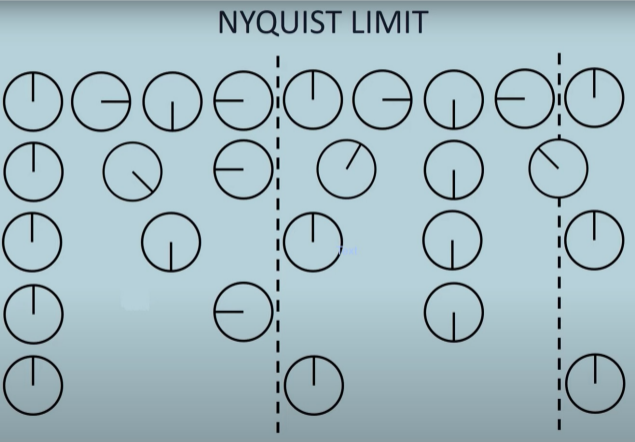
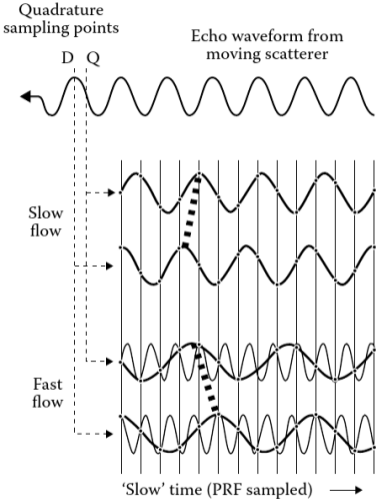
Speed uncertainty

Angle dependency



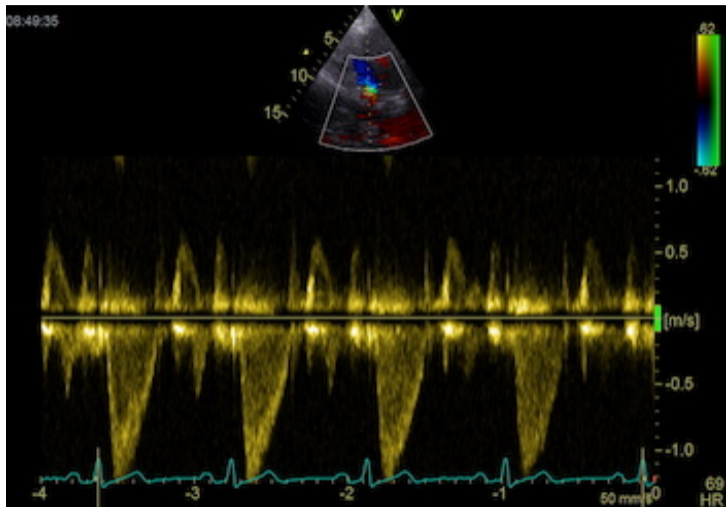
small angle \rightarrow higher number of cycles Q

Aliasing (PRF too low)



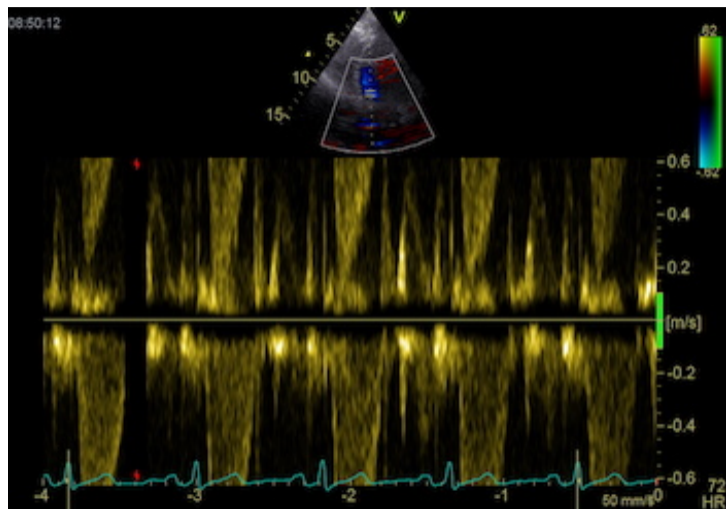
Nyquist → $f_d < f_p/2$

Aliasing example



B-mode+Doppler+velocity spectrum — high PRF f_p

Aliasing example



spectrum — low PRF f_p

Range-velocity tradeoff

$$f_d < f_p/2 \quad \longrightarrow \quad v_{\max} = \frac{f_p c}{4f_s}$$

$$z_{\max} = \frac{T_p c}{2} = \frac{c}{2f_p}$$

$$v_{\max} z_{\max} = \frac{c^2}{8f_s}$$

Range-velocity tradeoff

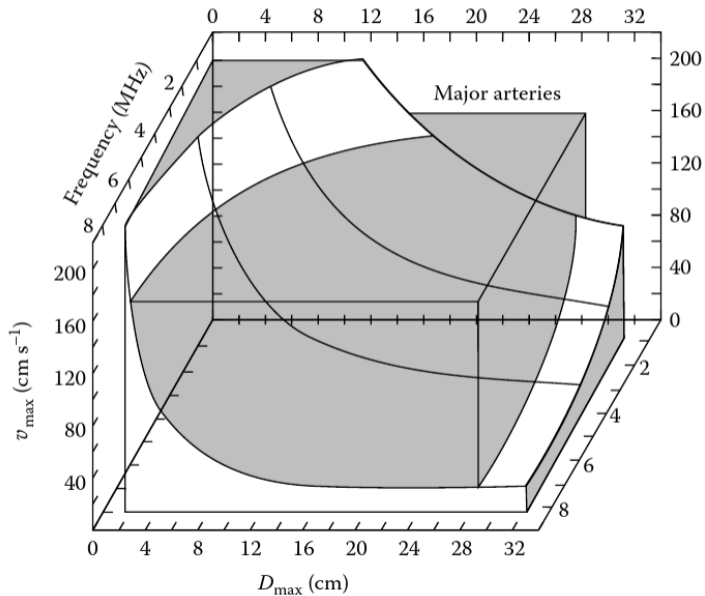
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Limitation is for $(v \cos \theta)_{\max}$.

Range-velocity tradeoff



Minimum velocity

Observe at least one period of f_d

$$T_d < NT_p$$

with N transmissions per line

$$f_d > \frac{f_p}{N}$$
$$v_{\min} = \frac{f_p c}{2Nf_s}$$

Minimum velocity

Observe at least one period of f_d

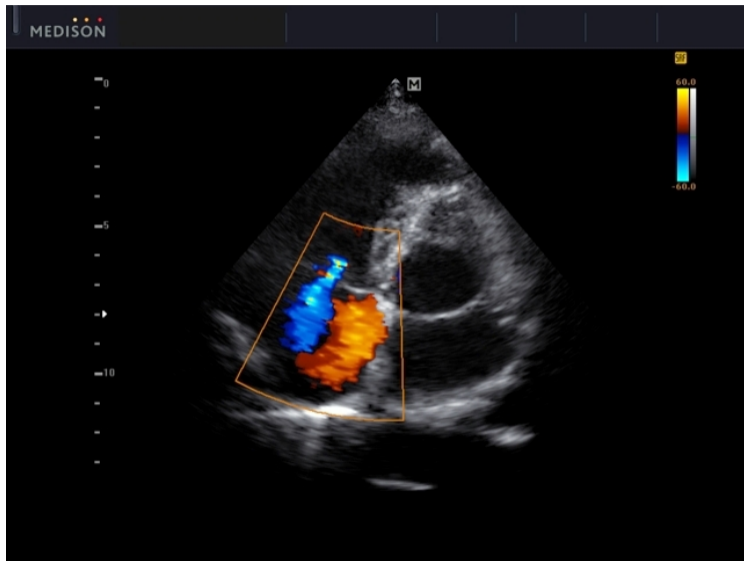
$$T_d < NT_p$$

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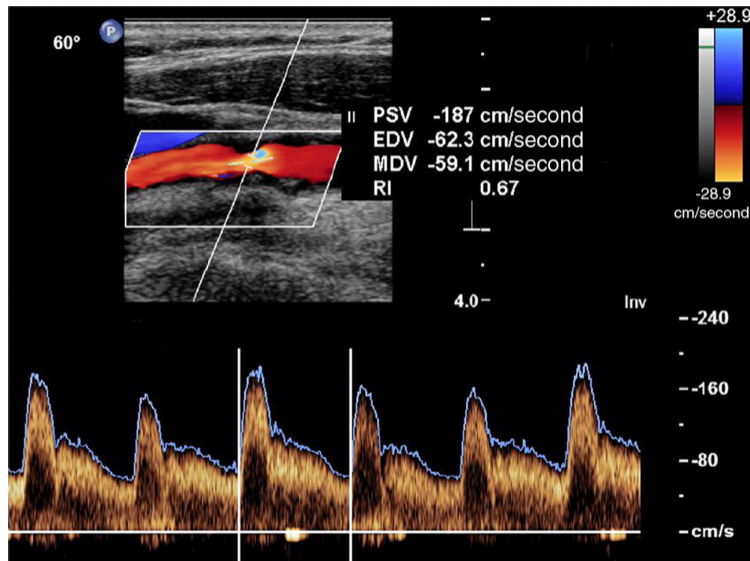
- ▶ Usually 2 ~ 3 cycles required
- ▶ $N = 5 \sim 10$ or more
- ▶ Temporal averaging
- ▶ \rightarrow slow f_p

Doppler US — examples



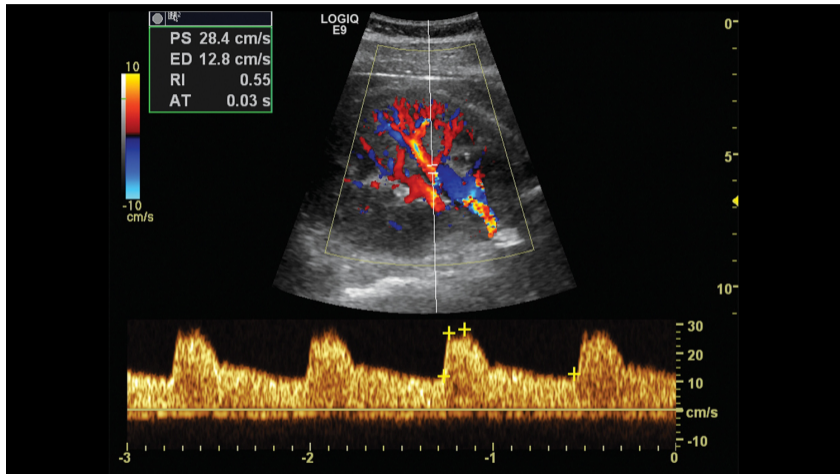
heart

Doppler US — examples



artery

Doppler US — examples



liver

Doppler ultrasound

US contrast agents

Harmonic imaging

3D US imaging

Contrast agents

- ▶ *1968, Gramiak, saline injection*
- ▶ Microbubbles ($2 \sim 5 \mu\text{m}$)
- ▶ Asymmetric compression/expansion
- ▶ Stabilization (synthetic polymers), up to 5 – 10 min.
- ▶ Injection
- ▶ Albunex, Optison, Echovist, Levovist. . .

Flash contrast imaging

US bubble destabilization.



normal

Flash contrast imaging

US bubble destabilization.



flash, bubbles broken

Flash contrast imaging

US bubble destabilization.



filling up

Myocardial perfusion evaluation.

Doppler ultrasound

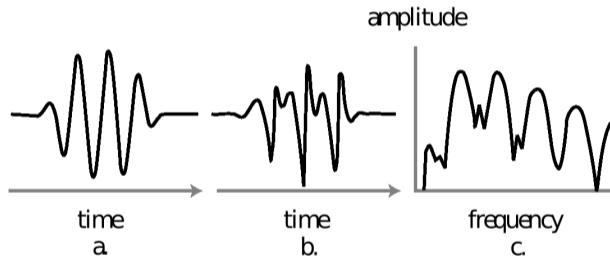
US contrast agents

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3D US imaging

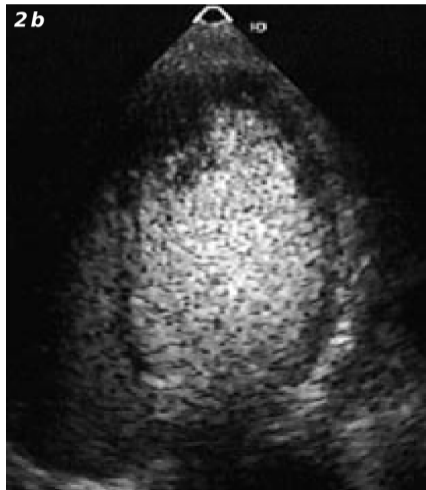
Nonlinear response

Assymmetric bubble compression



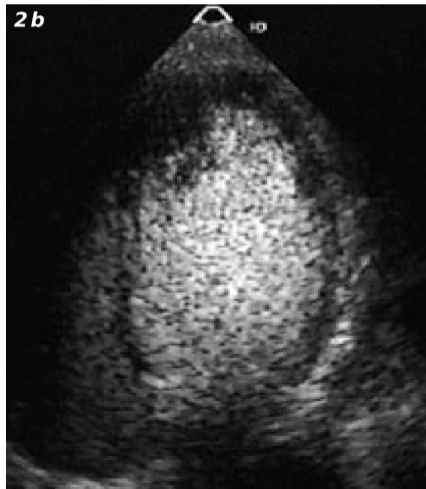
Harmonic imaging

- ▶ Transmit f_0 , receive $2f_0$



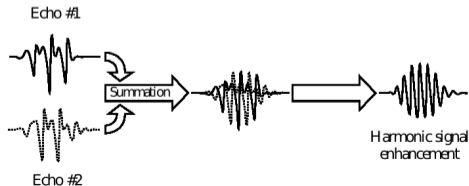
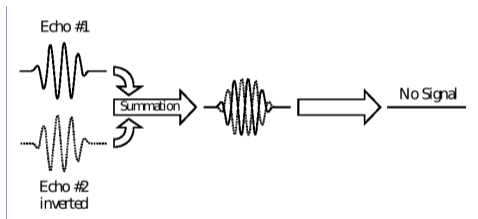
Harmonic imaging

- ▶ Transmit f_0 , receive $2f_0$
- ▶ Bandwidth limitation
- ▶ Bubbles not needed, tissue nonlinearity



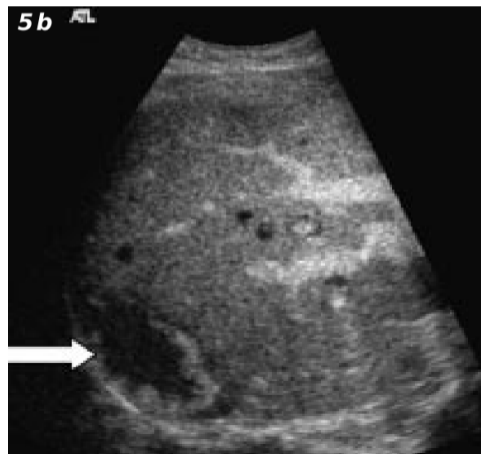
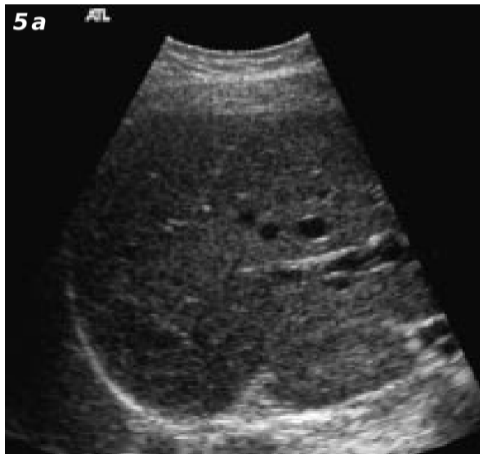
Pulse Inversion Harmonic Imaging

- ▶ Two pulses, second inverted
- ▶ Responses summed
- ▶ Filtration not needed



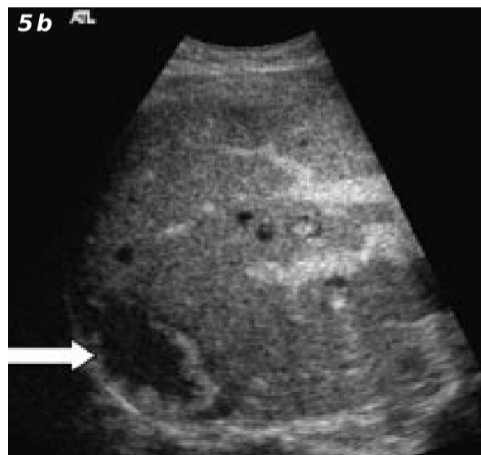
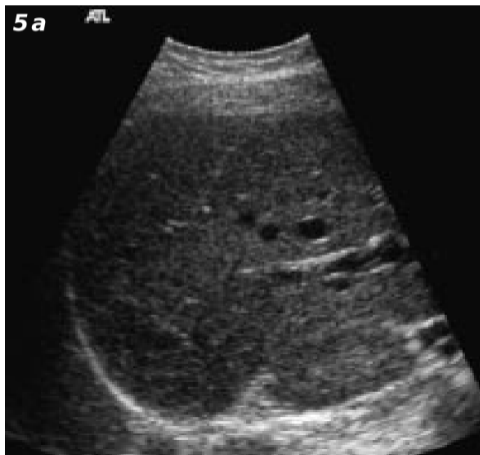
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Pulse Inversion Harmonic Imaging

- ▶ Two pulses, second inverted
- ▶ Responses summed
- ▶ Filtration not needed
- ▶ Several pulses (Power Pulse Inversion)



Doppler ultrasound

US contrast agents

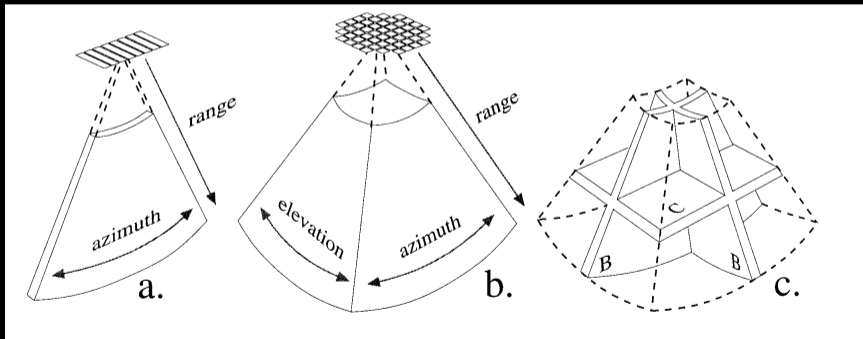
Harmonic imaging

3D US imaging

3D Reconstruction



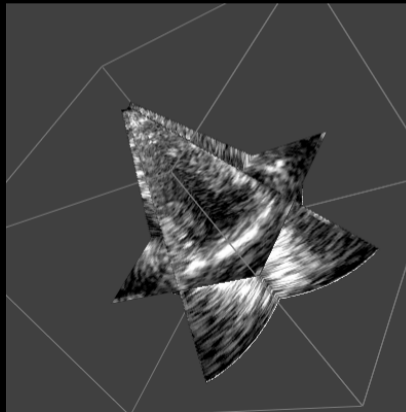
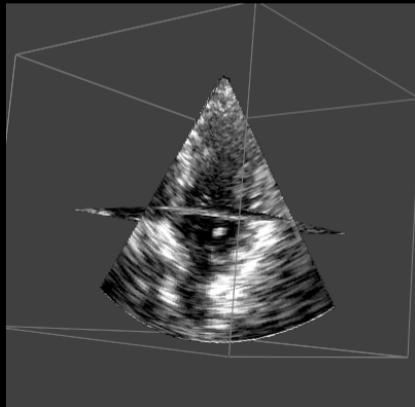
3D Ultrasound

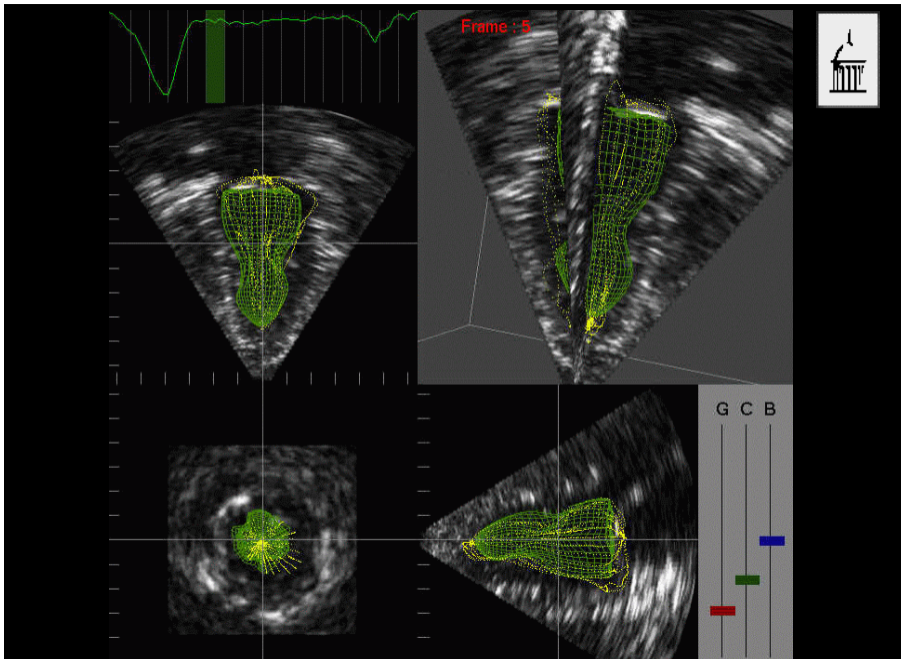


Traditional 2D

New 3D

Real-time 3D Ultrasound

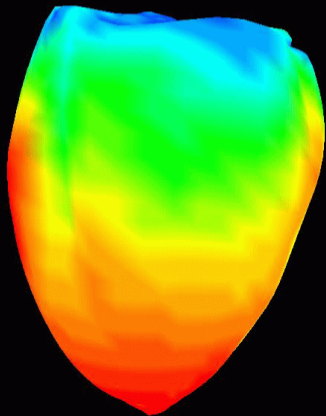




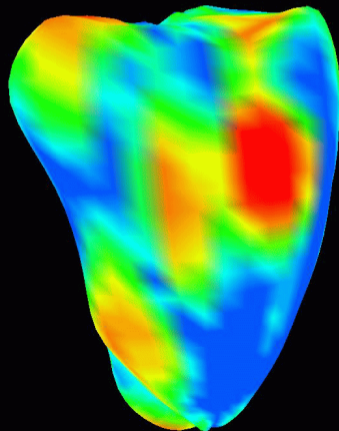
Frame : 5

G C B

Velocity of Contraction



N 1



A 1

Biological effects

- ▶ Thermal effects
 - ▶ 1.5 °C indefinitely or 6 °C for 1 min
 - ▶ highest risk in bones (transcranial imaging)
- ▶ Cavitation — growth/collapse of bubbles
 - ▶ for long pulse lengths or high pressure
 - ▶ may damage cells
 - ▶ unlikely to occur *in vivo*
- ▶ Radiation pressure — makes tissues/fluids move

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 - ▶ unlikely to occur *in vivo*
- ▶ Radiation pressure — makes tissues/fluids move
- ▶ Clinical studies found no harmful effects
- ▶ ... ultrasound power output is increasing.

Conclusions

- ▶ Non-invasive, affordable and portable imaging technique
- ▶ Excellent soft tissue imaging
- ▶ Lower image quality (wrt CT or MRI) due to speckle but improving
- ▶ Low penetration depth versus resolution
- ▶ Does not pass through air or gas
- ▶ Does not pass through bones, shadows
- ▶ Modern techniques — 3D, contrast agents, Doppler